KF-CE-EN
On 20/01/2019

CE Declaration of conformity

In accordance with of European parliament and Council Decision 93/42/CEE, annex VII concerning medical devices

We,

SAS KINVENT BIOMECANIQUE

Cap omega - cs 39521 Rond-point Benjamin Franklin
34000 Montpellier
France

guarantee and declare, under our sole responsibility, that the system:

KFORCE

Composed of the devices

	UMDNS NAME	***UMDNS CODE***	***EAN CODE***	***HS code***
K-FORCE Plates	Biomechanics Platforms	17242	*3770011995226*	9031808000
K-FORCE Muscle Controller	Dynamometer Exercise Systems, Computerized	17681	*3770011995103*	9031808000
K-FORCE Bubble	Dynamometer Exercise Systems, Computerized	17681	*3770011995448*	9031808000
K-FORCE Grip	Dynamometer Exercise Systems, Computerized	17681	*3770011995332*	9031808000
K-FORCE Sens	Sensors	13536	*3770011995554*	9031802000
K-FORCE Link	Dynamometer Exercise Systems, Computerized	17681	*377001199004*	9031808000

Class I pursuant to Rule 1 of Annex IX of the European Directive 93/42 / EEC complies with the applicable requirements of Directive 93/42 / EEC and the French Public Health Code.

This declaration is based on the following element:

• Technical file KF-CEDTv2-EN demonstrating compliance with the essential requirements of Directive 93/42 / EEC

This for the whole duration of the certification's validity, hence until the 01/01/2022 (included).

Made at MONTPELLIER, FRANCE
on the 20/01/2019

Athanase KOLLIAS, CEO