



Qualification Pack

Food Lab Instrumentation Specialist

QP Code: FIC/Q7610

Version: 1.0

NSQF Level: 5

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FIC/Q7610: Food Lab Instrumentation Specialist

Brief Job Description

A Lab Instrumentation Specialist is responsible for writing and executing of validation documents for laboratory instrumentation in compliance with Good Manufacturing Practices (GMP). An individual coordinates with instrument vendors and works with a cross-functional team to deliver all system lifecycle documents for the laboratory. In addition, writing Standard Operating Procedures (SOPs) to support the running of the laboratory and supporting containment verification of the laboratory. An individual also participates in setting up the general running of the laboratory and supporting contaminant verification of the lab. An individual also participates in setting up the general running of the lab- glassware, reagents, consumables, 5S, etc., and ensure the safety, consistency and reliability of chemical testing following the Good Laboratory Practices (GLP)

Personal Attributes

The job requires the individual to have strong technical lab and documentation skills to maintain accurate records in compliance with Good Manufacturing Practices (GMP) expectations. An individual must have good written and verbal communication skills along with effective interpersonal skills. She/he should have knowledge of GLP and Laboratory safety standards. An individual must be focused and demonstrate attention to detail. In addition to managing time efficiently, ability to manage and coordinate multiple projects should be exhibited. An individual must have good hand/eye coordination when handling samples and lab equipment.

Applicable National Occupational Standards (NOS)

Compulsory NOS:

1. [FIC/N7629: Planning, Organizing, and Setting up of Food Analysis Laboratory](#)
2. [FIC/N7630: Carry out Physical, Chemical, and Instrumental Analysis](#)
3. [FIC/N7631: Carry out maintenance and decommissioning of equipment](#)
4. [FIC/N9904: Ensure food safety at the workplace](#)
5. [DGT/VSQ/N0101: Employability Skills \(30 Hours\)](#)

Qualification Pack (QP) Parameters

| | |
|-------------------|-----------------|
| Sector | Food Processing |
| Sub-Sector | Generic |

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|---|--|
| Occupation | Quality Analysis/ Assurance |
| Country | India |
| NSQF Level | 5 |
| Credits | 18 |
| Aligned to NCO/ISCO/ISIC Code | NCO-2015/2113.0500 |
| Minimum Educational Qualification & Experience | <p>B.Sc ((Chemistry, Physics, or related Life Science) or Diploma in a Science related program or equivalent experience or training)</p> <p>OR</p> <p>Completed 1st year of UG (UG Certificate) (3year/4years UG) with 1-2 Years of experience relevant experience</p> <p>OR</p> <p>12th grade pass with 2 year NTC/ CITS/NAC (any combination of NTC/NAC/CITS or equivalent)</p> <p>OR</p> <p>12th grade pass with 1 year NTC/ NAC with 1-2 Years of experience relevant experience</p> <p>OR</p> <p>Completed 3 year diploma after 10th with 1-2 Years of experience relevant experience</p> <p>OR</p> <p>Previous relevant Qualification of NSQF Level (4.5) with 1-2 Years of experience 1.5 years of Experience</p> <p>OR</p> <p>Previous relevant Qualification of NSQF Level (4) with 2-3 Years of experience of relevant experience</p> |
| Minimum Level of Education for Training in School | |
| Pre-Requisite License or Training | NA |
| Minimum Job Entry Age | 20 Years |
| Last Reviewed On | NA |
| Next Review Date | 23/06/2026 |
| NSQC Approval Date | 23/06/2023 |
| Version | 1.0 |
| Reference code on NQR | QG-05-AG-00577-2023-V1-FICSI |
| NQR Version | 1.0 |

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FIC/N7629: Planning, Organizing, and Setting up of Food Analysis Laboratory

Description

This OS is about planning and setting up the Food analysis laboratory considering ISO guidelines and safety guidelines.

Scope

The scope covers the following :

- Analyze requirements for setting up a laboratory
- Comply with measures to ensure Laboratory Safety
- Maintenance of equipment
- Calibration of instruments/equipment

Elements and Performance Criteria

Analyze requirement for setting up a laboratory

To be competent, the user/individual on the job must be able to:

- PC1.** • ensure laboratories are design to meet both the generic and specialized activities
 - Generic activities - include wet chemistry, which requires extensive fixed benches with tprovision of water, power, sinks, cupboards, fume cupboards, reagent shelves, glassware cleaning and storage.
 - Specialized activities - include specialised rooms are required for clean air work or for work on substances, which need special care for reasons of safety (e.g., working with radioactive materials or storage or work
- PC2.** arrange dust-free facilities, both from environmental sources or from other samples, which is likely to be missed by normal quality control checks
- PC3.** ensure ventilation intakes and fume cupboard exhaust must be placed carefully so as to avoid re-circulation of laboratory air and associated risk of contamination of test materials and hazard to laboratory staff
- PC4.** manage that work surfaces and floor shall be made of impervious, smooth, easy to clean materials
- PC5.** arrange that all slabs and work surfaces are levelled and not inclined in any way
- PC6.** administer that there is 300-lux light intensity at working surfaces other than those required for specified tests
- PC7.** arrange for segregation of trace analysis from highly concentrated formulations and from pure substances used in preparing analytical standards. The segregation must apply to all facilities for washing/ cleaning equipment, washing, and storage of glassware, use of protective clothing etc.
- PC8.** organize the separation of preparation of media and media/ glassware sterilization areas from testing areas
- PC9.** manage entry in laboratory areas for reasons such as security, safety or sensitivity to contamination

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- PC10.** administer that eating, drinking, and smoking should be prohibited in the laboratory
- PC11.** arrange adequate fire safety measures at the laboratory
- PC12.** ensure a closed drainage system for all the discards and proper disposal procedure/system for all microbiological samples
- PC13.** manage proper gas/LPG supply system through pipelines to burners and a proper safety system for the same
- PC14.** ensure the appropriate negative air pressure is present in a lab compared to outside.

Comply measures to ensure Laboratory Safety

To be competent, the user/individual on the job must be able to:

- PC15.** administer that chemicals are not stored with an obscured or missing label, marked with the date of receipt before storing and arranged on the shelf in chemically compatible families, and not alphabetically in plastic or metal containers
- PC16.** arrange separate storage areas for highly toxic chemicals
- PC17.** manage empty and full cylinders storage in separate cages outside the laboratory on the ground floor
- PC18.** administer the supply of piped gas through clearly identifiable metal piping to instrument rooms and other equipment
- PC19.** monitor compressed gas cylinders must be firmly attached to a secure structure by a non-combustible material, such as a metal chain
- PC20.** organize the availability and use of a number and type of safety equipment in well-marked, highly visible, and easily accessible locations in or near all the laboratory rooms in the facility and must be maintained in working conditions
- PC21.**
 - arrange easy accessibility of all safety equipment
 - (Safety Equipment - emergency showers, eyewashes, first-aid kits, and spill kits)
- PC22.** validate electrical apparatus, telephones, thermostats, electrical control panels, or power sockets are not located within 0.5 m of the emergency shower or eyewash or within any area that may be considered as a splash or flood zone
- PC23.** categorize waste by its identity, constituents, and hazards to handle and manage safely
- PC24.** identify and label all chemical and waste containers in their respective sections/laboratories
- PC25.**
 - incorporate a procedure for proper disposal of chemical waste basis the category of chemicals used
 - Category - Chemicals that can be washed down the drains with excess water, such as CaCl₂, MgSO₄, Fine (TLC grade) silica, and alumina, etc.; Chemicals that cannot be washed down the drains, such as chloroform, dichloromethane, methanol, acetonitrile, Cyanides, and azides, etc.
- PC26.** arrange waste bins preferably color-coded and labeled for each laboratory
- PC27.** negotiate a contract with a commercial firm, licensed by the respective State Pollution Control Board, to remove and transport biological wastes to a designated disposal site for incineration
- PC28.** monitor all disposable petri-plates used for inoculation and enumeration of the microorganisms, should be autoclaved (steam sterilized) to inactivate the microorganisms for safety reasons. Once autoclaved, the wastes can be safely disposed off
- PC29.** administer that number of emergency exits must be in accordance with the building laws and codes
- PC30.** define an Emergency Evacuation Plan and route for all buildings floors, and areas, and post instructions in every laboratory section and corridor

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PC31. manage Laboratory Information Card at the entrance door of each laboratory and ensure Emergency exits shall be marked accordingly

Maintain instrumentation equipment used

To be competent, the user/individual on the job must be able to:

- PC32.** devise and plan all analytical work adequately to indicate the starting and intended finishing point of the particular task together with the strategy for achieving the desired aims, considering the scope of the amendment.
- PC33.**
- organize sampling plans to obtain quantitative or qualitative information, or to determine conformance or non-conformance with a specification
 - Sampling plans - may be random, systematic or sequential
- PC34.** select an appropriate sample or samples, from a larger amount of material for chemical analysis
- PC35.** document measurement uncertainty associated with sub-sampling etc. in test results separately from measurement uncertainty associated with the basic sampling process
- PC36.** monitor that all of the terms used are clearly defined and understandable to others
- PC37.**
- supervise that equipment used in laboratories should be of a specification, sufficient for an intended purpose, and kept in a state of maintenance and calibration consistent with its use
 - Equipment - General service equipment not used for making measurements or with minimal influence on measurements (e.g., hotplates, stirrers, nonvolumetric glassware and glassware used for rough volume measurements such as measuring cylinders) and laboratory heating or ventilation systems; Volumetric equip
- PC38.** validate that quality of reagents and other consumable materials must be appropriate for the intended use. Consideration needs to be given to the selection, purchase, reception, and storage of reagents
- PC39.**
- administer that grade of any critical reagent used (including water) should be stated in the method, together with guidance on any particular precautions which should be observed in its preparation, storage, and use
 - Precautions - include toxicity, flammability, stability to heat, air, and light; reactivity to other chemicals; reactivity to particular containers; and other hazards.
- PC40.** monitor reagents and reference materials prepared in the laboratory should be labelled to identify substance, strength, solvent (where not water), any special precautions or hazards, restrictions of use, and date of preparation and/or expiry
- PC41.**
- assess factors that make any analytical measurement result liable to deviate from true value as they influence the result
 - Factors - temperature effects on volumetric equipment, reflection and stray light in spectroscopic instruments, variations in electrical supply voltages, individual analysts' interpretation of specified methods and incomplete extraction recoveries
- PC42.** identify relevant sources of uncertainty and assignment of value to each significant contribution
- PC43.**
- administer the complete sequence of events necessary to achieve the purpose of the analysis in identifying relevant sources of uncertainty
 - Sequence - includes sampling and sub-sampling, sample preparation, extraction, cleanup, concentration or dilution, instrument calibration (including reference material preparation), instrumental analysis, raw data processing, and transcription of the output result

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- PC44.** • validate that the performance characteristics of a method are understood and capable of producing results and demonstrate that method is scientifically sound under the conditions in which it is to be applied.
- Performance characteristics - Selectivity & specificity (Description of the measurand), Measurement range Calibration and traceability, Bias, Linearity, Limit of detection (LOD)/ Limit of quantitation (LOQ), Ruggedness, Precision
- PC45.** administer that standard methods are developed and validated collaboratively by a group of experts and development should include consideration of all of the necessary aspects of validation and related uncertainty

PC1. administer that standard methods are developed and validated collaboratively by a group of experts and development should include consideration of all of the necessary aspects of validation and related uncertainty Conduct test and calibration of instruments / equipment

To be competent, the user/individual on the job must be able to:

- PC46.** work safely at all times, complying with health and safety, environmental and other relevant food and drink regulations, directives, and guidelines
- PC47.** obtain and use the correct issue of company and/or manufacturers' drawings and testing/calibration documentation
- PC48.** follow procedures for the use of tools and equipment to carry out the required tests and calibration
- PC49.** set up and carry out the tests and calibrations using organizational procedures and within agreed timescales
- PC50.** insert any relevant system trip defeats, including fire extinguishant, and emergency shutdown, in accordance with organizational procedures
- PC51.** • isolate instruments as recommended and per standards
- Instruments - including process, electrical, hydraulic, pneumatic, mechanical
- PC52.** check test equipment used is appropriate for the tests being carried out, is within current calibration dates, and is used within its specified range
- PC53.** maintain safe access and working arrangements for testing and calibration area to carry out required activities as per organisational procedures
- PC54.** administer electrostatic (ESD) precautions when handling sensitive components and circuit boards
- PC55.** re-connect and return equipment to service on completion of testing and calibration activities
- PC56.** record results of tests and calibrations in accordance with organisational procedures
- PC57.** review results and carry out further tests, if necessary, dispose of waste items and any spoilt products in a safe and environmentally acceptable manner, and leave the work area in a safe and clean condition in accordance with organisational procedures
- PC58.** maintain log book (digitally and manually) for each instrument /equipment and troubleshoot/adjust minor malfunctions in the instruments
- PC59.** account for any human or instrumental error in the test result
- PC60.** maintain repeatability and reproducibility of an analysis performed by him/her
- PC61.** • ensure precautionary instructions are adhered, to in the vicinity of an instrument
- Instructions - e.g. No mobile phones in the vicinity of NMR

Knowledge and Understanding (KU)

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The individual on the job needs to know and understand:

- KU1.** health and safety and environmental requirements of the area in which the installation activity is to take place, and the responsibility these requirements place on you not to compromise food safety
- KU2.** isolation and lock-off procedure or permit-to-work procedure that applies, including critical control points
- KU3.** specific health and safety food precautions to be applied during the installation procedure, and their effects on others
- KU4.** specific requirements of your customer/client specifications in relationship to the installation activities
- KU5.** responsibility in relationship to Hazard Analysis and Critical Control Points (HACCP), Threat Assessment and Critical Control Points (TACCP), Vulnerability Assessment and Critical Control Points (VACCP) during the installation activities
- KU6.** importance of wearing protective clothing and other appropriate safety equipment (PPE) during the installation process
- KU7.** responsibility in relationship to Hazard Analysis and Critical Control Points (HACCP), Threat Assessment and Critical Control Points (TACCP), Vulnerability Assessment and Critical Control Points (VACCP) during the installation activities
- KU8.** what constitutes a hazardous voltage and how to recognize victims of electric shock
- KU9.** knowledge of waste characteristics and constituents by laboratory personnel who conduct the process, procedure, or experiment
- KU10.** chemical Waste can be in the form of solvents, aqueous solutions, dry powders, and unwanted old chemicals
- KU11.** health and Safety information should be posted on the door of each laboratory indicating accurately the hazards that could be there in the laboratory, personal protection required, and the emergency contacts
- KU12.** never store highly reactive chemicals for more than 6 months
- KU13.** gas cylinders must be transported on purpose-built trolleys within the laboratory
- KU14.** oxygen cylinders, full or empty should not be stored in close proximity to the flammable gases
- KU15.** labs should be organized with the highest hazards (e.g., fume hoods) farthest from the entry door and the least hazardous elements (e.g., write-up stations) closest to the door
- KU16.** write-up desks and benches should be accessible without having to cross in front of fume hoods
- KU17.** all safety equipment, such as emergency showers, eyewashes, first-aid kits, and spill kits should be readily accessible
- KU18.** emergency center in a central location on each floor provides easy access for everyone. This center should have reagent neutralizers, spill kits, first aid kits, etc.
- KU19.** there should be at least one ABC fire extinguisher either inside the lab or in close proximity. Extinguishers should not be covered up or block access
- KU20.** there should be an eyewash unit, provided at least 10 seconds away from any analyst, and should supply a multi-stream cross flow of water at 20-25 degrees C (65- 75 degree Fahrenheit)

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- KU21.** safety showers should never be more than 100 ft. away from the analyst, along a clear and unobstructed path, and should be located in the corridor, clearly visible from the lab exits
- KU22.** that ISO/IEC 17025:1999 requires laboratories to evaluate their measurement uncertainty and to report measurement uncertainty under specific circumstances, for example, where it is relevant to the interpretation of the test result (which is often the case). Thus, statement of measurement uncertainty in test reports should become common practice in the future
- KU23.** elucidation of performance characteristics such as Selectivity & specificity (Description of the measurand), Measurement range Calibration and traceability, Bias, Linearity, Limit of detection (LOD)/ Limit of quantitation (LOQ), Ruggedness, Precision

Generic Skills (GS)

User/individual on the job needs to know how to:

- GS1.** read and interpret information such as product labels, safety data sheets, instructions, procedures, and specifications to determine activities in the calibration of instruments/equipment
- GS2.** write and document using appropriate terminology and in the specified format
- GS3.** communicate with others effectively
- GS4.** takes responsibility for planning, sequencing, and prioritizing tasks
- GS5.** analyse and report variances promptly

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Assessment Criteria

| Assessment Criteria for Outcomes | Theory Marks | Practical Marks | Project Marks | Viva Marks |
|---|--------------|-----------------|---------------|------------|
| <i>Analyze requirement for setting up a laboratory</i> | 7 | 7 | - | 7 |
| <p>PC1.</p> <ul style="list-style-type: none"> ensure laboratories are design to meet both the generic and specialized activities Generic activities - include wet chemistry, which requires extensive fixed benches with tprovision of water, power, sinks, cupboards, fume cupboards, reagent shelves, glassware cleaning and storage. Specialized activities - include specialised rooms are required for clean air work or for work on substances, which need special care for reasons of safety (e.g., working with radioactive materials or storage or work | 0.5 | 0.5 | - | 0.5 |
| <p>PC2. arrange dust-free facilities, both from environmental sources or from other samples, which is likely to be missed by normal quality control checks</p> | 0.5 | 0.5 | - | 0.5 |
| <p>PC3. ensure ventilation intakes and fume cupboard exhaust must be placed carefully so as to avoid re-circulation of laboratory air and associated risk of contamination of test materials and hazard to laboratory staff</p> | 0.5 | 0.5 | - | 0.5 |
| <p>PC4. manage that work surfaces and floor shall be made of impervious, smooth, easy to clean materials</p> | 0.5 | 0.5 | - | 0.5 |
| <p>PC5. arrange that all slabs and work surfaces are levelled and not inclined in any way</p> | 0.5 | 0.5 | - | 0.5 |
| <p>PC6. administer that there is 300-lux light intensity at working surfaces other than those required for specified tests</p> | 0.5 | 0.5 | - | 0.5 |
| <p>PC7. arrange for segregation of trace analysis from highly concentrated formulations and from pure substances used in preparing analytical standards. The segregation must apply to all facilities for washing/ cleaning equipment, washing, and storage of glassware, use of protective clothing etc.</p> | 0.5 | 0.5 | - | 0.5 |
| <p>PC8. organize the separation of preparation of media and media/ glassware sterilization areas from testing areas</p> | 0.5 | 0.5 | - | 0.5 |

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| Assessment Criteria for Outcomes | Theory Marks | Practical Marks | Project Marks | Viva Marks |
|--|--------------|-----------------|---------------|------------|
| PC9. manage entry in laboratory areas for reasons such as security, safety or sensitivity to contamination | 0.5 | 0.5 | - | 0.5 |
| PC10. administer that eating, drinking, and smoking should be prohibited in the laboratory | 0.5 | 0.5 | - | 0.5 |
| PC11. arrange adequate fire safety measures at the laboratory | 0.5 | 0.5 | - | 0.5 |
| PC12. ensure a closed drainage system for all the discards and proper disposal procedure/system for all microbiological samples | 0.5 | 0.5 | - | 0.5 |
| PC13. manage proper gas/LPG supply system through pipelines to burners and a proper safety system for the same | 0.5 | 0.5 | - | 0.5 |
| PC14. ensure the appropriate negative air pressure is present in a lab compared to outside. | 0.5 | 0.5 | - | 0.5 |
| <i>Comply measures to ensure Laboratory Safety</i> | 16 | 8.5 | - | 9 |
| PC15. administer that chemicals are not stored with an obscured or missing label, marked with the date of receipt before storing and arranged on the shelf in chemically compatible families, and not alphabetically in plastic or metal containers | 1 | 0.5 | - | 0.5 |
| PC16. arrange separate storage areas for highly toxic chemicals | 1 | 0.5 | - | 0.5 |
| PC17. manage empty and full cylinders storage in separate cages outside the laboratory on the ground floor | 0.5 | 0.5 | - | 0.5 |
| PC18. administer the supply of piped gas through clearly identifiable metal piping to instrument rooms and other equipment | 0.5 | 0.5 | - | 0.5 |
| PC19. monitor compressed gas cylinders must be firmly attached to a secure structure by a non-combustible material, such as a metal chain | 1 | 0.5 | - | 0.5 |

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| Assessment Criteria for Outcomes | Theory Marks | Practical Marks | Project Marks | Viva Marks |
|---|--------------|-----------------|---------------|------------|
| PC20. organize the availability and use of a number and type of safety equipment in well-marked, highly visible, and easily accessible locations in or near all the laboratory rooms in the facility and must be maintained in working conditions | 1 | 0.5 | - | 0.5 |
| PC21. <ul style="list-style-type: none"> • arrange easy accessibility of all safety equipment • (Safety Equipment - emergency showers, eyewashes, first-aid kits, and spill kits) | 1 | 0.5 | - | 0.5 |
| PC22. validate electrical apparatus, telephones, thermostats, electrical control panels, or power sockets are not located within 0.5 m of the emergency shower or eyewash or within any area that may be considered as a splash or flood zone | 1 | 0.5 | - | 0.5 |
| PC23. categorize waste by its identity, constituents, and hazards to handle and manage safely | 1 | 0.5 | - | 0.5 |
| PC24. identify and label all chemical and waste containers in their respective sections/laboratories | 1 | 0.5 | - | 0.5 |
| PC25. <ul style="list-style-type: none"> • incorporate a procedure for proper disposal of chemical waste basis the category of chemicals used • Category - Chemicals that can be washed down the drains with excess water, such as CaCl₂, MgSO₄, Fine (TLC grade) silica, and alumina, etc.; Chemicals that cannot be washed down the drains, such as chloroform, dichloromethane, methanol, acetonitrile, Cyanides, and azides, etc. | 1 | 0.5 | - | 0.5 |
| PC26. arrange waste bins preferably color-coded and labeled for each laboratory | 1 | 0.5 | - | 0.5 |
| PC27. negotiate a contract with a commercial firm, licensed by the respective State Pollution Control Board, to remove and transport biological wastes to a designated disposal site for incineration | 1 | 0.5 | - | 0.5 |
| PC28. monitor all disposable petri-plates used for inoculation and enumeration of the microorganisms, should be autoclaved (steam sterilized) to inactivate the microorganisms for safety reasons. Once autoclaved, the wastes can be safely disposed off | 1 | 0.5 | - | 1 |

