# **UW LRA WORKSHEET FOR HUMAN PATHOGENS AND TOXINS**

Completed by:	Date completed:	
Material description		
1. Name or description of the material being	handled. Briefly describe how you	intend to use the material.
2. Identify where the material will be used an		
Used	Stored	
	V	
3. Is the material considered pathogenic:   If Yes	Yes ☐ No If No	
Indicate Risk Group:   1 1 2 3	Indicate why it is considered no	on-pathogenic:
How was risk group determined?	☐ Material comes from an other	-
☐ Pathogen data sheet	individual	
☐ By supplier or other researcher	☐ Material comes from the env	vironment in an
☐ Pathogen risk assessment	unaltered state  Other:	
☐ Other:	□ Other:	
Personnel factors		
r Gi Sulli Gi I de Lui S		
<b>4.</b> Is a vaccine available?	f vaccine:	□ No □ N/A
5. Have all personnel working with or near th		☐ Yes ☐ No
any available vaccinations OR declined wit (Departments must have a record of the vac		
6. Is a medical surveillance plan in place and	documented? Please describe:	☐ Yes ☐ No
7. Is a Medical Contact Card required?		☐ Yes ☐ No
8. Instructed in signs and symptoms of infect	ion?	□ Yes □ No



## Factors associated with the specific work processes

Face shield   Safety glasses   N-95   Face mask   Back-closing gown at BSC   10. Frequency of contact with agent:   Routine/daily   Weekly   Random/monthly/yearly   11. Largest single volume used:   < 10 L   More than 10 L (if greater than 10L specify):   a. Indicate concentrations used (if concentrated, enter both before and after concentrations):   b. Indicate concentration required to cause infection:   N/A   12. Is all work with the active agent done in a BSC? (not required for CL1)   Yes   No     13. Is open bench work completed on agent (means not in hood)?   Yes   No     a. Describe in point form the techniques proposed on the open bench:    b. Describe your risk management techniques:   Yes   No     b. If yes, are you using safety-engineered sharps?   Yes   No     b. If not, please explain:   Shaking or vigorous mixing     Grinding   Flaming loops   Shaking or vigorous mixing     Grinding   Pipetting   Homogenizing     Opening containers with high internal pressures     Other procedures that may create an airborne exposure to a pathogen:   Yes   No     a. If YES, are sealed rotors, or sealed centrifuge safety cups available?   Yes   No     b. If NO to "16 a", do you only use screw -cap, non-glass tubes?   Yes   No     c. Will you open the tubes in the BSC after centrifuging?   Yes   No	9. PPE required when working with agent (check all that apply).  Note: lab coat, close-toed shoes, and gloves are all mandatory for microb	viological work!
11. Largest single volume used:   < 10 L   More than 10 L (if greater than 10L specify):   a. Indicate concentrations used (if concentrated, enter both before and after concentrations):   b. Indicate concentration required to cause infection:   N/A   12. Is all work with the active agent done in a BSC? (not required for CL1)   Yes   No     13. Is open bench work completed on agent (means not in hood)?   Yes   No     a. Describe in point form the techniques proposed on the open bench:   b. Describe your risk management techniques:   Yes   No     b. If yes, are you using safety-engineered sharps?   Yes   No     b. If not, please explain:   Centrifuging in open containers     Blending   Flaming loops   Shaking or vigorous mixing     Grinding   Pipetting   Homogenizing   Opening containers with high internal pressures     Other procedures that may create an airborne exposure to a pathogen:     16. Will your experiments involve centrifugation?   Yes   No     a. If YES, are sealed rotors, or sealed centrifuge safety cups available?   Yes   No     b. If NO to "16 a", do you only use screw -cap, non-glass tubes?   Yes   No     c. Will you open the tubes in the BSC after centrifuging?   Yes   No	$\square$ Face shield $\square$ Safety glasses $\square$ N-95 $\square$ Face mask $\square$ Back-closing gow	vn at BSC
