GRAIN SAMPLING METHODS TO ACHIEVE CONSUMER CONFIDENCE AND FOOD SAFETY

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GRAIN SAMPLING METHODS TO ACHIEVE CONSUMER CONFIDENCE AND FOOD SAFETY

by

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ABSTRACT

Grain is sampled for one, possibly, two reasons:

- to determine the average quality based on a representative sample
- to detect contaminants. This information may trigger rejection of a load or determine whether the levels of a contaminant in a bulk exceed a regulatory threshold.

This review examined published sampling regimes for a number of contaminants that now, or in the future, may affect food safety. These were: genetically modified organisms, mycotoxins, micro-organisms, heavy metals, agrochemical residues and arthropod pests.

A common, ISO 13690-based approach using 5-11 samples from 15-500t is used for some contaminants. This is reasonable when the contaminant (analyte) is homogeneously distributed, although sample size and number of samples/tonne are often poorly defined, standards used may have no scientific basis and there are rarely validation details. Sampling regimes for insects and in particular, mycotoxins, which are heterogeneously distributed, are very demanding.

An EU directive recommends taking 100 samples of 100g from 50-1500t for regulatory purposes for mycotoxin determination. Live insect contamination is best estimated using traps but the regulatory mycotoxin sampling may be adaptable for sampling insects and would also apply to dead insects.

It is recommended that the ISO-based sampling method be validated experimentally for a variety of analytes since it is widely cited and convenient to use. However, its effectiveness is unclear. A simplified method of sampling for mycotoxins should be developed that will give results comparable to the regulatory sampling. This would demonstrate ‘due diligence’ and avoid costs associated with the ‘regulatory’ sampling protocols. (If such a method proved ‘substantially equivalent’ to the regulatory sampling, it could be used for regulatory processes if there were cost savings). The pattern of mixing grain during outloading has considerable influence on the distribution of analytes. It therefore affects the sampling regime and merits study.
INTRODUCTION

The primary aim of the HGCA ‘Grain Sampling and Analysis Project’ (GSAP) is to develop and validate sampling protocols that are designed to test end-user specification requirements. The analytical harmonisation element of the project also focuses on these tests. Although these approaches should assist both seller and buyer of grain, they do not directly address consumer confidence and food safety. These concerns are currently driven by legislation and consumer pressure groups.

Because of these concerns and the way in which minor contaminants are distributed in raw agricultural products it is generally necessary to sample differently to obtain representative samples for a range of contaminants. Further homogenisation and sub-sampling is required to provide a meaningful analysts’ sample and the size of the sub-sample may vary with the analyte contaminant.

The main contaminants include:

- Genetically modified organisms
- Mycotoxins of field and storage origin
- Micro-organisms
- Heavy metals
- Agrochemical residues (including storage pesticides)
- Pests (including mites).

This desk study reviews relevant sampling protocols published in the UK and elsewhere. It analyses their scientific basis and comments on their validation, where any exist. The review concentrates on cereals (wheat, barley etc.) but, where appropriate (e.g. with GM oilseeds), reference is made to other commodities.

Where there are similarities between sampling requirements (based on sound science) for various analytes, their appropriateness with other analytes is considered. The aim is to identify as few sampling protocols as possible to deliver robust test data for the analytes considered. In addition, the potential and relevance of existing ‘quality’ sampling protocols to food safety testing is considered.

The review focuses on sampling in permanent storage situations, e.g. farm or co-operative storage (both flat and bin storage) and intake at buyers’ premises, where appropriate, defining lot size. Sampling of vessels is also considered to ensure coverage of conditions affecting imports and exports. Attention is also paid to the appropriateness of specifications for blending (homogenisation) and sub-sampling, to provide an analyst’s sample.
The review’s purpose is to recommend practical sampling procedures for surveillance testing of any or all the items listed. It also highlights areas where further information and experimentation is needed to develop or validate new protocols.

APPROACHES

This review commenced with a survey of BSI and ISO standards applying to grain sampling. Meetings with CSL experts in mycotoxins, GMOs, residues and pests produced much background material and provided valuable insights into protocols under development, as well as published information. Additional documents and information were sought using web searches which enabled access to many documents, e.g. EU directives and CODEX. Where information was particularly lacking, such as sampling to detect bacteriological contamination, conventional library literature reviews were conducted.

Most methodologies (e.g. CODEX, ISO) do not reference the sources or science behind protocols and rarely the expert committees, who agree them, so it is difficult to evaluate the appropriateness of recommendations.

SAMPLE / SAMPLING DEFINITIONS

Lot- An identified amount of grain (offered by an applicant for inspection). An identified quantity of material having characteristics presumed to be relatively uniform. This presumption may not be so for all the analytes covered by this review

Increment- A quantity of material taken at one time from a single point in a lot. Also known as ‘primary sample’.

Composite sample- A single sample composed of small portions taken from throughout a lot. Also known as bulk sample. A quantity of material obtained by combining and mixing all increments in the same lot.

Reduced sample- A representative part of the bulk sample obtained by the process of reduction in such a manner that the mass or volume approximates to the laboratory samples. Also known as ‘submitted sample’.

Laboratory sample- A sample representative of the quality and condition of the lot, obtained by reduction of the bulk sample and intended for analysis or other examination. Also known as ‘working sample’.
1. BACKGROUND TO SAMPLING

Grain is sampled for one, possibly, two reasons:

- to determine the average quality based on a representative sample
- to detect contaminants. This information may trigger rejection of a load or determine whether the levels of a contaminant in a bulk exceed a regulatory threshold.

The sampling regime adopted will depend on the aim. For instance, if a representative sample is required then a random or systematic sampling plan may be most appropriate but if the presence of an undesired quality such as a pest, for which there is no tolerance, is to be established, then the sampling plan will need to concentrate on the most likely position for the particular analyte.

Many standard reference works on sampling make no mention that different analytes may require different sampling regimes. Examples of these are the GIPSA (Grain Inspection, Packers and Stockyards Administration) Grain inspection handbook\(^\text{1.1}\) and the often quoted ISO 950, now superseded by ISO 13690\(^\text{1.2}\).

This review aims to identify one or more approaches to taking a representative sample for the analytes listed in the introduction.

Sampling protocols

GIPSA\(^\text{1.1}\), United States Department of Agriculture (USDA) discusses the mechanics of sampling and gives sampling plans for barges, trucks, hopper cars but which would apply equally to similar sized static bins. It suggests an acceptable sampling ration of 1250 g from 5,000 bushels (ca. 146t of wheat) of grain and that when containers contain inferior portions (e.g. musty, sour or heating) three separate samples must be drawn (one from the inferior portion, one from the entire lot and one from the remainder of the lot) producing a minimum of 2kg. The purpose of the sampling so described, whether for regulatory or market requirements, is not discussed.

