

CE

LVD

EMC

ROHS

1111 W. 35th Street, Chicago IL 60609, USA

DECLARATION OF CONFORMITY

Product(s) listed below meet the requirements of RoHS2 Directive 2011/65/EU as required by Article 7 by Decision 768/2008 and amended by 2015/863/EU (RoHS3). The RoHS2 directive (2011/65/EU) is an evolution of the original directive, which now includes the CE-Marking Directive and four new phthalate substances have been added to the restricted substances listed under Annex II of the EU RoHS2 Directive on March 31, 2015.

This declaration of conformity is issued under the sole responsibility of the manufacturer.

Type of equipment:	MEDICAL UPS
Model number(s):	SMX700HG SMX1200XLHG
Series number(s):	AGSM6833 AGSM6834
Applicable Council Directive(s):	LVD 2014/35/EU MED 93/42/EEC EMC 2014/30/EU RoHS3 2011/65/EU as amended by 2015/863/EU
RoHS Exemption(s):	6(a); 6(b); 6(c); 7(c)-II; 8(b)
Standards to which conformity is being declared:	EN 62040-1:2008+A1:2013 EN 60601-1+A1:2012 EN 60601-1-2:2015 EN 62040-2:2006 EN IEC 63000:2018
Manufacturer Representative in the EU:	Eaton I.F. 110 Rue Blaise Pascal 38330 Montbonnot St Martin France

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