



BIOSAFETY SELF-ASSESSMENT CHECKLIST

This checklist must be completed annually by the supervisor for each laboratory working with biologically hazardous materials (e.g. microorganisms / animals / plants / human specimens) to ensure that biosafety requirements are being implemented appropriately.

Inspection Team: _____

Laboratory Room No: _____

Laboratory Supervisor: _____

Inspection Date: _____

CRITERIA	YES	NO	N/A
RISK			
1. Have the risk groups for all biologically hazardous materials used in the laboratory been determined?	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
2. Has the UOW Risk Group Register been completed for this laboratory and does it include all the risk groups as per Step 1 above?	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
3. Has the completed Risk Group Register been filed with the school manager or equivalent?	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
SIGNAGE			
4. Has the correct signage been placed outside the laboratory entrances indicating physical containment levels, entry requirements and warning symbols?	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
5. Has the biohazard symbol been attached to equipment containing biological hazardous material? e.g. fridges, freezers, liquid nitrogen dewars.	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
OGTR			
6. If working with GMOs, has the correct OGTR signage been placed on the laboratory entrances?	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
7. Do Risk Assessments (RAs) and Safe Work Procedures (SWPs) for the OGTR PC2 laboratory identify reporting requirements to the OGTR for any unintentional release from the facility or during transportation from the facility?	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
8. Are all storage vessels containing GMOs labelled with contact details of the person responsible for the dealings?	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
9. Are there records of GMOs exact location?	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
10. Have the appropriate OGTR approvals for all recombinant DNA procedures undertaken in the laboratory been obtained?	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
Biosecurity / Department of Agriculture and Water Resources (Formally AQIS)			
11. If working with quarantineable material (deemed quarantineable by the DAWR as BC1 or BC2), is the facility certified?	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
12. Are DAWR work practices being followed?	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>

CRITERIA	YES	NO	N/A
WORK PRACTICES			
13. Have correct work practices been followed for the type of work being performed in the laboratory (e.g. PC1 or PC2 work practices)?	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
14. Have these work practices been documented on SafetyNet as an SWP?	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
15. If working in a PC2 facility, has entry been restricted to only authorised users, and has a register listing all authorized persons been posted at the laboratory entrance?	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
16. Does the laboratory have a safety manual specific to its needs?	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
TRAINING			
17. Have all users of the PC2 facilities been trained to ensure they can work safely in the facility?	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
18. Does the training include:			
▪ laboratory induction	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
▪ training in handling pathogens	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
▪ correct use of personal protective clothing and equipment (PPCE)	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
▪ immunisation requirements	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
▪ operation of specific equipment	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
▪ emergency plans, spill cleanups (includes correct spill kits for each lab)	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
▪ biosafety training	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
▪ completing the online quiz if working in an OGTR laboratory	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
▪ animal handling courses where necessary	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
▪ ethics requirements	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
RECORD KEEPING			
19. Have training records been kept?	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
20. Have immunisation records been kept (confidential material)?	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
PURCHASING			
21. Was the containment level of the facility suitable for the risk groups of biologically hazardous materials being purchased / obtained?	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
22. Have RAs been completed to identify and control any hazards before purchasing / obtaining biologically hazardous materials?	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
RISK ASSESSMENTS AND SAFE WORK PROCEDURES			
23. Were RAs readily available and do they demonstrate that all biologically hazardous materials are being adequately controlled?	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
24. Have RAs been approved by the laboratory supervisor?	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
25. Were SWPs available for all standardised procedures carried out in the laboratory (SWPs must also be written for any equipment that generates aerosols such as vortexing, sonicating, pipetting)?	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
26. Have SWPs been reviewed by competent personnel familiar with the procedures and approved by the laboratory supervisor?	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
27. Are SWPs listed on SafetyNet ?	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>

CRITERIA	YES	NO	N/A
IMMUNITY / HEALTH MONITORING			
28. Have laboratory personnel been informed of the potential risks when working with infectious or potentially infectious microorganisms while pregnant, considering pregnancy, immuno-compromised or immuno-suppressed?	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
29. Have laboratory personnel provided immunisation records or signed a Decline of Immunisation form?	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
30. Have workers working with Risk Group 3 and 4 human pathogens had an initial health assessment and follow up checks? Refer to the Air and Health Monitoring Guidelines.	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
31. Have pregnant workers been informed of the risks of working with certain pathogens?	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
32. Are staff working with animals? Have identified persons at risk of allergens been subject to ongoing health monitoring requirements? Refer to the Air and Health Monitoring Guidelines.	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
BIOLOGICAL WASTE			
33. Have correct disposal procedures been used for biologically hazardous waste?	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
34. Have Waste Tracking Logs been completed?	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
LABORATORY DECOMMISSIONING			
35. Were staff familiar with PC2 laboratory decommissioning procedures?	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
VISITORS AND CONTRACTORS			
36. Was there a laboratory induction for visitors and contractors to the laboratory?	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
37. Was PPE available for visitors?	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
TRANSPORT			
38. Were all laboratory personnel aware of the requirements for transporting biologically hazardous materials?	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
39. Were laboratory personnel able to identify the people qualified to pack biologically hazardous materials for transport by road, rail or air?	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
GENERAL			
40. Were all laboratory personnel aware of the current local biosafety representatives?	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
41. Were laboratory personnel aware of biosafety information available on the WHS website?	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
DEFINITIONS			
OGTR:Office of Gene Technology Regulator PC1/PC2:Physical Containment Level 1 or 2 GMO:Genetically Modified Organism AP:Approved Premises SWP:Safe Work Procedure RA:Risk Assessment DAWR:Department of Agriculture and Water Resources PPE:Personal Protective Equipment SafetyNet UOW online system for hazard and incident reporting			

The completed document must be filed in the local area [WHS Records Index](#) in accordance with the [WHS Records Handling Guidelines](#).