



Certificate of Registration

QUALITY MANAGEMENT SYSTEM - ISO 13485:2016

This is to certify that: CROSSJECT

6 Rue Pauline Kergomard

Dijon 21000 France

Holds Certificate Number: MD 735691

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

The design, development and manufacture of non-sterile medical device component within a drug device combination product (single use needle-free autoinjector) intended for medicinal product administration.

For and on behalf of BSI:

Graeme Tunbridge, Senior Vice President Global Regulatory & Quality

Original Registration Date: 2022-08-12 Effective Date: 2025-08-12 Latest Revision Date: 2025-07-15 Expiry Date: 2028-08-11

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...making excellence a habit."

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Certificate No: MD 735691

France

Location	Registered Activities
CROSSJECT 6 Rue Pauline Kergomard Dijon 21000 France	The design, development and manufacture of non-sterile medical device component within a drug device combination product (single use needle-free autoinjector) intended for medicinal product administration.
CROSSJECT Rue des Vernottes Arc-lès-Gray 70100 France	The manufacture of non-sterile medical device component within a drug device combination product (single use needle-free autoinjector) intended for medicinal product administration.
CROSSJECT Giranaux ZI les Giranaux 5015 Rue des Giranaux Arc-lès-Gray 70100	The manufacture of non-sterile medical device component within a drug device combination product (single use needle-free autoinjector) intended for medicinal product administration.

Original Registration Date: 2022-08-12 Effective Date: 2025-08-12 Latest Revision Date: 2025-07-15 Expiry Date: 2028-08-11

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