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[Delayed] Surgical Hemostat (TDM-621): Application for CE Marking

(This original disclosure in Japanese was released on May 20, 2013 at 16:00 (GMT+9))

The company hereby announces that our subsidiary, 3-D Matrix Europe SAS. submitted application documents to a third-party notified body of CE Marking on May 17, 2013, which is required to sell our surgical hemostat "TDM-621" in Europe.

CE Marking which our group applies for is a required indication to distribute medical device within the European market. A certification that the product and the quality management system are satisfied with EU legislation (Medical Device Directive) is needed to obtain CE Marking. Modular submission of application documents is approved by the notified body.

After CE Marking certification is obtained, our group can start to sell TDM-621 in EU and have a plan to perform clinical trial on a small scale in order to expand sales in EU market.

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