EUROLINK (EUROPE) LTD	***DECLARATION OF CONFORMITY***
Manufacturer:	The Lebanon Corporation 1700 N. Lebanon Street Lebanon Indiana 46052 USA
Conforming Apparatus:	Intraocular Pressure Reducer
Classification:	Class I - according to Annex IX Rule 5
Medicines and Healthcare products Regulatory Agency Apparatus Description:	Pressure Relief Device and Accessories
Medicines and Healthcare products Regulatory Agency Registration Reference:	CA 008803
Technical Document Reference No:	TF-0001
Harmonised EMC Standard(s) and Documents Referenced:	BS EN ISO 14971:2007 Medical Devices ï Application of Risk Management to Medical Devices Medical Devices Directive ï Annex I, Essential Requirements
EU Authorised Representative:	D.R.M. Green Eurolink (Europe) Ltd. Greyfriars Court Paradise Square Oxford Oxon OX1 1BE UK Tel: (44) 1793 784545 Fax: (44) 1793 784551

We certify that the apparatus identified above conforms to the requirements of Council Directive 93/42/EEC, as amended by Council Directive 2007/47/EC, on the approximation of the laws of the member state relating to medical devices.

Signed: 

D. R. M. Green

Date: 21 January 2015