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Fresh tissue is preferable for optimal RNA isolation. Alternatively, tissue should be submerged in RNA stabilization reagent immediately after dissection and stored at −80 °C. Add 1 ml TRIzol reagent per 100 mg fresh tissue, mince on ice using sterile scalpels, and homogenize with a sterile pellet pestle probe.

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Streaking for isolation on an agar plate. The pour plate method. Streaking for isolation on an agar plate involves the successive dilution of organisms until you have the cells at a low enough density that single cells are physically isolated spatially to give rise to recognizable individual colonies.

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