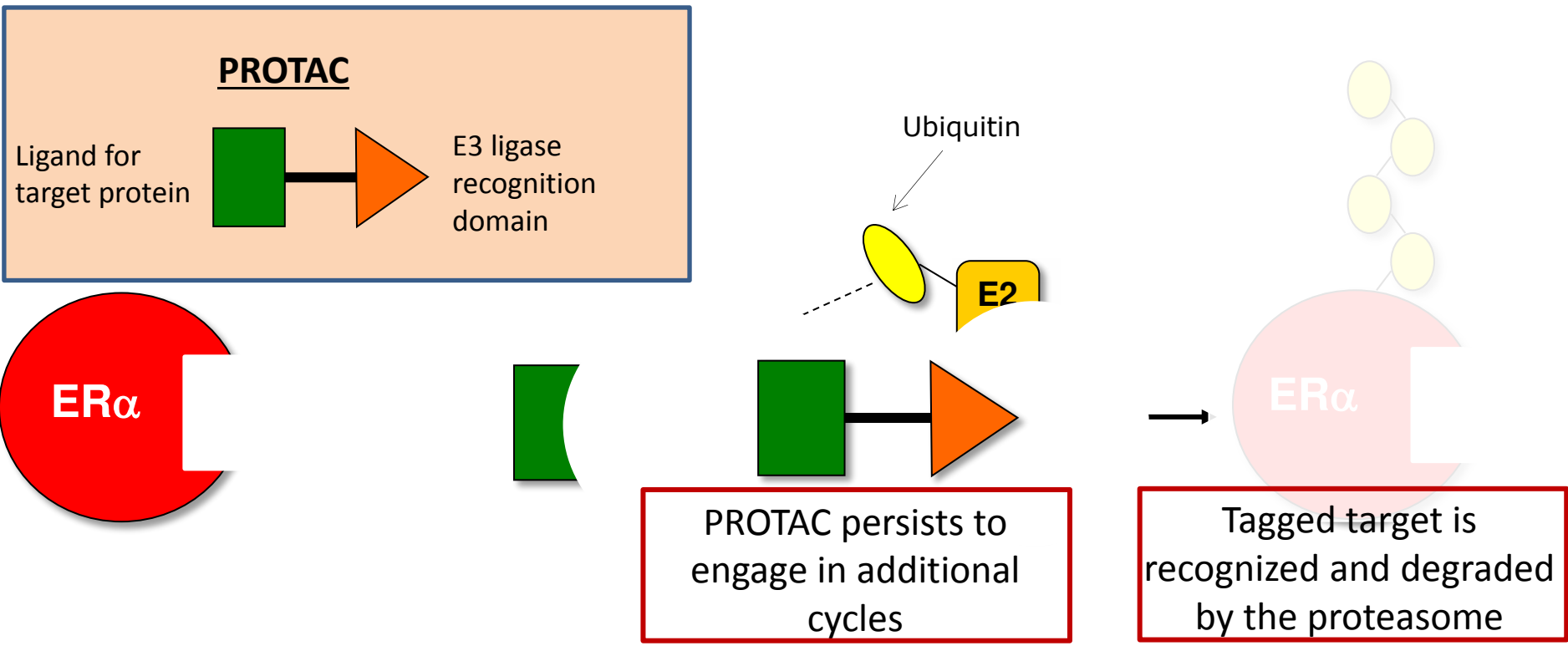


Targeted and Selective Degradation of ER α by PROTACs

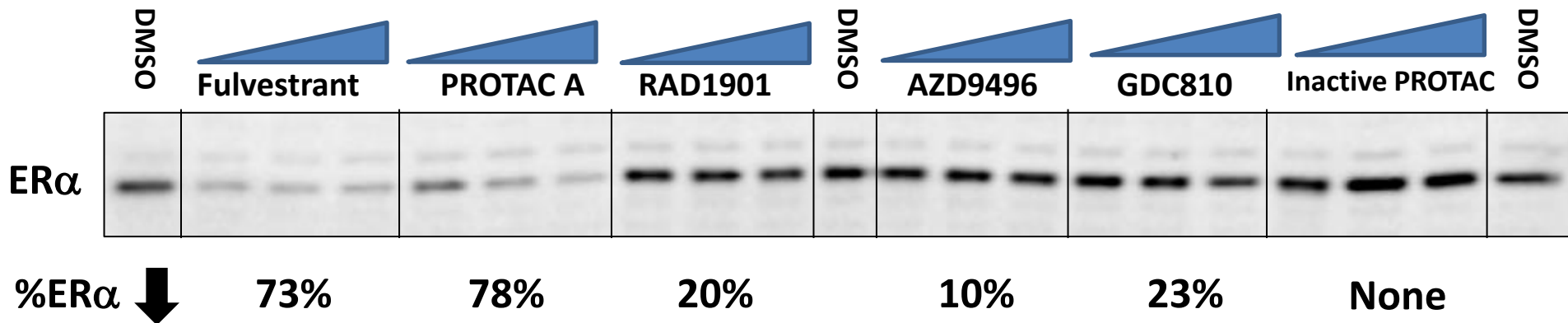
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PROTAC: PROteolysis Targeting Chimera

- Chimeric, small molecule PROTACs recruit E3 ligase to target proteins to promote polyubiquitination and induce their degradation via the proteasome

