



March 21, 2013

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ISO13485 Approval: International Standard of Quality Management System

The company hereby announces that the certificate of registration of ISO13485, an international standard of quality management system for medical devices, was received as of March 21, 2013, which contributes to global expansion of our business with hemostat (develop code: TDM-621).

ISO13485 of which the company acquired the approval is a quality management system to manufacture and provide medical devices that consistently meet safety and efficacy requirements. This approval of ISO13485 means that our development system, manufacture and marketing system, and quality assurance system relating to TDM-621 satisfies the international standard. Since the approval of ISO13485 is mostly required upon selling medical devices globally, our preparing to export of TDM-621 is promoted.

We will continue to make a contribution to medicine by developing and manufacturing high quality products using with advanced technology.

This approval does not influence the earning forecast of the company at this moment.