Environment Canada

Internal Quality Management Audit Report
Report No [insert number]

ISO 9001:2008

[Insert title and reference number of audited process]

[Insert date of audit]

Lead Auditor: _____

Process Owner: _____

Internal Quality Management Audit Report

Overarching Process:

Underlying Process name:

Process Owner:

Auditee management representative(s):

Lead Auditor:

Additional Audit Team members:

Location/address of audit:

Scope of audit:

Start Date: **End Date:**

Audit Findings

No issues or non-conformances were identified during this assessment. ☐

One or more non-conformances were identified. ☐

One or more opportunities for improvement were identified. ☐

Opening/Closing Meeting

Topics for Opening Meeting

Review scope and objectives of audit ☐
Review audit plan and methodology ☐
Health and Safety Concerns ☐
Time and Date for Closing Meeting ☐

Topics for Closing Meeting

Briefing on audit results ☐
Review non conformances ☐
Identify follow-up requirements ☐

(Insert name and number of process)
PROTECTED A once filled
http://ecollab.ec.gc.ca/theme/wesgms/quality%20manual/forms/allitems.aspx?pflg=1033&

(Insert date of audit)
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Audit Report
Opening Remarks
Report on setting of quality objectives and monitoring success:
Report on progress towards implementing continual improvement:
Report on customer feedback and responses (unsolicited or solicited):
Report on input to management review process and response to outcomes from that process:
Report on non-conformances identified in previous audit(s) and their current status:
(other reports at discretion of auditor)

(Insert name and number of process)
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Non-conformances

CPAR # see Note 1	Summary description	Process/ISO Clause	Internal QMS reference
			• Requirements (process or ISO): • Objective evidence: • Process or ISO reference:
			• Requirements (process or ISO): • Objective evidence: • Process or ISO reference:
			• Requirements (process or ISO): • Objective evidence: • Process or ISO reference:

Note 1: CPAR number to be assigned by QMS Office after review of preliminary audit report.

Additional detail or clarifications on above non-conformances:

(may be filled out by auditors a the time of audit or by QMS staff upon review of preliminary report)

(Insert name and number of process)
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Opportunities for Improvement

Summary description	Process/ISO Clause	Internal QMS reference

Additional detail on above opportunities:

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