



Strides Arcolab Limited,
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Press Release

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Strides Arcolab receives ANDA approval from US FDA for

Polyethylene Glycol 3350, Sodium Chloride, Sodium Bicarbonate and Potassium Chloride for Oral Solution

Product to be launched shortly

Bangalore, April 15, 2015, Strides Arcola Limited today announced that it has received approval from the United States Food & Drug Administration (USFDA) for Polyethylene Glycol 3350, Sodium Chloride, Sodium Bicarbonate, and Potassium Chloride for Oral Solution, 420 grams/11.2 grams/5.72 grams/1.48 grams/4 Liters.

According to IMS data, the US market for Polyethylene Glycol 3350, Sodium Chloride, Sodium Bicarbonate, and Potassium Chloride for Oral Solution is approximately USD 20 Million.

The product will be manufactured at the Company's Oral dosage facility at Bangalore and marketed by Strides in the US Market.

About Polyethylene Glycol 3350, Sodium Chloride, Sodium Bicarbonate, and Potassium Chloride for Oral Solution

It is used for bowel cleansing prior to colonoscopy. Polyethylene glycol 3350 is an osmotic laxative, and along with the electrolytes, the combination is used in preparation for colonoscopy in adults and pediatric patients aged six months or greater.

About Strides Arcola Limited

Strides Arcolab, listed on the Bombay Stock Exchange Limited (532531) and National Stock Exchange of India Limited (STAR), is a global pharmaceutical Company headquartered in Bangalore, India that develops and manufactures a wide range of IP-led niche pharmaceutical products. The Company has 8 manufacturing facilities presence in more than 75 countries in developed and emerging markets. Additional information is available at the Company's website at www.stridesarco.com.



For further information, please contact:

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