

Neuland Laboratories Limited 11th Floor, (5th Office Level), Phoenix IVY Building, Plot No. 573A-III, Road No.82, Jubilee Hills, Hyderabad-500033, Telangana, India

Tel : 040 67611600 / 67611700 Email : neuland@neulandlabs.com www.neulandlabs.com

February 13, 2023

To BSE Limited Phiroze Jeejeebhoy Towers, 25<sup>th</sup> Floor, Dalal Street, Mumbai - 400 001

The National Stock Exchange of India Ltd Exchange Plaza, Bandra Kurla Complex Bandra (E), Mumbai - 400 001

Scrip Code: 524558

Scrip Code: NEULANDLAB; Series: EQ

Dear Sir/Madam,

Sub: Outcome of Board Meeting - Un-audited Standalone & Consolidated Financial Results for the quarter and nine months ended December 31, 2022

Pursuant to Regulation 33 of SEBI (Listing Obligations & Disclosure Requirements) Regulations, 2015, we wish to inform you that the Board of Directors at their meeting held on even date, i.e. February 13, 2023, has *inter alia*, approved the Unaudited Financial Results (standalone & consolidated) of the Company for the quarter and nine months ended December 31, 2022.

A copy of the Unaudited Financial Results for the quarter and nine months ended December 31, 2022 together with the limited review reports (standalone & consolidated) by the Statutory Auditors of the Company and Press Release along with presentation to the Investors/ Analysts, on the Unaudited Financial Results of the Company for the quarter and nine months ended December 31, 2022 are enclosed herewith.

The above information will also be available on the website of the Company at www.neulandlabs.com.

The meeting of the Board of Directors of the Company commenced at 3:50 PM (IST) and concluded at 5:30 PM (IST).

This is for your information and records.

Thanking you,

Yours Sincerely For Neuland Laboratories Linited

Sarada Bhamidipatish Company Secretary

Encl: As above

Independent Auditor's Review Report on unaudited quarterly and year to date standalone financial results of Neuland Laboratories Limited pursuant to the Regulation 33 of the SEBI (Listing Obligations and Disclosure Requirements) Regulations, 2015, as amended

#### To The Board of Directors of Neuland Laboratories Limited

- 1. We have reviewed the accompanying statement of unaudited standalone financial results of **Neuland** Laboratories Limited ('the Company') for the quarter ended December 31, 2022 and the year to-date results for the period April 01, 2022 to December 31, 2022 ('the Statement') attached herewith, being submitted by the Company pursuant to the requirements of Regulation 33 of the Securities and Exchange Board of India (Listing Obligations and Disclosure Requirements) Regulations, 2015, as amended ('the Regulations').
- 2. This Statement, which is the responsibility of the Company's Management and approved by the Company's Board of Directors, has been prepared in accordance with the recognition and measurement principles laid down in Indian Accounting Standard 34 'Interim Financial Reporting' ('Ind AS 34'), prescribed under Section 133 of the Companies Act, 2013 read with relevant rules issued thereunder and other recognized accounting principles generally accepted in India and in compliance with the Regulations. Our responsibility is to express a conclusion on the Statement based on our review.
- 3. We conducted our review of the Statement in accordance with the Standard on Review Engagement (SRE) 2410, "Review of Interim Financial Information Performed by the Independent Auditor of the Entity" issued by the Institute of Chartered Accountants of India. This standard requires that we plan and perform the review to obtain moderate assurance as to whether the Statement is free of material misstatement. A review consists of making inquiries, primarily of persons responsible for financial and accounting matters, and applying analytical and other review procedures. A review is substantially less in scope than an audit conducted in accordance with Standards on Auditing and consequently does not enable us to obtain assurance that we would become aware of all significant matters that might be identified in an audit. Accordingly, we do not express an audit opinion.
- 4. Based on our review conducted as stated in paragraph 3 above, nothing has come to our attention that causes us to believe that the accompanying Statement prepared in accordance with the recognition and measurement principles laid down in Ind AS 34, prescribed under Section 133 of the Companies Act, 2013 read with relevant rules issued thereunder and other recognized accounting principles generally accepted in India has not disclosed the information required to be disclosed in terms of the Regulations including the manner in which it is to be disclosed, or that it contains any material misstatement.

For M S K A & Associates Chartered Accountants ICAI Firm Registration No. 105047W

Amit **Kumar Agarwa**l Partner Membership No. 214198

UDIN: 23214198BGXCOW8186 Place: Hyderabad, INDIA Date: February 13, 2023





Neuland Laboratories Limited 11th Floor, (5th Office Level), Phoenix IVY Building, Plot No. 573A-III, Road No.82, Jubilee Hills, Hyderabad-500033, Telangana, India

Tel : 040 6767 1600 / 6761 1700 Email : neuland@neulandlabs.com www.neulandlabs.com

