

BIODEX

DECLARATION OF CONFORMITY

PRODUCT IDENTIFICATION

Type of Equipment	Product Name(s)	Model/Number(s)
C-Arm Radiologic Table	Urology C-Arm Table-800	058-800, 058-805
	Brachytherapy C-Arm Table-810	058-810, 058-815
	3D Imaging C-Arm Table-820	058-820, 058-825
	Surgical C-Arm Table-840	058-840, 058-840-10 058-845, 058-845-10
	Surgical C-Arm Table-846	058-846, 058-846-10 058-847, 058-847-10
	Pain Management C-Arm Table –870	058-870, 058-870-10 058-875, 058-875-10

Serial Number [ENTER]

MANUFACTURER

Name of Company	Address	Representative
Biodex Medical Systems, Inc.	20 Ramsey Road Shirley, NY 11967-4704 USA	Paul Cadmus <i>General Manager, Senior VP Operations</i>

AUTHORIZED REPRESENTATIVE

Name of Company	Address	Telephone/Email
Emergo Europe B.V.	Prinsessegracht 20 2514 AP The Hague The Netherlands	Phone: +31.70.345.8570 Fax: +31.70.346.8570

CONFORMITY ASSESSMENT

Device Classification	Route to Compliance	Standards Application
Class I Rule 12	Annex VII of MDD 93/42/EEC Council Directive	IEC 60601-1:2005 (Third Edition) + CORR. 1:2006 + CORR. 2:2007 + A1:2012 ANSI/AAMI ES60601-1:2005 + A1:2012 + C1:2009 and A2:2010 CAN/CSA-C22.2 No. 60601-1:2014 IEC 60601-2-46 Edition 3.0 CAN/CSA-C22.2 No. 60601-2-46:12, IEC 60601-1-2:2014


QUALITY SYSTEM REGISTRAR

Intertek Services Certified Quality Management System:
ISO 13485:2016 Certificate: 0084059

Biodex Medical Systems, Inc. declares the above mentioned products meet the provision of the Council Directive 93/42/EEC for Medical Devices.

Company Representative: Paul Cadmus
Title: General Manager, Senior VP Operations
Date: March 25, 2021

No. 043 Rev. B

Signature: 

Biodex Medical Systems, Inc.

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