International Standards Organisation (ISO) 13690\(^\text{1.2}\) covers sampling of cereals from bags, from wagons, lorries, barges or ships and from silos, bins and warehouses. This suggests patterns of sampling (indicating position of sampling columns) from lorries etc., taking a minimum of 5 - 11 samples throughout the whole depth of the lot from lots varying from 15 – 500t (Table 1). It suggests a similar pattern should be used when sampling bulk grain. It recommends determining the number of increments by dividing the square root of the tonnage by two and rounding to the nearest whole number. Thus 12 samples would be taken from a 500t lot and 50 from 10,000t.
Table 1. Number of increments to be taken from large grain bulks (after ISO 13690\textsuperscript{1,2})

<table>
<thead>
<tr>
<th>Tonnage</th>
<th>500</th>
<th>1000</th>
<th>2000</th>
<th>4000</th>
<th>6000</th>
<th>8000</th>
<th>10000</th>
</tr>
</thead>
<tbody>
<tr>
<td>Increments</td>
<td>12</td>
<td>16</td>
<td>23</td>
<td>32</td>
<td>39</td>
<td>45</td>
<td>50</td>
</tr>
</tbody>
</table>

American Association of Cereal Chemists (AACC) Method 64-70\textsuperscript{1,3} refers to manual sampling of grains. This suggests sampling plans but does not recommend sampling rates, other than for moving streams where a suggestion of one sample of unspecified size from every 500 bushels (ca. 14.6t wheat) is made.

British Standard (BS) ISO 6644\textsuperscript{1,4} refers to automatic sampling of flowing grains but mentions no specific sample size or sampling rate. AACC Method 64-71\textsuperscript{1,5} recommends diverter sampler activation rates of once per 5.5t.

International Seed Testing Association (ISTA) (Bould, 1986)\textsuperscript{1,6} gives sampling frequencies for bags and other small containers, ranging from each container being sampled at 5 positions when there are 1-5 containers in the lot, to at least one in seven being sampled when there are more than 560 (Table 2). Sampling frequencies recommended for seed (including cereals) loose in the bulk vary from less than three from bulks up to 50kg, to one from every 700 kg from bulks of 28 – 40t. Guidance is also given relating size of primary samples taken by automatic samplers to the time taken to draw the sample, depending on flow rate.

Table 2. Sampling frequency for loose seed (after ISTA\textsuperscript{1,6})

<table>
<thead>
<tr>
<th>Maximum Lot size (kg)</th>
<th>50</th>
<th>1500</th>
<th>3000</th>
<th>5000</th>
<th>20000</th>
<th>28000</th>
<th>40000</th>
</tr>
</thead>
<tbody>
<tr>
<td>Min. No. samples</td>
<td>3</td>
<td>5</td>
<td>1/300kg</td>
<td>10</td>
<td>1/500kg</td>
<td>40</td>
<td>1/700kg</td>
</tr>
</tbody>
</table>

Blending and sub-sampling

ISO 13690\textsuperscript{1,2} gives a number of alternative methods of producing laboratory samples from the thoroughly mixed bulk sample. These include coning and quartering and dividers which may be multi-slot, conical or centrifugal. The number and size of the laboratory sample is determined by the requirements of the test but a general recommendation is made of a 1kg minimum size.

ISTA\textsuperscript{1,6} (Bould, 1986) discusses the same methods of producing a laboratory sample as ISO but additionally mentions the use of random cups, spoon and hand sampling which is restricted to very chaffy seeds, and a
rotary divider (for cleaner samples). ISTA also discusses the production of working samples, based on the number of seeds and thus on a pure seed content. This will affect certain parameters of the sample since the admixture component has a higher pesticide content and a more numerous microbiological component (Armitage et al., 1996).

**Validation / Principles**

The Codex Alimentarius Commission was created to develop food standards, guidelines and codes of practice under the Joint FAO/WHO Food Standards Programme. It publishes guidelines for sampling plans for food tested for compliance with particular Codex commodity standards (Codex, 2003). This lists 13 ISO standards which form the basis of the procedures discussed. It covers selection of sampling plans including those for the inspection of bulk material but the scope does not cover control of non-homogenous goods.

Drew et al. (1978) give tables of sample numbers required to give average quality of a lot to four levels of desired precision, based on data from American Society for Testing Material (ASTM) E122-58 (Table 3). Their approach was theoretical and intended to apply to a number of quality analytes, particularly in the context of surveys.

**Table 3. Number of sampling units required to achieve a given degree of precision (after Drew et al., 1978)**

<table>
<thead>
<tr>
<th>Range of results expected (%)</th>
<th>100</th>
<th>80</th>
<th>60</th>
<th>50</th>
<th>40</th>
<th>30</th>
<th>30</th>
<th>10</th>
<th>5</th>
</tr>
</thead>
<tbody>
<tr>
<td>Precision (+/-)</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>1 %</td>
<td>5625</td>
<td>3600</td>
<td>2025</td>
<td>1406</td>
<td>900</td>
<td>506</td>
<td>225</td>
<td>54</td>
<td>14</td>
</tr>
<tr>
<td>2 %</td>
<td>1406</td>
<td>900</td>
<td>507</td>
<td>351</td>
<td>225</td>
<td>126</td>
<td>57</td>
<td>14</td>
<td>4</td>
</tr>
<tr>
<td>5 %</td>
<td>225</td>
<td>144</td>
<td>81</td>
<td>56</td>
<td>36</td>
<td>20</td>
<td>9</td>
<td>2</td>
<td>-</td>
</tr>
<tr>
<td>10 %</td>
<td>57</td>
<td>36</td>
<td>21</td>
<td>14</td>
<td>9</td>
<td>5</td>
<td>3</td>
<td>-</td>
<td>-</td>
</tr>
</tbody>
</table>

Sellar (1994) gives the sample size required to detect different levels of infection for different probability levels (Table 4) based on a binomial probability distribution (appropriate for large consignments, such as grain bulks). Thus taking a sample of 300 items would give a 95% probability of detecting a 1% infection (or you would fail to detect it 5 times out of one hundred). The intention here was to detect the presence of undesirable contaminants, such as disease or GM seeds, rather than to determine ‘average’ quality.
Table 4. Levels of infection detected (at least one infected item found) with different sample sizes (after Sellar 1994)\(^{1.10}\)

<table>
<thead>
<tr>
<th>Sample size</th>
<th>50</th>
<th>100</th>
<th>150</th>
<th>300</th>
<th>500</th>
<th>1000</th>
<th>3000</th>
</tr>
</thead>
<tbody>
<tr>
<td>95% Probability</td>
<td>6.0%</td>
<td>3.0%</td>
<td>2.0%</td>
<td>1.0%</td>
<td>0.6%</td>
<td>0.3%</td>
<td>0.1%</td>
</tr>
<tr>
<td>99% probability</td>
<td>9.0%</td>
<td>4.6%</td>
<td>3.0%</td>
<td>1.5%</td>
<td>0.9%</td>
<td>0.46%</td>
<td>0.15%</td>
</tr>
</tbody>
</table>

ISTA\(^{1.6}\) discuss segregation patterns likely to occur, even in homogeneous bulks, as a result of filling containers and during transport. A cone of seed is likely to form at the bottom of bags and heavier seeds roll down the cone and to the edge while smaller seeds tend to remain in the centre (Hibbert and Woodwark, 1959\(^{1.11}\)). Chaff is picked up by upward air currents and ends up at the sides of bags (Debney, 1960\(^{1.12}\)) while weed seed (the example was red clover) is higher toward the centre. The tendency of seed to pick up moisture from the atmosphere at the edges and thus lose germinative capacity is also discussed.

