Declaration of Conformity

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

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**
Self-Certify	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: Donna T. Link

DATE: 2014/09/04
YYYY/MM/DD