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QUALITY MANAGEMENT SYSTEM
(TYPE 6)

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IC-DC-Q6
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FOREWORD

RMS COPYRIGHT AND USE OF THIS DOCUMENT

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When this document forms part of a deed

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BASE SPECIFICATION

This document is based on Specification RMS Q6 Edition 1 Revision 10.
RMS SPECIFICATION D&C Q6
QUALITY MANAGEMENT SYSTEM (TYPE 6)

1 GENERAL

1.1 SCOPE

The work to be executed under Specification RMS D&C Q6 (RMS Q) consists of:

(a) Developing and implementing a corporate Quality Management System in accordance with ISO 9001;

(b) Developing and implementing a PROJECT QUALITY PLAN to cover all your Work, whether permanent or temporary both on-site and off-site;

(c) Operating, maintaining and reviewing the PROJECT QUALITY PLAN and associated quality management system procedures; and

(d) Keeping Quality Records in accordance with RMS Q Clause 4.2.4.

1.2 STRUCTURE OF THE SPECIFICATION

This Specification includes a series of annexures that detail additional requirements.

1.2.1 Details of Work

Details of work are shown in Annexure Q/A.

1.2.2 (Not Used)

1.2.3 Schedules of HOLD POINTS and Identified Records

The schedules in Annexure Q/C list the HOLD POINTS that must be observed. Refer to Clause 1.4 for the definition of HOLD POINTS.

The Quality Records for the deed and referenced documents listed in Annexure Q/C must be created, when applicable, and located as specified in Annexure Q/C.

The records listed in Annexure Q/C are Identified Records for the purposes of Annexure Q/E.

1.2.4 Planning documents

The PROJECT QUALITY PLAN must include each of the documents and requirements listed in Annexure Q/D and must be implemented.

In all cases where RMS Q refers to manufacturers’ recommendations, these must be included in the PROJECT QUALITY PLAN.
1.2.5 Record Keeping

Records for the deed, including quality, environmental, OHS and other management records must comply with Annexure Q/E.

1.2.6 Referenced Documents

Standards, specifications and test methods are referred to in abbreviated form (e.g. AS 2350). For convenience, the full titles are given in Annexure Q/M.

1.3 SYSTEM NOT FORMATTED ON AS/NZS ISO 9001:2008

Where the Quality Management System documents are arranged differently to the format of AS/NZS ISO 9001:2008, include in the PROJECT QUALITY PLAN a matrix of how the Quality Management System addresses all the requirements of RMS Q and AS/NZS ISO 9001:2008.

1.4 TERMS AND DEFINITIONS

The definitions appearing in ISO 9000 and in ISO 9001 Clause 3 apply in the interpretation of the words and expressions appearing in the quality assurance provisions of the deed (except where the context otherwise requires).

“RMS Q” and “RMS D&C Q6” appearing in the deed documents means this Specification.

Additionally, the following words and expressions appearing in the deed have the meanings hereby assigned to them, except where the context otherwise requires:

"Hold Point": a point beyond which a work process must not proceed without the RMS Representative's express written authorisation;

"inspection records": the evidence of conformity specified in ISO 9001 Clauses 7.1 (d) and 8.2.4.

“inspection and test forms/ITP forms”: the forms that accompany the Inspection and Test Plan (ITP) and that are used for recording inspection/test results (e.g. verification checklists). If the ITP contains the facility to record inspection/test results, the ITP will also be regarded as an “ITP form” (refer RMS Q Clause 8.1.1).

“Project Testing”: testing, including sampling, carried out on the site, at concrete and asphalt batch plants, on aggregates and materials used for pavements and structures at off site locations and any other testing specified in Annexure Q/A to be Project Testing.

"Witness Point": a point in a work process where the Contractor must give prior notice to the RMS Representative and the option of attendance may be exercised by the RMS Representative.

2 (NOT USED)

3 (NOT USED)
4 QUALITY MANAGEMENT SYSTEM

4.1 GENERAL REQUIREMENTS

Develop a corporate Quality Management System that complies with all the requirements ISO 9001 and RMS Q. Implement and maintain the Quality Management System in accordance with ISO 9001 and RMS Q.

Apply the following quality assurance practices to the Contractor’s Work:

(a) ensure that purchased items conform to specification before incorporating them in the Project Works;
(b) plan and control work processes;
(c) plan and carry out inspection and testing (including identification and traceability) to verify that the work processes are effective and that all finished work complies with the deed;
(d) careful selection of subcontractors and confirmation that their work complies with the deed;
(e) where the Specifications require plans, procedures, methods and forms to be documented, use these documents in implementing the Quality Management System for the deed;
(f) acknowledge and rectify any nonconforming work and improve work processes to prevent recurrence of nonconformities;
(g) keep orderly records to demonstrate that the Contractor’s Work complies with the deed; and
(h) improve procedures and work practices when opportunities are identified to minimise errors, waste and product nonconformities.

4.2 DOCUMENTATION REQUIREMENTS

4.2.1 General

Those requirements in RMS Q that are additional to the requirements of ISO 9001 may be addressed within the corporate Quality Management System or in supplementary quality management system documentation applied to the deed. Where documentation of procedures is called for, it is acceptable to document the procedures either individually or combined with other procedures depending on how you choose to structure your quality management system.

4.2.2 Quality Management System Documents

4.2.2.1 QUALITY MANUAL

Establish and maintain a QUALITY MANUAL in accordance with ISO 9001 using HB90.3 Clause 4.2.2 for guidance.

4.2.2.2 Quality Management System Procedures

Document, maintain and implement procedures in accordance with ISO 9001 as part of the corporate Quality Management System to:

(a) control corporate and project documents (refer RMS Q Clause 4.2.3);
(b) manage quality records (refer RMS Q Clause 4.2.4);
(c) ensure personnel are competent and appropriately trained/qualified (refer RMS Q Clause 6.2.2);

(d) plan product realisation and preparation of the PROJECT QUALITY PLAN (refer RMS Q Clause 7.1);

(e) review customer requirements (refer RMS Q Clause 7.2);

(f) plan, resource and manage design and development (refer RMS Q Clause 7.3);

(g) control purchasing and subcontracted work to ensure conformity to specification requirements (refer RMS Q Clause 7.4);

(h) plan and implement process controls and monitor their effectiveness (refer RMS Q Clause 7.5.1);

(i) identify and trace products and work (refer RMS Q Clause 7.5.3);

(j) control inspection and testing activities (refer RMS Q Clauses 8.1.1, 8.2.4);

(k) plan and implement internal auditing (refer RMS Q Clause 8.2.2);

(l) identify, record, notify and control nonconforming products or services (refer RMS Q Clause 8.3);

(m) analyse nonconformities and implement corrective action (refer RMS Q Clause 8.5.2);

and

(n) implement preventive action (refer RMS Q Clause 8.5.3).

Procedures required by this Specification that are additional to the requirements of ISO 9001 may be included in your general corporate quality management system procedures or as supplementary corporate quality management system procedures to be applied on RMS deeds. The latter may be incorporated as part of a proforma PROJECT QUALITY PLAN for RMS deeds, controlled within your corporate Quality Management System.

The PROJECT QUALITY PLAN or QUALITY MANUAL must describe or reference the applicable quality management system procedures required by this Specification and show their revision status. Quality management system procedures referenced in the PROJECT QUALITY PLAN must be readily accessible to project personnel at their work locations.

4.2.2.3 PROJECT QUALITY PLAN

Prepare the PROJECT QUALITY PLAN to inform and direct your personnel about the specific quality practices, resources, sequence of activities, controls and checks that they have to implement during the deed. Include or reference in the PROJECT QUALITY PLAN the documents listed in Annexure Q/D plus any additional information nominated in the specifications for inclusion in the PROJECT QUALITY PLAN.

Associated technical documents that must be submitted with the PROJECT QUALITY PLAN include:

(a) Documentation required by the specifications or RMS Q Clause 7.5.1 to plan and implement controlled conditions; and

(b) Inspection and test plans and ITP forms that will be used by you to verify that the Project Works comply with the deed (refer RMS Q Clause 8.1.1).
4.2.2.4 Changes to the Project Quality Plan and Associated Documents

Immediately implement changes, where applicable, to the PROJECT QUALITY PLAN and corporate Quality Management System if the PROJECT QUALITY PLAN and associated quality management system documents:

(a) do not adequately address the Specification requirements; or
(b) are causing nonconformity; or
(c) have to be revised as a result of an audit; or
(d) no longer represent your current and/or appropriate practice.

Advise the RMS Representative promptly of any revisions to the PROJECT QUALITY PLAN or corporate Quality Management System and submit amended documentation detailing the revisions within 5 working days.

4.2.3 Control of Documents

Document a corporate quality management system procedure to address ISO 9001 Clause 4.2.3.

Describe in the PROJECT QUALITY PLAN how revisions to documents and data relevant to the Project Works and the Contractor's Work are to be identified in the document or appropriate attachments.

In addition to the documentation required by ISO 9001, copies of specifications, drawings, specified Test Methods, ISO 9001 and HB90.3 must be readily accessible on Site at all times. Copies of other documents referred to in the Specification must be available on site where required in the Specification. Ensure that copies of the remaining documents referred to in the Specification are accessible for reference by you either by the establishment of an on-site library, or by ready access to a library maintained elsewhere by you.

4.2.4 Control of Records

Document a corporate quality management system procedure to address ISO 9001 Clause 4.2.4.

The quality records must include all those shown in Annexure Q/E.

Implement a records management system in accordance with Clauses E1 and E2 of RMS Q. Use ISO 15489.1 and ISO 15489.2 for guidance in developing and implementing the records management system.