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| Assessment Criteria for Outcomes | Theory Marks | Practical Marks | Project Marks | Viva Marks |
|--|--------------|-----------------|---------------|------------|
| PC29. administer that number of emergency exits must be in accordance with the building laws and codes | 1 | 0.5 | - | 0.5 |
| PC30. define an Emergency Evacuation Plan and route for all buildings floors, and areas, and post instructions in every laboratory section and corridor | 1 | 0.5 | - | 0.5 |
| PC31. manage Laboratory Information Card at the entrance door of each laboratory and ensure Emergency exits shall be marked accordingly | 1 | 0.5 | - | 0.5 |
| <i>Maintain instrumentation equipment used</i> | 7 | 7.5 | - | 7 |
| PC32. devise and plan all analytical work adequately to indicate the starting and intended finishing point of the particular task together with the strategy for achieving the desired aims, considering the scope of the amendment. | 0.5 | 1 | - | 0.5 |
| PC33. <ul style="list-style-type: none"> • organize sampling plans to obtain quantitative or qualitative information, or to determine conformance or non-conformance with a specification • Sampling plans - may be random, systematic or sequential | 0.5 | 0.5 | - | 0.5 |
| PC34. select an appropriate sample or samples, from a larger amount of material for chemical analysis | 0.5 | 0.5 | - | 0.5 |
| PC35. document measurement uncertainty associated with sub-sampling etc. in test results separately from measurement uncertainty associated with the basic sampling process | 0.5 | 0.5 | - | 0.5 |
| PC36. monitor that all of the terms used are clearly defined and understandable to others | 0.5 | 0.5 | - | 0.5 |

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| Assessment Criteria for Outcomes | Theory Marks | Practical Marks | Project Marks | Viva Marks |
|--|--------------|-----------------|---------------|------------|
| <p>PC37.</p> <ul style="list-style-type: none"> supervise that equipment used in laboratories should be of a specification, sufficient for an intended purpose, and kept in a state of maintenance and calibration consistent with its use Equipment - General service equipment not used for making measurements or with minimal influence on measurements (e.g., hotplates, stirrers, nonvolumetric glassware and glassware used for rough volume measurements such as measuring cylinders) and laboratory heating or ventilation systems; Volumetric equip | 0.5 | 0.5 | - | 0.5 |
| <p>PC38. validate that quality of reagents and other consumable materials must be appropriate for the intended use. Consideration needs to be given to the selection, purchase, reception, and storage of reagents</p> | 0.5 | 0.5 | - | 0.5 |
| <p>PC39.</p> <ul style="list-style-type: none"> administer that grade of any critical reagent used (including water) should be stated in the method, together with guidance on any particular precautions which should be observed in its preparation, storage, and use Precautions - include toxicity, flammability, stability to heat, air, and light; reactivity to other chemicals; reactivity to particular containers; and other hazards. | 0.5 | 0.5 | - | 0.5 |
| <p>PC40. monitor reagents and reference materials prepared in the laboratory should be labelled to identify substance, strength, solvent (where not water), any special precautions or hazards, restrictions of use, and date of preparation and/or expiry</p> | 0.5 | 0.5 | - | 0.5 |
| <p>PC41.</p> <ul style="list-style-type: none"> assess factors that make any analytical measurement result liable to deviate from true value as they influence the result Factors - temperature effects on volumetric equipment, reflection and stray light in spectroscopic instruments, variations in electrical supply voltages, individual analysts' interpretation of specified methods and incomplete extraction recoveries | 0.5 | 0.5 | - | 0.5 |
| <p>PC42. identify relevant sources of uncertainty and assignment of value to each significant contribution</p> | 0.5 | 0.5 | - | 0.5 |

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| Assessment Criteria for Outcomes | Theory Marks | Practical Marks | Project Marks | Viva Marks |
|---|--------------|-----------------|---------------|------------|
| PC43. • administer the complete sequence of events necessary to achieve the purpose of the analysis in identifying relevant sources of uncertainty • Sequence - includes sampling and sub-sampling, sample preparation, extraction, cleanup, concentration or dilution, instrument calibration (including reference material preparation), instrumental analysis, raw data processing, and transcription of the output result | 0.5 | 0.5 | - | 0.5 |
| PC44. • validate that the performance characteristics of a method are understood and capable of producing results and demonstrate that method is scientifically sound under the conditions in which it is to be applied. • Performance characteristics - Selectivity & specificity (Description of the measurand), Measurement range Calibration and traceability, Bias, Linearity, Limit of detection (LOD)/ Limit of quantitation (LOQ), Ruggedness, Precision | 0.5 | 0.5 | - | 0.5 |
| PC45. administer that standard methods are developed and validated collaboratively by a group of experts and development should include consideration of all of the necessary aspects of validation and related uncertainty | 0.5 | 0.5 | - | 0.5 |
| <i>PC1. administer that standard methods are developed and validated collaboratively by a group of experts and development should include consideration of all of the necessary aspects of validation and related uncertainty Conduct test and calibration of instruments / equipment</i> | 8 | 8 | - | 8 |
| PC46. work safely at all times, complying with health and safety, environmental and other relevant food and drink regulations, directives, and guidelines | 0.5 | 0.5 | - | 0.5 |
| PC47. obtain and use the correct issue of company and/or manufacturers' drawings and testing/calibration documentation | 0.5 | 0.5 | - | 0.5 |
| PC48. follow procedures for the use of tools and equipment to carry out the required tests and calibration | 0.5 | 0.5 | - | 0.5 |

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| Assessment Criteria for Outcomes | Theory Marks | Practical Marks | Project Marks | Viva Marks |
|---|--------------|-----------------|---------------|------------|
| PC49. set up and carry out the tests and calibrations using organizational procedures and within agreed timescales | 0.5 | 0.5 | - | 0.5 |
| PC50. insert any relevant system trip defeats, including fire extinguishant, and emergency shutdown, in accordance with organizational procedures | 0.5 | 0.5 | - | 0.5 |
| PC51. <ul style="list-style-type: none"> isolate instruments as recommended and per standards Instruments - including process, electrical, hydraulic, pneumatic, mechanical | 0.5 | 0.5 | - | 0.5 |
| PC52. check test equipment used is appropriate for the tests being carried out, is within current calibration dates, and is used within its specified range | 0.5 | 0.5 | - | 0.5 |
| PC53. maintain safe access and working arrangements for testing and calibration area to carry out required activities as per organisational procedures | 0.5 | 0.5 | - | 0.5 |
| PC54. administer electrostatic (ESD) precautions when handling sensitive components and circuit boards | 0.5 | 0.5 | - | 0.5 |
| PC55. re-connect and return equipment to service on completion of testing and calibration activities | 0.5 | 0.5 | - | 0.5 |
| PC56. record results of tests and calibrations in accordance with organisational procedures | 0.5 | 0.5 | - | 0.5 |
| PC57. review results and carry out further tests, if necessary, dispose of waste items and any spoilt products in a safe and environmentally acceptable manner, and leave the work area in a safe and clean condition in accordance with organisational procedures | 0.5 | 0.5 | - | 0.5 |
| PC58. maintain log book (digitally and manually) for each instrument /equipment and troubleshoot/adjust minor malfunctions in the instruments | 0.5 | 0.5 | - | 0.5 |

Qualification Pack

| Assessment Criteria for Outcomes | Theory Marks | Practical Marks | Project Marks | Viva Marks |
|---|--------------|-----------------|---------------|------------|
| PC59. account for any human or instrumental error in the test result | 0.5 | 0.5 | - | 0.5 |
| PC60. maintain repeatability and reproducibility of an analysis performed by him/her | 0.5 | 0.5 | - | 0.5 |
| PC61. <ul style="list-style-type: none"> • ensure precautionary instructions are adhered, to in the vicinity of an instrument • Instructions - e.g. No mobile phones in the vicinity of NMR | 0.5 | 0.5 | - | 0.5 |
| NOS Total | 38 | 31 | - | 31 |

Qualification Pack

National Occupational Standards (NOS) Parameters

| | |
|----------------------------|---|
| NOS Code | FIC/N7629 |
| NOS Name | Planning, Organizing, and Setting up of Food Analysis Laboratory |
| Sector | Food Processing |
| Sub-Sector | Fruits and Vegetables, Food Grain Milling (Including oil seeds), Dairy Products, Meat and Poultry, Fish and Seafood, Bread and Bakery, Alcoholic Beverages, Aerated Water/Soft Drinks |
| Occupation | Quality Analysis/ Assurance |
| NSQF Level | 5 |
| Credits | 5 |
| Version | 1.0 |
| Last Reviewed Date | 23/06/2023 |
| Next Review Date | 23/06/2026 |
| NSQC Clearance Date | 23/06/2023 |

Qualification Pack

FIC/N7630: Carry out Physical, Chemical, and Instrumental Analysis

Description

This OS is about the activities carried out for physical, chemical and Instrumental analysis

Scope

The scope covers the following :

- Carry out Sampling and Sample Preparation
- Perform analysis through classical analytical techniques
- Carry out quantitative and qualitative analysis
- Carry out reporting and documentation

Elements and Performance Criteria

Carry out Sampling and Sample Preparation

To be competent, the user/individual on the job must be able to:

- PC1.** apply specific methods as per national or sectoral standards for sampling as appropriate
- PC2.** enforce sampling through experience or adapt methods from similar applications or be treated as heterogeneous
- PC3.** carry out sampling stages with an understanding of overall context of analysis
- PC4.** administer that large laboratory sample is reduced test portion is homogeneous, thus reducing the particle size by grinding or milling. If the laboratory sample is large, it may be necessary to subdivide it prior to grinding or milling
- PC5.**
- subdivide sample by following different mechanisms
 - Mechanisms - coning and quartering, riffing, or by means of a rotating sample divider or a centrifugal divider
- PC6.**
- take care of samples by avoiding cross-contamination, by ensuring equipment does not contaminate the sample and the composition of the sample is not altered during grinding
 - Equipment - metals, etc.
- PC7.**
- analyze problem necessitating sample extraction and design the sampling strategy considering the nature of problem
 - Nature of problem - average analyte concentration in the material is required; analyte profile across the material is required; material is suspected of contamination by a particular analyte; contaminant is heterogeneously distributed (occurs in hot spots) in the material; there may be other, non-analytical factors to consider, including the nature of the area under examination
- PC8.** administer the use and value of the rest of the original material once a sample has been removed for analysis.
- PC9.** maintain record of procedures followed in order that sampling process may be repeated exactly irrespective of strategy used for sampling

Perform analysis through classical analytical techniques

To be competent, the user/individual on the job must be able to:

- PC10.** identify whether the analyte is either organic or inorganic prior to qualitative analysis of any given compound

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- PC11.** perform classical qualitative analysis by adding one or a series of chemical reagents to the analyte
- PC12.** draw inferences about the identity of analyte by observing the chemical reactions and their products
- PC13.** administer the result of the assay between added chemical reagents and functional groups of organic molecules
- PC14.**
- identify molecules by following measurements in addition to chemical reactions between chemical reagents and functional groups
 - Measurements - including those of boiling points, melting points, and densities, etc.
- PC15.** examine during gravimetric analysis an excess of added reagent reacts with the analyte to form a precipitate
- PC16.** handle precipitate by filtering, drying, and weighing as its mass is used to calculate the concentration or amount of assayed substance in the analyte
- PC17.** examine during titrimetric (Volumetric) analysis, the reagent (the titrant) is added gradually or stepwise to the analyte from a buret
- PC18.**
- administer equivalence point of the titration, typically observed as a colour change
 - Point of titration - the point at which the quantities of the two reacting species are equivalent
- PC19.** supervise if no spontaneous colour change occurs during the titration, a small amount of a chemical indicator is added to the analyte prior to the titration
- PC20.** calculate the amount or concentration of the analyte by accessing the end point with the concentration of the titrant

Carry out quantitative and qualitative analysis

To be competent, the user/individual on the job must be able to:

- PC21.**
- instill the sample manually into carrier gas with a syringe, via a sample loop and analytical valve which are in-line with the carrier stream
 - Carrier Gas - nitrogen, helium, argon, and hydrogen or air
- PC22.** ensure in automated instruments, the carrier gas is switched in line with the sample loop for a precise, pre-determined period of time, injecting the sample onto the column administer the cycle is repeated continuously in the process Gas Chromatography (GC)/ Mass spectrometry (MS) analysis
- PC23.** mount columns in an oven with precise temperature and carrier gas flow control
- PC24.** monitor that under tightly controlled conditions, analysis is repeated; the same gas component will exit the column (elute) with the same timing as the previous analysis
- PC25.** regulate separated gasses leave (or elute from) the column(s), and pass through a detector which, in turn, responds with an output signal
- PC26.** observe that signal generates the characteristic GC peaks in a chromatogram and peaks are proportional in area to the concentration of gases of interest
- PC27.** instill a medium into which the target analytes can be preferentially partitioned followed by the separation of the analyte-containing phase from the remainder of the sample
- PC28.** administer sample is heated, have its pH adjusted, or be otherwise treated to make analytes more accessible to the partitioning medium

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- PC29.** • place solid analytes in solution for chromatographic separation irrespective of the extraction process
- Solution - additional pretreatments, such as liquid/liquid extraction (LLE) or solid-phase extraction (SPE), to remove substances that might interfere with the chromatographic separation and the detection of the target analytes
 - Extraction Process - Liquid extraction, Ultrasonic simulation, Steam distillation
- PC30.** employ Liquid-liquid extraction, on- and offline solid-phase extraction, and gel permeation chromatography (GPC) in the partitioning of target analytes in liquid samples or liquid extracts of solid samples
- PC31.** filter turbid or otherwise opaque liquids to remove particulates prior to extraction
- PC32.** install guard columns at the head of the analytical column to prevent contamination of the stationary phase by the sample matrix
- PC33.** ensure the guard column is connected to the precolumn by means of a transfer valve
- PC34.** administer automated sampling system instead of manual injectors to get highly reproducible injection volume, low carryover, and increased sample throughput
- PC35.** assemble HPLC pumps to deliver pulse-free flow with high precision over a wide flow rate range with low dead volume
- PC36.** maintain a limit of detection (LOD) and limit of quantification (LOQ) as two to three and 10 to 20 times the noise level. LOD for food additives can be as low as 100 parts per trillion
- PC37.** • administer other important detection parameters are appropriate, as per defined standards
- Parameters - dynamic range, calibration linearity, chromatographic selectivity, and qualitative information
- PC38.** automate derivatization and integrate online within the analysis to improve sensitivity and/or selectivity of an optical detector (UV, visible, or fluorescence) when analytes lack chromophores or an adequate optical response
- PC39.** administer to install of a lamp in Atomic Absorption (AA) unit for the analyte
- PC40.** regulate by nebulizing liquid sample and spray into an atomizer, or atom source
- PC41.** observe that the heat source dries the sample and atomizes it, breaking it down into individual elements. The light at a certain wavelength is passed through the atomized sample, where it is absorbed by elements of interest, and that energy absorption is measured
- PC42.** review in flame AA, the atomizer or atom source is in the form of an air/acetylene or nitrous oxide/acetylene flame
- PC43.** coordinate to introduce the sample as an aerosol into the flame, which is aligned so that the light beam passes through the flame, where the light is absorbed, measured, and reported by the instrument
- PC44.** facilitate Graphite Furnace Atomic Absorption (GFAA) by introducing the sample directly into a graphite tube, which is heated to remove liquid solvent and to atomize the remaining sample
- PC45.** observe that sample introduced to the tube is atomized, and the atoms are retained within the tube for an extended period
- PC46.** check light is passed through this tube and absorbance is measured and as a result of this improved analyte control sensitivity and detection limits are significantly improved
- PC47.** administer in Induction Coupled Plasma Mass Spectrometry (ICP-MS), the sample is atomized and ionized by the plasma source

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- PC48.** observe charged ions are passed into a mass spectrometer, where ions are split according to their mass before reaching the detector, which in turn identifies and quantifies the elements present in a sample based on their mass
- PC49.** facilitate significant improvement in sensitivity by following the ICP-MS approach
- Carry out reporting and documentation*
- To be competent, the user/individual on the job must be able to:
- PC50.** review and update the test methods and procedures as per the schedule
- PC51.** ensure the updation of logbooks, worksheets, calibration records, parameters of columns, reagents, volumetric solution, and working standards
- PC52.** account for defects/problems/incidents/quality issues/test results as applicable as per SOP
- PC53.** prepare analytical and quality control reports for detailed findings and recommendations as per SOPs
- PC54.** prepare reports of validations, deviations, and incidents to the production and quality assurance team in compliance with GLP and GMP
- PC55.** facilitate to document testing results and analysis accurately
- PC56.** maintain all original and controlled document files and quality records in a timely and accurate manner and made them available for inspection and auditory purposes

Knowledge and Understanding (KU)

The individual on the job needs to know and understand:

- KU1.** selection of an appropriate sample or samples, from a larger amount of material
- KU2.** procedures followed to select the test portion from the sample (or subsample) and include in-laboratory processing, mixing, reducing, coning & quartering, rifling, and milling & grinding
- KU3.** test portion which is actual material weighed or measured for the analysis
- KU4.** properties of an analyte such as volatility, sensitivity to light, thermal lability, and chemical reactivity are considered in designing sampling strategy and choosing equipment, packaging, and storage conditions
- KU5.** how to prepare a clear record of procedures for the sampling process
- KU6.** sampling procedures followed through inspection by attributes have risk associated with acceptance/rejection of non-conformities is predetermined by the acceptable quality level (AQL) or the limiting quality (LQ)
- KU7.** powerful integration software is used to quantify the peak size
- KU8.** GCs are designed with various detectors based on the analytical requirements, gas composition and required detection limits to help determine the detector used
- KU9.** process GCs typically utilize FID, PID, or TCD, due to their simple reliable designs
- KU10.** columns are tools used to separate the sample into its constituent components
- KU11.** equipment used for sampling, subsampling, sample handling, sample preparation, and sample extraction
- KU12.** significance of gravimetric or volumetric errors during sampling should be considered and any critical equipment calibrated

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- KU13.** poorly considered sampling, especially if destructive, may render the whole consignment valueless or inoperative
- KU14.** selection of the partitioning medium is determined by the nature of the sample matrix and the properties of the analytes
- KU15.** liquid extraction enables highly selective matching of solvent to analyte solubility. Extraction times are short, typically, minutes. It uses relatively small quantities of organic solvents, thereby reducing costs and facilitating disposal
- KU16.** ultrasonic stimulation is often applied in conjunction with the extraction solvent to drive the liquid medium into the interior of the matrix, ensuring intimate and energetic contact between solvent and analyte molecules
- KU17.** steam distillation can be utilized to selectively extract volatile compounds. Extraction times are long, however, and the method has a narrow range of use and is applied offline
- KU18.** light source is a hollow cathode lamp (HCL) or an electrodeless discharge lamp (EDL), with different lamps being used for different analytes (elements of interest)
- KU19.** Atomic Absorption Consumables are flame uses Lamps, Gases & O-rings, furnaces in addition need Graphite tubes Caps & Holders
- KU20.** ICP-MS Consumables are argon gas, glassware - torches, spray chambers, nebulizers and cones - skimmers and sampling
- KU21.** detection limits of atomic absorption and ICP-MS
- KU22.** procedures for reporting any unresolved issues and hazards
- KU23.** procedures for management of Standard Operating Procedures (SOPs), Standard Testing Procedures (STPs), protocols, Equipment documents, method validation protocols & reports
- KU24.** procedures for reporting non-conformance, deviations, validation results
- KU25.** procedure for reporting incidents where standard operating procedures are not followed
- KU26.** statistical concepts and application of statistical tools
- KU27.** guidelines for electronic records and signatures, audit trails, date, and time stamps
- KU28.** operating procedure of analytical instruments and equipment
- KU29.** concepts of organic and analytical chemistry

Generic Skills (GS)

User/individual on the job needs to know how to:

- GS1.** read and interpret information such as product labels, safety data sheets, instructions, procedures, and specifications to determine activities in the calibration of instruments/equipment
- GS2.** write and document using appropriate terminology and in the specified format
- GS3.** communicate with others effectively
- GS4.** takes responsibility for planning, sequencing, and prioritizing tasks
- GS5.** analyse and report variances promptly
- GS6.** check the sample integrity
- GS7.** maintain a laboratory notebook and follow the related rules
- GS8.** should be able to guide and lead sample handlers or other subordinates related to work
- GS9.** takes responsibility for planning, sequencing, and prioritising tasks



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- GS10.** analyse and report variances promptly
- GS11.** check the sample integrity
- GS12.** maintain a laboratory notebook and follow the related rules
- GS13.** should be able to guide and lead sample handlers or other subordinates related to work

Qualification Pack

Assessment Criteria

| Assessment Criteria for Outcomes | Theory Marks | Practical Marks | Project Marks | Viva Marks |
|---|--------------|-----------------|---------------|------------|
| <i>Carry out Sampling and Sample Preparation</i> | 6.5 | 4.5 | - | 4.5 |
| PC1. apply specific methods as per national or sectoral standards for sampling as appropriate | 0.5 | 0.5 | - | 0.5 |
| PC2. enforce sampling through experience or adapt methods from similar applications or be treated as heterogeneous | 0.5 | 0.5 | - | 0.5 |
| PC3. carry out sampling stages with an understanding of overall context of analysis | 1 | 0.5 | - | 0.5 |
| PC4. administer that large laboratory sample is reduced test portion is homogeneous, thus reducing the particle size by grinding or milling. If the laboratory sample is large, it may be necessary to subdivide it prior to grinding or milling | 1 | 0.5 | - | 0.5 |
| PC5. <ul style="list-style-type: none"> subdivide sample by following different mechanisms Mechanisms - coning and quartering, riffing, or by means of a rotating sample divider or a centrifugal divider | 1 | 0.5 | - | 0.5 |
| PC6. <ul style="list-style-type: none"> take care of samples by avoiding cross-contamination, by ensuring equipment does not contaminate the sample and the composition of the sample is not altered during grinding Equipment - metals, etc. | 0.5 | 0.5 | - | 0.5 |
| PC7. <ul style="list-style-type: none"> analyze problem necessitating sample extraction and design the sampling strategy considering the nature of problem Nature of problem - average analyte concentration in the material is required; analyte profile across the material is required; material is suspected of contamination by a particular analyte; contaminant is heterogeneously distributed (occurs in hot spots) in the material; there may be other, non-analytical factors to consider, including the nature of the area under examination | 0.5 | 0.5 | - | 0.5 |
| PC8. administer the use and value of the rest of the original material once a sample has been removed for analysis. | 1 | 0.5 | - | 0.5 |

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| Assessment Criteria for Outcomes | Theory Marks | Practical Marks | Project Marks | Viva Marks |
|---|--------------|-----------------|---------------|------------|
| PC9. maintain record of procedures followed in order that sampling process may be repeated exactly irrespective of strategy used for sampling | 0.5 | 0.5 | - | 0.5 |
| <i>Perform analysis through classical analytical techniques</i> | 10.5 | 5.5 | - | 5.5 |
| PC10. identify whether the analyte is either organic or inorganic prior to qualitative analysis of any given compound | 1 | 0.5 | - | 0.5 |
| PC11. perform classical qualitative analysis by adding one or a series of chemical reagents to the analyte | 1 | 0.5 | - | 0.5 |
| PC12. draw inferences about the identity of analyte by observing the chemical reactions and their products | 1 | 0.5 | - | 0.5 |
| PC13. administer the result of the assay between added chemical reagents and functional groups of organic molecules | 1 | 0.5 | - | 0.5 |
| PC14. <ul style="list-style-type: none"> identify molecules by following measurements in addition to chemical reactions between chemical reagents and functional groups Measurements - including those of boiling points, melting points, and densities, etc. | 1 | 0.5 | - | 0.5 |
| PC15. examine during gravimetric analysis an excess of added reagent reacts with the analyte to form a precipitate | 1 | 0.5 | - | 0.5 |
| PC16. handle precipitate by filtering, drying, and weighing as its mass is used to calculate the concentration or amount of assayed substance in the analyte | 1 | 0.5 | - | 0.5 |
| PC17. examine during titrimetric (Volumetric) analysis, the reagent (the titrant) is added gradually or stepwise to the analyte from a buret | 1 | 0.5 | - | 0.5 |
| PC18. <ul style="list-style-type: none"> administer equivalence point of the titration, typically observed as a colour change Point of titration - the point at which the quantities of the two reacting species are equivalent | 1 | 0.5 | - | 0.5 |

