CERTIFICATE OF REGISTRATION

REAGEN, INC.

7098 Miratech Drive Suite 110
San Diego, California 92121 USA

American Global Standards, Inc. (AGS) issues this certificate to the firm named above, having assessed and approved the firm's Medical Devices - Quality Management Systems - Requirements for Regulatory Purposes and finding the system conforms to the standards of:

ISO 13485:2016

The Medical Devices - Quality Management Systems - Requirements for Regulatory Purposes is applicable to the following:

Design, develop, manufacturing and distribution of in-vitro diagnostics test kits for infectious pathogens and clinical chemistry.

This approval is subject to the firm maintaining its system to the required standards, which will be monitored by AGS. In the issuance of this certificate, AGS assumes no liability to any party other than the firm named above, and then only in accordance with the agreed upon Quality System Assessment Agreement.

Certification Number:	AGS-US091224-I
Original Approval:	September 12, 2024
Date of Issue:	September 12, 2024
Date of Expiration:	September 11, 2027

For and On Behalf of American Global Standards, Inc.
Stephen Keneally, President

AIAO-BAR QMS ACCREDITED

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