

## ATTESTATION CE / EC CERTIFICATE

Examen de type / Type Examination

ANNEXE III de la Directive 93/42/CEE relative aux dispositifs médicaux

ANNEX III Directive 93/42/EEC concerning medical devices

Fabricant / Manufacturer

**ANIKA THERAPEUTICS S.R.L.**

**Corso Stati Uniti 4/U**

**CAP 35127 PADOVA (PD) ITALY**

Catégorie du(des) dispositif(s) / Device(s) category

**Support pour culture de cellules mesenchymateuses**

*Scaffold for mesenchymal stem cells*

Identification du(des) dispositif(s) / Identification of device(s)

**HYALOFAST**

**(GMDN 46426)**

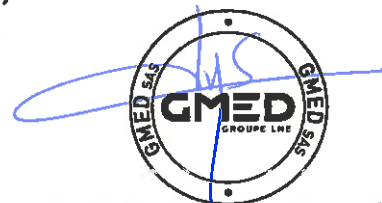
**voir addendum/see addendum**

GMED atteste qu'à l'examen des résultats figurant dans le(s) rapport(s) référencé(s) P601294/P169669-2, un échantillon représentatif de la production est conforme aux exigences de l'annexe I de la Directive 93/42/CEE

GMED certifies that, on the basis of the results contained in the file(s) referenced P601294/P169669-2, a representative sample of the production complies with the requirements of the Directive 93/42/EEC, annex I

Début de validité / Effective date : **October 3rd, 2019 (included)**

Valable jusqu'au / Expiry date : **October 14th, 2022 (Included)**



**On behalf of the President**

**Béatrice LYS**

**Technical Director**

**Identification des dispositifs / Identification of devices**

144714F	box containing 1 x (2 cm x 2 cm) pad
144712F	box containing 1 x (2 cm x 2 cm) pad (Turkey)
144720F	box containing 1 x (2 cm x 2 cm) pad (Spain)
144722F	box containing 1 x (5 cm x 5 cm) pad
144723F	box containing 1 x (5 cm x 5 cm) pad (Turkey)

**GMED 0459**



**On behalf of the President  
Béatrice LYS  
Technical Director**