



America

CERTIFICATE

No. QS2 17 03 83983 006

Certificate Holder: BioLegend, Inc.
9727 Pacific Heights Blvd
San Diego CA 92121
USA

Certification Mark:



Scope of Certificate: Design, Development, Manufacture and Distribution of Antibodies, Proteins, Cells and Cellular Extracts, and Chemical Reagents for Use in the Medical Device Industry

Standard(s): ISO 13485:2003

The Certification Body of TÜV SÜD America Inc. certifies that the company mentioned above has established and is maintaining a quality management system that meets the requirements of the listed standards.

Report No.: M6042

Effective Date: 2017-03-29

Expiry Date: 2019-02-28



Gary Minks
Vice President, Regulatory Affairs

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TÜV SÜD America Inc.
10 Centennial Drive
Peabody, MA 01960
USA

TÜV®





America

CERTIFICATE

No. QS2 17 03 83983 006

BioLegend, Inc.
 9727 Pacific Heights Blvd
 San Diego, CA 92121
 USA

Design, Development, and Manufacture of Antibodies, Cells and Cellular Extracts, and Chemical Reagents for Use in the Medical Device Industry

BioLegend, Inc.
 9717 Pacific Heights Blvd
 San Diego, CA 92121
 USA

Manufacture and Distribution of Antibodies, Proteins, and Chemical Reagents for Use in the Medical Device Industry

BioLegend, Inc.
 8959 Terman Court
 San Diego, CA 92121
 USA

Design, Development, and Manufacture of Antibodies, Proteins, and Chemical Reagents for Use in the Medical Device Industry

Effective Date: 2017-03-29
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 Vice President, Regulatory Affairs

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TÜV SÜD America Inc.
 10 Centennial Drive
 Peabody, MA 01960
 USA

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America

CERTIFICATE

No. QS2 16 12 96519 001

Certificate Holder: BioLegend, Inc
210 Rustcraft Road
Dedham MA 02026
USA

Certification Mark:



Scope of Certificate: Design, Development, Manufacturing and Distribution of Antibodies, Recombinant Proteins, Biomarker Immunoassays and Other Chemical Reagents for Use in Immunoassay and Functional Assays in the Medical Device Industry

Standard(s): ISO 13485:2003

The Certification Body of TÜV SÜD America Inc. certifies that the company mentioned above has established and is maintaining a quality management system that meets the requirements of the listed standards.

Report No.: M6971

Effective Date: 2016-12-23

Expiry Date: 2019-02-28



Gary Minks
Vice President, Regulatory Affairs

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