## Gompliancehelp iso consulting

**Audit Report for** 

### mBar Technologies Inc.

**Prepared for:** 

Todd Pack mBar Technologies Inc. 208-794-8666 tpack@mbarinc.com Audit Start Date:

2025-01-16

**Audit End Date:** 

2025-01-17

**Location of Audit:** 

Main Office, Boise ID

**Auditor Name:** 

Michael Stamm

No. of Audit Days: 1.5

# Thank you for letting us audit your awesome organization!!

#### **AUDIT SUMMARY**

mBar Technologies Inc. (mBar) underwent an internal audit of their Quality Management System (QMS) on January 16<sup>th</sup> and 17<sup>th</sup>, 2025. The scope consisted of the entire QMS as described in the company's Quality Manual and referenced procedures. The audit was based around the requirements of this QMS and the ISO IEC 17025:2017 standard for laboratories.

During this audit, all processes were reviewed, and process participants were interviewed. There were Number (0) non-conformances and Number (0) observations identified for an accumulative audit score of 100%.

Despite being a small organization with limited personnel resources, the on-site leader has shown a solid commitment to implementing the QMS and will be undergoing regular internal and third-party audits. The team was available and knowledgeable about their areas of responsibility and the importance of compliance with the standard and system and did a great job of retaining and providing records.

The company owner has shown outstanding commitment to the QMS and the continual improvement of processes and customer service using process improvement tools, metrics tracking, industry, and market knowledge, and maintaining sufficient evidence and accountability for quality performance.

We congratulate them on their excellent work to continue to maintain and improve the QMS as an integral part of the business. Great job, mBar!!!

Sincerely, Michael Stamm, Lead Auditor, ComplianceHelp Consulting

#### **SCOPE OF AUDIT**

ComplianceHelp will conduct an Internal Audit of the mBar Technologies Inc. QMS relative to ISO/IEC 17025:2017 standard.

#### STANDARD(S)

ISO/IEC 170254:2017

#### **Compliance Scores**

Would a great game ever become popular if there was no scoring? We love scoring. It can help an organization or department become laser focused on improvement.

Our scoring method is objective because it is based on evidence presented during the audit.

For greater insights, the auditor will also provide the Audit worksheets from this audit. The Audit worksheet will let you see, line by line, which areas of your organization are complying or lacking.

GROUP NAME	COMPLIANCE SCORE (%)
SECTION 1: Quality Manual	100.00
2.2 Quotations and Communications	100.00
2.2.1 Service to the Customer	100.00
2.3 Receipt of Calibration Items	100.00
2.4 Scheduling	100.00
2.5 Purchasing	100.00
2.6 Calibration Services	100.00
2.7 Post-Calibration Item/Product Management	100.00
3.1.1 Lab Conditions	100.00
3.2 Calibration Method Selection and Calibration Setup	100.00
3.2 calibration Method Selection and calibration Setup	100.00
3.3 Measurement of Calibration Uncertainty	100.00
3.4.1 Equipment Selection and Commissioning	100.00
3.4.2 Equipment Maintenance	100.00
3.4.3 Safe Handling and Care of Calibration Equipment	100.00

3.4.4 Calibration of Measuring Equipment	100.00
3.5 Ensuring the validity of results	100.00
4.1.1 Issues and Non-Conformances	100.00
4.1.2 Corrective Action	100.00
4.2.1 Scope and Master Document List	100.00
4.2.2 Development and Approval of New Documents	100.00
4.2.3 Document Review and Changes	100.00
4.2.4 Backing up Electronic Documents and Records	100.00
4.3 Records Control	100.00
4.3.2 Electronics Records	100.00
4.4 Personnel	100.00
4.4.2 Authorizations and Training	100.00
4.4.3 Employee Selection Process and Training/Supervision	100.00
4.4.4 Training Needs	100.00
4.4.5 Annual Reviews	100.00
4.5 Internal Audits	100.00
4.6 Management Review	100.00
4.6.1 Post Management Review	100.00
4.7 Risk Assessment	100.00
Average Compliance Score	100.0

#### **Audit Report**

If you scored 100% at the end of the Compliance Scores table above, then there's probably a good chance that this section is blank. Not to worry, it's not a glitch in our reporting, it's just that we could not find anything wrong, and that's not a bad thing.

And if there is anything reported in this section, hey, nobody's perfect. Step back, gather your thoughts, take a deep breath, and GET YOUR GAME ON.

By the way, the action code "NC" means non-conformance. A non-conformance either means you are not complying with your policies/procedures or the relevant standard. If you receive an NC, your organization will need to correct the situation and start conforming. The action code "OB" means Observation, or in other words a strong suggestion. You do not have to implement an observation, so we will leave that up to you to decide. Normally an observation is provided to help you improve.

Here is your Audit Report...

**Congratulations! There are no Findings** 

#### **Group Level Report**

Here are your Group-level actions and comments.

#### What next?

Audits are typically annual, so we look forward to seeing you next year. Here's the process for organizing next year's audits:

