JUBILANT PHARMA

ACTIVE PHARMACEUTICAL INGREDIENTS

Active Pharmaceutical Ingredients (APIs)

Jubilant API business is a preferred partner of choice for innovator and generic pharmaceutical companies across the globe. We have a prominent presence in markets such as North America, Europe, Latin America, Japan, Asia Pacific and Middle East.

In R&D, our focus continues to be on developing commercially competitive, IP compliant, robust and eco-friendly technologies. Our R&D thrives on 'green chemistry culture' and has developed various environmentally friendly & disruptive technologies such as continuous processes, incorporating optimum atom efficiencies, recycling and reuse of solvents/reagents/by-products targeting towards zero discharge of effluents, substitution/ minimization of hazardous chemicals and replacing chemical processes with enzymatic/ chemo catalysis processes.

JUBILANT EDGE

Business Overview

- Jubilant offers one of the broadest portfolio comprising **more than 90 different APIs** from therapeutic categories such as CNS, CVS, anti-infective, anti-diabetic, respiratory etc.
- Amongst the **world's largest capacities for APIs** such as Carbamazepine, Oxcarbazepine, Citalopram, Lamotrigine, Donepezil, Meclizine, Pinaverium, Escitalopram, Azithromycin, Risperidone & Valsartan.
- Proven expertise to operate **large scale chemical operations**, key to better cost efficiencies
- Diversified & large external customer base to drive growth across multiple regions
- Business sustainability with backward integration of key APIs with in-house KSMs

API R&D

State of the art R&D centres in India at Noida and Nanjangud, Mysore

Expertise in complex chemistries like Chiral separation, Low Temp reactions, Bio-transformation, Stereo-selective synthesis, Process intensification (continuous flow reactions)

API MANUFACTURING EXPERTISE

Our manufacturing facility at Nanjangud, Mysore, India is a state-of-the-art API manufacturing facility fully compliant with cGMP, quality & safety requirements.

Experienced team of over 900 personnel

Globally approved by major regulatory bodies including, US-FDA, PMDA- Japan, Health Canada, ANSM France, ANVISA-Brazil, TGA- Australia, COFEPRIS Mexico, KFDA and CDSCO.

267, 688 sq. mts. site with 33% built-up area, 6 multi-stream manufacturing blocks and a pilot plant having volumetric capacity of ~760 KL, 18 clean rooms and 170 reactors ranging from 0.1 KL to 15 KL.

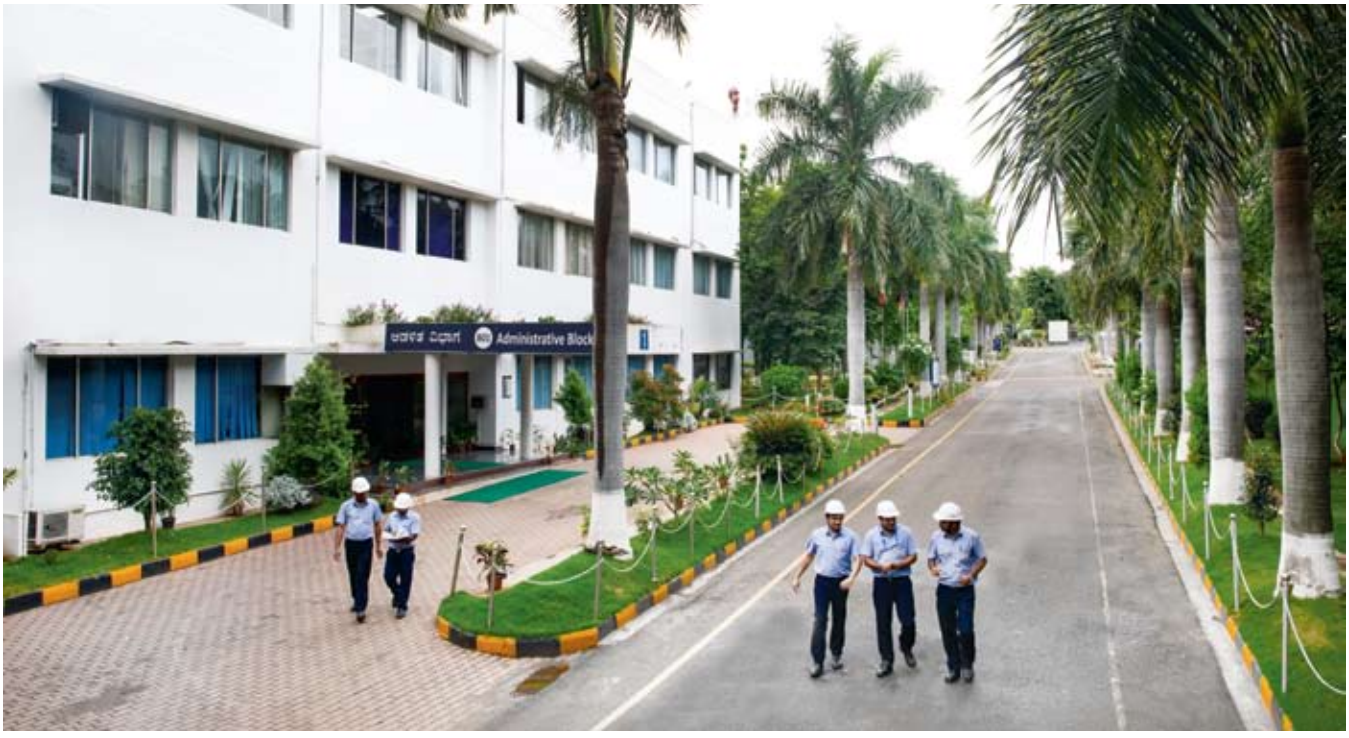
Dedicated DoE/ QbD cell for bringing quality and process robustness during the product development.

Team of more than 150 dedicated scientists including PhDs

Equipped with latest equipment's like LC HRMS, NMR, XRD, LCMS, GCMS, ICP-MS, etc.

Dedicated Project Manager for Cost & Time management.

Experts in polymorphic characterization and contamination, Genetoxic & Carry over studies, Impurity profiling etc.



Manufacturing facility at Nanjangud, Mysore, India

Office Locations:

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