CE

EC Declaration of Conformity

In accordance with the Directive 98/79/EC

Manufacturer: RheoMeditech Inc.
2nd Fl. 32 Anam-ro, Dongdaemun-gu,
Seoul, Republic of Korea 02578

Product Name **: Platelet Function Analyzer**
Brand Name **: ANYSIS Instrument**
Model Name **: ANYSIS-300, ANYSIS-300S**
Classification **: General *In Vitro* diagnostic device**

Conformity Assessment Route:

Safety : 61010-1 : 2010
EN 55011 : 2009 + A1 : 2010
EN 61326-1 : 2013
EN 61326-2-6 : 2013
EN61000-3-2 : 2014
EN61000-3-3 : 2014

Authorized Representative: **JaviTech e.K**
Sachsenhausener Str. 16, 65824
Schwalbach am Taunus, Germany

The undersigned herewith declare that the stated medical devices meet the transposition into national law, the provisions of council directive 98/79/EC concerning medical devices; All supporting documentation is retained at the premises of the manufacturer. We, the manufacturer, are exclusively responsible for the document.

This declaration is valid for all devices described in this document.

Signed: Eunah Rhee Date: 2021.05.31

Eunah Rhee
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