Prepare and submit a Records Management Plan (RMP) for your Work in accordance with Clauses E1 and E2 of RMS Q. The RMP must cover the record keeping practices, resources and sequence of activities required to meet all the requirements of RMS Q. The RMP must be consistent with the PROJECT QUALITY PLAN and include appropriate cross-referencing to the quality management system and PROJECT QUALITY PLAN.

Describe in the PROJECT QUALITY PLAN where the quality records shown in Annexure Q/E will be located and how they will be stored and maintained in accordance with the corporate quality management system procedure. Make the quality records available to the RMS Representative at all reasonable times. Where requested by the RMS Representative, permit the RMS Representative to make copies of quality records.
Prior to Construction Completion, provide the RMS Representative with any commissioning records and operation and maintenance manuals relevant to the Project Works.

Provide the RMS Representative with copies of any quality records within 14 days of a request by the RMS Representative.

4.2.5 Submission of Documents to RMS Representative

Submit documents in accordance with Annexure Q/A Table Q/A.1 and Table Q/A.2 and the deed.

When requested by the RMS Representative or the Project Verifier, for the purposes of quality audits, provide additional controlled copies of the QUALITY MANUAL, PROJECT QUALITY PLAN, RMP and associated quality management system documents.

4.2.6 (Not Used)

5 MANAGEMENT RESPONSIBILITY

5.1 MANAGEMENT COMMITMENT

No requirements additional to ISO 9001 Clause 5.1.

5.2 CUSTOMER FOCUS

No requirement additional to ISO 9001 Clause 5.2.

5.3 QUALITY POLICY

No requirements additional to ISO 9001 Clause 5.3.

5.4 PLANNING

5.4.1 Quality Objectives

Establish the Project Quality Objectives as part of the PROJECT QUALITY PLAN. The Project Quality Objectives must be relevant to the works required by the deed. Introduce the Project Quality Objectives to site management personnel (including relevant personnel of subcontractors) working on the project as part of the induction process (refer RMS Q Clause 6.2.2).

5.4.2 Quality Management System Planning

Plan the corporate Quality Management System using HB90.3 Clause 5.4.2 for guidance.

5.5 RESPONSIBILITY, AUTHORITY AND COMMUNICATION

5.5.1 Responsibility and Authority

Identify in the QUALITY MANUAL or corporate quality management system procedures those personnel who have responsibility for authorising corporate quality management system procedures.
Quality Management System (Type 6)  

The PROJECT QUALITY PLAN must:

(a) list the main responsibilities and authorities of personnel primarily responsible for upholding the quality management system provisions of the deed, including responsibilities for:
   (i) receiving, in-process and final (or acceptance) inspection and testing (refer RMS Q Clause 8.1);
   (ii) identifying/recording quality problems;
   (iii) initiating/recommending solutions through designated channels;
   (iv) ensuring that corrective action is implemented and effective;
   (v) communicating quality management requirements including solutions to problems;
   (vi) controlling further processing/delivery/installation of nonconforming product until deficiencies or unsatisfactory conditions have been corrected; and
   (vii) controlling monitoring and measurement devices.

(b) nominate the person responsible on site for main construction activities such as construction trials, placing concrete road base, concrete for reinforced structures, placing asphalt, bituminous spray sealing; and

(c) nominate the persons with the responsibility and authority for planning and implementing training and induction for the project, including establishing competence needed.

5.5.2 Management Representative

Nominate in the PROJECT QUALITY PLAN the Management Representative with corporate responsibility and authority for enacting ISO 9001 Clause 5.5.2.

The PROJECT QUALITY PLAN must:

(a) nominate your Project Quality Representative, directly responsible to top management and who has the defined authority and responsibility for ensuring that the requirements of the PROJECT QUALITY PLAN and associated quality management system procedures are implemented and maintained on the project; and

(b) where the Project Quality Representative is not your designated corporate Management Representative, indicate the relationship between the two persons.

Include in the PROJECT QUALITY PLAN the minimum qualifications and experience required of the Project Quality Representative. Include the actual qualifications of the Project Quality Representative in the training records (RMS Q Clause 6.2.2).

Establish the Project Quality Representative on site, if so specified in Annexure Q/A. If not required to be on site, the Project Quality Representative must be available for contact by telephone at all times when work is being carried out and be available to attend meetings on site within 24 hours of written or spoken notice by the RMS Representative.

5.5.3 Internal Communication

Implement internal communications, using HB90.3 Clause 5.5.3 for guidance.
5.6 MANAGEMENT REVIEW

Apply ISO 9001 Clause 5.6 for review of the corporate Quality Management System by top management, using HB90.3 Clause 5.6 for guidance.

The review must include the PROJECT QUALITY PLAN of this deed to confirm its continuing suitability and effectiveness for your Work. Include the agenda of items to be reviewed and proposed timetable for the reviews.

6 RESOURCE MANAGEMENT

6.1 PROVISION OF RESOURCES

No requirements additional to ISO 9001 Clause 6.1.

6.2 HUMAN RESOURCES

6.2.1 General

No requirements additional to ISO 9001 Clause 6.2.

6.2.2 Competence, Awareness and Training

Document a corporate quality management system procedure to address ISO 9001 Clause 6.2.2. Implement the procedure at corporate level and for project work.

Include in the PROJECT QUALITY PLAN a site-specific induction and training plan and induction and training procedures to describe the competence required, who is to be trained, when and how the training will be carried out.

Ensure that all on site personnel engaged on the project (including subcontractor’s personnel working under your Quality Management System) have undergone an appropriate induction program that explains how the Quality Management System is to be implemented on the project. At the request of the RMS Representative, make the induction program available to the RMS Representative staff.

6.3 INFRASTRUCTURE

No requirements additional to ISO 9001 Clause 6.3.

6.4 WORK ENVIRONMENT

No requirements additional to ISO 9001 Clause 6.4.
7 PRODUCT REALISATION

7.1 PLANNING OF PRODUCT REALISATION

Document a corporate quality management system procedure to address ISO 9001 Clause 7.1. Describe the method for preparing Project Quality Plans and associated documents. Include guidelines about how to determine project-specific requirements. Also include a template Project Quality Plan that complies with the Project Quality Plan requirements in RMS Q and that will be customised for RMS deeds.

7.2 CUSTOMER-RELATED PROCESSES

Document a corporate quality management system procedure to address ISO 9001 Clauses 7.2.1, 7.2.2 and 7.2.3 (c).

7.3 DESIGN AND DEVELOPMENT

Document a corporate quality management system procedure to address ISO 9001 Clause 7.3. Guidance is provided in HB90.3 Clause 7.3.

Include a template Design Plan to summarise the design planning process as described in ISO 9001 Clause 7.3.1.

Implement the design management procedure for design of the following, where required for the deed:
(a) Project Works and Temporary Works
(b) temporary structures and the checking of permanent structures for construction loadings;
(c) lifting devices for manufactured items;
(d) concrete mixes for structures and pavements and asphalt mixes for permanent works; and
(e) traffic management, temporary roadways and detours.

Include Design Plans in the PROJECT QUALITY PLAN for all design activities, including subcontractors engaged for design work. Where a design subcontractor does not have a Quality Management System conforming to the deed requirements, include the method of control and verification of the subcontractor's activities as part of the PROJECT QUALITY PLAN.

Notwithstanding RMS Q Clause 4.2 and Annexure Q/A, the submission of your Documents must comply with the deed requirements.

7.4 PURCHASING

7.4.1 Purchasing Process

Document a corporate quality management system procedure to address ISO 9001 Clause 7.4. Include a method to systematically plan and implement surveillance and inspection of subcontractors’ work.
Document in the PROJECT QUALITY PLAN how the subcontract requirements identified in Annexure Q/F will be included in subcontracts whenever they apply.

Where a supplier or subcontractor is to carry out work or provide services that require process validation (refer RMS Q Clause 7.5.2), evaluate the supplier or subcontractor on their capability to perform process validation. Document the method and results of this evaluation in the PROJECT QUALITY PLAN, including the details listed in Table Q/D.2.

### 7.4.2 Purchasing Information

The quality management system requirements detailed in RMS Q apply to all subcontracted products and services procured as part of your Work. This includes work process control documents and inspection/testing documents required by RMS Q Clauses 7.5.1 and 8.1.1.

Where any subcontractor is required to have RMS Prequalification or Registration, the subcontractor must use its quality management system that must conform to the quality management system requirements of the specified RMS Prequalification or Registration Category.

When a copy of a Subcontract is provided in accordance with the deed, include associated reference data (except price) and the applicable subcontract requirements listed in Annexure Q/F. When requested by the RMS Representative, also submit the evaluation of the subcontractor’s ability to meet subcontract requirements.

### 7.4.3 Verification of Purchased Product

Comply with RMS Q Clause 8.1(a) when receiving products from suppliers.

Include in the PROJECT QUALITY PLAN the subcontractors’ PROJECT QUALITY PLAN or process control documentation used to control processes and to verify purchased product.

Plan the extent of surveillance to be exercised for each subcontractor including management of information and records generated by subcontractors. When planning this surveillance, review the documents submitted by each subcontractor to ensure that all process control and inspection/testing requirements from the Specifications are adequately addressed. The surveillance process must include how nominated HOLD POINTS will be released and other activities to verify that the subcontractor’s output complies with the RMS Representative’s quality requirements.

Include in the PROJECT QUALITY PLAN the methods of surveillance that will be implemented for subcontracted work, in accordance with ISO 9001 Clause 7.4.3.

### 7.4.4 Use of Purchased Products

Ensure that purchased products are compatible with the other products and works and are handled, stored, combined with other products, installed and used in accordance with the manufacturer’s recommendations.

