## Prtcl\_Washing MiniON flow cell

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## There are 3 methods and choose one of them for your washing step

- ❖ Protocol 1: To store for later use
- 1. Open the priming cover and slowly remove the all of the previous buffer from the waste port
- 2. Add 150ul of Solution A to the Priming port (넣으면 밀려나옴.)
- 3. Incubation for 10 min
- 4. Add 500ul of Solution S to the Priming port
- 5. Close the priming cove and Store at 4 °C
- ❖ Protocol 2: Add the next library immediately
- 1. Open the priming cover and slowly remove the all of the previous buffer from the waste port
- 2. Add 150ul of Solution A to the Priming port (넣으면 밀려나옴.)
- 3. Incubation for 10 min
- 4. Priming port로 Add 150ul of Solution B 넣고
- 5. Ready for the next library
- ❖ Protocol 3: No washing kit but need to use flow cell immediately
- 6. Just remove the waste buffer
- 7. Add 800 ul of priming solution into the flow cell
- 8. Then directly add your next DNA library

https://www.youtube.com/watch?v=IRTdK0kl9v4