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Co-immunoprecipitation

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ABSTRACT

This protocol describes a common procedure to perform co-immunoprecipitation

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Protocol status: Working We use this protocol and it's working

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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 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.