



सहायक महाप्रबंधक Assistant General Manager र.मं.प.-निर्गम एवं सूचीबद्धता प्रभाग-2 / RAC-Division of Issues and Listing2 निगम वित्त विभाग / Corporation Finance Department

> SEBI/HO/CFD/RAC-DIL2/P/OW/2023/1400/1 January 11, 2023

Mr. Sameer Purohit
ICICI Securities Limited
ICICI Venture House
Appasaheb Marathe Marg, Prabhadevi
Mumbai - 400025

महोदय / महोदया, Madam / Sir,

### विषय / Sub: Proposed IPO of Innova Captab Limited

उपरोक्त से संबंधित प्रारूप प्रस्ताव दस्तावेज (ड्राफ्ट ऑफर डॉक्यूमेंट), भारतीय प्रतिभृति और विनिमय बोर्ड (सेबी) दवारा मांगे गए स्पष्टीकरणों और उसके संबंध में दिए गए उत्तरों के संदर्भ में, यह सूचित किया जाता है कि इनकी जाँच करने पर यह पाया गया है कि इनमें किमयाँ हैं / भारतीय प्रतिभृति और विनिमय बोर्ड [पूँजी का निर्गमन (इश्यू) और प्रकटीकरण अपेक्षाएँ] विनियम, 2018 [सेबी (इश्यू ऑफ कैपिटल एंड डिस्क्लोज़र रिक्वायरमेंटस) रेग्यूलेशन्स, 2018] के प्रावधानों और दिए गए अनुदेशों का पालन नहीं किया गया है, और आपके लिए यह जरूरी है कि आप स्टॉक एक्सचेंज और / या कंपनी रिजस्ट्रार के पास प्रस्ताव दस्तावेज दाखिल करने से पहले उन किमयों को दूर करें और संबंधित प्रावधानों तथा दिए गए अनुदेशों का पालन करें । उपरोक्त के संबंध में की गई टिप्पणियों का और जिन शर्तों आदि का पालन किया जाना है, उनका जिक्र संलग्नक 'I' और संलग्नक 'II' में किया गया है । कृपया यह भी नोट करें कि संलग्नक में जो किमयाँ बताई गई हैं / कुछ और प्रकटीकरण (डिस्क्लोज़र) करने की बात कही गई है, यह सब आपको केवल उदाहरण के तौर पर ही बताया गया है । यह सुनिश्चित करने की जिम्मेदारी आपकी है कि सभी और सही प्रकटीकरण किए जाएं।

With reference to the draft offer document in respect of captioned issue, clarifications sought by SEBI and the replies submitted therein, it is stated that on scrutiny of the same, deficiencies / instances of non-compliance of SEBI (Issue of Capital and Disclosure Requirements) Regulations, 2018 (hereinafter referred to as SEBI (ICDR) Regulations, 2018) and instructions have been observed, which are required to be rectified / complied with by you before filing the



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सेबी भवन, प्लॉट सं. सी 4-ए, "जी" ब्लॉक, बांद्रा-कुर्ला कॉम्प्लेक्स, बांद्रा (पूर्व), मुंबई - 400 051.

दूरभाष: 2644 9950 / 4045 9950 (आई.वी.आर.एस.), 2644 9000 / 4045 9000 फैक्स: 2644 9019 से 2644 9022 वेब: www.sebi.gov.in



offer document with the Stock Exchange and/ or ROC. Observations on the captioned issue and other conditions to be complied with are indicated in Annexure 'l' and 'll'. It may be noted that the deficiencies / requirement of additional disclosures listed in the Annexure are merely illustrative and not exhaustive. It is your responsibility to ensure full and true disclosures.

1. बुक रिनंग लीड मैनेजर होने के नाते, आप यह सुनिश्चित करेंगे कि स्टॉक एक्सचेंज / कंपनी रिजिस्ट्रार के पास प्रस्ताव दस्तावेज दाखिल करने से पहले संलग्नक में दी हुई टिप्पणियों / शर्तों आदि के अनुसार प्रस्ताव दस्तावेज में बदलाव कर लिए जाएं । कंपनी रिजिस्ट्रार / स्टॉक एक्सचेंज के पास अंतिम प्रस्ताव दस्तावेज दाखिल करने से पहले आपको हमें एक पत्र भेजकर इस बात की पुष्टि करनी होगी कि अपेक्षानुसार बदलाव कर लिए गए हैं और साथ ही यह भी बताना होगा कि प्रत्येक टिप्पणी / शर्त आदि के अनुसार बदलाव कैसे किए गए हैं । इसके अलावा और कोई भी बदलाव सेबी से लिखित सहमित लिए बिना नहीं किए जाएंगे ।

As Book Running Lead Manager (LM), you shall ensure that all changes are effected based on the observations / conditions contained in the Annexure before you file the offer document with the stock Exchange / ROC. A letter confirming these changes and explaining, in seriatim, the manner in which each observation / condition has been dealt with along with your comments should be submitted to us, before filing the final offer document with ROC / Stock Exchange. NO FURTHER CHANGES SHOULD BE EFFECTED WITHOUT SPECIFIC WRITTEN CONSENT OF SEBI.

2. यह स्पष्ट किया जाता है कि भारतीय प्रतिभूति और विनिमय बोर्ड (सेबी) के पास प्रस्ताव दस्तावेज (ऑफर डॉक्यूमेंट) दाखिल करने का अर्थ किसी भी तरह से यह न लगाया जाए कि सेबी दवारा इसे मंजूरी प्रदान कर दी गई है । सेबी न तो इस बात की कोई जिम्मेदारी लेता है कि जिस स्कीम या परियोजना (प्रोजेक्ट) के लिए निर्गम (इश्यू) लाए जाने का प्रस्ताव है उसकी वित्तीय स्थिति अच्छी है और न ही इस बात की जिम्मेदारी लेता है कि प्रस्ताव दस्तावेज में दी गई जानकारी या व्यक्त की गई राय सही है । अग्रणी प्रबंधकों (लीड मैनेजर्स) ने यह प्रमाणित किया है कि प्रस्ताव दस्तावेज में जो प्रकटीकरण (डिस्क्लोज़र) किए गए हैं वे मोटे तौर पर पर्याप्त हैं और जो प्रकटीकरणों (डिस्क्लोज़र) तथा निवेशक संरक्षण के संबंध में उस समय लागू सेबी के विनियमों के प्रावधानों के अनुसार किए गए हैं । अग्रणी प्रबंधक यह भी सुनिश्चित करेंगे कि ऐसा भारतीय प्रतिभूति और विनिमय बोर्ड [पँजी का निर्गमन (इश्यू) और प्रकटीकरण अपेक्षाएँ] विनियम, 2018 [सेबी (इश्यू ऑफ कैपिटल एंड डिस्क्लोज़र रिक्वायरमेंटस) रेग्यूलेशन्स, 2018] के अनुसार भी किया जाए । ऐसा करना इसलिए जरूरी है, तािक निवेशक प्रस्तािवत निर्गम (इश्यू) में निवेश करने के संबंध में सोच-समझकर निर्णय ले सकें।

It is to be distinctly understood that submission of offer document to SEBI should not in any way be deemed or construed that the same has been cleared or approved by SEBI. SEBI does not take any responsibility either for the financial soundness of any scheme or the project for which the issue is proposed to be made or for the correctness of the statements made or opinions expressed in the offer document. The LMs have certified that



the disclosures made in the offer document are generally adequate and are in conformity with SEBI regulations for disclosures and investor protection in force for the time being. The LMs are advised to ensure the same with respect to SEBI (ICDR) Regulations, 2018. This requirement is to facilitate investors to take an informed decision for making investment in the proposed issue.

3. यह भी पूरी तरह से स्पष्ट किया जाता है कि यदयपि इस बात की जिम्मेदारी मुख्य रूप से निर्गमकर्ता (इश्युअर) कंपनी की होती है कि प्रस्ताव दस्तावेज में समस्त जरूरी जानकारी प्रकट की जाए और जो सही और पर्याप्त हो, फिर भी अग्रणी प्रबंधकों (लीड मैनेजर्स) से अपेक्षित है कि वे यह सुनिश्चित करने के लिए पूरी तत्परता (इयू डिलिजेंस) बरतें कि कंपनी अपनी जिम्मेदारियाँ सही ढंग से निभाए, और इसी उद्देश्य से अग्रणी प्रबंधकों ने भारतीय प्रतिभूति और विनिमय बोर्ड (इश्यू ऑफ कैपिटल एंड डिस्क्लोज़र रिक्वायरमेंटस) रेग्यूलेशन्स, 2018 के अनुसार सेबी के पास पूरी तत्परता बरते जाने के संबंध में तारीख February 15, 2022 का प्रमाणपत्र (इयू डिलिजेंस सर्टिफिकेट) प्रस्तुत किया है।

It should also be clearly understood that while the Issuer Company is primarily responsible for the correctness, adequacy and disclosure of all relevant information in the offer document, the LMs are expected to exercise Due Diligence to ensure that the Company discharges its responsibility adequately in this behalf and towards this purpose, the LMs have furnished to SEBI a Due Diligence Certificate June 28, 2022 in accordance with SEBI (ICDR) Regulations, 2018.

4. हालाँकि, कंपनी प्रस्ताव दस्तावेज दाखिल कर देने से ही कंपनी अधिनियम, 2013 की धारा 34 के तहत दी गई किसी भी बाध्यता से मुक्त नहीं हो जाती या वह कानूनी प्रावधानों के अनुसार ली जाने वाली मंजूरी या ऐसी कोई अन्य मंजूरी लेने से मुक्त नहीं हो जाती, जो प्रस्तावित निर्गम के संबंध में लेनी जरूरी हो । हालाँकि, सेबी प्रस्ताव दस्तावेज में कोई अनियमितता या कमी पाए जाने पर कभी भी अग्रणी प्रबंधकों के खिलाफ कार्रवाई कर सकता है ।

The filing of offer document does not, however, absolve the company from any liabilities under Section 34 of the Companies Act, 2013 or from the requirement of obtaining such statutory or other clearances as may be required for the purpose of the proposed issue. SEBI further reserves the right to take up, at any point of time, with the LMs any irregularities or lapses in offer document.

5. किसी भी प्रचार सामग्री या विज्ञापन में ऐसा कुछ भी उल्लेख नहीं किया जाएगा, जो प्रारूप प्रस्ताव दस्तावेज (ड्राफ्ट ऑफर डाक्यूमेंट) में दी गई जानकारी से भिन्न हो । इस संबंध में आपका ध्यान विशेष रूप से कंपनी अधिनियम, 2013 की धारा 36 के प्रावधानों की ओर आकर्षित किया जाता है ।

Any publicity materials / advertisements should not contain matters extraneous to the information contained in the draft offer document. Attention is specifically drawn to the provisions of Section 36 of the Companies Act, 2013.





6. अग्रणी प्रबंधक यह सुनिश्चित करें कि भारतीय प्रतिभूति और विनिमय बोर्ड [पूँजी का निर्गमन (इश्यू) और प्रकटीकरण अपेक्षाएँ] विनियम, 2018 के विनियम 25(1) और अनुसूची- III के अनुसार उपरोक्त निर्गम (इश्यू) के संबंध में फाइलिंग फीस की गणना किस प्रकार की गई है उसका एक विस्तृत विवरण, यथास्थिति, कंपनी रिजिस्ट्रार के यहाँ प्रॉस्पेक्टस दाखिल किए जाने के सात दिनों के भीतर / स्टॉक एक्सचेंज के पास प्रस्ताव-पत्र (लेटर ऑफ ऑफर) दाखिल किए जाने के सात दिनों के भीतर, सेबी के पास प्रस्त्तत कर दिया जाए और साथ ही अब तक अदा की गई फाइलिंग फीस का ब्यौरा भी दिया जाए।

The LMs are advised to ensure that a detailed calculation of filing fees in relation to the captioned issue in terms of regulation 25(1) and Schedule III of the SEBI (ICDR) Regulations, 2018 is submitted to SEBI within seven days of filing the Prospectus with ROC/ within seven days of filing the Letter of Offer with the stock exchange, as the case may be, along with details of filing fees paid till date.

आपने जो फीस अदा की है, यदि वह वास्तव में अदा की जाने वाली फीस से कम हो, तो ऐसे में अग्रणी प्रबंधक यह सुनिश्चित करेंगे और इस बात की पुष्टि करेंगे कि सेबी को शेष फीस अदा किए जाने के संबंध में इन विनियमों की अन्सूची-III के प्रावधानों का पालन किया गया है।

If filing fees paid by you is less than the actual fees required to be paid, the LMs are advised to ensure and confirm compliance with the provisions of Schedule III of the said Regulations in regard to payment of the balance fees to SEBI.

आपने जो फीस अदा की है, यदि वह वास्तव में अदा की जाने वाली फीस से अधिक हो, तो ऐसे में आप सेबी को सूचित करेंगे कि कितनी फीस लौटाई जानी है, साथ ही आप यह भी बताएंगे कि आपने लौटाई जाने वाली फीस की रकम की गणना कैसे की है और सेबी को किसके नाम पर चेक जारी करना होगा।

If filing fees paid by you are more than the actual fees required to be paid, you are advised to inform SEBI about the amount to be refunded, along with detailed calculation of amount refundable and name of the person in whose favour, the cheque may be issued by SEBI.

7. प्रस्तावित निर्गम (इश्यू) इस अभिमत पत्र के जारी होने की तारीख से 12 महीनों के भीतर पैसा लगाने (अभिदान करने / सब्स्क्रिप्शन) के लिए खोला जा सकता है ।

The proposed issue can open for subscription within a period of 12 months from the date of issuance of this observation letter.

Place: Mumbai

Sonal Pednekar



#### **OBSERVATIONS**

- 1. Please refer to our email dated July 21, 2022, and the response from LM dated August 3, September 22,2022 and all other correspondences exchanged. Wherever the LM has undertaken to modify/delete/amend the risk factors or in other relevant section of the DRHP in its replies, the same shall be duly modified and incorporated in the UDRHP/RHP.
- 2. Clause 24 (3) of SEBI (ICDR) Regulations, 2018, requires LM to exercise due diligence and satisfy himself about all aspects of the issue including the veracity and adequacy of disclosures in the offer document. In view of the same, LM is advised to ensure that:
  - i. The offer document shall not contain any information where no responsibility is taken by the BRLMs or the Issuer Company / Expert.
  - ii. The "Industry Overview" section represents a fair and true view of the comparable industry scenario and the same is neither exaggerated nor have any underlying assumptions been omitted for investors to make an informed decision.
- 3. LM is advised to ensure that the processing fees for applications made by Retail Individual Bidders using the UPI Mechanism may be released to the remitter banks (SCSBs) only after such banks provide a written confirmation on compliance with SEBI Circular No: SEBI/HO/CFD/DIL2/P/CIR/2021/570 dated June 02, 2021 read with SEBI Circular No: SEBI/HO/CFD/DIL2/CIR/P/2021/2480/1/M dated March 16, 2021.
- 4. LM is advised to ensure that UDRHP is filed with SEBI not less than seven working days prior to submission of the draft price band advertisement.
- 5. LM is advised to take note of amendments to ICDR Regulations dated November 21,2022 with respect to Key Performance Indicators and ensure compliance with the same.
- 6. LM is advised to ensure that UDRHP contains necessary updated disclosures justifying the offer price under Section "Basis for offer price", "Risk Factors" etc., particularly emphasizing on appropriate Key Performance Indicators as applicable to the industry in which the issuer company operates, in quantitative terms, with corresponding suitable explanations so as to justify the offer price. (For illustration, P/E ratio in case DRHP is filed under Regulation 6 (1) of the ICDR Regulations (and /or) Market Cap / Total Revenue ratio in case DRHP is filed under Regulation 6 (2) of the ICDR Regulations)



- 7. LM is advised to disclose segregated amount of fresh issuances which shall be utilized for repayment of loans of the Company.
- 8. **Summary of the offer document:** LM is advised to include ratio of split and bonus issue dated April 4, 2022 and April 27,2022, respectively. Further, LM is advised to disclose that CCPS will be converted into equity shares prior to filing of RHP.
- 9. **Risk Factors** (i) Every risk factor shall be provided with a cross-reference to the detailed description of the facts / reasons in the DRHP, wherever applicable. (ii) In all risk factors, wherever either only percentages or the absolute values are mentioned, LM shall ensure to disclose both the absolute values and percentages adequately.
- 10. LM is advised to use only restated financial information under all the risk factors.
- 12. **Risk Factor 5**: LM is advised to remove the word 'leading' and include number of Indian pharmaceutical companies who are CDMO customers.

Further, LM is advised to modify stating that any reduction in number of CDMO customers and any adverse developments or inability to enter into or maintain relationships with these CDMO customers could have an adverse impact on the business, results of operations and financial conditions.

LM is also advised to add the percentage of revenue from operations from CDMO business in the same table.

- 13.LM is advised to include RF 5 and 11 sequentially.
- 14. Risk Factor 14: LM is advised to redraft the heading to reflect the risk explained in the third paragraph.
- 15. Risk Factor 16: LM is advised to include this RF as top 15.
- 16. **Risk factor 17**: LM is advised to include this RF as top 10 and modify the heading as: 'Our business is capital intensive. Any insufficient cash flows from our operations or inability to borrow to meet our working capital requirements may materially and adversely affect our business and results of operations'.



- 17. Risk Factor 19: LM is advised to delete 'which are routine in nature.'
- 18. Risk factor 21: LM is advised to include it under top 15 risk factors.
- 19. Risk Factor 22: LM is advised to include it under top 10 risk factors.
- 20. **Risk Factor 29**: LM is advised to redraft heading stating that export is from sale of goods and import for raw materials.
- 21. Risk Factor 34: LM is advised to include it under top 10 risk factors.
- 22. Risk Factor 44: LM is advised to disclose amount of outstanding debt and date of availing the same.
- 23. Risk Factor 57: LM is advised to remove future reference in the heading.
- 24. Risk Factor 58: LM is advised to redraft heading of the risk factor.
- 25.LM is advised to include a separate risk factor in top 10 that more than 90% of the raw materials are imported from one country (i.e. China).
- 26.LM is advised to include separate risk factor in top 15 regarding that nearly 30% of revenue is from related parties.
- 27.LM is advised to include separate risk factor in case there is any delay in expansion of existing plants.
- 28. Capital Structure: LM is advised to include a table showing pre-issue number of shares held and percentage holding for promoter, promoter group and selling shareholder.
- 29. Page 94: LM is advised to remove legal counsel to the BRLMs as to Indian Law.
- 30.LM is advised to ensure compliance with respect to provisions of independent directors as per SEBI LODR regulations.
- 31. Page 104: LM is advised to disclose reason for reduction in shareholding from 39.98% to 30.23% of Gian Parkash Aggarwal or alternatively, provide a table as given in 7(iii) for promoters on page 105.
- 32. Page 116/117: LM is advised to remove future projections.
- 33. Page 171: LM is advised to disclose break up of revenue for CDMO business and other business.





- 34. Page 207: LM is advised to disclose brief financial highlights for past 3 years and also include nature of activities carried out for all subsidiaries.
- 35. Page 237: LM is advised to provide details of promoters, brief financials for past 3 years and nature of activities undertaken by all group companies.
- 36.LM is advised to ensure that the main / sub-headings in the UDRHP / RHP do not have any abbreviations.
- 37.LM is advised that reference to name of any place mentioned in the offer document may be followed by name of City / State, as the case may be.
- 38.LM is advised to include updated audited financials at the time of filing UDRHP/RHP.
- 39. Page 413: LM is advised to disclose risk and the impact of recall of products under para (b).
- 40.LM is advised to asses and disclose the impact (if any), if the cases are not in favour of the Company/directors.
- 41. With respect to all the complaints received by LM / Company / forwarded by SEBI, LM is advised to ensure that there is adequate redressal of the complaint and relevant disclosures of the same are made in the Red Hearing Prospectus and other Offer related material along with the disclosures of the financial impact of the same, if any.
- 42.LM is advised to ensure that the disclosure of details of all the criminal matters initiated by or against the company, group, directors, promoters, subsidiaries which are at FIR stage and no / some cognizance has been taken by court, is incorporated in the UDRHP / RHP along with appropriate risk factors in this regard.
- 43.LM is advised to ensure following disclosures in the Issue advertisement for announcement of Price Band and all further advertisements as a box item below the price band:

### "Risks to Investors:

- i. The [to be disclosed] Merchant Bankers associated with the issue have handled [to be disclosed] public issues in the past three years out of which [to be disclosed] issues closed below the issue price on listing date."
- ii. Any adverse data in the basis for issue price should be disclosed. For example:



- "The Price/Earnings ratio based on diluted EPS for [latest full financial year] for the issuer at the upper end of the Price band is as high as [to be disclosed] as compared to the average industry peer group PE ratio of [to be disclosed]."
  - [if average industry peer group PE ratio is not available, then P/E of Nifty Fifty may be disclosed]
- "Average cost of acquisition of equity shares for the selling shareholders in IPO is [to be disclosed] and offer price at upper end of the price band is [to be disclosed]."
- "Weighted Average Return on Net Worth for [last three full financial years] is [to be disclosed] %."

The data on above disclosures shall be updated and disclosed prominently (in the same font size as the price band) in advertisements of Price Band and all further advertisements, website of the company and the stock exchange. Further, any adverse ratio / data in basis for issue price should be disclosed.

- 44.LM shall submit the draft advertisement for announcement of Price Band with SEBI before its publication in the newspapers for our comments, if any.
- 45. LM is advised to ensure compliance with the below email advisory sent to AIBI through email dated November 13 and November 15, 2021 and amendment to ICDR dated November 21,2022:
  - a. LM shall ensure that all issuer companies filing offer document should provide Price at which specified security was acquired in the last 3 years, by each of the promoters, promoter group, selling shareholders, shareholders entitled with right to nominate directors or any other rights. Following details may be disclosed for such transactions in tabular format name of acquirer, date of acquisition, number of shares acquired and acquisition price per share.
  - b. The portion pertaining to "Risks to Investors" shall constitute at least 33% of the price band advertisement space.
  - c. The risks to investors shall include weighted average cost of acquisition of all shares transacted in last 3 years, 18 months and 1 year, from the date of RHP, in the following format:

Period	Weighted Average Cost of Acquisition (in Rs.)	times the Weighted	Range of acquisition price: Lowest Price - Highest Price (in Rs.)
Last 1			
year			



	<u></u>	I CA CO I ATTOMICA
Last 18		
months		
Last 3		
years		

Range of acquisition should show lowest price of acquisition excluding gift/bonus.

- d. The font size for price band and "Risk to investors" should be increased to match the font of BID/Offer Programme.
- e. Matters related to ASBA and UPI may be brought subsequent to Price Band, Risks to Investors, Bid/ Offer Programme and other offer details, and can be of smaller font.

The portion pertaining to "BRLMs" shall not constitute more than 10% of the price band advertisement space.

46.LM is advised to suitably incorporate the comments of the stock exchanges, if any in the UDRHP/RHP.

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Annexure II

#### **General Observations**

- 1. LM is advised to ensure that prior to filing of RHP with Registrar of Companies, the Issuer Company has received crucial clearances / licenses / permissions / approvals from the required competent authority which are necessary for commencement of the activity for which the issue proceeds are proposed to be utilized.
- 2. LM is advised to ensure that the 'Observation Letter' issued by SEBI is included among the material contracts and documents for inspection.
- 3. LM is advised to ensure that prior to proceeding with the issue, "No Objection Certificates" are obtained from all the lenders with whom the company has entered into an agreement and the terms of such agreement require an approval to be taken.
- 4. LM is advised to ensure that adequate disclosures are made to disclose any material development which may have a material effect on the Issuer Company between the date of registering final prospectus or the RHP or the letter of offer, with the Registrar of Companies or designated stock exchange, as the case may be, and the date of allotment of specified securities, while ensuring compliance with Regulation 42 and Schedule IX of SEBI (ICDR) Regulations, 2018.
- 5. LM is advised to ensure that exact cross-referencing of page numbers is provided in the offer document instead of general cross-referencing.
- In terms of SEBI Circulars No. SEBI/CIR/ISD/03/2011, No. SEBI/CIR/ISD/05/2011 and SEBI/CIR/ISD/01/2012 dated June 17, 2011, September 30, 2011 and March 30, 2012 respectively, LM is advised to ensure that 100% promoter holding is in demat form prior to listing.
- 7. LM is advised to ensure that SCORES authentication is taken by the issuer company prior to listing.
- 8. In pursuance of Regulation 25 Sub-Regulation 9(a) of SEBI (ICDR) Regulations, 2018, LM is advised to certify while submitting the in-seriatim reply that all amendments, suggestions and observations advised by SEBI have been complied with and duly incorporated in the offer document, while also indicating the page number for the same.





#### 9. **ASBA**:

- i) LM is advised to ensure that sufficient number of Physical ASBA forms are printed and dispatched directly to all designated branches of SCSBs which are located in places of mandatory collection centers as specified in Schedule XII of SEBI (ICDR) Regulations, 2018, Syndicate Members and Registered Brokers of Stock Exchanges, the Registrars to an Issue and Share Transfer Agents (RTAs) and Depository Participants (DPs) registered with SEBI, at least two days before the opening of the issue. This shall be in addition to ASBA forms which shall be sent to controlling branch of SCSBs for sending to designated branches other than those located in mandatory collection center.
- ii) LM is advised to ensure that the ASBA mode of payment is highlighted in bold in all the advertisement / communication informing about the issue. Further, LM is also advised to ensure that the following is suitably incorporated in all advertisements / communications regarding the issue issued by the issuer:
  - a. The following may appear just below the price information of the issue as shown below:

"PRICE BAND: RS. XX TO RS. XX PER EQUITY SHARE OF FACE VALUE OF RS. XX EACH

THE FLOOR PRICE IS XX TIMES OF THE FACE VALUE AND THE CAP PRICE IS XX TIMES OF THE FACE VALUE

BID CAN BE MADE FOR A MINIMUM OF XX EQUITY SHARES AND IN MULTIPLES OF XX EQUITY SHARES THEREAFTER.

#### ASBA .

(APPLICATION SUPPORTED BY BLOCKED AMOUNT)

Simple, Safe, Smart way of Application !!!

Mandatory in public issue .No cheque will be accepted



now available in ASBA for retail individual investors.



\* ASBA is a better way of applying to issues by simply blocking the fund in the bank account.

For further details check section on ASBA below."

b. The following paragraph on ASBA may be inserted in the advertisement/Communications:

"ASBA has to be availed by all the investors except anchor investors. UPI may be availed by Retail Individual Investors.

For details on the ASBA and UPI process, please refer to the details given in ASBA form and abridged prospectus and also please refer to the section "Issue Procedure - Issue Procedure of ASBA Bidders" beginning on page xxx of the Red Herring Prospectus. The process is also available on the website of AIBI and Exchanges in the General Information Document."

ASBA bid-cum application forms can be downloaded from the websites of Bombay Stock Exchange and National Stock Exchange and can be obtained from the list of banks that is displayed on the website of SEBI at www.sebi.gov.in.\*\* List of banks supporting UPI is also available on the website of SEBI at www.sebi.gov.in.\*\*.

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SEBI registration no.: INM000010361 CIN: L67120MH1986PLC038784

December 6, 2023

#### **Securities and Exchange Board of India**

Corporation Finance Department Division of Issues and Listing - 2 SEBI Bhavan, Plot No. C4 A, 'G' Block Bandra Kurla Complex Bandra (East) Mumbai 400 051, Maharashtra, India

Kind Attention: Ms. Sonal Pednekar, Assistant General Manager

Dear Madam

Re: Proposed initial public offering of equity shares of face value of ₹ 10 each (the "Equity Shares") of Innova Captab Limited (the "Company") comprising of a fresh issue of Equity Shares aggregating up to ₹ 3,200.00 million ("Fresh Issue") and an offer for sale of up to 9,600,000 Equity Shares ("Offered Shares") aggregating up to ₹ [•] million, comprising of up to 3,200,000 Equity Shares aggregating up to ₹ [•] million by Manoj Kumar Lohariwala, up to 3,200,000 Equity Shares aggregating up to ₹ [•] million by Vinay Kumar Lohariwala (together with Manoj Kumar Lohariwala, referred to as the "Promoter Selling Shareholders") and up to 3,200,000 Equity Shares aggregating up to ₹ [•] million by Gian Parkash Aggarwal (the "Other Selling Shareholder", and together with the Promoter Selling Shareholders, referred to as the "Selling Shareholders", and such offer for sale of equity shares by the Selling Shareholders, referred to as the "Offer for Sale"). The Offer for Sale together with the Fresh Issue is referred to as the "Offer".

This is with reference to the letter bearing reference number SEBI/HO/CFD/RAC-DIL2/P/OW/2023/1400/1 dated January 11, 2023 (the "Final Observations"), issued by the Securities and Exchange Board of India ("SEBI"), in connection with the draft red herring prospectus of the Company dated June 28, 2022 (the "Draft Red Herring Prospectus" or "DRHP"), as amended by the addendum dated September 12, 2023 (the "Addendum"), each filed with SEBI in relation to the Offer.

The Company now wishes to file a draft of its red herring prospectus (the "UDRHP") with SEBI prior to filing of the red herring prospectus ("Red Herring Prospectus" or "RHP") with the Registrar of Companies, Maharashtra at Mumbai, ("RoC"), SEBI and the relevant stock exchanges ("Stock Exchanges").

In this regard, based on the confirmations, information and documents made available to us, representations of the Company and our discussions with representatives of the Company, please find enclosed the following:

- 1. In-seriatim response to the Final Observations, attached as **Annexure A**, along with references to the relevant page numbers of the UDRHP, where such observations have been addressed.
- 2. A draft of the UDRHP in clean mode, attached as **Annexure B**, which includes all the changes and updates made to the DRHP, and a blackline of the UDRHP against the DRHP, attached as **Annexure C**, to indicate the changes made to the DRHP. Please note that in the blackline version, all insertions in the UDRHP appear as underlined text while the deletions appear as strikethrough text. However, the updated financial information, including the updated proforma financial information, have been included, in the final form and without track changes, for ease of reference. The page number references in this letter relate to the



relevant pages in either the clean copy or the blackline copy of the UDRHP as indicated at the relevant column. Please note that where the page number of the blackline copy has been provided, the page number of the PDF file has been provided.

The Company has made a payment of ₹ 30,000.00 (Rupees thirty thousand only) plus GST at the rate of 18% aggregating to ₹ 35,400 (Rupees thirty five thousand and four hundred only) on December 5, 2023, as per mentioned below table towards additional fees as specified in as specified in Schedule III of the Securities and Exchange Board of India (Issue of Capital and Disclosure Requirements) Regulations, 2018, as amended ("SEBI ICDR Regulations") read with Regulation 25(6) and Schedule XVI of the SEBI ICDR Regulations, in respect of updates to certain sections in the UDRHP.

Particulars	Details
Fee paid for updates to the DRHP	₹ 30,000
GST paid	₹ 5,400
Total Additional Fee	₹ 35,400
Payment reference number/Tracking ID	2616004/113099015015
Payment Date	05-12-2023
GST Registration Number	02AABCH5082R2ZA
GST Registered Dealer Name	Innova Captab Limited
GST Registered Office Address	1281/1 Hill Top, Jharmajri, Baddi, District Solan, Himachal
	Pradesh
Location from where payment was discharged	Panchkula

The relevant extract as proof of payment being made is attached herewith as **Annexure D**.

We have set out below a tabular description of the compliance with the requirements specified in Schedule XVI, read with Regulation 25(6), of the SEBI ICDR Regulations:

#### (a) Changes which require fresh filing of the draft offer document with SEBI, along with fees:

S. No.	Nature of Update	Applicability
1.	Change in promoter of the issuer.	Not applicable
2.	Change in more than half of the board of directors of the issuer.	Not applicable
3.	Change in main object clause of the issuer.	Not applicable
4.	Any addition to objects of the issue resulting in an increase in estimated issue	Not applicable
	size or estimated means of finance by more than twenty per cent.	
5.	If there are grounds to believe that there is an exacerbation of risk on account of deletion of an object resulting in a decrease in issue size by more than twenty	Not applicable
	per cent.	
6.	<ul> <li>Any Increase or Decrease:</li> <li>(i) In case of a fresh issue: any increase or decrease in estimated issue size by more than twenty percent. of the estimated issue size; or</li> <li>(ii) In case of an offer for sale: any increase or decrease in either the number of shares offered for sale or the estimated issue size, by more than fifty per cent.; or</li> <li>(iii) In case of an issue comprising of both fresh issue and offer for sale: the</li> </ul>	Not applicable
	(iii) In case of an issue comprising of both fresh issue and offer for sale: the respective limits as above shall apply.	

