



Declaration of Conformity

| PRODUCT IDENTIFICATION | | |
|--|---|--|
| Product name | Model/number | |
| Cardiac Marker Test Devices | | |
| QuickProfile Troponin I Serum Test Card | 75001 | |
| QuickProfile Troponin I Whole Blood Test Card | 75002 | |
| QuickProfile Cardiac Panel Serum Test Card | 75003 | |
| QuickProfile Cardiac Panel Whole Blood Test Card | 75004 | |
| QuickProfile Myoglobin Serum Test card | 75005 | |
| QuickProfile Myoglobin Whole Blood Test Card | 75006 | |
| QuickProfile CK-MB Serum Test Card | 75007 | |
| QuickProfile CK-MB Whole Blood Test Card | 75008 | |
| QuickProfile Troponin I Strip | 75009 | |
| QuickProfile CK-MB Strip | 75010 | |
| QuickProfile Myoglobin Strip | 75011 | |
| MANUFACTURER | | |
| Name of company | Address | Representative |
| LumiQuick Diagnostics, Inc. | 2946 Scott Blvd. Santa Clara, CA 95054 USA | Jeff Wang |
| AUTHORIZED REPRESENTATIVE | | |
| Name of company | Address | Telephone/email |
| Emergo Europe | Prinsessegracht 20 2514 AP The Hague, Netherlands | +31.70.345.8570 - phone +31.70.346.7299 - fax europe@emergogroup.com |
| CONFORMITY ASSESSMENT | | |
| Device classification | Route to compliance | Standards applied |
| Class: Self-Certify | Annex III of IVDD 98/79/EC Council Directive | ISO 13485:2003 |

LumiQuick Diagnostics, Inc. declares that the above mentioned products meet the provision of the Council Directive 98/79/EC for In Vitro Diagnostic Medical Devices and Directive 98/79/EC as transposed in the national laws of the Member States.

COMPANY REPRESENTATIVE: Jeff Wang

TITLE: Quality Systems Manager

SIGNATURE:

DATE: 28/04/2017