Qualification Pack

| Assessment Criteria for Outcomes | Theory Marks | Practical Marks | Project Marks | Viva Marks |
|---|--------------|-----------------|---------------|-------------|
| PC19. supervise if no spontaneous colour change occurs during the titration, a small amount of a chemical indicator is added to the analyte prior to the titration | 0.5 | 0.5 | - | 0.5 |
| PC20. calculate the amount or concentration of the analyte by accessing the end point with the concentration of the titrant | 1 | 0.5 | - | 0.5 |
| <i>Carry out quantitative and qualitative analysis</i> | 23.5 | 14.5 | - | 14.5 |
| PC21. <ul style="list-style-type: none"> instill the sample manually into carrier gas with a syringe, via a sample loop and analytical valve which are in-line with the carrier stream Carrier Gas - nitrogen, helium, argon, and hydrogen or air | 1 | 0.5 | - | 0.5 |
| PC22. ensure in automated instruments, the carrier gas is switched in line with the sample loop for a precise, pre-determined period of time, injecting the sample onto the column administer the cycle is repeated continuously in the process Gas Chromatography (GC)/ Mass spectrometry (MS) analysis | 1 | 0.5 | - | 0.5 |
| PC23. mount columns in an oven with precise temperature and carrier gas flow control | 1 | 0.5 | - | 0.5 |
| PC24. monitor that under tightly controlled conditions, analysis is repeated; the same gas component will exit the column (elute) with the same timing as the previous analysis | 1 | 0.5 | - | 0.5 |
| PC25. regulate separated gasses leave (or elute from) the column(s), and pass through a detector which, in turn, responds with an output signal | 1 | 0.5 | - | 0.5 |
| PC26. observe that signal generates the characteristic GC peaks in a chromatogram and peaks are proportional in area to the concentration of gases of interest | 1 | 0.5 | - | 0.5 |
| PC27. instill a medium into which the target analytes can be preferentially partitioned followed by the separation of the analyte-containing phase from the remainder of the sample | 1 | 0.5 | - | 0.5 |

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| Assessment Criteria for Outcomes | Theory Marks | Practical Marks | Project Marks | Viva Marks |
|--|--------------|-----------------|---------------|------------|
| PC28. administer sample is heated, have its pH adjusted, or be otherwise treated to make analytes more accessible to the partitioning medium | 1 | 0.5 | - | 0.5 |
| PC29. <ul style="list-style-type: none"> place solid analytes in solution for chromatographic separation irrespective of the extraction process Solution - additional pretreatments, such as liquid/liquid extraction (LLE) or solid-phase extraction (SPE), to remove substances that might interfere with the chromatographic separation and the detection of the target analytes Extraction Process - Liquid extraction, Ultrasonic simulation, Steam distillation | 1 | 0.5 | - | 0.5 |
| PC30. employ Liquid-liquid extraction, on- and offline solid-phase extraction, and gel permeation chromatography (GPC) in the partitioning of target analytes in liquid samples or liquid extracts of solid samples | 1 | 0.5 | - | 0.5 |
| PC31. filter turbid or otherwise opaque liquids to remove particulates prior to extraction | 1 | 0.5 | - | 0.5 |
| PC32. install guard columns at the head of the analytical column to prevent contamination of the stationary phase by the sample matrix | 1 | 0.5 | - | 0.5 |
| PC33. ensure the guard column is connected to the precolumn by means of a transfer valve | 0.5 | 0.5 | - | 0.5 |
| PC34. administer automated sampling system instead of manual injectors to get highly reproducible injection volume, low carryover, and increased sample throughput | 0.5 | 0.5 | - | 0.5 |
| PC35. assemble HPLC pumps to deliver pulse-free flow with high precision over a wide flow rate range with low dead volume | 0.5 | 0.5 | - | 0.5 |
| PC36. maintain a limit of detection (LOD) and limit of quantification (LOQ) as two to three and 10 to 20 times the noise level. LOD for food additives can be as low as 100 parts per trillion | 1 | 0.5 | - | 0.5 |

Qualification Pack

| Assessment Criteria for Outcomes | Theory Marks | Practical Marks | Project Marks | Viva Marks |
|--|--------------|-----------------|---------------|------------|
| PC37. <ul style="list-style-type: none"> administer other important detection parameters are appropriate, as per defined standards Parameters - dynamic range, calibration linearity, chromatographic selectivity, and qualitative information | 1 | 0.5 | - | 0.5 |
| PC38. automate derivatization and integrate online within the analysis to improve sensitivity and/or selectivity of an optical detector (UV, visible, or fluorescence) when analytes lack chromophores or an adequate optical response | 1 | 0.5 | - | 0.5 |
| PC39. administer to install of a lamp in Atomic Absorption (AA) unit for the analyte | 1 | 0.5 | - | 0.5 |
| PC40. regulate by nebulizing liquid sample and spray into an atomizer, or atom source | 0.5 | 0.5 | - | 0.5 |
| PC41. observe that the heat source dries the sample and atomizes it, breaking it down into individual elements. The light at a certain wavelength is passed through the atomized sample, where it is absorbed by elements of interest, and that energy absorption is measured | 1 | 0.5 | - | 0.5 |
| PC42. review in flame AA, the atomizer or atom source is in the form of an air/acetylene or nitrous oxide/acetylene flame | 0.5 | 0.5 | - | 0.5 |
| PC43. coordinate to introduce the sample as an aerosol into the flame, which is aligned so that the light beam passes through the flame, where the light is absorbed, measured, and reported by the instrument | 0.5 | 0.5 | - | 0.5 |
| PC44. facilitate Graphite Furnace Atomic Absorption (GFAA) by introducing the sample directly into a graphite tube, which is heated to remove liquid solvent and to atomize the remaining sample | 1 | 0.5 | - | 0.5 |
| PC45. observe that sample introduced to the tube is atomized, and the atoms are retained within the tube for an extended period | 0.5 | 0.5 | - | 0.5 |
| PC46. check light is passed through this tube and absorbance is measured and as a result of this improved analyte control sensitivity and detection limits are significantly improved | 0.5 | 0.5 | - | 0.5 |

Qualification Pack

| Assessment Criteria for Outcomes | Theory Marks | Practical Marks | Project Marks | Viva Marks |
|--|--------------|-----------------|---------------|------------|
| PC47. administer in Induction Coupled Plasma Mass Spectrometry (ICP-MS), the sample is atomized and ionized by the plasma source | 0.5 | 0.5 | - | 0.5 |
| PC48. observe charged ions are passed into a mass spectrometer, where ions are split according to their mass before reaching the detector, which in turn identifies and quantifies the elements present in a sample based on their mass | 0.5 | 0.5 | - | 0.5 |
| PC49. facilitate significant improvement in sensitivity by following the ICP-MS approach | 0.5 | 0.5 | - | 0.5 |
| <i>Carry out reporting and documentation</i> | 3.5 | 3.5 | - | 3.5 |
| PC50. review and update the test methods and procedures as per the schedule | 0.5 | 0.5 | - | 0.5 |
| PC51. ensure the updation of logbooks, worksheets, calibration records, parameters of columns, reagents, volumetric solution, and working standards | 0.5 | 0.5 | - | 0.5 |
| PC52. account for defects/problems/incidents/quality issues/test results as applicable as per SOP | 0.5 | 0.5 | - | 0.5 |
| PC53. prepare analytical and quality control reports for detailed findings and recommendations as per SOPs | 0.5 | 0.5 | - | 0.5 |
| PC54. prepare reports of validations, deviations, and incidents to the production and quality assurance team in compliance with GLP and GMP | 0.5 | 0.5 | - | 0.5 |
| PC55. facilitate to document testing results and analysis accurately | 0.5 | 0.5 | - | 0.5 |
| PC56. maintain all original and controlled document files and quality records in a timely and accurate manner and made them available for inspection and auditory purposes | 0.5 | 0.5 | - | 0.5 |
| NOS Total | 44 | 28 | - | 28 |

Qualification Pack

National Occupational Standards (NOS) Parameters

| | |
|----------------------------|---|
| NOS Code | FIC/N7630 |
| NOS Name | Carry out Physical, Chemical, and Instrumental Analysis |
| Sector | Food Processing |
| Sub-Sector | Fruits and Vegetables, Food Grain Milling (Including oil seeds), Dairy Products, Meat and Poultry, Fish and Seafood, Bread and Bakery, Alcoholic Beverages, Aerated Water/Soft Drinks |
| Occupation | Quality Analysis/ Assurance |
| NSQF Level | 5 |
| Credits | 6 |
| Version | 1.0 |
| Last Reviewed Date | 23/06/2023 |
| Next Review Date | 23/06/2026 |
| NSQC Clearance Date | 23/06/2023 |

Qualification Pack

FIC/N7631: Carry out maintenance and decommissioning of equipment

Description

This OS unit is about carrying out the activities for preventive maintenance, repairing equipment and decommissioning of equipment along with documentation and reporting as per standards.

Scope

The scope covers the following :

- This unit/task covers the following:
- Carry out preventive planned maintenance, Overhaul instrumentation equipment, Decommissioning of equipment, Carry out documentation as per regulatory compliance standards

Elements and Performance Criteria

Carry out preventive planned maintenance

To be competent, the user/individual on the job must be able to:

- PC1.** work safely at all times, complying with health and safety, environmental and other relevant food and drink regulations, directives and guidelines
- PC2.** obtain and use the correct issue of company and/or manufacturers' drawings and maintenance documentation
- PC3.** plan and communicate the maintenance activities so as to minimise any disruption to normal working
- PC4.** follow the relevant maintenance schedules to carry out the required work
- PC5.** insert or override any relevant system trip defeats (including fire extinguishant, emergency shutdown) in accordance with organisational procedures
- PC6.** isolate instruments (including process, electrical, hydraulic, pneumatic, mechanical) in accordance with organisational procedures
- PC7.** provide and maintain safe access and working arrangements for the maintenance area
- PC8.** carry out the maintenance activities in accordance with organisational procedures within the limits of your personal authority
- PC9.** carry out functional tests and adjust equipment to specification
- PC10.** re-connect and return the system to service on completion of the maintenance activities
- PC11.** report any instances where the maintenance activities cannot be fully met or where there are identified defects outside the planned schedule
- PC12.** complete maintenance records and documentation in accordance with organisational requirements
- PC13.** dispose of waste materials in accordance with safe working practices and approved procedures

Overhaul instrumentation equipment

To be competent, the user/individual on the job must be able to:

- PC14.** obtain and use the correct equipment repair/overhauling documentation (including manuals, drawings, maintenance records)

