

Certificate of Conformity



Sartorius Biohit Liquid Handling Oy hereby confirm that the product line(s):

Picus & NxT Electronic pipettes+ Charging Stands
eLINE Electronic pipettes + Charging Stands
Biohit Midi Plus
Tacta Mechanical pipettes
mLINE Mechanical pipettes
Proline Plus Mechanical pipettes
Proline Mechanical pipettes

Are manufactured in: Sartorius Biohit Liquid Handling Oy Tietokatu 4, 87400 Kajaani, FINLAND

Country of origin is Finland.

All product lines are designed and manufactured according to the Essential Requirements listed in Annex I of EC Directive 98/79/EC (In Vitro Medical Devices).

This declaration is supported by following EC quality system approval certificates:

ISO 9001:2015 Certificate No. FI17/5101, issued by SGS, Finland, valid 14th December 2020

ISO 13485:2016 Certificate No. FI17/5103, issued by SGS, Finland, valid 14th December 2020

ISO 17025:2005 Certificate No. K041/A10/2016 issued by FINAS, valid 20th March 2020

ISO 14001:2015 Certificate No. FI17/5102, issued by SGS, Finland, valid 14th December 2020

Furthermore the Calibration Laboratory for Volumetric Instruments at Sartorius Biohit has been officially accredited by the Finnish Accreditation Service (FINAS) according to the **ISO/IEC 17025** (Cert. No. K041/2012).

All Sartorius Biohit products are individually tested according to the ISO 8655 and all delivered units have serial numbered QC-certificate included. The QC-certificate indicates that the unit has passed the given specifications.

Signed:

Helsinki 6th February 2018

Full Name: Seppo Riikonen

Title: Director, Quality & Process Development

Company: Sartorius Biohit Liquid Handling Oy, Laippatie 1, 00880 Helsinki, Finland
