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Diffusivity measurement with dual chamber diffusion cells

- 1. The dual chamber diffusion cell is constructed from acrylic. Each chamber is 38.1 mm (= 1.5 inch) in diameter with height 25.4 mm (= 1 inch). On the side, there is a ~10 mm (~0.4 inch) hole (throughout experiments, the hole is covered by ducktape to prevent evaporation except when pipetting) for depositing and withdrawing solutions.
- 2. The BSA solution is premixed to a concentration of 2 g/L and glycine 1.5 g/L.
- 3. 28 mL of 2 g/L BSA solution (or 1.5 g/L glycine solution) is added to one chamber and 28 mL pure DI water is quickly added in the other chamber.
- 4. The dual chamber diffusion cell sits on a shaking incubator at 250 RPM and 25 ℃.
- 5. For BSA diffusivity measurements, starting 2.5 hours after seeding the solution, 150 μ L of solution is taken from both chambers and measured at OD280. For the glycine diffusivity measurement, 150 μ L of solution is taken from both chambers and measured for OD200 every 30 min after seeding. Immediately after measurement, the 150 μ L solution is pipetted back into the chamber from which it was taken and the loss of solution is negligible.