<b><i>ICICI</i></b> Securities	JM FINANCIAL
ICICI Securities Limited	JM Financial Limited
ICICI Venture House	7th Floor, Cnergy
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Website: www.icicisecurities.com	Website: www.jmfl.com
SEBI Registration No: INM000011179	SEBI registration no.: INM000010361
CIN: 167120MH1995DLC086241	CIN: 167120MH1986DLC038784

S. No.	Nature of Update	Applicability
7.	Any increase in estimated deployment in any of the objects of the issue by more than 20%.	Not applicable
8.	Changes which may result in non-compliance with the provisions of the SEBI ICDR Regulations and the lead managers or the Company do not intend to seek relaxation under Regulation 303 of the SEBI ICDR Regulations.	

### (b) Changes which require filing of the updated offer document with SEBI, along with fees:

	nges which require filing of an updated offer cument with SEBI, along with fees	Applicability
Section 1: Risk Factors:	Any material development which may result in potential risk and require updation in this section.	The risk factors disclosed in the section titled "Risk Factors" of the UDRHP have been revised to reflect: (a) updates to the business of the Company since the date of filing of the DRHP and pursuant to inclusion of financial statements for the three months ended June 30, 2023, and the financial years ended March 31, 2023, and March 31, 2022, in the Restated Consolidated Financial Information, and the inclusion of the updated Proforma Condensed Consolidated Financial Information for the financial year ended March 31, 2023 to reflect the acquisition of Sharon Bio-Medicine Limited by the Company, (b) updated operational data pertaining to the business of the Company, (c) updates with respect to the legal proceedings involving the Company, (d) factual updates which have occurred post filing of the DRHP, and (e) changes in accordance with the responses to initial interim and Final Observations, and emails or other communications, received from SEBI, and the observations and emails received
Section 2: Capital Structure	An aggregate increase of 5% or more in the shareholding of the promoter or promoter group or an aggregate increase of 5% or more in the shareholding of the top ten shareholders.	from the Stock Exchanges in this regard.  Not applicable
Section 3: Issue Size	Any addition or deletion to the objects of the issue resulting in a change in the estimated issue size or estimated means of finance by more than 10% and not exceeding 20%.	Not applicable





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	nges which require filing of an updated offer cument with SEBI, along with fees	Applicability		
Section 4: Management	Appointment of any new director.	Not applicable		
Section 5: Promoter Group	Any addition to the promoter group or group companies.	Not applicable		
Section 6: Financial Statements	Any variation in net profits after tax or net loss after tax and / or extraordinary items in excess of 10% over the last updated financials submitted to SEBI.	Applicable.  The UDRHP has been updated to include the 'Restated Consolidated Financial Information' comprising the restated consolidated financial information of the Company and its Subsidiaries, as of and for the three months ended June 30, 2023, and as of and for the years ended March 31, 2023, March 31, 2022 and March 31, 2021.		
Section 7: Legal and other information	Any new litigation or any development about a pending litigation which is material in view of the merchant bankers	Applicable.  The section titled "Outstanding Litigation and Material Developments" beginning on page 424 of the UDRHP has been updated as follows: (i) to reflect changes in the status of the legal proceedings disclosed in the DRHP, where applicable; (ii) to include details of outstanding criminal proceedings, pending tax claims and other material litigation involving Sharon Bio-Medicine Limited, our Subsidiary acquired subsequent to the date of filing of the DRHP; and (iii) to include details of certain new outstanding litigation.		

Except as mentioned above, there are no other changes made to the DRHP which require payment of additional fees pursuant to paragraph 2(d) of Schedule III read with paragraph 2 of Schedule XVI of the SEBI ICDR Regulations.

We undertake to submit a detailed calculation of filing fees in terms of Regulation 25(6) and Schedule III of the SEBI ICDR Regulations within seven days of the filing of the Prospectus with the RoC, along with details of filing fees paid till date, in accordance with the Final Observations.

In addition, please see below the key updates since the date of filing of the DRHP in relation to the Company and the proposed Offer that have been included in the UDRHP, and shall be included in the RHP and the Prospectus.

#### Key updates





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Please find below the key updates since the date of the DRHP in relation to the Company and the proposed Offer that have been included in the UDRHP. The information contained in the DRHP has been updated to reflect the position of the Company as of a recent date and to address the observations, to the extent applicable, made by SEBI in the SEBI Observations (defined below):

### Amendments to the SEBI ICDR Regulations notified on July 25, 2022, November 21, 2022, January 13, 2023, and May 23, 2023, and changes introduced pursuant to the SEBI circular SEBI/HO/CFD/TPD1/CIR/P/2023/140 dated August 9, 2023

Amendments to the SEBI ICDR Regulations were notified on July 25, 2022, November 21, 2022, January 13, 2023, and May 23, 2023. Accordingly, updates to the UDRHP have been made pursuant to these amendments, to the extent applicable, including the identification of the Senior Management of the Company, pursuant to which the section titled "Our Management" has been updated in the UDRHP, and the disclosure of certain key performance indicators of the Company as required in terms of the SEBI ICDR Regulations, pursuant to which the section titled "Basis for the Offer Price" has been updated in the UDRHP.

Pursuant to SEBI circular SEBI/HO/CFD/TPD1/CIR/P/2023/140 dated August 9, 2023, the timeline of T + 6 days from the bid / offer closing date until the date on which the equity shares of the Company are listed on the Stock Exchanges has been mandatorily reduced to T + 3 days for public issues opening on or after December 1, 2023. Accordingly, updates to the UDRHP have been made to reflect that the Offer will be undertaken with the reduced T + 3 timelines, including to the section titled "Offer Procedure".

#### 2. Updates pursuant to the acquisition of Sharon Bio-Medicine Limited ("Sharon") by the Company

As highlighted in the Addendum, the Company acquired Sharon, a listed entity, through the corporate insolvency resolution process ("CIRP") under the Insolvency and Bankruptcy Code, 2016 ("IBC") before the Hon'ble National Company Law Tribunal, Mumbai Bench ("NCLT"). In accordance with the terms of the resolution plan approved by the NCLT, Univentis Medicare Limited ("UML"), a Subsidiary of the Company, infused ₹ 1,954.00 million into Sharon on June 26, 2023. The implementation of the plan was completed on June 30, 2023, the closing date as per the approved resolution plan, and subsequently, control and sole ownership over Sharon was established, pursuant to which Sharon became a wholly owned subsidiary of UML as of June 30, 2023. The UDRHP has been updated to include updates and additional disclosures as may be required under the SEBI ICDR Regulations in connection with the Company's acquisition of Sharon, including in the sections titled "Our Business", "Risk Factors", "History and Certain Corporate Matters", "Outstanding Litigation and Other Material Developments", "Government and Other Approvals" and "Management's Discussion and Analysis of Financial Condition and Results of Operations".

#### 3. Appointment of new Chief Financial Officer and a deputy Chief Financial Officer of the Company

At the time of filing the DRHP, the late Rishi Gupta was the chief financial officer of the Company. The Company has appointed Gaurav Srivastava as its new Chief Financial Officer on August 12, 2023. In addition, Mukesh Kumar Singh, who is the deputy chief financial officer of the Company, was designated as a Key Managerial Personnel with effect from June 30, 2023 pursuant to a resolution of the Board of Directors dated November 10, 2023. Accordingly, the UDRHP has been updated to reflect the appointment of a new Chief Financial Officer, including disclosures in connection with Gaurav Srivastava's appointment, his profile, the updated management organisation, etc., as required under the SEBI ICDR Regulations. The UDRHP has also been updated to reflect Mukesh Kumar Singh' details including his profile as a Key Managerial Personnel of the Company.





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## 4. Completion of Pre-IPO Placement and revision of the size of the Fresh Issue pursuant to the Pre-IPO Placement

As informed in our letters to SEBI dated July 8, 2023, and July 22, 2022, the Company, in consultation with the BRLMs, had initially undertaken a pre-IPO placement by way of a private placement of 1,412,430 compulsorily convertible preference shares ("CCPSs") at a price of ₹ 354.00 per CCPS (including a premium of ₹ 344.00) aggregating to ₹ 500.00 million. Thereafter, the Company, in consultation with the BRLMs, has undertaken another pre-IPO placement by way of a private placement of 669,642 Equity Shares with 334,821 Equity Shares allotted each to 360 One Special Opportunities Fund - Series 9 and 360 One Special Opportunities Fund - Series 10 at a price of ₹448.00 per Equity Share (including a premium of ₹438.00) aggregating to ₹300.00 million. Please note that the Company published advertisements dated July 8, 2023 and December 5, 2023, in all editions of Financial Express (a widely circulated English national daily newspaper), all editions of Jansatta (a widely circulated Hindi national daily newspaper) and the Mumbai edition of Navshakti (a widely circulated Marathi daily newspaper, Marathi being the regional language of Maharashtra where our Registered Office is located), with all the relevant details of the Pre-IPO Placements.

The size of the Fresh Issue of up to ₹ 4,000.00 million as disclosed in the DRHP has, in the aggregate, been reduced by ₹ 800.00 million, pursuant to the two pre-IPO placements discussed above ("Pre-IPO Placement") and, accordingly, the revised size of the Fresh Issue is up to ₹ 3,200.00 million. Appropriate disclosures in this regard have been updated in the UDRHP, including in the sections titled "Definitions and Abbreviations", "Summary of the Offer Document", "Capital Structure", "Objects of the Offer" and "Offer Structure".

Further, the Company had amended its Articles of Association on July 19, 2022, to enable the issuance of the CCPSs pursuant to the Pre-IPO Placement, as intimated to SEBI vide our letter dated July 22, 2022. However, pursuant to the e-mail dated June 28, 2023 received from SEBI, the Articles of Association have been further amended in order to ensure that no special rights are available to the Promoters / Shareholders of the Company in the Articles of Association, at the time of filing of the Red Herring Prospectus. As a result, Part-B of the Articles of Association have been deleted and Part-A of the Articles of Association have come in effect and in force. Accordingly, the Articles of Association as appearing in the section titled "Main Provisions of the Articles of Association" have been updated in the UDRHP to reflect the amendment to the Company's Articles of Association.

The CCPSs have been converted into 1,412,430 Equity Shares on December 1, 2023. Details in connection with the conversion of the CCPSs into Equity Shares have been updated in the UDRHP, including in the section titled "Capital Structure".

#### 5. Change in the registered office of the Company

Pursuant to the resolution of the shareholders of the Company passed on October 16, 2023, the Company has changed its registered office with effect from October 16, 2023, from Office No. 606, Ratan Galaxie – 6th Floor, Plot No. 1, J. N. Road, Mulund (W), Mumbai, Maharashtra 400 080, India to 601, Proxima, Plot No. 19, Sector 30 A, Vashi, Navi Mumbai, Maharashtra 400 705, India. Appropriate updates have accordingly been made to the UDRHP, including to the cover page and the sections titled "General Information" and "History and Certain Corporate Matters".





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#### 6. Updated financial information

The financial information of the Company has been updated to include (i) the restated consolidated financial information of the Company for the three months ended June 30, 2023, and the financial years ended March 31, 2023, March 31, 2022, and March 31, 2021, and (ii) the proforma condensed consolidated financial information of the Company, prepared on a voluntary basis, for the financial year ended March 31, 2023, to reflect the acquisition of Sharon Bio-Medicine Limited by the Company, comprising the proforma condensed consolidated statement of profit and loss which has been prepared as if the acquisition occurred immediately before the start of the financial year ended March 31, 2023, and the proforma condensed consolidated balance sheet which has been prepared as if the acquisition occurred as at March 31, 2023. Accordingly, the sections titled "Summary Financial Information", "Restated Consolidated Financial Information" and "Proforma Condensed Consolidated Financial Information" have been updated in the UDRHP. Further, pursuant to the updated financial information being available, appropriate updates have been made to the sections titled "Summary of the Offer Document", "Risk Factors", "Basis for the Offer Price", "Other Financial Information", "Capitalisation Statement", "Our Business", "Outstanding Litigation and Other Material Developments" and "Management's Discussion and Analysis of Financial Condition and Results of Operations", amongst others.

The section titled "Statement of Special Tax Benefits" has been updated in the UDRHP to include the revised statement of special tax benefits available to the Company, its shareholders and its Material Subsidiaries provided by B S R & Co. LLP, Chartered Accountants, the statutory auditors of the Company.

#### 7. Updates in business, industry and operational data

The operational data and key performances indicators disclosed in the DRHP have been updated in the UDRHP to include details as of a later date, including such as details in relation to the employee base of the Company which have been updated to reflect these details as on October 31, 2023, the details of outstanding dues to creditors of the Company which have been updated to reflect these details as on June 30, 2023, details of the total consolidated outstanding borrowings of the Company which have been updated to reflect these details as on October 31, 2023 and details of the outstanding borrowings of the Company proposed to be repaid / prepaid through the Net Proceeds which have been updated to reflect these details as on October 31, 2023, among others. In light of updated operational data and key performance indicators being available, the sections titled "Summary of the Offer Document", "Risk Factors", "Our Business", and "Management's Discussion and Analysis of Financial Position and Results of Operations" have also been updated.

Further, there have been updates to certain industry information, as detailed in the updated industry report titled "Assessment of Indian pharmaceutical and CDMO market" and dated October 2023, commissioned by the Company and issued by CRISIL Research, a division of CRISIL Limited. Pursuant to the updated industry report, the sections titled "Industry Overview", "Summary of the Offer Document", "Risk Factors", "Our Business", "Basis for the Offer Price" and "Management's Discussion and Analysis of Financial Position and Results of Operations" have also been updated.

#### 8. Updates in capital structure

As highlighted in paragraph 4 above, our Company has, following the filing of the DRHP, undertaken the Pre-IPO Placement of:





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E-mail: innova.ipo@icicisecurities.com Website: www.icicisecurities.com SEBI Registration No: INM000011179 CIN: L67120MH1995PLC086241 JM Financial Limited

7th Floor, Cnergy Appasaheb Marathe Marg Prabhadevi, Mumbai 400 025 Maharashtra, India

Tel: + 91 22 6630 3030 E-mail: innova.ipo@jmfl.com Website: www.jmfl.com

SEBI registration no.: INM000010361 CIN: L67120MH1986PLC038784

- (i) 1,412,430 CCPSs allotted to UTI Multi Opportunities Fund I at a price of ₹354.00 per CCPS (including a premium of ₹344.00) aggregating to ₹500.00 million. The CCPSs have been converted into 1,412,430 Equity Shares; and
- (ii) 669,642 Equity Shares with 334,821 Equity Shares allotted each to 360 One Special Opportunities Fund Series 9 and 360 One Special Opportunities Fund Series 10 at a price of ₹448.00 per Equity Share (including a premium of ₹438.00) aggregating to ₹300.00 million.

Details in connection with the Pre-IPO, including the conversion of the CCPSs into Equity Shares have been updated in the UDRHP, including in the section titled "Capital Structure".

#### 9. In-principle listing approvals

The Company has received 'in-principle' approvals from BSE Limited and the National Stock Exchange of India Limited for the listing of the Equity Shares, pursuant to letters dated September 16, 2022, and September 15, 2022, respectively, and appropriate updates have accordingly been made to the UDRHP to reflect this. Copies of such approval letters have been filed with your office through our letter dated September 16, 2022.

# 10. Appointment of the Escrow Collection Banks, Refund Bank, Public Offer Account Bank, Sponsor Bank, Syndicate Member, the Share Escrow Agent and the Monitoring Agency

For the purposes of the Offer, ICICI Bank Limited and HDFC Bank Limited shall act as the Escrow Collection Banks and the Sponsor Banks. Further, ICICI Bank Limited shall also act as the Public Offer Account Bank and the Refund Bank. In addition, JM Financial Services Limited shall act as the Syndicate Member, KFin Technologies Limited shall act as the Share Escrow Agent and CRISIL Ratings Limited shall act as the Monitoring Agency for the proposed Offer.

Details of the Bankers to the Offer, Syndicate Member, Share Escrow Agent and Monitoring Agency, for the purpose of the Offer, as required under the SEBI ICDR Regulations, have been updated at the appropriate places in the UDRHP. Further, the Cash Escrow and Sponsor Bank Agreement, the Share Escrow Agreement, the Syndicate Agreement and the Monitoring Agency Agreement ("RHP Stage Agreements"), will be signed prior to filing of the RHP with the RoC.

#### 11. Appointment of Designated Stock Exchange

The Company has appointed National Stock Exchange of India Limited ("**NSE**") as the designated stock exchange for the purposes of the Offer pursuant to the resolution passed by the Board of Directors dated November 10, 2023. Appropriate disclosures to this effect have been incorporated in the UDRHP.

#### 12. Price information of past issues

The price information of past issues of the BRLMs has been updated in the UDRHP to reflect updates since the date of filing of the DRHP. We further undertake to update this information, if applicable, in the RHP prior to filing with the RoC.

#### 13. Miscellaneous





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- (a) The Offer Agreement dated June 28, 2022 ("Offer Agreement"), entered between the Company, BRLMs and Selling Shareholders was amended on September 12, 2023, including to modify certain representations and warranties in connection with the acquisition of Sharon Bio-Medicine Limited by the Company. The Offer Agreement shall be further amended to reflect the update in the dispute resolution mechanism under the Offer Agreement, in light of the SEBI master circular dated July 31, 2023 bearing reference number SEBI/HO/OIAE/OIAE\_IAD-1/P/CIR/2023/145, as amended pursuant to the SEBI circular dated August 4, 2023 bearing reference number SEBI/HO/OIAE/OIAE\_IAD-1/P/CIR/2023/135 ("SEBI ODR Circulars").
- (b) The Registrar Agreement dated June 20, 2022, entered into between the Company, the Selling Shareholders and the Registrar to the Offer, shall be further amended to reflect the update in the dispute resolution mechanism under the Registrar Agreement, in light of the SEBI ODR Circulars.
- (c) The Company has obtained SCORES authentication and the disclosure under "Other Regulatory and Statutory Disclosures Disposal of Investor Grievances by our Company" has accordingly been modified in the UDRHP to reflect this.
- (d) Information set out in the DRHP, including in the sections titled "Certain Conventions, Use of Financial Information, Industry and Market Data and Currency of Presentation", "Summary of the Offer Document", "Risk Factors", "The Offer", "Summary Financial Information", "General Information", "Capital Structure", "Objects of the Offer", "Basis for the Offer Price", "Statement of Special Tax Benefits", "Industry Overview", "Our Business", "Key Regulations and Policies", "History and Certain Corporate Matters", "Our Management", "Our Group Companies", "Dividend Policy", "Restated Consolidated Financial Information", "Proforma Condensed Consolidated Financial Information", "Other Financial Information", "Capitalisation Statement", "Financial Indebtedness", "Management's Discussion and Analysis of Financial Position and Results of Operations", "Outstanding Litigation and Other Material Developments", "Government and Other Approvals", "Other Regulatory and Statutory Disclosures", "Terms of the Offer", "Offer Structure", "Offer Procedure" and "Material Contracts and Documents for Inspection", has been updated in the UDRHP to include factual changes since the filing of the DRHP, and reflect the updated position.

In addition to the above, the information contained in the UDRHP is updated to reflect the position of the Company as of a recent date and to address the observations, to the extent applicable, made by SEBI vide the letter dated July 21, 2022 (the "Initial Observations") and the email dated August 23, 2022 (the "Additional Observations", and together with the Initial Observations and the Final Observations, the "SEBI Observations"), wherein we have been advised to provide certain clarifications regarding the DRHP filed with SEBI in relation to the Offer, and other correspondences in relation to DRHP, and the Final Observations. The responses to the observations issued by SEBI have been included in the UDRHP (to the extent indicated in our responses) after taking into consideration factual updates as applicable.

Further, the following changes will be made to the RHP, before filing the RHP with the RoC:

a. The finalized schedule of the Offer and the corresponding dates, including the Bid / Offer Period and the Anchor Investor Bid / Offer Period, will be inserted at the appropriate places. All referenced page numbers will be inserted and updated in a uniform manner. The details of the finalized brokerage structure, processing fees payable to SCSBs, selling commission payable to Registered Brokers, RTAs, CDPs, Sponsor Bank(s) and Syndicate Member will be included prior to filing of the RHP with RoC.





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- b. Details of the Share Escrow Agreement, Cash Escrow and Sponsor Bank Agreement, Syndicate Agreement and Monitoring Agency Agreement to be executed on or before the filing of the RHP with the RoC, will be included in the RHP.
- c. The QR code for access to the RHP will be included on the cover page of the RHP.
- d. Details of the listed peers of the Company in the section titled "Basis for the Offer Price" will be updated in the RHP to reflect the updated details of the respective listed peers of the Company, as may be available, before the filing of the RHP with the RoC;
- e. The price information of past issues undertaken by the BRLMs has been updated in the draft of the UDRHP annexed herein, and shall be further updated, to the extent applicable, prior to filing of the RHP with RoC.
- f. The details of Equity Shares held by the Shareholders holding 1% or more of the paid-up Equity Share capital of the Company in the section titled "Capital Structure", shall, on the basis of the latest BENPOS statement, be updated, as may be required, prior to filing the RHP with the RoC.
- g. Any other material development which may require a change in disclosures in accordance with applicable law will be updated as appropriate in the RHP.

The Price Band, and the minimum Bid Lot will be advertised at least two Working Days prior to the Bid / Offer Opening Date in compliance with the SEBI ICDR Regulations. Further, we undertake to submit a draft copy of the price band advertisement to SEBI in advance before publication in the relevant newspapers.

We confirm that all amendments, suggestions and observations advised by SEBI and the Stock Exchanges have been incorporated in the UDRHP and will be incorporated in the RHP.

We further confirm that as on the date of the UDRHP, the clauses / covenants of Articles of Association of the Company are in compliance with the Companies Act, 2013, as amended, and the securities laws, as applicable.

Please note that no further changes, except formatting changes, changes to correct grammatical, typographical and other errors, changes as indicated in this letter and changes carried out in order to update any information that had been provided in the UDRHP, including any material development or update which may require a change in the disclosures in accordance with applicable law, until the date of filing of the RHP with the RoC, shall be made to the RHP without the specific written consent of SEBI.

All capitalised terms used in this letter (including the annexures and schedules) and not defined shall have the meaning assigned to such terms in the UDRHP.





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We request you to take our responses on record. Should you require any further information or clarifications, please feel free to contact any of the following officials of ICICI Securities Limited:

Contact Person	Telephone	Email
Prem D'cunha	+91 99302 86078	prem.dcunha@icicisecurities.com
Harsh Thakkar	+91 9920327474	harsh.thakkar@icicisecurities.com
Ashik Joisar	+91 7021907267	ashik.joisar@icicisecurities.com

Thanking you.

Sincerely,

Enclosed: Annexures as above.





#### Annexure A

### In-seriatim response to the Final Observations

S. No.	Observations	Response	UDRHP Page Number (clean)	UDRHP Page Number (redline)
		Annexure I – Observations		
1.	Please refer to our email dated July 21, 2022, and the response from LM dated August 3, September 22,2022 and all other correspondences exchanged. Wherever the LM has undertaken to modify/delete/amend the risk factors or in other relevant section of the DRHP in its replies, the same shall be duly modified and incorporated in the UDRHP/RHP.	Complied with and noted for compliance. For ease of reference, please see <b>Schedule A</b> and <b>Schedule B</b> . The relevant changes pursuant to the Initial Observations and our response dated August 3, 2022, and pursuant to the Additional Observations and our response dated September 22, 2022, have been included in the UDRHP after taking into consideration factual updates, as applicable.	-	-
2.	Clause 24 (3) of SEBI (ICDR) Regulations, 2018, requires LM to exercise due diligence and satisfy himself about all aspects of the issue including the veracity and adequacy of disclosures in the offer document. In view of the same, LM is advised to ensure that:  i. The offer document shall not contain any information where no responsibility is taken by the BRLMs or the Issuer Company / Expert.  ii. The "Industry Overview" section represents a fair and true view of the comparable industry scenario and the same is neither exaggerated nor have any underlying assumptions been omitted for investors to make an informed decision.	Complied with and noted for compliance.	-	-
3.	LM is advised to ensure that the processing fees for applications made by Retail Individual Bidders	Noted for compliance.	-	-





S. No.	Observations	Response	UDRHP Page Number (clean)	UDRHP Page Number (redline)
	using the UPI Mechanism may be released to the remitter banks (SCSBs) only after such banks provide a written confirmation on compliance with SEBI Circular No: SEBI/HO/CFD/DIL2/P/CIR/2021/570 dated June 02, 2021 read with SEBI Circular No: SEBI/HO/CFD/DIL2/CIR/P/2021/2480/1/M dated March 16, 2021.	We further submit that the disclosure appearing under the sections titled "Objects of the Offer", "Other Regulatory and Statutory Disclosures" and "Terms of the Offer" on pages 110, 424 and 436 of the DRHP, specifies that the processing fees for applications made by UPI Bidders using the UPI Mechanism may be released to the remitter banks (SCSBs) only after such banks provide a written confirmation on compliance with SEBI Circular No: SEBI/HO/CFD/DIL2/P/CIR/2021/570 dated June 2, 2021 read with SEBI Circular No: SEBI/HO/CFD/DIL2/CIR/P/2021/2480/1/M dated March 16, 2021, and SEBI circular no. SEBI/HO/CFD/DIL2/CIR/P/2022/51 dated April 20, 2022.		
4.	LM is advised to ensure that UDRHP is filed with SEBI not less than seven working days prior to submission of the draft price band advertisement.	Noted for compliance.	-	-
5.	LM is advised to take note of amendments to ICDR Regulations dated November 21, 2022, with respect to Key Performance Indicators and ensure compliance with the same.	Noted for compliance. The required updates with respect to Key Performance Indicators (" <b>KPIs</b> ") have been made to the UDRHP, including to the section titled "Basis for the Offer Price" of the UDRHP, pursuant to the amendment to the SEBI ICDR Regulations dated November 21, 2022.	-	-
6.	LM is advised to ensure that UDRHP contains necessary updated disclosures justifying the offer price under Section "Basis for offer price", "Risk Factors" etc., particularly emphasizing on appropriate Key Performance Indicators as applicable to the industry in which the issuer company operates, in quantitative terms, with corresponding suitable explanations so as to justify the offer price. (For illustration, P/E ratio in case DRHP is filed under Regulation 6 (1) of the ICDR Regulations (and /or) Market Cap / Total Revenue ratio in case DRHP is filed under Regulations)	Complied with and noted for compliance.	-	-
7.	LM is advised to disclose segregated amount of fresh issuances which shall be utilized for repayment of loans of the Company.	Noted for compliance. The table on page 106 of the UDRHP has been updated to include details of the amount of the Net Proceeds proposed to be utilized for the prepayment / repayment of each loan of the Company. For reference, the table has been reproduced in <b>Appendix A</b> below.	106	139





S. No.	Observations	Response	UDRHP Page Number (clean)	UDRHP Page Number (redline)
8.	Summary of the offer document: LM is advised to include ratio of split and bonus issue dated April 4, 2022 and April 27,2022, respectively. Further, LM is advised to disclose that CCPS will be converted into equity shares prior to filing of RHP.	Complied with and noted for compliance.  The ratio of the split and bonus issue dated April 4, 2022, and April 27, 2022, have been updated in the relevant footnotes to the tables (to the extent applicable) in the "Summary of the Offer Document" section of the UDRHP in the following manner:  "The cost of acquisition of equity shares, have been adjusted for the sub-division in the face value of the equity shares of our Company from ₹100 each to ₹10 each, such that each equity share of our Company of face value ₹100 was split into 10 Equity Shares of face value of ₹10 each, pursuant to a resolution of the Shareholders dated April 4, 2022, as applicable."  "Acquired pursuant to a bonus issuance of Equity Shares, in the ratio of three Equity Shares for every one Equity Share held, allotted on April 27, 2022."  Further, the CCPSs issued by the Company pursuant to the Pre-IPO Placement have been converted into 1,412,430 Equity Shares in the ratio of one Equity Share for every CCPS held, on December 1, 2023. Appropriate disclosures in this regard have been made in the UDRHP, including in the section titled "Capital Structure".	-	-
9.	Risk Factors – (i) Every risk factor shall be provided with a cross-reference to the detailed description of the facts/ reasons in the ORHP, wherever applicable. (ii) In all risk factors, wherever either only percentages or the absolute values are mentioned, LM shall ensure to disclose both the absolute values and percentages adequately.	Noted for compliance. We undertake to ensure that (i) every risk factor shall be provided with a cross-reference to the detailed description of the facts / reasons in the red herring prospectus/prospectus, wherever applicable; and (ii) in all risk factors, wherever either only percentages or the absolute values are mentioned, we will disclose both the absolute values and percentages adequately.	-	-
10.	LM is advised to use only restated financial information under all the risk factors.	We have not removed proforma financial information from the risk factor section of the UDRHP. The proforma financial information is important for investors as the restated financial information is not comparable from period to period.	-	-
11.	Risk Factor 1: LM is advised to delete 'In addition, because of their nature does not represent our	Noted for compliance. We have amended the heading of risk factor 1 in the UDRHP as follows:	33	40





S. No.	Observations			Response			UDRHP Page Number (clean)	UDRHP Page Number (redline)
	factual results of operations or financial condition' and include that Proforma Condensed Consolidated Financial Information has not been prepared in accordance with generally accepted accounting principles including accounting standards.	"Our Restated Conso to any future financi Our Company has us the acquisition of UN and for the nine mon 2020 and 2019, are prepare. In additional Information address of operations or fina	al results that we modertaken the acquingle of the acquire	ay prepare. sition of assets and white our Restated Co r 31, 2021 and as of, co each other and to a r nature, our Profor wation and, thereford	liabilities of the Inno onsolidated Financie and for the years end ny future financial i oma Condensed Coi e, does not represen	ova Partnership and al Information, as of, led, March 31, 2021, results that we may a nsolidated Financial of our factual results		
12.	Risk Factor 5: LM is advised to remove the word 'leading' and include number of Indian pharmaceutical companies who are CDMO customers.  Further, LM is advised to modify stating that any reduction in number of CDMO customers and any adverse developments or inability to enter into or maintain relationships with these CDMO customers could have an adverse impact on the business, results of operations and financial conditions.  LM is also advised to add the percentage of revenue from operations from CDMO business in the same table.	Noted for compliance comments) in the UI  "We depend on a ("CDMO") customer number of CDMO or relationships with the operations and final our CDMO business pharmaceutical process own brand names to increased from 108 is consolidated basis of December 31, 2021 CDMO customers on the consolidates of CDMO customers.	DRHP as follows:  limited number of strain including leading to stomers and advited condition.  is focused on providents for Indian pharms the end users. The in Fiscal 2019 to 164 and has increased from a proforma cons	contract developm g Indian pharmaceus erse developments ers could have an ad ding products and se maceutical companie number of CDMO co in the nine months of crom 155 in Fiscal 2 olidated basis. The f	nent and manufact vitical companies. A or inability to ente liverse effect on our  rvices across a divel es who market such p ustomers that we he ended December 31, 019 to 164 in the following table sets	turing organization ny reduction in the er into or maintain business, results of the er range of generic products under their ave represented has 2021 on a restated nine months ended	37	46





S. No.	Observations	Response	UDRHP Page Number (clean)	UDRHP Page Number (redline)
		In Fiscal 2023, we represented 182 customers on a pro forma consolidated basis.  Our business, results of operations and financial condition are dependent on our relationships with and continued supply to our Indian pharmaceutical customers. However, some of our customers may start manufacturing at their own facilities and may discontinue the use of our CDMO services and products. Further, we typically plan and incur capital expenditure for future periods. Delays in successfully entering into contracts for utilization of upcoming capacity may result in lack of proportionate increase in our revenues and results of operations, vis-à-vis an installed capacity increase. In addition, there can be no assurance that we will be able to maintain historic levels or increased levels of business with our significant customers. If we are unable to maintain relationships with the Indian pharmaceutical companies on existing or favourable terms and conditions and if there is delay in replacing these discontinuations with our new products or new customers or maximize utilisation of our installed capacities, it could have an adverse impact on our business, results of operations, margins and financial condition.  The table set forth below provides our revenue from operations from our CDMO business on a restated consolidated basis from our top ten customers and such revenue as a percentage of our operations for the years and period indicated.		





