



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

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[www.neulandlabs.com](http://www.neulandlabs.com)

December 8, 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: NEULANLAB; Series: EQ**

Dear Sir/Madam,

**Sub: Press Release**

We are enclosing Press Release on the sale of shares by Dr. Davuluri Rama Mohan Rao, Founder & Promoter and Executive Chairman of Neuland Laboratories Limited.

This is for your information and records.

Thanking you,

Yours Sincerely,

**For Neuland Laboratories Limited**

**Sarada Bhamidipati**  
**Company Secretary**

*Encl: As above*



**Hyderabad, India, December 8, 2023** - Neuland Laboratories Limited (NLL) (NSE: NEULANLAB; BSE:524558), a pharmaceutical manufacturer providing active pharmaceutical ingredients (APIs), complex intermediates and custom manufacturing solutions services to customers located in around 80 countries, has been informed that Dr. Davuluri Rama Mohan Rao, Founder & Promoter and Executive Chairman of Neuland Laboratories Limited (the Company) has sold 4,00,000 equity shares of the Company (i.e. 3.12 % stake) today through open market at an average price of Rs. 5,012.45.

Dr. Rao has informed that he intends to use the funds generated from this sale for personal purposes. Davuluri family does not plan to sell any further shares in the foreseeable future and are fully committed and invested in the future of Neuland Laboratories Limited.

### **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 100 APIs and has filed over 950+ Regulatory filings in the US (65 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](http://www.NeulandLabs.com).

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

***IR Department at Neuland***

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***Ravi Udeshi, EY IR***

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