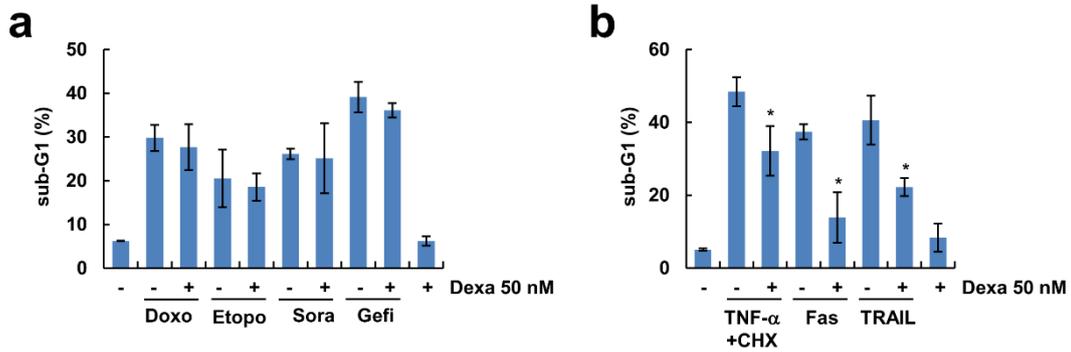


1 *Supplementary Materials*

2 **Dexamethasone Inhibits TRAIL-Induced Apoptosis**
3 **through c-FLIP(L) Upregulation and DR5**
4 **Downregulation by GSK3 β activation in Cancer Cells**

5 **Mi-Yeon Jeon, Seon Min Woo, Seung Un Seo, Sang Hyun Kim, Ju-Ock Nam, Shin Kim,**
6 **Jong-Wook Park, Peter Kubatka, Kyoung-jin Min, and Taeg Kyu Kwon**

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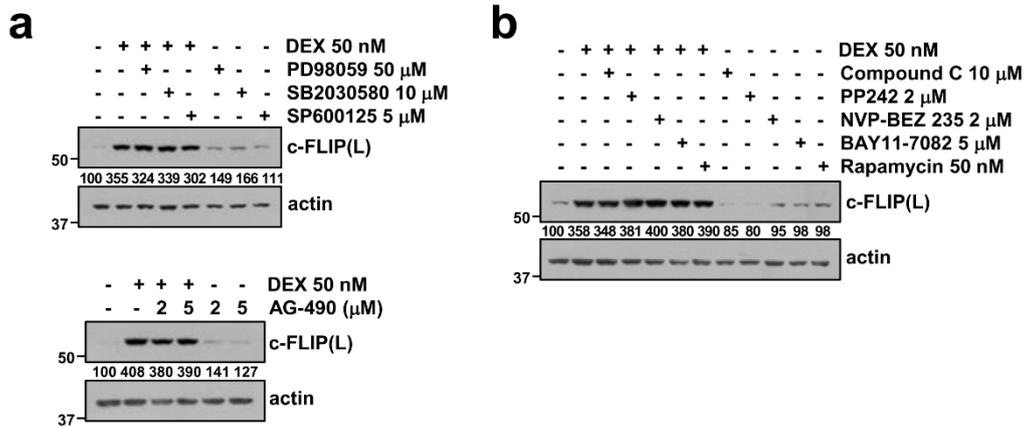
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Figure S1. Effect of DEX on anti-cancer drugs-induced cell death. **(a,b)** Caki-1 cells were pretreated with the indicated concentrations of DEX for 1 h, and then treated with anti-cancer drugs (doxorubicin, etoposide, sorafenib and gefitinib) **(a)** or death receptor-associated drugs (TNF- α +CHX, Fas and TRAIL) **(b)** for 24 h. Apoptosis was determined by flow cytometry.



