

**\*\*REVISED\*\* 12/26/2024****HSC.34357 Nucleic Acid Extraction/Isolation/Purification****Phase II**

**Nucleic acids are extracted, isolated, and purified by methods reported in the literature, by an established commercially available kit or instrument, or by a method validated by the laboratory.**

*NOTE: Extraction procedures may combine purification or isolation of nucleic acids according to the level of purity needed for downstream applications.*

**Evidence of Compliance:**

- ✓ Records to support nucleic acid extraction/isolation/purification is performed by a validated method

**REFERENCES**

- 1) Sambrook J, *et al.* Molecular cloning: A laboratory manual, second edition. Cold Spring Harbor, NY: Cold Spring Harbor Laboratory Press, 1989:E.3-E.4

**HSC.34544 Reverse Transcription****Phase II**

**For RNA amplification methods, appropriate controls are used for reverse transcription.**

**\*\*REVISED\*\* 12/26/2024****HSC.34731 Specimen Preservation/Storage****Phase II**

**The laboratory uses established methods for specimen preservation and storage before testing.**

*NOTE: Patient samples may be stored in a frost-free freezer only if protected from thawing.*

- Repeated freeze-thaw cycles contribute to biomolecular degradation and are detrimental to biospecimen quality.
- It is prudent to avoid freeze-thaw altogether by aliquoting specimens before freezing.
- Peripheral blood specimens should not be frozen, unless otherwise validated, because induced hemolysis can result in PCR inhibition through the presence of contaminating hemoglobin.

**REFERENCES**

- 1) Rainen L, *et al.* Stabilization of mRNA expression in whole blood samples. *Clin Chem.* 2002;48:1883-1890

**HSC.34760 Dedicated Pipettes****Phase II**

**Dedicated pipettors are used for pre-amplification procedures.**

**\*\*REVISED\*\* 12/26/2024****HSC.34918 Nucleic Acid Quantity and Quality Determination****Phase II**

**The quantity and quality of nucleic acids are determined, when appropriate.**

*NOTE: The quantity and quality of nucleic acids (DNA or RNA) must be measured prior to use in a procedure whose success depends on accurately determining the quantity, concentration, integrity, and/or purity of the nucleic acids. Techniques commonly used to assess nucleic acid quantity and/or quality include electrophoresis, UV/VIS spectrophotometry and fluorescence spectroscopy.*

**Evidence of Compliance:**

- ✓ Defined conditions under which quantity and/or quality of nucleic acid are measured **AND**
- ✓ Records of nucleic acid quantity and/or quality determinations