

# Clarification\_ QM 58 Category 4 Quality Requirements KZN Province

## **Catering**

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## Content of this presentation



- 1. Quality Requirement Category 4
- 2.Section A document submission
- 3.Section B document submission
- 4.Complete Section E:Form A

## 1- Quality Requirement - Category 3



#### **SECTION A:**

A.1 QMS Manual or a document that defines and describes the QMS and its **scope** or Quality Method statement based on scope. (**The manual to be aligned to ISO 9001:2015**)

#### Clauses of ISO 9001:2015

- Clause 4 Context of organization
- Clause 5 Leadership
- Clause 6 Planning
- Clause 7 Support
- Clause 8 Operation
- Clause 9 Performance Evaluation
- Clause 10 Improvement

## 1- Quality Requirement - Category 3



#### **SECTION A:**

#### A.2 Quality Policy Approved by top management

 Commitment to: meet customer requirements / expectations; continual improvement; satisfying all applicable requirements including statutory requirements.

#### A.4 Control of documented information

- Describe how documents are approved, distributed, retrieved, reviewed. Control of obsolete and external origin.
- Describe storage(how records are kept), protection, retrieval, retention time and disposal of records

## 1- Quality Requirement - Category 3



#### **SECTION A:**

#### A.6 Documented information for Non-Conformance and Corrective Action

- Describe elimination of detected non-conformance
- Describe process of Corrective Action Prevention of recurrence (ID root cause)

#### A.7 Documented information for Internal audit

 Planned interval, conducted by trained and independent personnel, who will conduct, how will they conduct and intervals. Reporting and correction of audit findings

#### Section B



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#### **B.1 Documented information for defined roles, responsibilities and authorities**

- Submits either of these documents
  - Organogram
  - Job Profile
  - Responsibility Matrix etc.

Please ensure that the Quality Roles / Function is detailed

#### Section B



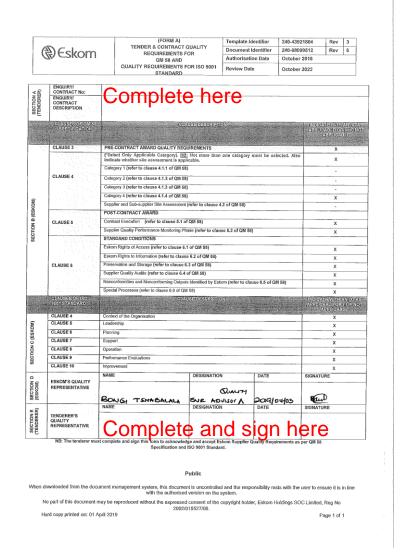
## B.2 <u>Documented information for Control of Externally Provided Processes, Products and Services</u>

- Describe how you ensure that work completed/supplied by external party (Sub-Contractor, Suppliers) compiles with the requirements.
  - Selection
  - Evaluation
  - On going monitoring
  - Inspection
  - Product acceptance /approval / final Inspection
  - Controls that you have in place

## Section E - Form A completion and signing



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#### **Omissions**



#### Section A

- Documents not compliant to ISO 9001:2015 requirements (A1,A2,A4,A6,A7)
- Documents compliant to ISO 9001:2015 but reference made to ISO 9001:2008
- Quality Policy not approved

#### Section B

- B.1 Documented information for defined roles, responsibilities and authorities
  - Quality role not included
- B.2 Documented information for Control of Externally Provided Processes, Products and Services
  - Documentation control process only listed no indication of other process e.g supplier/contractor management

#### Section E

- Form A enquiry number and description omitted
- Not signed by Quality Rep





Thank you