medical.dyaco.com
+886-4-7977-888

Declaration of Conformity

According to Medical Device Directive 93/42/EEC

Manufacturer:

Dyaco International Inc.
No.1, Gong 1st Road, Hemei Township,
Changhua County 50843, Taiwan

Conformity Assessment Procedure:

Annex II excluding (4) of the Medical Device Directive
MDD 93/42/EEC

Notified Body:

TUV SUD
Ridlerstrasse 65, DE-80339 Munich, GERMANY CE 0123

Product Identification

Category:	Rehabilitation Bikes and Recumbent Seated Stepper
Brand:	Dyaco
Mode No:	7.0R
Lot/Batches/Serial No:	D70R

MDD Directive:

Class Im, Rule 12, MDD
93/42/EEC

The medical device compliance with the essential equirements in accordance with Annex II of the Medical Device Directive 93/42/EEC.

We declare under our sole responsibility that the products, to which this declaration relates, are in conformity with the Essential Requirements Annex II of the above directive.
This DOC is Valid until May 26th 2024 - EC certificate validity date.
Taiwan, 2021/8/11
Carl D. Mebane, Director of Regulatory Affairs

Signature