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- PC15.** follow the relevant repair/overhauling schedules to carry out the required work
- PC16.** provide and maintain safe access and working arrangements for the repair/overhauling area
- PC17.** follow decontamination procedures for instruments that have been used with hazardous (including toxic, corrosive, inflammable, explosive, radioactive) substances
- PC18.** apply electrostatic discharge (ESD) protection procedures as per procedures
- PC19.** establish and mark/label components to aid in re-assembly
- PC20.** check that equipment is maintained free from damage and foreign objects
- PC21.** carry out repair/overhaul to the agreed level, using correct tools and techniques as per standards
- PC22.** identify and store all removed components in accordance with organisational procedures
- PC23.** report any instances where repair/overhauling activities cannot be fully met, or where there are identified defects outside planned repair/overhauling schedule
- PC24.** dispose of unwanted components, waste materials and substances, in accordance with safe working practices approved procedures

Decommissioning of equipment

To be competent, the user/individual on the job must be able to:

- PC25.** conduct equipment survey or inspection before planning for decommissioning
- PC26.** identify and select equipment/facility to be decommissioned for recycling, reuse or resale
- PC27.** record manufacturer, serial number, equipment specs, part number, suite and rack location/area and photos, etc. before initiating the process
- PC28.** coordinate with facility managers to finalize equipment inventory for review and approval
- PC29.** administer decommissioning of approved equipment ensuring site security
- PC30.** direct wiped, decommissioned and disconnected equipment towards the circular repository
- PC31.** prepare transfer notes and reports for audit compliance

Carry out documentation as per regulatory compliance standards

To be competent, the user/individual on the job must be able to:

- PC32.** identify procedures, records and workplace documentation needed for audit
- PC33.** document evidence collection methods and sources along with checklists developed as per specifications
- PC34.** identify and categorized non - conformities observed from prior audit trails and plan actionable steps
- PC35.** maintain records of effectiveness of corrective actions

Knowledge and Understanding (KU)

The individual on the job needs to know and understand:

- KU1.** health and safety and environmental requirements of the area in which the repair/overhaul activity is to take place, and the responsibility these requirements place on you not to compromise food safety
- KU2.** isolation and lock-off procedure or permit-to-work procedure that applies to the equipment, including the critical control points

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- KU3.** specific health and safety food and drink precautions to be applied during the repair/overhaul activity, and their effects on other
- KU4.** responsibilities in relationship to Hazard Analysis and Critical Control Points (HACCP), Threat Assessment and Critical Control Points (TACCP), Vulnerability Assessment and Critical Control Points (VACCP) during the repair/overhaul activities
- KU5.** what constitutes a hazardous voltage and how to recognize victims of electric shock
- KU6.** how to reduce the risks of a phase to earth shock (including insulated tools, rubber mating and isolating transformers)
- KU7.** protective equipment that you need to use for both personal protection (PPE) and protection of the instrumentation and control equipment being repaired
- KU8.** hazards associated with repairing/overhauling instrumentation and control equipment, and with the tools and equipment used, and how to minimise them and reduce any risks
- KU9.** how to extract and use information from equipment manuals, history/maintenance reports, charts, circuit and physical layouts, specifications, symbols used in instrumentation and control circuits, and other documents needed in the repair/overhaul process
- KU10.** terminology used with instrumentation and control equipment, and the use of system diagrams and associated symbols
- KU11.** basic principles of operation of the instrumentation and control equipment being repaired/overhauled, and the performance
- KU12.** how to make sensory checks (by sight, sound, smell, touch) where to obtain, and how to interpret drawings, schematic and physical diagrams, specifications, flow charts, manufacturers' manuals, maintenance schedules and other documents required for the maintenance activities the various planned maintenance schedules that are generally used (including condition-based maintenance, scheduled maintenance and total preventative maintenance (TPM))
- KU13.** schedules and methods to be followed, in compliance with company procedures for planned maintenance on instrumentation and control equipment
- KU14.** basic principles of how the system functions, its operating sequence, the working purpose of individual units/components and how they interact
- KU15.** principles of the equipment's design feature for safe operation in a food or drink environment including minimising the chance of contaminants or foreign bodies in the final product
- KU16.** equipment operating and control procedures, and how to apply them along with the planned maintenance procedures
- KU17.** reasons for making sure that control systems are isolated or put into manual control, and appropriate trip locks, keys or program overrides are inserted, before removing any sensors or instruments from the system
- KU18.** the identification of instrument sensors (including how to identify their markings, calibration information, operating parameters and working range)
- KU18.** methods of checking and calibrating instruments, and the type and range of equipment that can be used
- KU19.** testing methods and procedures to be used to check that system conforms within acceptable limits
- KU20.** procedure for obtaining consumables and 'lived' or consumable items that will require replacing during the planned maintenance activity
- KU21.** company policy on the repair/replacement of components during the maintenance

Qualification Pack

- KU22.** problems that can occur whilst carrying out planned maintenance activities, and how they can be avoided
- KU23.** organisational procedure to be adopted for the safe disposal of waste of all types of materials including any spoilt food or drink products
- KU24.** extent of your own authority and to whom you should report if you have problems that you cannot resolve
- KU25.** legislation, regulations, orders, codes and standards applicable to the areas being audited
- KU26.** communication methods relevant to different groups and audience
- KU27.** legislation that impacts on acceptable communication methods and conduct including anti-discrimination, anti-harassment and privacy legislation
- KU28.** methods used to identify Critical Control Points and establish critical limits, suited to the nature of the hazard, the requirements of the audit and the industry sector
- KU29.** the underlying principles of risk-based approaches to controlling food safety hazards including HACCP as described in the Codex Alimentarius Guidelines
- KU30.** vocabulary and terms relating to food safety, including terms and jargon to describe technical processes, industry standards and common biological and chemical terms
- KU31.** activities involved in decommissioning such as, wiping data and migrating services, checking processes for backup and recovery are fully functional, identifying any SLAs, licenses and warranties that may need to be cancelled, disconnecting utilities, considering the fire safety of the vacated facility or area, ensuring the site remains secure during and post-decommissioning to prevent trespassing

Generic Skills (GS)

User/individual on the job needs to know how to:

- GS1.** read and interpret information such as product labels, safety data sheets, instructions, procedures, and specifications to determine activities in honey extraction processes
- GS2.** write and document using appropriate terminology and in specified format
- GS3.** communicate with others effectively
- GS4.** takes responsibility for planning, sequencing, and prioritising tasks
- GS5.** analyse and report variances promptly

Qualification Pack

Assessment Criteria

| Assessment Criteria for Outcomes | Theory Marks | Practical Marks | Project Marks | Viva Marks |
|---|--------------|-----------------|---------------|------------|
| <i>Carry out preventive planned maintenance</i> | 19 | 13 | - | 6.5 |
| PC1. work safely at all times, complying with health and safety, environmental and other relevant food and drink regulations, directives and guidelines | 1 | 1 | - | 0.5 |
| PC2. obtain and use the correct issue of company and/or manufacturers' drawings and maintenance documentation | 1 | 1 | - | 0.5 |
| PC3. plan and communicate the maintenance activities so as to minimise any disruption to normal working | 2 | 1 | - | 0.5 |
| PC4. follow the relevant maintenance schedules to carry out the required work | 1 | 1 | - | 0.5 |
| PC5. insert or override any relevant system trip defeats (including fire extinguishant, emergency shutdown) in accordance with organisational procedures | 2 | 1 | - | 0.5 |
| PC6. isolate instruments (including process, electrical, hydraulic, pneumatic, mechanical) in accordance with organisational procedures | 1 | 1 | - | 0.5 |
| PC7. provide and maintain safe access and working arrangements for the maintenance area | 2 | 1 | - | 0.5 |
| PC8. carry out the maintenance activities in accordance with organisational procedures within the limits of your personal authority | 1 | 1 | - | 0.5 |
| PC9. carry out functional tests and adjust equipment to specification | 2 | 1 | - | 0.5 |
| PC10. re-connect and return the system to service on completion of the maintenance activities | 1 | 1 | - | 0.5 |
| PC11. report any instances where the maintenance activities cannot be fully met or where there are identified defects outside the planned schedule | 2 | 1 | - | 0.5 |

Qualification Pack

| Assessment Criteria for Outcomes | Theory Marks | Practical Marks | Project Marks | Viva Marks |
|--|--------------|-----------------|---------------|------------|
| PC12. complete maintenance records and documentation in accordance with organisational requirements | 1 | 1 | - | 0.5 |
| PC13. dispose of waste materials in accordance with safe working practices and approved procedures | 2 | 1 | - | 0.5 |
| <i>Overhaul instrumentation equipment</i> | 17 | 11 | - | 5.5 |
| PC14. obtain and use the correct equipment repair/overhauling documentation (including manuals, drawings, maintenance records) | 1 | 1 | - | 0.5 |
| PC15. follow the relevant repair/overhauling schedules to carry out the required work | 1 | 1 | - | 0.5 |
| PC16. provide and maintain safe access and working arrangements for the repair/overhauling area | 1 | 1 | - | 0.5 |
| PC17. follow decontamination procedures for instruments that have been used with hazardous (including toxic, corrosive, inflammable, explosive, radioactive) substances | 2 | 1 | - | 0.5 |
| PC18. apply electrostatic discharge (ESD) protection procedures as per procedures | 1 | 1 | - | 0.5 |
| PC19. establish and mark/label components to aid in re-assembly | 2 | 1 | - | 0.5 |
| PC20. check that equipment is maintained free from damage and foreign objects | 2 | 1 | - | 0.5 |
| PC21. carry out repair/overhaul to the agreed level, using correct tools and techniques as per standards | 2 | 1 | - | 0.5 |
| PC22. identify and store all removed components in accordance with organisational procedures | 1 | 1 | - | 0.5 |
| PC23. report any instances where repair/overhauling activities cannot be fully met, or where there are identified defects outside planned repair/overhauling schedule | 2 | 1 | - | 0.5 |

Qualification Pack

| Assessment Criteria for Outcomes | Theory Marks | Practical Marks | Project Marks | Viva Marks |
|--|--------------|-----------------|---------------|------------|
| PC24. dispose of unwanted components, waste materials and substances, in accordance with safe working practices approved procedures | 2 | 1 | - | 0.5 |
| <i>Decommissioning of equipment</i> | 7 | 7 | - | 4 |
| PC25. conduct equipment survey or inspection before planning for decommissioning | 1 | 1 | - | 0.5 |
| PC26. identify and select equipment/facility to be decommissioned for recycling, reuse or resale | 1 | 1 | - | 0.5 |
| PC27. record manufacturer, serial number, equipment specs, part number, suite and rack location/area and photos, etc. before initiating the process | 1 | 1 | - | 0.5 |
| PC28. coordinate with facility managers to finalize equipment inventory for review and approval | 1 | 1 | - | 0.5 |
| PC29. administer decommissioning of approved equipment ensuring site security | 0.5 | 1 | - | 1 |
| PC30. direct wiped, decommissioned and disconnected equipment towards the circular repository | 1.5 | 1 | - | 0.5 |
| PC31. prepare transfer notes and reports for audit compliance | 1 | 1 | - | 0.5 |
| <i>Carry out documentation as per regulatory compliance standards</i> | 4 | 4 | - | 2 |
| PC32. identify procedures, records and workplace documentation needed for audit | 1 | 1 | - | 0.5 |
| PC33. document evidence collection methods and sources along with checklists developed as per specifications | 1 | 1 | - | 0.5 |
| PC34. identify and categorized non - conformities observed from prior audit trails and plan actionable steps | 1 | 1 | - | 0.5 |
| PC35. maintain records of effectiveness of corrective actions | 1 | 1 | - | 0.5 |
| NOS Total | 47 | 35 | - | 18 |

Qualification Pack

National Occupational Standards (NOS) Parameters

| | |
|----------------------------|--|
| NOS Code | FIC/N7631 |
| NOS Name | Carry out maintenance and decommissioning of equipment |
| Sector | Food Processing |
| Sub-Sector | Generic |
| Occupation | Quality Analysis/ Assurance |
| NSQF Level | 5 |
| Credits | 5 |
| Version | 1.0 |
| Last Reviewed Date | 23/06/2023 |
| Next Review Date | 23/06/2026 |
| NSQC Clearance Date | 23/06/2023 |

Qualification Pack

FIC/N9904: Ensure food safety at the workplace

Description

This unit is about performing various tasks for ensuring food safety at the workplace.

