

BONESUPPORT announces - De Novo application for CERAMENT G has been submitted to the FDA

Lund, Sweden, 17.30 CET, 17 April 2020 — BONESUPPORT™, an emerging leader in orthobiologics for the management of bone injuries today announced that the company has submitted a De Novo application to the US Food and Drug Administration (FDA) to obtain a market approval for the company's antibiotic-releasing product CERAMENT® G. The application is specified for the indication osteomyelitis (bone infection) and can potentially result in an approval at the latter end of 2020.

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About BONESUPPORT™

BONESUPPORT (Nasdaq Stockholm: BONEX) develops and commercializes innovative injectable bio-ceramic bone graft substitutes that remodel to the patient's own bone and have the capability of eluting drugs. BONESUPPORT's bone graft substitutes are based on the patented technology platform <u>CERAMENT</u>. The company is conducting several clinical studies to further demonstrate the clinical and health economic benefits that its products deliver and a Premarket approval filing with the FDA (USA) for <u>CERAMENT G</u> is planned in 2021. The company is based in Lund, Sweden, and the net sales amounted to SEK 155 million in 2019. Please visit <u>www.bonesupport.com</u> for more information.

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