### 7.5 PRODUCTION AND SERVICE PROVISION

#### 7.5.1 Control of Production and Service Provision

Document a corporate quality management system procedure to address ISO 9001 Clause 7.5.1. Apply the procedure to plan, document, implement and monitor the controlled conditions for each work process. Consider the following (as appropriate) when planning work process controls:
Quality Management System (Type 6)  

(a) sequence of operations;
(b) types of equipment required, capability, maintenance, calibration;
(c) any special working environment aspects;
(d) competency and skills of personnel;
(e) work methods and materials to be used;
(f) product characteristics, tolerances and workmanship standards to be met;
(g) use of process control charts, when specified;
(h) inspection, test and control points;
(i) how the process will be monitored to ensure its continuing suitability;
(j) records to be kept as evidence that the work process controls remain effective; and
(k) defining responsibility for implementing and monitoring work process controls and rectifying any deficiencies.

Include the work process control documents in the PROJECT QUALITY PLAN (refer RMS Q Clause 4.2.2.3).

Treat and carry out survey as a separate application of work process control in accordance with RMS D&C G71.

7.5.2 Validation of Processes for Production and Service Provision

Identify in the PROJECT QUALITY PLAN any work processes (including subcontracted work) where the resulting output cannot be verified by subsequent monitoring and measurement. In such cases, control of the work processes must be documented and implemented in accordance with ISO 9001 Clause 7.5.2.

7.5.3 Identification and Traceability

Document in the PROJECT QUALITY PLAN how identification and traceability will be dealt with in accordance with ISO 9001 Clause 7.5.3.

Subdivide your Work into Lots or discrete work areas and control work in accordance with Annexure Q/L. Document in the PROJECT QUALITY PLAN the method(s) for subdividing the work into Lots or discrete work areas and for allocating Lot numbers (refer RMS Q Annexure Q/L Clause L1) to uniquely identify each Lot.

Any Lot that is visually non-homogeneous and/or non-representative must be rejected.

Identify all samples and test results with the field locations and Lot number, as applicable, to which they relate.

Maintain a register that identifies every Work Lot established for the deed.

Describe in the PROJECT QUALITY PLAN how traceability of the materials specified in RMS Q Annexure Q/G will be maintained.
7.5.4 Customer Property

Describe in the PROJECT QUALITY PLAN how ISO 9001 Clause 7.5.4 will be implemented for any property supplied by the RMS Representative for the Project Works or Contractor’s Work.

7.5.5 Preservation of Product

Describe in the PROJECT QUALITY PLAN how ISO 9001 Clause 7.5.5 will be implemented for transport, identification, handling, packaging, storage and protection on site to prevent damage or deterioration.

7.6 CONTROL OF MONITORING AND MEASURING DEVICES

Describe in the PROJECT QUALITY PLAN how ISO 9001 Clause 7.6 will be implemented for monitoring and measuring devices used to set out, construct or check the Contractor’s Work and to monitor the work environment (as appropriate), including selection of devices capable of the necessary accuracy and precision for the intended applications.

Monitoring and measuring devices include measuring equipment for production purposes (such as a concrete batching plant).

For laboratory testing equipment, NATA certification will be accepted as satisfying the requirements of ISO 9001 Clause 7.6.

Verify through the audit process, the control of laboratory equipment supplied and operated by subcontractors. Identify all inspection, measuring and test equipment (other than laboratory equipment) maintained and calibrated by subcontractors, which is used or proposed to be used for your Work. For all on-site activities where inspection, measuring and test equipment is maintained and calibrated by a subcontractor, ensure that the subcontractor holds, at the locations where the subcontracted work is being carried out, a valid calibration certificate or a copy of the subcontractor's equipment register showing the calibration status of the equipment.

8 MEASUREMENT, ANALYSIS AND IMPROVEMENT

8.1 GENERAL

In demonstrating conformity of the product to specified requirements, carry out inspection and testing:

(a) before any supplied product is used in the Project Works (receiving inspection and testing);
(b) progressively during construction of the Project Works (in-process inspection and testing); and
(c) as a final check that all inspection and testing necessary to demonstrate conformity of the Project Works to specified requirements has been carried out (final or acceptance inspection and testing).

8.1.1 Inspection and Test Planning

Document a corporate quality management system procedure to address ISO 9001 Clause 8.1a and RMS Q Clause 8.1. Prepare template Inspection and Test Plans (ITPs) and ITP forms.
Document ITPs and ITP forms for all inspection and testing required by the Specifications. Include these documents in the PROJECT QUALITY PLAN (refer RMS Q Clause 4.2.2.3).

The ITPs and/or ITP forms must indicate:

(a) who performs the receiving, in-process and final inspections or testing and at what stage of the work;

(b) how the inspection or test is to be carried out and recorded (e.g. as a documented testing procedure or by reference to a Standard test method);

(c) the acceptance criteria and frequency of inspection and testing. The detail for the acceptance criteria and frequency of inspection and testing and must replicate the nominated requirements of the Specifications. Reference to a specification clause alone is unacceptable;

(d) who reviews inspection/test results, evaluates whether work conforms, determines what to do next if work does not pass a required inspection or test and closes out Work Lots;

(e) when statistical analysis of test results is required (refer RMS Q Clause 8.2.4);

(f) when nonconformity control is addressed (refer RMS Q Clause 8.2.4.2) including closing out Work Lots (refer RMS Q Clause 8.2.4.3);

(g) who performs final review of all inspection/test results to confirm that all inspections and tests have been carried out to completely verify conformity for each Lot;

(h) the time limits for testing, time constraints for submission, and Hold and Witness Points that are nominated in the Specifications; and

(i) the requirements of RMS Q Clause 7.5.3 for Identification and Traceability and the sampling methods as required by RMS Q Clause L1.

8.1.2 Frequency of Testing

The frequency of testing must be appropriate to verify conformity and must not be less than that stated in the Specifications. Nominate appropriate frequencies even where no minimum frequency of inspection or testing is stated in the relevant Specification.

Include in the management review of the PROJECT QUALITY PLAN a review of the appropriateness of the frequency of testing nominated in the Inspection and Test Plan(s). Take into account the frequency of nonconformity detected, including nonconformities remedied by simple reworking.

The RMS Representative may conditionally agree to a proposal by you to reduce the specified minimum frequency of testing by up to 50% or as defined in the relevant Specification. The proposal must be supported by a statistical analysis verifying consistent process capability and product characteristics.

The specified minimum frequency of testing must be restored when a nonconformity is detected and until the RMS Representative agrees to a new proposal by you to reduce the specified minimum frequency. The RMS Representative may vary or restore the specified minimum frequency of testing, either selectively or permanently, at any time.

8.1.3 Inspection and Test Status

Describe in the PROJECT QUALITY PLAN, the method to be used for identifying and controlling the inspection and test status of all product and Project Works, including product and work which is incorporated in the Project Works prior to being verified as conforming.
If inspection/test records (such as a Lot register) do not clearly show the inspection and test status of each Lot or work area, Lots must be physically marked in the field to show whether they conform.

8.2 MONITORING AND MEASUREMENT

8.2.1 Customer Satisfaction

Describe in the PROJECT QUALITY PLAN, if specified in Annexure Q/A, the methods to be used to assess customer satisfaction during the project, in accordance with ISO 9001 Clause 8.2.1 (refer RMS Q Clause 5.2).

8.2.2 Internal Audit

Document a corporate quality management system procedure to address ISO 9001 Clause 8.2.2.

8.2.2.1 Audit Schedule

Incorporate in the PROJECT QUALITY PLAN an Audit Schedule for the project that identifies the following types of audit:

(a) audits of the operation of the Quality Management System, to evaluate the effectiveness of the Quality Management System as applied to the project;

(b) product or service audits, to assess the conformity of the product or service with the specified technical requirements; and

(c) audits of work process control, to evaluate how effectively work process controls are implemented in practice.

Include in audits the activities of subcontractors engaged on the project.

8.2.2.2 Adjustment to Audit Schedule

Adjust the audit schedule:

(a) when the results of previous audits indicate the need for a higher (or lower) audit frequency;

(b) when significant changes are made to functional areas of the quality management system, including reorganisations and revisions to procedure(s);

(c) when safety, performance or reliability of the product is in jeopardy, or suspected to be in jeopardy, due to nonconformity in the Quality Management System;

(d) when necessary to verify that the required corrective/preventive action has been taken; or

(e) when required due to changes in your Contract Program.

8.2.3 Monitoring and Measurement of Processes

Describe in the PROJECT QUALITY PLAN how ISO 9001 Clause 8.2.3 will be implemented to monitor the effectiveness of the work processes used for your Work.

Review each work process control and the associated documents and inspections and tests while that work process is in progress to monitor whether the planned controls are effective in achieving product conformity.
8.2.4 Monitoring and Measurement of Product

Implement the Inspection and Test Plans for the project, as established in RMS Q Clause 8.1.1.

Where acceptance characteristics are described in the Specifications in terms of characteristic values, apply statistical techniques to analyse test results in accordance with Annexure Q/L Clause L3 (refer RMS Q Clause 8.1.1(e)).

Document and maintain a method to confirm that all products or work Lots requiring inspection and/or testing are so inspected and/or tested (refer RMS Q Clause 7.5.3) at the required testing frequency. Include this method in the PROJECT QUALITY PLAN or ITP documentation.

Arrange sampling and testing to be performed in accordance with Annexure Q/L.

8.2.4.1 Hold Points

Describe in the PROJECT QUALITY PLAN the method of arranging for the release of HOLD POINTS by the Nominated Authority.

Do not proceed beyond a HOLD POINT until the Nominated Authority has released that HOLD POINT. Make suitable arrangements to notify the Nominated Authority when a HOLD POINT will be reached.

8.2.4.2 Inspection and Test Records

The inspection, test and verification records for each Lot or work area (refer RMS Q Clause 7.5.3) must:

(a) clearly show or reference the actual results obtained for any inspection and/or test and demonstrate conformity with the specified requirements;

(b) be progressively maintained as results are achieved; and

(c) indicate that control of nonconformity is addressed.

Make inspection, test and verification records available for evaluation by the Project Verifier and RMS Representative. Inspection and test records which are to be held on site, (refer Annexure Q/A), must be stored in a room reasonably accessible to the RMS Representative and Project Verifier with facilities for the inspection of the records. Access must not be limited by your other management activities.

8.2.4.3 Close Out of Work Lots and Release of Products

Work Lots must not be closed out nor product released, dispatched, used or installed until you have fully verified their conformity and incorporated the required inspection and/or test results/reports, including the documentation referred to in RMS Q Clause L2.2, into your records.

Work Lots and products must not be covered up until their conformity has been fully verified, except as permitted below in this Clause.

The RMS Representative recognises that some specified compliance testing might take over 48 hours to complete. In such circumstances, Work Lots/products may be covered up before the Lot is closed out, subject to the following conditions:

(a) you document an effective traceability/closure method in the PROJECT QUALITY PLAN that nominates the person or position responsible and describes how work or
product that may need to be covered up will be identified, traced, recorded and promptly verified and what action will be taken if full conformity is not achieved;

(b) you apply the traceability method on each occasion that a Work Lot or product is covered up under such circumstances;

(c) any specified verification survey has demonstrated conformity before covering up the work;

(d) you demonstrate on the basis of past work that it is highly unlikely that the work will fail to pass the specified compliance testing; and

(e) you apply the closure method and only close out the Work Lot after verifying that the work or product has passed the specified compliance testing.

Where product or work fails to pass any inspection and/or test, the Work Lot must not be closed out until the nonconformity has been rectified and closed out in accordance with RMS Q Clause 8.3.

8.3 CONTROL OF NONCONFORMING PRODUCT

Document a corporate quality management system procedure to address ISO 9001 Clause 8.3. Prepare a standard form for use as a Nonconformity Report.

Describe in the PROJECT QUALITY PLAN how the additional requirements of RMS Q Clause 8.3 will be implemented for the deed.

Identify and control all products or services that fail to pass any inspection or test in accordance with the defined acceptance criteria. Where conformity may be achieved by simple reworking or repair (that is, without reference to the Project Verifier and RMS Representative), record the required action in a format to suit your continual improvement procedures (refer RMS Q Clause 8.5).

Where conformity cannot be achieved by simply reworking with the original process, notify the Project Verifier and RMS Representative of the nonconformity and record it on an appropriate register. Submit a Nonconformity Report within 2 working days of detection of the nonconformity indicating the proposed rectification method, and when the rectification is to be undertaken.

If surveillance or an audit by the Project Verifier or RMS Representative indicates a nonconforming product that has not been addressed by a Nonconformity Report, the Project Verifier or RMS Representative will issue a ‘Nonconforming Product Notification’. This nonconforming product must be dealt with in the same manner as if you had identified it.

For any nonconforming product, do not proceed with the rectification work, or cover up, or further build on it until the proposed rectification method has been accepted by the Project Verifier. Where the proposed rectification method would result in the product remaining nonconforming with the deed, a Variation, pursuant to the provisions of Clause 15.6 or Clause 15.7 of the deed, or an amendment to the deed must be obtained by you from RMS to make the non-conforming product conforming. In evaluating the proposed rectification method, the Project Verifier or the RMS Representative may require additional supporting documentation, including engineering calculations or the opinion of a recognised technical expert in the field under consideration.

Acceptance of the proposed rectification method will be at the discretion of the Project Verifier.
8.4 ANALYSIS OF DATA

No requirements additional to ISO 9001 Clause 8.4.

8.5 IMPROVEMENT

8.5.1 Continual Improvement

No requirements additional to ISO 9001 Clause 8.5.1.

8.5.2 Corrective Action

Document a corporate quality management system procedure to address ISO 9001 Clause 8.5.2.

Establish and maintain a Corrective Action Register to record a summary of corrective actions or list those records that demonstrate corrective actions.

If surveillance or an audit by the Project Verifier or RMS Representative indicates that your Quality Management System does not comply with the provisions of the deed or that a condition adverse to quality exists, the Project Verifier or RMS Representative may issue a Corrective Action Request.

Rectify any nonconformity or condition adverse to quality notified by the Project Verifier or RMS Representative. Take corrective/preventive action to prevent recurrence of the nonconformity or remove the condition adverse to quality and return the completed Corrective Action Request, all within 7 days after the Corrective Action Request is given to you.

Address the Corrective Action Request in accordance with your arrangements for handling customer complaints as required by ISO 9001 Clause 7.2.3(c).

HOLD POINT
(Where required by the Nominated Authority)
Process Held: The Process referred to in the Corrective Action Request.
Submission Details: Details of the corrective action.
Release of Hold Point: The Nominated Authority will consider the submitted documents prior to authorising the release of the Hold Point.
Enter details of the developed corrective action onto the Nonconformity Report or Corrective Action records, as appropriate.

8.5.3 Preventive Action

Document a corporate quality management system procedure to address ISO 9001 Clause 8.5.3. Include identification and communication of opportunities for improvement to the quality management systems of subcontractors, the Project Verifier and RMS Representative.

Establish and maintain a Preventive Action Register to record a summary of preventive actions or list those records that demonstrate preventive actions.

You may nominate to the RMS Representative and the Project Verifier those preventive actions records, or parts thereof, which are commercially sensitive and restrict access to them, as agreed with the RMS Representative.

9 RMS REPRESENTATIVE AND PROJECT VERIFIER SURVEILLANCE AND AUDITS

9.1 GENERAL

All testing by the Project Verifier or RMS Representative associated with surveillance and audits will be conducted by a laboratory with NATA accreditation for the test methods specified. The results of such testing will be recorded on NATA endorsed test reports. If NATA has not accredited a laboratory for a test, the test must be carried out at a laboratory approved by the RMS Representative and the Project Verifier.

9.2 QUALITY MANAGEMENT SYSTEM, PROCESS QUALITY AND PRODUCT QUALITY AUDITS AND SURVEILLANCE.

Quality management system audits by the Project Verifier or RMS Representative may be conducted on a scheduled basis on all aspects of your Quality Management System and will be performed in accordance with recognised audit procedures.

The Project Verifier or RMS Representative will give you at least 5 days notice that a quality management system audit is to be conducted.

Surveillance, process quality audits and product quality audits by the Project Verifier or RMS Representative may be conducted at any time.

If surveillance or an audit indicates a significant nonconformity of a product or service, the Project Verifier or RMS Representative is entitled to conduct a quality management system audit at twenty four hours notice to you.

Make suitable facilities available at the site to accommodate an audit team of three persons.

9.3 (NOT USED)
ANNEXURE Q/A – DETAILS OF WORK

A1 PROJECT SPECIFIC REQUIREMENTS

<table>
<thead>
<tr>
<th>Clause</th>
<th>Description</th>
<th>Requirement</th>
</tr>
</thead>
<tbody>
<tr>
<td>5.5.2</td>
<td>Establish the Project Quality Representative on site:</td>
<td>Yes</td>
</tr>
<tr>
<td>8.2.1</td>
<td>Describe in the PQP the methods to assess customer satisfaction:</td>
<td>Yes</td>
</tr>
<tr>
<td>Annexure Q/C Clause C2</td>
<td>Site records more than 35 days old must be stored on site:</td>
<td>Yes/No</td>
</tr>
</tbody>
</table>

Project Testing also applies to the following tests:  
(Annexure Q/L Clause L2.2.1)

A2 DOCUMENT SUBMISSION REQUIREMENTS

Documents and records must be submitted in accordance with this matrix. Refer to the Specifications for full details of submission requirements.

Table Q/A.1 – Document and Record Submission Matrix

<table>
<thead>
<tr>
<th>Document</th>
<th>Number of Copies to be Submitted</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>First Stage</td>
</tr>
<tr>
<td></td>
<td>14 days prior to use</td>
</tr>
<tr>
<td>Quality Manual and quality policy</td>
<td>1</td>
</tr>
<tr>
<td>PROJECT QUALITY PLAN</td>
<td>2</td>
</tr>
<tr>
<td>Applicable quality management system procedures</td>
<td>1</td>
</tr>
<tr>
<td>Process control procedures</td>
<td>2</td>
</tr>
<tr>
<td>Inspection and test plans and record forms</td>
<td>2</td>
</tr>
<tr>
<td>Quality records</td>
<td>(3)</td>
</tr>
<tr>
<td>Index of quality records</td>
<td>(4)</td>
</tr>
</tbody>
</table>

Notes:
(1) Copies will be returned on request.
(2) Also required within 35 days of acceptance of tender.
(3) Copy must be provided for RMS Representative’s records as specified or as directed.
(4) Prior to Construction Completion (RMS Q Clause E1.4.1).
ANNEXURE Q/B – (NOT USED)

ANNEXURE Q/C – SCHEDULES OF HOLD POINTS AND IDENTIFIED RECORDS

Refer to RMS Q Clause 1.2.3.

C1 SCHEDULE OF HOLD POINTS

<table>
<thead>
<tr>
<th>Clause</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td>8.3</td>
<td>Covering up of rectified work</td>
</tr>
<tr>
<td>8.5.2</td>
<td>The Process referred to in the Corrective Action Request</td>
</tr>
</tbody>
</table>

Note: The imposition of a Hold Point is at the direction of the Project Verifier or RTA’s Representative

C2 SCHEDULE OF QUALITY RECORDS AND IDENTIFIED RECORDS

The Quality Records and reference documents must include the following. Until the Date of Construction Completion, the originals or copies must be at the following locations, unless otherwise agreed by the RMS Representative. These records must be made available to the RMS Representative.

Records located with the RMS Representative (indicated by “R” in the table) are Identified Records for the purposes of RMS Q Annexure Q/E.

