EU DECLARATION OF CONFORMITY

Manufacturer: RESI Třeboň spol. s r.o.
Registered office: Novohradská 1153, 379 01 Třeboň, Czech Republic
Company ID: 25178989
Manufacturer's registration number: 009259

This EU Declaration of Conformity is issued on the sole responsibility of the manufacturer.

Medical device name: JORDAN F
Device models: F2e, F2h, F2e Face, F2h Face, F3e, F3 Prima, F3h, F3e Face, F3h Face, F4e, F4h, F4e face, F4h Face, F5e, F5h, F5e Face, F5h Face
Medical device code: 85942014506005Q
Catalogue number of the medical device: 1006
Designated purpose: The JORDAN F tables are intended for use in medical facilities as examination beds for basic examination of patients in surgeries, they are also used as rehabilitation beds that are intended for use by patients of rehabilitation facilities, they are used in physiotherapy under the constant supervision by medical staff, especially by a rehabilitation and myoskeletal specialist. It is suitable for massage and spa procedures.
The JORDAN F tables can also be used for ambulance surgery interventions without anaesthesia when the patient is treated in the state of consciousness.
The JORDAN tables are not intended for use in an explosive atmosphere, are not suitable for use in the presence of inflammable anaesthetics and air mixtures, or inflammable anaesthetics and oxygen mixtures, or monoxide nitrogen.

Risk class of the medical device: I

The JORDAN F medical device to which the existing declaration applies complies with Regulation of the European Parliament and of the Council (EU) 2017/745 of 5 April 2017 on medical devices, amending Directive 2001/83/EC, Regulation (EC) No. 178/2002 and Regulation (EC) No. 1223/2009 and repealing Council Directives 90/385/EEC and 93/42/EEC.

In Třeboň on 04.05.23

RESI Třeboň spol. s r.o.

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Executive