Discussion

In many ways, the generalised protocols for sampling reflect the overall problems of the more specific protocols which assume that a sample taken a certain way will be suitable for determination of all qualities or analytes. This is most unlikely and certainly unproven. Associated statistics often assume homogeneity of the quality examined that is not validated with practical studies. The ISTA documents\(^ {1.6}\) highlight one particular issue which is the segregation of grain based on seed weight during conveying. This little-considered issue could affect sampling static bulks both for quality such as specific weight or seed variety but would also affect issues such as pest distribution and/or fungal content and mycotoxins which are often higher in the fraction which contains chaff and small seeds.

References

1.1 USDA GIPSA (Grain Inspection, Packers and Stockyards Administration) Grain inspection handbook – Book 1 Sampling.

http://www.ciiitltd.com/grain_inspection.htm


1.4 British Standards BS ISO 6644. 2002. Flowing cereals and milled cereal products – Automatic sampling by mechanical means. 9pp

http://www.aaccnet.org/ApprovedMethods/toct.htm


2. GENETICALLY MODIFIED ORGANISMS (GMOs)

Currently the GM commodities likely to be encountered in trade are mainly maize (corn), rapeseed and soya but not wheat or barley. Sampling for GM material may be to confirm that a GM bulk is of the specified trait, to determine contamination of a GM bulk by another GM trait or to detect contamination of conventional bulk grain by GM material.

Regulatory levels

Kay and Paoletti (2001)\(^2\) indicate three EU legislative tools:

1. EU directive 2001/18 which requires that GMO content in raw material must be indicated (shown).
2. Regulation 258/97 that requires that the content of GM material in the primary ingredient or final food product must be indicated
3. Regulation 1139/98 requires that adventitious presence of certain GM material must be indicated above a given threshold.

Sampling protocols

The purpose of sampling for GMOs is usually to detect and reject the contaminant. A draft standard operating procedure (SOP) produced on behalf of the European Enforcement Project (Reed, 2002)\(^2\) suggests a sampling regime (based on ISO 950), superseded by ISO 13690. Samples are taken from 5 –11 points for lorries of 15-50t to total 1kg from each load for rapeseed or twice this for soya or maize. Samples from ships holds, docksides, stores or silos should be from lots of up to 500t. The suggested rate from moving grain (ISO 950) is 1kg/100t for cereals (wheat and barley) but double this for maize and soya. For static bulks the ISO 950 approach is to take 10 samples totalling about 5kg per 100t from different depths and positions.

Kay and Paoletti (2001)\(^2\) suggest sampling should take place according to ISO 13690 (cereals) or ISO 542 (rapeseed) for static grain or ISO 6644 for flowing grain. The bulk sample should comprise 30 incremental samples and be >20 times larger than the laboratory sample although non-uniform GM distribution demands an increase in the number of incremental samples. No documents describe the error associated with the creation of the bulk sample but it is assumed to be negligible compared with the creation of the laboratory sample (see table below).

USFDA\(^2\) give industry guidelines for sampling corn for Cry9C protein residues from ‘Starlink’ GM corn. These recommend taking a sample of at least 2400 kernels from each vehicle or carrier using GIPSA recommended protocols.
Blending and sub-sampling

The European draft SOP\textsuperscript{2,2} suggests a working sub-sample of a minimum 3000 seeds prepared to ISTA methods.

Kay and Paolleti (2001)\textsuperscript{2,1} note this should be achieved by random sampling produced by thorough mixing of the bulk sampling followed by division into halves, quarters, eighths etc for instance using sample dividers. They show the number of grains required for a laboratory sample to detect GMO percentages for probability levels of 90, 95 and 99% based on binomial distribution (Table 5).

\textit{Table 5. Upper limit of GMO content (%) for selected range of sample sizes and probability levels (after Kay and Paolleti\textsuperscript{2,1})}

\begin{tabular}{lccc}
  No Grains & 90\% & 95\% & 99\% \\
  \hline
  100 & 2.28 & 2.95 & 4.50 \\
  200 & 1.14 & 1.49 & 2.28 \\
  300 & 0.76 & 0.99 & 1.52 \\
  400 & 0.57 & 0.75 & 1.14 \\
  800 & 0.29 & 0.37 & 0.57 \\
  1200 & 0.19 & 0.25 & 0.38 \\
  2000 & 0.12 & 0.15 & 0.23 \\
  2500 & 0.09 & 0.12 & 0.18 \\
  3000 & 0.08 & 0.10 & 0.15 \\
  6000 & 0.04 & 0.05 & 0.08 \\
  10000 & 0.02 & 0.03 & 0.05 \\
  \hline
\end{tabular}

Validation

A USDA GIPSA website (2000)\textsuperscript{2,4} discusses the theory of sampling for the detection of ‘Biotech’ grains, suggesting that to detect Biotech samples of 0.1-5% would require sample sizes between 881 and 18g for 95% success or 27-1354g for 99% success (Table 6).
Table 6. Sample sizes such that lots containing given concentration levels in corn are detected (and rejected) 95% or 99% of the time\textsuperscript{2,4}.

| Biotech. Concentration (%) | No. kernels |   | Approx. wt (g) |   |
|---------------------------|-------------|-----------------|-------------|
|                           | 95%         | 99%             | 95%         | 99%         |
| 0.1                       | 2995        | 4603            | 881         | 1354        |
| 0.5                       | 598         | 919             | 176         | 271         |
| 1.0                       | 299         | 459             | 88          | 135         |
| 2.0                       | 149         | 228             | 44          | 68          |
| 3.0                       | 99          | 152             | 30          | 45          |
| 4.0                       | 74          | 113             | 22          | 34          |
| 5.0                       | 59          | 90              | 18          | 27          |

Guidelines from various EU and USA sources for detection of GM are summarised in a draft document by Paoletti et al. (2002)\textsuperscript{2,5} which argues that most of these are based on the false assumption of homogeneity or random distribution. This being so, a validation protocol was proposed to identify appropriate sampling strategies by simulation studies and by testing of these strategies based on data from EU member states. An interesting detail of the proposal is the necessity of developing a software tool to test various sampling strategies based on different population structures. This approach would have benefit elsewhere but would of course depend on existing understanding of the population structures. In most cases, these would have to be derived by observation and experiment.

Discussion

The details of sampling for GMOs detailed here indicate the ideal approach to sampling for this particular analyte. A draft EU protocol has been drawn up and an experimental protocol for validation of this has been proposed. It is to be hoped, but is by no means certain, that EU funding routes will enable completion of this project. This sampling is intended for detection of contamination rather than determination (of average quality) and may therefore require a different approach than for most analytes (with the possible exception of insects).

