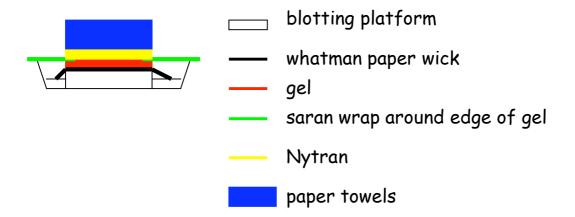
## NORTHERN BLOTTING

- 1. Run RNA on formaldehyde gel.
- 2. Measure fluorescence on Biorad Multi-fluor and photograph gel.
- 3. Soak gel for 15' at room temperature in  $20 \times SSC$ .
- 4. Cut a piece of Nytran and two pieces of filter paper the size of the gel. Wet Nytran in  $2 \times SSC$ .
- 5. Set up blotting tray with  $20 \times SSC$  in tray as follows:



- 6. Leave overnight at room temperature.
- 7. Remove filter and x-link in Stratalinker.
- 8. Photograph filter.
- 9. Either use immediately or store at room temperature between whatman filter paper.

## 20 x 55C (2L)

3 M NaCl 351 g 0.3M sodium citrate 176 g pH to 7.6 with HCl