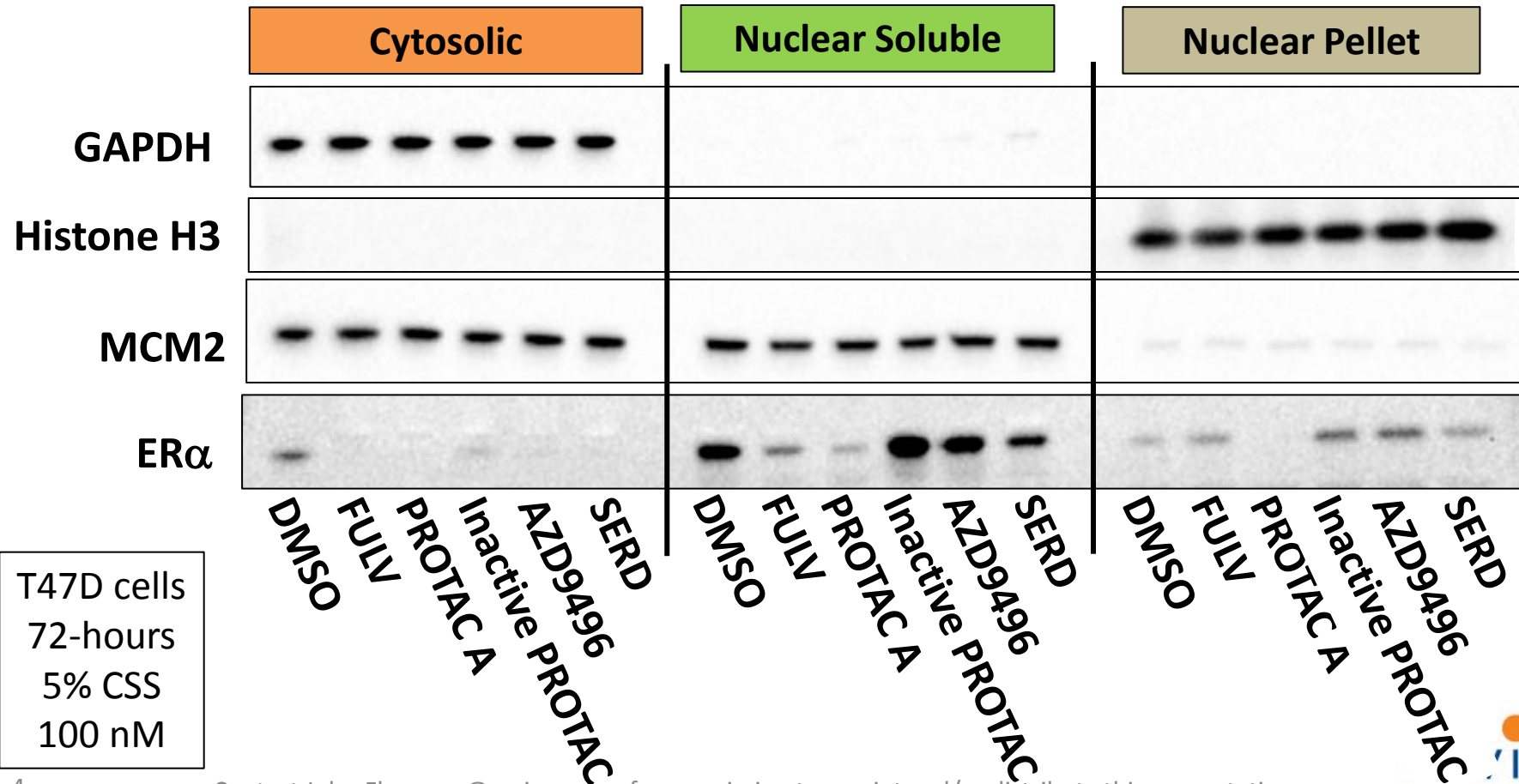


PROTAC potently reduces ER α levels in T47D cells

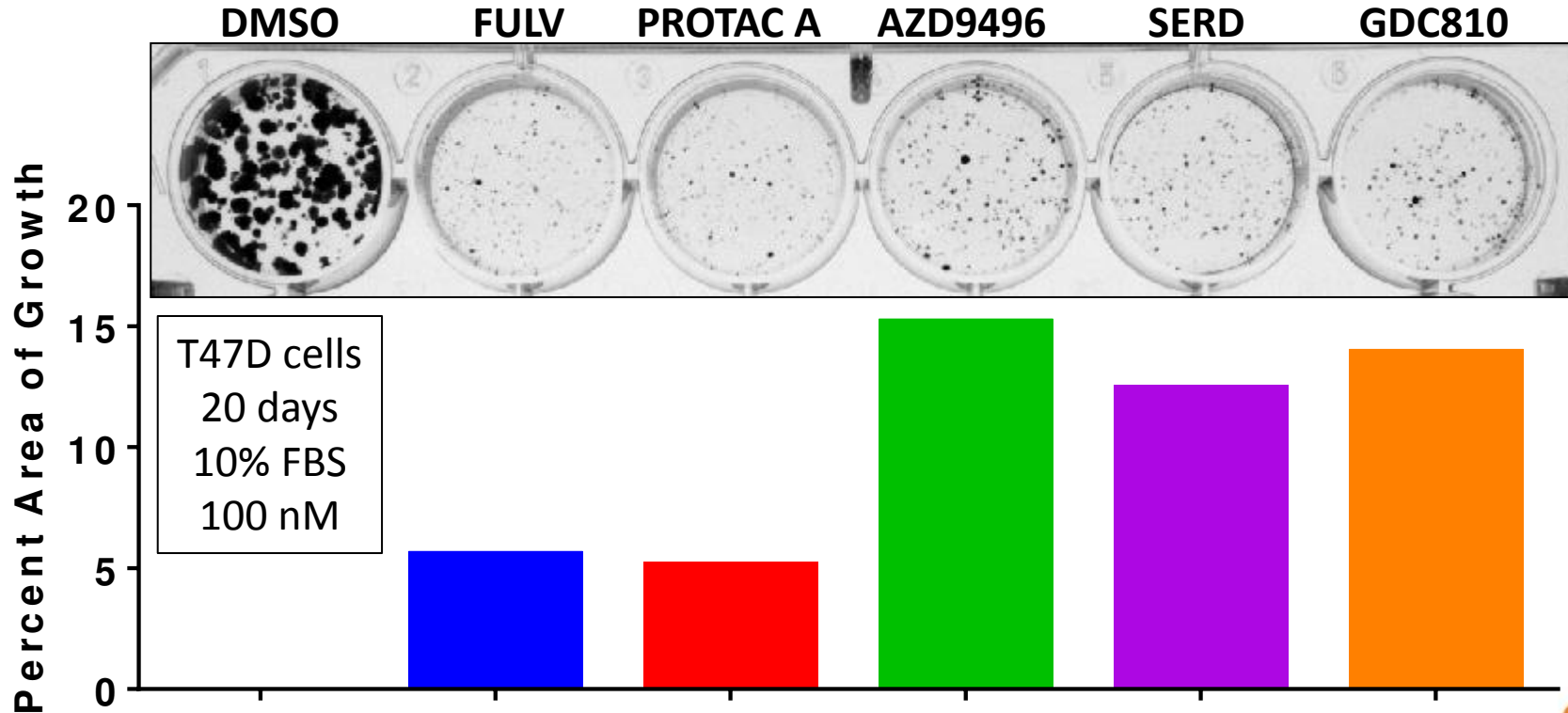


- Robust degradation of ER α with