References

2.2 Reed, P. 2002. Protocol for bulk grain sampling and testing for GM presence. Draft SOP produced on behalf of European Enforcement Project. CSL, York, UK. 8pp

2.3 Anon, 2001. FDA recommendations for sampling and testing yellow corn and dry-milled yellow corn shipments intended for human food use for Cry9C protein residues USFDA, Centre for Food Safety and applied nutrition, 8pp
http://vm.cfsan.fda.gov/~dms/starguid.html

2.4 Anon, 2000. Sampling for the detection of Biotech grains.
http://www.usda.gov/gipsa/biotech/sample1.htm

2.5 Paoletti, C., Donatelli, M., Kay, S., van den Ede G. 2002. Simulating kernel lot sampling; the effect of heterogeneity on the detection of GMO contaminations. Seed Science and Technology.
3. MYCOTOXINS

Mycotoxins are fungal metabolites produced to give certain species of fungi competitive advantage over others. Unfortunately some have been proven to have ill effects on human health and also genotoxic carcinogens. Their production is usually related to certain moisture and temperature conditions, related to the growth requirements of different species. Mycotoxins in cereals have two origins, either being formed in the field during the growth or in store. The former include fumonisins, trichothecenes (e.g. deoxynivalenol) and zearalenone and are derived from *Fusarium* spp. Although there is conflicting opinion, it is generally accepted that these are not usually produced in storage, except perhaps during advanced decay. (However, propionic acid treatment of damp grain may select for *Fusarium* spp. according to Burrell et al, 1983) During storage, different fungi predominate and the main concerns then centre on toxins produce by the relatively xerophilic fungal genera that prefer lower moisture contents; *Aspergillus* (*Eurotium*) and *Penicillium* which may produce aflatoxins and ochratoxins respectively. It is the latter group that are currently of greatest concern in the UK since temperatures and moisture contents here do not favour the former. It is generally considered (eg Cahagnier et al, in press) that the mycotoxins are formed during mycelial growth and may be related to ergosterol production. It has also been noted that ochratoxin A production rarely occurs when colony counts, which are mainly conidial, are below 1,000 per g (Olsen, National Food Administration, Sweden, personal communication).

The origin of mycotoxin formation will naturally have an effect on the sampling process. It may be assumed that field mycotoxins may be mixed in transport and during conveying as the grain goes into store and therefore may be relatively homogeneous. On the other hand, storage mycotoxins are likely to be locally distributed and will only be formed when the grain is stored for some time above the threshold for growth of the mycotoxin-producing fungi. This is only likely to occur on un-dried grain during harvest backlogs and during the ambient-air drying process when grain at the top of the bin remains close to its original moisture content until the drying front passes through. In this case, the mycotoxins will be primarily at the top of the bin (or floor store) if the grain is not ‘turned’ after drying.

**Regulatory levels**

A CSL internal database lists current regulatory levels of numerous mycotoxins on specified commodities and will be widely available soon.


EU regulatory levels for the storage mycotoxins are quite simple: 4µg/kg total aflatoxins of which aflatoxin B1 must not exceed 2µg/kg in cereals (including buckwheat, *Fagopyrum* sp.), with the exception of maize, to be subjected to a sorting, or other physical treatment before their human consumption or their use as an ingredient in foodstuffs (Commission Regulation (EC) No. 466/2001).
5 µg/kg ochratoxin A (OA) in raw cereal grains (including raw rice and buckwheat) (Commission Regulation (EC) No. 472/2002)\(^{1,4}\) (amending Commission regulation (EC) No 466/2001)

However, harmonised EU regulatory levels do not currently exist for the field mycotoxins. For other individual countries they vary widely depending on consumer (e.g. infants, adults, animal species) and commodity.

**Sampling protocols**

For OA, EU Commission directive 2002/26/EC\(^{3,5}\) recommends 100 samples of 100g be taken from bulks between 50 and 1500t resulting in aggregate samples of 10kg (Table 7). Similar guidelines are laid down for sampling for aflatoxins\(^{3,6}\).

The sampling and analysis directives which accompany the regulations are addressed to enforcement agencies (i.e. local authorities and port health authorities) to enable them to enforce the regulations adequately. These agencies are required to use these sampling regimes but there is no direct requirement for others to do so. How others ensure that the commodities they produce and sell comply with the regulations is a matter for them to determine and to provide evidence of in their due diligence defence if an official control sample finds them wanting. The official sampling regime is onerous and the directives state that it is acceptable as a defence if an alternative sampling regime is used that can be shown to be "substantially equivalent" to the official one.

**Table 7. No. of incremental samples to be taken for OA determination depending on lot size (after CD 2002/26/EC\(^{3,5}\))**

<table>
<thead>
<tr>
<th>Lot wt (t)</th>
<th>Wt or no of sub-lots</th>
<th>No of incremental samples</th>
<th>Aggregate sample weight (kg)</th>
</tr>
</thead>
<tbody>
<tr>
<td>≥ 1500</td>
<td>500 t</td>
<td>100</td>
<td>10</td>
</tr>
<tr>
<td>&gt;300 &lt;1500</td>
<td>3 sub-lots</td>
<td>100</td>
<td>10</td>
</tr>
<tr>
<td>≥ 50 ≤ 300</td>
<td>100 t</td>
<td>100</td>
<td>10</td>
</tr>
<tr>
<td>&gt;20- &lt;50</td>
<td></td>
<td>100</td>
<td>10</td>
</tr>
<tr>
<td>&gt;10 ≤ 20</td>
<td></td>
<td>60</td>
<td>6</td>
</tr>
<tr>
<td>&gt;3 - ≤10</td>
<td></td>
<td>40</td>
<td>4</td>
</tr>
<tr>
<td>&gt;1 - ≤3</td>
<td></td>
<td>20</td>
<td>2</td>
</tr>
<tr>
<td>≤ 1</td>
<td></td>
<td>10</td>
<td>1</td>
</tr>
</tbody>
</table>
USFDA recommendations\textsuperscript{3,7} on mycotoxin sample sizes suggest merely 10 samples of 1lb (0.454kg) to be taken from bulks of small grains. A GIPSA document\textsuperscript{3,8} on mycotoxin/aflatoxin sampling notes that optimum sample size would be 10lbs but established reduced sizes of 2 and 3lbs from trucks and rail cars in response to industry concerns of increased costs of inspection. Sampling patterns indicated are similar to those in ISO 13690.

American Association of Cereal Chemists (AACC) methodology\textsuperscript{3,9} notes that the USFDA has determined that one infected kernel of corn containing 100,000µg/kg aflatoxin may bring a sample of 1500g (ca. 5000 kernels) to the FDA action level.

Recommendations for sampling wheat to determine deoxynivalenol (DON) levels produced by \textit{Fusarium graminearum} are to be found in the Michigan State University extension material (Hart and Schenberger 1997\textsuperscript{3,10}). These suggest probe sample of 1-2lbs (0.45-0.9kg) be taken from random positions or at random times during stream sampling.

\textbf{Blending and sub-sampling}

The EU requirement\textsuperscript{3,5} is that the entire aggregate sample should be ground using a process demonstrated to achieve complete homogenisation and the replicate laboratory/working sample be taken from the homogenised material. Three replicate samples are taken for enforcement purposes by the trade (defence) referee and compliance judged based on analysis of one enforcement sample. The size of samples is not given. GIPSA\textsuperscript{3,8} recommends a minimum 200g sample which is ground and from this a 50g sub-sample is obtained. In view of the EU requirement and the supporting research, this 200g sample would appear to be rather inadequate.

Michigan Sate DON recommendations (Hart and Schenberger, 1997\textsuperscript{3,10}) are that 5-15\% of the total sample be milled and used for analysis.

\textbf{Validation}

A computer programme ‘Mycotox’ developed by RHM Technology for MAFF (now DEFRA) produces sampling plans for a selected lot acceptance mycotoxin level. It gives three choices of sampling plan: ‘routine’ allowing the user to select 4-100 incremental samples, and ‘pragmatic’ which has 6 incremental samples and an upper limit of a 10kg aggregate sample, suitable for current industrial equipment. The final, ‘regulatory’ plan sets the number of samples based on the load size and thus requires a larger sample weight than the equivalent ‘routine’ plan. Although this appears a useful tool, it does not seem to have been validated by experiment.
A critical study (Wilson et al., 1999) compared the suitability of four aflatoxin sampling plans to detect OA by sampling coffee and wheat. These were the USDA plan, the Dutch code of practice and the UK plan for groundnuts and the EC plan for aflatoxin. Only the EC sampling plan for aflatoxins, based on taking 20-100 incremental samples of 30g, depending on lot size, produced representative samples for OA in wheat.

