

# NVLAP ON-SITE ASSESSMENT SUMMARY

Please complete this summary and attach it to the original On-Site Assessment Report.

**DO NOT LEAVE THIS SUMMARY WITH THE LABORATORY.**

Laboratory Name: \_\_\_\_\_

Lab Code: \_\_\_\_\_

Fields of Accreditation: \_\_\_\_\_

Assessor Name(s): \_\_\_\_\_

Date of **Pre-assessment** Review of Quality Manual: \_\_\_\_\_

Date(s) of On-Site Assessment: \_\_\_\_\_

This report contains changes to the laboratory's Scope of Accreditation:  additions;  deletions;  modifications. (Please describe in the On-Site Narrative Summary.)

## SUMMARY AND RECOMMENDATIONS:

- The laboratory has no nonconformities and no written response to NVLAP is required.
- The laboratory has nonconformities in the following area(s). I have notified the laboratory of these nonconformities and the requirement to respond to NVLAP in writing about their resolution.

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| <input type="checkbox"/> <b>4 General requirements</b>                         | <input type="checkbox"/> 7.8 Reporting of results  |
| <input type="checkbox"/> 4.1 Impartiality                                      | <input type="checkbox"/> 7.9 Complaints  |
| <input type="checkbox"/> 4.2 Confidentiality                                   | <input type="checkbox"/> 7.10 Nonconforming work   |
| <input type="checkbox"/> <b>5 Structural requirements</b>                      | <input type="checkbox"/> 7.11 Control of data and information management                           |
| <input type="checkbox"/> 5.1 to 5.7  | <input type="checkbox"/> <b>8 Management system requirements</b>                                   |
| <input type="checkbox"/> <b>6 Resource requirements</b>                        | <input type="checkbox"/> 8.1 Options   |
| <input type="checkbox"/> 6.1 General   | <input type="checkbox"/> 8.2 Management system documentation                                       |
| <input type="checkbox"/> 6.2 Personnel   | <input type="checkbox"/> 8.3 Control of management system documents                                |
| <input type="checkbox"/> 6.3 Facilities and environmental conditions           | <input type="checkbox"/> 8.4 Control of records  |
| <input type="checkbox"/> 6.4 Equipment   | <input type="checkbox"/> 8.5 Actions to address risks and opportunities                            |
| <input type="checkbox"/> 6.5 Metrological traceability                         | <input type="checkbox"/> 8.6 Improvement   |
| <input type="checkbox"/> 6.6 Externally provided products and services         | <input type="checkbox"/> 8.7 Corrective actions  |
| <input type="checkbox"/> <b>7 Process requirements</b>                         | <input type="checkbox"/> 8.8 Internal audits   |
| <input type="checkbox"/> 7.1 Review of requests, tenders and contracts         | <input type="checkbox"/> 8.9 Management reviews  |
| <input type="checkbox"/> 7.2 Selection, verification and validation of methods | <input type="checkbox"/> Annex A. Referencing NVLAP accreditation                                  |
| <input type="checkbox"/> 7.3 Preventive action                                 | <input type="checkbox"/> Annex B. Implementation of traceability policy in accredited laboratories |
| <input type="checkbox"/> 7.4 Handling of test or calibration items             | <input type="checkbox"/> Annex E. Use of the Accredited Laboratory Combined ILAC MRA Mark          |
| <input type="checkbox"/> 7.5 Technical records                                 | <input type="checkbox"/> Other (Specify) _____   |
| <input type="checkbox"/> 7.6 Evaluation of measurement uncertainty             |  |
| <input type="checkbox"/> 7.7 Ensuring the validity of results                  |  |

- Based on my findings regarding nonconformities, staff competence, and laboratory procedures, I recommend that another on-site assessment be performed before this laboratory is granted accreditation.

Signature of Lead Assessor: _____	Date: _____
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