

Full Quality Assurance System
Directive 93/42/EEC on Medical devices, Annex II excluding (4)

CE Certiso Ltd. (NB 2409) certifies that the following manufacturer's quality management system concerning to the listed devices and device categories meets the requirements of the related requirements of the directive.

Name of the manufacturer:

Juves Sarl

Headquarters: **52 Boulevard Sébastopol, 75003, Paris, France**

Scope:

Dermal fillers

The certificate covers the following devices:

Description of the device	Type	Intended use	Model	Risk class
calcium hydroxyapatite dermal implant	Simetro	sub-dermal and deep dermal use	Simetro packed with or without needle	III*
crosslinked hyaluronic acid dermal filler	FigurHA	sub-dermal and deep dermal use	FigurHA Initio FigurHA Intenso FigurHA Volumo FigurHA LipContour FigurHA Eyelights	III*

*In case of devices in Class III, this certificate independently does not authorize the manufacturer for the use of CE mark on the devices.

This certificate is valid only in case of successfully conducted annual surveillance audits.

ID number of the related audit report: 140-CE-190212

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