S. No.	Observations		Response								Nu	HP Page mber lean)	UDRHP Page Number (redline)
			Fiscal 2019		Fiscal 2020		Fiscal 2021		Nine Months ended December 31, 2021				
		Revenue from Operations	₹ million	% of revenue from operations from CDMO busines	₹ million	% of revenue from operations from CDMO busines	₹ million	% of revenue from operations from CDMO busines	₹ million	%-of revenue from operati ons from CDMO busines			
		Top Ten Customers  The table set proforma consoperations for	solidated l	<del>basis from</del>	our top to	en custom		-					





S. No.	Observations		Response									UDRHP Page Number (redline)
			Fiscal	<del>2019</del>	Fiscal	<del>2020</del>	Fiscal 2021		Nine Months ended December 31, 2021			
		Revenue from Operations	₹ million	% of revenue from operati ons from CDMO busines	₹ million	% of revenue from operati ons from CDMO busines	₹ million	% of revenue from operations from CDMO busines	<b>₹</b> million	%-of revenue from operations from CDMO business		
		<del>Top Ten</del> <del>Customers</del>	<del>2,276.2</del> <del>6</del>	<del>57.66</del>	<del>2,158.0</del> <del>6</del>	<del>55.18</del>	<del>2,267.4</del> <del>8</del>	<del>55.98</del>	<del>2,491.1</del> <del>5</del>	<del>54.25</del>		
		Our revenue fr	et forth b	elow.				-	·	<del>(in ₹ million,</del>	<b>,</b>	
		Top To Customers (1)	en Fisco	<del>d 2019</del>	Fiscal 2	2020	Fiscal 20		Vine Month December 3			
		4		654.64		619.27		580.70		622.00		
		<del>2</del> <del>3</del>		180.17 228.75		<del>275.26</del> <del>235.21</del>		<del>325.06</del>		<del>- 621.46</del> <del>- 161.30</del>		
		4		209.71		194.97		185.35		153.97		
		5	<del> </del>	95.07		<del>111.19</del>		<del>163.31</del>		339.85		
		6		<del>240.16</del>		<del>153.78</del>		221.32		80.64		
		7		<del>129.03</del>		121.57		113.07		181.81		





S. No.	Observations			Respons	se		UDRHP Page Number (clean)	UDRHP Page Number (redline)
		8	<del></del>	<del>48.92</del>	<del>66.75</del>	<del></del>		
		9	48.81	90.01	98.06	<del>151.30</del>		
		<del>10</del>	<del></del>	98.10	<del>85.11</del>	<del>78.41</del>		
		<del>Total</del>	<del>1,992.38</del>	<del>1,948.28</del>	2,022.01	<del>2,491.15</del>		
		basis for Fiscal 2	2019, Fiscal 2020, Fiscal i	2 <mark>021 and the nine mo</mark> r	nths ended December 3	ribution on a restated consoli 1, 2021. Isis for the years and po		
		indicated are set		<del>петs он а ргојотн</del>	<del>na consonautea bu</del>	in ₹ millio		
		<del>Top Ten</del> <del>Customers (1)</del>	Fiscal 2019	Fiscal 2020	Fiscal 2021	Nine Months ended December 31, 2021		
		<del>1</del>	<del>654.64</del>	<del>619.27</del>	<del>580.70</del>	<del>622.00</del>		
		2	<del>220.13</del>	<del>297.43</del>	<del>221.19</del>	<del>621.46</del>		
		3	<del>228.75</del>	<del>235.21</del>	<del>325.06</del>	<del>161.30</del>		
		4	<del>269.46</del>	<del>241.72</del>	240.74	<del>153.97</del>		
		5	300.08	<del>197.80</del>	282.58	<del>80.64</del>		
		6	<del>132.42</del>	149.92	<del>187.31</del>	339.85		
		7	<del></del>	<del>157.82</del>	<del>159.74</del>	<del></del>		
		8	<del>227.51</del>	66.48	<del>81.42</del>	<del>100.41</del>		
		9	48.81	90.01	98.06	<del></del>		
		<del>10</del>	20.45	<del>102.40</del>	90.67	<del>78.41</del>		
		<del>Total</del>	<del>2,276.26</del>	<del>2,158.06</del>	<del>2,267.48</del>	<del>2,491.15</del>		
		(1) The top ten cu	ustomers provided are	our top ten custom	ners in terms of rever	nue contribution on a prof	orma	





S. No.	Observations		Response								UDRHP Page Number (clean)	UDRHP Page Number (redline)
		<del>consolid</del>	<del>ated basis f</del>	<del>for Fiscal 2019</del>	<del>), Fiscal 202(</del>	<del>), Fiscal 2021</del>	and the nin	<del>e months en</del>	<del>ded Decemb</del>	e <del>r 31, 2021.</del>		
		consolidate	the table set forth below provides our revenue from operations from our CDMO business on a restated consolidated basis from our top ten customers and such revenue as a percentage of our operations or the years and period indicated.									
		Revenue from										
		Operatio ns	₹ million	% of revenue from operatio ns from CDMO business	₹ million	% of revenue from operatio ns from CDMO business	₹ million	% of revenue from operatio ns from CDMO business	₹ million	% of revenue from operatio ns from CDMO business		
		Top Ten Custome rs	2,022.01	54.52%	3,341.18	48.66%	3,825.40	56.29%	1,136.63	68.39%		
		The table so forma cons	olidated b for Fiscal 2	pasis from c 2023.	our top ten	customers	-					
		Revenue fr	om Operati	ions	Fiscal 2023	3		% of revi	enue from	onerations		
					₹ million			from CDM	•			
		Top Ten Cus	stomers				3,825.40			56.29%		
		indicated a	ur revenue from our top 10 customers on a restated consolidated basis for the fiscal years and period dicated are set forth below.  (in ₹ million)									
		Top Ten Cu (1)		Fiscal 202	21	Fiscal 2022	Fi	scal 2023	е	Months nded 30, 2023		





S. No.	Observations		UDRHP Page Number (clean)	UDRHP Page Number (redline)				
		1	580.70	770.71	1,001.04	365.37		
		2	325.06	728.07	643.80	186.45		
		3	221.32	462.24	467.66	162.08		
		4	185.35	283.24	412.20	100.12		
		5	183.27	240.25	238.98	70.14		
		6	163.31	229.56	229.05	68.83		
		7	113.07	198.67	225.20	60.82		
		8	98.06	154.73	221.50	44.60		
		9	85.11	144.48	198.15	39.51		
		10	66.76	129.23	187.82	38.71		
		Total	2,022.01	3,341.18	3,825.40	1,136.63		
		below.	en Customers (1)		Fiscal 2023	(in ₹ million)		
		1	en Customers (1)		FISCUI 2025	1,001.04		
		2				643.80		
		3				467.66		
		4				412.20		
		5				238.98		
		6				229.05		
		7				225.20		
		8				221.50		
		9				198.15		
		10				187.82		
		Total				3,825.40		
		consolidated ba	stomers provided are our to sis for Fiscal 2023. arious long-term agree					





S. No.	Observations	Response	UDRHP Page Number (clean)	UDRHP Page Number (redline)
		change, sometimes significantly based on the expected forecast volume required by our customers. In addition, certain of our agreements may be terminated by the customer without notice. While, in the recent past, none of our agreements have been terminated without notice there can be no assurance that such instances will not occur in future. In addition, the amount of customer spending on pharmaceutical development and manufacturing, particularly the amount our customers choose to spend on outsourcing CDMO services and products, has a large impact on our sales and profitability. Consolidation in the pharmaceutical industry may also impact such spending as customers integrate acquired operations, including research and development departments and manufacturing operations. Any reduction in customer spending on outsourcing CDMO services and products as a result of these and other factors could have a material adverse effect on our business, results of operations and financial condition."	27	
13.	LM is advised to include RF 5 and 11 sequentially.  Risk Factor 14: LM is advised to redraft the heading to reflect the risk explained in the third paragraph.	Noted for compliance. We have included risk factor 5 and risk factor 11 sequentially in the UDRHP.  Noted for compliance. We have amended the heading of the risk factor 14 in the UDRHP as follows:  "Reforms in the healthcare industry and the uncertainty associated with pharmaceutical pricing, reimbursement and related matters could adversely affect the pricing and demand for our products as well as the consumer demand for the products we manufacture for our customers, which may significantly influence our business, results of operations and financial condition. Further, our business and results of operations may be adversely impacted due to the price ceiling imposed by the Government".	37 44	46 55
15.	Risk Factor 16: LM is advised to include this RF as top 15.	Noted for compliance. We have included risk factor 16 in the top 15 risk factors in the UDRHP.	43	54
16.	Risk factor 17: LM is advised to include this RF as top 10 and modify the heading as: 'Our business is capital intensive. Any insufficient cash flows from our operations or inability to borrow to meet our working capital requirements may materially and adversely affect our business and results of operations'.	Noted for compliance. We have included risk factor 17 in the top 10 risk factors in the UDRHP. In addition, we have amended the heading of the risk factor 17 in the UDRHP as follows:  "Our business is working capital intensive. If we experience Any insufficient cash flows from our operations or inability are unable to borrow to meet our working capital requirements, it may materially and adversely affect our business and results of operations."	42	53
17.	Risk Factor 19: LM is advised to delete 'which are routine in nature.'	Noted for compliance. We have amended risk factor 19 (as already modified pursuant to previous SEBI comments) in the UDRHP as follows:	49	67





S. No.	Observations	Response	UDRHP Page Number (clean)	UDRHP Page Number (redline)
_	Observations	"Our proposed capacity expansion plans relating to our manufacturing facilities are subject to the risk of unanticipated delays in implementation and cost overruns.  We have made and intend to continue making investments to expand the capacity of our manufacturing facilities to aid our growth efforts. We intend to construct a new 240,916 sq. ft facility in Jammu ("Jammu Facility"), which may be utilised for manufacturing tablets, capsules, dry syrups and injections. The estimated total project cost for this new Jammu Facility is ₹3,551.72 million, as certified by Ravinder K. Sharma & Co. Chartered Accountants.  Further, expansion of manufacturing facilities requires governmental, statutory and other regulatory approvals, licenses, permits and registrations to be obtained from various authorities and we cannot assure you that we will be able to obtain or renew such approvals, licenses, permits and registrations in a timely manner, or at all. If we fail to obtain or renew such licenses, approvals, registrations and permits in a timely manner, our commissioning date for the expansion plans may be delayed, which could adversely affect our business and results of operations.  Construction of our new Jammu Facility will be subject to the potential problems and uncertainties that construction projects face including cost overruns or delays. In addition, construction and operation of our new Jammu Facility will require us to obtain various approvals. There can be no assurance that we will be able to obtain these registrations and approvals in a timely manner or at all.  As on November 15, 2023, we have made the following progress on construction of our new Jammu Facility:  Land has been acquired and possession has been taken;		Number
		<ul> <li>Construction is ongoing;</li> <li>Orders for plant and machinery are ongoing;</li> <li>Construction contracts are being finalized;</li> <li>Purchase orders for plant, equipment and other fixed assets, both imported and indigenous,</li> </ul>		





S. No.	Observations	Response	UDRHP Page Number (clean)	UDRHP Page Number (redline)
		amounting to ₹2,946.97 million have been placed;  • An amount of ₹2,498.65 million has already been incurred on the project out of which ₹1,061.48 million has funded by through our internal accruals and the remaining ₹1,437.16 million has been disbursed by HDFC bank / State Bank of India;  • Out of the purchase orders placed for imported machinery and equipment, 4 sets of blow fill seal machines having invoice value of CHF 13.50 million (₹1,302.87 million) have been received at the facility;  • Acknowledgment of our intent to establish a manufacturing enterprise has been received from the office of the General Manager of District Industries Centre, Kathua;  • GST registration has been received; and  • Consent to Establish received from the Jammu and Kashmir State Pollution Control Board."  Construction of our new Jammu Facility will be subject to the potential problems and uncertainties that construction projects face including cost overruns or delays. In addition, construction and operation of our new Jammu Facility will require us to obtain various approvals, which are routine in nature. There can be no assurance that we will be able to obtain these registrations and approvals in a timely manner or at all.  As on August 31, 2022 we have made the following progress on construction of our new Jammu facility:  • Land has been acquired and possession taken;  • Orders for plant and machinery are ongoing;  • Construction contracts are being finalized;  • Term loan for the Jammu Facility has been sanctioned;  • Acknowledgment of our intent to establish a manufacturing enterprise has been received from the office of the General Manager of District Industries Centre; and  • Application has been made for consent to establish with the Pollution Control Committee."		





S. No.	Observations		Response		UDRHP Page Number (clean)	UDRHP Page Number (redline)				
18.	Risk factor 21: LM is advised to include it under top 15 risk factors.	Noted for compliance. We have	oted for compliance. We have included risk factor 21 in the top 15 risk factors in the UDRHP.							
19.	Risk Factor 22: LM is advised to include it under top 10 risk factors.	Noted for compliance. We have	included risk factor 22 in the t	op 10 risk factors in the UDRHP.	40	52				
20.	Risk Factor 29: LM is advised to redraft heading stating that export is from sale of goods and import for raw materials.	"We face foreign exchange risk	ted for compliance. We have amended the heading of the risk factor 29 in the UDRHP as follows:  We face foreign exchange risks that could adversely affect our results of operations as a portion of a revenue is from exports and a portion of our expenditure is from imports for raw material, both							
21.	Risk Factor 34: LM is advised to include it under top 10 risk factors.	Noted for compliance. We have		op 10 risk factors in the UDRHP.	40	52				
22.	Risk Factor 44: LM is advised to disclose amount of outstanding debt and date of availing the same.	"Any downgrade of our de  As of October 31, 2023, we had consolidated basis. For details,	oted for compliance. We have amended risk factor 44 (as already modified pursuant to previous EBI comments) in the UDRHP as follows:  "Any downgrade of our debt ratings could adversely affect our business.  so of October 31, 2023, we had total outstanding borrowings of ₹4,811.91 million on a restated pursuant business. For details, see "Financial Indebtedness" beginning on page []. As per the credit atting letter dated September 27, 2023, we have received the following credit ratings on our debt and dedit facilities.							
		Instrument or Rating Type	Rating Agency	Ratings						
		Long Term Bank Facilities	CARE Ratings Limited	A- (Negative)						
		Short Term Bank Facilities	CARE Ratings Limited	A2+						
		ability to meet financial comm three years due to the non-sub- required data could again resu assurance that these ratings wil	nese ratings assess our overall financial capacity to pay our obligations and are reflective of our polity to meet financial commitments as they become due. Our credit ratings were suspended for ree years due to the non-submission of data to the rating agencies and any such failure to provide quired data could again result in the suspension of our credit ratings. Further, there can be no surance that these ratings will not be revised or changed by the above rating agencies due to various ctors. Any downgrade in our credit ratings may increase interest rates for refinancing our							





S. No.	Observations	Response	UDRHP Page Number (clean)	UDRHP Page Number (redline)
		outstanding debt, which would increase our financing costs, and adversely affect our future issuances of debt and our ability to raise new capital on a competitive basis."		
23.	Risk Factor 57: LM is advised to remove future reference in the heading.	Noted for compliance. We have amended the heading of the risk factor 57 in the UDRHP as follows:  "Information relating to the installed manufacturing capacity of our two manufacturing facilities included in this Draft-Red Herring Prospectus are based on various assumptions and estimates and future production and capacity may vary."	64	91
24.	Risk Factor 58: LM is advised to redraft heading of the risk factor.	Noted for compliance. We have amended the heading of the risk factor 58 in the UDRHP as follows:  "Certain sections of this Draft Red Herring Prospectus contain information from the CRISIL Report which we commissioned and purchased and any reliance on such information for making an investment decision in the Offer is subject to inherent risks. We commissioned and purchased the CRISIL Report. This Draft Red Herring Prospectus contains information from the CRISIL Report and such information is subject to inherent risks and limitations."	65	92
25.	LM is advised to include a separate risk factor in top 10 that more than 90% of the raw materials are imported from one country (i.e., China).	Noted for compliance. We note that new risk factor 9 (as previously provided to SEBI in earlier responses to SEBI Observations) provides a discussion of the risk related to the Company's dependence on China. New risk factor 9 (as previously provided to SEBI in earlier responses to SEBI Observations) has been added to the UDRHP as follows:  "Our dependence on China, China SEZ and Hong Kong for our raw material supplies exposes us to political, economic and social conditions in greater China.  We are dependent on the import raw materials from China, China SEZ and Hong Kong.  The table set forth below provides our imported raw materials from China, China SEZ and Hong Kong on a restated consolidated basis as a percentage of our cost of imported raw materials and as a percentage of total raw materials purchases for the fiscal years/period indicated.	41	53





S. No.	Observations	Response				UDRHP Page Number (clean)	UDRHP Page Number (redline)			
		Restated Consolidated	For the three months ended June							
			2021 2	2022 2023	30, 2023					
		Imported raw materials from China, China SEZ and Hong Kong as a percentage of our cost of imported raw materials	91.85%90	0.03% 75.41%	% 100%					
		Imported raw materials from China, China SEZ and Hong Kong as a percentage of total raw material	173 17% 17 7 7 8 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1							
		political, economic and social conditions in greater China. Further adversely impacted by the economic downturn in their national catheir banking and financial systems, economic weakness, unfavoinflation, lowering of spending power and customer confidence, a	Our dependence on China, China SEZ and Hong Kong for our raw material supplies exposes us to political, economic and social conditions in greater China. Further, our raw material suppliers may be adversely impacted by the economic downturn in their national or regional economies, disruption in their banking and financial systems, economic weakness, unfavourable government policies, rising inflation, lowering of spending power and customer confidence, and political uncertainty."							
26.	LM is advised to include separate risk factor in top 15 regarding that nearly 30% of revenue is from related parties.	Noted for compliance. We note that a revised risk factor 15 (as p responses to SEBI Observations) provides a discussion that nearl from related parties. Revised risk factor 15 (as previously provided Observations) has been added to UDRHP as follows:	previously rly 30% of t	provided to the Compa	SEBI in earlier ny's revenue is	44	56			
		"We have in the past entered into related party transactions and which may potentially involve conflicts of interest with the equi in the three months ended June 30, 2023, we derived 10.72% and from operations on a restated consolidated basis from related p								
		We have in the course of our business entered into, and will continu with our related parties including Key Managerial Personnel. The transactions as a percentage of revenue from operations for the p								
			r the year ei arch 31, 202							





S. No.	Observations		Response								
							June 30, 2023				
		Related part transactions as percentage revenue from operations	a of 29	9.05%	27.57%	10.72%	8.88%				
		UML (which is no finished pharmac as related party respectively, in Fis as per Ind AS 24,	eutical products transactions. Re scal 2021, Fiscal .								
		Nature of Transaction	Name of the related party	For the year ended March 31, 2021	For the y ended March : 2022	d ended 31, March 31,	months ended				
		Revenue from operations (net of returns)	Univentis Medicare Limited	16.42%	13	3.07% 11.94	13.19%				
		accordance with a we could not have with related parts of the effect on our finant to enter into relate enter into post-lic Companies Act a shareholders and arrangements in individually or in	nile we believe that all such related party transactions are conducted on an arms' length basis in cordance with the Companies Act and other applicable regulations, there can be no assurance that could not have achieved more favourable terms if such transactions had not been entered into the related parties or that such transactions, individually or in aggregate, will not have an adverse exect on our financial condition and results of operations. Furthermore, it is likely that we will continue enter into related party transactions in the future. All such related party transactions that we may see into post-listing, will be subject to board or shareholder approval, as necessary under the impanies Act and the SEBI Listing Regulations, in the interest of our Company and our minority areholders and in compliance with the SEBI Listing Regulations, we cannot assure you that these angements in the future, or any future related party transactions that we may enter into, lividually or in the aggregate, will not have an adverse effect on our business, financial condition, ults of operations, cash flows and prospects."								





S. No.	Observations	Response	UDRHP Page Number (clean)	UDRHP Page Number (redline)
27.	LM is advised to include separate risk factor in case there is any delay in expansion of existing plants.	Noted and complied with. We note that risk factor 19 provides a discussion of the risk in delay in expansion of existing plants. Risk factor 19 (as previously modified by SEBI comments) reads as follows:  "Our proposed capacity expansion plans relating to our manufacturing facilities are subject to the risk of unanticipated delays in implementation and cost overruns.  We have made and intend to continue making investments to expand the capacity of our manufacturing facilities to aid our growth efforts. We intend to construct a new 240,916 sq. ft facility in Jammu ("Jammu Facility"), which may be utilised for manufacturing tablets, capsules, dry syrups and injections. The estimated total project cost for this new Jammu Facility is ₹3,551.72 million, as certified by Ravinder K. Sharma & Co. Chartered Accountants.  Further, expansion of manufacturing facilities requires governmental, statutory and other regulatory approvals, licenses, permits and registrations to be obtained from various authorities and we cannot assure you that we will be able to obtain or renew such approvals, licenses, permits and registrations in a timely manner, or at all. If we fail to obtain or renew such licenses, approvals, registrations and permits in a timely manner, our commissioning date for the expansion plans may be delayed, which could adversely affect our business and results of operations.  Construction of our new Jammu Facility will be subject to the potential problems and uncertainties that construction projects face including cost overruns or delays. In addition, construction and operation of our new Jammu Facility will require us to obtain various approvals. There can be no assurance that we will be able to obtain these registrations and approvals in a timely manner or at all.  As on November 15, 2023, we have made the following progress on construction of our new Jammu Facility:  Land has been acquired and possession has been taken;  Construction is ongoing:	49	67





S. No.	Observations	Response	UDRHP Page Number (clean)	UDRHP Page Number (redline)
		<ul> <li>Orders for plant and machinery are ongoing;</li> <li>Construction contracts are being finalized;</li> <li>Purchase orders for plant, equipment and other fixed assets, both imported and indigenous, amounting to ₹2,946.97 million have been placed;</li> <li>An amount of ₹2,498.65 million has already been incurred on the project out of which ₹1,061.48 million has funded by through our internal accruals and the remaining ₹1,437.16 million has been disbursed by HDFC bank / State Bank of India;</li> <li>Out of the purchase orders placed for imported machinery and equipment, 4 sets of blow fill seal machines having invoice value of CHF 13.50 million (₹1,302.87 million) have been received at the facility;</li> <li>Acknowledgment of our intent to establish a manufacturing enterprise has been received from the office of the General Manager of District Industries Centre, Kathua;</li> <li>GST registration has been received; and</li> <li>Consent to Establish received from the Jammu and Kashmir State Pollution Control Board."</li> <li>Construction of our new Jammu Facility will be subject to the potential problems and uncertainties that construction projects face including cost overruns or delays. In addition, construction and operation of our new Jammu Facility will require us to obtain various approvals, which are routine in nature. There can be no assurance that we will be able to obtain these registrations and approvals in a timely manner or at all.</li> <li>As on August 31, 2022 we have made the following progress on construction of our new Jammu Facility:</li> </ul>		
		<ul> <li>Land has been acquired and possession taken;</li> <li>Orders for plant and machinery are ongoing;</li> <li>Construction contracts are being finalized;</li> <li>Term loan for the Jammu Facility has been sanctioned;</li> <li>Acknowledgment of our intent to establish a manufacturing enterprise has been received from the office of the General Manager of District Industries Centre; and</li> <li>Application has been made for consent to establish with the Pollution Control Committee."</li> </ul>		





S. No.	Observations		Respo	onse		UDRHP Page Number (clean)	UDRHP Page Number (redline)		
28.	Capital Structure: LM is advised to include a table showing pre-issue number of shares held and percentage holding for promoter, promoter group and selling shareholder.	The pre-Offer "Capital Structhe Company" Further, the promoter Gromanner:  "(vi) Equity Some Sequity Shares Promoter Grown Pro	"(vi) Equity Shareholding of the Promoter Group  As on the date of this Draft Red Herring Prospectus, except for Vandana Lohariwala who holds 4,000 Equity Shares and Chhavi Lohariwala who holds 4,000 Equity Shares, none of the members of our Promoter Group hold any Equity Shares in our Company. As on the date of this Red Herring Prospectus, except as disclosed below, and other than our Promoters, none of the members of our Promoter Group						
		S. No.	Name of the Shareholder	Pre-Offer Equity No. of Equity Shares	Share Capital % of total Shareholding				
		1. 2.	Vandana Lohariwala Chhavi Lohariwala						
		Details of the	nareholding of the Selling Shareholders shareholding of the Selling Shareholders are as set out below:	ed					





S. No.	Observations			UDRHP Page Number (clean)	UDRHP Page Number (redline)						
				Name of the Cl	l l . l		e-Offer Equity S				
			S. No.	Name of the SI	narenolaer		of Equity hares	% of total Shareholding			
			1.	Manoj Kumar Lohariv	vala		19,036,000	38.0	01		
			2.	Vinay Kumar Lohariw	ala		14,436,000	28.8	32		
			<i>3</i> .	Gian Parkash Aggarw	al		14,512,000	28.9	<del>8</del>		
			Total 47,984,000 95.81								
		[]"									
29.	Page 94: LM is advised to remove legal counsel to the BRLMs as to Indian Law.	Note	d for con	npliance.						-	-
30.	LM is advised to ensure compliance with respect to provisions of independent directors as per SEBI LODR regulations.	Comp	olied wit	n.		-	-				
31.	Page 104: LM is advised to disclose reason for reduction in shareholding from 39.98% to 30.23% of Gian Parkash Aggarwal or alternatively, provide a table as given in 7(iii) for promoters on page 105.	the U	IDRHP:	npliance. We have u  f the Selling Shareh				'Capital Struct	ure' section of	98	134
		in the the b	he build-up of the shareholding of our Promoter Selling Shareholders since the incorporation of out ompany is set out in "- Details of shareholding of our Promoters and members of the Promoter Group of the Company - Build-up of the Promoters' shareholding in our Company" above on page [•]. Further the build-up of the shareholding of Gian Parkash Aggarwal since the incorporation of our Company is est out below:								
		allo tra tran	Date of allotment/       Nature of transaction       No. of equity       Face value per equity       Issue price/ transfer price       Percentage of pre-Offer       Percentage of pre-Offer         transfer / transmission       shares       share (₹)       per equity per equity equity share capital (%)       capital (%)								
		June	16, 2009	Transfer from Vasanthi Muppidathy Thevar	2,500	100	20.00	0.05	[•]		





S. No.	Observations				UDRHP Page Number (clean)	UDRHP Page Number (redline)				
			Transfer from Muppidathy Sivan Thevar	19,900	100	20.00	0.40	[•]		
		January 10, 2011	Further Allotment	97,600	100	100.00	1.95	[•]		
		June 15, 2011	Further Allotment	76,000	100	100.00	1.52	[•]		
		December 26, 2011	Further Allotment	196,000	100	100.00	3.91	[•]		
		March 17, 2012	Further Allotment	88,000	100	100.00	1.76	[•]		
		May 31, 2018	Transfer to Archit Aggarwal	(100)	100	-	(0.00)	[•]		
			Transfer to Veena Devi	(100)	100	-	(0.00)	[•]		
		January 18, 2022	Transfer to Vinay Kumar Lohariwala	(117,000)	100	1,666.67	(2.34)	[•]		
		each were sub-d	solution passed by our Si livided into equity shares ne split of equity shares w	of face value ₹10	each. Pursuan	t to the corporate a				
		April 27, 2022	Bonus issue	10,884,000	10	-	21.73	[•]		
		Total				14,512,000	28.98	[•]		
32.	Page 116/117: LM is advised to remove future projections.	key assumption requirements	ease note that the projected working capital requirements for Fiscal 2023 and 2024 along with the y assumptions were disclosed on page 116 and 117 of the DRHP to comply with the applicable quirements under Schedule VI of the SEBI ICDR Regulations. These disclosures shall be further dated to reflect the projected working capital requirements for Fiscals 2024 and 2025 in the UDRHP.							150
33.	Page 171: LM is advised to disclose break up of revenue for CDMO business and other business.		te for compliance. We have amended the "Overview" subsection of the section "Our Business" on ge 171 of the DRHP to add the break up of CDMO and other businesses in the UDRHP as follows:							268
			forth below sets for operations on a rest ted.							
		Re	estated Consolidated	Financial Inform	nation					





S. No.	Observations		Response							UDRHP Page Number (clean)	UDRHP Page Number (redline)	
		Business Area	Fisca	2021	Fisca	1 2022	Fisca	l 2023		nths ended 80, 2023		
		Alex	₹ million	% of revenue from operations	₹ million	% of revenue from operations	₹ million	% of revenue from operations	₹ million	% of revenue from operatio ns		
		CDMO services and products	3,708.71	90.31%	6,866.94	85.78%	6,795.56	73.36%	1,662.10	71.26%		
		Domestic branded generics	0.00	0.00%	370.51	4.63%	1,661.61	17.94%	422.53	18.12%		
		Internati onal branded generics	397.91	9.69%	767.81	9.59%	806.63	8.71%	247.80	10.62%		
		Sharon	0.00	0.00%	0.00	0.00%	0.00	0.00%	0.00	0.00%		
		Revenue from Operatio ns	4,106.62	100.00%	8,005.26	100.00%	9,263.80	100.00%	2,332.43	100.00%		
34.	Page 207: LM is advised to disclose brief financial highlights for past 3 years and also include nature of activities carried out for all subsidiaries.	Univentis F	of busines oundation, nancial higl	ss that the are engage nlights of UI	Company's	en disclose	d on page 2	207 of the D	RHP.	" <b>UML</b> ") and the UDRHP,	223	295





S. No.	Observations		Response			UDRHP Page Number (clean)	UDRHP Page Number (redline)			
		"1. Univentis Medicare Limited ("	I. Univentis Medicare Limited ("UML")							
			] Brief financial highlights of UML for the financial years ended March 31, 2023, March 31, 202 and March 31, 2021 are set out below. (₹ in milli							
		Particulars Particulars	Fiscal 2023	Fiscal 2022	Fiscal 2021					
		Reserves (excluding revaluation reserve)         524.12         418.94         298.84           Sales         1783.32         1687.77         1579.49								
		Sales	1579.49							
		Profit after tax								
		Earnings per share								
		Diluted earnings per share	690.78	798.20	859.48					
		Net asset value	524.12	420.44	300.34					
		[]"								
		"2. Univentis Foundation  [] Brief financial highlights of the and March 31, 2022 are set out be		ne financial years en	ded March 31, 2023 (in ₹)					
		Particulars	Fiscal 2023	Fi	scal 2022					
		Reserves (excluding revaluation reserve)	68	87,390	3,518					
		Sales		NA	NA					
		Profit after tax	<u> </u>	NA	NA					
		Earnings per share								
		Diluted earnings per share								
		Net asset value								
		[]"								





S. No.	Observations		Response			UDRHP Page Number (clean)	UDRHP Page Number (redline)
		Subsequent to the filing of the ("Sharon"). Sharon is a wholly on Company. Appropriate disclosures Corporate Matters" section of the engaged in by Sharon and its brief  "3. Sharon Bio-Medicine Limited ("" [] Sharon is engaged in the busing ingredients as well as finished dose [] Brief financial highlights of Sharon March 31, 2021 are set out be	wned Subsidiary of UMI is in this regard have be UDRHP. The relevant distinancial highlights are resonant of manufacturing of ages.	L, and accordingly, is a seen included in the "Haclosures regarding the eproduced below for your fintermediates and actions."	Subsidiary of the istory and Certain nature of activities our reference:  we pharmaceutical  3, March 31, 2022,		
		Particulars Particulars	Fiscal 2023	Fiscal 2022	(₹ in million) <b>Fiscal 2021</b>		
		Reserves (excluding revaluation reserve)	(6,418.27)	(6,515.37)	(6,707.85)		
		Sales	1,922.16	1,883.91	1,728.62		
		Profit after tax	93.32	176.75	140.47		
		Earnings per share	16.21	30.71	24.40		
		Diluted earnings per share	16.21	30.71	24.40		
		Net asset value	(6,000.18)	(6,092.20)	(6,254.69)		
		[]"					
35.	Page 237: LM is advised to provide details of	Noted for compliance.				256	330
	promoters, brief financials for past 3 years and	·					
	nature of activities undertaken by all group	We have updated the section titled	d "Group Companies" in	the following manner ir	the UDRHP:		
	companies.	"1. Nugenic Pharma Private Limito	ed ("Nugenic Pharma")				
		The registered office of Nugenic Ph Baddi, Solan 173 205, Himachal Pi			•		





S. No.	Observations		Response			UDRHP Page Number (clean)	UDRHP Page Number (redline)
		Lohariwala, Manoj Kumar Lo Lohariwala and Bindu Devi Loh manufacturing of packaging ma Brief financial highlights of Nuge Fiscals 2023, 2022 and 2021 are	nariwala. Nugenic Pharr terials. nic Pharma based on the	na is currently engage	ed in the business of		
		Particulars	Fiscal 2023	Fiscal 2022	Fiscal 2021		
		Reserves (excluding revaluation reserve)	165.13	157.05	155.59		
		Total Revenue	807.93	639.47	605.12		
		Profit /(Loss) for the year	8.25	1.46	8.90		
		Earnings per share	4.41	0.78	4.75		
		Diluted earnings per share	4.41	0.78	4.75		
		Net asset value	183.86	175.77	174.32		
		Nugenic Pharma Private Limiter statements for Fiscals 2021, 202 Company at www.innovacaptab  2. Azine Healthcare Private Lim  The registered office of Azine He BSNL Tower, Bavla, Ahmedabad Lohariwala, Shyamsunder Agravengaged in the business of manual	20 and 2019 This information.com/investor-relations. ited ("Azine Healthcare" althcare Private Limited 382 200, Gujarat, India. In val and Shyamsunder Lo	ation is also available o ) is situated at Plot No. 4 The promoters of Azine hariwala HUF. Azine H	on the website of our  101, Kerela GIDC, Opp  Healthcare are Rekha		
		Brief financial highlights of Azine Fiscals 2023, 2022 and 2021 are Particulars	set out below:	e audited standalone fir	nancial statements for (₹ in million) <b>Fiscal 2021</b>		
		Reserves (excluding revaluation	Fiscal 2023 45.63	44.80	43.52		
		reserve)	45.05	44.80	45.52		





S. No.	Observations		Response			UDRHP Page Number (clean)	UDRHP Page Number (redline)
		Total Revenue	268.18	367.03	274.41		
		Profit /(Loss) for the year	0.83	1.28	0.94		
		Earnings per share	3.45	5.27	3.90		
		Diluted earnings per share	3.45	5.27	3.90		
		Net asset value	48.05	47.22	45.94		
		Azine Healthcare Private Limited statements for Fiscals 2021, 2020. Company at www.innovacaptable.  3. DMS Electronics Private Limit.  The registered office of DMS Electronics Private Limit.  The registered office of DMS Electronics Private Limit.  The registered office of DMS Electronics Private Limit.  Rangarao Colony, Dattagally, Mare Gian Parkash Aggarwal, Vij. Rakesh Kumar Aggarwal and Mof manufacturing of printed circular private private Limit.  Brief financial highlights of DMS Fiscals 2023, 2022 and 2021 are	20 and 2019 This information.com/investor-relations.  Ited ("DMS Electronics")  Electronics Private Limited Mysore 570 022, Karnatako Mysore 570 029, Karnatako Mysore 570 020, Karnatako Mysore 570 020	tion is also available on I is situated at 11, A1 a, India. The promoters kit Aggarwal, Lalit Jinda tronics is currently enga	the website of our  Block, Vasu Layout of DMS Electronics II, Archit Aggarwal, ged in the business		
		Particulars Particulars	Fiscal 2023	Fiscal 2022	Fiscal 2021		
		Reserves (excluding revaluation reserve)	(28.50)	(23.91)	0.06		
		Total Revenue	4.93	221.08	151.75		
		Profit /(Loss) for the year	(4.59)	(24.06)	4.27		
		Earnings per share	(1.83)	(10.32)	4.27		
		Diluted earnings per share	(1.83)	(10.32)	4.27		
		Net asset value	(3.50)	1.09	10.06		
		DMS Electronics Private Limited statements for Fiscals 2021, 20 Company at www.innovacaptab	<del>20 and 2019</del> This informa				





S. No.	Observations	Response	UDRHP Page Number (clean)	UDRHP Page Number (redline)
36.	LM is advised to ensure that the main / sub-headings in the UDRHP / RHP do not have any abbreviations.	Complied with and noted for compliance.	-	-
37.	LM is advised that reference to name of any place mentioned in the offer document may be followed by name of City I State, as the case may be.	Complied with and noted for compliance.	-	-
38.	LM is advised to include updated audited financials at the time of filing UDRHP/RHP.	Noted for compliance to the extent applicable. We shall include the updated Restated Consolidated Financial Information, as of, and for the three months ended June 30, 2023, and as of, and for the years ended, March 31, 2023, 2022 and 2021 in the UDRHP, the RHP and the Prospectus.	260	334
39.	Page 413: LM is advised to disclose risk and the impact of recall of products under para (b).	Noted for compliance. We undertake to further amend Risk Factor 10 (as already modified pursuant to previous SEBI comments) as follows:  "Failure to comply with the quality requirements and technical specifications prescribed by our customers may lead to loss of business from such customers and could negatively impact our business, results of operations and financial condition, including cancellation of existing and future orders which may expose us to warranty claims.  Our products and manufacturing processes are subject to stringent quality standards and specifications, typically specified by our CDMO customers in their respective agreements, and any deviations from the required specifications by our Company or failure to comply with the technical specifications of our customers regarding the composition of drugs or any alterations in manufacturing process or method or raw material, may lead to a recall of products or cancellation of the orders placed by our customers or may require prior intimation or consent from the customer.  Some agreements also require us to furnish quality assurance and compliance certificates to the customers certifying that the quality of the products is as per the agreed specifications. As per the terms and conditions of the respective agreements, our customers have the right to reject the products in case of, inter alia, manufacturing defects, and discrepancy with respect to prescribed specifications, and we are responsible to replace such products free of any additional cost within a stipulated timeframe along with indemnity to the customer for losses arising from breach of obligations,	39	48





S. No.	Observations	Response	UDRHP Page Number (clean)	UDRHP Page Number (redline)
		specification of raw material used and manufacturing defect.  During Fiscal 2019, Fiscal 2020 and Fiscal 2021 and during the nine months ended December 31, 2021, we have received certain complaints from our CDMO customers. The nature of these complaints was mainly related to packaging material of the products. There have not been any financial implications related to these complaints. We undertake corrective and preventive actions on these complaints on a regular basis.		
		While we believe we undertake the necessary measures and engage internal and external experts to ensure that our facilities comply with the applicable standards as imposed by our customers, any failure on our part to maintain the applicable standards and manufacture products according to prescribed specifications, may lead to cancellation of the order, loss of customers, loss of reputation and goodwill of our Company. Additionally, it could expose us to indemnity, warranty claims, monetary liability and/or litigation. Our CDMO customers are typically provided the right to audit our manufacturing facilities, processes or systems, under such agreements, after providing a certain period of notice. While we have not received any adverse observations in the past from our customers pursuant to such audits, there can be no assurance that such audits would not result in any adverse observations in the future or that our customers will necessarily engage us for their outsourcing operations. The finished product delivered by us is further subject to laboratory validation by certain customers. Occurrence of any event on account of errors and omission could result in damage to our reputation and loss of customers, which could adversely affect our business, operations, our cash flows and financial condition.		
		In cases of recall of a product manufactured by us, our CDMO agreements typically require us to bear all the expenses and costs of such recall either upfront or by way of deduction from our bills, and the customers may also opt to terminate the agreement on account of such recall. During the period from April 1, 2018, to December 31, 2021, we had four cases of goods returned due to product recalls, and our liabilities on account of such product recalls is estimated at ₹0.05 million.  In addition, a complaint dated September 11, 2018 was filed by the Drugs Inspector, representing the state of Tamil Nadu, before the Court of Judicial Magistrate No. VII, Coimbatore, against our Company and four of our Directors, namely, Jayant Vasudeo Rao, Gian Parkash Agarwal, Manoj Kumar		





S. No.	Observations	Response	UDRHP Page Number (clean)	UDRHP Page Number (redline)
		Lohariwala and Vinay Kumar Lohariwala, for contravening the provisions of section 18 (a)(i) of the Drugs and Cosmetics Act, 1940 by manufacturing, selling and distributing the 'Not of Standard Quality' drug ventoxol expectorant, since the sample taken did not confirm to the label claim with respect to one of its contents. Our Company has, pursuant to such complaint, recalled the drug from the vendors and distribution channels. The matter is currently pending. The impact of this drug recall did not have a material effect on our business or results of operations.		
		Further, a complaint dated February 3, 2022 was filed by the Drugs Inspector, Srikakulum, representing the state of Andhra Pradesh, before the Court of Additional Judicial First Class Magistrate, Srikakulum District, against our Company and one of our Directors namely, Jayant Vasudeo Rao, for contravening the provisions of section 18 (a)(i) read with section 16 (1)(a) and the second schedule (1) of the Drugs and Cosmetics Act, 1940 by manufacturing, selling and distributing the 'Not of Standard Quality' drug pantaprazol sodium tablets with brand name Pantofresh - 40, since the sample taken had failed in the dissolution test as per the standards laid down by the Indian Pharmacopoeia Commission. Our Company has recalled the drug pursuant to such complaint. The matter is currently pending. The impact of this drug recall did not have a material effect on our business or results of operations."		
40.	LM is advised to assess and disclose the impact (if any), if the cases are not in favour of the Company/directors.	Complied with. A risk factor titled "There is pending litigation against our Company, Promoters, Subsidiaries and certain of our Directors. Any adverse decision in such proceedings may render us/them liable to liabilities/penalties and may adversely affect our business, results of operations and financial condition" has been disclosed in the "Risk Factors" section of the DRHP, on page 52 of the DRHP.  This risk factor has been suitably updated in the UDRHP to include details of the outstanding litigation required to be disclosed in terms of the SEBI ICDR Regulations as on the date of the UDRHP.	53	72
41.	With respect to all the complaints received by LM / Company / forwarded by SEBI, LM is advised to ensure that there is adequate redressal of the complaint and relevant disclosures of the same are made in the Red Hearing Prospectus and other Offer related material along with the disclosures of the financial impact of the same, if any.	Noted for compliance. Neither the BRLMs nor the Company have received any complaints, nor have any complaints been forwarded by SEBI, in connection with the DRHP or the Offer.	-	-





S. No.	Observations	Response	UDRHP Page Number (clean)	UDRHP Page Number (redline)
42.	LM is advised to ensure that the disclosure of details of all the criminal matters initiated by or against the company, group, directors, promoters, subsidiaries which are at FIR stage and no / some cognizance has been taken by court, is incorporated in the UDRHP / RHP along with appropriate risk factors in this regard.	Complied with and noted for compliance, to the extent applicable.  Details of all outstanding criminal proceedings involving the Company, Subsidiaries, Promoters and Directors, including proceedings at the FIR stage, if any, have been disclosed in the UDRHP, in accordance with the SEBI ICDR Regulations.  Further, in terms of the SEBI ICDR Regulations, only those proceedings involving the Group Companies which have a material impact on the Company are required to be disclosed in the UDRHP. Accordingly, details of all criminal proceedings involving the Group Companies which have a material impact on the Company, including proceedings at the FIR stage, if any, have been disclosed in the UDRHP, in accordance with the SEBI ICDR Regulations. Details in connection with any other criminal proceedings involving the Group Companies are not sought to be included.	424	460
43.	LM is advised to ensure following disclosures in the Issue advertisement for announcement of Price Band and all further advertisements as a box item below the price band:  "Risks to Investors:  i. The [to be disclosed] Merchant Bankers associated with the issue have handled [to be disclosed] public issues in the past three years out of which [to be disclosed] issues closed below the issue price on listing date."  ii. Any adverse data in the basis for issue price should be disclosed. For example:  • "The Price/Earnings ratio based on diluted EPS for [latest full financial year] for the issuer at the upper end of the Price band is as high as [to be disclosed] as	Noted for compliance.	-	-





S. No.	Observations	Response	UDRHP Page Number (clean)	UDRHP Page Number (redline)
	compared to the average industry peer group PE ratio of [to be disclosed]."  [if average industry peer group PE ratio is not available, then PIE of Nifty Fifty may be disclosed]  • "Average cost of acquisition of equity shares for the selling shareholders in /PO is [to be disclosed] and offer price at upper end of the price band is [to be disclosed].  • "Weighted Average Return on Net Worth for [last three full financial years] is [to be disclosed] %."  The data on above disclosures shall be updated and disclosed prominently (in the same font size as the price band) in advertisements of Price Band and all further advertisements, website of the company and the stock exchange. Further, any adverse ratio/ data in basis for issue price should be disclosed.			
44.	LM shall submit the draft advertisement for announcement of Price Band with SEBI before its publication in the newspapers for our comments, if any.	Noted for compliance.	-	-
45.	LM is advised to ensure compliance with the below email advisory sent to AIBI through email dated November 13 and November 15, 2021 and amendment to ICDR dated November 21,2022:-	Complied with and noted for compliance.	-	-





S. No.	Observations	Response	UDRHP Page Number (clean)	UDRHP Page Number (redline)
	a. LM shall ensure that all issuer companies filing offer document should provide — Price at which specified security was acquired in the last 3 years, by each of the promoters, promoter group, selling shareholders, shareholders entitled with right to nominate directors or any other rights. Following details may be disclosed for such transactions in tabular format — name of acquirer, date of acquisition, number of shares acquired and acquisition price per share.  b. The portion pertaining to "Risks to Investors" shall constitute at least 33% of the price band advertisement space.  c. The risks to investors shall include weighted average cost of acquisition of all shares transacted in last 3 years, 18 months and 1 year, from the date of RHP, in the following format:			
	Period Weighted Average (X' times the Acquisition Price: Acquisitio Average Lowest Price (In Rs.)  Last 1 year  Last 18 month S			





S. No.	Observations	Response	UDRHP Page Number (clean)	UDRHP Page Number (redline)
	Range of acquisition should show lowest price of acquisition excluding gift/bonus.  d. The font size for price band and "Risk to investors" should be increased to match the font of BID/Offer Programme.  e. Matters related to ASBA and UPI may be brought subsequent to Price Band, Risks to Investors, Bid/Offer Programme and other offer details, and can be of smaller font.  The portion pertaining to "BRLMs" shall not constitute more than 10% of the price band advertisement space.			
46.	LM is advised to suitably incorporate the comments of the stock exchanges, if any in the UDRHP/RHP.	Noted for compliance.	-	-
		Annexure II – General Observations		
1.	LM is advised to ensure that prior to filing of RHP with Registrar of Companies, the Issuer Company has received crucial clearances I licenses / permissions / approvals from the required competent authority which are necessary for commencement of the activity for which the issue proceeds are proposed to be utilized.	Not applicable. The Net Proceeds of the Offer are not being utilised by the Company for the commencement of any activity which would require approval from any statutory or regulatory authority.	-	-
2.	LM is advised to ensure that the 'Observation Letter' issued by SEBI is included among the material contracts and documents for inspection.	Complied with. It is confirmed that the observation letter issued by SEBI has been included in the "Material Contracts and Documents for Inspection" section of the UDRHP.	531	571





S. No.	Observations	Response	UDRHP Page Number (clean)	UDRHP Page Number (redline)
3.	LM is advised to ensure that prior to proceeding with the issue, "No Objection Certificates" are obtained from all the lenders with whom the company has entered into an agreement and the terms of such agreement require an approval to be taken.	Complied with. The Company has obtained the necessary consents as required under the relevant loan documentation for undertaking activities in relation to the Offer, from the relevant lenders from whom such consent is required.	-	-
4.	LM is advised to ensure that adequate disclosures are made to disclose any material development which may have a material effect on the Issuer Company between the date of registering final prospectus or the red herring prospectus or the letter of offer, with the Registrar of Companies or designated stock exchange, as the case may be, and the date of allotment of specified securities, while ensuring compliance with Regulation 42 and Schedule IX of SEBI (ICDR) Regulations, 2018.	Noted for compliance.	-	-
5.	LM is advised to ensure that exact cross- referencing of page numbers is provided in the offer document instead of general cross- referencing.	Complied with and noted for compliance.	-	-
6.	In terms of SEBI Circulars No. SEBI/CIR/ISD/03/2011, No. SEBI/CIR/ISD/05/2011 and SEBI/CIR/ISD/01/2012 dated June 17, 2011, September 30, 2011 and March 30, 2012 respectively, LM is advised to ensure that 100% promoter holding is in demat form prior to listing.	Complied with. The Equity Shares held by the Promoters are in dematerialized form as on the date of the UDRHP. Further, the disclosure in this regard has been included under the heading "Capital Structure" in the UDRHP.	97	133
7.	LM is advised to ensure that SCORES authentication is taken by the issuer company prior to listing.	Complied with. The Company has obtained SCORES authentication and the disclosure under "Other Regulatory and Statutory Disclosures – Disposal of Investor Grievances by our Company" has accordingly been modified in the UDRHP to reflect this.	457	497
8.	In pursuance of Regulation 25 Sub-Regulation 9(a) of SEBI (ICDR) Regulations, 2018, LM is advised to	Complied with to the extent applicable. We certify that all amendments, suggestions and observations advised by SEBI have been complied with and duly incorporated in the UDRHP.	-	-





S. No.	Observations	Response	UDRHP Page Number (clean)	UDRHP Page Number (redline)
	certify while submitting the in-seriatim reply that all amendments, suggestions and observations advised by SEBI have been complied with and duly incorporated in the offer document, while also indicating the page number for the same.			
9.	i) LM is advised to ensure that sufficient number of Physical ASSA forms are printed and dispatched directly to all designated branches of SCSBs which are located in places of mandatory collection centres as specified in Schedule XII of SEBI (ICDR) Regulations, 2018, Syndicate Members and Registered Brokers of Stock Exchanges, the Registrars to an Issue and Share Transfer Agents (RTAs) and Depository Participants (DPs) . registered with SEBI, at least two days before the opening of the issue. This shall be in addition to ASBA forms which shall be sent to controlling branch of SCSBs for sending to designated branches other than those located in mandatory collection center.  ii) LM is advised to ensure that the ASBA mode of payment is highlighted in bold in all the advertisement / communication informing about the issue. Further, LM is also advised to ensure that the following is suitably incorporated in all advertisements / communications regarding the issue issued by the issuer:	Noted for compliance.	-	





S. No.	Observations	Response	UDRHP Page Number (clean)	UDRHP Page Number (redline)
	a. The following may appear just below the price information of the issue as shown below:			
	"PRICE BAND: RS. xx TO RS. xx PER EQUITY SHARE OF FACE VALUE OF RS. xx EACH			
	THE FLOOR PRICE IS XX TIMES OF THE FACE VALUE AND THE CAP PRICE IS XX TIMES OF THE FACE VALUE			
	BID CAN BE MADE FOR A MINIMUM OF XX EQUITY SHARES AND IN MULTIPLES OF XX EQUITY SHARES THEREAFTER.			
	ASBA .  (APPLICATION SUPPORTED BY BLOCKED AMOUNT)  Simple, Safe, Smart way of Application !!!  Mandatory in public issue .No cheque will be accepted			
	UNIFIED FAUNCHITS INTERFACE  now available in ASBA for retail individual investigation of the control of the co			
	*ASBA is a better way of applying to issues by simply blocking the fund in the bank account.			
	For further details check section on ASBA below. "			





S. No.	Observations	Response	UDRHP Page Number (clean)	UDRHP Page Number (redline)
	<ul> <li>b. The following paragraph on ASBA may be inserted in the advertisement/</li> <li>Communications:</li> </ul>			
	"ASBA has to be availed by all the investors except anchor investors. UPI may be availed by Retail Individual Investors.			
	For details on the ASBA and UPI process, please refer to the details given in ASBA form and abridged prospectus and also please refer to the section "Issue Procedure - Issue Procedure of ASBA Bidders" beginning on page xxx of the Red Herring Prospectus. The process is also available on the website of AIBI and Exchanges in the General Information Document."			
	ASBA bid-cum application forms can be downloaded from the websites of Bombay Stock Exchange and National Stock Exchange and can be obtained from the list of banks that is displayed on the website of SEBI at www.sebi.gov.in.**List of banks supporting UPI is also available on the website of SEBI at www.sebi.gov.in**			





## Appendix A

Name of the lender	Nature of borrowing	Date of original sanction letter	Sanctione d amount (in ₹ million)	Outstanding amount as at October 31, 2023 (in ₹ million)	Repayment date / schedule	Interest rate (p.a.) as at October 31, 2023	Purpose of raising the loan	Pre-payment penalty, if any	Amount proposed to be funded through the Net Proceeds (in ₹ million)
State Bank of India	Cash credit / working capital demand loan*	July 14, 2021	1,250.00	769.30	Repayable on demand.	Cash credit: (6 month MCLR + 0.10%) = 8.50% Working capital demand loan: (3 month TB + 1%) = 7.72%	Working capital requirements.	Nil, in case of prepayment from own sources / 2.00%, in case of takeover of limits by other banks or financial institutions.	656.82
	Term loan	September 8, 2023	800.00	88.29	Maximum tenure of 96 months (including a moratorium of 16 months)	(3 month MCLR + 0.00%) = 8.15%	To set up a new facility at Jammu	Nil, in case of prepayment from own sources / 2.00%, in case of takeover of limits by other banks or financial institutions.	-
Yes Bank Limited	Cash credit / working capital demand loan**	April 29, 2022	650.00	583.28	Maximum tenure of 12 months.	Cash credit: (1 month MCLR + 0.05%) = 9.10% Working capital demand loan: (1 month TB + 1.43%) = 7.90%	Working capital requirements.	Nil, in case of prepayment from own sources / 2.00%, in case of takeover of limits by other banks or financial institutions.	583.28
The Hong Kong and Shanghai Banking Corporation	Cash credit/ Working capital demand loan***	August 19, 2020	100.00	60.54	Repayable on demand.	Cash credit: 8.50% Working capital demand loan: (3 month TB + 1.43%) = 8.46%	Working capital requirements.	Nil, in case of prepayment from own sources / 2.00%, in case of takeover of limits by other banks or financial institutions.	60.54
	Term loan	March 19, 2021	200.00	98.08	84 months (including a moratorium of 6 months).	(3 month TB + 1.50) = 8.31%	Acquisition of capital equipment.	Nil, in case of prepayment from own sources / 2.00%, in case of takeover of limits by other banks or financial institutions.	98.08
		March 19, 2021		45.28	84 months (including a moratorium of 6 months).	(3 month TB + 1.50) = 8.31%	Acquisition of capital equipment.	Nil, in case of prepayment from own sources / 2.00%, in case of takeover of limits by other banks or financial institutions.	45.28
HDFC Bank Limited	Cash credit/ working capital demand loan****	September 26, 2022	200.00	120.26	Repayable on demand.	Cash credit: (3 month TB + 1.46%) = 8.17% Working capital demand loan: (3	Working capital requirements	2% per annum over and above agreed rate of interest.	-





Name of the lender	Nature of borrowing	Date of original sanction letter	Sanctione d amount (in ₹ million)	Outstanding amount as at October 31, 2023 (in ₹ million)	Repayment date / schedule	Interest rate (p.a.) as at October 31, 2023	Purpose of raising the loan	Pre-payment penalty, if any	Amount proposed to be funded through the Net Proceeds (in ₹ million)
						month TB + 1.46%) = 8.52%			
	Term loan	September 26, 2022	2,300.00	1,309.83	120 months (including a moratorium of 24 months)	3 month TB + 1.04% = 7.75%	To set up a new facility at Jammu	2% per annum over and above agreed rate of interest.	-

<sup>\*</sup> Working Capital demand loan amounting to ₹650 million is within overall cash credit limit of ₹1,250 million

**Note:** In accordance with Clause 9(A)(2)(b) of Part A of Schedule VI of the SEBI ICDR Regulations which requires a certificate from the statutory auditor certifying the utilization of loan for the purposed availed, our Company has obtained the requisite certificate dated [•] from our Statutory Auditors, B S R & Co. LLP, Chartered Accountants, wherein the Statutory Auditors have certified that nothing has come to their attention that causes them to believe that the loans that are proposed to be repaid or pre-paid out of Net Proceeds have not been utilized for the purposes for which these were availed.

<sup>\*\*</sup> Working Capital demand loan and cash credit amounting to ₹650 million are within overall cash credit limit of ₹650 million.

<sup>\*\*\*</sup> Working Capital demand loan amounting to ₹100 million and cash credit limit of ₹100 million are within overall combined fund based limit of ₹100 million.

<sup>\*\*\*\*</sup> Working Capital demand loan amounting to ₹200 million is within overall cash credit limit of ₹200 million.





## Schedule A

## Compliance with in-seriatim response to the Initial Observations

S.	Observation	Response	RHP Page	RHP Page
No.			Number (clean)	Number (redline)
	With reference to the Draft Red He	rring Prospectus of captioned IPO filed with SEBI, you are advised to clarify on the following points:	(Clean)	(rediffe)
GENI	ERAL CLARIFICATIONS:	This Prospectus of Captioned IPO fried with SEBI, you are advised to clarify on the following points.		
1.	LM is advised to examine the applicability of SEBI (Framework for Rejection of Draft Offer Documents) Order, 2012, to the DRHP of Issuer Company and confirm whether the said general order is applicable to the instant public issue/DRHP. LM is further advised to provide a para-wise reply on the applicability of	We have set forth a para-wise confirmation with respect to the non-applicability of each criteria specified under the SEBI (Framework for Rejection of Draft Offer Documents) Order, 2012 dated October 9, 2012 to the DRHP and/or the proposed Offer, as <b>Schedule I</b> .	=	=
2.	aforesaid order.  LM is advised to examine the applicability of SEBI (Issuing Observations on Draft Offer Documents Pending Regulatory Actions) Order, 2020, dated February 05, 2020, to the DRHP of Issuer Company and confirm whether the said general order is applicable to the instant public issue/DRHP. LM is further advised to provide a para-wise reply on the applicability of aforesaid order.	We have set forth a para-wise confirmation with respect to the non-applicability of each criteria specified under the Securities and Exchange Board of India (Issuing Observations on Draft Offer Documents Pending Regulatory Actions) Order, 2020 dated February 5, 2020 to the DRHP and/or the proposed Offer, as <b>Schedule II</b> .	-	-
3.	LM is also advised to disclose whether any action has been	Complied with to the extent applicable. Details of all pending actions by statutory or regulatory authorities against the Promoters and Directors of the Company have been disclosed in the section titled "Outstanding Litigation and Other"	424	460





S. No.	Observation	Response	RHP Page Number (clean)	RHP Page Number (redline)
	taken / is pending against the promoter / promoter group/director of the issuer / Group Companies etc. by any regulatory authority in India or overseas.	Material Developments" on page 412 of the DRHP. Further, as on the date of the DRHP, there were no pending litigation proceedings involving the Group Companies which will have a material impact on the Company, and a disclosure in this regard has been included on pages 238 and 419 of the DRHP.  The updated draft red herring prospectus ("UDRHP") and RHP will be updated with any further developments in such matters and to include further matters that may have been initiated since the date of filing of the DRHP, if any.  Further, in terms of the SEBI ICDR Regulations, there is no requirement for the Company to disclose details of any litigation involving members of the Promoter Group. Accordingly, no disclosures for litigation involving members of the Promoter Group are sought to be made.		
4.	LM is advised to submit whether there has been any instance of issuance of equity shares in the past by the issuer Company, the Group Companies or entities forming part of the Promoter Group to more than 49 investors in violation of:  a) Section 67(3) of Companies Act, 1956; or b) relevant section(s) of Companies Act, 2013, including Section 42 and the rules notified thereunder; or c) the SEBI Regulations; or d) the SEBI (Disclosure and Investor Protection) Guidelines, 2000, as applicable.	There have been no instances in the past of issuance of equity shares by the Company, the Group Companies or entities forming part of the Promoter Group to more than 49 / 200 investors, as applicable, in violation of:  Section 67(3) of Companies Act, 1956; or  relevant section(s) of Companies Act, 2013, including Section 42 and the rules notified thereunder, each as amended; or  the SEBI ICDR Regulations; or  the SEBI (Disclosure and Investor Protection) Guidelines, 2000, as applicable.	-	-
5.	It has been observed that in various instances, disclosures	The phrase 'we believe' has been used primarily in the sections titled "Our Business", "Risk Factors", "Our Objects" and "Management's Discussion and Analysis of Financial Position and Results of Operations" in the DRHP, and in the	-	-





S. No.	Observation	Response	RHP Page Number	RHP Page Number
	have been made in the offer document stating 'we believe and /or we are' LM is advised to provide the basis for making such disclosures in the offer document while also explaining compliance with Regulation 24 (1) and Regulation 25 (2) (b) of SEBI (Issue of Capital and Disclosure Requirements) Regulations, 2018 ("ICDR Regulations").	context of disclosures relating to the business and operations of the Company and the industry in which it operates, to the extent that such disclosures are qualitative in nature and not entirely quantifiable. The term 'we believe' has been used in certain statements while describing the expected effect or possible outcome of a particular factor which cannot be quantifiably expressed or where the effect of a singular factor could not be distinguished and expressed in isolated terms from other factors that may have also affected a particular outcome. Further, such statements have also been used in these sections to describe the Company's belief in the strategies adopted by the Company in various aspects of its operations and its impact on the growth of its business.  Therefore, to the extent that any disclosures are qualitative in nature and not entirely quantifiable or ascribable to a particular cause, such disclosures have been qualified by the phrase 'we believe' in order to highlight to potential investors that such statements, while based on reasonable assumptions, cannot be confirmed in tangible terms or by an independent source.  Further, the phrase 'we believe' has not been used in relation to the Company's market standing or position relative to its competitors. Any statements in relation to the market standing of the Company or its position relative to its competitors have been included based on the statements derived from the CRISIL Report.  Accordingly, the DRHP is in compliance with Regulation 24(1) of the SEBI ICDR Regulations, to the extent applicable. Statements containing 'we believe' which will be retained in the UDRHP, RHP and the Prospectus to be filed in connection with the Offer will continue to be those which meet the rationale set out above.  In compliance with Regulation 25(2)(b) of the SEBI ICDR Regulations, the BRLMs have submitted a due diligence certificate dated June 28, 2022, along with (i) a due diligence process note as required under Form A of Schedule V of the SEBI ICDR Regulat	(clean)	(redline)
6.	LM is advised to refrain from using adjectives and replace the words such as largest, biggest etc. used at various places in the draft offer document.	Noted for compliance.	-	-
7.	LM is advised to ensure that all the cross references given in the offer document are correct, leading to the exact page, instead of referring to the beginning of the section. LM may provide the exact	Noted for compliance to the extent applicable.	_	_





S.	Observation	bservation Response		RHP Page
No.			Number (clean)	Number (redline)
	risk factor number instead of			
	giving cross referencing of the			
	page no., in all the references in			
	the document.			
SPEC	IFIC CLARIFICATIONS:			
8. Fo	rward looking statements			
	It may be noted that point (e)	Complied with.	24	29
	under Instructions of Part A of			
	Schedule VI of ICDR Regulations,	Paragraph (11)(I)(C)(ii) of Part A of Schedule VI of the SEBI ICDR Regulations requires that factors that may affect the		
	2018, states that the offer	results of operations of an issuer shall be disclosed in its offer documents. Further, Paragraph (10)(B)(2) of Part A of		
	document should not make any	Schedule VI of the SEBI ICDR Regulations requires that statements about the business strategy of an issuer shall be		
	forward looking statements that	disclosed in its offer documents, without any forecast of projections relating to the financial performance of the issuer.		
	cannot be substantiated. In view Further, in accordance with Paragraph (5)(D)(1) of Part A of Schedule VI of the SEBI ICDR Regulations, details of risks			
	of the same, you are advised to	envisaged by an issuer are required to be disclosed in its offer documents.		
	confirm and explain compliance			
	with aforesaid provision with	In light of these disclosure requirements, certain forward-looking statements have been included in the DRHP to		
	respect to all such forward looking	describe, among other things, the business strategies of the Company, plans, the potential effect or outcome of certain		
	statements made in the DRHP.	risks which cannot be quantified, and to indicate certain significant factors which could potentially have an impact on		
		the results of operations or financial condition of the Company. In terms of paragraph (e) under Instructions of Part A		
		of Schedule VI of the SEBI ICDR Regulations, there is a restriction to include in the offer documents, any forward-		
		looking statements, unless such statements can be substantiated. As set out in the section titled "Forward-Looking		
		Statements" on page 24 of the DRHP, the assumptions based on which such forward-looking statements have been		
		made in the DRHP are reasonable and reflect the views of the Company's management based on available information		
		as of the date of the DRHP. Accordingly, such statements included in the DRHP are in compliance with paragraph (e)		
		under Instructions of Part A of Schedule VI of the SEBI ICDR Regulations.		
9. Ris	sk Factors			
a)	LM is advised to rearrange the risk	Complied with and noted for compliance.	-	-
	factors based on materiality.			
		The risk factors included in the DRHP were arranged and disclosed in order of materiality as perceived by the Company,		
		in terms of (i) the potential impact of the relevant risk on the financial condition, cash flows and results of operations		
		of the Company, and (ii) the likelihood of the occurrence of the relevant risk event described.		