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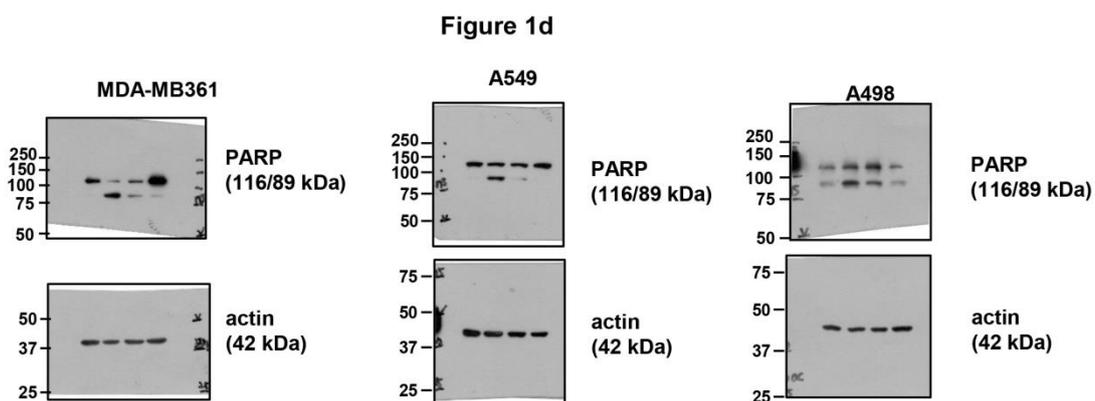
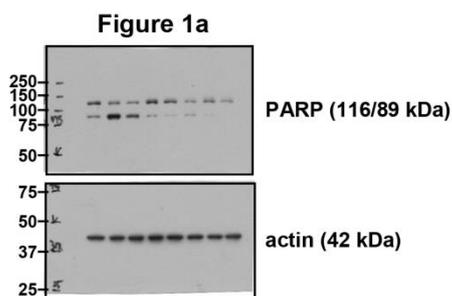
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Figure S2. Effect of various multiple kinases inhibitors on DEX-mediated c-FLIP upregulation. **(a,b)** Caki-1 cells were pretreated with MAPKs inhibitors [PD98059 (MEK inhibitor), SB203580 (p38 MAPK inhibitor), and SP600125 (JNK inhibitor)], JAK/STAT inhibitor (AG-490) **(a)**, AMPK inhibitor (compound C), mTOR kinase inhibitors (PP242, NVP-BEZ235 and Rapamycin) and NF- κ B inhibitor (BAY11-7082) **(b)** for 1 h, and then treated with 50 nM DEX for 24 h. Protein expression was determined by western blotting.

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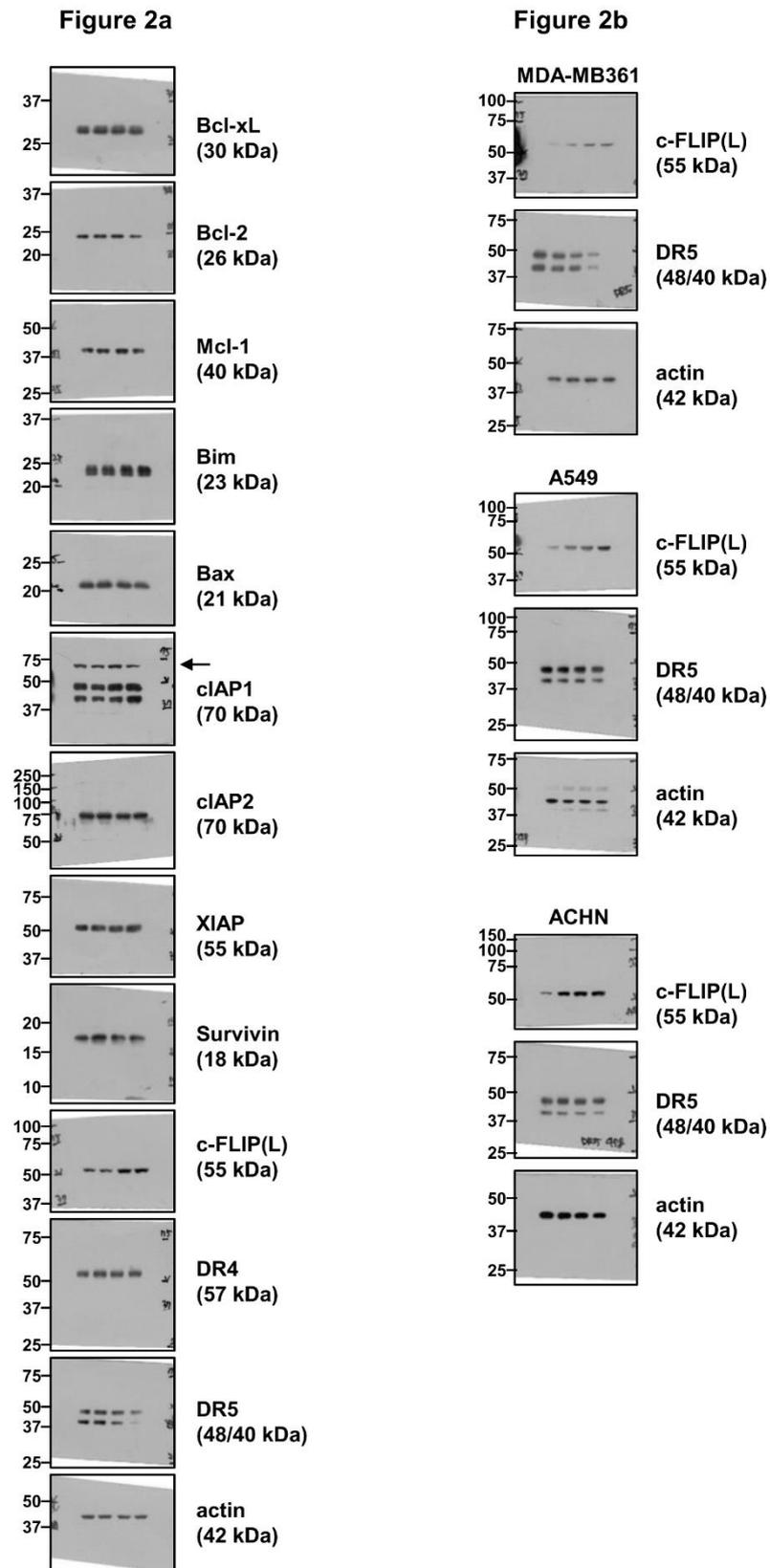
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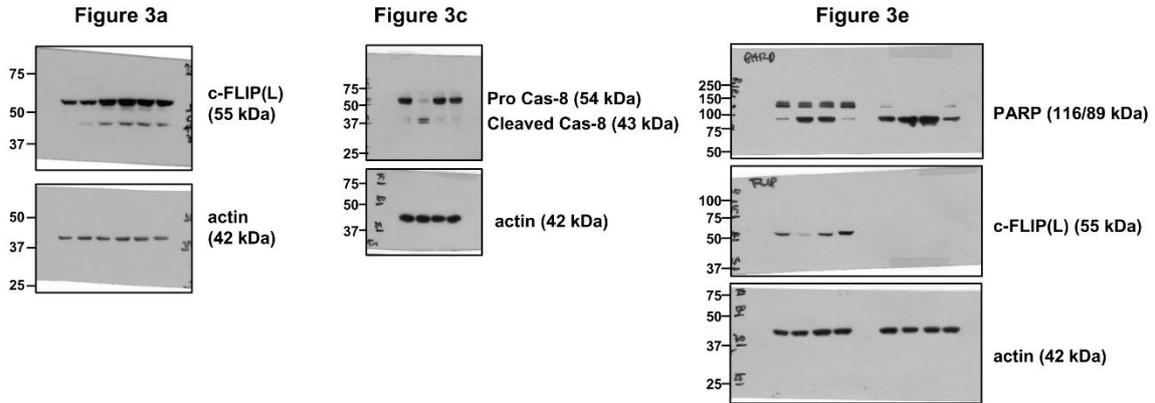
Figure S3. Uncropped western blots for Figure 1.



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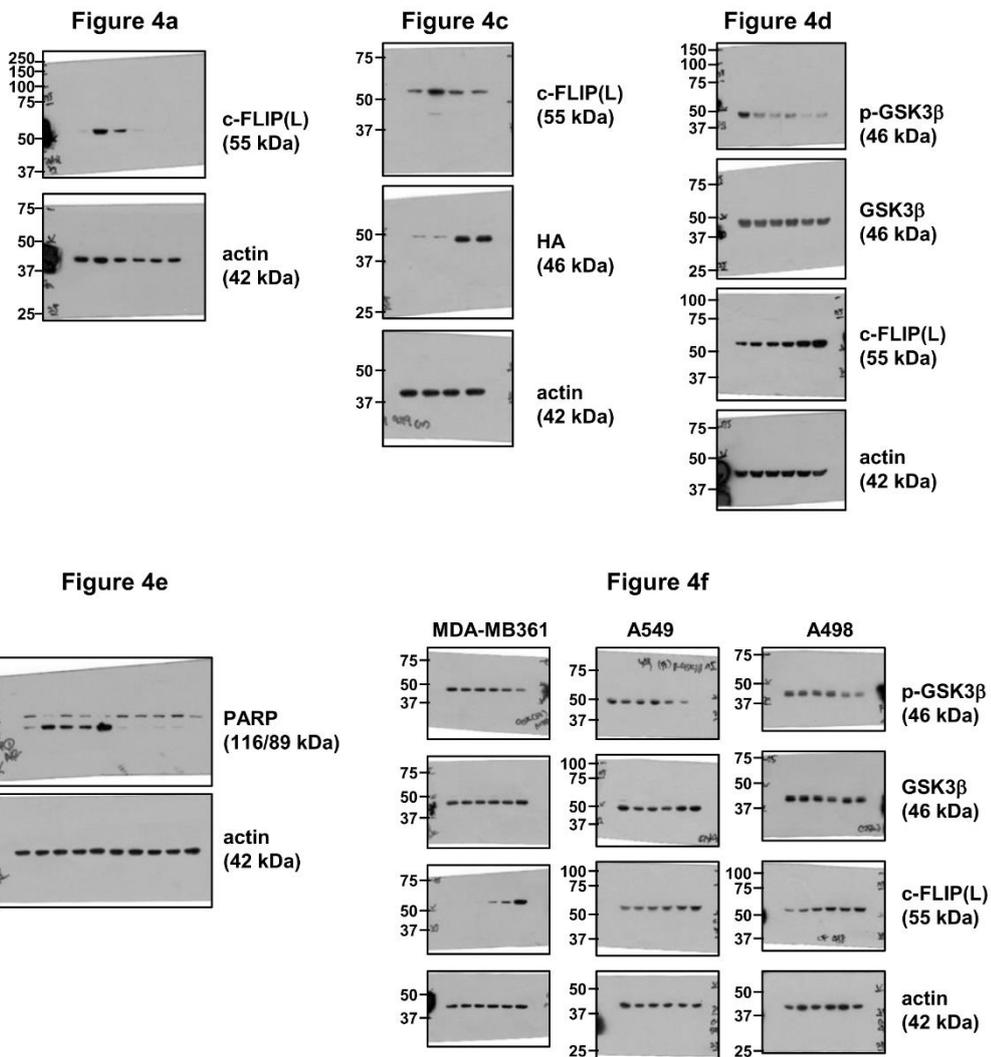
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Figure S4. Uncropped western blots for Figure 2.



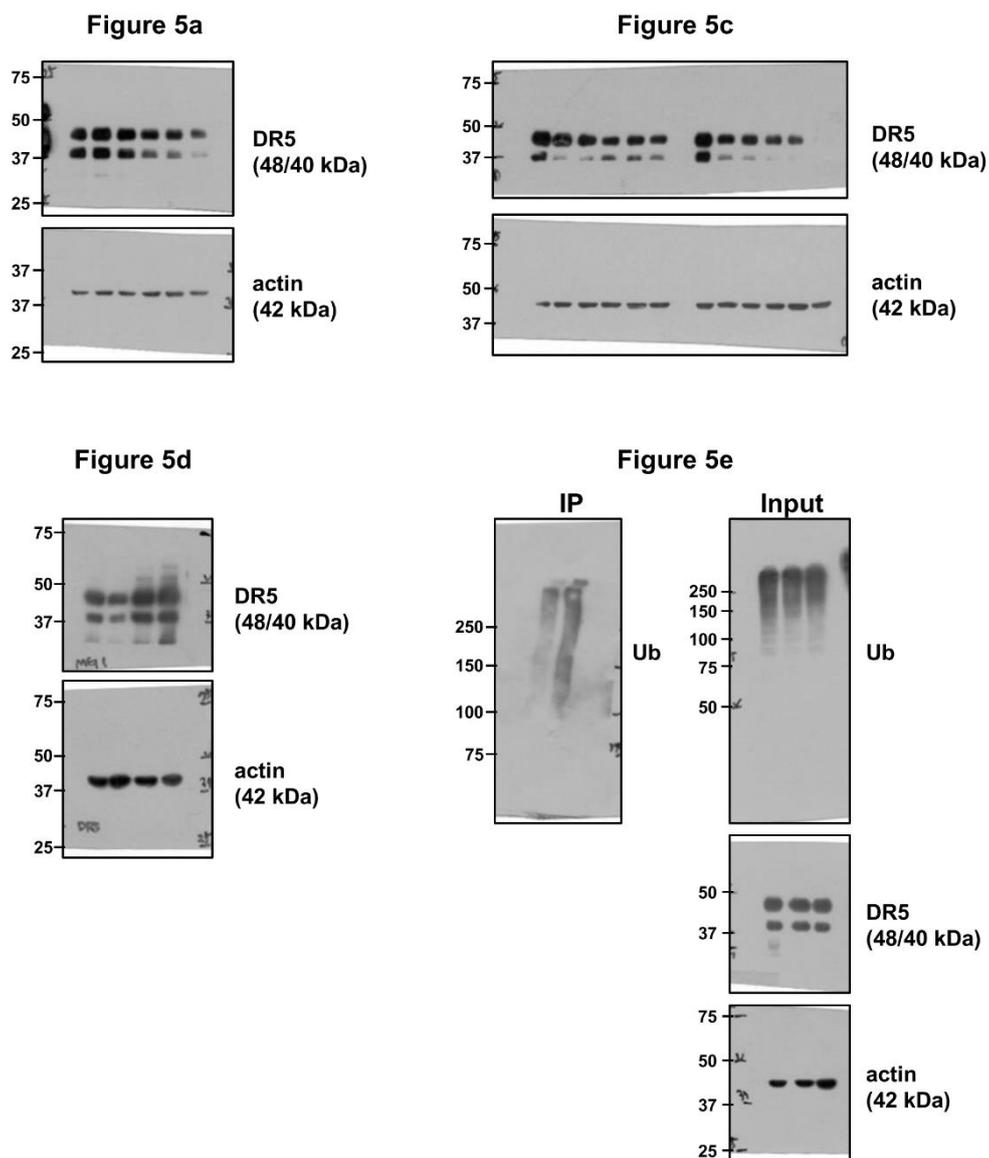
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Figure S5. Uncropped western blots for Figure 3.



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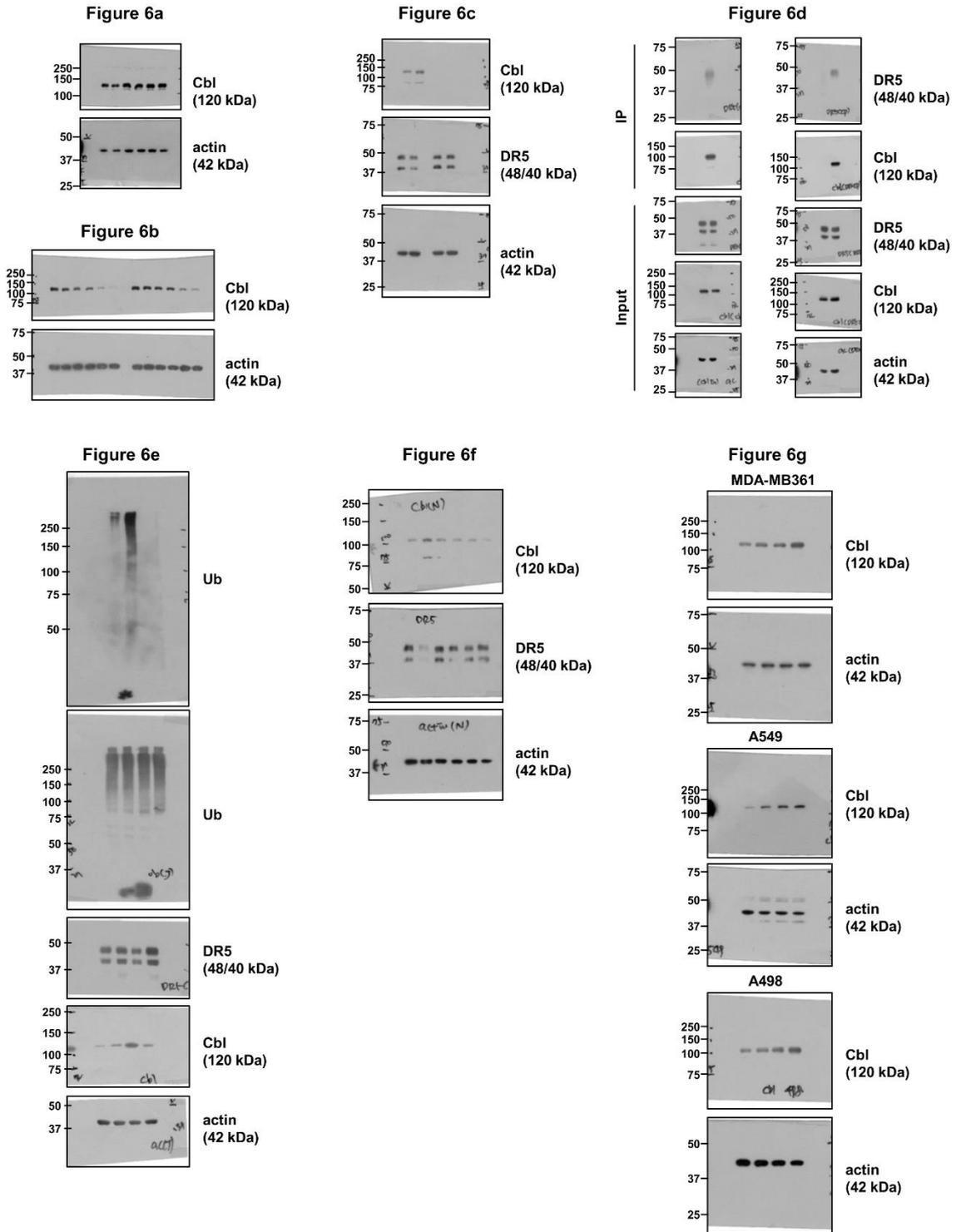
Figure S6. Uncropped western blots for Figure 4.



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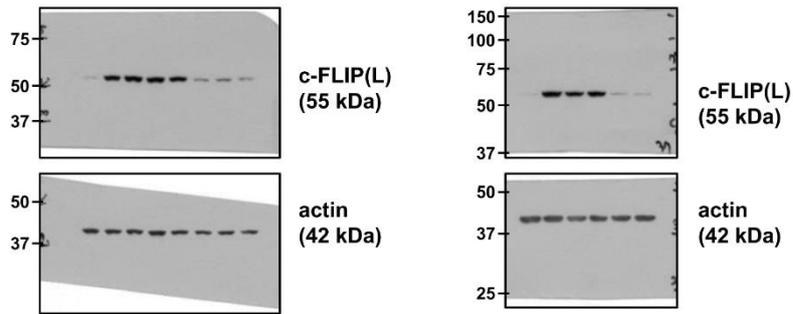
Figure S7. Uncropped western blots for Figure 5.



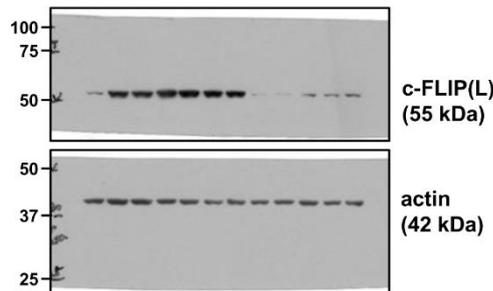
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Figure S8. Uncropped western blots for Figure 6.

Supplementary Figure 2a



Supplementary Figure 2b



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36 **Figure S9.** Uncropped western blots for Figure S2.

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