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Indian Pharmaceuticals Company Taj Pharmaceuticals receives FDA approval for Generic Nortriptyline

Mumbai - 19th June 2009 - Long Awaited approval for Nortriptyline given to Taj Pharmaceuticals from FDA to manufacture generic version



Taj Pharma Group announces
USFDA tentative approval for Various
generic Medicines

19th June 2009, Indian Pharmaceuticals Company Taj Pharmaceuticals based in Mumbai announced FDA approval for Nortriptyline Generic drug manufacturing for regulatory market and non-regulated market.

The company is said to have approvals for different combinations of the drug, i.e. Nortriptyline hcl 25mg.

The generic manufacturing strategies of New Drug Nortriptyline and the product-process of Taj Pharmaceuticals Ltd. are compared and collaborated using data from global Pharmaceuticals manufacturing plants in the UK(Newcastle), US, Korea, Italy, Germany and Japan.

The Indian company is eminent manufacturer, supplier and exporter of Nortriptyline and various pharmaceuticals generics medicines in Tablets, capsules, Vial (injections), ampule, Strips and schedule packs etc.

The companies integrated research team facilitates the affordable production of the drug for even non-regulated markets of the world. i.e. Middle East, Thailand, Bangladesh etc.

The company is also involved in manufacturing of Active Ingredient of various pharmaceuticals finished formulations in IP /BP / USP grades for all markets.

The approvals bring a positive uplift in the share market and public perception for Pharmaceuticals investors.



About Taj Pharmaceutical Limited

Taj Pharmaceuticals Limited is a pharmaceutical company founded and based in India. The company manufacturers pharmaceutical formulations and API for India and other countries of world. The company was established in 1995 as an enterprise and in 2004 became a public limited company. As per Mumbai pharmaxil and Chemixil association the company manufacturers and exports to countries like Albania, Argentina, Austria, Chile and Iraq. In 1995 pharmaceuticals wing only has a schedule M certification for pharmaceuticals products manufacturing in India. Taj Pharmaceuticals established its manufacturing unit in Gujarat because of government policies in 1999 with WHO / GMP licence. The company in 2003 revived all the old manufacturing units and approached the **FDA Gujarat for 4000 new pharmaceuticals drug permissions** for the first time in India.

According to the Indian Trade Mark the company owns about **450 brands and 4600 generic** manufacturing permissions in India. According to the export data analysis the company was the largest exporter of generic medicines to the Europe and Middle East countries.

www.tajpharma.com

The company medicines are present in France, Georgia, Egypt and CIF countries.

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