AKRA DERMO JET		SMQ	
Forme 17	Num. Enr.	Date 25/11/2019	Nombre de Pages 1
Titre/Objet	Declaration of CE conformity		

The company :

AKRA DERMOJET 23 Bis Rue Louis Barthou
64000 PAU - FRANCE

Hereby certifies that the medical device (Class IIa – rule 11):

The first CE mark was opposed in May, 25, 1998 for all Dermojet references

Descriptive summary	References	Year-to-market[1]
DERMOJET Standard *DERMOJET HR* *DERMOJET Polymedical*	100 200 500	1960 1989 1992

Meets the provisions of (Annex II of Directive 93/42/EEC. The certificate does not cover examination of the medical device design in accordance with Annex II Clause IV of directive 93/42/EEC

The approved quality system established at the manufacturer is subject to regular surveillance audits by the notified body in accordance with Annex II clause 5 of directive 93/42/EEC.

This declaration is based on the full quality assurance system certification (Annex II of Directive 93/42 EEC) (certificate 1144C04191101) delivered by HTCERT (Notified body number 2803),

Date/*Dated*: 25/11/2019

Responsable Qualité/*Quality Assurance Mgr*
Ungar Régis