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12. Is all work with the active agent done in a BSC? (not required for CL1)   Yes   No   13. Is open bench work completed on agent (means not in hood)?   Yes   No   14. Are you using sharps?   Yes   No   15. Identify the processes that increase exposure potential (check all that apply):   Cell sorting   Sonication   Centrifuging in open containers   Blending   Flaming loops   Shaking or vigorous mixing   Grinding   Pipetting   Homogenizing   Opening containers with high internal pressures   Other procedures that may create an airborne exposure to a pathogen:  16. Will your experiments involve centrifugation?   Yes   No   16. Will your experiments involve centrifuge safety cups available?   Yes   No   17. Identify the processes that increase exposure potential (check all that apply):   Pipetting   Homogenizing   Opening containers with high internal pressures   Other procedures that may create an airborne exposure to a pathogen:  16. Will your experiments involve centrifugation?   Yes   No   17. Identify the processes that increase exposure exposure to a pathogen:   Yes   No   18. Identify the processes that increase exposure exposure to a pathogen:   Yes   No   19. Identify the processes that increase exposure exposure to a pathogen:   Yes   No   19. Identify the processes that increase exposure exposure to a pathogen:   Yes   No   19. Identify the processes that increase exposure exposure to a pathogen:   Yes   No   19. Identify the processes that increase exposure exposure to a pathogen:   Yes   No   19. Identify the processes that increase exposure	a. Indicate concentrations used (if concentrated, enter both before and	after concentrations):
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14. Are you using sharps?	13. Is open bench work completed on agent (means not in hood)?	□Yes □No
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a. If yes, are you using safety-engineered sharps?	14. Are you using sharps?	□ Yes □ No
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<ul> <li>Cell sorting</li></ul>	b. If not, please explain:	
<ul> <li>□ Grinding □ Pipetting □ Homogenizing</li> <li>□ Opening containers with high internal pressures</li> <li>□ Other procedures that may create an airborne exposure to a pathogen:</li> <li>16. Will your experiments involve centrifugation? □ Yes □ No</li> <li>a. If YES, are sealed rotors, or sealed centrifuge safety cups available? □ Yes □ No</li> <li>b. If NO to "16 a", do you only use screw -cap, non-glass tubes? □ Yes □ No</li> <li>c. Will you open the tubes in the BSC after centrifuging? □ Yes □ No</li> </ul>	☐ Cell sorting ☐ Sonication ☐ Centrifuging in open contain	• • • •
a. If YES, are sealed rotors, or sealed centrifuge safety cups available?  b. If NO to "16 a", do you only use screw –cap, non-glass tubes?  c. Will you open the tubes in the BSC after centrifuging?  yes No	<ul><li>☐ Grinding</li><li>☐ Pipetting</li><li>☐ Homogenizing</li><li>☐ Opening containers with high internal pressures</li></ul>	:
b. If NO to "16 a", do you only use screw –cap, non-glass tubes?  c. Will you open the tubes in the BSC after centrifuging?  Yes No	16. Will your experiments involve centrifugation?	☐ Yes ☐ No
c. Will you open the tubes in the BSC after centrifuging?	a. If YES, are sealed rotors, or sealed centrifuge safety cups available?	☐ Yes ☐ No
	•	
d. It NO to (b) or (c), explain how you will protect against exposure:	<ul><li>c. Will you open the tubes in the BSC after centrifuging?</li><li>d. If NO to (b) or (c), explain how you will protect against exposure:</li></ul>	☐ Yes ☐ No

## **Disinfection and waste disposal**

- 17. At what stage of your work will the infectious agent be inactivated or lysed?  $\square$  N/A Note: N/A should only be used if there is no infectious agent.
- 18. Specify disinfectants and decontaminants and decontamination procedures in use:  $\square$  N/A

Disinfectant	Working Concentration	Contact Time (min)	Preparation Frequency	Indicate where used (surface, equipment, tools, etc)