Site records more than 35 days old (if permitted in RMS Q Annexure Q/A) may be stored off site, but must be available on site within 24 hours of notice given by the RMS Representative. All records must be made available to the RMS Representative upon request until the Date of Construction Completion.

<table>
<thead>
<tr>
<th>Clause</th>
<th>System Requirement</th>
<th>Required Record or Reference</th>
<th>Location</th>
</tr>
</thead>
<tbody>
<tr>
<td>4.2.2</td>
<td>Quality Management System documents</td>
<td>QUALITY MANUAL, quality policy and PROJECT QUALITY PLAN, Quality Management System procedures when applicable</td>
<td>S &amp; R</td>
</tr>
<tr>
<td>4.2.3</td>
<td>Control of documents</td>
<td>List of who holds issued documents  Register of current document issue/revisions</td>
<td>S  S</td>
</tr>
<tr>
<td>4.2.4</td>
<td>Control of records</td>
<td>Index of all quality records (prior to Construction Completion)</td>
<td>R</td>
</tr>
<tr>
<td>5.6</td>
<td>Management review</td>
<td>Records of management reviews for the project</td>
<td>S</td>
</tr>
<tr>
<td>6.2.2</td>
<td>Competence, awareness and training</td>
<td>Personnel qualifications/skills records  Induction and training records</td>
<td>P  S</td>
</tr>
<tr>
<td>6.4</td>
<td>Work environment</td>
<td>Records of work environment controls, where applicable</td>
<td>S</td>
</tr>
<tr>
<td>7.2.2</td>
<td>Review of requirements</td>
<td>Minutes of tender/deed reviews</td>
<td>M</td>
</tr>
<tr>
<td>Clause</td>
<td>System Requirement</td>
<td>Required Record or Reference</td>
<td>Location</td>
</tr>
<tr>
<td>--------</td>
<td>--------------------</td>
<td>-----------------------------</td>
<td>----------</td>
</tr>
<tr>
<td>7.3</td>
<td>Design and development</td>
<td>Design Plan, inputs/outputs, changes, verification/review/ validation records</td>
<td>D &amp; P</td>
</tr>
<tr>
<td>7.4</td>
<td>Purchasing</td>
<td>Evaluation of subcontractors and suppliers</td>
<td>P &amp; S</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Surveillance, audit of subcontractors</td>
<td>S</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Subcontractor supplied documentation</td>
<td>P &amp; S</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Certificate of testing by suppliers</td>
<td>S</td>
</tr>
<tr>
<td>7.5.1</td>
<td>Control of production and service provision</td>
<td>Procedures describing how to control work processes</td>
<td>P &amp; R</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Records demonstrating effectiveness of work process controls</td>
<td>S</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Records of process validation when applicable</td>
<td>S</td>
</tr>
<tr>
<td>7.5.3</td>
<td>Identification and traceability</td>
<td>Product batch/traceability records</td>
<td>S</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Lot Identification Register</td>
<td>S</td>
</tr>
<tr>
<td>7.5.4</td>
<td>Customer property</td>
<td>Contractor’s verification records/reports</td>
<td>S</td>
</tr>
<tr>
<td>7.5.5</td>
<td>Preservation of product</td>
<td>Delivery dockets</td>
<td>S</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Product preservation control/inspection records</td>
<td>S</td>
</tr>
<tr>
<td>7.6</td>
<td>Control of monitoring and measuring devices</td>
<td>Register of equipment</td>
<td>P</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Calibration frequency and certificates</td>
<td>P</td>
</tr>
<tr>
<td>8.1.1</td>
<td>Inspection and test planning</td>
<td>Inspection and test plans</td>
<td>S &amp; R</td>
</tr>
<tr>
<td>8.2.1</td>
<td>Customer satisfaction</td>
<td>Customer satisfaction records and actions taken to improve customer satisfaction</td>
<td>P</td>
</tr>
<tr>
<td>8.2.2</td>
<td>Internal audit</td>
<td>Audit reports</td>
<td>S</td>
</tr>
<tr>
<td>8.2.4.2</td>
<td>Inspection and test records</td>
<td>Records/checklists of inspection and testing</td>
<td>P</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Conformity reports for each completed Lot</td>
<td>S</td>
</tr>
<tr>
<td>8.3</td>
<td>Control of nonconforming product</td>
<td>Nonconformity reports</td>
<td>S &amp; R</td>
</tr>
<tr>
<td></td>
<td></td>
<td>RMS Representative and Project Verifier</td>
<td>S &amp; R</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Nonconforming Product Notifications</td>
<td>S &amp; R</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Nonconformity Register</td>
<td>P</td>
</tr>
<tr>
<td>8.4</td>
<td>Analysis of data</td>
<td>Records of analysis of data generated during the deed</td>
<td>S</td>
</tr>
<tr>
<td>8.5.2</td>
<td>Corrective action</td>
<td>Corrective action reports and Register</td>
<td>S</td>
</tr>
<tr>
<td></td>
<td></td>
<td>RMS Representative and Project Verifier</td>
<td>S &amp; R</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Corrective Action Requests</td>
<td>S &amp; R</td>
</tr>
<tr>
<td>8.5.3</td>
<td>Preventive action</td>
<td>Preventive action reports and Register</td>
<td>S &amp; R</td>
</tr>
</tbody>
</table>

Legend of Document Location: D = Office of designer; R = RMS Representative; S = Site; M = Office of Management Representative with executive responsibility; P = Principal place where the document is used.
ANNEXURE Q/D – PLANNING DOCUMENTS

Refer to RMS Q Clause 1.2.4.

The PROJECT QUALITY PLAN and its references must, as a minimum, include the following, when applicable.

Table Q/D.1 – PROJECT QUALITY PLAN

<table>
<thead>
<tr>
<th>Clause</th>
<th>Required Planning Document or Reference</th>
</tr>
</thead>
<tbody>
<tr>
<td>1.3</td>
<td>Matrix explaining the quality management system if it is not based on AS/NZS ISO 9001:2008</td>
</tr>
<tr>
<td>4.2.2.2</td>
<td>RMS specific procedures when they are not incorporated into the corporate system procedures</td>
</tr>
<tr>
<td>4.2.2.2</td>
<td>Description of applicable corporate quality management system procedures.</td>
</tr>
<tr>
<td>4.2.3</td>
<td>Description of how changes to documents and data relevant to your Work are to be identified in the document or appropriate attachments</td>
</tr>
<tr>
<td>4.2.4</td>
<td>Description of how quality records will be stored and maintained</td>
</tr>
<tr>
<td>5.4.1</td>
<td>Project Quality Objectives established when specified in Annexure Q/A</td>
</tr>
<tr>
<td>5.5.1</td>
<td>List of main responsibilities and authorities of personnel primarily responsible for quality assurance activities in your Work</td>
</tr>
<tr>
<td>5.5.2</td>
<td>Nominate Management Representative and Project Quality Representative. Describe reporting relationship, if these positions are held by different people</td>
</tr>
<tr>
<td>5.5.2</td>
<td>Minimum qualifications and experience required of the Project Quality Representative</td>
</tr>
<tr>
<td>6.2.2</td>
<td>Site specific induction and training plan and procedures</td>
</tr>
<tr>
<td>7.3</td>
<td>Design Plans for all design activities, including subcontractors engaged for design work</td>
</tr>
<tr>
<td>7.4.1</td>
<td>Describe how Annexure Q/F requirements will be included in subcontracts (when applicable)</td>
</tr>
<tr>
<td>7.4.1</td>
<td>Document method and results of subcontractor evaluation for work processes that require process validation (when applicable)</td>
</tr>
<tr>
<td>7.4.1</td>
<td>Purchase planning details (refer Table Q/D.2)</td>
</tr>
<tr>
<td>7.4.2</td>
<td>Subcontractor's PROJECT QUALITY PLAN or process control documentation for each subcontract</td>
</tr>
<tr>
<td>7.4.3</td>
<td>Method of surveillance for subcontracted work</td>
</tr>
<tr>
<td>7.5.1</td>
<td>Work process control documents</td>
</tr>
<tr>
<td>7.5.2</td>
<td>Identification of work processes where the resulting output cannot be verified by subsequent monitoring and measurement</td>
</tr>
<tr>
<td>7.5.3</td>
<td>Dealing with identification and traceability</td>
</tr>
<tr>
<td>7.5.3</td>
<td>Method of maintaining traceability for materials listed on Annexure Q/G</td>
</tr>
<tr>
<td>7.5.3</td>
<td>Methods for subdividing the work into Lots and allocating Lot numbers</td>
</tr>
<tr>
<td>7.5.4</td>
<td>Description or how RMS supplied property will be dealt with</td>
</tr>
<tr>
<td>Clause</td>
<td>Required Planning Document or Reference</td>
</tr>
<tr>
<td>--------</td>
<td>----------------------------------------</td>
</tr>
<tr>
<td>7.5.5</td>
<td>Method of preserving products</td>
</tr>
<tr>
<td>7.6</td>
<td>Method of control of monitoring and measuring devices</td>
</tr>
<tr>
<td>8.1.1</td>
<td>Inspection and Test Plans, ITP Record Forms for all inspection and testing required by the Specifications</td>
</tr>
<tr>
<td>8.1.3</td>
<td>Method for identifying and controlling inspection and test status</td>
</tr>
<tr>
<td>8.2.1</td>
<td>Methods to assess customer satisfaction</td>
</tr>
<tr>
<td>8.2.2.1</td>
<td>Audit schedule</td>
</tr>
<tr>
<td>8.2.3</td>
<td>Method of monitoring the effectiveness of processes for your Work</td>
</tr>
<tr>
<td>8.2.4</td>
<td>Method for verifying that all inspection and/or testing is carried out at the required testing frequency</td>
</tr>
<tr>
<td>8.2.4.1</td>
<td>Method for release of Hold Points</td>
</tr>
<tr>
<td>8.2.4.3</td>
<td>Traceability/closure method for close-out of Work Lots</td>
</tr>
<tr>
<td>8.3</td>
<td>Additional requirements of RMS Q Clause 8.3</td>
</tr>
<tr>
<td></td>
<td>Any other documents or information required by the specifications to be included in the PROJECT QUALITY PLAN</td>
</tr>
</tbody>
</table>

NOTE: Where the above documents are liable to change, they may be referenced in the PROJECT QUALITY PLAN and attached as an annexure to expedite their revision.

