

## Green DNA/RNA Dry Transport

Cat No R1141

## (I) Sample Preparation

- Perform all steps at room temperature and centrifugation at 10,000-16,000 x g for 30 seconds, unless specified.
- 1. Mix sample 1:1 with DNA/RNA Shield<sup>™</sup> (2X Concentrate).

Example: Mix 50 µl sample and 50 µl DNA/RNA Shield<sup>™</sup> (2X Concentrate).

Add an equal volume of ethanol (95-100%) and mix well.

Example: Add 100 µl ethanol.

- Transfer the sample to the Zymo-Spin<sup>™</sup> IC Column in a Collection 3. Tube and centrifuge. Discard the flow-through.
- 4. Add 700 µl Transport Wash Buffer to the column and centrifuge. Discard the flow-through.
- 5. Remove the column and place it into a new collection tube. Seal the column (parafilm).
- 6. Transport/store at ambient temperature.

## (II) Sample Elution

- Perform all steps at room temperature and centrifugation at 10,000-16,000 x g for 30 seconds, unless specified.
- 1. Add 700 µL ethanol (95-100%) to the column and centrifuge. Carefully transfer the column into a DNase/RNase-free tube (not provided).
- Add ≥ 15 µL DNase/RNase-Free Water directly to the column matrix. and centrifuge.

Alternatively, for highly concentrated samples use  $\geq 6 \ \mu l$  elution. The eluate can be used immediately or stored frozen. No clean-up required.



