Aurobindo receives Tentative approval for Quinapril Hydrochloride and Hydrochlorothiazide Tablets from USFDA

Aurobindo Pharma Limited is pleased to announce that the company has received tentative approval from the USFDA for Quinapril Hydrochloride and Hydrochlorothiazide tablets 10mg/12.5mg, 20mg/12.5mg and 20mg/25mg. With this, Aurobindo's product basket for the USA market has gone up to forty five.

This is indicated for the treatment of hypertension.

Quinapril Hydrochloride and Hydrochlorothiazide tablets 10mg/12.5mg, 20mg/12.5mg and 20mg/25mg is the generic equivalent of ACCURETIC™ tablets (Pfizer Pharmaceuticals Ltd.,). This is the fifth anti hypertensive (Medicines regulating blood pressure) of Aurobindo bagging US FDA's nod.

About Aurobindo Pharma Limited:

Aurobindo Pharma Limited, headquartered at Hyderabad, India, manufactures generic pharmaceuticals and active pharmaceutical ingredients. The company has received approvals for most of the targeted API and formulation facilities from leading regulatory agencies like US FDA, UK MHRA, WHO, Health Canada, MCC South Africa.

The company's robust product portfolio is spread over 6 major therapeutic/product areas encompassing Antibiotics, Anti-Retrovirals, CVS, CNS, Gastroenterologicals, and Anti-Allergics.

The formulation portfolio is built on six technology platforms such as immediate release generics, SR/CR generics, and Orally disintegrating/Mouth dissolving generics, Combination products, Sterile/Lyophilized generics and Liquids/Dry syrups. Aurobindo has filed over 110 DMFs and 90 ANDAs for the USA market alone in addition to filings in other countries.

The pace of filings is matched by rapid product approvals from various markets. Till date Aurobindo received 45 ANDA approvals (both final and tentative) from US alone. Aurobindo operates in over 100 countries and markets over 180 APIs and 250 formulations.