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## Co-immunoprecipitation

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**Protocol status:** Working

We use this protocol and it's working

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## Abstract






This protocol describes a common procedure to perform co-immunoprecipitation

## Troubleshooting



## Co-immunoprecipitation

1h 10m

- 1 Media fractions were collected and centrifuged at 1000×g for  00:10:00 
- 2 The supernatant fractions were collected and concentrated (20-fold) using a 10 kDa Amicon filter.
- 3 Concentrated media fractions (1 ml) were incubated with 20 µl of anti-FLAG M2 affinity gel for  01:00:00 at  4 °C 
- 4 After washing 5 times with lysis buffer , SDS-PAGE sample loading buffer was added to the beads (twice the beads volume) and samples were processed for SDS-PAGE and immunoblot.