

# **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 .  
Application of Risk Management to Medical Devices  
Medical Devices Directive . Annex I, Essential  
Requirements

**EU Authorised Representative:**

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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:**

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D. R. M. Green

**Date:** 3 February 2012

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