#### NEULAND LABORATORIES LIMITED 11th Floor (5th Office Level), Phoenix IVY Building, Plot No. 573A-III, Road No 82, Jubilee Hills, Hyderabad - 500033 STATEMENT OF STANDALONE FINANCIAL RESULTS FOR THE QUARTER AND NINE MONTHS ENDED 31 DECEMBER 2022 (Amount in lakhs of ₹, unless otherwise stated) Year Ended Quarter Ended Nine Months Ended 31.03,2022 31.12.2022 30.09.2022 31.12.2021 31 12 2022 31.12.2021 SI. No. Particulars (Audited) (Unaudited) (Unaudited) (Unaudited) (Unaudited) (Unaudited) 1 Revenue 95,107.66 26,925.00 29.370.55 23.646.98 78.412.54 69.520.41 Revenue from operations (a) 93.31 190.01 168.40 424.25 207.67 23.91 (b) Other income 95,315.33 23,836.99 78,580.94 69,944.66 27,018.31 29.394.46 Total income Expenses 2 12,156.92 9,584.52 34,981.78 33,984.03 43.755.36 Cost of materials consumed 13,200.18 (a) (1,541.47) 1,026,90 (3,530.89) (2,644.72) (1.076.59) (4.482.70) Changes in inventories of finished goods and work-in-progress (b) 13,332.28 17,575.85 4,569.27 4,988.89 5,055.70 14,705.46 Employee benefits expense (c)317.54 332.16 1,072.44 1,349.44 274,47 877.81 $\{d\}$ Finance costs 4,903.64 1.277.48 3,658.41 1.333.62 1.317.17 3.930.63 Depreciation and amortisation expense (e) 9,899.99 13.136.31 3,388.00 3.539.91 3,387.99 10,013.57 Manufacturing expenses rß 1,848.86 5,766.59 7,962.22 2,598.29 2,777.71 8,035.36 (g) Other expenses 87,141.35 64.182.85 Total expenses 23,138.73 24.088.36 22.027.18 68.061.91 5,761.81 8,173.98 3,879.58 5,306,10 1,809.81 10,519.03 Profit before tax (1-2) 3 4 Tax expense 1,433.24 662.15 444.31 2,505.38 1.339.15 2.042.19 Current tax (a) 251.98 (221.55) 173.80 39.65 92.27 152.76 Deferred tax (b) 6,353.34 4.170.68 7.860.89 Profit for the period / year (3-4) 3.043.63 3.833.21 1,273,23 5 Other comprehensive income (net of taxes) 6 (a) items that will not be reclassified to profit or loss (81.20) (148.44) (396.90) Re-measurement gains/(losses) on defined benefit plans (49,48) (15.07) (302.16) Equity instruments through other comprehensive income (167.78) 2.97 0.66 (166.32) (301.97)37.36 99.89 Tax on items that will not be reclassified to profit or loss 20.44 12.46 3.79 3.757.63 5,754.17 2.875.85 3.775.41 7.683.29 1.236.87 Total comprehensive income 7 Paid-up Equity Share Capital 1,290.05 1,290.05 1,290.05 1,290.05 1,290.05 1.290.05 82,813.49 Other equity (excluding revaluation reserve) ß 9 Earnings Per Share (of ₹10 each) (In absolute ₹ terms) 23.72 29.88 9,92 61.27 32.51 49.52 Basic (refer note 4) (a) 32.51 Diluted (refer note 4) 23.72 29.88 9.92 61.27 49.52 {b) See accompanying notes to the financial results





Registered Office : 11th Floor, (5th Office Level), Phoenix IVY Building, Plot No. 573A-III, Road No.82, Jubilee Hiljs, Hyderabad-500033, Telangana, India | CIN No. L85195TG1984PLC004393



NOT	TES:
ŧ	The financial results for the quarter and nine months ended 31 December 2022 have been reviewed by the Audit Committee and approved by the Board of Directors at the respective meetings held on 13th February 2023.
3	The financial results have been prepared in accordance with the Indian Accounting Standards (Ind AS) prescribed under Section 133 of the Companies Act, 2013, and othe accounting principles generally accepted in India and in terms of Regulation 33 of the SEBI (Listing Obligations and Disclosure Requirements) Regulations, 2015.
3	The operations of the Company are predominantly related to the manufacture and sale of active pharmaceutical ingredients and allied services. As such there is only or primary reportable segment as per Ind AS 108 "Operating Segments".
4	The EPS for quarters has not been annualised.
5	The previous period figures have been regrouped/nearranged wherever necessary to make it comparable with the current period.
	For Neuland Laboratories Limited
	HYDERABAD T
	Place: Hyderabad Dr. D R Rao Executive Chairman
	Date: 13 February 2023 (DIN 09107737)



### MSKA & Associates Chartered Accountants

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Hyderabad

Independent Auditor's Review Report on Consolidated Unaudited Quarterly and year to date financial results of Neuland Laboratories Limited pursuant to the Regulation 33 of the SEBI (Listing Obligations and Disclosure Requirements) Regulations, 2015, as amended

### To The Board of Directors Neuland Laboratories Limited

- 1. We have reviewed the accompanying statement of consolidated unaudited financial results of Neuland Laboratories Limited ('the Holding Company') and its subsidiaries, (the Holding Company and its subsidiaries together referred to as the 'Group') and for the quarter ended December 31, 2022 and the year to-date results for the period from April 01, 2022 to December 31, 2022 ('the Statement'), being submitted by the Holding Company pursuant to the requirements of Regulation 33 of the Securities and Exchange Board of India (Listing Obligations and Disclosure Requirements) Regulations, 2015, as amended ('the Regulations').
- 2. This Statement, which is the responsibility of the Holding Company's Management and approved by the Holding Company's Board of Directors, has been prepared in accordance with the recognition and measurement principles laid down in Indian Accounting Standard 34 'Interim Financial Reporting' ('Ind AS 34'), prescribed under Section 133 of the Companies Act, 2013 read with relevant rules issued thereunder and other recognized accounting principles generally accepted in India and in compliance with the Regulations. Our responsibility is to express a conclusion on the Statement based on our review.
- 3. We conducted our review of the Statement in accordance with the Standard on Review Engagement (SRE) 2410, 'Review of Interim Financial Information Performed by the Independent Auditor of the Entity' issued by the Institute of Chartered Accountants of India. This standard requires that we plan and perform the review to obtain moderate assurance as to whether the Statement is free of material misstatement. A review consists of making inquiries, primarily of persons responsible for financial and accounting matters, and applying analytical and other review procedures. A review is substantially less in scope than an audit conducted in accordance with Standards on Auditing and consequently does not enable us to obtain assurance that we would become aware of all significant matters that might be identified in an audit. Accordingly, we do not express an audit opinion.

We also performed procedures in accordance with the circular issued by the SEBI under Regulation 33 (8) of the SEBI (Listing Obligations and Disclosure Requirements) Regulations, 2015, as amended, to the extent applicable.

Sr. No	Name of the Entity	Relationship with the Holding Company
1	Neuland Laboratories K.K., Japan	Wholly Owned Subsidiary
2	Neuland Laboratories Inc., USA	Wholly Owned Subsidiary

4. This Statement includes the results of the Holding Company and the following entities:

5. Based on our review conducted and procedures performed as stated in paragraph 3 above, nothing has come to our attention that causes us to believe that the accompanying Statement prepared in

accordance with the recognition and measurement principles laid down in Ind AS 34, prescribed under Section 133 of the Companies Act, 2013 read with relevant rules issued thereunder and other recognized accounting principles generally accepted in India has not disclosed the information required to be disclosed in terms of the Regulations including the manner in which it is to be disclosed, or that it contains any material misstatement.

