

# **SAMPLE COLLECTION, HANDLING AND PREPARATION**

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# Sample collection

- ⦿ The adequacy and condition of the sample or specimen received for examination are of primary importance
- ⦿ If samples are improperly collected: the laboratory results will be meaningless
- ⦿ Sampling protocol should be clearly defined
  - ⦿ Start with description of primary food product

# Sample collection

- Identity of the food
  - Common/alternative name
    - E.g. Maize, Nigeria beans
  - Scientific name (Genus, species, variety)
    - E.g. Zea mays, Vigna unguiculata
  - Plant food (entire plant/part e.g. roots)
  - Animal food (entire animal/part)
  - State of maturity (ripe immature)
  - Other details

# Sample collection

- Need to know:
  - Number and size of sample to be collected
  - Distribution of samples
  - Stratification to be used
- Sample label should be permanently attached to the sample
  - Common name of food
  - Sample code number
  - Date of receipt in Lab.

# Sample collection

- During sample collection:

- Collection details

- Date and time of collection
- Name of collector
- Place of origin
- Sampling point/addresses (roadside stall, farm, market)
- Condition of cultivation (feed regime, altitude, irrigation)
- Purchase price
- Graphical record (Photograph, visual record with scale)
- Transport conditions ( mode and conditions of transport)

# Sample collection

- ◉ Description of sample collected: after sample collection
  - Food type (Legume, fruit juice, milk product)
  - Local use of foods (Famine. Festivals)
  - State of food sample (solid, semisolid, viscous, or liquid)
  - Process and preservation methods (canned smoked)
  - Preparation method (cooking)
  - Extent of preparation (raw, fully cooked, reheated)

# Sample collection

- ◉ Description of sample collected: after sample collection
  - Extent of preparation (raw, fully cooked, reheated)
  - Packing medium (brine, oil)
  - Container or wrapping (can glass)
  - Contact surface (can , glass)
  - Label or list of ingredients (estimated by inspection)

# Sample collection

- ◉ Description of sample collected: after sample collection
  - Batch number
  - Weight of food collected/individual items
  - Number of items
  - Weight of common measure or portion

# Sample collection

## ○ Things to note

- Deliver samples to the laboratory promptly with the original conditions maintained as nearly as possible
- If products are in bulk: storage procedures, choice of containers, modes of transport should be considered
- Use containers that are clean, dry, leak-proof, wide-mouthed, sterile, and of a size suitable for samples of the product.

# Sample Transportation

## ○ Things to note

- Whenever possible, avoid glass containers, which may break
- For dry materials, use sterile metal boxes, cans, bags, or packets with suitable closures.
- Identify each sample unit (defined later) with a properly marked strip of masking tape.
- Transport frozen or refrigerated products in approved insulated containers of rigid construction

# Sample Handling

## ○ During Handling

- Aim: To protect the sample from changes in composition and contamination
  - Things to note
    - Weight and nature of edible/inedible matter (Prior to further processing (outer wilted leaves))
    - Method of preparation (Cooking or not, time, temperature of preparation)
    - Weight before/after cooking
    - Ingredients added if any

# Sample Handling

## ◉ During Handling

- Method of mixing and reduction (grinding, homogenization)
- Types of storage (addition of preservatives, temp of storage)
- Methods used of take analytical samples
- Storage of analytical samples or further processing
- Name and signature of person completing record
- Date of record
- Other details

# Sample Preparation

- Preparation of analytical portions
  - If the particle size or bulk is too large for analysis, it must be reduced in bulk or size for analysis
  - Documentation of sample preparation is very important
  - Separate edible/inedible portions, record descriptions and weigh all parts
  - Measure portion sizes, weights, volumes, density etc.

# Sample Preparation

## ○ Homogeneous foods

### ■ Solids:

- *Friable: crumble and mix.*
- *Sticky: freeze and crush at low temperature.*
- *Hygroscopic: take portions rapidly into pre-weighed sealable containers for weighing.*

## ○ Emulsions.

- Take by weight rather than volume; warm and mix.

## ○ Liquids with suspended solids.

- Homogenize, or sample during gentle mixing.