Coker et al (1998) validated sampling plans for aflatoxin B1 in animal feedstuffs but noted that this was based on a reasonably homogeneous distribution of the toxin and should not be applied to whole commodities where a highly skewed distribution pattern was to be expected. They assessed 10 sampling plans based on 1-8 composite samples each comprising 10 or 20 25g incremental samples. The coefficient of variation for aflatoxin B1 varied between 4.0 and 12.5%.

GIPSA notes that, since a single kernel of corn may contain 400,000µg/kg of DON the accuracy of a representative sample is critical. Only 1-3% of kernels may contain mycotoxin and these are not evenly distributed. They cite data to show how the variability increases with decreasing sample size. For a truckload of wheat containing 20µg/kg aflatoxin, variability falls from 1-46.9 for 1lb (0.45kg) samples, to 11.6-28.4 for 10lb (4.5kg) samples. They note that a 1998 study indicates that increasing sample size did not decrease the variability of DON results and refer to a Michigan State University study (Hart and Schenberger, 1997) indicating variability was significantly higher if less than four probes were taken per truck (Table 8).

<table>
<thead>
<tr>
<th>Sample size (lbs)</th>
<th>10</th>
<th>5</th>
<th>2.5</th>
<th>1</th>
</tr>
</thead>
<tbody>
<tr>
<td>Kernels</td>
<td>30000</td>
<td>15000</td>
<td>7500</td>
<td>3000</td>
</tr>
<tr>
<td>Variability (µg/kg)</td>
<td>11.6-28.4</td>
<td>8.1-31.9</td>
<td>3.2-38.8</td>
<td>1-46.9</td>
</tr>
</tbody>
</table>

Hart and Schenberger (1997) show the relationship between the number of sample probes and the one or two sided boundaries for DON concentration at the 95% confidence level (Table 9). For instance, to be within 0.86-1.0ppm of the upper limit of the mean DON level (1 sided) would require analysis of four probes but to be within 1.0ppm either side of the mean would require analysis of five probes (2 sided). This was based on analysis in an epidemic year.
Table 9. The relationship between number of samples required and one or two sided boundaries at the 95% confidence level.

<table>
<thead>
<tr>
<th>No. Probes</th>
<th>Sub-sample of milled wheat</th>
<th>Whole sample of milled wheat</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>95% 2 sided</td>
<td>95% 1 sided</td>
</tr>
<tr>
<td>2</td>
<td>15.30</td>
<td>3.80</td>
</tr>
<tr>
<td>3</td>
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<td>1.43</td>
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<tr>
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<td>2.11</td>
<td>0.81</td>
</tr>
<tr>
<td>6</td>
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<td>0.70</td>
</tr>
<tr>
<td>7</td>
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<td>0.63</td>
</tr>
<tr>
<td>8</td>
<td>1.42</td>
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</tr>
<tr>
<td>9</td>
<td>1.31</td>
<td>0.53</td>
</tr>
<tr>
<td>10</td>
<td>1.22</td>
<td>0.49</td>
</tr>
<tr>
<td>12</td>
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<tr>
<td>14</td>
<td>0.98</td>
<td>0.40</td>
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<tr>
<td>16</td>
<td>0.91</td>
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<td>18</td>
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<td>0.21</td>
</tr>
<tr>
<td>50</td>
<td>0.48</td>
<td>0.20</td>
</tr>
</tbody>
</table>

http://www.cips.msu.edu/people/hartp/vomitoxin.pdf

Discussion

The situation with sampling for the storage mycotoxins, ochratoxin A and the aflatoxins indicates one extreme solution - a thorough labour-intensive plan which does not, as yet, have any full scale associated validation experiments. The options of compliance, benefits and costs are discussed by the UK Food Standards Agency (FSA). There it is noted that there is no requirement for businesses to carry out additional sampling other than the need to observe ‘due diligence’, discussed above. Costs may include cost of sampling and analysis, additional staff, purchasing additional material to replace rejected consignments,
price premiums to cover cost of rejections, additional port charges. An example shows re-occurring costs of £2-3,000 for a typical miller. However, in view of these costs, it may be worthwhile considering the necessity of such intensive sampling of bulks of grain that have been harvested below the critical moisture content for storage mycotoxin production. Some difficulties in sampling are indicated above. The homogeneity of toxins in animal feedstuffs, presumably a result of mixing and formulation, are not reflected by the distribution in the raw products such as grain. If experience with the field mycotoxin, DON is anything to go by, storage mycotoxins are likely to originate in one or two individual, possible dead or weakened grains which makes them very difficult to detect when sampling static bulks. The origin of storage mycotoxins has still not been decisively located but since they depend on relatively high moisture contents, there are only two likely situations where they can occur. The first is in harvest backlogs where damp grain is left for some time before being passed through a hot-air dryer, the other is during ambient-air drying when a drying front passes through the grain from the bottom up and thus the highest mycotoxin risk will be found in un-dried grain toward the top of the bulk. Both situations will require a different sampling regime since in the first case, the highest level is likely to occur in the centre of the heating bulk of damp grain and in the second, highest toxin levels will occur toward the surface of the grain bulk. It could be argued that, while looking for likely areas of OA occurrence may be a health safeguard, it will be counter to producing a representative sample. The question to be resolved is whether sampling in this context should be detection or determination.

References


3.7 Anon, Chapter 4- Chart 6 Mycotoxin sample sizes. Investigations Operations manual USFDA Office of regulatory affairs.

3.8 Anon. Grain fungal diseases and mycotoxin reference. Chapter 4 pp 24-30 sampling.

3.9 American Association of Cereal Chemists. AACC Method 45-01. Sampling grain for mycotoxins, 1 p

http://msue/msu/edu/msue/imp/modlab/26309701.html


http://www.food.gov.uk/foodindustry/regulation/ria/71631
4. MICRO-ORGANISMS

This term covers both fungi and bacteria. Neither will grow under normal storage conditions; the former requires a minimum rh of 65% while bacteria require over 90% rh to flourish. The main sources of contamination from bacteria are from vertebrate faeces or from soil contamination of the grain and grain store, for instance from soil picked up during harvesting and the tyres of vehicles that drive into the grain store during loading or unloading.

Regulatory levels

In the UK, MAFF (now DEFRA) in the codes of practice for salmonella relating to feedstuffs\(^{4.1}\), note that MAFF, the storekeeper, vendor and purchaser are notified of isolation of salmonella, the occurrence should be investigated and curative procedures instituted. The codes of practice note that the Zoonoses order of 1989 requires that all isolations of salmonella from feedstuffs and ingredients be reported to MAFF.

Many intake inspection procedures, particularly of cereals for human consumption, e.g. the malting and milling industries, reject samples with rodent or bird droppings, presumably because of the assumption of bacterial contamination.

