



Product Service

# EC Certificate

## EC Design-Examination Certificate

Directive 93/42/EEC on Medical Devices (MDD), Annex II (4)  
(Devices in Class III)

No. G7 17 11 32283 052

**Manufacturer:** **Biomatlante SA**  
**ZA les Quatre Nations**  
5, Rue Edouard Belin  
44360 Vigneux de Bretagne  
FRANCE



**Product:** **Bone substitutes**

**Model(s):** **Injectable paste (dental / maxillofacial)**

**Parameters:** see attachment

The Certification Body of TÜV SÜD Product Service GmbH declares that a design examination has been carried out on the respective devices in accordance with MDD Annex II (4). The design of the devices conforms to the requirements of this Directive. For marketing of these devices an additional Annex II certificate is mandatory. See also notes overleaf.

**Report no.:** 713105279

**Valid from:** 2017-12-19

**Valid until:** 2022-11-26

**Date,** 2017-12-19

Stefan Preiß



TÜV SÜD Product Service GmbH is Notified Body with identification no. 0123

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