**Table Q/D.2 – Purchase Planning Details**

Purchase planning details to be included in the PROJECT QUALITY PLAN (refer RMS Q Clause 7.4.1) must include the following:

<table>
<thead>
<tr>
<th>Required Document or Reference</th>
</tr>
</thead>
<tbody>
<tr>
<td>(a) Types of product or service subcontracted</td>
</tr>
<tr>
<td>(b) Purchasing schedule which states the timing of procurement of the product or service to be subcontracted and includes provision for the approval process</td>
</tr>
<tr>
<td>(c) Your method of assessment of the subcontractor's ability to meet the subcontract requirements including the quality management system requirements specified</td>
</tr>
<tr>
<td>(d) Your plan for inspection and surveillance of the subcontractor to verify the operation of the quality management system and product conformity requirements</td>
</tr>
<tr>
<td>(e) All specified inspection and testing shown in the subcontractor's Inspection and Test Plans</td>
</tr>
</tbody>
</table>
ANNEXURE Q/E – RECORD KEEPING AND IDENTIFIED RECORDS

Refer to RMS Q Clauses 1.2.5 and 4.2.4.

E1 RECORD KEEPING

E1.1 General

The work to be executed under Clauses E1 and E2 of RMS Q consists of:

(a) Development and implementation of a RECORDS MANAGEMENT PLAN (RMP);

(b) Operation, maintenance and review of the RMP during the Contractor’s Work and thereafter as required by RMS Q; and

(c) Secure storage of project records and delivery of Identified Records.

E1.2 Record Keeping Definitions

The terms and definitions in ISO 15489.1 apply to RMS Q. In addition the following definitions are applicable:

**Project Record.** Any record generated to document the execution of the project but does not include records which are commercial in confidence or relate to staff confidential matters.

**Quality Record.** A record used to demonstrate conformity to specified requirements and effective operation under AS/NZS ISO 9001 or required by RMS Q.

**Identified Record.** Any project record that is named as an Identified Record in the deed.

E1.3 Contractor’s Records Management Plan (RMP)

**E1.3.1 Scope of RMP**

Include procedures for the following in the RMP:

(a) the systematic control of the creation, registration, indexing, filing, maintenance, storage, movement, retrieval and disposal of project records related to the deed;

(b) providing to the RMS Representative the information required under the deed;

(c) submission and delivery of Identified Records as specified;

(d) disaster recovery plan in accordance with Clause E1.3.5; and

(e) providing a list of Identified Records relevant to the deed which must be progressively handed over to the RMS.

Include an index of project records in the RMP that is consistent with the records management system. Keep the index up-to-date until the end of the Landscaping Maintenance period and provide the index to the RMS Representative whenever the index is revised.

Include a list of Identified Records relevant to the deed in the RMP.

Revise the RMP and implement more appropriate record keeping practices if the original record keeping practices prove not to be fully effective. Apply Clause 4.2.3 of RMS Q whenever the RMP is revised.
When requested by the RMS Representative or the Project Verifier for the purposes of audit, provide up to three additional controlled copies of the RMP and relevant documented procedures. These documents will be returned when no longer required by the RMS Representative or the Project Verifier.

**E1.3.2 Record Keeping Policy**

Establish a record keeping policy as part of the RMP. The record keeping policy must define the record keeping objectives of the project and be relevant to the works required by the deed. Introduce the policy to all appropriate personnel working on the project (including appropriate personnel in Subcontractors' workforces) as part of the induction process (refer to Clause 6.2.2 of RMS Q).

**E1.3.3 Project Management**

Nominate in the RMP a full time member of your site management team to be the authorised contact person for communications with the RMS Representative and the Project Verifier on record keeping matters. Detail their responsibilities in the RMP, include filing, indexing, storage, movement, retrieval and disposal of records. This person must be fully conversant with the RMP, your record keeping system and progress with hand-over of Identified Records and must promptly provide access to or copies of records to the RMS Representative or the Project Verifier as required.

**E1.3.4 Monitoring and Compliance**

The RMP must state the requirements for review of the RMP which must be carried out by a senior officer of your management. Confirm the continuing suitability and effectiveness of the RMP for your Work.

Undertake compliance audits of the RMP at intervals of not more than six months, preferably in conjunction with quality management system audits.

**E1.3.5 Disaster Recovery Plan**

The records management system must address disaster preparedness to ensure that risks are identified and mitigated.

Records that are particularly critical for business continuity may require additional methods of protection and duplication to ensure accessibility in the event of a disaster.

Develop a disaster recovery plan that defines an organised and prioritised response to the disaster, planning for the continuance of regular business operations during the disaster and making appropriate plans for recovery after the disaster.

Integrity must be demonstrably maintained during and after recovery from disaster.

In the development of a disaster recovery plan, you may refer to Appendix B of AS 4390.6-1996 (superseded).

**E1.4 Project Records: Basic Record Keeping Requirements**

**E1.4.1 General**

Project records include the following:
(a) Quality Records as shown in Clause C2 of RMS Q;
(b) Records of Environmental Activities in accordance with RMS D&C G36, Clause 4.6;
(c) Records related to Work Health and Safety activities in accordance with RMS D&C G22;
(d) Contract Programs, as required by the deed;
(e) Identified Records that are a sub-set of the project records – see Clause E2 of RMS Q.

The project records must be:

(i) sufficiently comprehensive to demonstrate compliance with deed Specifications. This includes subcontractor and supplier records, where relevant;

(ii) accurate, legible and fully completed;

(iii) kept in order, particularly in the case of multi-page records;

(iv) filed in such a way that individual records can be readily retrieved;

(v) filed promptly after they are generated or received;

(vi) securely maintained to prevent unauthorised access, alteration, removal, deterioration, damage or loss;

(vii) kept track of where authorised removal or transfer of records within the company is permitted; and

(viii) entered on a register which shows what records are handed over to the RMS Representative, the Project Verifier or sent to other parties, including the date and method of hand-over/dispatch.

Make all project records available to the RMS Representative or the Project Verifier at all reasonable times. When requested by the RMS Representative or the Project Verifier, permit the RMS Representative or the Project Verifier to copy such records.

By the end of the Landscaping Maintenance period, provide the up-to-date index of all project records to the RMS Representative. Following the provision of that index, provide copies of any project records within 14 days of a request by the RMS Representative.

E1.4.2 Form of Records

Keep records as paper files or in electronic form in accordance with Clause 9 of ISO 15489.1 and Clause 4 of ISO 15489.2 unless otherwise provided in the deed or agreed with the RMS Representative.

Store records on media that ensure their useability, reliability, authenticity and preservation in accordance with the deed. The media and formats used for making records must be in accordance with Clause 9 of ISO 15489.1.

E1.4.3 Storage

Store and maintain project records or copies thereof such that they are readily retrievable, in facilities that provide a suitable environment to minimise deterioration or damage, and to prevent loss.

Store project records:
Quality Management System (Type 6) D&C Q6

(a) Prior to the end of the Landscaping Maintenance Period, at the location or locations specified in the deed or, failing such specified location, at the principal place where the respective records are used or such other place as may be agreed with the RMS Representative.

(b) After the end of the Landscaping Maintenance Period, at a location within the RMS Region in which your Work was carried out or at a location within the Sydney metropolitan area, as agreed with the RMS Representative. Inform the RMS Representative of the street address of the location. The location must not be changed without the approval of the RMS Representative.

E1.4.4 Retention Period

Keep all project records for a minimum period of five years after Final Completion. This requirement continues to apply even though the records or copies of the records may have been given to the RMS Representative and the RMS Representative may have taken copies of the records.

E1.4.5 Disposal of Records

The records must be pulped, shredded or burned in industrial facilities when disposing of records after the expiration of the retention period. Dumping of project records is not permitted.

E2 IDENTIFIED RECORDS

E2.1 General

Deliver Identified Records to the RMS Representative in accordance with Clauses E1 and E2 of RMS Q.

Each RMS Specification includes as an Annexure a list of Identified Records applicable to that Specification. Compile a list of all Identified Records for all RMS Specifications included in the deed and include the list in the RMP.

The RMS Representative may direct, as a variation, that records be added to or deleted from the list. Deal with those records as Identified Records as from the time of receipt of the RMS Representative direction.

For the purposes of the definition of Construction Completion in the deed, Identified Records which are then in existence, or must be in existence to comply with the provisions of the deed, are deemed to be essential for the operation, use and maintenance of your Work.

E2.2 Form of Identified Records

Unless otherwise provided in the deed or agreed with the RMS Representative, all Identified Records, except for work-as-executed drawings, must be delivered to the RMS Representative in hard copy on paper size A3 or A4 in good quality file housing/covering.

The submission of work-as-executed drawings must comply with the requirements in Scope of Works and Technical Criteria.
The paper must be premium bond paper of minimum weight 80 grams per square metre. The colour of the paper used must be white, unless the record is normally maintained in some other colour for ease of identification, in which case that particular colour may be used.

The use of thermal paper is not acceptable.

With the agreement of the RMS Representative, Identified Records may be provided in electronic form if it can be demonstrated that:

(a) the records have been properly captured into an electronic record keeping system and the records remain accessible, authentic, reliable and useable through any kind of system change until Final Completion and for the entire retention period; and

(b) the records are compatible with the RMS’ electronic records management system.

