### Microorganisms

See also CLEAPSS Student Safety Sheet 76, Bioreactors and Fermenters

<table>
<thead>
<tr>
<th>Source</th>
<th>Hazard</th>
<th>Comment</th>
</tr>
</thead>
<tbody>
<tr>
<td>Samples from the environment</td>
<td>BIOHAZARD</td>
<td>Air, water and soil samples could be used, but do not sample from high-risk areas, e.g., toilets or the floors of changing rooms. All environmental samples could be contaminated with pathogens (organisms which cause disease).</td>
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<tr>
<td>Samples from humans</td>
<td>BIOHAZARD</td>
<td>‘Finger dabs’ could be used. Samples may, however, be contaminated with pathogens (see above).</td>
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<tr>
<td>Foods</td>
<td>BIOHAZARD</td>
<td>Any uncooked animal product (eggs, meat, cheese etc) may be contaminated with bacteria, especially Salmonella and Escherichia coli (E. coli) from the gut, which can cause food poisoning. Take care to prevent cross contamination between cooked and uncooked foods. Thorough cooking will destroy bacteria.</td>
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<tr>
<td>Purchased cultures (ie, samples of microorganisms bought from suppliers)</td>
<td>BIOHAZARD</td>
<td>Cultures bought from reputable suppliers (but not those from hospitals, etc) should be safe but may have become contaminated. E. coli is often studied in schools, but this is not the same strain of bacterium that causes food poisoning.</td>
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</tbody>
</table>

### Typical control measures to reduce risk

- When culturing, if possible, grow bacteria and fungi on solids (agar) rather than liquids (broth) to avoid spills and aerosol formation. Choose culture media, eg, nutrient agar, that do not favour the growth of pathogens. Incubate at temperatures which do not encourage growth of pathogens (avoid temperatures 30°C to 42°C).
- Do not seal cultures completely before incubation (otherwise hazardous anaerobic bacteria are encouraged) but make sure they cannot be opened accidentally.
- Incubated cultures taken from the environment or humans must never be opened.
- Use sterile equipment and aseptic technique (eg, by flaming loops and mouths of bottles, etc).
- Avoid draughts (from open windows and doors) which could contaminate cultures and cause spores from fungi (eg, mould) to spread.
- Work near Bunsen-burner flames so that the updraught helps to prevent contamination of cultures.
- After work is complete, treat surfaces using a suitable disinfectant for a sufficient length of time.
- Dispose of all cultures (including mould on food eg, mouldy bread) by sterilisation in an autoclave (pressure cooker).
- Always wash hands after using/handling cultures and before handling food.
- Wear a clean lab coat or overall to protect cultures and food from microbes on the skin, clothing, etc.
- In cooking, ensure that food is heated to at least 70 °C for at least 2 minutes.
- Do not reheat cooked food; prepare, store and display cooked and uncooked foods separately.

### Assessing the risks

- What are the details of the activity to be undertaken? What are the hazards?
- What is the chance of something going wrong? eg, could a food or a culture be, or become, contaminated? Could microorganisms or their spores escape?
- How serious would it be if something did go wrong? eg, could material pathogenic to humans be released? Could food poisoning result?
- How can the risk(s) be controlled for this activity? eg can it be done safely? Does the procedure need to be altered?

### Emergency action

In all emergency situations, alert the responsible adult immediately. Be aware that actions may include the following:

- Spilt on the floor, bench, etc For spills of cultures, place paper towels over the spill, pour disinfectant (eg, Virkon) on top and leave for at least 15 minutes. Bleach is usually suitable in the home.