

TEE Complete Probe Reprocessing Infection Prevention Rounding Checklist

OBSERVATION HEADER
Date:
Department:
Survey Team:
Staff Interviewed:
Equipment Observed:
Notes/Instructions:

1. Staff can show where IFUs are accessed and how they are built into the workflow, as needed.
Compliant? <input type="radio"/> YES or <input type="radio"/> NO
2. PPE is readily and easily available where reprocessing activities are taking place.
Compliant? <input type="radio"/> YES or <input type="radio"/> NO
3. Hand hygiene products are available near where the work is being performed and easily accessible.
Compliant? <input type="radio"/> YES or <input type="radio"/> NO
POINT OF USE CLEANING
4. Probe is immediately wiped down after the procedure to remove blood, body fluid and to prevent biofilm with an enzymatic applicator or detergent.
Compliant? <input type="radio"/> YES or <input type="radio"/> NO
INSPECTION
5. Staff can demonstrate how they would inspect the probe for damage, including cracks, degradation, tears, sharp edges and bite marks.
Compliant? <input type="radio"/> YES or <input type="radio"/> NO
6. Staff can verbalize the process if they feel, after inspection, the probe needs to be taken out of service.
Compliant? <input type="radio"/> YES or <input type="radio"/> NO
SOILED TRANSPORT
7. TEE probe is transported in a leak proof, puncture proof container that easily shows it is a biohazard, and secures the probe in place to prevent damage while separating the probe (HLD) and the cord & electrical pack (LLD).
Compliant? <input type="radio"/> YES or <input type="radio"/> NO or <input type="radio"/> N/A (choose N/A if cleaning and disinfection occur in same room as procedure.)
***If automating cleaning and HLD skip to Electrical Leakage Testing

MANUAL CLEANING

The following steps are taken per the IFUs:

8. Correct temperature is maintained for the duration of soaking.

Compliant? YES or NO

9. Probe is soaked for the correct amount of time.

Compliant? YES or NO

10. Correct amount of detergent to water is observed.

Compliant? YES or NO

11. Probe is positioned in a way that only the section that should be under water is under the water/detergent solution.

Compliant? YES or NO

12. Rinsing is performed per the IFUs.

Compliant? YES or NO

13. Probes are dried with a single-use, single-packaged low-linting cloth after cleaning is complete.

Compliant? YES or NO

14. Staff dons and doffs appropriate PPE during manual cleaning, per hospital policy, correctly and safely.

Compliant? YES or NO

MANUAL HIGH-LEVEL DISINFECTION

The following steps are taken per the IFUs:

15. Correct temperature is maintained for the duration of soaking

Compliant? YES or NO

16. MRC testing is performed via test strip and verified.

Compliant? YES or NO

17. MRC testing result is documented for all probes undergoing HLD.

Compliant? YES or NO

18. Expiration date on test strip bottle is observed and validated.

Compliant? YES or NO

19. Staff dons and doffs appropriate PPE during manual cleaning, per hospital policy, correctly and safely.

Compliant? YES or NO

20. Probe is soaked for the correct amount of time

Compliant? YES or NO

21. Probe is positioned in a way that only section that should be underwater is under the water/disinfectant solution

Compliant? YES or NO

22. Rinsing per the IFU
Compliant? <input type="radio"/> YES or <input type="radio"/> NO
23. Probes are dried with a single-use, single-packaged low-linting cloth after removal from the HLD machine or manual HLD solution.
Compliant? <input type="radio"/> YES or <input type="radio"/> NO
24. Documentation is observed to verify all steps of the manual HLD cycle have been performed and were either successful or failed.
Compliant? <input type="radio"/> YES or <input type="radio"/> NO
25. Traceability is verified for patient identification.
Compliant? <input type="radio"/> YES or <input type="radio"/> NO
ELECTRICAL LEAKAGE TESTING
26. Electrical leak testing is performed on each probe after use.
Compliant? <input type="radio"/> YES or <input type="radio"/> NO
27. Results for electrical leak testing are documented for all probe reprocessing cycles (review logbook to confirm).
Compliant? <input type="radio"/> YES or <input type="radio"/> NO
28. Staff can clearly explain the process if electrical leakage testing fails.
Compliant? <input type="radio"/> YES or <input type="radio"/> NO
AUTOMATED HIGH-LEVEL DISINFECTION
The following steps are taken per the IFUs:
29. Probes are dried with a single-use, single-packaged low-linting cloth after removal from the HLD machine or manual HLD solution.
Compliant? <input type="radio"/> YES or <input type="radio"/> NO
30. Documentation is observed to verify all steps of the automated HLD cycle have been performed and either were successful or failed.
Compliant? <input type="radio"/> YES or <input type="radio"/> NO
31. Traceability is verified for patient identification.
Compliant? <input type="radio"/> YES or <input type="radio"/> NO
LOW-LEVEL DISINFECTION
32. Transducer cord, handle and electrical pack are disinfected with a hospital-approved low-level disinfectant.
Compliant? <input type="radio"/> YES or <input type="radio"/> NO
33. Staff can speak to the contact time of the disinfectant used in the prior step.
Compliant? <input type="radio"/> YES or <input type="radio"/> NO
DRYING
34. Probes are dried with a single-use, single-packaged low-linting cloth after removal from the HLD machine or manual HLD solution.
Compliant? <input type="radio"/> YES or <input type="radio"/> NO

STORAGE

35. TEE probe is hung vertically with the distal tip hanging without touching anything. Storage should be in a clean, well-ventilated space (Check the bottom of the cabinet for stains or towels/pads to indicate probes were not dry when hung).

Compliant? YES or NO

36. Staff can speak to the process and frequency of cleaning the probe storage cabinet.

Compliant? YES or NO

37. Storage cabinet filter change is performed per IFU with documentation present.

Compliant? YES or NO

38. Staff can verbalize the maximum hang time or storage time per facility policy before reprocessing in between disinfection events.

Compliant? YES or NO

ENVIRONMENTAL

39. Work flows from dirty to clean.

Compliant? YES or NO

40. Staff can verbalize what to do in the event of a chemical spill in the reprocessing space (check status of spill kit).

Compliant? YES or NO

41. Ceiling tiles are free from stains and damage.

Compliant? YES or NO

42. Flooring is in good repair, free from cracks and damage.

Compliant? YES or NO

43. Reprocessing space shows evidence of negative pressure airflow.

Compliant? YES or NO

MISCELLANEOUS

44. Staff can verbalize how they would trace a probe back to a patient if needed.

Compliant? YES or NO

45. Competencies are performed annually by a qualified person, as well as when there is new equipment, supplies or processes.

Compliant? YES or NO



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