

Support protocol

 $\label{eq:relation} Filtration \cdot Rapid \ \mbox{Tests} \cdot Water \ \mbox{Analysis} \cdot Chromatography \cdot Bioanalysis \\ Filtration \cdot Schnellteste \cdot Wasseranalytik \cdot Chromatographie \cdot Bioanalytik \\$

Using NucleoSpin[®] RNA XS for difficult-to-lyse tissue (Rev. 03)

This support protocol can be tried for RNA isolation from difficult-to-lyse tissue, such as cerebellum, heart, or skeletal muscle, using the NucleoSpin[®] RNA XS kit.

1. Add **120** μ L of **Buffer RA1** to the sample. Homogenize with **NucleoSpin[®] Filter**, homogenizer, or with a syringe and a needle.

- 2. Incubate 5 min at room-temperature, then vortex well.
- 3. Centrifuge at 14,000 x g for 5 min to remove debris.
- 4. Carefully transfer supernatant to a new tube. Avoid pipetting pelleted material.

5. Add 90 µL of 96 - 100% ethanol to the sample, mix very well by vortexing.

6. Centrifuge at 14,000 x g for 10 min. Discard as much supernatant as possible.

7. Air-dry.

- 8. Add 15 µL of RNase-free water to the pellet and resuspend completely.
- **9.** Add **100** µI of **Buffer RA1**, mix by pipetting and light vortexing.

Continue with step 2 of the standard protocol.

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