PROTACs and fulvestrant in other ER+ cell lines (BT474, HCC1428, MDA-MB-361, MCF7, MDA-MB-134-VI)
- Inactive PROTAC binds to ER α but not to any E3 ligase
- Concentrations tested were 6, 25, and 100 nM

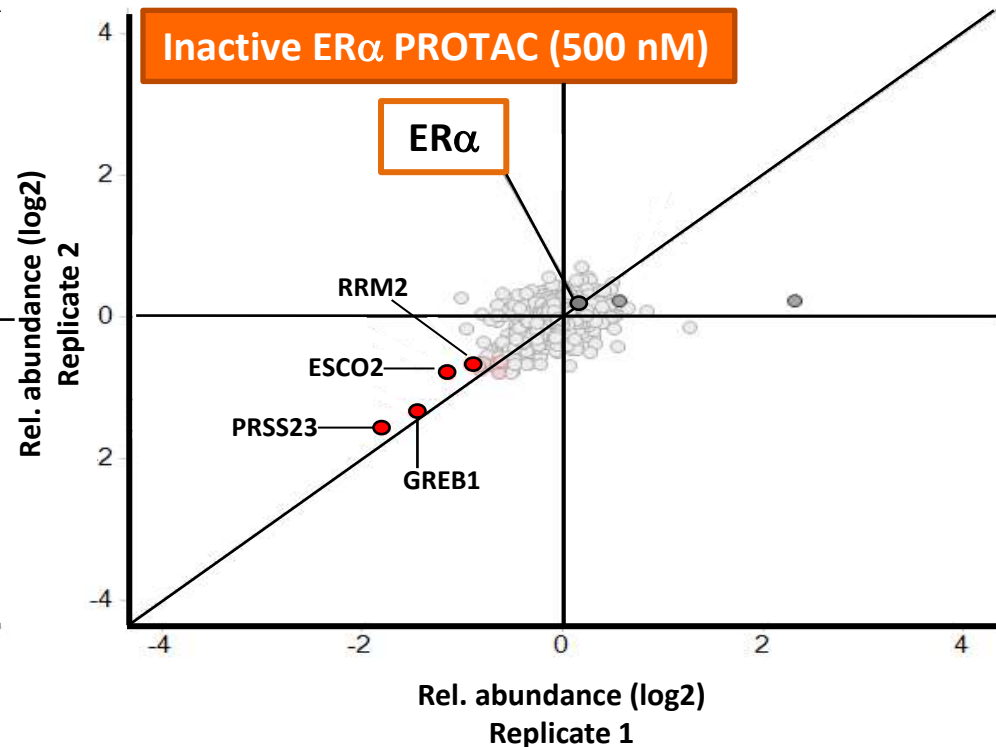
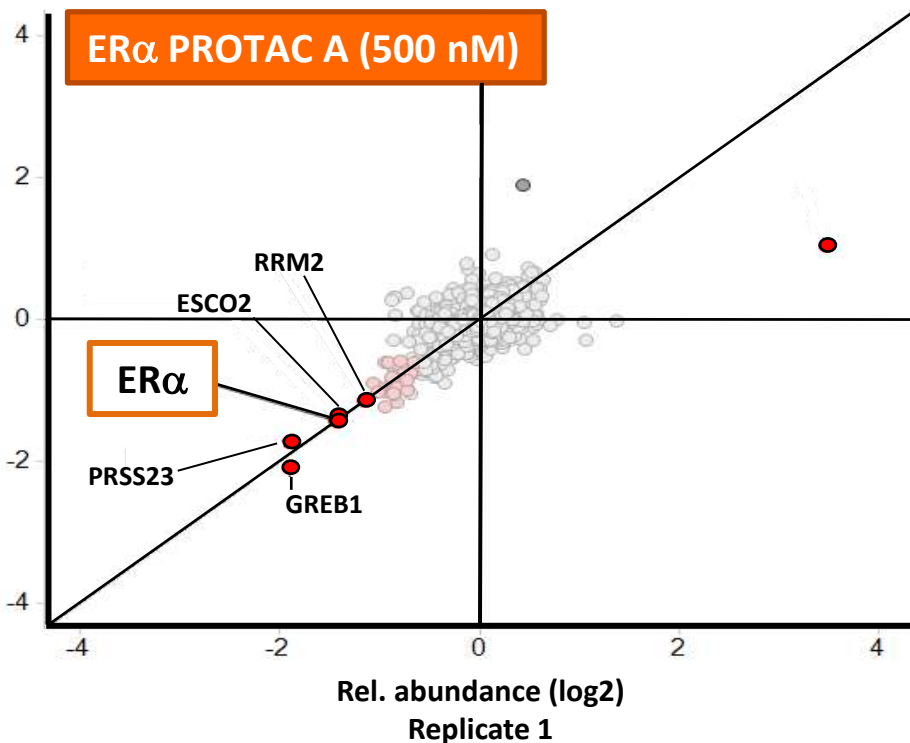
PROTAC reduces ER α levels in nuclear fractions



