

■ Announcements

- ❖ Sign up for journal club (D8 only)
- ❖ FNT: draft introduction, new methods.

■ Lab Quiz

■ Pre-lab Lecture

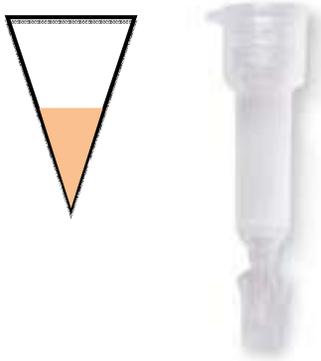
- ❖ Today in Lab: M1D4

M1D4 Workflow

**Remember to use RNase-free equipment and technique!
Keep all RNA on ice when not in use.**

1. DNase treatment (30'), prepare spin columns

We will check your HW calcs
and let you know if correct.



2. Measure [RNA] using spec.

Calculate if you have enough
to proceed – talk to us if not!



3. Dilute and denature RNA Goal: start by ~ 2:30-3



4. Incubate RNA with beads Goal: start by ~ 3-3:30

Add tRNA to “cover” agarose
(RNA binds non-specifically)



5. Run RNA through column 6. Begin RNA precipitation

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20.109 Laboratory Fundamentals in Biological Engineering
Spring 2010

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