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PRESS RELEASE

CAPLIN POINT GETS EU GMP APPROVAL FOR STERILE INJECTABLE SITE

CHENNAI, JULY 08, 2015: Caplin Point has received EU GMP approval for its small volume sterile injectable liquids facility at Chennai. The site was inspected by the EU authorities in March 2015 and has been certified as compliant with the guidelines stated in Art. 111 (5) of Directive 2001/83/EC. The facility is capable of manufacturing Liquid Injectables in Vials, Ampoules, Lyophilized Vials and Ophthalmic dosages.

Commenting on the EU certification, Mr.C.C.Paarthipan, Chairman, Caplin Point said "This is our first approval and we're expecting more regulatory GMP approvals in future. The global injectable market is projected to grow to US\$70 billion by 2020 and is frequently plagued by shortages. This presents an exciting opportunity for us to carve out a niche."

EU approval grants access not only to the European Union, but also several other countries in the Regulated domain through mutual recognition process.

ABOUT CAPLIN POINT LABORATORIES LIMITED:

Caplin Point Laboratories Limited is a niche pharmaceutical company with a unique business model catering predominantly to emerging markets of Latin America and Africa.

Caplin Point has four state of the art manufacturing facilities that cater to a complete range of finished dosage forms. The Company currently holds over 1600 product licenses across Latin America and Africa with another 400 in the pipeline. The company will be filing their first ANDAs by end of 2015.

Company has also recently added Soft Gelatin Capsules, Suppositories and Carbapenem lines to its existing facilities, for emerging markets.



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