

DECLARATION OF CONFORMITY QUALITY CERTIFICATE and CERTIFICATE OF ORIGIN

18 06 2024

Labtech Kft. (4031 Debrecen, Vág út 4, Hungary), as the manufacturer of the noninvasive cardiologic measuring equipment classified as **Class IIa** (according to Annex IX Rule 10 of the 93/42/EEC directive) including:

**EC type Holter systems with 'CARDIOSPY' software including:
EC-2H, EC-3H, EC-12H, EC-ABP, EC-3H/ABP models**

**EC type Holter Systems recorders including:
EC-2H, EC-3H, EC-12H, EC-ABP, EC-3H/ABP models**

and

**EC type resting and stress Test ECG systems with 'CARDIOSPY' software including:
EC-12R, EC-12RM, EC-12S, EC-12R/S, EC-3RT, EC-12RT, EC-12LT models**

**EC type resting and stress Test ECG recorders including:
EC-12R, EC-12RM, EC-12S, EC-12R/S, EC-3RT, EC-12RT, EC-12LT models**

hereby states that the country of origin of the medical devices is Hungary. The devices and all the accessories meet the requirements of the **Article 120** of the Regulation (EU) 2017/745 of the European Parliament and of the Council (**MDR**) regarding the Transitional provisions for the legacy devices certified based on 93/42/EEC directive and those of below listed international standards:

EN60601-1:2006 + A1:2013 + A12:2014

EN 60601-1-2:2015

EN 60601-1-6:2010

EN 80601-2-30:2010/A1:2015

EN 60601-2-25:1995

EN 60601-2-51:2003

EN 60601-2-47:2001 and AAMI / ANSI / ISO 60601-2-47:2012

EN ISO 15223-1:2016

EN 62304:2006

EN 62366:2008

Serial number:

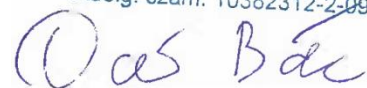
The quality management system of Labtech Kft. meets the **ISO 13485:2016** (Certificate No.: 4-601-135-2406) standard requirements, and the noninvasive cardiologic measuring equipments are entitled to bear the **"CE" mark** (Certificate No.: 5-850-200-1806)

The attesting certificates were issued by the NEOEMKI National Medical Device Conformity Assessment and Certification LLC (1011), (H-1097 Budapest, Albert Flórián út 3/a., Telephone: +36 20 268 7595)

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