



**REGISTERED OFFICE**

**GRANULES INDIA LTD.**, 2nd Floor, 3rd Block, My Home Hub,  
Madhapur, Hyderabad – 500 081, Telangana, INDIA.

Tel: +91 40 69043500, Fax: +91 40 23115145, mail@granulesindia.com, www.granulesindia.com

CIN: L24110TG1991PLC012471

**Dated December 30, 2021**

To,  
National Stock Exchange of India Limited  
BSE Limited  
Symbol: NSE: GRANULES; BSE: 532482

**Sub: Press Release**

Dear Sir,

We are herewith enclosing the press release given by the Company.

This is for your information and dissemination to the members of the exchange.

Thanking You.

Yours sincerely,

**For GRANULES INDIA LIMITED**

*G. Chaitanya.*



**CHAITANYA TUMMALA  
(COMPANY SECRETARY &  
COMPLIANCE OFFICER)**



**Press Release**

**For Immediate Release**

## **Granules Pharmaceuticals, Inc. Receives ANDA Approval for Amphetamine Mixed Salts (IR Tablets)**

**Hyderabad, 30 December 2021:** Granules India Limited announced today that the US Food & Drug Administration (US FDA) has approved the Abbreviated New Drug Application (ANDA) filed by Granules Pharmaceuticals, Inc. (GPI), a wholly owned foreign subsidiary of Granules India Limited, for Amphetamine Mixed Salts, USP 1.25mg, 1.875mg, 2.5mg, 3.125mg, 3.75mg, 5mg, 7.5mg Immediate-Release Tablets. It is bioequivalent to the reference listed drug product (RLD), *Adderall*® of Teva Women's Health Inc. The product would be available for the US market shortly.

Commenting on the approval **Ms. Priyanka Chigurupati, Executive Director, GPI**, said *“We are pleased to receive the approval of the controlled substance product and will surely be a valuable addition to our growing product portfolio in the US market. We will be launching the product shortly.”*

Granules now have a total of 47 ANDA approvals from the US FDA (45 Final approvals and 2 tentative approvals).

The current annual U.S. market for Amphetamine Mixed Salts, USP 1.25mg, 1.875mg, 2.5mg, 3.125mg, 3.75mg, 5mg, 7.5mg Immediate Release Tablets strengths is approximately \$ 335 Million, according to MAT, Oct 2021 IQVIA/IMS Health.

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## **About Granules India Ltd. (BSE: 532482, NSE: GRANULES)**

Granules India Limited, incorporated in 1991 is a vertically integrated fast growing Indian pharmaceutical company headquartered at Hyderabad with best in class facilities and commitment to operational excellence, quality, and customer service. We are among the few pharmaceutical companies in the world to be present in the manufacturing of entire value chain – from Active Pharmaceutical Ingredients (APIs), Pharmaceutical Formulation Intermediates (PFIs) and Finished Dosages (FDs). Our products are being distributed to over 300+ customers in regulated and semi-regulated markets with a global presence extending to over 75+ countries with offices across India, U.S. and U.K. The Company has 7 manufacturing facilities out of which 6 are located in India and 1 in USA and has regulatory approvals from US FDA, EDQM, EU GMP, COFEPRIS, WHO GMP, TGA, K FDA, DEA, MCC and HALAL.

**Safe Harbor:** This document is to provide the general background information about the Company's activities as at the date of the release. The information contained herein is for general information purposes only and based on estimates and should not be considered as a recommendation that any investor should subscribe / purchase the company shares. The Company makes no representation or warranty, express or implied, as to, and does not accept any responsibility or liability with respect to, the fairness, accuracy, completeness or correctness of any information contained herein. This release may include certain "forward looking statements". These statements are based on current expectations, forecasts and assumptions that are subject to risks and uncertainties which could cause actual outcomes and results to differ materially from these statements. Important factors that could cause actual results to differ materially from our expectations include, amongst others general economic and business conditions in India, ability to successfully implement our strategy, our research and development efforts, our growth and expansion plans and technological changes, changes in the value of the Rupee and other currencies, changes in the Indian and international interest rates, change in laws and regulations that apply to the Indian and global pharmaceuticals industries, increasing competition, changes in political conditions in India or any other country and changes in the foreign exchange control regulations in India. Neither the company, nor its directors and any of the affiliates or employees have any obligation to update or otherwise revise any forward-looking statements. The readers may use their own judgment and are advised to make their own calculations before deciding on any matter based on the information given herein. No part of this release may be reproduced, quoted or circulated without prior written approval from Granules India Limited.

### **Investor Contacts:**

<b>Krishna Raghunathan</b>	<b>Chaitanya Tummala</b>
Vice President – Finance and Investor Relations	Company Secretary
040-69043573	040-69043614
<a href="mailto:krishna.raghunathan@granulesindia.com">krishna.raghunathan@granulesindia.com</a>	<a href="mailto:chaitanya.tummala@granulesindia.com">chaitanya.tummala@granulesindia.com</a>