

# EC DECLARATION OF CONFORMITY

## TEARLAB™ OSMOLARITY SYSTEM

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We, TearLab Corporation, with offices at 7360 Carroll Road, Suite 200, San Diego, California 92121, USA, declare under our own responsibility that the TearLab™ Osmolarity System:

- P/N 100000, TearLab™ Osmolarity System, Western Europe
- P/N 100016, TearLab™ Osmolarity System, Eastern Europe

is in conformity with all the applicable provisions of the following Directives:

- IN VITRO DIAGNOSTIC MEDICAL DEVICES DIRECTIVE 98/79 EC

DEVICE CLASSIFICATION: General In Vitro Diagnostic (IVD) device

QUALITY SYSTEM: TearLab Corporation's quality system is certified to ISO 13485: 2003

TECHNICAL FILE: Part (A) of the Technical File for this product shall be available from TearLab Corporation's Authorized Representative for the EU; Part (B) of the Technical File shall reside at TearLab Corporation's corporate headquarters.

NAME AND ADDRESS OF AUTHORIZED REPRESENTATIVE FOR THE EU:

- Cavendish Scott Ltd, PO Box 107, SG5 1FW, England

SIGNED ON BEHALF OF TEARLAB CORPORATION:



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SIGNATURE

Michael S. Berg  
NAME

VP, Clinical & Regulatory Operations  
TITLE

June 15, 2010  
DATE