Scope

The scope covers the following :

- Ensure food safety at the workplace

Elements and Performance Criteria

Ensure food safety at the workplace

To be competent, the user/individual on the job must be able to:

- PC1.** identify the biological, chemical, and physical hazards at various stages of food processing. Stages: procurement of raw material; production, manufacturing, distribution, delivery of finished product, etc.
- PC2.** implement food safety procedures and regulatory policies at the food processing workplace. Policies: Visitor's Policy, Health declaration policy, Jewellery policy, Quality, and safety policy
- PC3.** ensure that the materials are adequately isolated to prevent them from contamination. Materials: raw materials, processed materials, finished goods, etc. Contamination: Physical, Chemical, Biological & shop floor environment
- PC4.** establish and follow Good Manufacturing Practices (GMPs) laid down in applicable Food Safety and Standards Authority of India (FSSAI) guidelines. Good Manufacturing Practices (GMPs): location and layout(ergonomics), cleaning and sanitation, equipment and containers, pest control, facilities (lighting, water supply, drainage and waste disposal, air quality and ventilation), food storage, transportation, and distribution etc.
- PC5.** establish and follow allergen management system for handling and storage of raw materials
- PC6.** establish and follow monitoring systems like Hazard Analysis Critical Control Point (HACCP), product information and consumer awareness, product recall and withdrawal, and traceability HACCP: Hazard identification, identification of critical control points, establish critical limits, corrective and preventive action. Product information and consumer awareness: Product labelling and consumer education. Traceability: forward and backward traceability
- PC7.** take appropriate action in instances such as VACCP (Vulnerability Assessment Critical Control Points) and TACCP (Threat Assessment Critical Control Points)
- PC8.** plan, conduct, manage, consolidate outcomes, and close corrective actions of workplace audit on food safety as per FSSAI guidelines, address the non-conformance with root cause analysis (RCA), corrective action preventive action(CAPA)
- PC9.** address issues pertaining to food safety and quality reported by the team members
- PC10.** record information such as food safety regulations followed, inspections done, faults observed, etc. as per standard procedure
- PC11.** organize trainings and workshops on food safety aspects such as Good Manufacturing Practices (GMP), HACCP, VACCP, TACCP, etc.

Qualification Pack

Knowledge and Understanding (KU)

The individual on the job needs to know and understand:

- KU1.** various processes that take place in a food industry
- KU2.** biological, chemical, and physical hazards in a food industry
- KU3.** types of food contaminations, their causes, and ways to prevent it
- KU4.** organisational policy and procedures for ensuring food safety (such as Visitor's Policy, Health declaration policy, Jewelry policy, Quality, and safety policy)
- KU5.** applicable regulations for ensuring food safety as listed in 'The Food Safety and Standards Act, 2006'
- KU6.** role of HACCP in food industry, its constituents and procedure to implement it in an organisation
- KU7.** VACCP and TACCP and how to implement it effectively
- KU8.** how to conduct workplace food safety audits
- KU9.** types of allergen and allergen management at workplace
- KU10.** key observations and corrective actions to be applied for ensuring food safety
- KU11.** various issues that can arise during production and other processes as faced by team
- KU12.** information to be recorded in the work process
- KU13.** how to do root cause analysis and perform corrective action and preventive actions
- KU14.** how to conduct training of workforce on various food safety procedures such as GMP, HACCP, information to be shared, ways to report accidents, escalation of issues beyond own scope, etc.

Generic Skills (GS)

User/individual on the job needs to know how to:

- GS1.** write an accident/incident report in local language or English
- GS2.** read and comprehend basic content to read labels, charts, signages and symbols
- GS3.** read and comprehend basic English to read product manuals for safe operation
- GS4.** question coworkers appropriately in order to clarify instructions and other issues
- GS5.** make appropriate decisions pertaining to the concerned area of work regarding the work objective, span of authority, responsibility, laid down procedure and guidelines
- GS6.** plan and organize the work schedule, work area, tools, equipment, and materials for improved productivity
- GS7.** identify probable solutions to the problems in hand
- GS8.** evaluate proposed solution with respect to key priorities and considerations
- GS9.** seek official and authorised sources of help and guidance to resolve problems that cannot be solved at one's level of authority
- GS10.** identify cause and effect relations in their area of work to anticipate potential problems and their solution
- GS11.** analyse the problem, suggest corrective actions and implement workable solutions

Qualification Pack

Assessment Criteria

| Assessment Criteria for Outcomes | Theory Marks | Practical Marks | Project Marks | Viva Marks |
|---|--------------|-----------------|---------------|------------|
| <i>Ensure food safety at the workplace</i> | 30 | 70 | - | - |
| PC1. identify the biological, chemical, and physical hazards at various stages of food processing. Stages: procurement of raw material; production, manufacturing, distribution, delivery of finished product, etc. | - | - | - | - |
| PC2. implement food safety procedures and regulatory policies at the food processing workplace. Policies: Visitor's Policy, Health declaration policy, Jewellery policy, Quality, and safety policy | - | - | - | - |
| PC3. ensure that the materials are adequately isolated to prevent them from contamination. Materials: raw materials, processed materials, finished goods, etc. Contamination: Physical, Chemical, Biological & shop floor environment | - | - | - | - |
| PC4. establish and follow Good Manufacturing Practices (GMPs) laid down in applicable Food Safety and Standards Authority of India (FSSAI) guidelines. Good Manufacturing Practices (GMPs): location and layout(ergonomics), cleaning and sanitation, equipment and containers, pest control, facilities (lighting, water supply, drainage and waste disposal, air quality and ventilation), food storage, transportation, and distribution etc. | - | - | - | - |
| PC5. establish and follow allergen management system for handling and storage of raw materials | - | - | - | - |
| PC6. establish and follow monitoring systems like Hazard Analysis Critical Control Point (HACCP), product information and consumer awareness, product recall and withdrawal, and traceability HACCP: Hazard identification, identification of critical control points, establish critical limits, corrective and preventive action. Product information and consumer awareness: Product labelling and consumer education. Traceability: forward and backward traceability | - | - | - | - |

Qualification Pack

| Assessment Criteria for Outcomes | Theory Marks | Practical Marks | Project Marks | Viva Marks |
|--|--------------|-----------------|---------------|------------|
| PC7. take appropriate action in instances such as VACCP (Vulnerability Assessment Critical Control Points) and TACCP (Threat Assessment Critical Control Points) | - | - | - | - |
| PC8. plan, conduct, manage, consolidate outcomes, and close corrective actions of workplace audit on food safety as per FSSAI guidelines, address the non-conformance with root cause analysis (RCA), corrective action preventive action(CAPA) | - | - | - | - |
| PC9. address issues pertaining to food safety and quality reported by the team members | - | - | - | - |
| PC10. record information such as food safety regulations followed, inspections done, faults observed, etc. as per standard procedure | - | - | - | - |
| PC11. organize trainings and workshops on food safety aspects such as Good Manufacturing Practices (GMP), HACCP, VACCP, TACCP, etc. | - | - | - | - |
| NOS Total | 30 | 70 | - | - |

Qualification Pack

National Occupational Standards (NOS) Parameters

| | |
|----------------------------|-------------------------------------|
| NOS Code | FIC/N9904 |
| NOS Name | Ensure food safety at the workplace |
| Sector | Food Processing |
| Sub-Sector | Generic |
| Occupation | Generic |
| NSQF Level | 5 |
| Credits | TBD |
| Version | 1.0 |
| Last Reviewed Date | 23/06/2023 |
| Next Review Date | 23/06/2026 |
| NSQF Clearance Date | 23/06/2023 |

Qualification Pack

DGT/VSQ/N0101: Employability Skills (30 Hours)

Description

This unit is about employability skills, Constitutional values, becoming a professional in the 21st Century, digital, financial, and legal literacy, diversity and Inclusion, English and communication skills, customer service, entrepreneurship, and apprenticeship, getting ready for jobs and career development.

Scope

The scope covers the following :

- Introduction to Employability Skills
- Constitutional values - Citizenship
- Becoming a Professional in the 21st Century
- Basic English Skills
- Communication Skills
- Diversity & Inclusion
- Financial and Legal Literacy
- Essential Digital Skills
- Entrepreneurship
- Customer Service
- Getting ready for Apprenticeship & Jobs

Elements and Performance Criteria

Introduction to Employability Skills

To be competent, the user/individual on the job must be able to:

PC1. understand the significance of employability skills in meeting the job requirements

Constitutional values - Citizenship

To be competent, the user/individual on the job must be able to:

PC2. identify constitutional values, civic rights, duties, personal values and ethics and environmentally sustainable practices

Becoming a Professional in the 21st Century

To be competent, the user/individual on the job must be able to:

PC3. explain 21st Century Skills such as Self-Awareness, Behavior Skills, Positive attitude, self-motivation, problem-solving, creative thinking, time management, social and cultural awareness, emotional awareness, continuous learning mindset etc.

Basic English Skills

To be competent, the user/individual on the job must be able to:

PC4. speak with others using some basic English phrases or sentences

Communication Skills

To be competent, the user/individual on the job must be able to:

PC5. follow good manners while communicating with others

PC6. work with others in a team

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Diversity & Inclusion

To be competent, the user/individual on the job must be able to:

PC7. communicate and behave appropriately with all genders and PwD

PC8. report any issues related to sexual harassment

Financial and Legal Literacy

To be competent, the user/individual on the job must be able to:

PC9. use various financial products and services safely and securely

PC10. calculate income, expenses, savings etc.