6. The consolidated unaudited financial results includes the interim financial information of two subsidiaries (mentioned in paragraph 4 above) which have not been reviewed or audited by their auditors, whose interim financial information reflect total revenue (before consolidation adjustments) of Rs. 293.90 lakhs and 824.19 lakhs, total profit after tax (before consolidation adjustments) of Rs. 11.19 lakhs and Rs. 36.50 lakhs and total other comprehensive income (before consolidation adjustments) of Rs. 24.82 lakhs and 73.38 lakhs for the quarter ended December 31, 2022 and for the period from April 01, 2022 to December 31, 2022 respectively as considered in the statement. According to the information and explanations given to us by the Management, these interim financial information are not material to the Group.

Our conclusion on the Statement is not modified in respect of our reliance on the interim financial information certified by the Management.

For M S K A & Associates Chartered Accountants ICAI Firm Registration No. 105047W

Amit Kumar Agarwal Partner Membership No. 214198

UDIN: 23214198BGXCOX2378 Place: Hyderabad, INDIA Date: February 13, 2023





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Tel : 040 6761 1600 / 6761 1700 Email : neuland@neulandlabs.com www.neulandlabs.com

#### NEULAND LABORATORIES LIMITED 11th Floor (5th Office Level), Phoemix IVY Building, Plot No. \$73A-III, Road No 82, Jubilee Hills, Hyderabad - 500033 STATEMENT OF CONSOLIDATED FINANCIAL RESULTS FOR THE QUARTER AND NINE MONTHS ENDED 31 DECEMBER 2022

			Quarter Ended		Nine Months Ended		Year Ended	
51. No.	Particulars	31,12,2022 (Unaudited)	30.09.2022 (Unaudited)	31.12,2021 (Unaudited)	31.12.2022 (Unaudited)	31,12,2021 (Unaudited)	31.03.2022 (Audited)	
1	Revenue						25 427 44	
	(a) Revenue from operations	26,925.00	29,370.55	23,646.98	78,412.54	69,520.41	95,107.66	
	(b) Other income	93.31	23.91	190.01	168.40	424.25	207.6	
	Total Income	27,018.31	29,394.46	23,836.99	78,580.94	69,944.66	95,315.34	
2	Expenses					72 00 / 02	43 365 3	
	(a) Cost of materials consumed	13,200.18	12,156.92	9,584.53	34,981.78	33,984.03	43,755.3	
	(b) Changes in inventories of finished goods and work-in-progress	{2,644.72}	(1,076.59)	1,026.90	(4,482.70)	(3,530.89)	(1,541.4)	
	(c) Employee benefits expense	5,162.34	5,233,22	4,803,67	15,225.53	13,850.27	18,247.2	
	(d) Finance costs	274.49	317.54	332.18	877.84	1,072.48	1,349.5	
	(e) Depreciation and amortisation expense	1,333.63	1,317.19	1,277.48	3,930.75	3,658.61	4,904.0	
	(f) Manufacturing expenses	3,387.99	3,539.90	3,387.99	10,013.56	9,699.99	13,136.3	
	(g) Other expenses	2,410.89	2,587.42	1,600.65	7,475.90	5,221.44	7,248.1	
	Total expenses	23,124.80	24,075.60	22,013.40	68,022.66	64,155.94	87,099.0	
3	Profit before tax (1-2)	3,893.51	5,318.86	1,823.59	10,558.28	5,788.72	B,216.2	
4	Tax expense							
	(a) Current tax	664,89	1,433.24	437.76	2,508.14	1,332.61	2,042.9	
	(b) Deferred tax	173.80	39.65	92.27	152.77	251.99	(208,8	
5	Profit for the period / year (3-4)	3,054.82	3,845.97	1,293.56	7,897.37	4,204.13	6,382.2	
6	Other comprehensive income (net of taxes)							
	(a) items that will not be reclassified to profit or loss							
	Re-measurement gains/ (losses) on defined benefit plans	•	(81.20)	(49,48)	(15.07)		(396.9)	
	Equity Instruments through other comprehensive income	(167.78)	2.97	0.66	(166.32)		(302.1)	
	Tax on items that will not be reclassified to profit or loss		20.43	12.46	3.79	37.36	99.8	
	(b) items to be reclassified to profit or loss							
	Exchange differences in translating the financial statements of a foreign operations	13.63	14.30	2.64	36.87	3.44	6.1	
	Total comprehensive income	2,900.67	3,802.47	1,259.84	7,756.64	3,794.52	5,789.2	
7	Paid-up Equity Share Capital	1,290.05	1,290.05	1,290.05	\$,290.05	1,290.05	1,290.0	
8	Other equity (excluding revaluation reserve)						83,676.1	
9	Earnings Per Share (of ₹10 each) (in absolute ₹ terms)							
	(a) Basic	23.81	29.97	10.08	61.55	32.77	49.7	
	(b) Diluted	23.81	29.97	10,08	61.55	32.77	49.7	
	See accompanying notes to the financial results							





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For Neuland Laborat

Dr. D R Rao Executive Chair

(DIN 00107737)

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TYDERABAD

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#### NOTES:

- 1 The financial results for the quarter and nine months ended 31 Dec 2022 have been reviewed by the Audit Committee and approved by the Board of Directors at their respective meetings held on 13th February 2023.
- 2 The financial results have been prepared in accordance with the Indian Accounting Standards (ind AS) prescribed under Section 133 of the Companies Act, 2013, and other accounting principles generally accepted in India and in terms of Regulation 33 of the SEBI (Listing Obligations and Disclosure Requirements) Regulations, 2015.
- The consolidated financial results include results of the following wholly owned subsidiaries:
  (a) Neuland Laboratories Inc., USA
  (b) Neuland Laboratories KK., Japan.
- 4 The operations of the Company and its subsidaries are predominantly related to the manufacture and sale of active pharmaceutical ingredients and allied services. As such there is only one primary reportable segment as per ind AS 108 "Operating Segments".
- 5 The EPS for quarters has not been annualised.
- 6 The previous period figures have been regrouped/rearranged wherever necessary to make it comparable with the current period.