Greater anti-proliferative effects with ER α degraders



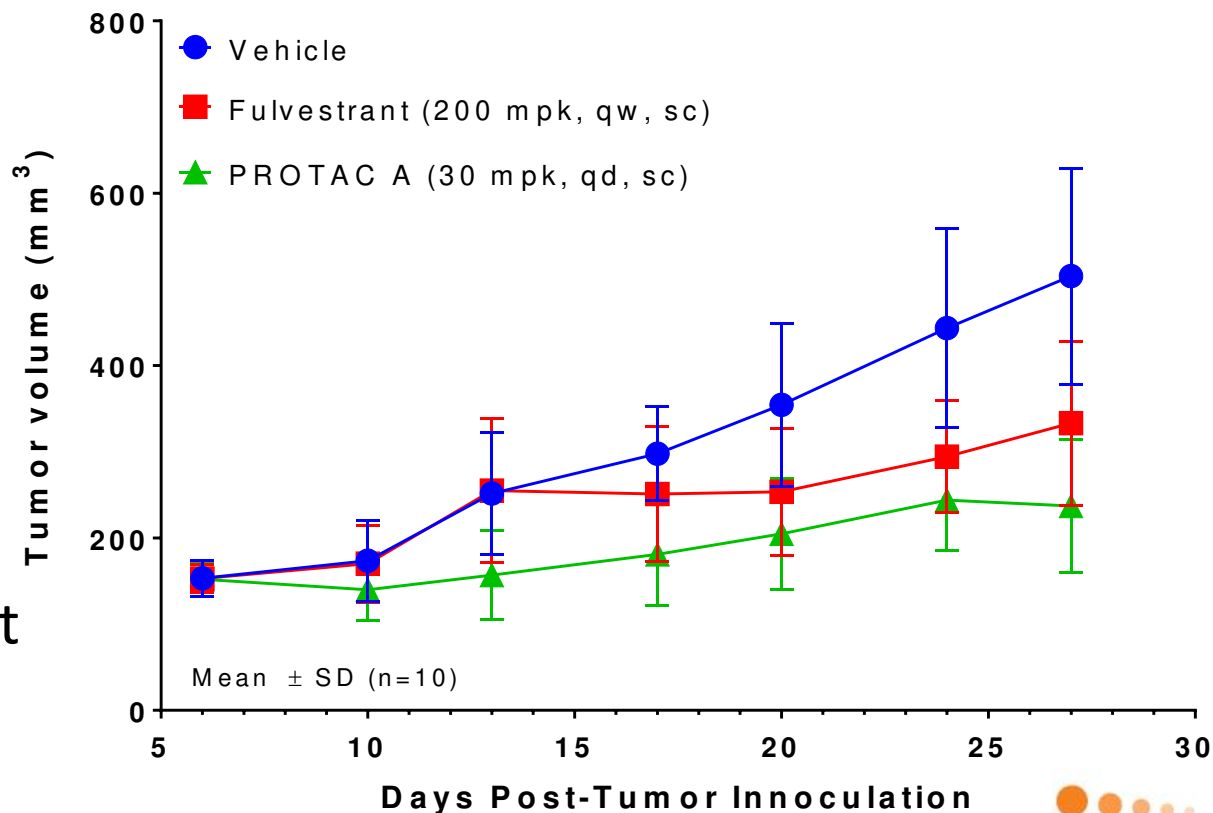
Selective degradation of ER α by PROTACs



