

# EC Declaration of Conformity

**IVD**

In accordance with Directive 98/79/EC of the European Parliament and of the Council on in vitro diagnostic medical devices.

## Manufacturer Information

Manufacturer:

**LOMINA Superbio a.s.**

Na Radosti 184/59, Prague 5, 158 00, Czech Republic

ID: CZ07420099, www.lomina.ch, SÚKL reg.: 061583



## Medical Device (Product) Identification Data.

Name: Lomina Pregnancy Enhanced Sensitivity Test

**REF** La-HCG-Pro

Intended use:

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The The Lomina Pregnancy Enhanced Sensitivity Test is a rapid chromatographic immunoassay for the qualitative detection of human chorionic gonadotropin in urine to aid in the early detection of pregnancy.

## Version for Health Care Professionals

**Category of in vitro diagnostic medical device: IVD OTHER**

The manufacturer declares that the properties of the above in vitro diagnostic medical device fulfil all the requirements laid down in Directive 98/79/EC, and that the in vitro diagnostic medical device will perform in accordance with its intended purpose. The manufacturer further declares that he has taken measures to ensure compliance of the medical device placed on the market with the essential requirements and the manufacturer's technical documentation pursuant to Annex III of Directive 98/79/EC.

## **Harmonized standards:**

EN ISO 13485:2016, EN ISO 15223-1:2016, EN ISO 14971:2019, EN 13641: 2002,  
EN ISO 18113-1:2011, EN ISO 18113-2:2011, EN 13612:2002, EN ISO 23640:2015

Name: Michal Horáček, PMP, MBA  
Position: CEO  
Valid from: 2022/05/05

Signature:

