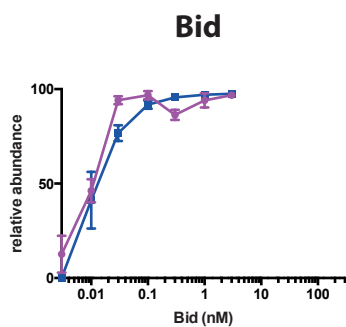
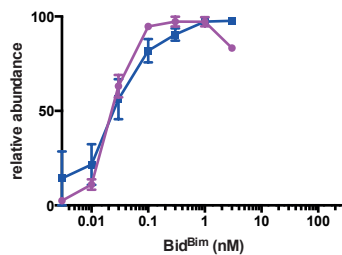
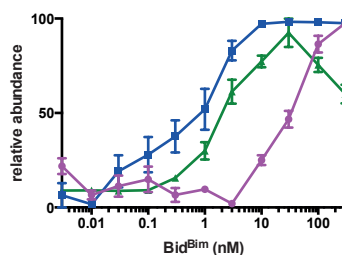
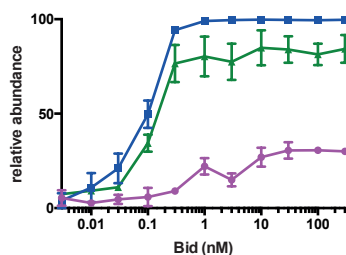


Titration

- Mcl-1

**Bid<sup>Bim</sup>**

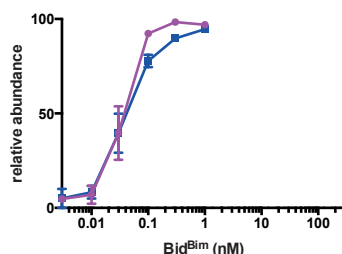
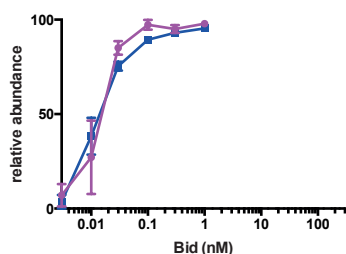
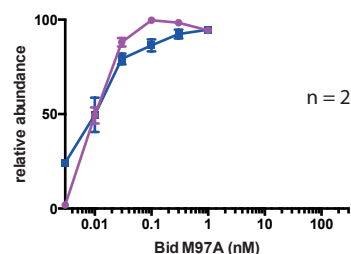
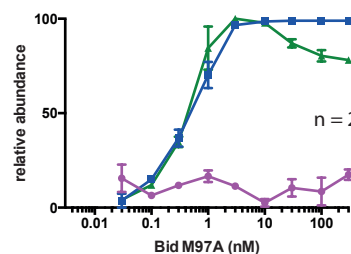
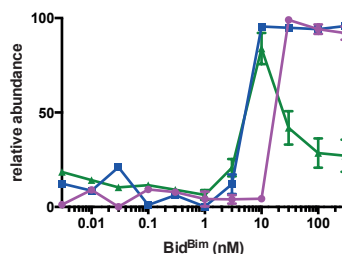
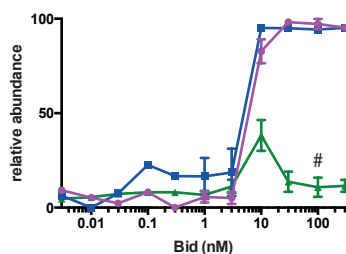
+ Mcl-1



**legend:**

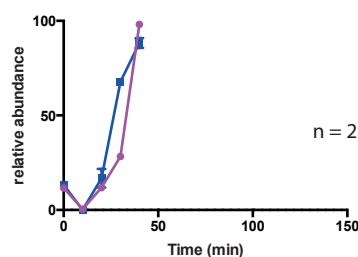
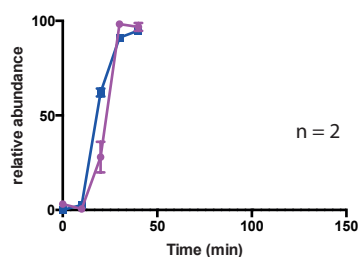
- cytochrome c release
- Bak activation (PK)
- Bak:Mcl-1 (IP)

Titration

- Bcl-x<sub>L</sub>**Bid M97A**+ Bcl-x<sub>L</sub>

# Bid 100 nM IP n = 2

- Mcl-1



mean +/- SEM  
n = 3 except where indicated

Cytochrome c release calculated separately from supernatant and pellet as for Direct Activation paper, but then averaged before plotting.

Bak activation is calculated as loss of the non-activated Bak band (not measured by gain of lower bands)

Bak:Mcl-1 interaction is measured by the relative intensity of the IP: Mcl-1 WB: Bak blots. In all cases there were comparison bands on different blots so the relative IP is consistent across plots from the same type of experiment.

Time course

+ Mcl-1