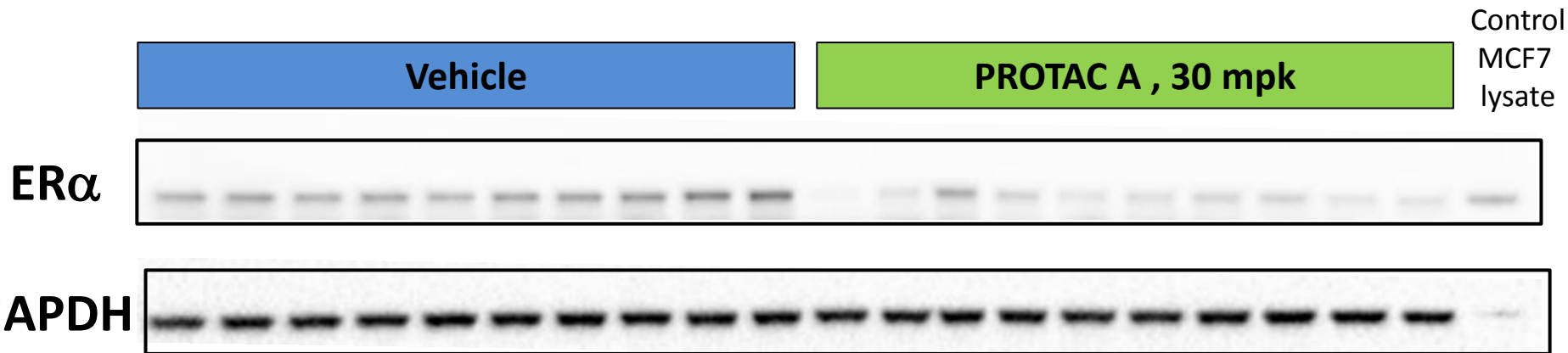
LC/MS quantitation of >7600 proteins after overnight incubation
of MCF7 cells with PROTACs

ER α PROTAC inhibits growth of MCF7 xenografts

- 21-day growth study
- 68% TGI achieved with daily 30 mpk PROTAC (subcutaneous)
- Estradiol required for model to grow so PROTAC anti-endocrine activity is likely better at post-menopausal E2 levels

