Decleration of CE conformity

According to Annex VII to Directive 93/42 / EEC on medical devices

We,

SAS CONDOR

38 avenue du Languedoc RD6113

11200 Lézignan-Corbières – France

Guarantee and declare, under our sole responsibility, that the device:

CondorScan

Class I pursuant to Regulations 5 and 12 of Annex IX to European Directive 93/42 / EEC

Complies with the applicable requirements of Directive 93/42 /

This statement is based on the following:

- Technical dossier demonstrating compliance with the essential requirements of Directive 93/42 / EEC

This for the period of validity of the certificate, that is until: 01/04/2020.

Created at Lézignan-Corbières, on 13/07/2016

François Duret, President of Condor SAS

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