**E2.3 Filing and Indexing**

File Identified Records as required to be prepared under the relevant RMS Specification. They must be uniquely numbered and filed in chronological order under the heading of the relevant RMS Specification by which they were created. Include an index for quick reference. Alternatively, you may submit for the RMS Representative approval an alternative means of filing and indexing which provides the information required by this Clause E2.3 and forms part of your standard record keeping system.

**E2.4 Delivery**

Deliver Identified Records to the RMS Representative progressively during the course of the deed at the times specified in the deed or, if not so specified, at such times or within such periods as may be agreed with the RMS Representative.

The RMS Representative may withhold Final Completion until such time as all the Identified Records have been delivered. Make arrangements with the RMS Representative for the delivery of any outstanding Identified Records.

In the event that the RMS Representative elects to exercise the power conferred on him by the deed to terminate the deed, all project records, including Identified Records, then in your possession become the property of the RMS Representative forthwith and must be handed over to the RMS Representative.

**E2.5 Records to be Kept**

Notwithstanding that project records and Identified Records have been delivered to the RMS Representative or the RMS Representative has taken copies of project records or Identified Records, retain the originals of those records or, where originals are not held by you, good quality copies of the records, for the period specified in Clause E1.4.4 of RMS Q.

Where electronic records are provided, keep a copy of those records in accordance with the provisions of RMS Q.
Subcontracts for products and services provided to the RMS Representative (refer to RMS Q Clause 7.4.2) must include the following requirements, as applicable.

<table>
<thead>
<tr>
<th>Subcontract Requirements</th>
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<tbody>
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<td>(i)</td>
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<td>(ii)</td>
</tr>
<tr>
<td>(iii)</td>
</tr>
<tr>
<td>(iv)</td>
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<td>(xiii)</td>
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<td>(xiv)</td>
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<td>(xv)</td>
</tr>
<tr>
<td>(xvi)</td>
</tr>
</tbody>
</table>
Refer to RMS Q Clause 7.5.3 – Apply traceability to the following products:

<table>
<thead>
<tr>
<th>Specification</th>
<th>Product</th>
<th>Requirement</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Concrete batches used in bridge components, cast-in-place box culverts</td>
<td>The trace must start at the batch plant and finish at the location where the material is incorporated in the Project Works. Records must be kept of the batch quantities and time, testing details and location of placement.</td>
</tr>
<tr>
<td></td>
<td>and retaining walls.</td>
<td></td>
</tr>
<tr>
<td></td>
<td>Concrete batches used in road pavement sub-base and base.</td>
<td>The trace must start at the batch plant and finish at the location where the material is incorporated in the Project Works. Records must be kept of the batch quantities and time, testing details and location of placement.</td>
</tr>
<tr>
<td></td>
<td>Stabilised material used in road pavement.</td>
<td>The trace must start at the batch plant and finish at the location where the material is incorporated in the Project Works. Records must be kept of the batch quantities and time, testing details and location of placement.</td>
</tr>
<tr>
<td></td>
<td>Asphalt used in wearing courses, intermediate courses and drainage</td>
<td>The trace must start at the batch plant and finish at the location where the material is incorporated in the Project Works. Records must be kept of the batch quantities and time, testing details and location of placement.</td>
</tr>
<tr>
<td></td>
<td>layers.</td>
<td></td>
</tr>
<tr>
<td></td>
<td>Steel plate in bridge girders and bridge columns.</td>
<td>The trace must start at the steelworks and finish at the location of the plate in the girder or column. Records must be kept of the steel heat number, testing details and location of the plate in the girder or column.</td>
</tr>
</tbody>
</table>
ANNEXURES Q/H TO Q/J – (NOT USED)

ANNEXURE Q/K – SURVEY PROCEDURES

Refer to RMS Q Clause 7.5.1. Carry out survey in accordance with RMS D&C G71.

ANNEXURE Q/L – TESTING PROCEDURES

This Annexure provides details on the content of sampling and testing procedures to be included in the PROJECT QUALITY PLAN, when applicable.

L1 IDENTIFICATION OF WORK LOTS

Refer to RMS Q Clause 7.5.3.

Your Work must be subdivided into Lots or discrete work areas. A Lot must consist of a continuous portion of homogeneous and/or representative material or end product produced under essentially constant conditions. Discrete portions of a Lot that are visually non-homogeneous and/or non-representative must be excluded from the Lot and must be either treated as separate Lots or reworked to achieve conformity to the Specification.

The size of a Lot must not exceed one day's output except that this period may be extended by agreement with the Project Verifier where full conformity cannot be achieved in one day.

Describe in the PROJECT QUALITY PLAN how the Lot is to be identified in the field.

Determine the bounds of each Lot before sampling. Set the bounds of each Lot so that each Lot is represented by a single tested sample, except where statistical methods (which require several tested samples to represent a Lot) are used. Each acceptance criterion may have different Lot boundaries. Demonstrate the relationship of the boundaries of all adjacent Lots to confirm that the Lots represent the total work.

Give each Lot a unique Lot number. Use this Lot number as an identifier on all quality records. The Lot numbering system must be compatible with any activity numbering system used for your Contract Program. Record the Lot number on an appropriate register that indicates the three-dimensional location of the Lot. Include in the PROJECT QUALITY PLAN details of the Lot numbering system and the place where the Lot register is kept. Record chainages of start and finish, lateral location and layer location. When the Lot number does not indicate the location of the Lot, agree the method for identification of the Lot with the Project Verifier.

L2 TESTING SERVICES

L2.1 General

L2.1.1 National Association of Testing Authorities (NATA) Accreditation

Apply the requirements of Clause L2 to all laboratories verifying conformity of materials and work used for the deed. Ensure that all suppliers and subcontractors required to use or supply
tested materials or work are informed of and implement the requirements of Clause L2 and the testing requirements in the relevant specifications.

If NATA has not accredited a laboratory for a test, the test must be carried out by a laboratory accredited for the test by an organisation mutually recognised by NATA and approved by the RMS Representative and the Project Verifier. The test must be carried out and results endorsed in accordance with accreditation conditions or by a laboratory approved by the RMS Representative and the Project Verifier with results reported in a format also acceptable to the RMS Representative and the Project Verifier.

**L2.1.2 Laboratory Independence**

You and the laboratories used for the testing must ensure objectivity and impartiality in sampling, testing and reporting of results. The laboratories must act independently of you, any of your subcontractors and the RMS Representative in conducting the sampling and testing.

Independence, for the purposes of this clause, means “management and personnel are free from any undue internal and external, commercial, financial and other pressures and influences that may adversely affect quality of their work” (refer AS ISO/IEC 17025 Clause 4.1.5(b)).

**L2.1.3 Sampling Personnel**

Sampling, including selection of locations, must be conducted by personnel either accredited by NATA for that sampling procedure or who are from a NATA accredited laboratory or approved by the RMS Representative and the Project Verifier and who have been assessed as proficient for that sampling procedure and must be supervised by an officer having NATA signatory approval for that process.

**L2.1.4 Test Certificates**

Test results for each Lot must be reported in NATA endorsed documentation.

**L2.2 Project Testing**

**L2.2.1 Project Laboratories**

In addition to Clause L2.1, Clause L2.2 applies for all samples and tests carried out on the site, at concrete and asphalt batch plants, on aggregates and materials used for pavements and structures at off site locations and any other testing specified in Annexure Q/A to be Project Testing.

Engage one or more Project Testing laboratories that hold NATA accreditation at your own cost.

As a condition of appointment by you (refer RMS Q Clause 7.4), laboratories, material suppliers or subcontractors who have NATA accreditation must provide to you, the RMS Representative and the Project Verifier, on request, copies of NATA audits, relevant to the type of tests carried out for the deed.

The RMS Representative and the Project Verifier may forward to NATA copies of test records, certificates, reports of surveillance, performance and audits of any laboratory used for sampling and testing the conformity of materials and work.

The same Project Testing laboratory responsible for testing the sample must undertake the sampling, unless otherwise approved by the RMS Representative and the Project Verifier.
Should you propose sampling and/or testing by personnel other than from the laboratory, you must, prior to sampling and/or testing commencing, submit a proposal for the RMS Representative and the Project Verifier's approval showing the sampling and/or testing to be performed, by whom, their experience and the measures to ensure the integrity of the sampling and testing.

You must ensure that the sampler has NATA accreditation for the specified sampling methods (where applicable) and understands the requirements for independent random and unbiased sampling. If NATA accreditation does not apply for any sampling or testing method, you must ensure that the sampler and/or tester is suitably trained and competent.

Should a Project Testing laboratory need to subcontract testing then, prior to sampling and testing commencing, you must submit a proposal for the RMS Representative and the Project Verifier's approval showing the tests to be performed, by whom, their experience and the measures to ensure the integrity of the sampling and testing.

1.2.2.2 Inspection and Test Plans

Provide the Project Testing laboratory with all information, including relevant parts of the deed, Specifications, Inspection and Test Plans (ITP’s), and ensure that laboratory performs sampling and testing in accordance with the deed.

Project Testing laboratories that provide on-site testing services must independently review your Inspection and Test Plans (and/or your subcontractors) to confirm that:

(a) all conformity tests are identified, and
(b) sampling and test methods, acceptance criteria and frequency of testing conform to the deed and Specifications.

Any discrepancies must be resolved between you and the Project Testing laboratory and amended Inspection and Test Plans issued, where appropriate. The Project Testing laboratory, for each staged submission of ITP’s (submitted in accordance with RMS Q Clause 4.2.5), must supply to the RMS Representative and the Project Verifier, prior to commencement of any sampling and testing, a written report describing the outcome of this review.