Place: Hyderabad Date: 13 February 2023





### Neuland Q3FY23 income at Rs.270.2 crore, up 13.3% YoY

### EBITDA at Rs. 54.9 crore, up 60.5% YoY

Hyderabad, India, February 13, 2023 - Neuland Laboratories Limited (NLL) (NSE: NEULANDLAB; BSE:524558), a pharmaceutical manufacturer providing active pharmaceutical ingredients (APIs), complex intermediates and custom manufacturing solutions services to customers located in around 80 countries, today announced financial results for the third quarter and nine months ended December 31, 2022.

Commenting on the performance Mr. Sucheth Davuluri, Vice-Chairman and Chief Executive Officer of the Company said, "Over the 9-month period, we recorded a healthy revenue increase led by continued growth in the high margin Specialty and CMS business. Further, the margin improvement reflects the operating leverage playing out. In line with our strategy, we are seeing a significant improvement in the quality of business which is reflected in heathy cashflows. We expect the current business momentum to continue going forward."

In addition, Mr. Saharsh Davuluri, Vice Chairman and Managing Director, Neuland Labs added "Our investments in Unit 3 as well our emphasis on R&D and Project Execution has resulted in our profitability seeing a marked increase. This was a significant quarter for CMS as we saw the impact of molecules transitioning from development to commercial leading to the highest ever commercial revenues. We expect the CMS business to scale greater heights over the foreseeable future led by steady additions to the pipeline."

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Particulars	Q3FY23	Q2FY23	QoQ Growth (%)	Q3FY22	YoY Growth (%)	9MFY23	9MFY22	YoY Growth (%)
Total Income	270.2	293.9	(8.1)%	238.4	13.3%	785.8	699.4	12.3%
EBITDA	54.9	69.4	(20.9)%	34.2	60.5%	153.3	104.9	46.1%
EBITDA margin (%)	20.3%	23.6%	(330) bps	14.3%	600 bps	19.5%	15.0%	450 bps
PAT	30.4	38.3	(20.6)%	12.7	139.0%	78.6	41.7	88.5%
PAT margin (%)	11.3%	13.0%	(170) bps	5.3%	600 bps	10.0%	6.0%	400 bps
EPS (Basic) Rs.	23.7	29.9	(20.6)%	9.9	139.0%	61.3	32.5	88.5%

### Financial Summary

Rs. crore





### Q3 FY23 Earnings Call

The company will conduct a one-hour Earnings call at 16:30 hrs. IST on Tuesday, February 14, 2023 where the management will discuss the Company's performance and answer questions from participants. To participate in this conference call, please register on the link below:

**Diamond Pass Registration Link** 

Please note that the transcript of the conference call will be uploaded on the company website in due course.

### About Neuland Laboratories Limited

For over 39 years, Neuland Labs has been at the forefront of manufacturing APIs through its cGMP manufacturing facilities, working with customers in close to 80 countries. Neuland Labs has developed more than 300 processes and 75 APIs and has filed over 903+ Regulatory filings in the US (60 active US DMFs), the European Union (EU) and other geographies. Its manufacturing facilities are inspected and approved by the U.S. FDA and other leading regulatory agencies. Its record of quality manufacturing and reliability is highlighted by cGMP certifications that include the U.S. FDA, TGA (Australia), EDQM (EU), German Health Authority, ANVISA (Brazil), EMA (EU), Cofepris (Mexico), KFDA (Korea), PMDA (Japan), CFDA (China), FSI "SID &GP" Russia, Health Canada, ISO 9001, ISO14001, OHSAS18001 and ISO 27001. For more information, visit www.NeulandLabs.com.

If you have any questions or require further information, please feel free to contact

IR Department at Neuland Tel: +91 40 6761 1600 Email: <u>ir@neulandlabs.com</u> Ravi Udeshi, EY IR Email: <u>ravi.udeshi@in.ey.com</u>



# Earnings Presentation

BSE CODE : 524558 | NSE SYMBOL : NEULANDLAB | BLOOMBERG: NLL:IN | REUTERS: NEUL.NS

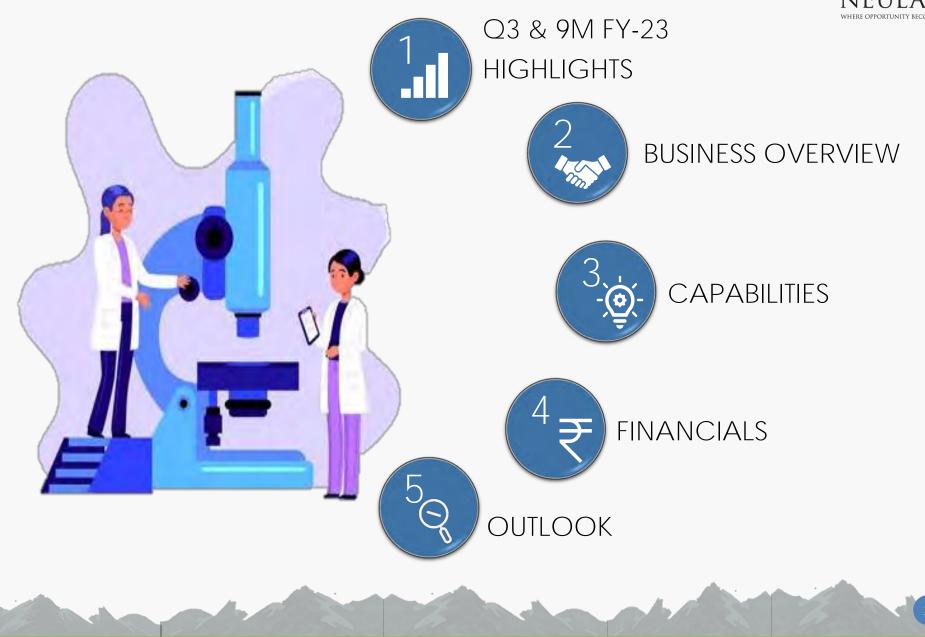
# Safe Harbour



Except for the historical information contained herein, statements in this presentation and the subsequent discussions, which include words or phrases such as "will", "aim", "will likely result", "would", "believe", "may", "expect", "will continue", "anticipate", "estimate", "intend", "plan", "contemplate", seek to", "future", "objective", "goal", "likely", "project", "should", "potential", "will pursue", and similar expressions of such expressions may constitute "forward-looking statements". These forward-looking statements involve a number of risks, uncertainties and other factors that could cause actual results to differ materially from those suggested by the forward-looking statements. These risks and uncertainties include but are not limited to our ability to successfully implement our strategy, our growth and expansion plans, obtain regulatory approvals, our provisioning policies, technological changes, investment and business income, cash flow projections, our exposure to market risks as well as other risks. The Company does not undertake any obligation to update forward-looking statements to reflect events or circumstances after the date thereof.

