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Absorbable Hemostat PuraStat®
Product Registration Obtained in Mexico

The 3-D Matrix group is working towards commercialization of absorbable hemostat PuraStat® globally. The company hereby announces that the subsidiary, 3-D Matrix Europe SAS., (3DMEU) has received notification that the medical device product registration approval in Mexico has been granted by the Mexican regulatory body, COMISIÓN FEDERAL PARA LA PROTECCIÓN CONTRA RIESGOS SANITARIOS (COFEPRIS).

The group has obtained CE marking for the absorbable hemostat PuraStat® 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, marketing and sales activities in Mexico can start.

3DMEU had already entered into a partnership agreement including minimum purchase amount with Genelife S.A. (“Genelife”, head office located at Miguel Laurent 17, Del Valle, Benito Juarez, Mexico City DF, Mexico, Mexico) for sales and marketing of PuraStat® in Mexico as of February 18, 2016. With this COFEPRIS approval, 3DMEU will start sales through Genelife this fourth quarter of FY2015. The 3-D Matrix group will conduct marketing activities jointly with Genelife to expand product sales in Mexico.

This announcement does not influence the earning forecast of the company at this moment. However, we will make necessary announcements when it is deemed that there is any influence.