

## **Stripping westerns**

To probe the same immunoblot with a different primary antibody:

1. Strip the blot in 50-100 ml stripping buffer (0.2M glycine pH 2) for 20 min (room temperature), with agitation.
2. Wash x3 in PBS
3. Blocked in 5% Marvel/PBS and hybridize with 1° and 2° antibodies as per normal procedure.