

**For Immediate Press Release**

**Inogen Laboratories clears the Pharmaceuticals and Medical Devices Agency (PMDA), Japan Audit**

---

Hyderabad, August 21 2012: Inogen Laboratories (Inogen), a GVK BIO associate company announced today that its GMP compliant facilities in Hyderabad were audited and approved by the Japanese regulator, **Pharmaceuticals and Medical Devices Agency (PMDA)**. PMDA conducts scientific reviews of marketing authorization application of pharmaceuticals and medical devices and monitoring of their post-marketing safety.

Kiran Kumar R, President, Inogen Laboratories said “A four member team from PMDA, Japan visited and audited Inogen’s facility, processes and quality management system and approved us without any major observations. We are pleased with the outcome of this audit and expect greater penetration into the Japanese market.”

**About Inogen Laboratories Private Limited**

Inogen Laboratories Private Limited is ISO 9001 & WHO cGMP certified and is a GVK Biosciences (GVK BIO) associate company based out of Hyderabad, India. Inogen offers an attractive mix of Services (Process R&D and Custom Chemical Synthesis and Manufacturing) and Products (APIs and Intermediates) to its clients. The Company has state-of-the-art process and analytical R&D laboratories, pilot plant, kilo lab, custom synthesis plant and two production blocks. Inogen works with pharmaceutical, generic and biotech companies who have audited and approved its manufacturing and scale up facilities.

For any further information, please contact:

Sharada Alvakonda  
Dy. General Manager – Corporate Communications  
GVK Biosciences Private Limited  
E [sharada@gvkbio.com](mailto:sharada@gvkbio.com)