# Sample Preparation

## ○ Reduction by quartering

- The principle is that the quarter should be representative of the whole
- Any symmetrical food should be cut into quarters, and one-quarter of each batch taken for processing for analysis
- Large items, if symmetrical, can be reduced in size by this technique
- Oval or elongated foods (e.g. potato or cucumber) should be cut into eighths, and two-eighths taken for a quarter,

# Sample Preparation

## ○ Reduction by quartering

- Food lots of small items (flour, rice, legumes, small fruits, chopped mixed units).
  - The bulk is tipped into a uniform pile on a clean, inert surface
  - Turned over several times with a polythene or glass spatula.
  - The pile is leveled and then divided into four equal segments.
  - Two opposing segments are taken and the other two discarded.
  - The remaining segments are mixed and further reduced in the same way

# Sample Preparation

## ○ Reduction by quartering

- Foods consisting of fairly large, separate, but similar portions, such as loaves of bread or joints of a meat, should be quartered and sampled then processed for analysis.
- Segmented foods sampling e.g. packets of biscuits, cartons of eggs, batches of bread rolls.
  - Take every fourth item to form a composite sample.
  - For sliced loaves, take every fourth slice and one end slice, which then must be thoroughly crumbed before further reduction.

# Sample Preparation

## ○ Examples of analytical sample preparations

### ○ Nuts.

- Batches of nuts should be ground separately with a pestle and mortar, then mixed together thoroughly in a bowl.
- An analytical portion should be taken for inorganic analyses and the remaining mixture should be homogenized mechanically for further analyses.

### ○ Eggs:

- - Fresh. Fresh eggs should be shelled and mixed briskly with a fork; after analytical portions are taken for inorganic analyses, the remainder is homogenized mechanically.
- - *Dried. Dried eggs should be handled as flour.*

# Sample Preparation

## ○ Examples of analytical sample preparations

### ■ Fruit.

- Large fruits (e.g. pineapples or watermelons) and medium-sized ones (e.g. apples) must be quartered.
- Small fruits (e.g. cherries) should be quartered by the method used for particulate foods.
- Quarters should be coarsely chopped and combined, and unhomogenized analytical portions should be taken for immediate vitamin C and inorganic analyses.
- The remaining mixture can then be homogenized to produce an analytical sample for other analyses.

# Sample Preparation

## ○ Examples of analytical sample preparations

- Meats and fish (raw, cooked and processed).
  - The fat and muscle of some meats are more conveniently analysed separately and the results combined to produce the final values.
  - The edible portion of each unit is chopped coarsely with a sharp knife (fish is flaked with a fork) and mixed thoroughly in a bowl with a spatula.
  - A portion is removed, frozen and crushed in a polythene bag, and used for inorganic analyses.
  - The remainder of the analytical sample is minced and mixed thoroughly again; portions are taken for further analyses.
  - Care must be taken to avoid fat separation during mixing

# Sample Preparation

- **Examples of analytical sample preparations**
  - *Leafy vegetables and vegetable inflorescences.*
    - *Small leafy vegetables* should be mixed together in a bowl, chopped coarsely and mixed again briefly.
    - A large portion should be taken for inorganic analysis and another portion into metaphosphoric acid for vitamin C analysis.
    - Large tight-leaved vegetables (e.g. cabbage, iceberg lettuce) must be quartered.

# Sample Preparation

- **Examples of analytical sample preparations**
  - All large leafy vegetables must be chopped coarsely and mixed, and this must be done very quickly
  - After the mixing, analytical portions should be taken for analyses of vitamin C, vitamin A, carotenes, vitamin E and inorganic nutrients
  - The remainder can be chopped further. Stalks are often difficult to reduce and may have to be chopped separately and reintegrated into the food sample.

# Sample Preparation

- **Examples of analytical sample preparations**
  - Prepared composite foods and dishes.
    - This is the form in which most foods are consumed.
    - Items should be briefly homogenized, carefully mixed, then rehomogenized.
    - It can be assumed that laboratory homogenization will not introduce any contamination greater than that arising during domestic or commercial food preparation.

