Electric Mobility Euro Ltd.
Canal Way Ilminster
Somerset TA19 9DL
Company Registration in England No. 2419231

E C DECLARATION OF CONFORMITY

The Products Covered by this Declaration

Electric Mobility (Euro) Model: T4KE/ Veo and Veo X scooters.

The EC Directives covered by this Declaration

- Council Directive 93/42/EEC of 14th June 1993 concerning Medical Devices .
- Restriction of Hazardous Substances (RoHS) Directive 2011/65/EU

The Basis on which Conformity is Declared

The Class of the Scooter: as defined in the Directive Annex IX, is Class I
The product identified above complies with the Essential Requirements of the Medical Devices Directive.

The following Normative European and International Standards have been referenced to meet the Essential Requirements of the Directive/s:

EN 12184: 2009	Electrically powered wheelchairs, Scooters and their chargers
EN ISO 14971:2007	Medical Devices Risk Management
EN 1041 : 2008	Information supplied by the manufacturer with medical devices
EN 12182:1999	Technical aids for disabled persons – General Requirements and test methods
EN60601-1-2:2007	EMC – Scooter Unit
EN 55011:2007+A2:2007	EMC – Scooter Unit
BS EN 1021-1/2:2006	Seat Fabric Flammability (In lieu of ISO: 7176-16 below)
EN 60335-2-29	Electrical Safety – Battery Chargers (or EN 60601-1)
EN 55022	EMC compliance- Battery Charger
ISO 7176-1 1999	Wheelchairs Static Stability
ISO 7176-2 2001	Wheelchairs Dynamic Stability
ISO 7176-3 2003	Wheelchairs Brakes
ISO 7176-4 1997	Wheelchairs Energy Consumption
ISO 7176-5 2008	Wheelchairs Overall Dimensions
ISO 7176-6 2001	Wheelchairs Speed and acceleration
ISO 7176-7 1998	Wheelchairs Dimensions of seating etc
ISO 7176-8 1998	Wheelchairs Static Impact and fatigue strength
ISO 7176-9 2009	Wheelchairs Climatic Testing
ISO 7176-10 2008	Wheelchairs Obstacle climbing
ISO 7176-14 2008	Wheelchairs Power and control systems
ISO 7176-15 1996	Wheelchairs Labelling and information
ISO 7176-16 1997	Wheelchairs Resistance to ignition
ISO 7176-21 2009	Wheelchairs EMC

The technical documentation required to demonstrate that the requirements of the Medical Devices Directive has been compiled by the signatory below and is available for inspection by the relevant enforcement authorities.

The CE Mark was first applied on: 12/2012

I hereby certify that the products described above comply with the essential requirements of the Medical Devices Directive 93/42/EEC of 14th June1993:

Signed: J.H.Heath **Date:** 12/2012

Name: S.H.HEARTH **Authority:** M.D.