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	Prepared By	John Adcock		Date	July 2012	

DECLARATION OF CONFORMITY
European Communities Council Directives 93/42/EEC and 2007/42/EC Concerning Medical Devices
Manual Wheelchairs

Date	Issue No.	Amendment Description
17 th February 2012	1.0	New document; Replaces earlier declarations that related to specific products.
28 th March 2012	2.0	New Luna model added.
12 th July 2012	2.1	Minor correction

The undersigned declares that the medical devices described below meet the provisions of the Council Directives which apply and the CE Mark may therefore be affixed. The Products are registered with the UK Competent Authority. All supporting documentation is retained at the premises of the manufacturer.

Product Description	Manual wheelchairs manufactured by this company, see Annex A
Manufacturer	PDG - Product Design Group, Unit 103-318 East Kent Ave. South, Vancouver, BC, Canada V5X 4N6.
Intended Use	Highly adjustable mobility aids for specified patient groups as described in the appropriate instructions for use for each design.
MDD Classification	Class I (one) - Annex IX, Rule 1
Notified Body	Not applicable
EU Authorised Representative	Advena Ltd. Registered office; Thorne Widgery House, 33 Bridge Street, Hereford HR4 9DQ, UK
Assessment route	Self certification by Annex VII; EC Declaration of Conformity and Article 14; Registration of persons responsible for placing devices on the market.
European registration	By the UK Medicines and Healthcare Products Regulatory Agency (MHRA); number assigned CA 004986
Applicable product standards.	As shown in document AS 101

Signed  Date JULY 16/2012
Name (printed) MIKE JALMARSON
Position PRODUCT MANAGER





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ANNEX A

Product Variations;

- Eclipse
- Bentley
- Stellar
- Fuze T50
- Fuze T20
- Fuze JR
- Luna Bariatric

This declaration covers all sizes, configurations, and the spares and accessories made available with these wheelchairs.