

EC DECLARATION OF CONFORMITY

TEARLAB™ OSMOLARITY CONTROL SOLUTIONS

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 Control Solutions:

- P/N 100012, TearLab™ Osmolarity Normal Control Solutions
- P/N 100013, TearLab™ Osmolarity High Control Solutions

are 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:



SIGNATURE

Michael S. Berg
NAME

VP, Clinical & Regulatory Operations
TITLE

June 15, 2010
DATE