Sampling protocols

The salmonella codes of practice\(^{4.1}\) refer to microbiological monitoring of raw materials in the production of feedstuffs. In the case of storage, handling and transport of raw materials for feedstuffs, owners of raw material entering stores (other than intervention or on-farm) should collect aggregate samples over a period of one month or less. For imports, this sampling occurs where the cargo discharges. The frequency should vary in accordance with the risk factor. For products despatched from stores, it mentions that on each day material is despatched, at least one sample should be taken for aggregation and that these aggregate samples should contain no more than 40 incremental samples collected over a one month period.

The USFDA in their Salmonella Sampling Plan\(^{4.2}\) categorises whole grain as a food item that would normally be subjected to a process lethal to Salmonella between sampling and consumption. It suggests taking 15 sub-samples of at least 100g based on their categorisation of commodities but it is clear from the comments (e.g. ‘The usual sub-sample is a consumer size container of a product) that their main concern is grain as human food. It comments additionally that ‘more than one sub-sample may be collected from large institutional or bulk containers when the number of sub-samples required exceeds the number of containers in the lot’ (see definitions).
The International Commission on Microbiological Specifications for foods\textsuperscript{4.3} also comment on sampling plans for cereals and cereal products. This discusses field and storage fungi, Salmonellae and \textit{Bacillus cereus} as potential contaminants but despite the title do not discuss any details of sampling cereals for these contaminants.

\textbf{Blending and sub-sampling}

Nothing found

\textbf{Validation}

Nothing found

\textbf{Discussion}

Only the Salmonella code of practice contains sampling recommendations appropriate to cereal storage. The description of a single sample of unknown size to be taken for aggregation into a monthly composite seems unlikely to be adequate in view of the need to investigate the occurrence of Salmonella, not least because it will not be possible to identify the source (e.g. bin within the store).

\textbf{References}

\textsuperscript{4.1} Anon 1995. Codes of practice for the control of salmonella During the storage, handling and transport of raw materials intended for incorporation into, or direct use as animal feedingstuffs. MAFF publications, London.19 pp.

\textsuperscript{4.2} Anon, Chapter 4- Chart 1 Salmonella sampling plan. Investigations Operations manual USFDA Office of regulatory affairs.

5. HEAVY METALS

Heavy metals in soil may be of natural occurrence or via fertilisers, sewage or manure application or even atmospheric deposition. In recent years, contamination of soil has increased due to the application of biosolids and this can lead to uptake by the growing crops and transmission into the human food chain through consumption of cereals and their products.

**Regulatory levels**

A draft EU regulation has proposed concentration limits of 0.2mg/kg for lead and cadmium in wheat and lead in barley but only 0.1mg/kg for cadmium in barley.

**Sampling protocols**

EU Commission directive 2001/22/EC lays down sampling methods (and methods of analysis) for the official control of levels of lead, cadmium, mercury and 3-MCPD in foodstuffs. It states that 3, 5 or 10 incremental samples should be taken from lots of <50kg, 50-500kg and > 500kg. It also mentions that sampling should take place at the point where the commodity enters the food chain so it is clear, once again, that the procedure is not directly applicable to cereals in storage or in transit.

**Blending and sub-sampling**

If the lot consists of individual packages, then the directive recommends that one incremental sample should be taken where there are 1-25 packages or units in the lot and above this, about 5% should be taken. The incremental samples should be bulked to form the aggregate sample which should be at least 1kg (unless where not practical, e.g. where only one package has been sampled). It suggests laboratory samples should be large enough for at least duplicate analysis. However, no guidance is given on how to produce the laboratory sample.

**Validation**

Nothing found.

**Discussion**

This appears to be an area where there has been very little work and only one sampling protocol was found. Heavy metals in grain depend on uptake by the growing crop. Therefore, one can reasonably assume homogeneous distribution of heavy metals in stored grain in any one bulk since it will usually represent a
single variety grown in one or more fields that have been treated in a similar way. In some cases only selected field may have been treated with bio-solids and here it is clearly of value to follow HGCA sampling protocols\(^5.2\) to ensure crops from known fields are sorted into defined bins, or bays in a flat store. However, a study to confirm the normal range of heavy metals in a bulk of stored grain and this may be required.

References


6. AGROCHEMICAL RESIDUES (INCLUDING STORAGE PESTICIDES)

The commonest encountered residues on cereal grains tend to be chlormequat and pirimiphos-methyl (Anon, 1999). The former is a plant growth regulator (gibberellin biosynthesis inhibitor) that shortens and strengthens cereal stems, thus reducing ‘lodging’. The latter is an OP pesticide commonly applied in-store as a fabric treatment or admixed directly onto the grain as an admixture protectant against infestation. Diatomaceous earths (DEs) are beginning to be used as a replacement for OP dusts as a top-dressing for stored grain and the recommended levels of application are similar to the levels permitted for their addition as food additives.

Regulatory levels

The maximum residue levels (MRLs) are as follows:
5 ppm for pirimiphos-methyl, 2 ppm for chlormequat
These MRLs are laid down in UK statutory instruments and EU legislation.

Sampling protocols

The Codex Alimentarius suggests sampling procedures should follow ISO 950 recommendations and includes a table suggesting only 1 primary sample is required from a well-mixed or homogeneous bulk and that 3, 5 and 10 samples should be taken respectively from lots of < 50, 50-500 and > 500kg. It also suggests that the minimum size of a laboratory sample should be 1kg.

The USFDA recommendations are that five to ten 0.9kg sub-samples making up a total sample size of 4.5-9kg should be taken from lot sizes between 12 or less and over 190 (cases, crates, boxes etc).

Blending and sub-sampling

Nothing found.

Validation

Nothing found.

Discussion

Where an entire bulk has been admixed with a storage pesticide or where a bulk originates from a field which has been treated before harvest with a pesticide, herbicide or other chemical agent, it is reasonable to
expect the distribution of such a chemical agent to be homogeneous. The ‘grain passport system’ whereby paperwork indicating pesticide treatments accompanies loads of grain should also guard against multiple treatments and uneven distribution. Therefore, in these circumstances the sample guidance highlighted here may well be appropriate.

However, it is often suggested that treatment of the bin fabric may lead to local contamination of cereals close to the bin wall (although mixing may occur during outloading and conveying). Similarly, top-dressing of stored grain with OP dust formulations has become part of an integrated pest management regime based on cooling and drying. While OP dusts are no longer available on the market, top-dressing still remains an option e.g. by spraying with an emulsifiable concentrate as the last tonne goes into a silo or by raking in a diatomaceous earth dust. (Although this may need a label amendment). Where fabric treatments or top-dressing is used specific sampling regimes may be required. However, this is an area that does not appear to have been addressed.

References

6.1 Anon, 1999. Annual report of the working party on pesticide residues. Suppl. To Pesticides Monitor 1999. MAFF, PSD, HSE.

