## [BACK TO THE LAB]



As you return to the lab and bring your systems back on-line, follow these simple steps for proper startup:

### LC SYSTEM START-UP REFERENCE

- Check the state of the instrument, connections and waste, and ensure it is calibrated/within the preventative maintenance period
- Discard all aqueous/buffered mobile phases that have been opened, left out, or are greater than one (1) month old
- 3. Discard any samples that have been left in the sample manager or autosampler
- 4. Remove the analytical column from the system and replace with an appropriate union
- 5. Place all solvent lines, including wash and purge lines, into a clean bottle containing 80–90% organic (MeCN or MeOH)/10–20% water. If salt buffer were left on the instrument, prime with 100% water prior to adding any organic MP to the system
- 6. Cycle the power to each module
  - a. Allow each module to go through it's start up diagnostics before powering on the next unit
  - b. Check all communications between systems/ workstation and reestablish as needed.
    - i. Includes remapping any shared network drives
- 7. Prime each solvent/wash line for 5 min with the above mobile phase

- 8. Set flow to 0.5 mL/min and allow to flow the system with no column for 20–30 min
- 9. Where applicable, run System Diagnostic tests. For example:
  - a. Pump leak tests
  - b. Sample manager diagnostics and leak tests
- Create fresh working mobile phase(s) and wash solvents and route the LC solvent lines to the appropriate solvent containers
- 11. Prime each line for 5 min
- 12. Connect the column to the inlet of the system (Do not connect the column to the detector)
  - a. Gradually start the flow, collecting the mobile phase effluent in a beaker/waste container
    - i. Increase the flow gradually to the desired flow rate
    - ii. Observe the pressure, ensure the pressure reaches a steady state for 5 min
- 13. Stop the flow and now connect the column to the detector
  - a. Gradually start the flow
    - i. Increase the flow gradually to the desired rate
    - ii. Observe the pressure and detector signal, ensure the pressure and detector signal reach a steady state for 5 min





## [BACK TO THE LAB]

### MASS SPECTROMETER START-UP REFERENCE

 Check the state of the MS detector, vacuum pump(s), connections and waste and ensure it is calibrated/within the preventative maintenance period

2. If required, pump down the MS and wait until appropriate vacuum level is achieved before proceeding

- 3. Check the operations manual; where applicable, run MS Diagnostic tests
- 4. Key parameters to check prior to operation (may vary with MS model)
  - a. Vacuum readbacks
  - b. Gas gauge readbacks
  - c. Voltages switch on and readbacks are correct
  - d. Source temperature setpoint is achieved
  - e. Sensitivity and resolution benchmarks are tuned properly
- 5. Ensure the entire system is working properly
  - a. Run a system qual check and compare to last known results to make sure the system is working properly
    - i. Example: system suitability standard injection or a OC Reference Material



If you are looking for additional information or need assistance with bringing your instrumentation back up after long-term storage,

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