

Certificate of Registration

QUALITY MANAGEMENT SYSTEM - ISO 13485:2016

This is to certify that: Adaptive Biotechnologies Corporation
1165 Eastlake Ave E
Seattle
Washington
98109
USA

Facility ID Number: F008675

Holds Certificate No: **MDSAP 836620**

The company listed on this certificate has been audited to and found to conform with ISO 13485:2016 including the following country specific requirements:

Australia: Therapeutic Goods (Medical Devices) Regulations, 2002, Schedule 3 Part 1 (excluding Part 1.6) - Full Quality Assurance Procedure

Canada: Medical Devices Regulations - Part 1 - SOR 98/282

USA: 21 CFR 820, 21 CFR 803, 21 CFR 806, 21 CFR 807 - Subparts A to D

The design, development and manufacture of in-vitro diagnostic software and reagents used in the aid of detection of minimal residual disease status in the field of oncology.

For and on behalf of BSI:

Graeme Tunbridge, Senior Vice President Global Regulatory & Quality

Original Registration Date: 2026-06-16

Effective Date: 2026-06-16

Expiry Date: 2029-06-01



BSI Group America Inc. is an MDSAP recognised auditing organization

Certificate No: **MDSAP 836620**

Location	Registered Activities
Adaptive Biotechnologies Corporation 1165 Eastlake Ave E Seattle Washington 98109 USA Facility ID Number: F008675	Design and development, design validation.
Adaptive Biotechnologies Corporation 19910 North Creek Parkway Bldg. F Bothell Washington 98011 USA Facility ID Number: F008675	Main production facility and supply chain management center for the whole organization.



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Page: 2 of 2

This certificate remains the property of BSI and shall be returned immediately upon request.
An electronic certificate can be authenticated [online](#). Printed copies can be validated at www.bsigroup.com/ClientDirectory
To be read in conjunction with the scope above or the attached appendix.

America Headquarters: BSI Group America Inc. 1950 Opportunity Way, Suite 900, Reston, VA 20190.
A Member of the BSI Group of Companies.