6.2 Statutory Instrument No 1767. The Pesticides (Maximum Levels in Crops, Food and Feeding Stuffs) (England and Wales) (Amendment) Regulations 2002 and The Pesticides (Maximum Levels in Crops, Food and Feeding Stuffs) (Scotland) Amendment Regulations 2002 (Statutory Instrument No 271

6.3 Statutory Instrument No 2723. The Pesticides (Maximum Levels in Crops, Food and Feeding Stuffs) (England and Wales) (Amendment)(No.2) Regulations 2002 and The Pesticides (Maximum Residue Levels in Crops, Food and Feeding Stuffs) (Scotland) amendment (No. 2) Regulations 2002 (Statutory Instrument No 489)


7. PESTS (INCLUDING MITES).

Insects and mites are of importance because of the direct damage they cause to grain. They are able to transmit disease organisms including mycotoxigenic fungi, due to their allergenic properties and because of their ability to increase rapidly under favourable conditions and to survive for long periods under unfavourable conditions. However, being mobile, they present a different challenge to other contaminants and sampling regimes need to take that into account and/or exploit their mobility. For this reason, there are two sampling procedures usually applied to insects. Grain samples can be taken and sieved to remove the arthropods, alternatively a variety of patented traps are available to sample over an extended period. As this document is intended to investigate a common sampling procedure for all analytes, the trapping aspect will only be lightly touched on here. An added complication concerns ‘hidden infestations’ of the non-mobile stages of species such as the grain weevil, *Sitophilus granarius* (L.) that complete their development inside the grain. Clearly these will not be detected by traps since they are not mobile and samples will need to be incubated to allow them to complete their development to adult. Alternatively, a rapid detection method such as egg-plug staining, x-ray or acoustic detection can be employed.

**Regulatory levels**

In the UK and most of Europe as well as Australia, there is no tolerance of live insects in trade. However, in the USA, the permitted threshold is 2 insects/kg. Thresholds for mites vary and informal limits for many petfoods are below 10/kg and for Intervention, below 250/kg.

**Sampling protocols**

The UK SOP for the phytosanitary certification of grain for export\textsuperscript{2,1} requires that 3kg of grain be taken from each lorry from at least 3 dispersed points and from different depths (unless an automatic core sampler is used). Whilst loading from silo/store to ship, 4kg of grain must be collected every 10 minutes from the grain stream.

ISO 6639/2\textsuperscript{2,2} deals with sampling for determination of hidden insect infestation. For grain flowing at rates of up to 100t/h, sampled lots should be 1–5t with increments of 1kg/t from a point there the grain is in free-fall. (This is because samples from conveyer belts are deemed less representative than those taken from points of free fall).

For sampling from static bulks, it notes insects are sought only in regions where they most commonly occur, which is near the surface, outlet spouts and ventilation and aeration openings and at shallow depths of 2-3m (for weevils). When the air temperature is above 15°C, surface layers 100 mm deep are sampled taking an
increment of 1 kg for every tonne of surface grain using a hand scoop and when the air temperature is below 15°C, a layer 250mm deep is sampled. In the former case, the number of increments is calculated by multiplying the bulk density by the surface area and dividing by 1,000 and in the latter case the division factor is by 400. It is also suggested that increments of 1kg be taken from the bottom of the bin by running grain from the outlet. For sampling beneath the surface, it merely states that increments of 1kg should be taken ‘at regular intervals’. This is clearly an example of ‘focused sampling’ – aimed at detection rather than determination.

Guidance on the detection of live invertebrate infestation by trapping is given in draft ISO 60027.3. However, this does not define trap spacing in the context of bulk grain, merely recommending probe traps be placed in pairs; in the top 10-30cm of grain and at 200cm depth. This contrasts with manufacturers’ recommendations for pitfall cone traps which recommend pairs be used at the surface and at 8-15cm in a 5-6m grid.

Blending and sub-sampling

ISO 6639/2 states that unless increments are submitted as laboratory samples, they should be combined and mixed and reduced as suggested by ISO 950 to provide a laboratory sample of no less than 1kg.

Validation

Wilkin (1991) showed that even a sampling rate of 4kg /20t, four times the industry (ISO) norm there was only a 50% chance of detecting insect with a single sample even with 5 insects/kg. In this case, the insects were evenly distributed while in practice, this is only sometimes the case, so even these estimates are likely to be optimistic.

Hagstrum et al. (no date) give tables to show the probability (at 95% level) of catching insects at different densities, in relation to the number of 1kg samples taken per 1000 bushels (ca 29t wheat) (Table 10). In the same publication they also give the number of samples needed to estimate populations to an accuracy of +/- the value of the mean (Table 11).
Table 10. Probability of detecting insects based on sample numbers and insect density (after Hagstrum et al.7.5)

<table>
<thead>
<tr>
<th>Insects/kg</th>
<th>kg samples/1000 Bushels</th>
<th>0.02</th>
<th>0.06</th>
<th>0.2</th>
<th>0.6</th>
<th>2.0</th>
<th>6.0</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td></td>
<td>0.02</td>
<td>0.06</td>
<td>0.19</td>
<td>0.43</td>
<td>0.76</td>
<td>0.95</td>
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<td>2</td>
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</tr>
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<td>1.00</td>
</tr>
</tbody>
</table>

Table 11. 95% confidence intervals in relation to sample number and insect density (after Hagstrum et al.7.5)

<table>
<thead>
<tr>
<th>Insects/kg</th>
<th>kg samples/1000 Bushels</th>
<th>0.02</th>
<th>0.06</th>
<th>0.2</th>
<th>0.6</th>
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<th>6.0</th>
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<tr>
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<td>0.07</td>
<td>0.15</td>
<td>0.31</td>
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</table>

In the US, the distribution of different insect species has been shown to be similar (Hagstrum et al., 19887.6) but experience in aerated bins in the UK suggests different species behave and distribute themselves differently. Whilst grain weevils (*Sitophilus granarius* L.) are unlikely to move much from their point of introduction, sawtoothed grain beetles (*Oryzaephilus surinamensis* L.) are more active and move rapidly away from cooling fronts to the surface (Armitage et al., 1983, 1994, 20027.7, 7.8, 7.9).

A most thorough synthesis of the statistics of sampling for insects by trap or by withdrawn incremental samples is given by Subramanyam and Hagstrum (1996)7.10 (Table 13). The advantages in reduced effort and accuracy of insect traps over taking samples is amply illustrated in the table below.
Table 12. The probability of detecting 0.2 Cryptolestes ferrugineus Steph. per kg (after Subramanyam and Hagstrum (1996)7,10).

<table>
<thead>
<tr>
<th>Probe traps</th>
<th>Duration (days)</th>
<th>Samples</th>
<th>Detection Probability</th>
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</thead>
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<td>No.</td>
<td></td>
</tr>
<tr>
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</tr>
<tr>
<td>5</td>
<td>1</td>
<td>5</td>
<td>0.64</td>
</tr>
<tr>
<td>5</td>
<td>5</td>
<td>25</td>
<td>0.99</td>
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</table>

The distribution of mites in UK grain has been examined by sampling bulks (Armitage 1984, Armitage et al., 1994, 20027,9, 7.10, 7.11) and been shown to be related to storage moisture content and is fairly homogeneous within layers. It may be summarised briefly by saying that when grain is held at or below the recommended moisture content of 14.5%, pest mites will only be found at the grain surface during winter. Thus sampling in the right place is vital in gaining an idea of their population density.

Discussion.

