



SCOPE OF ACCREDITATION TO ISO/IEC 17025:2005

VIVITRO LABS, INC.
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MECHANICAL

Valid To: January 31, 2020

Certificate Number: 3224.01

In recognition of the successful completion of the A2LA evaluation process, accreditation is granted to this laboratory to perform the following tests on medical products:

<u>Test:</u>	<u>Test Method(s):</u>
Cardiovascular Implants-Cardiac Valve Prostheses Steady Forward-Flow Testing Steady Back-Flow Leakage Testing	BS EN ISO 5840:2005 (Annex L.2 and L.3); IS EN ISO 5840:2009 (Annex L.2 and L.3); IS EN ISO 5840-2:2015 (Annex F.2 and F.3); IS EN ISO 5840-3:2013 (Annex N.2 and N.3); QV-504
Pulsate-Flow Testing	BS EN ISO 5840:2005 (Annex L, <i>Excluding 4.3.5 and 4.3.6</i>); IS EN ISO 5840:2009 (Annex L, <i>Excluding 4.3.5 and 4.3.6</i>); IS EN ISO 5840-2:2015 (Annex F, <i>Excluding 4.3.5 and 4.3.6</i>); IS EN ISO 5840-3:2013 (Annex N, <i>Excluding 4.3.5 and 4.3.6</i>); QV-501
Durability Testing	BS EN ISO 5840:2005 (Annex M); IS EN ISO 5840:2009 (Annex M); IS EN ISO 5840-2:2015 (Annex G); IS EN ISO 5840-3:2013 (Annex O); QV-502



Accredited Laboratory

A2LA has accredited

VIVITRO LABS, INC.

Victoria, British Columbia, CANADA

for technical competence in the field of

Mechanical Testing

This laboratory is accredited in accordance with the recognized International Standard ISO/IEC 17025:2005 *General requirements for the competence of testing and calibration laboratories*. This accreditation demonstrates technical competence for a defined scope and the operation of a laboratory quality management system (refer to joint ISO-ILAC-IAF Communiqué dated 8 January 2009).



Presented this 15th day of January 2018.

A handwritten signature in black ink, written over a horizontal line.

President & CEO
For the Accreditation Council
Certificate Number 3224.01
Valid to January 31, 2020

For the tests to which this accreditation applies, please refer to the laboratory's Mechanical Scope of Accreditation.