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Absorbable Hemostat "TDM-621" Product Registration Obtained in Brazil

The 3-D Matrix group is working towards commercialization of absorbable hemostat "TDM-621" globally. The company hereby announces that the subsidiary, 3-D Matrix Europe SAS., has received notification today on November 4th that the medical device product registration approval in Brazil has been granted by the Brazilian regulatory body, Agência Nacional de Vigilância Sanitária (ANVISA).

The group has obtained CE marking for the absorbable hemostat "TDM-621" on January 14, 2014. This CE marking can be leveraged by using it as a reference regulatory agency approval in various countries in Asia-Pacific and Latin America. The product can be commercially marketed once approval is obtained in each country.

With this medical device product registration approval, our commercialization activities in Brazil can start. The company is selecting the local partner to distribute "PuraStat®" through the partner to start commercializing by the end of this fiscal year.

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