

Certificate of Registration

QUALITY MANAGEMENT SYSTEM - ISO 13485:2016

This is to certify that:

FREYR SOFTWARE SERVICES
PRIVATE LIMITED
H08, Phoenix Avance Gachibowli
Hyderabad
Telangana
500 081
India

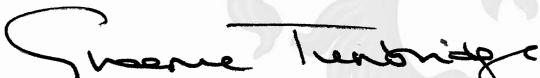
Holds Certificate Number:

MD 811683

and operates a Quality Management System which complies with the requirements of ISO 13485:2016 for the following scope:

Provision of Global Consulting Services for Product Registrations, Regulatory Affairs, Clinical & Medical Writing, Quality Management System, Labelling & Artwork Services, Post Market Surveillance Services, Importer Services and Authorized Representation Services, such as EAR, Swiss Rep, UKRP to the Medical Device and IVD Industry.

For and on behalf of BSI:



Graeme Tunbridge, Senior Vice President Global Regulatory & Quality

Original Registration Date: 2025-01-30

Effective Date: 2025-01-30

Latest Revision Date: 2025-12-03

Expiry Date: 2028-01-29



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Printed copies can be validated at www.bsigroup.com/ClientDirectory

Information and Contact:

BSI Group The Netherlands B.V., John M. Keynesplein 9, 1066 EP Amsterdam, The Netherlands | Tel: +31 20 3460 780
BSI Group The Netherlands B.V. is registered in The Netherlands under number 33264284 | A Member of the BSI Group Holdings B.V.

Certificate No: MD 811683

Location	Registered Activities
Freyr Software Services Pvt Ltd H08, Phoenix Avance Gachibowli Hyderabad Telangana 500 081 India	Provision of Global Consulting Services for Product Registrations, Regulatory Affairs, Clinical & Medical Writing, Quality Management System, Labelling & Artwork Services, Post Market Surveillance Services, Importer Services and Authorized Representation Services, such as EAR, Swiss Rep, UKRP to the Medical Device and IVD Industry.
Freyr Regulatory GmbH Group Company of Freyr Software Services Pvt Ltd Marie-Curie- Straße 8 Lörrach 79539 Germany	Provision of global consulting services for product registrations, regulatory affairs, clinical & medical writing, Quality Management system, labelling & artwork, Post Market Surveillance activities, Importer services and Authorized representation services to the medical devices and IVD industry.



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