



Certificate

No. Q5 115293 0001 Rev. 01

Holder of Certificate: **Biovica International AB**
Uppsala Science Park
Dag Hammarskjölds väg 54B
752 37 Uppsala
SWEDEN

Certification Mark:



Scope of Certificate: **Design and development, manufacturing and distribution of in vitro diagnostic reagents for oncology based on enzyme activity methods.**

The Certification Body of TÜV SÜD Product Service GmbH certifies that the company mentioned above has established and is maintaining a quality management system, which meets the requirements of the listed standard(s). All applicable requirements of the testing and certification regulation of TÜV SÜD Group have to be complied with. For details and certificate validity see: [www.tuvsud.com/ps-cert?q=cert:Q5 115293 0001 Rev. 01](http://www.tuvsud.com/ps-cert?q=cert:Q5_115293_0001_Rev_01)

Report No.: 75957501

Valid from: 2023-09-05

Valid until: 2026-09-04

Date, 2023-09-05



Christoph Dicks
Head of Certification/Notified Body



Product Service

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Applied Standard(s): EN ISO 13485:2016
Medical devices - Quality management systems -
Requirements for regulatory purposes
(ISO 13485:2016)
DIN EN ISO 13485:2016

Facility(ies): **Biovica International AB**
Uppsala Science Park, Dag Hammarskjölds väg 54B, 752 37
Uppsala, SWEDEN

See Scope of Certificate

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