S. No.	Observation	Response	RHP Page Number (clean)	RHP Page Number (redline)
		In addition, while determining the materiality of a risk, in accordance with Paragraph (5)(C) of Schedule VI of the SEBI ICDR Regulations, the following factors were also considered:  • some risks may not be material individually but may be material when considered collectively;  • some risks may have an impact which is qualitative though not quantitative; and  • some risks may not be material at present but may have a material impact in the future	-	
b)	<b>Risk Factor 4</b> : LM is advised to disclose top 10 CDMO customers based on the revenue from operations.	Complied with and noted for compliance. We undertake to amend Risk Factor 4 in the UDRHP and RHP as follows:  "4. We depend on a limited number of contract development and manufacturing organization ("CDMO") customers, including leading Indian pharmaceutical companies. Any adverse developments or inability to enter into or maintain relationships with these CDMO customers could have an adverse effect on our business, results of operations and financial condition.	37	46
		Our CDMO business is focused on providing products and services across a diverse range of generic pharmaceutical products for Indian pharmaceutical companies who market such products under their own brand names to the end users. In Fiscal 2019, Fiscal 2020 and Fiscal 2021 and in the nine months ended December 31, 2021, revenue from our CDMO business on a restated consolidated basis were ₹3,313.68 million, ₹3,429.19 million, ₹3,708.71 million and ₹5,400.64 million, respectively, and accounted for 93.13%, 91.86%, 90.31% and 92.46%, respectively, of revenue from operations on a restated consolidated basis. In Fiscal 2019, Fiscal 2020 and Fiscal 2021 and in the nine months ended December 31, 2021, revenue from our CDMO business on a proforma basis were ₹3,947.68 million, ₹3,911.00 million, ₹4,050.31 million and ₹4,591.57 million, respectively, and accounted for 79.98%, 72.25%, 65.98% and 72.45%, respectively, of revenue from operations on a proforma consolidated basis. Further, the number of CDMO customers that we have represented has increased from 108 in Fiscal 2019 to 164 in the nine months ended December 31, 2021 on a restated consolidated basis and has increased from 155 in Fiscal 2019 to 164 in the nine months ended December 31, 2021 on a proforma consolidated basis.		
		31, 2021 on a proforma consolidated basis. Our business, results of operations and financial condition are dependent on our relationships with and continued supply to our Indian pharmaceutical customers. However, some of our customers may start manufacturing at their own facilities and may discontinue the use of our CDMO services and products. Further, we typically plan and incur capital expenditure for future periods. Delays in successfully entering into contracts for utilization of upcoming capacity may result in lack of proportionate increase in our revenues and results of operations, vis-à-vis an installed capacity increase. In addition, there can be no assurance that we will be able to maintain historic levels of business with our significant customers. If we are unable to maintain relationships with the Indian pharmaceutical companies on existing terms and conditions and if there is delay in replacing these		





S. No.	Observation			Response	Response					
		operations and financia our reputation, it could	l condition. Further, if a have a follow-on effect	any such customer rela t on our ability to enga	ntionship terminations age with new custome	t on our business, results of result in adverse impact on rs.  2021, our top 10 customers				
		₹2,491.15 million, respections from our CD on a restated consolidarespectively, which reprour CDMO business on cended December 31, ₹2,276.26 million, ₹2,155.18%, 55.98% and 5 consolidated basis; ar ₹2,963.18 million, ₹2,606.69%, 66.78% and 6 consolidated basis.	ontributed revenues on a restated consolidated basis of ₹1,992.38 million, ₹1,948.28 million, ₹2,022.01 million and 2,491.15 million, respectively, which represented 60.13%, 56.81%, 54.52% and 46.13%, respectively, of revenue from perations from our CDMO business on a restated consolidated basis; and our top 20 customers contributed revenues in a restated consolidated basis of ₹2,478.74 million, ₹2,270.02 million, ₹2,365.33 million and ₹3,035.77 million, respectively, which represented 74.80%, 66.20%, 63.78% and 56.21%, respectively, of revenue from operations from our CDMO business on a restated consolidated basis. In Fiscal 2019, Fiscal 2020 and Fiscal 2021 and in the nine months and the description of the nine months and the description of the nine months of 2,276.26 million, ₹2,158.06 million, ₹2,267.48 million and ₹2,491.15 million, respectively, which represented 57.66%, 5.18%, 55.98% and 54.25%, respectively, of revenue from operations from our CDMO business on a proforma consolidated basis; and our top 20 customers contributed revenues on a proforma consolidated basis of 2,963.18 million, ₹2,608.10 million, ₹2,704.73 million and ₹3,007.05 million, respectively, which represented 75.06%, 66.9%, 66.78% and 65.49%, respectively, of revenue from operations from our CDMO business on a proforma							
		Top Ten Customers (1)	Fiscal 2019	Fiscal 2020	Fiscal 2021	(in ₹ million)  Nine Months ended				
			FISCUI 2019	FISCUI 2020	FISCUI 2021	December 31, 2021				
		1	654.64	619.27	580.70	622.00				
		2	180.17	275.26	183.27	621.46				
		3								
		4	4 209.71 194.97 185.35 153.97							
		5	95.07	111.19	163.31	339.85				
		6	240.16	153.78	221.32	80.64				
		7	129.03	121.57	113.07	181.81				





S. No.	Observation			Response			RHP Page Number (clean)	RHP Page Number (redline)		
		8	188.35	48.92	66.75	100.41				
		9	48.81	90.01	98.06	151.30				
		10	17.68	98.10	85.11	78.41				
		Total	1,992.38	1,948.28	2,022.01	2,491.15				
		basis for Fiscal 20	1) The top ten customers provided are our top ten customers in terms of revenue contribution on a restated consolidated basis for Fiscal 2019, Fiscal 2020, Fiscal 2021 and the nine months ended December 31, 2021.  Evenue from our top 10 customers on a proforma consolidated basis for the periods indicated are set forth below.  (in ₹ million)							
		Top Ten Customers (1)	Fiscal 2019	Fiscal 2020	Fiscal 2021	Nine Months ended December 31, 2021				
		1	654.64	619.27	580.70	622.00				
		2	220.13	297.43	221.19	621.46				
		3	228.75	235.21	325.06	161.30				
		4	269.46	241.72	240.74	153.97				
		5	300.08	197.80	282.58	80.64				
		6	132.42	149.92	187.31	339.85				
		7	174.00	157.82	159.74	181.81				
		8	227.51	66.48	81.42	100.41				
		9	48.81	90.01	98.06	151.30				
		10	20.45	102.40	90.67	78.41				
		Total	2,276.26	2,158.06	2,267.48	2,491.15				
		* *	· ·	op ten customers in term. 021 and the nine months (	•	on on a proforma consolidated 121.				
c)	Risk Factor 5: LM is advised to disclose top 10 export (final product) and import (raw material) jurisdictions and their	• •			to Risk Factor 6. Wo	e undertake to amend Risk	34	42		





S. No.	Observation		RHP Page Number (clean)	RHP Page Number (redline)		
	share in the total export and import, respectively.					
d)	Risk Factor 5: LM is advised to disclose market share of the Company in India and abroad for the segments in which Company operates. Any increase / decrease of the market share, in these segments, in past 3 financial years to be disclosed. LM is also advised to provide names of key competitors for the segments in which the Company operates.	Noted for compliance. We understand that the of share and competitors in the CDMO formulation. We undertake to amend Risk Factor 6 in the UDF.  "6. We operate in a market that is highly companies in India and other jurisdictions. In products of other suppliers in India and other jurisdictions. In products of other suppliers in India and other jurisdictions. In products of other suppliers in India and other jurisdictions. In products of other suppliers in India and other jurisdictions. In products of other suppliers in India and other jurisdictions. In products of other suppliers in India and other jurisdictions. In products of other suppliers in India and other jurisdictions. In products of other suppliers in India and other jurisdictions. In products of other suppliers in India and other jurisdictions. In products of other suppliers in India and other jurisdictions. In products of other suppliers in India and other jurisdictions. In products of other suppliers in India and other jurisdictions. In products of other suppliers in India and other jurisdictions. In products of other suppliers in India and other jurisdictions. In products of other suppliers in India and other jurisdictions. In products of other suppliers in India and other jurisdictions. In products of other suppliers in India and other jurisdictions. In products of other jurisd	Institutions was available from the and RHP as set forth below the products. We compete to products, particularly for addition, our branded generalised companies in the CDMO in acceutical outsourcing or CDM contract manufacturers proving an acceuting services to fill the mented in terms of both, number tely 15,000 unorganised platition, with large number of or acceuting players is lowered owing CDMO segment include Akumacceuticals Limited, Tirupati Manacceuticals Limited Manacceuticals Limi	the industry expert, CRISIL Research.  A provide outsourced pharmaceutical or formulations, to pharmaceutical or formulations, competition in the CDMO of the	34	42
		Company Name	Date of Incorporation	Registered office location		
		Acme Formulation Private Limited				
		Akums Drugs and Parmaceuticals Ltd	Akums Drugs and Parmaceuticals Ltd 2004 Delhi			
		Innova Captab Limited	2005	Mumbai		
		Synokem Pharmaceuticals Limited	1983	Delhi		
		Theon Pharmaceuticals Limited	2005	Chandigarh		





S. No.	Observation		Response		RHP Page Number (clean)	RHP Page Number (redline)
		Tirupati Medicare Ltd	2005	Delhi		
		Windlas Biotech Ltd	2001	Dehradun		
		Note: The list of competitors is an indicative list and in Sources: MCA, company websites and filings, CRISIL Re (Source: CRISIL Report).  In addition, in Europe and Asia, there are a large serve only their local or national markets. Also, I of their manufacturing capacity, and any such	esearch re number of privately owned, a large pharmaceutical companies	s have been seeking to divest portions		
		(Source: CRISIL Report, May 2022). We compete portfolio and novelty of new offerings), of suppl time delivery and manufacturing flexibility) and result in a decrease in the fees paid for our service and manufacturing services, which could have financial condition.  Our revenue from exports to our top 10 export services outside India on a restated consolidated 31, 2021.	primarily on the basis of produly (quality, regulatory compliant cost-effective manufacturing. Costs and reduced demand for out a material adverse effect on outdestinations as a percentage of basis are set forth below for the	ct portfolio (range of existing product ce and financial stability), service (on- tompetition may, among other things, sourced pharmaceutical development ar business, results of operations and four revenue from sale of goods and the nine month period ended December		
		Export Country (1)		Nonths ended nber 31, 2021		
		Kenya	Decen	17.27%		
		Venezuela		12.92%		
		Sri Lanka		12.73%		
		Tanzania		11.66%		
		Uganda		9.14%		
		Ethiopia		8.79%		
		Ghana		8.04%		
		Nigeria		5.94%		
		Myanmar		5.41%		
		Republic of the Congo		1.42%		
		Total		93.32%		





S. No.	Observation		Response			RHP Page Number (clean)	RHP Page Number (redline)
		(1) The top ten export countries provid consolidated basis for the nine mont	ths ended December 31, 202	21.			
		Our cost of raw materials from our top in restated consolidated basis are set forth b			- <del>-</del>		
		Import Country (1)		Nine Months end December 31, 20			
		China			20.29%		
		China-SEZ			16.63%		
		Hong Kong			54.88%		
		<u>Netherlands</u>			8.20%		
		Total			100.00%		
		The following table indicates market size our share of the global CDMO formulation		indicated.	MO formulation market and in ₹ billion, except percentages)		
		Particulars Particulars	Fiscal 2019	Fiscal 2020	Fiscal 2021		
		Market size of Global CDMO market (API+ Formulation)	6,874.92	7,315.87	8,186.63		
		Market size of Global CDMO Formulation market	1,230.49	1,330.32	1,525.36		
		Revenue from CDMO services for Innova Captab Limited (1)	3.95	3.91	4.05		
		Market share of Innova Captab Limited in the global CDMO formulation market	0.32%	0.29%	0.27%		
		(1) Market share arrived at using revenu (Source: CRISIL Report).	ue on a proforma consolida	ted basis.			
		The following table indicates market size our share of the Indian CDMO formulation			MO formulation market and		





S. No.	Observation		Response			RHP Page Number (clean)	RHP Page Number (redline)
				(1	in ₹ billion, except percentages)		
		Particulars Particulars	Fiscal 2019	Fiscal 2020	Fiscal 2021		
		Market size of Global CDMO market (API+ Formulation) (1)	821.14	898.44	1,014.08		
		Market size of Global CDMO Formulation market	416.03	449.22	512.11		
		Revenue from CDMO services for Innova Captab Limited (2)	3.95	3.91	4.05		
		Market share of Innova Captab Limited in the global CDMO formulation market  (1) Include domestic and export operations	0.95%	0.87%	0.79%		
		(2) Market share arrived at using rever (Source: CRISIL Report).  For our domestic branded generics busin product categories, and within each cate players are adding generic products to a markets, we compete with local compant that are engaged in manufacturing and resulting business, we expect competition from most Some of our competitors may have substanced from the Greater financial, marketing, technical or demand faster with new, alternative or even expense, our business, results of operation or extent of our customer requirements which could have a material adverse effe	ress, we compete with congory, upon dosage strengtheir portfolio. (Source: dies, multinational corportinarketing generic pharmajor international generic antially greater financial, or other resources may allower ging technologies. If the sand financial conditions and render our service ct on our business, result	ompanies in the Indian magths and drug delivery. It can be companies from the endian mage and companies from the end of t	Many of the pharmaceutical 2). Further, in international om other emerging markets is we grow our international other resources than we do as spond to changes in market inificant market share at our ected. Changes in the nature bsolete or non-competitive, incial condition."		
e)	<b>Risk Factor 7</b> : LM is advised to disclose liabilities on account of product recall in the past.	"7. Our CDMO agreements impose se contractual obligations and/or our custo consequent damage to our reputation v financial condition.	veral contractual obligo omers perceive any defic	ations upon us. If we iency in our service we n	are unable to meet these nay face legal liabilities and		46





S. No.	Observation	Response	RHP Page Number (clean)	RHP Page Number (redline)
		Our CDMO agreements are typically long-term in nature where the tenure of the contract ranges mostly between two to five years, with the option of renewal on mutually agreed terms. Some of these agreements provide that the quality, quantity and specifications for the products shall be approved by the customer and be in accordance with the requirements specified in the relevant agreements. Some agreements also require us to furnish quality assurance and compliance certificates to the customers certifying that the quality of the products is as per the agreed specifications. As per the terms and conditions of the respective agreements, our customers have the right to reject the products in case of, inter alia, manufacturing defects, and discrepancy with respect to prescribed specifications, and we are responsible to replace such products free of any additional cost within a stipulated timeframe along with indemnity to the customer for losses arising from breach of obligations, specification of raw material used and manufacturing defect. In addition, for any changes in the product specifications, analytical methods, batch manufacturing reports or raw material used, some agreements require us to obtain prior consent from our customers. Certain agreements also require us to make best efforts to incorporate suggestions received by the customers. With respect to the proposed changes. We are also responsible for the procurement of raw materials, including APIs and excipients, and packaging materials in accordance with the specifications provided by the customer and in certain cases, the third-party vendor should be approved by the customer. Further, certain of our agreements require customers to provide periodic forecasts/ estimates indicating the quantities of the product they intend to purchase; however, certain portions of such forecasts/ estimates are non-binding in nature. We are, in certain of our agreements, required to assure that the product wall always have a minimum shelf life ranging between 85% to 90% of th		





S. No.	Observation	Response	RHP Page Number (clean)	RHP Page Number (redline)
f)	Risk Factor 8: LM is advised to disclose instances of contractual and product liability claims pursuant to inspection by regulatory authorities in the past three years.	"8. The pharmaceutical market is subject to extensive regulation and failures to comply with the existing and future regulatory requirements in any pharmaceutical market could adversely affect our business in that market, results of operations and financial condition.  We operate in a highly regulated industry and our operations are subject to extensive regulation governing the pharmaceutical market. The development, testing, manufacturing, marketing and sale of pharmaceutical products are subject to extensive regulation in India and other countries where we export our products. We are required to comply with the regulatory requirements of various local, state, provincial and national regulatory authorities, such as the Drugs Controller General of India, Central Drugs Standard Control Organization, State Dugs Controller, Ministry of Health and Family Welfare, Controlling cum Licensing Authority, and for certain facilities involved in producing products for exports, international regulatory authorities, such as regulatory authorities in the Africa and Asia. We are subject to international and national guidelines and regulations concerning development, testing, manufacturing processes, equipment and facilities, including the WHO GMP as well as the Schedule M of the Drugs and Cosmetic Rules, 1945 ("Schedule M"). Further, as we expand our operations and geographic scope, we may be exposed to more complex and new regulatory and administrative requirements and legal risks, any of which may require expertise in which we have limited experience as well as impose significant compliance costs on us.  These regulatory requirements impact many aspects of our operations, including manufacturing, developing, storage, distribution, import and export and record keeping related to our products. Regulatory agencies may, for instance, delay, limit or deny approval for many reasons, including:  • changes to the regulatory approval process, including new data requirements for product candidates in thosejurisdictions in which we or	45	57
		• the manufacturing processes, facilities, systems or personnel may not meet the applicable GMP guidelines.		





S. No.	Observation	Response	RHP Page Number (clean)	RHP Page Number (redline)
		Inspections by regulatory authorities that identify any deficiencies could result in remedial actions, production stoppages or facility closure, which would disrupt the manufacturing process and supply of products to our customers. In addition, Although there have been no instances of contractual and product liability claims pursuant to inspection by regulatory authorities in Fiscal 2019, Fiscal 2020 or Fiscal 2021 or the nine month period ended December 31, 2021, any such future failure to comply could expose us to contractual and product liability claims, including claims by customers or recall or other corrective actions, the cost of which could be significant.		
		In addition, we believe applicable regulations have become increasingly stringent and if new legislation or regulations are enacted or existing legislation or regulations are amended or are interpreted or enforced differently, we may be required to obtain additional approvals or operate according to different manufacturing or operating standards. This may require a change in our development and manufacturing techniques or additional capital investments in our facilities. Any related costs may be significant. For example, some of the reforms mentioned in the Draft Pharma Policy, 2017 such as discontinuation of loan licensing (contract manufacturing), regulating marketing practices, banning of brand names, if implemented, will negatively disrupt the domestic pharmaceuticals industry. (Source: CRISIL Report, May 2022). In addition, in September 2018, the Ministry of Health and Family Welfare, GoI, banned over 325 fixed-dose combination drugs, following the recommendations of an expert committee, which found that the combinations lacked "therapeutic justification". If we fail to comply with applicable regulatory requirements in the future, then we may be subject to warning letters and/or civil or criminal penalties and fines, suspension or withdrawal of regulatory approvals, product recalls, seizure of products, restrictions on the import and export of our products, debarment, exclusion, disgorgement of profits, operating restrictions and criminal prosecution and the loss of contracts and resulting revenue losses."		
g)	Risk Factor 9: LM is advised to disclose comparable benchmarks (Industry or comparable peers) for R&D expenses.	Noted for compliance. We undertake to amend Risk Factor 9 in the UDRHP and RHP as set forth below. Please note that R&D expenditure for the Company was not available, and the Company is in the process of estimating the same. Once this data is available, it will be furnished to CRISIL Research, and we undertake to update the information provided in Risk Factor 9 below.  "9. We are dependent on our R&D activities for our future success. If we do not successfully develop new products or continue our generic product portfolio expansion in a timely and cost-effective manner, our business, results of operations and financial condition may be adversely affected.	47	59
		The pharmaceutical and healthcare industry is characterised by technological advancements, introduction of		





S. No.	Observation		Response			RHP Page Number (clean)	RHP Page Number (redline)
		innovative products, price fluctuations resources towards our R&D activities. manufacturing facility at Baddi, Himac generic products, manufacturing procelaboratory is focused on developing ey While we have made significant investmactivities will yield proportionate result developed or launched as a result of sur The following table sets forth the R&D CDMO formulation players and Indian of	We have a dedicated F hal Pradesh. We are incresses and technologies for the ments in R&D activities, this of substantial commercial R&D activities.	R&D laboratory and piceasingly engaged in R&diverse therapeutic segmanufacture of upconcere can be no assurance ial value or that commensus	lot equipment located at our D activities to develop various gments. In particular, our R&D ning patent expired products. In the text our expenditure on R&D ercially viable products may be		
					(in ₹ million, except percentages)		
		Company Name	Total Income	R&D expenditure	R&D expenditure as a % of total income		
		Indian CDMO formulation players			oj total meeme		
		Akums Drug and Pharmaceuticals Ltd.	27,438.85	280.31	1.02%		
		Innova Captab Limited (1)	6,162.54	[ <b>•</b> ]*	[•]*		
		Synokem Pharmaceuticals Ltd.	5,576.06	70.20	1.26%		
		Windlas Biotech Ltd.	4,306.96	36.06	0.84%		
		Theon Pharmaceuticals Ltd.	4,005.06	9.90	0.25%		
		Acme Formulations Private Itd.	4,127.53	47.98	1.16%		
		Indian domestic formulation players					
		Abbott India	43,909.20	10.80	0.02%		
		Alembic Pharma	54,031.40	7,330.00	13.57%		
		Aurobindo Pharma Ltd.	251,554.70	15,993.80	6.36%		
		Biocon Ltd	73,603.00	5,531.00	7.51%		
		Cipla Ltd	194,255.80	8,667.00	4.46%		
		Dr.Reddy's Laboratories Ltd.	193,389.00	17,333.00	8.96%		





S. No.	Observation		Response			RHP Page Number (clean)	RHP Page Number (redline)
		GlaxoSmithKline	30,361.84	18.00	0.06%		
		Glenmark Pharmaceuticals Ltd.	109,941.45	13,187.00	11.99%		
		Ipca Labs	54,828.30	1,266.70	2.31%		
		Lupin Ltd#	152,992.50	14,324.20	9.36%		
		Panacea Biotech Ltd.	6,347.82	247.70	3.90%		
		Sun Pharmaceuticals Industries Ltd.	343,336.60	21,443.30	6.25%		
		Torrent Pharmaceuticals Ltd	80,614.80	4,870.00	6.04%		
		Wockhardt Ltd.	28,405.70	2,655.00	9.35%		
		(Source: CRISIL Report).  The success of our new generic product anticipate customer needs and preference and customers; customize our productio effective manner; and successfully marke of new products is characterised by significationing regulatory approvals, building manufacturing and R&D facilities and expert of cost overruns, without a propose CDMO and generic segments may have gobetter position to identify market trends, of if we do not successfully develop new promanner that is attractive to our customers affected.  Our future results of operations also dependent of the continue our product portfolio expansions in the continue our capacities in capabilities in order to ensure continued.	es; obtain timely regulatory and an capacities, develop and and tand sell our products. In accant upfront costs, including ginventory and sales and suipment for future expansionate increase in revenue reater financial, research and adapt to changes in industry oducts or continue our products, our business, results of operation in a timely and cost-efficient product portfolio by enterexisting products as well and cost-efficient degree, or existing products as well and cost-efficient degree.	reprovals; establish collaboration and products didition, the development of costs relating to product of marketing. Our planned on could result in higher of sections and offer innovative new part portfolio expansion in our ability to successfully fective manner. Further, as expanding and strengas expanding and strengas and strengand strengand on the could be successfully fective manner.	crations with suppliers in a timely and cost- and commercialisation development activities, in investments in new costs, especially in the four competitors in the They may also be in a products. Accordingly, a timely, cost-effective ition may be adversely develop new products is part of our business we chains. In addition, othering our research		





S. No.	Observation	Response	RHP Page Number (clean)	RHP Page Number (redline)
		success. The development and commercialisation of new products (whether ours or our customers' products) are complex, time-consuming, costly and involves a high degree of business risk. We may be unable to successfully create these new products or encounter unexpected delays in the launch of these products and even if launched as planned, such products may not perform as we expect."		
h)	<b>Risk Factor 10</b> : LM is advised to disclose complaints received in the past 3 years and its financial implications.	Complied with and noted for compliance. We undertake to amend Risk Factor 10 in the UDRHP and RHP as follows:  "10. We are subject to strict technical specifications, quality requirements, regular inspections and audits by our CDMO customers including leading Indian pharmaceutical companies. Our failure to comply with the quality standards and technical specifications prescribed by such customers may lead to loss of business from such customers and could negatively impact our business, results of operations and financial condition, including cancellation of existing and future orders which may expose us to warranty claims.	39	48
		Our products and manufacturing processes are subject to stringent quality standards and specifications, typically specified by our CDMO customers in their respective agreements. Adherence to quality standards is a critical factor in our production process as any deviations from the required specifications by our Company or failure to comply with the technical specifications of our customers regarding the composition of drugs, may lead to a recall of products or cancellation of the orders placed by our customers. Further, for any change in the product specifications, manufacturing process, manufacturing site, manufacturing method or raw material used, we are required to inform or obtain prior consent from some of our CDMO customers.		
		During Fiscal 2019, Fiscal 2020 and Fiscal 2021 and during the nine months ended December 31, 2021, we have received certain complaints from our CDMO customers. The nature of these complaints was mainly related to packaging material of the products. There have not been any financial implications related to these complaints. We undertake corrective and preventive actions on these complaints on a regular basis.		
		While we believe we undertake the necessary measures and engage internal and external experts to ensure that our facilities comply with the applicable standards as imposed by our customers, any failure on our part to maintain the applicable standards and manufacture products according to prescribed specifications, may lead to cancellation of the order, loss of customers, loss of reputation and goodwill of our Company. Additionally, it could expose us to indemnity, warranty claims, monetary liability and/or litigation. Our CDMO customers are typically provided the right to audit our manufacturing facilities, processes or systems, under such agreements, after providing a certain period of notice. While we have not received any adverse observations in the past from our customers pursuant to such audits, there can be		





S. No.	Observation	Response	RHP Page Number (clean)	RHP Page Number (redline)
		no assurance that such audits would not result in any adverse observations in the future or that our customers will necessarily engage us for their outsourcing operations. The audit may involve inspection of, inter alia, our manufacturing facility and equipment, quality control procedures, review of the manufacturing processes and raw materials and packaging. The finished product delivered by us is further subject to laboratory validation by certain customers. This is an extensive and stringent process undertaken by our customers. Occurrence of any event on account of errors and omission could result in damage to our reputation and loss of customers, which could adversely affect our business, operations, our cash flows and financial condition. In the past, we have received certain complaints from our customers for which our Company has undertaken corrective measures on a regular basis as appropriate, and there can be no assurance that we would not receive such complaints in the future as well.		
		One of our manufacturing facilities is Schedule M compliant. If we fail to comply with applicable quality standards specified by our customers or if the relevant accreditation institute or agency declines to certify our products, or if we are otherwise unable to obtain such quality accreditations in the future, within time or at all, our business, results of operations and financial conditions will be materially and adversely affected. The quality of our products is critical to the success of our business and depends on the effectiveness of our quality assurance system, which, in turn, depends on a number of factors, including the design of our facility, our training program, and the checks and balances implemented at stage of development/ manufacturing and testing processes in line with the current GMP guidelines. While other than incidents in the ordinary course of business, there has not been any failure or deterioration of quality systems in the past, any significant failure or deterioration of our quality system in future could result in defective or substandard products, which, in turn, may result in delays in the delivery of our products and the need to replace defective or substandard products. As a result, our business, results of operations and financial condition could be materially and adversely affected."		
i)	Risk Factor 12: LM is advised to disclose any material impact of changes in environmental, health and safety or labour laws, in past 3 financial years.	Noted for compliance. We undertake to amend Risk Factor 12 in the UDRHP and RHP as follows:  "12. Our operations are subject to environmental and workers' health and safety laws and regulations. We may have to incur material costs to comply with these regulations or suffer material liabilities or damages in the event of an incidence or non-compliance of environmental and other similar laws and regulations which may have a material adverse effect on our reputation, business, financial condition and results of operations.  Our operations are subject to extensive environmental and hazardous waste management laws and regulations in India, including the Environment (Protection) Act, 1986 and the Environment Protection Rules, 1986, the Air (Prevention and Control of Pollution) Act, 1981, the Water (Prevention and Control of Pollution) Act, 1974, the	45	57





S. No.	Observation	Response	RHP Page Number (clean)	RHP Page Number (redline)
		Hazardous and Other Wastes (Management and Transboundary Movement) Rules, 2016, and other rules and regulations promulgated by the Ministry of Environment, Forest and Climate Change, Government of India ("MoEF") and various other statutory and regulatory authorities and agencies in India. The pharmaceutical industry is subject to strict regulations with respect to a range of environmental matters including limitations on land use, licensing requirements, management of materials used in manufacturing activities, the storage of inflammable and hazardous substances and associated risks, the storage, treatment and disposal of wastes, remediation of contaminated soil and groundwater, air quality standards, water pollution and discharge of hazardous materials into the environment. For details of the key regulations applicable to our business in India, see "Key Regulations and Policies" on page 198. The discharge or emission of chemicals, dust or other pollutants into the air, soil or water that exceed permitted levels and cause damage to others may give rise to liabilities towards the government and third parties and may result in our incurring costs to remedy any such discharge or emissions.		
		Environmental laws and regulations in India have become and continue to be more stringent, and the scope and extent of new environmental regulations, including their effect on our operations, cannot be predicted with any certainty. In case of any change in environmental or pollution regulations, we may be required to invest in, among other things, environmental monitoring, pollution control equipment, and emissions management and other expenditure to comply with applicable environmental standards. Any failure on our part to comply with any existing or future regulations applicable to us may result in legal proceedings, including public interest litigation, being commenced against us, third party claims or the levy of regulatory fines. Further, any violation of the environmental laws and regulations may result in fines, criminal sanctions, revocation of operating permits, or shutdown of our manufacturing facilities.		
		In addition, as a responsible corporate entity, we are continuously endeavouring to comply with Environmental, Social & Governance compliance requirements covering aspects of legal compliance, ethics & business conduct, quality & patient safety, human rights, labour and employment, health safety and well-being of employees, sustainability & environmental responsibility and quality management systems.		
		We are also subject to the laws and regulations governing employees in areas such as minimum wage and maximum working hours, overtime, working conditions, hiring and termination of employees, contract labour and work permits. There is a risk that we may fail to comply with such regulations, which could lead to enforced shutdowns and other sanctions imposed by the relevant authorities, as well as the withholding or delay in receipt of regulatory approvals for our new products. While there have been no instances where we have failed to comply with regulations that has resulted in a shutdown or other sanctions being imposed on us, we cannot assure you that we will not be involved in future litigation or other proceedings or be held liable in any litigation or proceedings including in relation to safety,		





