Jubilant Life Sciences Ltd.
1A, Sector 16A, Noida – 201301, India
Tel.: +91 120 4361000
www.jubl.com

Jubilant Life Sciences receives ANDA approvals for Valsartan Tablets

Noida (UP), India, Tuesday, January 06, 2015

Jubilant Life Sciences Ltd, an integrated Pharmaceuticals and Life Sciences Company, announced today that it has received Abbreviated New Drug Application (ANDA) final approval from the US Food and Drug Administration (US FDA) for **Valsartan Tablets USP, 40 mg, 80 mg, 160 mg and 320 mg**, the generic version of Diovan® (of Novartis), used as an anti-hypertensive. We expect to launch this product immediately. The current annualized US market size for Valsartan Tablets USP, 40 mg, 80 mg, 160 mg, and 320 mg as per IMS is USD 2 Billion.

As on September 30, 2014, Jubilant Life Sciences had a total of 781 filings for formulations of which 322 have been approved in various regions globally. This includes 72 ANDAs filed in the US and 46 Dossier filings in Europe.

About Jubilant Life Sciences

Jubilant Life Sciences Limited is a global Pharmaceutical and Life Sciences Company engaged in manufacture and supply of APIs, Generics, Specialty Pharmaceuticals and Life Science Ingredients. It also provides Services in Contract Manufacturing and Drug Discovery. The Company's strength lies in its unique offerings of Pharmaceutical and Life Sciences products and services across the value chain. With 10 world-class manufacturing facilities in India, US and Canada and a team of about 6,200 multicultural people across the globe, the Company is committed to deliver value to its customers spread across over 100 countries. The Company is well recognized as a 'Partner of Choice' by leading pharmaceuticals and life sciences companies globally.
For more info: www.jubl.com.

For more information please contact:

For Investors

Ravi Agrawal
Jubilant Life Sciences Limited
Ph: +91-120 436 1002
E-mail: ravi_agrawal@jubl.com

Anupam Jain
Jubilant Life Sciences Limited
Ph: +91-120 436 1021
E-mail: anupam_jain@jubl.com

Siddharth Rangnekar
Citigate Dewe Rogerson
Tel: +91 22 6645 1209
E-mail: siddharth@cdr-india.com

Karl Kolah
Citigate Dewe Rogerson
Tel: +91 22 6645 1220
E-mail: karl@cdr-india.com

For Media

Neha Garg
Jubilant Life Sciences Limited
Ph: +91-120 436 1067
E-mail: neha_garg@jubl.com

Saurabh Gupta
Perfect Relations
Ph: +91 9818075578
E-mail: sgupta@perfectrelations.com

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