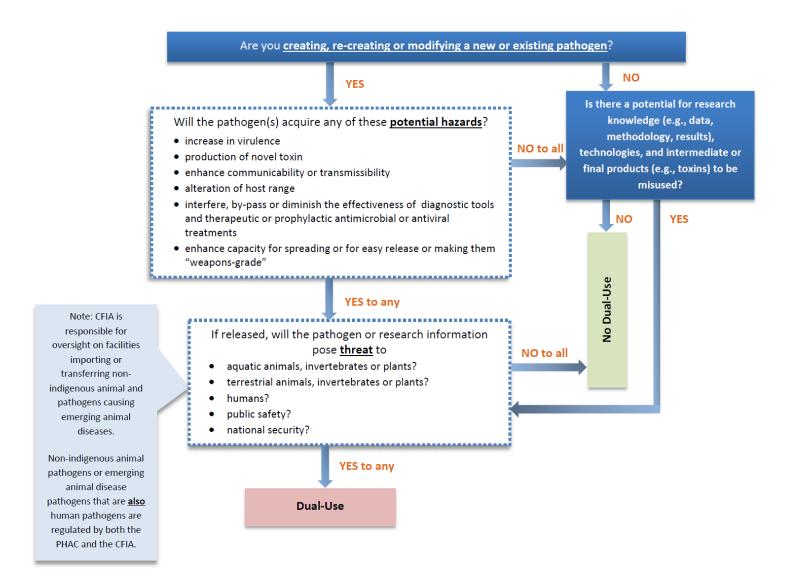
**19.** Complete the table to identify how biohazardous wastes generated by your research are treated. Any autoclaving and direct disposal requires weekly efficacy logs.

	Waste Generated and disinfection process	Disposed by (select one)
□Yes	Solid waste contaminated with biohazardous	☐ Biowaste bin (UW Disposal Service)
□No	material and all microbial and eukaryotic cell	☐ Autoclaving:
□N/A	cultures, including broth cultures	Temp: °C
		Time: min.
□Yes	Needle and syringe assemblies (sharps)	☐ Biowaste sharps bin (UW Disposal Service)
□No		☐ Autoclaving
□N/A		Temp: °C
		Time: min.
□Yes	Used glass and hard plastic pipettes and	☐ Biowaste sharps bin (UW Disposal Service)
□No	Pasteur pipettes will be:	☐ Autoclaved and disposed as regular waste
□N/A		Temp: °C
		Time: min.
□Yes	Liquid waste contaminated with	☐ Biowaste bin (UW Disposal Service)
□No	biohazardous material	☐ Autoclaving
□N/A		Temp: °C
		Time: min.
		☐ Chemically
		Chemical name:
		Contact time:
□Yes	Other, specify:	
□No		
□N/A		

### **Dual use potential**

20. Review chart below and identify dual use potential.

**Does dual use potential exist?** ☐ Yes ☐ No If yes, please describe:



#### **Summary**

All individuals in this lab must review the following University SOPs and guidance documents. Please go to the <u>biosafety website</u> to access them.

- UW Emergency Response Guide for Biologicals - Exposures and Spills
- Movement and Transportation of Biological Materials
- Guidance on Disinfection
- Vacuum Aspiration Guidance
- Proper Pipetting Techniques

<b>Example</b> – SOP 734 – Purification of xxx by centrifugation	n
List the names of all workers on this project:	
I acknowledge that work on this project will not be  1. All workers on this project have reviewed th	<u> </u>
- · · · ·	e mandatory documents listed above.
<ol> <li>All workers on this project have reviewed th</li> <li>All workers have completed the University's</li> <li>The Safety Office has approved this project.</li> </ol>	e mandatory documents listed above.
<ol> <li>All workers have completed the University's</li> <li>The Safety Office has approved this project.</li> </ol> Supervisor name:	e mandatory documents listed above.
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1. All workers on this project have reviewed th 2. All workers have completed the University's 3. The Safety Office has approved this project.  Supervisor name:  Signature:  Safety Office determination  Project may proceed as proposed  Yes  No	e mandatory documents listed above. online <u>Biosafety training module</u> .