S. No.	Observation	Response	RHP Page Number (clean)	RHP Page Number (redline)
		As a consequence of unanticipated regulatory or other developments, future environmental and regulatory related expenditures may vary substantially from those currently anticipated. Although there has been no material impact on our business, results of operations or financial condition due to changes in environmental, health and safety or labour laws in Fiscal 2019, Fiscal 2020 or Fiscal 2021 or the nine month period ended December 31, 2021, We we cannot assure you that our costs of complying with current and future environmental laws and other regulations will not adversely affect our business, results of operations or financial condition. In addition, we could incur substantial costs, our products could be restricted from entering certain markets, and we could face other sanctions, if we were to violate or become liable under environmental laws or if our products become non-compliant with applicable regulations. Our potential exposure includes fines and civil or criminal sanctions, third-party property damage or personal injury claims and clean-up costs. The amount and timing of costs under environmental laws are difficult to predict."		
j)	Risk Factor 13: LM is advised to disclose top 10 third party contract research organizations which conduct clinical trials.	Noted for compliance. The Company consulted CRISIL Research, the industry expert, and was able to identify contract research organizations providing clinical trials in the global and Indian market, but CRISIL Research was not able rank these organizations. We undertake to amend Risk Factor 13 in the UDRHP and RHP as follows:  "13. Any failure of the third parties, on whom we rely for clinical trials, in performing their obligations and complying with regulatory standards could result in a delay in receiving regulatory approval and adversely affect our business, financial condition and results of operations.	46	58
		We depend on third party qualified contract research organisations to conduct clinical trials and studies of our new products and expect to continue to do so. We rely on such parties for successful execution of our clinical trials and studies, however, we do not control many aspects of their activities.  Set forth below are key pharmaceutical contract research organization providing clinical trial solutions in the global and Indian market as of July 2022.		
		Company Name Key products/services  Contract research organizations in global market		
		IQVIA Holdings Inc.  Technology & analytics solutions, research & development solution, contract sales & medical solutions		





S. No.	Observation		Response					
		Icon PLC	Clinical research services, molecule development consulting, functional service provision					
		Parexel International Corporation	Clinical research services, outsourcing services, medical communications					
		Thermo Fisher Scientific Inc.	Drug discovery and development, pre-clinical and clinical drug testing, drug formulation manufacturing					
		Labcorp	Drug development consulting, clinical development, clinical testing					
		Syneos Health	Clinical development, consulting					
		Contract research organizations in Indian	market					
		Syngene International Ltd.	Drug discovery, drug development and drug manufacturing					
		Vimta Labs Ltd.	Pre-clinical research, clinical research					
		Veeda Clinical Research Ltd.	Pre-clinical research and development, clinical research and development					
		Jubilant Biosys Ltd.	Drug discovery and contract research services					
		Aragen Life Sciences Pvt. Ltd.	Drug discovery, drug development and drug manufacturing					
		Lambda Therapeutic Research Ltd.	Pre-clinical research, clinical research					
		Clininvent Research Pvt. Ltd.	Drug discovery, drug development and drug manufacturing					
		Siro Clinpharm Private Limited	Clinical operations, medical writing					
		Diagnosearch Life Sciences Pvt.Ltd.	Clinical operations, consulting					
		Note: The list of players is an indicative list Sources: company websites and filings, CRIS (Source: CRISIL Report).						
		under such agreements include protocol documents and monitoring the study, of may not conduct our studies in accordation for confirming that each of our clinical protocol. While, other than in the ordinate have defaulted or not complied with the	til completion of the service stipulated in the agreement. Our responsibilities col review, supply of investigational products, provision of the study related amongst others. Third parties may also not complete activities on schedule or note with applicable trial, plans and protocols. Nonetheless, we are responsible at trials is conducted in accordance with its general investigational plan and pary course of business, there have not been any instances where third parties are robligations, if the third parties fail to carry out their obligations in the future, mmercialisation could be delayed or prevented or an enforcement action could					





S. No.	Observation	Response	RHP Page Number (clean)	RHP Page Number (redline)
		Our reliance on these third parties does not relieve us of our responsibility to comply with the applicable regulations and standards of the regulatory authorities related to good clinical practices. In particular, these third-parties must comply with regulatory standards and their failure to do so could result in warning or deficiency letters from regulatory authorities, which could interfere with or disrupt their ability to complete our studies on time, thereby affecting our product approval process or even forcing a withdrawal of our product which may adversely affect our business, financial condition and results of operations."		
k)	Risk Factor 14: LM is advised to quantify the impact of reforms in the healthcare industry.	Noted for compliance. The Company has indicated that it is difficult to quantify the impact of reforms in the healthcare industry with any certainty. The Company undertakes to provide further information on how price controls in the pharmaceutical industry work and the number of drugs affected. We undertake to amend Risk Factor 14 in the UDRHP and RHP as follows:  "14. Reforms in the healthcare industry and the uncertainty associated with pharmaceutical pricing, reimbursement and related matters could adversely affect the pricing and demand for our products as well as the consumer demand for the products we manufacture for our customers, which may significantly influence our business, results of operations and financial condition.  The healthcare industry has changed significantly over time, including, amongst others, healthcare reform, adverse changes in government or private funding of healthcare products and services, legislation or regulations governing the privacy of patient information or patient access to care, or the delivery, pricing or reimbursement of pharmaceuticals and healthcare services or mandated benefits. Such changes may cause the healthcare industry participants to reduce the number of our services and products that they purchase from us or the price they are willing to pay for our services and products. Changes in the healthcare industry's pricing, selling, inventory, distribution or supply policies or practices could also significantly reduce our revenue and profitability. As a result, our success will depend in part on the extent to which government and health administration authorities regulate the maximum retail price of the products manufactured by us and marketed by our customers. The role of third-party private health insurers and other third-party payers is also becoming important in in-patient settings in hospitals. Increasing expenditures for healthcare has been the subject of considerable public attention in almost every jurisdiction where we conduct business. Both private and go	44	55





S. No.	Observation	Response	RHP Page Number (clean)	RHP Page Number (redline)
		price controls can limit the profit our customers can earn from the market and thereby have a follow-on effect in them seeking a lower price manufacturing supplier for that product. Significant changes in price control limits set by regulators can change the profitability both for our customers as well as for us.  The Government of India has been taking various steps in order to control the prices of drugs and make it more affordable to consumers. The Gol's new Pharmaceutical Policy, notified in 2012, was put out as the final price notification in May 2013, bringing 348 essential drugs in the National List of Essential Medicines ("NLEM") under price control. (Source: CRISIL Report). Between Fiscal 2014 and Fiscal 2015, the industry saw drug prices being regulated for more than 500 medicines under the Drug Price Control Order ("DPCO"), thereby negatively impacting the industry. Further, the revised National List of Essential Medicines NLEM 2015 added more than 100 new drugs under price control, with many high-value chronic drugs from anti-diabetes and HIV being covered under it. As of November 2016, the National Pharmaceutical Pricing Authority ("NPPA") has notified ceiling prices for 540 drugs. The NLEM 2015 contains about 870 scheduled drug formulations. (Source: CRISIL Report).  Under Gol's policy, the ceiling price for each drug under control would be fixed as the simple average price of brands having more than 1 per cent market share (by value) in the sales (MAT - Moving Annual Turnover) of that particular molecule. (Source: CRISIL Report). Thus, prices of brands which are higher than this ceiling will need to be lowered. The ceiling prices will be allowed an annual increase as per the Wholesale Price Index. (Source: CRISIL Report). Prices will be recalculated using MAT only once in five years or when the NLEM is updated. Drugs under the NLEM comprised estimated 15-20% of the overall domestic formulation market in Fiscal 2021. (Source: CRISIL Report).	(clean)	(redline)
		Therefore, the Government's firm stance on pricing even in future might have a negative impact on the profitability for some pharmaceutical companies, which are selling branded generics at a high premium price. Currently, prices for approximately 900 to 1,000 scheduled formulations have been fixed so far. Due to the drop in realizations of pharmaceutical formulations, margins of contract manufacturing players have reduced as well. Therefore, both companies that market pharmaceutical formulations and CDMOs are equally impacted due to the price ceiling imposed by the Government. If the prices of more of our products or our customers products are administered or determined by the DPCO or NPPA or other similar authorities outside India, it would have an adverse impact on our profitability."		
1)	Risk Factor 22: LM is advised to clearly disclose the attrition rate of employees involved in the R&D. Further provide the percentage of employees involved in the R&D	Noted for compliance. We undertake to amend Risk Factor 22 in the UDRHP and RHP as follows:  "22. We are dependent upon the experience and skill of our management team and a number of key managerial personnel as well as on our ability to attract and retain personnel with technical expertise. If we are unable to	50	61





S. No.	Observation			RHP Page Number (clean)	RHP Page Number (redline)							
	vis-à-vis the total employees, in past 3 financial years.	attract or retain such condition.  We are dependent of			, ,,		-					
		experience, along wi	tion and managing our business. We believe that the inputs received from our senior management and th rience, along with the expertise, experience and services of our Promoters and Executive Directors are valuable development of business and operations and the strategic directions taken by our Company. For furth mation, see "Our Management" on page 211. Our ability to meet continued success and future business challeng									
		depends on our abilion number of skilled em relationships with a	inds on our ability to attract, recruit and retain experienced, talented and skilled professionals. Without a sufficient ber of skilled employees, our operations and manufacturing quality could suffer. Our sales team has also developed ionships with a number of distributors and stockists that would be difficult to replace. Competition for qualified nical personnel and operators as well as sales personnel with established dealer relationships is intense, both in									
		retaining our existin pharmaceutical com attract qualified ind	aining our existing employees and when replacing or finding additional suitable employees. Competition among armaceutical companies for qualified employees, particularly R&D personnel, is intense and the ability to retain and ract qualified individuals is critical to our success. During Fiscal 2019, Fiscal 2020 and Fiscal 2021 and the nine anths ended December 31, 2021, there has been only one change in KMP.									
		In addition, we may technical expertise to compensation more expertise that our bu	erminate their emp rapidly than in the µ siness requires. The	loyment with us. Woast to remain completes of services of s	e may also be requosetitive in attracting uch personnel could	uired to increase our g and retaining perso d have an adverse eff	levels of employee onnel with technical ect on our business,					
		results of operations periods indicated										
		Period	Period Un-Skilled Semi-Skilled Skilled Highly Skilled* Total									
		Fiscal 2019										
		Fiscal 2020										
		Fiscal 2021										
		Nine months ended December 31, 2021	Nine months 32.88% 25.88% 7.60% 9.30% 23.20% aded December									





S. No.	Observation		Response		RHP Page Number (clean)	RHP Page Number (redline)					
		* including assistant manager and above	uding assistant manager and above								
		able to recruit unskilled personnel to	above attrition rates reflect the high attrition levels in our business for unskilled employees. While we have been to recruit unskilled personnel to fill vacancies in the past, if we are unable to recruit the requisite unskilled sonnel in the future our business, results of operations and financial condition could be adversely affected.								
		Our attrition rate for R&D employees a periods indicated is set forth below.	ur attrition rate for R&D employees and the number of R&D employees as a percentage of total employees for the priods indicated is set forth below.								
		Period	Attrition Rate % of R&D employees	% of R&D employees of total employees							
		Fiscal 2019	25.00%	2.37%							
		Fiscal 2020	52.94%	2.57%							
		Fiscal 2021	22.22%	2.16%							
		Nine months ended December 31, 2021	7.69%	2.23%							
		personnel or our inability to manage the on our financial results and business properties our ability to expand our business may.  As we intend to continue to expand our retain experienced management, R&D of compensation more rapidly than in the assurance that our competitors will not skilled personnel. Further, as on the depolicies. In the event that we are not business, or if we experience high attrituments.	operations and develop new products, wand sales personnel. We may also be requipast to remain competitive in attracting toffer better compensation packages, into of this Draft Red Herring Prospectusable to attract and retain talented emploion levels which are largely out of our coiness, results of operations and financial	ategories may have an adverse effect nal qualified personnel or retain them, we will need to continue to attract and ired to increase our levels of employee a suitable employees. There can be no centives and other perquisites to such a, we do not have key man insurance loyees as required for conducting our partrol, or if we are unable to motivate							





S. No.	Observation			Response			RHP Page Number (clean)	RHP Page Number (redline)				
m)	Risk Factor 23: LM is advised to	Complied with and note	d for compliance. We ι	undertake to amend Ri	sk Factor 23 in the UI	DRHP and RHP as follows:	51	70				
	disclose the receivables as a											
	percentage of revenues.	•	Our inability to collect receivables and default in payment from our customers could result in the reduction of rofits and affect our cash flows.									
		30 to 90 days. While we limit the credit, we externation and payment I while we maintain what based upon our historic accurate.  Our trade receivables of	najority of our sales are to customers on an open credit basis, with standard payment terms of generally between 90 days. While we generally monitor the ability of our customers to pay these open credit arrangements and the credit, we extend to what we believe is reasonable based on an evaluation of each customer's financial ition and payment history, we may still experience losses because of a customer being unable to pay. As a result, we maintain what we believe to be a reasonable allowance for doubtful receivables for potential credit losses of upon our historical trends and other available information, there is a risk that our estimates may not be rate.  Trade receivables on a restated consolidated basis and our trade receivables as a percentage of revenue from ations on a restated consolidated basis for the periods indicated are set forth below.  (in ₹ million, except percentages)									
		Particulars	Fiscal 2019	Fiscal 2020	Fiscal 2021	Nine Months ended December 31, 2021						
		Trade receivables	908.29	867.69	1,385.53	2,006.82						
		Trade receivables as a			,	<u> </u>						
		percentage of revenue	25.53%	23.24%	33.74%	34.36%						
			our trade receivables on a proforma consolidated basis and our trade receivables as a percentage of revenue from perations on a proforma consolidated basis for the periods indicated are set forth below.									
					(in	₹ million, except percentages)						
		Particulars	December 31, 2021									
		Trade receivables	1,319.14	1,373.62	1,487.31	2,006.82						
		Trade receivables as a										
		percentage of revenue from operations	26.73%	25.37%	24.23%	31.66%						
1		ji oili operutions	20.73%	23.37%	24.25%	31.00/0	1	1				





S. No.	Observation	Response	RHP Page Number	RHP Page Number
			(clean)	(redline)
		In Fiscal 2019, Fiscal 2020 and Fiscal 2021 and in the nine months ended December 31, 2021, our trade receivables		
		on a restated consolidated basis were ₹908.29 million, ₹867.69 million, ₹1,385.53 million and ₹2,006.82 million,		
		respectively, and our receivable turnover days on a restated consolidated basis were 93 days, 85 days, 123 days and		
		94 days, respectively, in the same periods. In Fiscal 2019, Fiscal 2020 and Fiscal 2021 and in the nine months ended		
		December 31, 2021, trade receivables on a proforma consolidated basis were ₹1,319.14 million, ₹1,373.62 million,		
		₹1,487.31 million and ₹2,006.82 million, respectively, and receivable turnover days on a proforma consolidated basis		
		were 98 days, 93 days, 88 days and 87 days, respectively, in the same periods. Any increase in our receivable turnover		
		days will negatively affect our business. If we are unable to collect customer receivables or if the provisions for doubtful		
		receivables are inadequate, it could have a material adverse effect on our business, results of operations and financial condition.		
		Macroeconomic conditions could also result in financial difficulties, including insolvency or bankruptcy, of our		
		customers, and as a result could cause customers to delay payments to us, request modifications to their payment		
		arrangements, that could increase our receivables or affect our working capital requirements, or default on their		
		payment obligations to us. An increase in bad debts or in defaults by our customer, may compel us to utilize greater		
		amounts of our operating working capital and result in increased interest costs, thereby adversely affecting our results		
		of operations and cash flows."		
10. C	apital Structure			
a)	LM is advised to ensure that the	Noted for compliance. As required under Regulation 20 of the SEBI ICDR Regulations, the Company shall ensure that	-	-
	Company will comply with the	the details of the Equity Shares locked-in are recorded by the relevant Depository, and the same has been disclosed		
	lock-in provisions specified in the	on page 107 of the DRHP.		
	SEBI ICDR Regulations.			
	ndustry Overview			
a)	It is observed that in the offer	Noted for compliance, to the extent applicable.	-	-
	document, the issuer has relied			
	upon the data contained in the	Please note that the industry and market data set forth in the Draft Red Herring Prospectus has been obtained from		
	report prepared by CRISIL Limited	the report titled 'Assessment of Indian pharmaceutical and CDMO market', dated May 2022, issued by CRISIL Research,		
	at the request of the issuer,	a division of CRISIL Limited. CRISIL Research will update their report before the Company files the RHP and industry		
	however the data has not been provided as on date. LM is advised	information will be updated in the RHP as of March 31, 2022 or another recent date to the extent such data is available.		
	•			
	to maintain consistency in the data periodicity and ensure that			
	uata periodicity and ensure that			





S. No.	Observation	Response	RHP Page Number (clean)	RHP Page Number (redline)
	the data relied in the report is not older than six months as on the date of filing of offer document with ROC.		(cicuit)	(reduite)
b)	LM is advised to disclose whether the Company, its promoters/directors/KMPs are connected with CRISIL Limited.	Complied with. Based on the confirmation received from CRISIL Research, a division of CRISIL Limited, we have disclosed on page 21 of the DRHP that CRISIL Research is an independent agency and is not related to the Company, Directors, Promoters or Key Managerial Personnel.	21	26
a)	LM is advised to disclose data of past three years at the relevant places and ensure that at all applicable places, where data points are being compared, the same is disclosed in a tabular format.	Complied with and noted for compliance. The section titled "Our Business" disclosed all comparable data for past three years. We undertake to such disclose data in tabular format to the extent possible. We undertake to amend parts of the section "Our Business" as follows:  "Our Business  []  Raw Materials and Procurement  We purchase APIs and other materials such as, excipients and impurities from third party suppliers domestically. In addition, we purchase certain APIs from third party international suppliers. In Fiscal 2019, Fiscal 2020 and Fiscal 2021 and in the nine months ended December 31, 2021, our cost of materials consumed on a restated consolidated basis was ₹2,784.93 million, ₹2,886.90 million, ₹3,014.60 million and ₹4,321.31 million, respectively, and represented 84.07%, 85.64%, 82.44% and 83.78%, respectively, of total expenses on a restated consolidated basis in the same periods. In Fiscal 2019, Fiscal 2020 and Fiscal 2021 and in the nine months ended December 31, 2021, cost of materials consumed on a proforma consolidated basis was ₹3,686.83 million, ₹3,910.54 million, ₹4,118.11 million and ₹4,315.95 million, respectively, and represented 82.63%, 82.10%, 76.75% and 78.58%, respectively, of total expenses on a proforma consolidated basis in the same periods.  The table set forth below provides on a restated consolidated basis our cost of materials consumed and as a percentage of total expenses for the periods indicated.	208, 209, 210	278, 279, 281





S. No.	Observation				Re	esponse					RHP Page Number (clean)	RHP Page Number (redline)
			Fisco	al <b>201</b> 9	Fisca	Fiscal 2020 Fiscal 2			Nine months ended December 31, 2021			
		Particulars	₹ million	% of total expenses	₹ million	% of total expenses	₹ million	% of total expenses	₹ million	% of total expenses		
		Cost of materials consumed	2,784.93	84.07%	2,886.90	85.64%	3,014.60	82.44%	4,321.31	83.78%		
		The table set forth percentage of total	•	•	-	nsolidated b	asis our cos	st of materic	als consume	ed and as a		
			Fisca	I 2019	Fisca	l 2020	Fiscal 2021		Nine months ended December 31, 2021			
		Particulars	₹ million	% of total expenses	₹ million	% of total expenses	₹ million	% of total expenses	₹ million	% of total expenses		
		Cost of materials consumed	3,686.83	82.63%	3,910.54	82.10%	4,118.11	76.75%	4,315.95	78.58%		
		[]										
		Logistics										
		Each of our facilities five depots in major			arehouse, e	nabling smod	oth function	ing of our op	perations. W	/e also have		
		In Fiscal 2019, Fisca							, , ,			
		per our Restated Col respectively, and rep		•			•	,		,		
		Consolidated Finance	<del>cial Informa</del>	ition in the s	<del>ame periods</del>	s. In Fiscal 20	19, Fiscal 2	<del>020 and Fisc</del>	<del>al 2021 and</del>	in the nine		
		months ended Dece million, ₹31.98 millio of total expenses on	<del>on and ₹27.</del>	34 million, re	espectively, c	<del>and represent</del>	<del>ed 0.30%, 0</del>					





S. No.	Observation		Response										
			e table set forth below provides our freight charges as per our Restated Consolidated Financial Information and as percentage of our total expenses as per our Restated Consolidated Financial Information for the periods indicated.										
			Fisco	al 2019	Fisca	ıl 2020	Fisca	l 2021		nths ended er 31, 2021			
		Particulars	₹ million	% of total expenses	₹ million	% of total expenses	₹ million	% of total expenses	₹ million	% of total expenses			
		Freight charges	1.08	0.03%	2.70	0.08%	6.27	0.17%	5.75	0.11%			
			total expenses for the periods indicated.  Fiscal 2019 Fiscal 2020 Fiscal 2021 Nine months ended December 31, 2021										
		Particulars	₹ million	% of total expenses	₹ million	% of total expenses	₹ million	% of total expenses	₹ million	% of total expenses			
		Freight charges	13.23	0.30%	24.33	0.51%	31.98	0.60%	27.34	0.50%			
		[]											
		Utilities											
		We consume fuel as power grid Addition supply of power. ##	onally, we ho on Fiscal 2019 oenses as pe	ave also insto <del>), Fiscal 2020</del> r our Restate	alled genera <del>and Fiscal :</del> d Consolidat	ntors in our n <del>2021 and in t</del> t <del>ed Financial</del>	nanufacturii t <del>he nine mo</del> Information	ng facilities t <del>nths ended E</del> were ₹41.75	o ensure un December 31 million, ₹5!	interrupted 1 <del>, 2021, our</del> 5.41 million,			
		₹54.78 million and total expenses as p											





S. No.	Observation				Re	sponse					RHP Page Number (clean)	RHP Page Number (redline)	
		million, ₹67.17 mil respectively, of tota The table set forth b	the nine months ended December 31, 2021, power and fuel expenses on a proforma consolidated basis were ₹54.40 llion, ₹67.17 million, ₹67.95 million and ₹58.04 million, and accounted for 1.22%, 1.41%, 1.27% and 1.06%, spectively, of total expenses on a proforma consolidated basis.  e table set forth below provides our power and fuel expenses as per our Restated Consolidated Financial Information d as a percentage of total expenses as per our Restated Consolidated Financial Information for the periods indicated.										
			Fisco	al 2019	Fisca	I 2020	Fisca	l 2021		oths ended er 31, 2021			
		Particulars	₹ million	% of total expenses	₹ million	% of total expenses	₹ million	% of total expenses	₹ million	% of total expenses			
		Power and fuel expenses	41.75	1.26%	55.41	1.64%	54.78	1.50%	57.43	1.11%			
		The table set forth be of total expenses fo			orma consoli	dated basis o	our power ar	nd fuel expens	ses and as a	percentage			
			Fisca	l 2019	Fisca	l 2020	Fisca	l 2021		nths ended er 31, 2021			
		Particulars	₹ million	% of total expenses	₹ million	% of total expenses	₹ million	% of total expenses	₹ million	% of total expenses			
		Power and fuel expenses	54.40	1.22%	67.17	1.41%	67.95	1.27%	58.04	1.06%			
		[]"											
	listory and Certain Corporate Matter												
a)	LM is advised that it is categorically disclosed in the	Noted for complian	ce.								227	299	
												l i	
	DRHP under section "History and Certain Corporate Matters" of the												





S. No.	Observation	Response	RHP Page Number	RHP Page Number
	special rights available to the Promoters / Shareholders (except for nominee/nomination rights and information rights) would survive post listing of the Equity Shares of the Company and same shall cease to exist or shall expire / waived off immediately before or on the date shares are allotted to public shareholders in IPO, without requiring any further action.		(clean)	(redline)
b)	LM is advised to make disclosures if special rights for nominee/nomination rights and information rights are available to certain Promoters / Shareholders that would continue post listing and if yes, then details of the same may be clearly disclosed under section "History and Certain Corporate Matters".	Not applicable. None of the Promoters or Shareholders hold any nominee / nomination rights and information rights that will survive post listing of the Equity Shares on the Stock Exchanges.	227	299
c)	Further, LM shall specifically disclose that special rights, post listing shall be subject to approval of the Shareholders by way of a special resolution, in the first general meeting of the Company held post listing of the Equity Shares.	Not applicable. None of the Shareholders hold any special rights that will survive post listing of the Equity Shares on the Stock Exchanges.	227	299





S. No.	Observation		Response		RHP Page Number (clean)	RHP Page Number (redline)
d)	LM is advised to ensure that special rights which will continue post listing are not prejudicial or adverse to the interest of the minority / public shareholders.	We confirm that none of the Shareho the Stock Exchanges.	olders hold any special rights that w	vill survive post listing of the Equity Shares on	227	299
14. C	our Management					
(a)	LM is advised to disclose the reasons for resignation of the directors on the board, in the past 3 financial years.  Noted for compliance. We undertake to update the section titled "Our Management – Changes to our Board in the last three years" on page 218 of the DRHP, in the following manner in the UDRHP and RHP:  "Except as mentioned below, there have been no changes in our Board of Directors in the last three years:				238	310
		Name of director	Date of appointment / change in designation / cessation	Reason		
		Chhavi Lohariwala	September 27, 2019	Appointment as Director		
		Jayant Vasudeo Rao	February 28, 2020	Change in designation from Additional Director to Whole-time Director		
		Manoj Kumar Lohariwala	March 18, 2022	Change in designation from Whole-time Director to Chairman and Whole-time Director		
		Vinay Kumar Lohariwala	March 18, 2022	Change in designation from Whole-time Director to Managing Director		
		Chhavi Lohariwala	April 1, 2022	Cessation as Director due to personal and unavoidable circumstances		
		Gian Parkash Aggarwal	April 1. 2022	Cessation as Non- Executive Director due to personal and unavoidable circumstances		
		Anup Agarwal	April 1, 2022	Cessation as Non-Executive Independent Director due to personal and unavoidable circumstances		
		Pradosh Kumar	April 1, 2022	Cessation as Non-Executive Independent Director due to personal and unavoidable circumstances		





S. No.	Observation	Response			RHP Page Number (clean)	RHP Page Number (redline)
		Archit Aggarwal	April 1, 2022	Appointment as Non- Executive Director		
		Sudhir Kumar Bassi	April 1, 2022	Appointment as Non- Executive		
				Independent Director		
		Priyanka Dixit Sibal	April 1, 2022	Appointment as Non- Executive		
				Independent Director		
		Shirish Gundopant Belapure	April 1, 2022	Appointment as Non- Executive		
				Independent Director		
		Mahender Korthiwada	April 1, 2022	Appointment as Non- Executive		
		Note: This table does not include d		Independent Director		
		<b>Note:</b> This table does not include d	etalis of regularisations of ac	iditional Directors.		
15. F	inancial Information					
(a)	LM is further advised to disclose					
, ,	the reasons for the following in					
	the Management Discussion and					
	Analysis section:					
	i. Increase in Property, plant and	-		d "Management's Discussion and Analysis of Financial	-	-
	equipment, Right of use asset,	Condition and Results of Operation	s" to include the following:			
	goodwill as on March 31, 2021	"Destated energy dated are set	1.111.11.11			
	and December 31, 2021;	"Restated consolidated assets and	Ilabilities			
		Assets				
		Non-Current assets:				
		Property, plant and equipment				
		Our property, plant and equipmen	t increased from ₹763.59 m	illion as at March 31, 2021 to ₹1,544.13 million as at		
				ment is mainly due to ₹34.70 million (net) of additions		
		arising out of acquisition of UML as	of December 31, 2021. Furti	her, during the nine months ended December 31, 2021,		
				ent on account of capitalization of assets from capital		
		work in progress and ₹157.04 millio	on of other additions (net) m	ainly in land, building and plant and equipment.		