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# Q3 & 9M FY-23 HIGHLIGHTS

# Management Speak





### SUCHETH DAVULURI

"Over the 9-month period, we recorded healthy revenue increase led by continued growth in the high margin Specialty and CMS business. Further, the margin improvement reflects the operating leverage playing out. In line with our strategy, we are seeing a significant improvement in the quality of business which is reflected in heathy cashflows. We expect the current business momentum to continue going forward."



### SAHARSH DAVULURI

"Our investments in Unit 3 as well our emphasis on R&D and Project execution has resulted in our profitability seeing a marked increase. This was a significant quarter for CMS as we saw the impact of molecules transitioning from development to commercial leading to the highest ever commercial revenues. We expect the CMS business to scale greater heights over the foreseeable future led by steady additions to the pipeline."

# Key Highlights

23 FY

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### **Business Highlights**

- Specialty business driven by Apixaban and Ezetimibe
- CMS business driven by commercial molecules
- Prime segment revenues recovered backed by Mirtazapine, Ciprofloxacin, Labetalol



### Financial Highlights

- Total income was Rs. 270.2 crores in Q3FY23, an increase of 13.3% YoY
- EBITDA was Rs. 54.9 crore in Q3FY23, an increase of 60.5% YoY
- EBITDA margin increased by 600 bps YoY to 20.3% in Q3 FY23 from 14.3% due to
  - Easing of input prices
  - Shift towards higher margin products
- PAT increased by 139.0% YoY to Rs. 30.4 crores
- Debt: Equity stood at 0.13x due to Rs. 80 crores borrowings retired in Q3FY23

# Key Highlights

PM FY2

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### **Business Highlights**

- Specialty business growth driven by Apixaban, Ezetimibe, Paliperidone, and Donepezil
- CMS revenues driven by commercial molecules some of which have transitioned recently
- In Prime segment Mirtazapine and Labetalol were the key molecules
- Sustainability rating of Silver given by ECOVADIS during the period



### **Financial Highlights**

- Total income was Rs. 785.8 crore in 9MFY23, an increase of 12.3% YoY
- EBITDA was Rs. 153.3 crore in 9MFY23, an increase of 46.1% YoY
- EBITDA margin increased by 450 bps YoY to 19.5% in 9M FY23 from 15.0% due to
  - Shift towards higher margin products
- PAT increased by 88.5% to Rs. 78.6 crores
- Debt: Equity stood at 0.13x due to retirement of Rs. 111 crores borrowings in 9MFY23



# Profit & Loss Snapshot (Standalone)



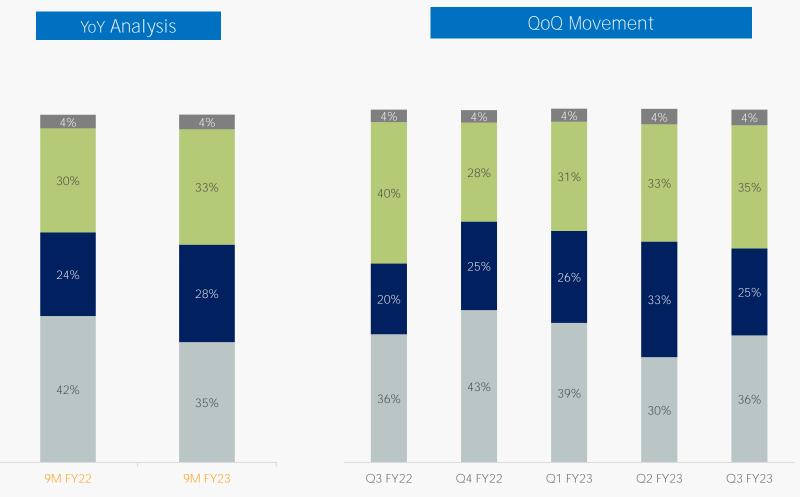
			QoQ		YoY			YoY
Particulars (Rs. Cr)		Q2FY23	(%)	Q3FY22	(%)		9MFY22	(%)
Total Income	270.2	293.9	(8.1)%	238.4	13.3%	785.8	699.4	12.3%
EBITDA	54.9	69.4	(20.9)%	34.2	60.5%	153.3	104.9	46.1%
EBITDA Margin	20.3%	23.6%	(330) bps	14.3%	600 bps	19.5%	15.0%	450 bps
Profit Before Tax	38.8	53.1	(26.9)%	18.1	114.4%	105.2	57.6	82.6%
Profit Before Tax Margin	14.4%	18.1%	(370) bps	7.6%	680 bps	13.4%	8.2%	520 bps
Profit After Tax	30.4	38.3	(20.6)%	12.7	139.0%	78.6	41.7	88.5%
Profit After Tax Margin	11.3%	13.0%	(170) bps	5.3%	600 bps	10.0%	6.0%	400 bps
Earnings Per Share (Rs.)	23.7	29.9	(20.6)%	9.9	139.0%	61.3	32.5	88.5%

# Financials (Standalone)



# Key Operating Metrics



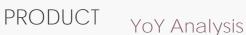


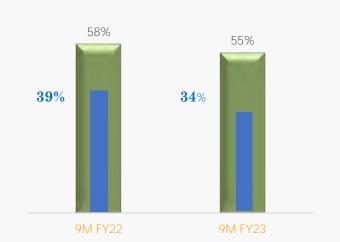
■ Prime ■ Specialty ■ CMS ■ Others

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# Business Salience (Overall Company)





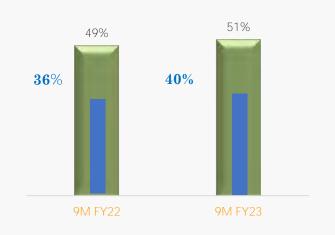




**QoQ** Movement

TOP 10 TOP 5

### CUSTOMER YoY Analysis



### QoQ Movement



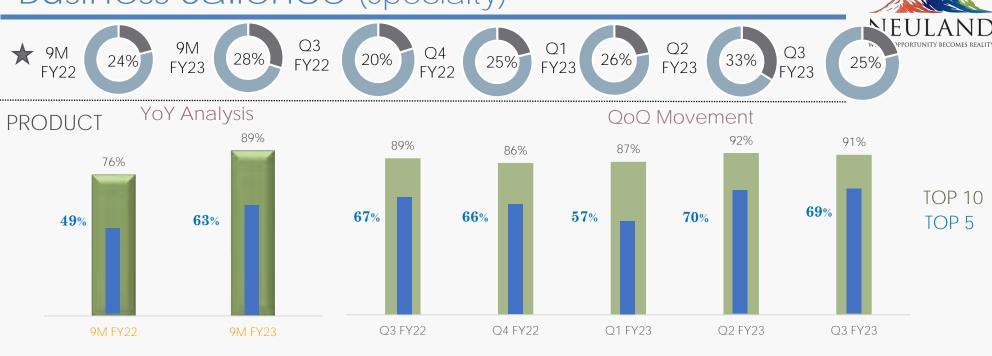
### Business Salience (Prime)



\* % represents Prime % from Overall revenue for respective Quarter / YTD

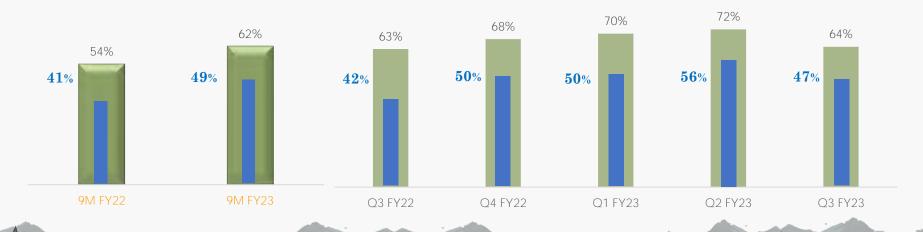
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### Business Salience (specialty)



CUSTOMER YoY Analysis

### QoQ Movement



X % represents Niche / Speciality % from Overall revenue for respective Quarter / YTD

### Business Salience (CMS)



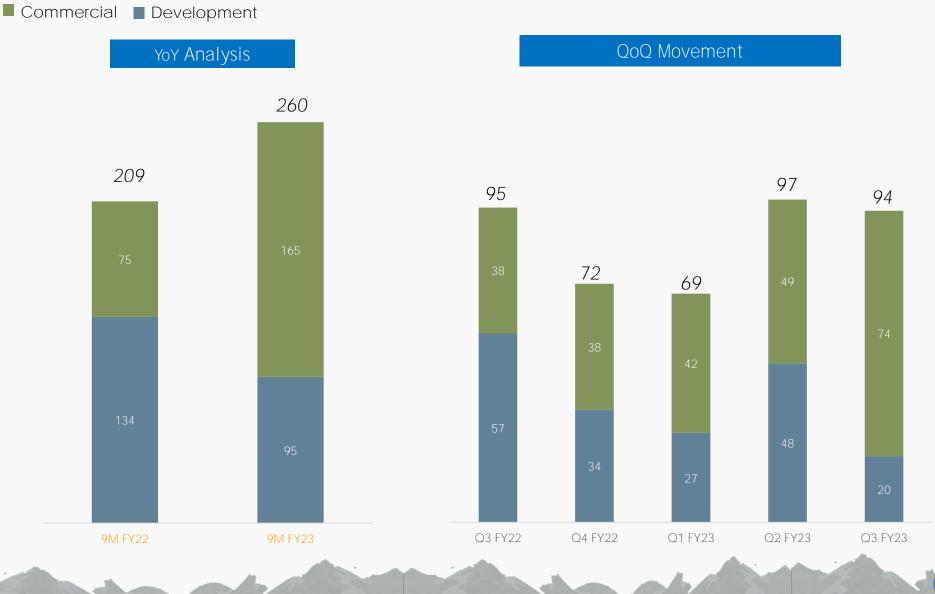
% represents CMS % from Overall revenue for respective Quarter / YTD

# Key Operating Metrics - CMS Revenue Split





In Cr



### Number of Active CMS Projects



Q3 FY-23	Pre-Clinical	P-1	P-2	P-3	Development	Commercial	Grand Total
API	17	4	7	5	8	9	50
Intermediate	10	4	4	2	7	12	39
Grand Total	27	8	11	7	15	21	89
Q3 FY-22	Pre-Clinical	P-1	P-2	P-3	Development	Commercial	Grand Total
API	15	3	8	5	10	7	48
Intermediate	7	5	2	0	8	11	33
Grand Total	22	8	10	5	18	18	81
Q3 FY-21	Pre-Clinical	P-1	P-2	P-3	Development	Commercial	Grand Total
API	14	4	6	3	10	6	43
API Intermediate	14 7	4	6 2	3 3	10 8	6 9	43 33
Intermediate	7	4	2	3	8	9	33
Intermediate Grand Total	7 21	4 8	2 8	3 6	8 18	9 15	33 76
Intermediate Grand Total Q3 FY-20	7 21 Pre-Clinical	<b>4</b> <b>8</b> P-1	2 8 P-2	<b>3</b> 6 P-3	8 18 Development	9 15 Commercial	33 76 Grand Total



# BUSINESS OVERVIEW

# Our Journey – Key Milestones





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# Generic Drug Substance (GDS)