PC11. approach the concerned authorities for any exploitation as per legal rights and laws

Essential Digital Skills

To be competent, the user/individual on the job must be able to:

PC12. operate digital devices and use its features and applications securely and safely

PC13. use internet and social media platforms securely and safely

Entrepreneurship

To be competent, the user/individual on the job must be able to:

PC14. identify and assess opportunities for potential business

PC15. identify sources for arranging money and associated financial and legal challenges

Customer Service

To be competent, the user/individual on the job must be able to:

PC16. identify different types of customers

PC17. identify customer needs and address them appropriately

PC18. follow appropriate hygiene and grooming standards

Getting ready for apprenticeship & Jobs

To be competent, the user/individual on the job must be able to:

PC19. create a basic biodata

PC20. search for suitable jobs and apply

PC21. identify and register apprenticeship opportunities as per requirement

Knowledge and Understanding (KU)

The individual on the job needs to know and understand:

KU1. need for employability skills

KU2. various constitutional and personal values

KU3. different environmentally sustainable practices and their importance

KU4. Twenty first (21st) century skills and their importance

KU5. how to use basic spoken English language

KU6. Do and dont of effective communication

KU7. inclusivity and its importance

KU8. different types of disabilities and appropriate communication and behaviour towards PwD

KU9. different types of financial products and services

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- KU10.** how to compute income and expenses
- KU11.** importance of maintaining safety and security in financial transactions
- KU12.** different legal rights and laws
- KU13.** how to operate digital devices and applications safely and securely
- KU14.** ways to identify business opportunities
- KU15.** types of customers and their needs
- KU16.** how to apply for a job and prepare for an interview
- KU17.** apprenticeship scheme and the process of registering on apprenticeship portal

Generic Skills (GS)

User/individual on the job needs to know how to:

- GS1.** communicate effectively using appropriate language
- GS2.** behave politely and appropriately with all
- GS3.** perform basic calculations
- GS4.** solve problems effectively
- GS5.** be careful and attentive at work
- GS6.** use time effectively
- GS7.** maintain hygiene and sanitisation to avoid infection

Qualification Pack

Assessment Criteria

| Assessment Criteria for Outcomes | Theory Marks | Practical Marks | Project Marks | Viva Marks |
|--|--------------|-----------------|---------------|------------|
| <i>Introduction to Employability Skills</i> | 1 | 1 | - | - |
| PC1. understand the significance of employability skills in meeting the job requirements | - | - | - | - |
| <i>Constitutional values - Citizenship</i> | 1 | 1 | - | - |
| PC2. identify constitutional values, civic rights, duties, personal values and ethics and environmentally sustainable practices | - | - | - | - |
| <i>Becoming a Professional in the 21st Century</i> | 1 | 3 | - | - |
| PC3. explain 21st Century Skills such as Self-Awareness, Behavior Skills, Positive attitude, self-motivation, problem-solving, creative thinking, time management, social and cultural awareness, emotional awareness, continuous learning mindset etc. | - | - | - | - |
| <i>Basic English Skills</i> | 2 | 3 | - | - |
| PC4. speak with others using some basic English phrases or sentences | - | - | - | - |
| <i>Communication Skills</i> | 1 | 1 | - | - |
| PC5. follow good manners while communicating with others | - | - | - | - |
| PC6. work with others in a team | - | - | - | - |
| <i>Diversity & Inclusion</i> | 1 | 1 | - | - |
| PC7. communicate and behave appropriately with all genders and PwD | - | - | - | - |
| PC8. report any issues related to sexual harassment | - | - | - | - |
| <i>Financial and Legal Literacy</i> | 3 | 4 | - | - |
| PC9. use various financial products and services safely and securely | - | - | - | - |

Qualification Pack

| Assessment Criteria for Outcomes | Theory Marks | Practical Marks | Project Marks | Viva Marks |
|---|--------------|-----------------|---------------|------------|
| PC10. calculate income, expenses, savings etc. | - | - | - | - |
| PC11. approach the concerned authorities for any exploitation as per legal rights and laws | - | - | - | - |
| <i>Essential Digital Skills</i> | 4 | 6 | - | - |
| PC12. operate digital devices and use its features and applications securely and safely | - | - | - | - |
| PC13. use internet and social media platforms securely and safely | - | - | - | - |
| <i>Entrepreneurship</i> | 3 | 5 | - | - |
| PC14. identify and assess opportunities for potential business | - | - | - | - |
| PC15. identify sources for arranging money and associated financial and legal challenges | - | - | - | - |
| <i>Customer Service</i> | 2 | 2 | - | - |
| PC16. identify different types of customers | - | - | - | - |
| PC17. identify customer needs and address them appropriately | - | - | - | - |
| PC18. follow appropriate hygiene and grooming standards | - | - | - | - |
| <i>Getting ready for apprenticeship & Jobs</i> | 1 | 3 | - | - |
| PC19. create a basic biodata | - | - | - | - |
| PC20. search for suitable jobs and apply | - | - | - | - |
| PC21. identify and register apprenticeship opportunities as per requirement | - | - | - | - |
| NOS Total | 20 | 30 | - | - |

Qualification Pack

National Occupational Standards (NOS) Parameters

| | |
|----------------------------|---------------------------------|
| NOS Code | DGT/VSQ/N0101 |
| NOS Name | Employability Skills (30 Hours) |
| Sector | Cross Sectoral |
| Sub-Sector | Professional Skills |
| Occupation | Employability |
| NSQF Level | 2 |
| Credits | 1 |
| Version | 1.0 |
| Last Reviewed Date | 23/06/2023 |
| Next Review Date | 23/06/2026 |
| NSQC Clearance Date | 23/06/2023 |

Assessment Guidelines and Assessment Weightage

Assessment Guidelines

1. Criteria for assessment for each Qualification Pack will be created by the Sector Skill Council. Each Performance Criteria (PC) will be assigned marks proportional to its importance in NOS. SSC will also lay down proportion of marks for Theory and Skills Practical for each PC.
2. The assessment for the theory part will be based on knowledge bank of questions created by the SSC.
3. Assessment will be conducted for all compulsory NOS, and where applicable, on the selected elective/option NOS/set of NOS.
4. Individual assessment agencies will create unique question papers for theory part for each candidate at each examination/training centre (as per assessment criteria below).
5. Individual assessment agencies will create unique evaluations for skill practical for every student at each examination/training centre based on this criterion.
6. To pass the Qualification Pack, every trainee should score a minimum of 70% of aggregate marks to successfully clear the assessment.
7. In case of unsuccessful completion, the trainee may seek reassessment on the Qualification Pack.

Qualification Pack

Minimum Aggregate Passing % at QP Level : 70

(Please note: Every Trainee should score a minimum aggregate passing percentage as specified above, to successfully clear the Qualification Pack assessment.)

Minimum Passing % at NOS Level: 70

(Please note: A Trainee must score the minimum percentage for each NOS separately as well as on the QP as a whole.)

Assessment Weightage

Compulsory NOS

| National Occupational Standards | Theory Marks | Practical Marks | Project Marks | Viva Marks | Total Marks | Weightage |
|--|--------------|-----------------|---------------|------------|-------------|------------|
| FIC/N7629.Planning, Organizing, and Setting up of Food Analysis Laboratory | 38 | 31 | - | 31 | 100 | 25 |
| FIC/N7630.Carry out Physical, Chemical, and Instrumental Analysis | 44 | 28 | - | 28 | 100 | 25 |
| FIC/N7631.Carry out maintenance and decommissioning of equipment | 47 | 35 | - | 18 | 100 | 25 |
| FIC/N9904.Ensure food safety at the workplace | 30 | 70 | - | - | 100 | 15 |
| DGT/VSQ/N0101.Employability Skills (30 Hours) | 20 | 30 | - | - | 50 | 10 |
| Total | 179 | 194 | - | 77 | 450 | 100 |

Qualification Pack

Acronyms

| | |
|-------------|--|
| NOS | National Occupational Standard(s) |
| NSQF | National Skills Qualifications Framework |
| QP | Qualifications Pack |
| TVET | Technical and Vocational Education and Training |
| AA | Assessment Agency |
| AB | Awarding Body |
| ISCO | International Standard Classification of Occupations |
| NCO | National Classification of Occupations |
| NCrF | National Credit Framework |
| NOS | National Occupational Standard(s) |
| NQR | National Qualification Register |
| NSQF | National Skills Qualifications Framework |
| OJT | On-the-Job Training |
| NOS | National Occupational Standard(s) |
| NSQF | National Skills Qualifications Framework |
| QP | Qualifications Pack |
| TVET | Technical and Vocational Education and Training |

Qualification Pack

Glossary

| | |
|--|--|
| Sector | Sector is a conglomeration of different business operations having similar business and interests. It may also be defined as a distinct subset of the economy whose components share similar characteristics and interests. |
| Sub-sector | Sub-sector is derived from a further breakdown based on the characteristics and interests of its components. |
| Occupation | Occupation is a set of job roles, which perform similar/ related set of functions in an industry. |
| Job role | Job role defines a unique set of functions that together form a unique employment opportunity in an organisation. |
| Occupational Standards (OS) | OS specify the standards of performance an individual must achieve when carrying out a function in the workplace, together with the Knowledge and Understanding (KU) they need to meet that standard consistently. Occupational Standards are applicable both in the Indian and global contexts. |
| Performance Criteria (PC) | Performance Criteria (PC) are statements that together specify the standard of performance required when carrying out a task. |
| National Occupational Standards (NOS) | NOS are occupational standards which apply uniquely in the Indian context. |
| Qualifications Pack (QP) | QP comprises the set of OS, together with the educational, training and other criteria required to perform a job role. A QP is assigned a unique qualifications pack code. |
| Unit Code | Unit code is a unique identifier for an Occupational Standard, which is denoted by an 'N' |
| Unit Title | Unit title gives a clear overall statement about what the incumbent should be able to do. |
| Description | Description gives a short summary of the unit content. This would be helpful to anyone searching on a database to verify that this is the appropriate OS they are looking for. |
| Scope | Scope is a set of statements specifying the range of variables that an individual may have to deal with in carrying out the function which have a critical impact on quality of performance required. |

Qualification Pack

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| Knowledge and Understanding (KU) | Knowledge and Understanding (KU) are statements which together specify the technical, generic, professional and organisational specific knowledge that an individual needs in order to perform to the required standard. |
| Organisational Context | Organisational context includes the way the organisation is structured and how it operates, including the extent of operative knowledge managers have of their relevant areas of responsibility. |
| Technical Knowledge | Technical knowledge is the specific knowledge needed to accomplish specific designated responsibilities. |
| Core Skills/ Generic Skills (GS) | Core skills or Generic Skills (GS) are a group of skills that are the key to learning and working in today's world. These skills are typically needed in any work environment in today's world. These skills are typically needed in any work environment. In the context of the OS, these include communication related skills that are applicable to most job roles. |
| Electives | Electives are NOS/set of NOS that are identified by the sector as contributive to specialization in a job role. There may be multiple electives within a QP for each specialized job role. Trainees must select at least one elective for the successful completion of a QP with Electives. |
| Options | Options are NOS/set of NOS that are identified by the sector as additional skills. There may be multiple options within a QP. It is not mandatory to select any of the options to complete a QP with Options. |
| National Occupational Standard | NOS defines the measurable performance outcomes required from an individual engaged in a particular task. They list down what an individual performing that task should know and also do. |
| Qualifications | A formal outcome of an assessment and validation process is obtained when a competent body determines that an individual has achieved learning outcomes to given standards |
| Qualification File | A Qualification File is a template designed to capture necessary information of a Qualification from the perspective of NSQF compliance. The Qualification File will be normally submitted by the awarding body for the qualification. |
| Sector | A grouping of professional activities on the basis of their main economic function, product, service, or technology. |
| Long Term Training | Long-term skilling means any vocational training program undertaken for a year and above. |
| Sector | Sector is a conglomeration of different business operations having similar business and interests. It may also be defined as a distinct subset of the economy whose components share similar characteristics and interests. |

Qualification Pack

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| Sub-sector | Sub-sector is derived from a further breakdown based on the characteristics and interests of its components. |
| Occupation | Occupation is a set of job roles, which perform similar/ related set of functions in an industry. |
| Job role | Job role defines a unique set of functions that together form a unique employment opportunity in an organisation. |
| Occupational Standards | OS specify the standards of performance an individual must achieve when carrying out a function in the workplace, together with the Knowledge and Understanding (KU) they need to meet that standard consistently. Occupational Standards are applicable both in the Indian and global contexts. |
| Performance Criteria (PC) | Performance Criteria (PC) are statements that together specify the standard of performance required when carrying out a task. |
| National Occupational Standard | NOS are occupational standards which apply uniquely in the Indian context. |
| Qualification Pack(QP) | QP comprises the set of OS, together with the educational, training and other criteria required to perform a job role. A QP is assigned a unique qualifications pack code. |
| Unit Code | Unit code is a unique identifier for an Occupational Standard, which is denoted by an 'N' |
| Unit Title | Unit title gives a clear overall statement about what the incumbent should be able to do. |
| Description | Description gives a short summary of the unit content. This would be helpful to anyone searching on a database to verify that this is the appropriate OS they are looking for. |
| Scope | Scope is a set of statements specifying the range of variables that an individual may have to deal with in carrying out the function which have a critical impact on quality of performance required. |