S. No.	Observation	Response	RHP Page Number (clean)	RHP Page Number (redline)
		Right-of-use assets		
		Our right-of-use assets increased from $\ 323.37$ million as at March 31, 2021 to $\ 92.07$ million as at December 31, 2021. The increase in right-of-use assets in property is mainly due to $\ 17.46$ million towards additions on account of acquisition of UML and further additions (net) of $\ 51.24$ million in the nine months ending December 31, 2021.		
		Capital work-in-progress		
		Capital work in progress decreased from ₹72.64 million as at March 31, 2021, to ₹3.21 million as at December 31, 2021, due to a transfer of assets from capital work-in-progress to plant and equipment, building, electrical equipment, lab equipment, furniture and fixtures, computer and printer and office equipment on account of capitalisation during the nine months ending December 31, 2021.		
		Goodwill		
		Our goodwill increased from NIL as at March 31, 2021 to ₹166.94 million as at December 31, 2021. Our goodwill as at December 31, 2021 is attributable to additions on account of acquisition of UML as a wholly owned subsidiary.		
		Other intangible assets		
		Our other intangible assets decreased from ₹4.44 million as at March 31, 2021 to ₹3.78 million as at December 31, 2021.		
		Financial assets		
		Our financial assets decreased from $\stackrel{?}{\sim}34.95$ million as at March 31, 2021 to $\stackrel{?}{\sim}5.61$ million as at December 31, 2021, which was mainly due to a reduction in security deposits by $\stackrel{?}{\sim}29.97$ million as at December 31, 2021 and an increase in bank-deposit accounts by $\stackrel{?}{\sim}0.63$ million as at December 31, 2021.		
		Income tax assets (net)		





S. No.	Observation	Response	RHP Page Number (clean)	RHP Page Number (redline)
		Our income tax assets (net) increased from ₹13.32 million as at March 31, 2021 to ₹43.14 million as at December 31, 2021, which was due to an increase in the amount of advance income tax and tax deducted at source (net of provision for income tax).		
		Other non-current assets		
		Other non-current assets decreased from ₹79.23 million as at March 31, 2021 to ₹15.41 million as at December 31, 2021 mainly due to reduction in capital advances by ₹65.42 million as at December 31, 2021 and an increase in prepaid expenses by ₹1.60 million as at December 31, 2021.		
		Current assets:		
		Inventories		
		Our inventories increased from $\[3914.45\]$ million as at March 31, 2021 to $\[31,447.47\]$ million as at December 31, 2021. Our inventories increased due to an increase in inventory of stock in trade by $\[3279.97\]$ million mainly on account of acquisition of UML by $\[3277.44\]$ million and other additions of $\[320.53\]$ million, and increases in inventory of raw materials by $\[341.44\]$ million, work in progress by $\[385.71\]$ million, packing material by $\[363.94\]$ million and stores and spares by $\[380.02\]$ million. Correspondingly, finished goods reduced by $\[380.02\]$ million.		
		Financial Assets		
		Our financial assets increased from ₹1,531.35 million as at March 31, 2021 to ₹2,081.16 million as at December 31, 2021 for the reasons set forth below.		
		<ul> <li>Trade receivables increased from ₹1,385.53 million as at March 31, 2021 to ₹2,006.82 million as at December 31, 2021, mainly due to acquisition of UML as of December 31, 2021. Also, the revenue from operations increased from ₹4,106.62 million in the year ended March 31, 2021 to ₹5,841.21 million in the nine months period ended December 31, 2021.</li> <li>Cash and cash equivalents decreased from ₹47.95 million as at March 31, 2021 to ₹7.69 million as at</li> </ul>		
		December 31, 2021. Bank balances other than cash & cash equivalents decreased from ₹70.99 million as at March 31, 2021 to ₹19.90 million as at December 31, 2021. The net decrease in bank balances / cash and cash equivalents as at December 31, 2021 was mainly due to payments made for acquisition of business on		





S. No.	Observation	Response	RHP Page Number	RHP Page Number
		<ul> <li>account of slump sale amounting to ₹542.50 million in respect of the Innova Partnership and payments made for acquisition of UML amounting to ₹50.00 million.</li> <li>Loans (which primarily constitutes loan to employees) marginally increased from ₹4.65 million as at March 31, 2021 to ₹4.69 million as at December 31, 2021.</li> <li>Other financial assets increased from ₹22.23 million as at March 31, 2021, to ₹42.06 million as at December 31, 2021. This increase was primarily due to an increase in security deposits by ₹13.02 million as at December 31, 2021, IPO expenses recoverable by ₹3.57 million as at December 31, 2021 and other recoverable by ₹9.46 million as at December 31, 2021, offset by reduction in interest accrued but not due on fixed deposits by ₹6.20 million as at December 31, 2021.</li> <li>Other current assets</li> </ul>	(clean)	(redline)
		Other current assets increased from ₹258.82 million as at March 31, 2021 to ₹340.59 million as at December 31, 2021 mainly due to increase as at December 31, 2021 in balance with government authorities by ₹63.13 million, prepaid expenses by ₹9.94 million, right to return asset by ₹9.31 million and advances to employees by ₹1.54 million. Further, advances to suppliers reduced by ₹2.15 million as at December 31, 2021."		
	ii. Decrease in capital work-in- progress, other financial assets, other non-current assets as on March 31, 2021 and December 31, 2021;	Noted for compliance. We undertake to amend the section titled "Management's Discussion and Analysis of Financial Condition and Results of Operations" as per our response to observation 15(a)(i) above.	-	-
	iii. Increase in trade receivable, loans, other financial assets, other current assets and decrease in bank balances as on March 31, 2021 and December 31, 2021;	Noted for compliance. We undertake to amend the section titled "Management's Discussion and Analysis of Financial Condition and Results of Operations" as per our response to observation 15(a)(i) above.	-	-
	iv. Increase in borrowings (current as well as non-current) and lease liabilities, provisions, deferred tax liabilities as on	Noted for compliance. We undertake to amend the section titled "Management's Discussion and Analysis of Financial Condition and Results of Operations" to include the following:  "Restated consolidated assets and liabilities	-	-





S. No.	Observation	Response	RHP Page Number (clean)	RHP Page Number (redline)
	March 31, 2021 and December 31, 2021;	[]	(accust)	(commo)
		Liabilities		
		Non-current liabilities:		
		Financial liabilities		
		Our financial liabilities increased from ₹63.53 million as at March 31, 2021 to ₹381.84 million as at December 31, 2021 for the reasons set forth below.		
		<ul> <li>Our borrowings increased from ₹60.00 million as at March 31, 2021 to ₹374.80 million as at December 31, 2021, due to an increase in term loans as at December 31, 2021.</li> </ul>		
		<ul> <li>Our lease liabilities increased from ₹3.53 million as at March 31, 2021, to ₹7.04 million as at December 31, 2021, mainly due to additions on account of the acquisition of UML and other additions during the year.</li> </ul>		
		Provisions		
		Our provisions increased from ₹12.34 million as at March 31, 2021 to ₹ 22.89 million as at December 31, 2021, due to an increase in provisions for employee benefits.		
		Deferred tax liabilities (net)		
		Our deferred tax liabilities increased from ₹19.26 million as at March 31, 2021 to ₹45.26 million as at December 31, 2021, mainly due to deferred tax impact of increase in excess depreciation as per Income Tax Act, 1961 over book values.		
		Other non-current liabilities		
		Our other non-current liabilities decreased from ₹1.29 million as at March 31, 2021 to ₹0.96 million as at December 31, 2021.		
		Current liabilities:		





S. No.	Observation	Response	RHP Page Number (clean)	RHP Page Number (redline)
		Financial liabilities  Our financial liabilities increased from ₹2,096.08 million as at March 31, 2021 to ₹3,137.39 million as at December 31, 2021 for the reasons set forth below.  • Our current borrowings increased from ₹390.26 million as at March 31, 2021 to ₹1,053.08 million as at December 31, 2021, due to an increase in cash credit limit from banks by ₹297.94 million as at December 31, 2021, an increase in current maturities of term loans by ₹47.00 million as at December 31, 2021 and increase in unsecured borrowings by ₹317.88 million as at December 31, 2021.  • Our lease liabilities increased from ₹1.18 million as at March 31, 2021 to ₹3.57 million as at December 31, 2021, mainly due to additions on account of the acquisition of UML and other additions during the year.  • Our trade payables increased from ₹1,122.33 million as at March 31, 2021 to ₹1,416.64 million as at December 31, 2021. This increase was primarily due to an increase in total outstanding dues of creditors other than micro and small enterprises from ₹1,087.51 million as at March 31, 2021 to ₹1,407.58 million as at December 31, 2021, which was primarily on account of acquisition of UML as on December 31, 2021.  • Our other financial liabilities increased from ₹582.31 million as at March 31, 2021, to ₹664.10 million as at December 31, 2021, mainly due to an increase in interest accrued but not due on borrowings by ₹5.53 million, employee related payables by ₹20.48 million, capital creditors by ₹20.98 million, security deposits by ₹27.30 million and amount payable on account of acquisition of UML by ₹550.00 million. The increase was offset by reduction in amount payable on acquisition of assets and liabilities of Innova Partnership on account of slump sale by ₹542.50 million.  Our other financial liabilities increased from ₹33.71 million as at March 31, 2020, to ₹582.31 million as at March 31, 2021, mainly due to increase in liability of amount payable for acquisition of business on account of the slump sale of the Innova Partnership amount	(clean)	(realine)
		Our other current liabilities increased from ₹50.11 million as at March 31, 2021 to ₹150.82 million as at December 31, 2021, mainly due to an increase in contract liabilities by ₹61.99 million, statutory dues by ₹5.53 million, refund liability by ₹12.09 million and deferred government grant by ₹21.10 million.		





S. No.	Observation	Response	RHP Page Number (clean)	RHP Page Number (redline)
		Provisions  Our provisions decreased from ₹5.34 million as at March 31, 2021 to ₹3.37 million as at December 31, 2021, mainly due to a reduction in the current portion of the provision for employee benefits.  Current tax liabilities (net)	(clean)	(rediffie)
		Our current tax liabilities (net) increased from NIL as at March 31, 2021 to ₹45.68 million as at December 31, 2021, mainly due to provision for income tax (net of advance tax)."		
	v. Increase in other financial liabilities as on March 31, 2021 and March 31, 2020;	Noted for compliance. We undertake to amend the section titled "Management's Discussion and Analysis of Financial Condition and Results of Operations" as per our response to observation 15(a)(iv) above.	-	-
	vi. Total outstandings dues	Noted for compliance. We undertake to amend the section titled "Management's Discussion and Analysis of Financial Condition and Results of Operations" as per our response to observation 15(a)(iv) above.	-	-
	vii.Analysis of various rations	Complied with and noted for compliance.  A detailed analysis of various ratios applicable to the Company's business are disclosed on a restated consolidated basis and proforma consolidated basis in the section entitled "Management's Discussion and Analysis of Financial Condition and Results of Operations – Key Performance Indicators and Non-GAAP Financial Measures" beginning on page 349 of the DRHP.	372	373
(b)	LM is also advised to disclose the auditor qualification, if any, in the offer document and the Issuer's management comments on the same.	Complied with and noted for compliance.  As disclosed on page 407 of the DRHP, there have been no reservations or qualifications or adverse remarks of the Statutory Auditors in the nine months ended December 31, 2021, and in Fiscals 2021, 2020 and 2019.	420	454
16. R	Related Party Transactions (RPTs)			
(a)	i. to quantify the RPTs, in brief, that the company has entered into in tabular form; and	Complied with and noted for compliance.  A summary of the related party transactions that the Company has entered into, as derived from the Restated Consolidated Financial Information, in the nine months ended December 31, 2021 and in Fiscals 2021, 2020 and 2019,	28	34





S. No.	Observation	Response	RHP Page Number (clean)	RHP Page Number (redline)
	ii. to disclose the RPTs as a percentage of revenue in the offer document	including the quantification of the related party transactions, has been included in the section titled "Summary of the Offer Document – Summary of related party transactions" on page 29 of the DRHP.  Further, we undertake to include the following statement in the section titled "Summary of the Offer Document – Summary of related party transactions" in the UDRHP and RHP, disclosing details of the related party transactions set out in the table described above as a percentage of revenue from operations:  "The related party transactions covered under our profit and loss account as a percentage of our revenue from		
		operations constituted 32.05%, 29.05%, 27.01% and 20.75% for the nine months ended December 31, 2021, and the financial years ended March 31, 2021, March 31, 2020 and March 31, 2019."		
	inancial Indebtedness			
(a)	LM is advised to disclose the outstanding amount / default in making payment obligation for the last three years.	Noted for compliance to the extent applicable. We undertake to include the following disclosure in the section titled "Financial Indebtedness" in the UDRHP and RHP:  "As on March 31, 2021, March 31, 2020 and March 31, 2019, our total borrowings as per our Restated Consolidated Financial Information were ₹450.26 million, ₹535.12 million, ₹720.54 million."  Further, we confirm that the Company has not defaulted in making payment obligations owed to its lenders in the last	422	457
		three years. A negative confirmation to this effect has been included in the section titled "History and Certain Corporate Matters - Defaults or rescheduling/restructuring of borrowings with financial institutions/banks" on page 206 of the DRHP.		
(b)	i. Name of the lenders ii. Any restrictive covenants entered into with the lenders iii. Instance of past default iv. Whether all the loans are secured or unsecured v. Whether any approval	<ul> <li>(i) We undertake to include the names of the lenders of the Company as on the date of the RHP, in the section "Financial Indebtedness" of the RHP.</li> <li>(ii) An indicative list of the restrictive covenants applicable to the Company and its Subsidiary, UML, under the documentation entered into with their lenders has been included in the section "Financial Indebtedness" on page 410 of the DRHP.</li> <li>(iii) A negative confirmation to this effect has been included in the section titled "History and Certain Corporate"</li> </ul>	422	457
	required from the lenders for the proposed IPO	<ul> <li>(iii) A negative commutation to this effect has been included in the section titled "instity und certain corporate Matters - Defaults or rescheduling/restructuring of borrowings with financial institutions/banks" on page 206 of the DRHP.</li> <li>(iv) Details of the total secured and unsecured outstanding borrowings of the Company on a consolidated basis as on May 15, 2022, have been disclosed in the section titled "Financial Indebtedness" on page 410 of the DRHP.</li> </ul>		





S. No.	Observation	Response	RHP Page Number (clean)	RHP Page Number (redline)
		(v) As disclosed on page 410 of the DRHP, the Company has obtained the necessary consents required under the relevant loan documentation for undertaking activities in relation to the Offer.		
(c)	LM is advised to disclose if consent from all the lenders, in writing, has been obtained for the proposed IPO.	Complied with. Please see our response to 17(b)(v) above.	422	457
(d)	LM is advised to provide financial statements for the year ended March 31,2022.	Noted for compliance. The restated consolidated financial statements for the year ended March 31, 2022, will be included in the UDRHP and RHP.	260	334
18. L	egal and other information			
(a)	LM is advised to provide full details of pending lawsuits against the Company and its subsidiaries and associates, promoters/promoter group, directors and LM is advised to quantify the financial impact of the same, wherever possible.	Complied with and noted for compliance, to the extent applicable.  In accordance with the SEBI ICDR Regulations, details of: (i) all outstanding criminal litigation involving the Company, its Subsidiaries, its Directors and its Promoters; (ii) all outstanding actions by regulatory authorities and statutory authorities involving the Company, its Subsidiaries, its Directors and its Promoters; (iii) disciplinary action including penalty imposed by SEBI or stock exchanges against the Promoters in the last five financial years including outstanding action; (iv) claims related to direct and indirect taxes involving the Company, its Subsidiaries, its Directors and its Promoters, in a consolidated manner, giving the number of cases and total amount, as well as summary disclosures for those tax proceedings exceeding the materiality threshold as per the materiality policy adopted by the board of directors of the Company on June 19, 2022; and (v) other pending litigation involving the Company, its Subsidiaries, its Directors and its Promoters, as per the materiality policy adopted by the board of directors of the Company on June 19, 2022, have been disclosed in the section "Outstanding Litigation and Other Material Developments" beginning on page 412 of the DRHP.	424	460
		The UDRHP and RHP will be updated with any further developments in such matters and to include further matters that may have been initiated since the date of filing of the DRHP, if any.  Further, in terms of the SEBI ICDR Regulations, there is no requirement for the Company to disclose details of any litigation involving associates and members of the Promoter Group. Accordingly, no disclosures for litigation involving associates and members of the Promoter Group or are sought to be made. Further, the Company does not have any associates.		





S.	Observation	Response	RHP Page	RHP Page
No.			Number (clean)	Number (redline)
(b)	LM shall update the details of status of litigation with the	Noted for compliance, to the extent applicable.	-	-
	latest/updated position of litigations against	Please note that in terms of the SEBI ICDR Regulations, there is no requirement for the Company to disclose details of any litigation involving members of the Promoter Group or companies promoted by the Company. Accordingly, no		
	promoter/promoter group entities/company and the companies promoted by the	disclosures for litigation involving members of the Promoter Group or companies promoted by the Company are sought to be made.		
	issuer.	With respect to litigation against the Promoters and the Company, please see our response to 18(a) above.		
(c)	LM is advised to confirm that the existing litigations are not so major that the issuer's survival is dependent on the outcome of the pending litigation.	We confirm that the existing litigations involving the Company are not so major that the Company's survival is dependent on the outcome of the pending litigation.	-	-
(d)	LM is advised to ensure the disclosures of all actions taken by statutory and regulatory authorities.	Complied with. Details of all pending actions by regulatory authorities and statutory authorities involving the Company, its Directors and its Promoters, have been disclosed in the section "Outstanding Litigation and Other Material Developments" beginning on page 412 of the DRHP. Further, as on the date of the DRHP, there were no pending actions initiated by statutory or regulatory authorities against the Subsidiaries, as has been disclosed on page 415 of the DRHP. The UDRHP and RHP will be updated with any further developments in such matters and to include further pending regulatory or statutory proceedings that may have been initiated against the Company, Directors, Promoters and Subsidiaries since the date of filing of the DRHP, if any.	424	460
(e)	LM is advised to disclose the current status of outstanding dues to creditors and its impact.	Noted for compliance. In compliance with Clause 12(A)(2) of Part I of Schedule VI of the SEBI ICDR Regulations, details of outstanding dues to creditors of the Company (i.e., its material creditors, MSMEs and other creditors), including amounts due, as disclosed in the section titled "Outstanding Litigation and Material Developments" on pages 419 and 420 of the DRHP will be updated in the UDRHP and RHP.	439	477
19. N	/liscellaneous	·		
(a)	Any risk on pricing of the issue / basis of issue price, track record of BRLMs, average cost of acquisition of shares by selling shareholders etc. which is proposed to be included in the issue advertisement may be submitted.	Noted for compliance.	_	-





S.	Observation	Response	RHP Page	RHP Page
No.		·	Number	Number
			(clean)	(redline)
(b)	LM is advised to ensure following	Noted for compliance.	-	-
	disclosures in the Issue			
	advertisement for announcement			
	of Price Band and all further			
	advertisements as a box item			
	below the price band:			
	"Risks to Investors:			
	a)The [to be disclosed] Merchant			
	Bankers associated with the			
	issue have handled [to be			
	disclosed] public issues in the			
	past three years out of which			
	[to be disclosed] issues closed			
	below the issue price on listing			
	date."			
	b) Any adverse data/ noting in the			
	basis for issue price should be			
	disclosed. For example:			
	<ul> <li>"The Price/Earnings ratio</li> </ul>			
	based on diluted EPS for			
	[latest full financial year] for			
	the issuer at the upper end			
	of the Price band is as high			
	as [to be disclosed] as			
	compared to the average			
	industry peer group PE ratio			
	of [to be disclosed]."			
	<ul> <li>"Average cost of acquisition</li> </ul>			
	of equity shares for the			
	selling shareholders in IPO			
	is [to be disclosed] and offer			
	price at upper end of the			





S.	Observation	Response	RHP Page	RHP Page
No.			Number	Number
			(clean)	(redline)
	price band is [to be			
	disclosed]."			
	<ul> <li>"Weighted Average Return</li> </ul>			
	on Net Worth for [last three			
	full financial years] is [to be			
	disclosed] %."			
	The data on above disclosures			
	shall be updated and disclosed			
	prominently (in the same font size			
	as the price band) in			
	advertisements of Price Band and			
	all further advertisements,			
	website of the company and the			
	stock exchange. Further, any			
	adverse ratio / data in basis for			
	issue price should also be			
	disclosed. LM shall submit the			
	draft advertisement for			
	announcement of Price Band with			
	SEBI before its publication in the			
	newspapers for our comments			





## **SCHEDULE I**

# Para wise compliance with the Securities and Exchange Board of India (Framework for Rejection of Draft Offer Documents) Order, 2012

S.	Rejection Criteria	Response
No.		·
1.1	Where Capital Structure involves any of the following	
(i)	Existence of circular transactions for building up the capital / net worth of the issuer.	Not applicable
(ii)	Ultimate promoters are unidentifiable.	Not applicable
(iii)	Promoters' contribution not complying with SEBI (Issue of Capital and Disclosure Requirements) Regulations, 2018 in letter or in spirit.	Not applicable
1.2	Where Object of the Issue	
(i)	Is vague for which a major portion of the issue proceeds are proposed to be utilized.	Not applicable
(ii)	Is repayment of loan or inter corporate deposit or any other borrowing of similar nature, and the issuer is not in a position to disclose the ultimate purpose for which the loan was taken or demonstrate utilization of the same for the disclosed purpose.	Not applicable
(iii)	Is such where the major portion of the issue proceeds is proposed to be utilized for the purpose which does not create any tangible asset for the issuer, such as, expenses towards brand building, advertisement, payment to consultants, etc., and there is not enough justification for creation of such assets in terms of past performance, experience and concrete business plan of the issuer.	Not applicable
(iv)	Is to set up a plant and the issuer has not received crucial clearances / licenses / permissions / approvals from the required competent authority which is necessary for commencement of the activity and because of such non-receipt of clearances / licenses / permissions / approvals, the issue proceeds might not be utilized towards the stated objects of the issue.	Not applicable
(v)	Is such where the time gap between raising the funds and proposed utilization of the same is unreasonably long.	Not applicable
1.3	Where business model of an issuer is	
	Exaggerated, complex or misleading and the investors may not be able to assess the risks associated with such business models.	Not applicable
1.4	Where scrutiny of Financial Statements shows	
(i)	Sudden spurt in the business just before filing the draft offer document and reply to clarifications sought is not satisfactory. This will include spurt in line items such as Income, Debtors/Creditors, intangible assets, etc.	Not applicable
(ii)	Qualified audit reports or the reports where auditors have raised doubts / concerns over the accounting policies. This would also be applicable for the subsidiaries, joint ventures and associate companies of the issuer which significantly contributes to the business of the issuer. This would also be applicable for the entities where the issue proceeds are proposed to be utilized.	Not applicable
(iii)	Change in accounting policy with a view to show enhanced prospects for the issuer in contradiction with accounting norms.	Not applicable
(iv)	Majority of the business is with related parties or where circular transactions with connected / group entities exist with a view to show enhanced prospects of the issuer.	Not applicable
1.5	Where there exists litigation including regulatory action	





S.	Rejection Criteria	Response
No.		
(i)	Which is so major that the issuer's survival is dependent on the	Not applicable
	outcome of the pending litigation.	
(ii)	Which is wilfully concealed or covered.	Not applicable
1.6	Other General Criteria	
(i)	Failure to provide complete documentation in terms of requirements	Not applicable
	of SEBI (Issue of Capital and Disclosure Requirements) Regulations,	
	2018.	
(ii)	Non-furnishing of information or delay in furnishing of information	Not applicable
	or furnishing of incorrect / vague / misleading / incomplete / false /	
	non satisfactory information to the Board.	
(iii)	Failure to resolve conflict of interest, whether direct or indirect,	Not applicable
	between the issuer and Merchant Banker appointed by the issuer to	
	undertake the book building process. Quantification of conflict of	
	interest may not always be possible but it would largely depend upon	
	the Board's assessment on whether such conflict of interest may	
	affect the judgment and ability of the Merchant Banker in conducting	
	due diligence activity of issuer.	





## **SCHEDULE II**

# Para wise compliance of the Securities and Exchange Board of India (Issuing Observations on Draft Offer Documents Pending Regulatory Actions) Order, 2020

S.	Rejection Criteria	Response
No.		
A.	Treatment where there is a probable cause for investigation or enquiry or when a enquiry is in progress against the entities	
1.	Where there is a probable cause for investigation, examination or enquiry against the entities, the observations on the draft offer document filed by the issuer with the Board shall be kept in abeyance for a period of thirty days after such probable cause arises or the date of filing of the draft offer document with the Board, whichever is later.	Not applicable
2.	Where the Board is unable to conclude such investigation, examination or enquiry against the entities due to the reasons beyond its control or due to the conduct of the parties other than the entities, the observations on the draft offer document shall be kept in abeyance for a further period of thirty days.	Not applicable
3.	Where the Board is unable to conclude such investigation, examination or enquiry against the entities due to the conduct of the entities, the observations on the draft offer document shall be kept in abeyance till the time such investigation, examination or enquiry is concluded.	Not applicable
В.	Treatment where show cause notice has been issued.	Nick coult cold
1.	Where a show cause notice has been issued to the entities in an adjudication proceeding, the Board may process the draft offer document and issue observations and advice the entities to make necessary disclosures and statements in respect of such proceedings and the possible adverse impact of an order on the entities, in the offer document.	Not applicable
2.	Where a show cause notice has been issued in respect of proceedings under subsection (4) of section 11 or section 11B(1), the Board shall keep in abeyance the issuance of observations for a period of ninety days from the date of filing of the draft offer document with the Board.	Not applicable
3.	Where the Board is unable to conclude the proceedings as referred to sub-clause (2) due to the reasons beyond its control or due to the conduct of the parties other than the entities, the observations on the draft offer document shall be kept in abeyance for a further period of forty five days.	Not applicable
4.	Where the Board is unable to conclude the proceedings as referred to sub-clause (2) due to the conduct of the entities, the observations on the draft offer document shall be kept in abeyance till the time such proceedings are concluded.	Not applicable
5.	Where no order is passed within the time period specified in clause (3), the Board may process the draft offer document and issue observations and advise the entities to make necessary disclosures and statements in respect of such proceedings and the possible adverse impact of an order on the entities, in the offer document.	Not applicable
C.	Treatment where recovery proceedings have been initiated or an order for disgorge penalty has not been complied with or in case of non-compliance with any direct Board.	
1.	Where the Board has initiated proceedings for recovery against the entities or when an order for disgorgement or monetary penalty passed against the entities is not complied with or in case of non-compliance with any direction issued by the Board, the observations on the draft offer document filed by the issuer with the Board shall be kept in abeyance till such proceedings are concluded or until the directions are complied with.	Not applicable
D.	Reconsideration of proceedings pursuant to remand by the Securities Appellate Trib	
1.	Where proceedings has been remanded by the Securities Appellate Tribunal or a court, the same shall in effect be treated as proceedings covered under this Order,	Not applicable





S.	Rejection Criteria	Response
No.		
	and the Board may take appropriate action in respect of the draft offer document under the provisions of this general order, subject to any order passed by the Securities Appellate Tribunal or a court, as the case may be, while remanding the matter.	
E.	Issuance of observations when the issuer is restrained by a court from making a pub offer document.	lic issue or filing of
1.	Where the issuer has been restrained by a court or tribunal from making an issue of securities or from issuing offer document to the public, the Board may examine the offer document and issue its observations thereof with a qualification that said observations are issued in accordance with the regulatory powers conferred on the Board and that the public issue or issuance of the offer document to the public by the issuer shall be subject to the orders of such court or tribunal or authority.	Not applicable





# Schedule B

# Compliance with in-seriatim response to the Additional Observations

S. No.	Observation	Response	RHP Page Number (Clean)	RHP Page Number (Redline)
Gene	eral Clarifications:			
1.	SEBI circular SEBI/HO/CFD/SSEP/CIR/P/2022/14 dated February 04, 2022, inter-alia, specifies that the issuer company/ MBs shall insert a Quick Response (QR) code on the front page of the documents such as front outside cover page, abridged prospectus, price band advertisement, etc. as deemed fit by them. The scan of QR code shall lead to downloading of prospectus, abridged prospectus and price band advertisement as applicable. It is observed that QR provided in DRHP does not provide a direct access to DRHP of the company. LM is advised to re-assess the compliance of the aforesaid clause and submit an undertaking that the particular clause of the circular is duly complied with.	Complied with.  The quick response (QR) code provided on the outside cover page of the DRHP leads to a general disclaimer page, and accepting the disclaimer leads to a webpage on the website of the left lead BRLM on which the DRHP, RHP, Prospectus, abridged prospectus, price band advertisement etc. will be available for downloading. Request you to please note that the disclaimer page is important to draw attention to the fact the information contained on the webpage is not intended to be viewed by persons not resident in India.	Cover Page	Cover Page
2.	LM is advised to ensure that at all applicable places, where data points are being compared, the same is disclosed in a tabular format and updated as on March 31, 2022.	Noted for compliance to the extent possible.	-	-





S. No.	Obse	rvation		Response					RHP Page Number (Redline)	
Othe	r Financial Informa	ition:								
3.	LM is advised to	orovide ke	y ratios in	The key ratios of the Company, along with peer comparisons, for F	Fiscal 2021 have bee	en disclosed on page 16	5 of	177	239	
	tabular form	along w	ith peer	the DRHP. The table will be updated to reflect the key ratios of the		with peer comparisons,	for			
	comparison for th	e past 3 y	ears.	Fiscal 2022 in the updated draft red herring prospectus ("UDRHP")	) and RHP.					
Obje	cts of the Offer									
4.	LM is advised amount of fres utilized for repayr Company in the fo	sh issuan ment of lo	ce being ans of the	Details of the borrowings of the Company as on May 15, 2022, rais and more than 12 months prior to the filing of the DRHP, net of the out below:	•			-	-	
				Particulars of utilisation	Amount raised 0-12	Amount raised more				
	Particulars of	Amount	Amount		months prior to	than 12 months prior				
	utilisation	raised	raised		filing of DRHP (₹	to filing of DRHP (₹				
		0-12	more		million) net of the	million) net of the				
		months	than 12		amount repaid in	amount repaid in this				
		prior to	months		this period	period				
		filing of	prior to	Working capital	1,006.12	357.18				
		DRHP	filing of DRHP	Expansion of existing plant production capacity	-51.94	102.61				
		(₹ million)	ORHP (₹ million)	Setting up of new block within the existing unit and reimbursement of capital expenditure done through own sources	-15.30	200.00				
	Working capital Expansion of			Purchase of land and construction of corporate office and R&D centre	70.00	-				
	existing plant			Acquisition of capital equipment	49.64	139.10				
	production capacity									
	Setting up of									
	new block									
	within the									
	existing unit									
	Reimbursement									
	of capital									





S. No.	Observation	Response	RHP Page Number	RHP Page Number
			(Clean)	(Redline)
	expenditure			
	done through			
	own sources			
	Purchase of			
	land and			
	construction of			
	corporate office			
	and R&D centre			
	Acquisition of			
	capital			
	equipment			
5.	LM is advised to include loan sanction	Noted for compliance. The table providing details of certain borrowings availed by the Company as on May 15,	106	142
	date in the table providing details of	ate in the table providing details of 2022, on page 113 of the DRHP, will be updated in the UDRHP and RHP in the manner set out in Schedule I. We		
	certain borrowings availed by our	further undertake to update the information in this table to a more recent date in the UDRHP and RHP.		
	Company as on May 15, 2022 at page			
	no. 113.			
_	Factors:			
6.	LM is advised to include a risk factor that Innova Partnership and UML	Noted for compliance. We will add the following risk factor as risk factor 51.	56	77
	were owned by promoters and disclose amount paid for such acquisitions.	"51. Our Company acquired the assets and liabilities of the business of the Innova Partnership and acquired		
		Effective as of March 31, 2021, our Company acquired the assets and liabilities of the business of Innova Partnership as a going concern through a slump sale from persons including our Promoters. The total consideration paid by our Company for the business of Innova Partnership was ₹542.50 million. Effective as of December 31, 2021, our Company acquired UML as a wholly-owned subsidiary from persons including our Promoters for a total purchase consideration of ₹600.00 million. We acquired the assets and liabilities of the Innova Partnership and		
		acquired UML (both acquisitions, the "Acquisition Transactions") to take advantage of the manufacturing and economic synergies with our Company. For further details, please see "History and Certain Corporate Matters – Details regarding material acquisition or divestment of business or undertakings in the last 10 years" on page 208.		
		However, we may not be able to effectively integrate the businesses that we acquire or we may experience difficulties arising from coordinating and consolidating corporate and administrative functions, including		





S. No.	Observation	Response		RHP Page Number (Redline)
		integration of internal controls and procedures with our ongoing operations. A failure to successfully integrate an acquired business or inability to realize the anticipated benefits of acquisition could adversely affect our results of operations and financial condition"		
7.	LM is advised to disclose a separate risk factor in top 10 that more than 90% of the raw materials are imported from once country (i.e. China)	Noted for compliance. We will add the following risk factor as risk factor 56.  "56. In Fiscal 2019, Fiscal 2020 and Fiscal 2021 and in the nine months ended December 31, 2021, imported raw materials from China, China SEZ and Hong Kong on a restated consolidated basis as a percentage of our cost of imported raw materials were 100.00%, 100.00%, 91.85% and 91.80%, respectively and as a percentage of total raw materials purchases were 3.27%, 7.24%, 8.90% and 13.10%, respectively. Our dependence on China, China SEZ and Hong Kong for our raw material supplies exposes us to political, economic and social conditions in greater China.	41	53
		In Fiscal 2019, Fiscal 2020 and Fiscal 2021 and in the nine months ended December 31, 2021, imported raw materials from China, China SEZ and Hong Kong on a restated consolidated basis as a percentage of our cost of imported raw materials were 100.00%, 100.00%, 91.85% and 91.80%, respectively, and as a percentage of total raw materials purchases were 3.27%, 7.24%, 8.90% and 13.10%,, respectively. In Fiscal 2019, Fiscal 2020 and Fiscal 2021 and in the nine months ended December 31, 2021, imported raw materials from China, China SEZ and Hong Kong on a proforma consolidated basis as a percentage of our cost of imported raw materials were 100.00%, 100.00%, 95.86% and 91.80%, respectively, and as a percentage of total raw materials purchases were 2.34%, 10.34%, 13.11% and 13.10%, respectively. Our dependence on China China, China SEZ and Hong Kong for our raw material supplies exposes us to political, economic and social conditions in greater China. Further, our raw material suppliers may be adversely impacted by the economic downturn in their national or regional economies, disruption in their banking and financial systems, economic weakness, unfavourable government policies, rising inflation, lowering of spending power and customer confidence, and political uncertainty."		
8.	LM is advised to include risk factor 6 as risk factor 2.	Noted for compliance. We will include risk factor 6 as risk factor 2.	67	94
9.	LM is advised to include risk factor 18 in the top 10 risk factors.	Noted for compliance. We will include risk factor 18 in the top 10 risk factors.	49	66
10.	LM is advised to include separate risk factor in top 15 regarding that nearly 15% of revenue is from related parties. LM is also advised to disclose that UML and Azine Healthcare	Noted for compliance. We will include the following risk factor in the top 15 risk factors:  "15. We derive a significant portion of our revenue from operations on a restated consolidated basis from related party transactions. Azine Healthcare Private Limited and UML (which is now a wholly owned subsidiary	44	56