# We started as a Prime API manufacturer...

# ..Added complex molecules for Speciality products..



### Capability

3 US FDA and EU GMP compliant manufacturing facilities

✓ Collective capacity: ~860 KL

### **Business Approach**

- Work on molecules either with a business leadership approach or partnership with client
- Ensure uninterrupted supply with quality commitment



### Strategy Forward

- Maintain leadership position in key molecules
- Work on process optimization to improve yields, productivity and thus margins



### Capability

High end complex chemistry capabilities

- Backend support by R&D department
- Experience of hurdle free scale up



### **Business Approach**

 Work with leading companies and help them to meet their technical requirements while being competitive



### Strategy Forward

- Focus on niche APIs with complex chemistry
- ✓ File IP for non infringing processes

Robust manufacturing base placed on the foundation of quality and pureplay API commitment

# On path to being a preferred partner in CMS..





- Manufacturing API to customer specifications
- Designing and developing manufacturing processes
- Process optimization for competitiveness
- ✓ Complete CMC partner for the API
- ✓ Patent protection for processes

### **Business Approach**

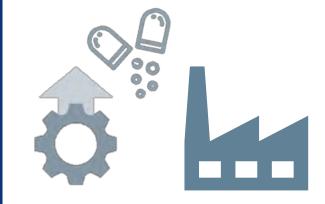
- ✓ Local presence in US, Europe and Japan with technical as well as commercial employees
- Consultative approach on customer relationships
- ✓ Business targeted on Neuland's technology capabilities and perceived customer needs leading to increased traction



### Strategy Forward

- Add depth in technical capabilities
- Investment in QBD labs, process engineering and foray into new areas of customer solutions
- ✓ Work effectively on customer relationships and leverage on portfolio expansion
- ✓ Targeting molecules in the later stages of the clinical cycle

Create a sustainable CMS business that is driven by technology and strong customer relationships



# CAPABILITIES

### Scaled up Manufacturing Facilities over the years





Bonthapally, Hyderabad

UNIT-2 363 KL Pashamylaram, Hyderabad

Gaddapotharam, Hyderabad

Year of Establishment	1986	1994	2017*
Blocks	Block - 1, 2, 3, 4, H, KL & S	34 Block-1, 2, 3, FC, NMSM, Mini plant 6	Block - 1, 2, 4, 5
Hydrogenation Reaction Volume	7.4KL	6 KL	Facility creation under process
Solvent Recovery System	100KLD	20KLD	50KLD
Cryogenic Reaction Volume	25KL	15 KL	15KL
Regulatory	USFDA, EDQM, CFDA, PMDA, et. al	USFDA, EDQM, PMDA, ANVISA et. al	Desktop Inspection by USFDA in 2020; ANVISA (Brazil) 2022

Adding capacities for new CMS projects and growth of key GDS molecules

# Backed up by sound R&D capabilities





### Infrastructure

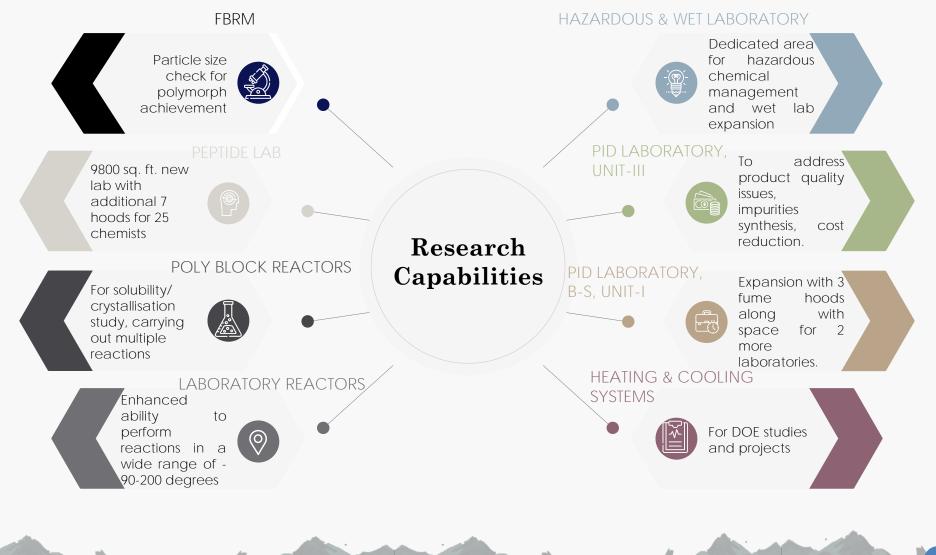
- 15 Development Labs with space for expansion
- 60 Fume hoods
- Analytical Labs
- Dedicated kilo Lab for Scale up
- Dedicated Labs for Peptides
- Separate facility for D2 analogues

### Significant R&D Achievements

- Several NCE APIs added in NDA or commercial stage
  drugs
- Support for multiple APIs each year in Phase 2
  and Phase 3 clinical candidates
- Generic API business -
- ✓ 900+ DMFs filed
- ✓ 300+ API processes developed
- ✓ 204+ patents filed. Received USPTO patent for improved process synthesis of Paliperidone Palmitate