# Sample Preparation

- **Examples of analytical sample preparations**
  - Care is required to blend in the individual pieces of muscle, fat, vegetables, etc., which may be found in mixed prepared foods.
  - Portions for vitamin C assay are best taken from the mixed homogenate before it is rehomogenized.
  - If the prepared foods are hot, speed is essential to prevent moisture loss.
  - Total meals or diets can be handled in the same way.

# Sample Preparation

- ◉ Some practical equipment requirements for handling and preparation of laboratory and analytical samples
  - **General:**
    - Trays (for carrying foods)
    - Chopping boards (polythene, wood)
    - Oven thermometer, meat thermometer
    - Waring blender
    - Pestle and mortar
    - Ball mill
    - Hammer mill

# Sample Storage

- Keep ground samples in glass or plastic containers with air and water tight covers.
- Samples not analyzed immediately should be left in cold storage to minimise spoilage and other chemical reactions.
- Samples for lipid analysis - store under nitrogen at low temperature to prevent oxidation and unsaturated lipids

# Sample Storage

- ◉ Light may initiate oxidation so store in dark containers.
- ◉ For lipid analysis, antioxidants may be added if they wont interfere with the analysis
- ◉ It is therefore desirable to store a number of identical analytical samples
- ◉ Minimize the number of staff involved in taking portions from them.

# Sample Storage

Effects of sample storage and preparation on nutrient content and precautions required to minimize them

| Effects            | Potential Changes  | Nutrients Affected                             | Precaution   |
|--------------------|--|--|--|
| Drying out         | Loss of water  | All nutrients                                  | Design of protocol, Keep samples sealed, weigh food at start and during preservation       |
| Absorption         | Gain of water  | All nutrients                                  | Design of protocol, keep samples in sealed container                                       |
| Microbial activity | Degradation/ autolysis/ synthesis                        | Loss of CHO, proteins, gain in thiamin, Vit B6 | Storage at low temperature, pasteurization or addition of inhibitors                       |
| Oxidation          | Destruction of unsaturated fatty acids, loss of vitamins | Alterations in profile of fats                 | Store at -30C in sealed containers under nitrogen. Add antioxidants, bacteriostatic agents |
| Acid               | Hydrolysis   | Loss of sucrose and higher oligosaccharides    | Store at low temperatures<br>Neutralize acids  |

# Sample Storage

Effects of sample storage and preparation on nutrient content and precautions required to minimize them

| Effects                                       | Potential Changes                | Nutrients Affected                                      | Precaution   |
|---|----------------------------------|---|--|
| Alkaline                                      | Destruction                      | Loss of thiamine  | Avoid alkaline conditions and SO <sub>2</sub>                                |
| Light   | Photo degradation                | Loss of riboflavin                                      | Protect from light   |
| Contamination during sampling                 | From cooking vessels, soil, dust | Increase inorganic nutrients                            | Design protocol to minimize contamination, gently rinse with distilled water |
| Contamination from metallic blades, glassware | Increase in inorganic nutrients  | Increase in major trace elements                        | Select apparatus with care<br>Clean all utensils<br>Store in plastic bags    |
| Separation                                    | Separation of fats               | Changes in compositional<br>Alteration in fibre content | Avoid over vigorous mixing and thaw/freeze cycles                            |

# Sources of errors in sampling

- ⦿ It is essential that all those involved in the sampling process are familiar with the objectives of the work and are clear about their roles.
- ⦿ This will identify aspects that are unclear or impracticable and require modification to avoid errors.

# Sources of errors in sampling

## Major Sources of errors in sampling

| Source                        | Examples   | Precaution  |
|-------------------------------|--|---|
| Food sample identification    | Poor labeling of samples   | Maintenance of documentation throughout sampling and analytical process |
| Nature of sample              | Samples do not conform to the defined sampling protocol                      | Explicit instructions in sampling protocol, training of sample staff    |
| Transport and handling        | Samples contaminated, degraded or depleted during transport, loss of samples | Protocol specifies condition to be maintained, supervision              |
| Analytical sample preparation | Incorrect mixing or homogenization   | Proper supervision in laboratory<br>Laboratory quality assurance        |
| Analytical sample storage     | Incorrect storage of samples   | Proper laboratory techniques and supervision                            |

**Thank you**