S. No.	Observation		Re	sponse				RHP Page Number (Clean)	RHP Page Number (Redline)
	undertake marketing and sale of finished pharmaceutical products of the Company and revenue from their operations is considered as related party transaction.	In the nine months ended a derived from the Restated consolidated basis were 32 UML (which is now a which pharmaceutical products of transactions. Related part UML, respectively, in the notes of the second	Company) undertake marketing and sale of finished pharmaceutical products of the Company and the from their operations are considered as related party transactions.  In the Restated December 31, 2021 and Fiscals 2021, 2020 and 2019 our related party transactions as a from the Restated Consolidated Financials as a percentage of revenue from operations on a restated dated basis were 32.05%, 29.05%, 27.01% and 20.75%, respectively. Azine Healthcare Private Limited and which is now a wholly owned subsidiary of the Company) undertake marketing and sale of finished acceutical products of the Company and revenue from their operations are considered as related party citions. Related party transactions entered into by our Company and Azine Healthcare Private Limited and despectively, in the nine months ended December 31, 2021 and Fiscals 2021, 2020 and 2019 as per Ind AS ived from the Restated Consolidated Financial Information is detailed below.						
		Nature of transaction	Name of the related party	For the period ended December 31, 2021	For the year ended March 31, 2021	For the year ended March 31, 2020	For the year ended March 31, 2019		
		Revenue from operations	Univentis Medicare Limited	809.07	674.30	509.57	304.87		
		(net of returns)  Revenue from operations (net of returns)	Azine Healthcare Private Limited	18.28	5.84	2.74	12.47		
11.	Risk factor 6: LM is advised to disclose market share using revenue on a restated consolidated basis.	"6. We operate in a mark manufacturing services of companies in India and or products of other supplier.  We compete to provide services and products in manufacturers focusing on	inplied with and noted for compliance. We undertake to amend Risk Factor 6 in the UDRHP and RHP as set forth low. Amendments also include changes as set forth in our response letter dated August 3, 2022.  We operate in a market that is highly competitive. We compete to provide outsourced pharmaceutical anufacturing services or CDMO services and products, particularly for formulations, to pharmaceutical impanies in India and other jurisdictions. In addition, our branded generic products compete with generic inducts of other suppliers in India and other jurisdictions.  Compete to provide services to pharmaceutical companies in the CDMO industry. Our competition in the CDMO vices and products includes full-service pharmaceutical outsourcing or CDMO companies; contract mufacturers focusing on a limited number of dosage forms; contract manufacturers providing multiple dosage ins; and large pharmaceutical companies offering third-party manufacturing services to fill their excess capacity.					34	42





S. No.	Observation		Response				
		with 300 to 400 organised players and approxima characterized by high fragmentation and compet As a result, the bargaining power of contract manu CRISIL Report, May 2022).	The domestic formulations industry is highly fragmented in terms of both, number of manufacturers and products, with 300 to 400 organised players and approximately 15,000 unorganised players. Contract manufacturing is also characterized by high fragmentation and competition, with large number of organized and unorganized players. As a result, the bargaining power of contract manufacturing players is lowered owing to high competition. (Source: CRISIL Report, May 2022).  The following table sets forth the key players across the Indian CDMO industry for the Fiscal 2021.				
		Company Name	Date of Incorporation	Registered office location			
		Acme Formulation Private Limited	2004	Himachal Pradesh			
		Akums Drugs and Parmaceuticals Ltd	2004	Delhi			
		Innova Captab Limited	2005	Mumbai			
		Synokem Pharmaceuticals Limited	1983	Delhi			
		Theon Pharmaceuticals Limited	2005	Chandigarh			
		Tirupati Medicare Ltd	2005	Delhi			
		Windlas Biotech Ltd	2001	Dehradun			
		Note: The list of competitors is an indicative list and no Sources: MCA, company websites and filings, CRISIL Res (Source: CRISIL Report).  In addition, in Europe and Asia, there are a large that serve only their local or national markets. Als portions of their manufacturing capacity, and an industry. (Source: CRISIL Report, May 2022). We existing product portfolio and novelty of new off stability), service (on-time delivery and manufact may, among other things, result in a decrease in the pharmaceutical development and manufacturing business, results of operations and financial conditional control of the control	rearch  I number of privately owned, I nage pharmaceutical comply such divested businesses m I compete primarily on the borerings), of supply (quality, reguring flexibility) and cost-effecte fees paid for our services are services, which could have tion.	panies have been seeking to divest ay increase competition in CDMO asis of product portfolio (range of gulatory compliance and financial ective manufacturing. Competition and reduced demand for outsourced a material adverse effect on our our revenue from sale of goods and			





S. No.	Observation	Response				RHP Page Number (Clean)	RHP Page Number (Redline)
		Export Country (1)		Nine Months er December 31, 2			
		Kenya			17.27%		
		Venezuela			12.92%		
		Sri Lanka			12.73%		
		Tanzania			11.66%		
		Uganda			9.14%		
		<u>Ethiopia</u>			8.79%		
		Ghana			8.04%		
		Nigeria			5.94%		
		Myanmar			5.41%		
		Republic of the Congo Total			1.42% <b>93.32%</b>		
		(1) The top ten export countries provid	dad are our ton ton owners	auntries in terms of revenue			
		Our cost of raw materials from our top i a restated consolidated basis are set for		onth period ended Decen	nber 31, 2021.		
		Import Country (1)		Nine Months e December 31, 2			
		China			20.29%		
		China-SEZ			16.63%		
		Hong Kong			54.88%		
		Netherlands			8.20%		
		Total			100.00%		
		The following table indicates market size of the global CDMO market and the global CDMO formulation market and our share of the global CDMO formulation market on a restated consolidated basis for the periods indicated.					
		Particulars Particulars	(in ₹ billion, except percentages)  Particulars Fiscal 2019 Fiscal 2020 Fiscal 2021				
			1.500. 2015	7.550.7 2020	7.5547.2522		
		Market size of Global CDMO market (API+ Formulation)	6,874.92	7,315.87	8,186.63		





S. No.	Observation		RHP Page Number (Clean)	RHP Page Number (Redline)					
		Market size of Global CDMO Formulation							
		market	1,230.49	1,330.32	1,525.36				
		Revenue from CDMO services for	2.24	2.42	2.74				
		Innova Captab Limited (1) Market share of Innova Captab Limited in	3.31	3.43	3.71				
		the global CDMO formulation market	0.27%	0.26%	0.24%				
		(1) Market share arrived at using reve (Source: CRISIL Report).  The following table indicates market siz and our share of the Indian CDMO forms							
		(in ₹ billion, except percentages)							
		Particulars	Fiscal 2019	Fiscal 2020	Fiscal 2021				
		Market size of Global CDMO market (API+ Formulation) (1)	821.14	898.44	1,014.08				
		Market size of Global CDMO Formulation market	416.03	449.22	512.11				
		Revenue from CDMO services for Innova Captab Limited (2)	3.31	3.43	3.71				
		Market share of Innova Captab Limited in the global CDMO formulation market	0.80%	0.76%	0.72%				
		<ul> <li>(1) Include domestic and export operations</li> <li>(2) Market share arrived at using revenue on a restated consolidated basis.</li> <li>(Source: CRISIL Report).</li> <li>The following table indicates market size of the global CDMO market and the global CDMO formulation market and our share of the global CDMO formulation market on a proforma consolidated basis for the periods indicated.</li> <li>(in ₹ billion, except percentages)</li> </ul>							
		Particulars	Fiscal 2019	Fiscal 2020	Fiscal 2021				
					7.000, 2022				
		Market size of Global CDMO market (API+ Formulation)	6,874.92	7,315.87	8,186.63				





S. No.	Observation		RHP Page Number (Clean)	RHP Page Number (Redline)							
		Market size of Global CDMO Formulation market	1,230.49	1,330.32	1,525.36						
		Revenue from CDMO services for Innova Captab Limited (1)	3.95	3.91	4.05						
		Market share of Innova Captab Limited in the global CDMO formulation market	0.32%	0.29%	0.27%						
		<ul> <li>(1) Market share arrived at using revenue on a proforma consolidated basis.</li> <li>(Source: CRISIL Report).</li> <li>The following table indicates market size of the Indian CDMO market and the Indian CDMO formulation market and our share of the Indian CDMO formulation market on a proforma consolidated basis for the periods indicated</li> </ul>									
			f billion, except percentages)								
		Particulars	Fiscal 2019	Fiscal 2020	Fiscal 2021						
		Market size of Global CDMO market (API+ Formulation) (1)	821.14	898.44	1,014.08						
		Market size of Global CDMO Formulation market	416.03	449.22	512.11						
		Revenue from CDMO services for Innova Captab Limited (2)	3.95	3.91	4.05						
		Market share of Innova Captab Limited in the global CDMO formulation market	0.95%	0.87%	0.79%						
		(1) Include domestic and export open (2) Market share arrived at using reve (Source: CRISIL Report).									
		For our domestic branded generics be therapeutic product categories, and with									
		pharmaceutical players are adding gene	= : :								
		in international markets, we compete w	•	•	•						
		emerging markets that are engaged in grow our international business, we exp	, ,		·						





		Number (Clean)	Number (Redline)
	Some of our competitors may have substantially greater financial, marketing, technical or other resources than we do. Greater financial, marketing, technical or other resources may allow our competitors to respond to changes in market demand faster with new, alternative or emerging technologies. If our competitors gain significant market share at our expense, our business, results of operations and financial condition could be adversely affected. Changes in the nature or extent of our customer requirements may render our service and product offerings obsolete or non-competitive, which could have a material adverse effect on our business, results of operations and financial condition."		
Risk factor 9: LM is advised to include table of attrition rate as given in reply to RF22. Further, LM is advised to explain reason for high attrition. LM is also advised to clarify reason for non-availability of R& D expenditure of the Company and how the company is going to estimate the same.	Complied with and noted for compliance. We undertake to amend Risk Factor 9 in the UDRHP and RHP as set forth below. Amendments also include changes as set forth in our response letter dated August 3, 2022. Our R&D expenditure as set forth in our amendments below has been prepared on the basis of expenditure incurred on R&D department under the head of salaries, remuneration, power & electricity, material consumption, clinical trial expenses and product registration and licence fees.  "9. We are dependent on our R&D activities for our future success. If we do not successfully develop new products or continue our generic product portfolio expansion in a timely and cost-effective manner, our business, results of operations and financial condition may be adversely affected.  The pharmaceutical and healthcare industry is characterised by technological advancements, introduction of	47	59
	innovative products, price fluctuations and intense competition. We have invested substantial effort, funds and other resources towards our R&D activities. We have a dedicated R&D laboratory and pilot equipment located at our manufacturing facility at Baddi, Himachal Pradesh. We are increasingly engaged in R&D activities to develop various generic products, manufacturing processes and technologies for diverse therapeutic segments. In particular, our R&D laboratory is focused on developing efficient processes for the manufacture of upcoming patent expired products. In the nine months ended December 31, 2021 and Fiscal 2021, our R&D expenditure on a restated consolidated basis was ₹38.51 million and ₹21.45 million, respectively. While we have made significant investments in R&D activities, there can be no assurance that our expenditure on R&D activities will yield proportionate results of substantial commercial value or that commercially viable products may be developed or launched as a result of such R&D activities.  The following table sets forth the R&D expenditure and R&D expenditure as a percentage of total income of Indian CDMO formulation players and Indian demostic formulation players as of Fiscal 2021.		
	table of attrition rate as given in reply to RF22. Further, LM is advised to explain reason for high attrition. LM is also advised to clarify reason for non-availability of R& D expenditure of the Company and how the company is	market demand faster with new, alternative or emerging technologies. If our competitors gain significant market share at our expense, our business, results of operations and financial condition could be adversely affected. Changes in the nature or extent of our customer requirements may render our service and product offerings obsolete or non-competitive, which could have a material adverse effect on our business, results of operations and financial condition."  Risk factor 9: LM is advised to include table of attrition rate as given in reply to RF22. Further, LM is advised to be expenditured as given in reply to RF22. Further, LM is advised to explain reason for high attrition. LM is also advised to clarify reason for non-availability of R& D expenditure of the Company and how the company is going to estimate the same.  "9. We are dependent on our R&D activities for our future success. If we do not successfully develop new products or continue our generic product portfolio expansion in a timely and cost-effective manner, our business, results of operations and financial condition may be adversely affected.  The pharmaceutical and healthcare industry is characterised by technological advancements, introduction of innovative products, price fluctuations and intense competition. We have invested substantial effort, funds and other resources towards our R&D activities. We have a dedicated R&D laboratory and pilot equipment located at our manufacturing facility at Baddi, Himachal Pradesh. We are increasingly engaged in R&D activities to develop various generic products, manufacturing process and technologies for diverse therapeutic segments. In particular, our R&D laboratory is focused on developing efficient processes for the manufacture of upcoming patent expired products. In the nine months ended December 31, 2021 and Fiscal 2021, our R&D expenditure on a restated consolidated basis was 348.51 million and 321.45 million, respectively. While we have made significant investments in R&D activities, there can be no	market demand faster with new, alternative or emerging technologies. If our competitors gain significant market share at our expense, our business, results of operations and financial condition could be adversely affected. Changes in the nature or extent of our customer requirements may render our service and product offerings obsolete or non-competitive, which could have a material adverse effect on our business, results of operations and financial condition."  Risk factor 9: LM is advised to include table of attrition rate as given in reply to RF22. Further, LM is advised to explain reason for high attrition. LM is advised to explain reason for high attrition. LM is also advised to clarify reason for non-availability of R&D expenditure of the Company and how the company is going to estimate the same.  **B.**D department under the head of salaries, remuneration, power & electricity, material consumption, clinical trial expenses and product registration and licence fees.  **9. We are dependent on our R&D activities for our future success. If we do not successfully develop new products or continue our generic product portfolio expansion in a timely and cost-effective manner, our business, results of operations and financial condition may be adversely affected.  The pharmaceutical and healthcare industry is characterised by technological advancements, introduction of innovative products, price fluctuations and intense competition. We have invested substantial effort, funds and other resources towards our R&D activities. We have a dedicated R&D laboratory and pilot equipment located at our manufacturing facility at Baddi, Himachal Pradesh. We are increasingly engaged in R&D activities to develop various generic products, manufacturing processes and technologies for diverse therapeutic segments. In particular, our R&D laboratory is focused on developing efficient processes for the manufacture of upcoming patent expired products, manufacturing processes and technologies for diverse therapeutic segments. In particula





S. No.	Observation		Response			RHP Page Number (Clean)	RHP Page Number (Redline)
				(in ⁼	₹ million, except percentages)		
		Company Name	Total Income	R&D expenditure	R&D expenditure as a % of total income		
		Indian CDMO formulation players			o,		
		Akums Drug and Pharmaceuticals Ltd.	27,438.85	280.31	1.02%		
		Innova Captab Limited (1)	6,162.54	21.45	0.35%		
		Synokem Pharmaceuticals Ltd.	5,576.06	70.20	1.26%		
		Windlas Biotech Ltd.	4,306.96	36.06	0.84%		
		Theon Pharmaceuticals Ltd.	4,005.06	9.90	0.25%		
		Acme Formulations Private Itd.	4,127.53	47.98	1.16%		
		Indian domestic formulation players					
		Abbott India	43,909.20	10.80	0.02%		
		Alembic Pharma	54,031.40	7,330.00	13.57%		
		Aurobindo Pharma Ltd.	251,554.70	15,993.80	6.36%		
		Biocon Ltd	73,603.00	5,531.00	7.51%		
		Cipla Ltd	194,255.80	8,667.00	4.46%		
		Dr.Reddy's Laboratories Ltd.	193,389.00	17,333.00	8.96%		
		GlaxoSmithKline	30,361.84	18.00	0.06%		
		Glenmark Pharmaceuticals Ltd.	109,941.45	13,187.00	11.99%		
		Ipca Labs	54,828.30	1,266.70	2.31%		
		Lupin Ltd#	152,992.50	14,324.20	9.36%		
		Panacea Biotech Ltd.	6,347.82	247.70	3.90%		
		Sun Pharmaceuticals Industries Ltd.	343,336.60	21,443.30	6.25%		
		Torrent Pharmaceuticals Ltd	80,614.80	4,870.00	6.04%		
		Wockhardt Ltd.	28,405.70	2,655.00	9.35%		
		(1) R&D expenditure and total income (Source: CRISIL Report).	e on a proforma consolidate	d basis.			





S. No.	Observation		RHP Page Number (Clean)	RHP Page Number (Redline)		
		anticipate customer needs and prej suppliers and customers; customize and cost-effective manner; and succommercialisation of new products is development activities, obtaining reginvestments in new manufacturing a costs, especially in the event of cost our competitors in the CDMO and resources. They may also be in a betinnovative new products. According portfolio expansion in a timely, cost-operations and financial condition mour future results of operations also products and continue our product pour business strategies, we intend to chains. In addition, we intend to expandition our research capabilities in order to determinant for our success. The dicustomers' products) are complex, till unable to successfully create these in and even if launched as planned, suc	depend, to a significant degree, on our portfolio expansion in a timely and cost-of further diversify our product portfolion and our capacities in existing products as ensure continued product innovation. evelopment and commercialisation of me-consuming, costly and involves a higher products or encounter unexpected definition of the products may not perform as we expect and the number of R&D employees as	rovals; establish collaborations with manufacture our products in a timely. In addition, the development and its, including costs relating to product and sales and marketing. Our planned ture expansion could result in higher use in revenues. Additionally, some of mancial, research and technological dapt to changes in industry and offer two products or continue our product our customers, our business, results of the product was a successfully develop new deffective manner. Further, as part of the by entering into new product value well as expanding and strengthening and Innovation continues to be the key new products (whether ours or our hidegree of business risk. We may be elays in the launch of these products of the second in the launch of these products of the second in the launch of these products of the second in the launch of these products of the second in the launch of these products of the second in the launch of these products of the second in the launch of these products of the second in the launch of these products of the second in the launch of these products of the second in the launch of these products of the second in the launch of these products of the second in the launch of these products of the second in the launch of the s		
		Period				
		Fiscal 2019	25.00%	2.37%		
		Fiscal 2020	52.94%	2.57%		
		Fiscal 2021	22.22%	2.16%		





S. No.	Observation		Nine months ended December 31, 2.23% 2021  Our R&D department's average strength is around 20 employees and due to a relatively lower base, attrition percentage rate reflects a higher number even in case of normal attrition of 4-5 employees a year. The attrition during the Fiscal 2019 was due to attrition of employees at the unskilled and semi-skilled level. During Fiscal 2020, our entire senior staff of then R&D department left our company and started their own R&D Centre. This also led								
		Our R&D department's average streed percentage rate reflects a higher number during the Fiscal 2019 was due to attr									
13.	Risk factor 13: LM is advised to disclose top research organizations with whom the issuer has agreement.	to high attrition in the entire R&D dep Noted for compliance. The Company Organisation; (2) ICBio Clinical Resear	igh attrition in the entire R&D department during Fiscal 2021."  ed for compliance. The Company has third party clinical trial agreements with (1) AnaCipher Clinical Research anisation; (2) ICBio Clinical Research Pvt. Ltd.; and (3) Synergen Bio Pvt. Ltd. The Company, however, does not e consent to disclose the name of these companies in the RHP so no additional amendment to Risk Factor 13								
14.	Risk factor 20: LM is advised to disclose progress of the upcoming facility in Jammu and to also include the updates in the section on major events and milestones of the Company.	is proposed. Complied with and noted for compliance. We undertake to amend Risk Factor 20 in the UDRHP and RHP as set forth below. Complied with and noted for compliance. We undertake to amend Risk Factor 20 in the UDRHP and RHP as set forth below. Complied with and noted for compliance. We undertake to amend Risk Factor 20 in the UDRHP and RHP as set forth below. Complied with and noted for compliance. We undertake to amend Risk Factor 20 in the UDRHP and RHP as set forth below. Complied with and noted for compliance with a set forth below. Complied with and noted for compliance with a set forth below. Complied with and noted for compliance with a set forth below. Complied with and noted for compliance with a set forth below. Complied with and noted for compliance with a set forth below. Complied with and noted for compliance with a set forth below. Complied with and noted for compliance with a set forth below. Complied with and noted for compliance with a set forth below. Complied with a set forth below. C									
		1		utory and other regulatory approvals, and we cannot assure you that we will							





S. No.	Observation	Response	RHP Page Number (Clean)	RHP Page Number (Redline)
		be able to obtain or renew such approvals, licenses, permits and registrations in a timely manner, or at all. If we fail to obtain or renew such licenses, approvals, registrations and permits in a timely manner, our commissioning date for the expansion plans may be delayed, which could adversely affect our business and results of operations.  There can be no assurance that the proposed capacity additions and expansions will be completed as planned or on schedule, and if they are not completed in a timely manner, or at all, our budgeted costs may be insufficient to meet our proposed capital expenditure requirements. If our actual capital expenditures significantly exceed our budgets, or even if our budgets were sufficient to cover these projects, we may not be able to achieve the intended economic benefits of these projects, which in turn may materially and adversely affect our financial condition, results of operations, cash flows, and prospects. There can be no assurance that we will be able to complete the aforementioned expansion and additions in accordance with our proposed plans and any delay could have an adverse impact on our growth, prospects, cash flows and financial condition.  Construction and operation of our new Jammu Facility will require us to obtain various approvals, which are routine in nature. There can be no assurance that we will be able to obtain these registrations and approvals in a timely manner or at all.  As on August 31, 2021, we have made the following progress on construction of our new Jammu Facility:  Land has been acquired and possession taken;  Orders for plant & machinery are ongoing;  Construction contracts are being finalized;  Term loan for the Jammu Facility has been sanctioned;  Acknowledgment of our intent to establish a manufacturing enterprise has been received from the office		
		<ul> <li>of the General Manager of District Industries Centre; and</li> <li>Application has been made for Consent to Establish with the Pollution Control Committee."</li> </ul>		
15.	Risk Factor 23: LM is advised to clarify reason for y-o-y increase in trade receivables.	Complied with and noted for compliance. We undertake to amend Risk Factor 23 in the UDRHP and RHP as set forth below. Amendments also include changes as set forth in our response letter dated August 3, 2022.	51	70
		"23. Our inability to collect receivables and default in payment from our customers could result in the reduction of our profits and affect our cash flows.		





S. No.	Observation		Response							
		The majority of our sale between 30 to 90 days arrangements and limit customer's financial conunable to pay. As a result for potential credit losse estimates may not be account of the contraction of the								
		operations on a restated consolidated basis for the periods indicated are set forth below.  (in ₹ million, except percentages								
		Particulars	Fiscal 2019	Fiscal 2020	Fiscal 2021	Nine Months ended				
		Trade receivables	908.29	867.69	1,385.53	2,006.82				
		Trade receivables as a percentage of revenue from operations	25.53%	23.24	33.74%	34.36%				
		Our trade receivables on operations on a proform			are set forth below.	entage of revenue from illion, except percentages)				
		Particulars	Fiscal 2019	Fiscal 2020	Fiscal 2021	Nine Months ended				
		Trade receivables	1,319.14	1,373.62	1,487.31	2,006.82				
		Trade receivables as a percentage of revenue from operations	26.73%	25.37%	24.23%	31.66%				
		On a restated consolidated \$867.69 million as at Mosconsolidated basis also	arch 31, 2020. Trade re	ceivables as a percentag	ge of revenue from o	-				





S. No.	Observation	Response	RHP Page Number (Clean)	RHP Page Number (Redline)
		Trade receivables increased from ₹867.69 million as at March 31, 2020 to ₹1,385.53 million as at March 31, 2021 on a restated consolidated basis, mainly due to acquisition of assets and liabilities of Innova Partnership on account of slump sale effective as of March 31, 2021. Also, the revenue from operations increased from ₹4,106.62 million for Fiscal 2021 to ₹5,841.21 million on a restated consolidated basis for the nine months ended December 31, 2021. Trade receivables as a percentage of revenue from operations on a restated consolidated basis increased from 23.24% for Fiscal 2020 to 33.74% for Fiscal 2021 mainly due to the reason that because Innova Partnership assets and liabilities were acquired on March 31, 2021, the trade receivables include Innova Partnership's trade receivables but the revenue from operations for the Fiscal 2021 does not include Innova Partnership's revenue from operations for the Fiscal 2021.		
		Trade receivables on a restated consolidated basis increased from ₹1,385.53 million as at March 31, 2021 to ₹2,006.82 million as at December 31, 2021, mainly due to acquisition of UML as of December 31, 2021. Trade receivables on a restated consolidated basis as a percentage of revenue from operations increased from 33.74% for Fiscal 2021 to 34.36% for the nine months ended December 31, 2021 mainly due to the reason that because UML was acquired on December 31, 2021, the trade receivables include UML's trade receivables but the revenue from operations for the nine months ended December 31, 2021 does not include UML's revenue from operations for the nine months ended December 31, 2021.		
		Trade receivables on a proforma consolidated basis marginally increased from ₹1,319.14 million as at March 31, 2019 to ₹1,373.62 million as at March 31, 2020 and to ₹1,487.31 million as at March 31, 2021. However, trade receivables as a percentage of revenue from operations on a proforma consolidated basis reduced from 26.73% for Fiscal 2019 to 25.37% for Fiscal 2020 and to 24.23% for Fiscal 2021.		
		Trade receivables on a proforma consolidated basis increased from ₹1,487.31 million as at March 31, 2021 to ₹2,006.82 million as at December 31, 2021 mainly due to increase in revenue from operations on a proforma consolidated basis from ₹6,138.39 million for Fiscal 2021 (12 months) to ₹6,337.96 million (9 months). Also, receivable turnover days on a proforma consolidated basis were 88 days and 87 days respectively for the same periods. Trade receivables as a percentage of revenue from operations on a proforma consolidated basis increased from 24.23% for Fiscal 2021 to 31.66% for the nine months ended December 31, 2021 due to the reason that the calculation of trade receivables as a percentage of revenue from operations for the nine months ended December 31, 2021 has not been annualized.		





	Niaalaa	
	Number	Number
	(Clean)	(Redline)
In Fiscal 2019, Fiscal 2020 and Fiscal 2021 and in the nine months ended December 31, 2021, our receivable turnover days on a restated consolidated basis were 93 days, 85 days, 123 days and 94 days, respectively, in the same periods. In Fiscal 2019, Fiscal 2020 and Fiscal 2021 and in the nine months ended December 31, 2021, receivable turnover days on a proforma consolidated basis were 98 days, 93 days, 88 days and 87 days, respectively, in the same periods. Any increase in our receivable turnover days will negatively affect our business. If we are unable to collect customer receivables or if the provisions for doubtful receivables are inadequate, it could have a material adverse effect on our business, results of operations and financial condition.  Macroeconomic conditions could also result in financial difficulties, including insolvency or bankruptcy, of our customers, and as a result could cause customers to delay payments to us, request modifications to their payment arrangements, that could increase our receivables or affect our working capital requirements, or default on their payment obligations to us. An increase in bad debts or in defaults by our customer, may compel us to utilize greater amounts of our operating working capital and result in increased interest costs, thereby adversely affecting our	(e.cu.i)	(neume)
	same periods. In Fiscal 2019, Fiscal 2020 and Fiscal 2021 and in the nine months ended December 31, 2021, receivable turnover days on a proforma consolidated basis were 98 days, 93 days, 88 days and 87 days, respectively, in the same periods. Any increase in our receivable turnover days will negatively affect our business. If we are unable to collect customer receivables or if the provisions for doubtful receivables are inadequate, it could have a material adverse effect on our business, results of operations and financial condition.  Macroeconomic conditions could also result in financial difficulties, including insolvency or bankruptcy, of our customers, and as a result could cause customers to delay payments to us, request modifications to their payment arrangements, that could increase our receivables or affect our working capital requirements, or default on their payment obligations to us. An increase in bad debts or in defaults by our customer, may compel us to utilize greater	same periods. In Fiscal 2019, Fiscal 2020 and Fiscal 2021 and in the nine months ended December 31, 2021, receivable turnover days on a proforma consolidated basis were 98 days, 93 days, 88 days and 87 days, respectively, in the same periods. Any increase in our receivable turnover days will negatively affect our business. If we are unable to collect customer receivables or if the provisions for doubtful receivables are inadequate, it could have a material adverse effect on our business, results of operations and financial condition.  Macroeconomic conditions could also result in financial difficulties, including insolvency or bankruptcy, of our customers, and as a result could cause customers to delay payments to us, request modifications to their payment arrangements, that could increase our receivables or affect our working capital requirements, or default on their payment obligations to us. An increase in bad debts or in defaults by our customer, may compel us to utilize greater amounts of our operating working capital and result in increased interest costs, thereby adversely affecting our





## Schedule I

Name of the lender***	Nature of borrowing	Date of original sanction letter	Sanctione d amount (in ₹ million)	Outstanding amount as at May 15, 2022 (in ₹ million)	Repayment date / schedule	Interest rate (p.a.) as at May 15, 2022	Purpose of raising the loan	Pre-payment penalty, if any
State Bank of India	Cash credit / export packaging credit*	July 14, 2021	550.00	536.70	Repayable on demand.	7.15% (3 month MCLR plus 0.30%)	For working capital requirements.	Nil, in case of prepayment from own sources / 2.00%, in case of takeover of limits by other banks or financial institutions.
	Term loan	August 9, 2016	300.00	44.00	Maximum tenure of 84 months.	7.15% (MCLR 3 month plus 0.40%)	For expansion of existing plant production capacity.	Nil, in case of prepayment from own sources / 2.00%, in case of takeover of limits by other banks or financial institutions.
Yes Bank Limited	Cash credit / working capital demand loan / export packaging credit**	April 29, 2022	800.00	632.76	Maximum tenure of 12 months.	7.35% (1 month MCLR plus 0.05 %)	For working capital requirements.	Nil, in case of prepayment from own sources / 2.00%, in case of takeover of limits by other banks or financial institutions.
	Tourism	August 11, 2016	350.00	3.40	Maximum tenure of 72 months (including a moratorium period of 12 months) from the date of the first disbursement.	7.50% (3 month MCLR plus 0.50%)	For expansion of existing plant production capacity.	Nil, in case of prepayment from own sources / 2.00%, in case of takeover of limits by other banks or financial institutions.
	Term loan	August 11, 2016	- 250.00	3.27	Maximum tenure of 72 months (including a moratorium period of 12 months) from the date of the first disbursement.	7.50% (3 month MCLR plus 0.50%)		Nil, in case of prepayment from own sources / 2.00%, in case of takeover of limits by other banks or financial institutions.
	Term loan	February 9, 2021	200.00	184.70	Maximum tenure of 84 months.	7.50% (3 month MCLR plus 0.05%)	For setting up of new block within the existing unit, and for reimbursement to be done for the capital expenditure already done through own sources.	Nil, in case of prepayment from own sources / 2.00%, in case of takeover of limits by other banks or financial institutions.
	Term loan	September 4, 2021	120.00	70.00	Principal amount to be repaid in 78 equal instalments, after an 18 month moratorium.	7.40% (1 month MCLR plus 0.010%)	For purchase of land and construction of corporate office and R&D center and for reimbursement which is to be done for the capex already done through own sources.	Nil, in case of prepayment from own sources / 2.00%, in case of takeover of limits by other banks or financial institutions.





Name of the lender***	Nature of borrowing	Date of original sanction letter	Sanctione d amount (in ₹ million)	Outstanding amount as at May 15, 2022 (in ₹ million)	Repayment date / schedule	Interest rate (p.a.) as at May 15, 2022	Purpose of raising the loan	Pre-payment penalty, if any
The Hong Kong and Shanghai	Cash credit	August 19, 2020	100.00	83.77	Repayable on demand.	6.30%	To finance working capital requirements.	Nil, in case of prepayment from own sources / 2.00%, in case of takeover of limits by other banks or financial institutions.
Banking Corporation	Term loan	March 19, 2021	200.00	130.18	84 months (including a moratorium of 6 months).	6.50%	Acquisition of capital equipment.	Nil, in case of prepayment from own sources / 2.00%, in case of takeover of limits by other banks or financial institutions.
	Termioan	March 19, 2021	200.00	58.56	84 months (including a moratorium of 6 months).	6.50%	Acquisition of capital equipment.	Nil, in case of prepayment from own sources / 2.00%, in case of takeover of limits by other banks or financial institutions.

<sup>\*</sup> Export packaging credit limit amounting to ₹100.00 million is within overall fund based working capital of ₹550.00 million.

Note: In accordance with Clause 9(A)(2)(b) of Part A of Schedule VI of the SEBI ICDR Regulations which requires a certificate from the statutory auditor certifying the utilization of loan for the purposed availed, our Company has obtained the requisite certificate dated June 27, 2022, from our Statutory Auditors, B S R & Co. LLP, Chartered Accountants, wherein the Statutory Auditors have certified that nothing has come to their attention that causes them to believe that the loans that are proposed to be repaid or pre-paid out of Net Proceeds have not been utilized for the purposes for which these were availed.

<sup>\*\*</sup> Working capital demand loan and export packaging credit amounting to ₹750.00 million and ₹300.00 million respectively are within overall cash credit limit of ₹750.00 million.

<sup>\*\*\*</sup> Additionally, our Company may avail additional loan facilities or draw down existing facilities from time to time to meet our business requirements. Accordingly, our Company may utilise the Net Proceeds for repayment of any such refinanced facilities (including any prepayment fees or penalties thereon), any additional facilities obtained by our Company or working capital facilities outstanding at the time of utilisation of Net Proceeds.