1.2.2.3 Selection of Sampling Locations

You must define the Lots for sampling, and the Project Testing laboratory must select the sampling locations in accordance with the deed and Specifications. Sampling must not be restricted to locations dimensioned or otherwise defined for setting out your Work in the Design Documentation drawings or Specification, but must be undertaken in a random or unbiased manner at any location within your Work.

1.2.2.4 Test Certificates and Declarations

The test report for each Lot (or sub-Lot) must include the following details:

(a) Identification of work and materials with the relevant Lot number;

(b) Where sampling is performed by personnel other than from the laboratory undertaking the testing,
   (i) declaration from the sampler that the sampling was carried out in accordance with Annexure Q/L and the specified sampling methods. Detail all samples taken as part of the Lot.
(ii) declaration (in a format acceptable to the RMS Representative and the Project Verifier) by an officer having NATA signatory approval for the testing performed, that the test results, and statistical analysis where applicable, conform with the specified criteria. This declaration must reference your Inspection and Test Plan and the sampler’s declaration.

(c) Where sampling is performed by personnel from the laboratory undertaking the testing, declaration (in a format agreed by the RMS Representative and the Project Verifier) by an officer having NATA signatory approval for the sampling and testing performed, that the sampling was carried out in accordance with Annexure Q/L and the specified sampling methods, test results, and statistical analysis (where applicable), conform to the Inspection and Test Plan. This declaration must reference and indicate the issue number or date of the Inspection and Test Plan.

(d) Declaration that no samples have been abandoned or untested, or details of any samples that have been abandoned or untested for any reason.

L2.2.5 Availability of Sampling and Testing Records

Sampling and testing records shown in Annexure Q/F to be held on site must be stored in a room readily accessible to the RMS Representative and the Project Verifier with facilities for inspection of the records. Access must not be limited by the Laboratory’s other management activities.

The laboratory, on request, must independently provide to the RMS Representative and the Project Verifier, concurrently with submission to you, the test certificates and declarations in Clause L2.1, including preliminary results forwarded to you.

The RMS Representative must be given physical access to sites and personnel in conjunction with or through you. You must nominate a member of the Project Testing laboratory team to be the authorised contact person for communications with the RMS Representative and the Project Verifier in sampling and testing matters. This person must be fully conversant with the relevant parts of the specifications, specified test methods, the test carried out and testing records and must promptly provide, when requested, information on testing and access to, or copies of, testing records including worksheets to the RMS Representative and the Project Verifier.

L2.3 Protection of Sampled Work

Samples removed from your Work must be replaced, unless otherwise specified, with similar material placed and finished in accordance with the relevant specification requirements, within 7 days of sampling and prior to the use, deterioration, contamination or covering up of the sampled work.

L3 Statistical Techniques

Use statistical techniques in accordance with the following sub-Clauses where required in the Specifications. By agreement with the RMS Representative, areas that are not generally rectangular may be notionally rearranged to suit the method of determining sampling locations in RMS Q Clause L3.1.
L3.1 Sampling and Testing

The number of samples per Lot (n) must be not less than:

<table>
<thead>
<tr>
<th>Specified Relative Compaction (%)</th>
<th>Minimum Testing Frequency for Lot Area of:</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>&gt; 5000 m²</td>
</tr>
<tr>
<td>≤ 90.0</td>
<td>1 per 3000 m²</td>
</tr>
<tr>
<td>&gt; 90.0 ≤ 95.0</td>
<td>1 per 2000 m²</td>
</tr>
<tr>
<td>&gt; 95.0 ≤ 98.0</td>
<td>1 per 2000 m² (min. 6)</td>
</tr>
<tr>
<td>&gt; 98.0 ≤ 100.0</td>
<td>1 per 2000 m² (min. 6)</td>
</tr>
<tr>
<td>&gt; 100.0</td>
<td>1 per 1000 m² (min. 10)</td>
</tr>
</tbody>
</table>

NOTES:
1. Where the sampler/tester can assure that the work is homogeneous and has been carried out within the same day under homogeneous conditions, and:
   (a) where the minimum specified compaction is below 100.0%, work in separate areas, up to a total area of 1000 m², may be considered as one Lot; or
   (b) where the minimum specified compaction is below 98.0%, layers may be covered before testing and may be considered as one Lot, subject to the following:
   
   - Sum Total Area of layers: < 100 m², 101-500 m², 501-1000 m²
   - Maximum number of layers: 5, 3, 2
   - Maximum thickness of Lot: 600 mm, 600 mm, 600 mm
   - Minimum number of Tests: 1, 2, 3

   The tests must be evenly distributed throughout the layers and areas of the Lot.

2. Lots less than 2 m wide must not be longer than 150 m.

3. Except as stated in 1(b) above or specifically allowed by the relevant specification, the Lot will only be one layer.

Sampling locations must be determined by the sampling personnel in a random or unbiased manner (refer RMS Q Clause L2.2.3) as follows, unless directed otherwise by the RMS Representative:

(a) Representing the Lot as a rectangle, subdivide the Lot lengthwise into equal-area sub-Lots in accordance with the number of samples selected (n);
(b) Establish six equally spaced grid lines within the Lot, as illustrated in Figure Q/L.1;
(c) Where the width of Lot is between 0.5 m and 2.5 m, the number of grid lines may be reduced such that the distance between adjacent grid lines (equally spaced) does not exceed 400 mm;
(d) Where the Lot is less than 500 mm wide, the offset locations must be randomly selected;
(e) Determine the order of sampling of the six lines by selecting a six digit number from Table Q/L.1. A starting point on the table (e.g. 1st number, block 6D (= 415236)) will be advised by the RMS Representative prior to the commencement of testing. The numbers are to be used...
sequentially down the Table until further notice from the RMS Representative, starting at the point advised by the RMS Representative, and selecting a new number for each Lot tested;

(f) Where there are less than 6 grid lines in the Lot, delete from the random number selected from Table Q/L.1, the numerals that exceed the number of gridlines in the Lot (e.g. in the above example, where there are only 4 grid lines, the sequence is 4123);

(g) If for any reason the starting point has not been advised then it must be the first number in the block determined, by the following method, from the date on which sampling is first undertaken:

<table>
<thead>
<tr>
<th>Select column:</th>
<th>A</th>
<th>B</th>
<th>C</th>
<th>D</th>
</tr>
</thead>
<tbody>
<tr>
<td>For:</td>
<td></td>
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<tr>
<td>January</td>
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<td>September</td>
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Day: select a row on the basis of:

1st, 11th, 21st, 31st = Row 1;
2nd, 12th, 22nd = Row 2; etc.

(h) For each block in Table Q/L.1, use the Fraction R at the right of the relevant random number. Length coordinate for sample location in sub-Lot 1 = RL/n;

(i) Record the Lot number on Table Q/L.1 to the right of the applicable random number and indicate the date of the sampling on the Table;

(j) For the sample location in the next sub-Lot:

Add L/n to the previous length coordinate.

Go to the next line as indicated by the six-digit number
(e.g. if the number is 415236 the first line tested is 4, followed by 1, 5, 3, 2 and 6 and the sample locations are as shown in Figure Q/L.1);

(k) If the Lot requires more than six sampling locations, repeat the sequence using the same Grid Line Sequence and Fraction R to provide as many additional locations as are required.
Sublot 1
L/n

Sublot 2
L/n

Sample Location

L = length of lot

200 mm

Figure Q/L.1  Sampling Locations for a Rectangular Lot
### Table Q/L.1 – Random Grid Line Sequences (and Random Fraction R)

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<th>Sequence</th>
<th>R</th>
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L3.2 Method for Statistical Calculation for Conformity of Lots

When acceptance criteria specify a maximum and/or minimum characteristic value of attribute (Q), $Q_U$ and/or $Q_L$ must be used to determine $Q$.

The calculation of the characteristic value of attribute ($Q$) for the Lot must be as follows:

(a) Sample Size = 1

$$Q_U = Q_L = \text{Test result}$$

(b) Sample Size = 2

$$Q_U = \text{highest test result}$$

$$Q_L = \text{lowest test result}$$

(c) Sample Size > 2

$$Q_U = \bar{x} + ks$$

$$Q_L = \bar{x} - ks$$

where $\bar{x} =$ arithmetic mean of attribute test results for all sub-Lots

$s =$ standard deviation of sub-Lot attribute test results

$$s = \sqrt{\frac{\sum_{i=1}^{n} (x_i - \bar{x})^2}{n-1}}$$

$k =$ acceptance constant from Table Q/L.2 (based on 10% producer's risk)

Table Q/L.2 – Acceptance Constant $k$

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<th>7</th>
<th>8</th>
<th>9</th>
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<th>15 - 19</th>
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A Lot achieves conformity if:

$$Q_U \leq \text{the specified upper limit for characteristic value of the attribute; and}$$

$$Q_L \geq \text{the specified lower limit for characteristic value of the attribute.}$$

If:

$$Q_U \text{ is more than the specified upper limit for characteristic value; or}$$

$$Q_L \text{ is less than the specified upper/lower limit for characteristic value,}$$

and reworking is subsequently undertaken, the complete Lot must be resampled and retested to verify conformity.
ANNEXURE Q/M – REFERENCED DOCUMENTS

Refer to RMS Q Clause 1.2.6.

**RMS Specifications**

| RMS D&C G22   | Work Health and Safety (Construction Works) |
| RMS D&C G36   | Environmental Protection                   |
| RMS D&C G71   | Construction Surveys                       |

**Australian Standards and Handbooks**

| AS 4390.6  | AS 4390:1996, Records management Part 6: Storage (superseded) |
| ISO 9000   | AS/NZS ISO 9000:2006, Quality management systems - Fundamentals and vocabulary |
| ISO 9001   | AS/NZS ISO 9001:2008, Quality management systems - Requirements |
| ISO/IEC 17025| General requirements for the competence of testing and calibration laboratories |