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RoHS 2 Declaration of Conformity

The EU Directive 2011/65/EU (referred to as RoHS 2 Directive) restricts the use of certain hazardous substances in electronic and electric equipment. It came into effect on 21 July 2011. This directive substantially updated and amended the original RoHS 1 Directive 2002/95/EC. Medical Devices are under Category 8 and were initially excluded from meeting the requirements. This exemption has been removed from the revised directive meaning Medical Devices must be compliant with RoHS 2 from 22nd July 2014.

MGE declares that all of its electrical and electronic Medical Products are compliant with EU Directive 2011/65/EU on the Restriction of Hazardous Substances (RoHS 2).

Basis of Compliance

Substance amounts, if present, are below the required limits in the following table.

Substance	Allowable Limit
Lead (Pb)	0.1% (1000 ppm)
Mercury (Hg)	0.1% (1000 ppm)
Cadmium (Cd)	0.01% (100 ppm)
Hexavalent Chromium (Cr 6+)	0.1% (100 ppm)
Polybrominated Biphenyls (PBB)	0.1% (100 ppm)
Polybrominated Diphenyl Ether (PBDE)	0.1% (100 ppm)

This Declaration of Conformity is based on the RoHS Declarations of Conformity received from MGE component suppliers. MGE makes every effort to ensure the correctness of supplier Declarations of Conformity and accepts no responsibility or liability as a result of inaccuracies of component supplier's declarations.

MGE remain dedicated to ensuring the continued and successful implementation of this program

Graham Martin
Managing Director & Quality Manager