STANDARD AND SPECIAL MICROBIOLOGICAL PRACTICES

(For assistance, please contact EHS at (402) 472-4925, or visit our web site at http://ehs.unl.edu/)

References
The information in this SOP is extracted from the following sources:
1. NIH Guidelines for Research Involving Recombinant DNA Molecules (NIH Guidelines), National Institutes of Health
2. Biosafety in Microbiological and Biomedical Laboratories, Centers for Disease Control and National Institutes of Health

In accordance with the above referenced sources, certain standard and special microbiological practices must be observed in laboratories working with certain biological agents. This SOP summarizes those practices and can be used as a training and information tool. This SOP (or equivalent) should be incorporated into laboratory-specific biosafety manuals and/or training materials. In several cases, the practices listed below are re-worded to harmonize between the various cited sources. Principal Investigators (PIs) are advised to supplement these standard and special practices with laboratory or procedure specific guidance, as appropriate and applicable.

Scope
This SOP applies to all work at UNL that is subject to the UNL Biosafety Guidelines and which is conducted at biosafety containment levels 1 or 2.

Standard and Special Practices
Depending on the containment level approved for the project by the Institutional Biosafety Committee (IBC), the following standard practices must be observed:
1. Laboratory access is restricted. In accordance with UNL procedure:
   a. Laboratory doors are equipped with current and accurate hazard warning placards. See EHS SOP, Door Postings for Potentially Hazardous Locations.
   b. Laboratory doors are closed at all times, and locked when not occupied.
   c. If the containment level is BSL-2 or higher, only persons who have been adequately trained, advised of the potential hazards, meet specific entry requirements (e.g., vaccination, other medical surveillance or qualification, serum banking, etc., as specified in the project specific protocol), and who comply with all entry and exit procedures are allowed to enter the laboratory.
   d. If the containment level is BSL-2 or higher, access for minors may be restricted or prohibited in accordance with Human Resources policies and/or labor laws.
2. Good personal hygiene practices are observed.
   a. Eating, drinking, smoking, handling contact lenses, and applying cosmetics, and
      storing food for human consumption are not permitted in the work area.
   b. Persons wash their hands: after handling potentially infectious materials, rDNA
      molecules, and/or animals, and; before exiting the laboratory.
   c. Mechanical pipetting devices are used; mouth pipetting is prohibited.

3. Appropriate Personal Protective Equipment (PPE) is available and used.
   a. If the containment level is BSL-2 or higher, laboratory coats, gowns, smocks, or
      uniforms are worn while in the laboratory, and removed before exiting the
      laboratory.
   b. Appropriate protective gloves are used when contact with rDNA, infectious
      materials, and/or animals is likely. Gloves are changed when contaminated,
      integrity has been compromised, or when otherwise necessary. Hands are
      washed after removing gloves, before touching clean surfaces, before exiting the
      laboratory, and before donning new gloves. Disposable gloves are not reused
      and are managed as potentially contaminated laboratory waste.
   c. For HIV and HBV laboratories, solid front or wrap-around gowns, scrub suits, or
      coveralls are worn in the laboratory. Laboratory clothing is not worn outside the
      laboratory, and it is decontaminated prior to laundering and disposal.
   d. Eye protection is used. Goggles may be required if there is substantial risk of
      splashes and/or aerosols. Additional PPE may be required for special tasks
      (e.g., face shield, respirator, goggles). Eye and face protection must be disposed
      of with other contaminated laboratory waste or decontaminated before reuse, as
      appropriate.

4. Disinfection and decontamination.
   a. Work surfaces are decontaminated once a day and after any spill of potentially
      infectious materials with appropriate disinfectant (as specified in the applicable
      protocol).
   b. Laboratory equipment is routinely decontaminated as well as after spills,
      splashes, or other potential contamination. Equipment is decontaminated before
      repair, maintenance, or removal from the laboratory.
   c. Spills involving infectious materials must be promptly contained, decontaminated,
      and cleaned up. Laboratory staff are trained in spill clean-up and have
      appropriate spill clean-up materials readily available. See EHS SOP, *Spill and
      Exposure Response for Biohazardous Materials (including Recombinant
      Nucleic Acids)*. All contaminated liquid or solid wastes are decontaminated
      before disposal. Contaminated materials that are to be decontaminated at a site
      away from the laboratory are placed in a durable leak-proof container which is
      closed before being removed from the laboratory. Materials to be transported by
      public highways or roads for off-site decontamination are packed in accordance
      with applicable local, state, and federal regulations.

5. All procedures are performed carefully to minimize the creation of splashes and
   aerosols. For BSL-2, all activities that are anticipated to present a splash or aerosol
   hazard are conducted in biological safety cabinets or other physical containment
   devices. For HIV and HBV labs, all work with open vessels is be conducted in a
   biosafety cabinet (not on the open bench).

6. Use of sharps (e.g., needles, scalpels, pipettes, glassware) is avoided. Plasticware is
   substituted for glassware whenever possible.
   a. When absolutely necessary, the following general precautions are taken.
      i. Needles are not bent, sheared, broken, recapped, removed from
         disposable syringes, or otherwise manipulated by hand before disposal.
ii. Used sharps are carefully placed in puncture-resistant containers.

iii. Broken glassware is not handled directly. Instead, it is handled using mechanical assistance (e.g., brush and dustpan, tongs, forceps).

iv. Effective hand protection is used (e.g., molded guards, use of forceps when using scalpels, etc.).

b. If the containment level is BSL-2 or higher, hypodermic needles and syringes are used only for parenteral injection and aspiration of fluids from laboratory animals and diaphragm bottles. Only needle-locking syringes or disposable syringe-needle units (i.e., needle is integral to the syringe) are used. Extreme caution is used when handling needles and syringes to avoid autoinoculation and the generation of aerosols during use and disposal. Needles are not bent, sheared, replaced in the needle sheath or guard, or removed from the syringe following use.

7. If the containment level is BSL-2 or higher, spills and accidents that result in overt exposures are reported immediately to the Principal Investigator (PI) and EHS. Medical evaluation is sought immediately. See EHS SOP, On the Job Injuries. In some cases, reporting to regulatory authorities may be required. See EHS SOP, Incident Reporting – National Institutes of Health (NIH) Guidance.

8. An effective rodent and insect control program is observed. Signs of infestation are immediately reported to the Principal Investigator (PI) and action taken to impede incursion. If the containment level is BSL-2, animals and plants unrelated to the work are not permitted in the work area.

9. For HIV and HBV laboratories, vacuum lines are protected with high efficiency particulate air/HEPA filters and liquid disinfection traps, which are checked routinely and maintained or replaced as necessary.

10. Laboratory personnel demonstrate proficiency in standard and special microbiological practices, receive appropriate training regarding their duties, the necessary precautions to prevent exposures, and exposure evaluation procedures. Personnel receive annual updates or additional training when procedural or policy changes occur. Personnel are advised that personal health status may impact an individual’s susceptibility to infection, and their ability to receive immunizations or prophylactic interventions. Personnel are encouraged to self-report conditions that may impact their immune competence or predispose them to infection so that they may seek counseling and guidance from UNL’s healthcare provider.