



Declaration of Conformity

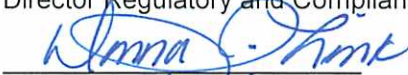
PRODUCT IDENTIFICATION PARASITOLOGY FAMILY OF PRODUCTS	
Product name	Model/number
<i>GIARDIA II</i>	PT5012, 30405
<i>CRYPTOSPORIDIUM II</i>	PT5014, 30406
<i>E. HISTOLYTICA II</i>	T5017, 30404
<i>GIARDIA/CRYPTOSPORIDIUM CHEK®</i>	T5031, 30401, T30401SKG
<i>GIARDIA/CRYPTOSPORIDIUM QUIK CHEK</i>	T30407
<i>TRI-COMBO PARASITE SCREEN</i>	T30408
<i>E. HISTOLYTICA QUIK CHEK</i>	T30409
Kits listed include all of the corresponding components.	

MANUFACTURER		
Name of company	Address	Representative
TECHLAB®	2001 Kraft Dr Blacksburg, VA 24060 U.S.A.	Donna T. Link – Director Regulatory and Compliance and Quality Management Representative

AUTHORIZED REPRESENTATIVE		
Name of company	Address	Telephone/email
Emergo Europe	Molenstraat 15 2513 BH The Hague, Netherlands	+31.70.345.8570 - phone +31.70.346.7299 - fax service@emergogroup.com

CONFORMITY ASSESSMENT		
Device classification	Route to compliance	Standards applied
<i>Self-Certify</i>	Annex III of IVDD 98/79/EEC Council Directive	ISO 13485:2003

TECHLAB® declares that the above mentioned products meet the provision of the Council Directive 98/79/EEC for *in vitro* diagnostic Medical Devices.

COMPANY REPRESENTATIVE: Donna T. Link
 TITLE: Director Regulatory and Compliance, QMR
 SIGNATURE: 
 DATE: 2014/09/04
 YYYY/MM/DD