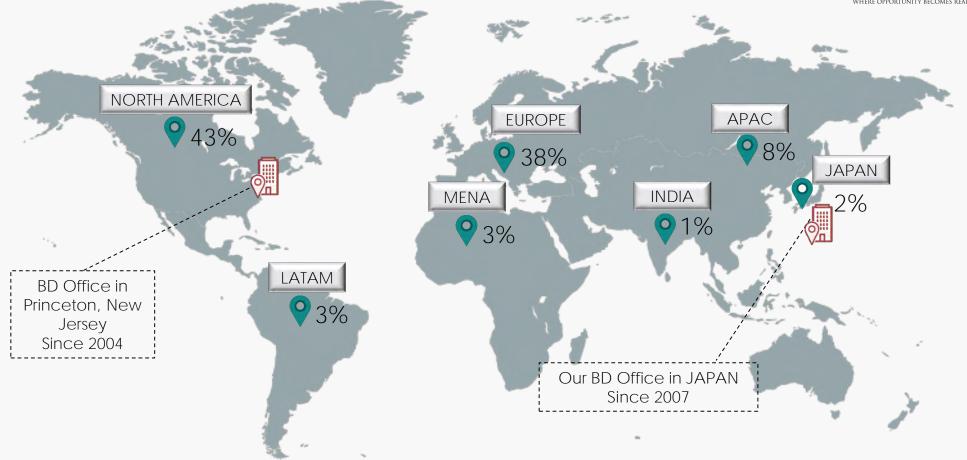
# New capabilities built





# **Global** Presence



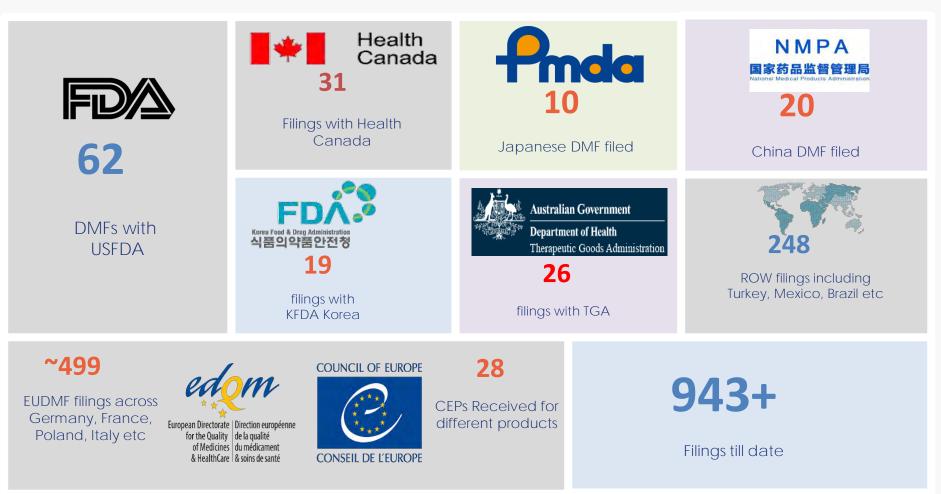


% Refers to 9M FY23 Sales by End market

25

# Regulatory Filings Across Geographies





\*\* The numbers on this slide reflect the number of filings, the number of active filings could vary as geographic filings are merged and changes in product portfolio.



# FINANCIALS

# Continuous Growth...



### Rs. In Cr



### FINANCIAL PERFORMANCE HIGHLIGHTS

UOo.

- Revenue CAGR of 12.5% for FY 19-22 led by growth in all 3 businesses
- EBITDA growth of 33.0% CAGR in FY 19-22 due to balanced contribution from both GDS and CMS business
- Shift to CMS and Specialty in overall revenue mix along with resource efficiency steps accelerated profitability

\*\* FY21 included other income of Rs. 13.09 crores towards profit on sale of investment property

\* This was after a one-time tax charge of Rs. 23.2 Cr in Q4FY20 that the Company chose to exercise under Section 115BAA of the IT act

# Stable Balance Sheet..

NEULAND WHERE OPPORTUNITY RECOMES REALITY

Particulars (Rs. Cr)	Mar-20	Mar-21	Mar-22	Dec-22
Shareholde rs' funds	706	782	836	906
Net Debt	214	152	212	72
Investments	8	7	4	2
Tangible Assets	391	438	497	508
Intangible				





Tangible Assets	391	438	497	508
Intangible Assets (Excluding Goodwill)	2	3	2	2
Working Capital	289	309	382	391



### Debt to Equity (x)



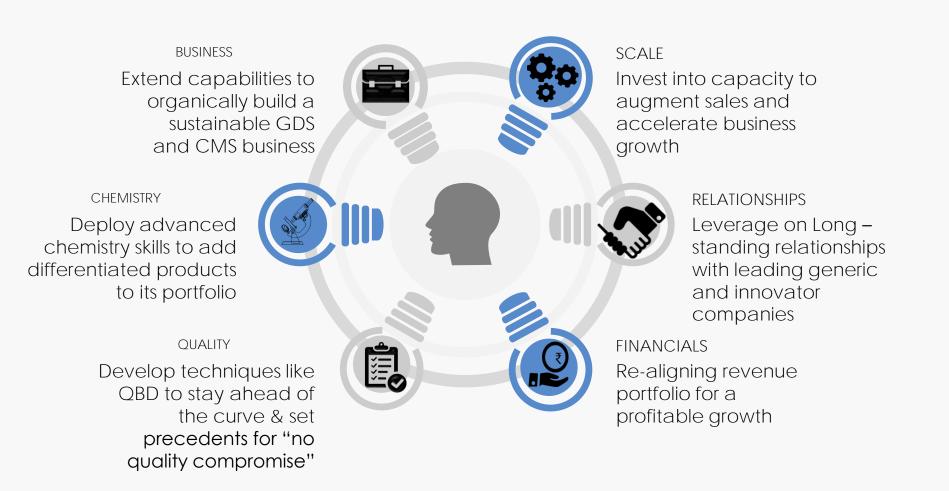
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# ..Laying Foundation for our Growth Strategy

### CREATE AN ORGANIZATION THAT RESULTS IN VALUE FOR ALL STAKEHOLDERS



**9 NEULAN** WHERE OPPORTUNITY BECOMES RE

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# Contact Us



For over 39 Years, Neuland Laboratories Ltd. (BSE:524558, NSE: NEULANDLAB) has been at the forefront of manufacturing APIs through its cGMP manufacturing facilities, working with customers in close to 80 countries.

Neuland Labs has developed more than 300 processes and 75 APIs and has filed over 900+ Regulatory filings in the US (62 active US DMFs), the European Union (EU) and other geographies. Its manufacturing facilities are inspected and approved by the U.S. FDA and other leading regulatory agencies. Its record of quality manufacturing and reliability is highlighted by cGMP certifications that include the U.S. FDA, TGA (Australia), EDQM (EU), German Health Authority, ANVISA (Brazil), EMA (EU), Cofepris (Mexico), KFDA (Korea), PMDA (Japan), CFDA (China), FSI "SID &GP" Russia, Health Canada, ISO 9001, ISO14001, OHSAS18001 and ISO 27001. For further information contact

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Thank You

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