While sampling grain for mites is fairly straightforward and predictable, since mites are largely distributed in relation to grain moisture content, sampling for insects is an altogether more complex issue, comparable in difficulty to the mycotoxin question. Although it is clear that traps are appropriate to monitor insect populations in farm-stored grain, they are not appropriate for incorporation into a common sampling plan where instant diagnosis is required at the point of sale. Some accommodation would need to be reached about the acceptance of on-farm monitoring results at the point of sale. In addition, traps do not indicate the presence of dead arthropods or their fragments that may be important allergenic contaminants. The suggestions in ISO 6639/27.2 are frankly bizarre and over-complicated and it is not clear what science has led to these suggestions. They are clearly intended, not to produce a representative sample, but to detect any infestation. Reduction of the bulk sample, as is suggested in ISO 6639/C will only decrease the already low chances of detecting infestation so would be inappropriate. In this case, sieving the entire bulk sample before sub-division and analysis for other analytes may be appropriate. This has been facilitated by the commercial production of inclined sieves (e.g. Wilkin et al., 19947.12) which can process 5kg at a time. This approach would only be useful for detecting free-roaming adult insects. If a rapid assessment of hidden infestation was required, it is most likely that acoustic detection would be the only applicable method, capable of giving a near instant result.
References

7.1 DEFRA. 2000 Phytosanitary certification of grain for export. Form GSOP 1 Plant Health Division, York, UK.


7.5 Hagstrum, D, Flinn, P. and Fargo, S., How to sample grain for insects Chapter 11 in ‘Stored product management’ Publication E-912 Oklahoma State University / USDA. http://www.okstate.edu/OSU_Ag/agedcm4h/pearl/e912/ch11.pdf


OVERVIEW AND RECOMMENDATIONS

UK industry would benefit from a single validated sampling protocol for all the analytes discussed, which would ensure representative results from analysed laboratory samples. However, it is clear that the distribution of each analyte is different; some are heterogeneous, others homogeneous and others unknown or the opinion thereof held by different authorities is contradictory. In addition this report shows the purpose of sampling to vary. Some samples are taken for detection (GMO, Salmonella, insects) while in most cases the aim is for determination to gain a representative sample that predicts average quality.

Not all analytes have formal sampling protocols and even where protocols exist, these have rarely been validated by experiment. The number of samples required by sampling protocol varies by analyte and there is little consistency in defining size of samples and in expressing the sampling frequency in a consistent way (e.g. number and size of samples per unit of bulk) or even using SI units (see Table 13).

Specific recommendations for heavy metal sampling, for GM and for pesticide residues are ISO-based and although as yet unvalidated may prove to be largely appropriate. However, it is clear that sampling for the very heterogeneous biological analytes, e.g. arthropods and fungi and their metabolites, mycotoxins require more detailed protocols and sampling regimes.

One approach to a ‘one size fits all’ solution, would be to adopt the most rigorous regulatory sampling requirement, which is undoubtedly that for storage mycotoxins, and divide the resulting bulk sample to provide the necessary laboratory samples for other analytes. (In the case of insects, there would be considerable advantage in examining the entire bulked sample, for instance by passing it over an inclined sieve).

However, before such an approach is adopted, it would be highly desirable to simplify the statutory sampling plan for storage mycotoxins which has the potential of considerable extra costs for industry. Design of such a plan would depend on a greater understanding of mycotoxin formation during drying or storage of damp grain, than currently exists. Anecdotal information received from ‘Euromalt’ suggests so far that only the EU regulatory sampling for mycotoxins reliably detects the regulatory mycotoxin level but there is a definite need to find a simplified sampling regime that is substantially equivalent to the regulatory procedure and thus demonstrate ‘due diligence’.

Another approach would be to adopt the ISO standard for those analytes that are likely to be homogeneous - pesticides, heavy metals, GMOs, mites, microbial contamination. Sampling for OA could be carried out at a lesser frequency to demonstrate due diligence. Should surveys by FSA, DEFRA etc. show the inadequacies of this approach, a greater frequency of OA sampling would be required.
HGCA sampling recommendations are to take a sample from 1kg from each trailer (or 1kg/t from driers) and make a composite sample to represent every 50t. This does not specify a rate of sampling per tonne since a trailer can be anything up to 20t. This has been shown to be adequate for determining market quality but needs to be validated to see if it is also suitable for determination of homogeneously distributed analytes such as pesticides, or heavy metals or to see if it can be adapted for heterogeneous analytes such as mycotoxins.

However, adopting any of these approaches would first require experimental validation and testing of the applicability to practice by sampling from farm and commercial stores. The first approach would require estimation of the size of sub-sample to be taken for each analyte.

Finally, the pattern of mixing of grain during unloading of bins or floor stores has considerable influence on the distribution of analytes and ultimately therefore upon the sampling regime. For instance, during the unloading of hopper-bottom bins, the grain empties as a column above the bin centre and grain from the surface is some of the earliest to be discharged. In view of the importance of the grain surface as a primary source of some contaminants such as mycotoxins and arthropods, due to the fluctuating moisture contents there, it is easy to see that by failing to sample this, earliest discharging grain, one might also miss the prime source of these contaminants. Equally, this pattern of discharge suggests only limited mixing of the surface grain might occur. Since this factor has such an important effect on the assumed and actual distribution of analytes, it requires its own experimental study.

ACKNOWLEDGEMENTS

This report was compiled with the guidance of several CSL experts in their fields; Christine Henry and James Blackburn (GMOs), Susan MacDonald (Mycotoxins), John Chambers (insects and mites) and Stuart Reynolds (Pesticides). Thanks are also due to Ken Wildey and Simon Hook, HGCA for helpful discussions.
Table 13. Summary of current sampling recommendations for different analytes.

<table>
<thead>
<tr>
<th>Analyte / Purpose</th>
<th>Authority</th>
<th>Samples Incremental</th>
<th>Nos/t</th>
<th>Bulk/Aggregate</th>
<th>kg/t</th>
<th>Laboratory</th>
</tr>
</thead>
<tbody>
<tr>
<td>Usually Determination</td>
<td>ISO 13690</td>
<td>5-11 from 15-500t</td>
<td>1/3-4.5t</td>
<td>1/5-11</td>
<td>1.25kg from 5000 bushels</td>
<td>1kg/116t</td>
</tr>
<tr>
<td>GM Detection</td>
<td>EEP As ISO</td>
<td>30</td>
<td>1/3-4.5t</td>
<td>1/20 bulk</td>
<td>3000 kernels</td>
<td></td>
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<tr>
<td>Mycotoxin Determination</td>
<td>EU 2002/26/EC</td>
<td>100 from 50-1500t</td>
<td>1/0.5-150t</td>
<td>10 kg</td>
<td></td>
<td></td>
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<tr>
<td>Salmonella Detection</td>
<td>USFDA</td>
<td>15 x 100g</td>
<td>2-3 lbs</td>
<td>200g lab, 50g sub-sample</td>
<td></td>
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<tr>
<td>Heavy metals Determination</td>
<td>EU 2001/271 EC</td>
<td>3-10 from &lt;50-&gt;500kg</td>
<td>1/16-50t</td>
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<td></td>
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<tr>
<td>Pesticides Determination</td>
<td>Codex</td>
<td>3-10 from &lt;50-&gt;500kg</td>
<td>1/15-50t</td>
<td></td>
<td></td>
<td></td>
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<tr>
<td>Arthropods Detection</td>
<td>DEFRA</td>
<td>3 kg from 3 pts/lorry</td>
<td>1kg/7t</td>
<td></td>
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<tr>
<td></td>
<td>Phyto. Certificate</td>
<td>4kg/10 min</td>
<td>1kg/t</td>
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<tr>
<td></td>
<td>ISO 6639/C</td>
<td>1kg/t</td>
<td>1 kg</td>
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