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Agarose gel for RNA and DNA

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ABSTRACT

General method for running agarose gels for RNA and DNA samples

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Protocol status: Working We use this protocol and it's working

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Preparing the agarose gel and plate

4m 30s

- 1 A 1 g agarose in A 100 mL of
 - X TAE (Tris-Acetate-EDTA) buffer, 1x Contributed by users
- Microwave solution in a flask for 00:03:00 until the agarose is dissolved. Microwave for 00:01:00 thereafter 00:00:30

4m 30s

Allow to cool before adding A 1 µL

Ethidium bromide, 10mg/mL, 10mL **Amresco Catalog #X328- 10ML**

in the gel medium

before completely cool. Then pour into the gel mold

3.1 Make sure that your plate is balanced and the teeth are in. Decide if you need a long ladder or a short weather.

Preparing the chamber

Add Δ 5 μL Ethidium bromide, 10mg/mL, 10mL Amresco Catalog #X328- in the

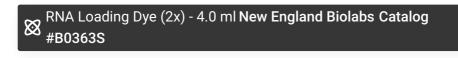
gel medium within the chromatography chamber

Preparing RNA samples

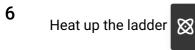
5m

V1 = (10)(50)/RNA conc. (from nanodrop)

Add the V1 sample of RNA to 1:1 ratio of



to each sample



RiboRuler High Range RNA Ladder Thermo Fisher Catalog #SM1821

and RNA samples to

Preparing DNA samples

7 SyberSafe DNA Gel Stain Invitrogen - Thermo Fisher Catalog #S33101

is already in the

samples for the PCR; then use the

GeneRuler 1 kb Plus DNA Ladder Thermo Fisher Catalog #SM1331

Setting up the sample gels

1h 50m

5m

- 8 Add \bot 5 μ L of ladder and samples to gel wells
- 9 For RNA: 80V for (5) 01:00:00 For DNA: 80V for 👏 00:40:00

1h 40m

10 *Note: can check gel after 00:10:00 to see if there is product showing on the gel 10m