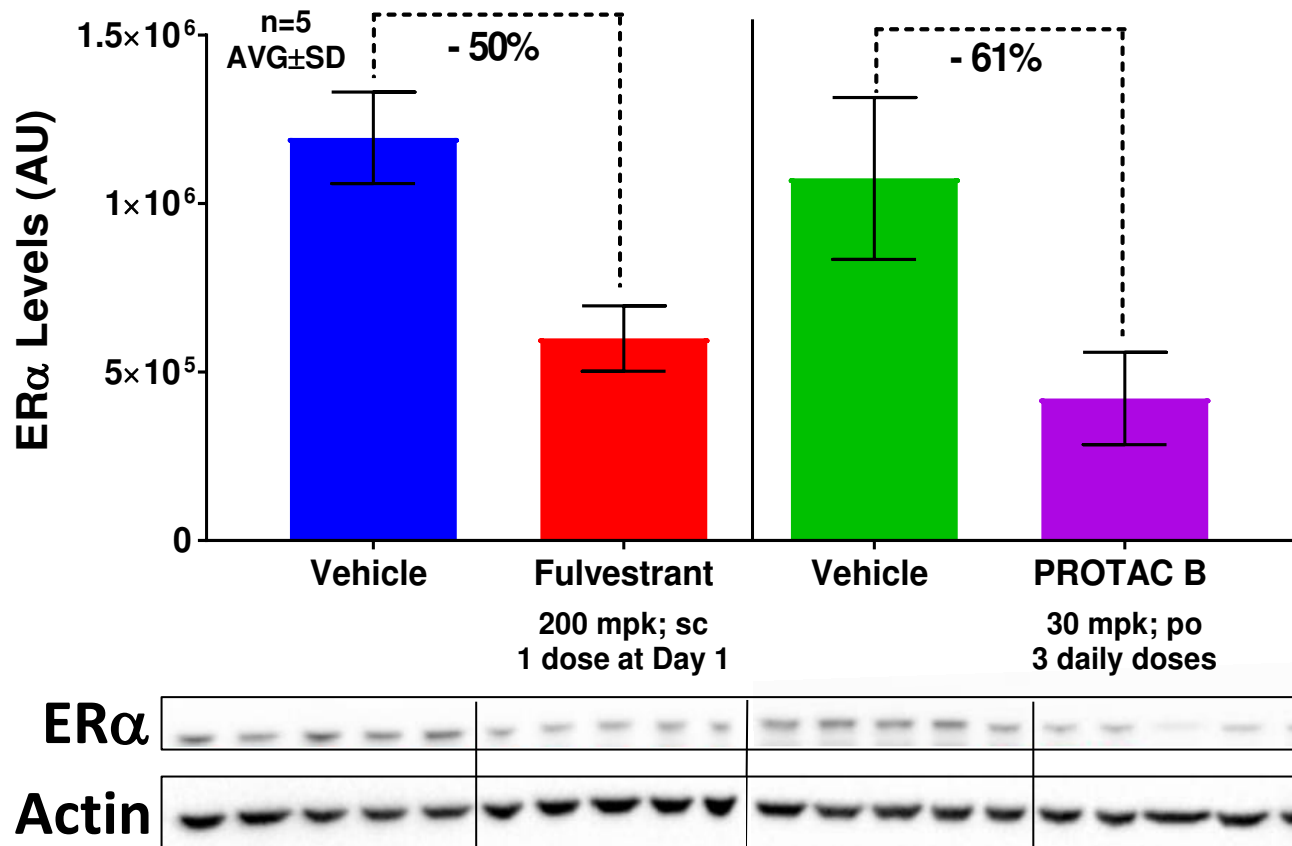


PROTAC decreases ER α levels in MCF7 xenografts



- ~60% reduction in ER α levels with PROTAC A (30 mpk, s.c.)
- Tumors were harvested four hours post-dose

ER α degradation after oral administration of PROTAC



- MCF7-estradiol deprived cells implanted in adult female mice
- Similar reduction of ER α also seen in immature rat uteri

Summary

- PROTAC-mediated degradation provides a distinct mechanism for reducing ER α when compared to fulvestrant and current clinical SERDs
- PROTACs reduce ER α levels in all ER+ cell lines tested
- Oral administration of PROTACs reduced ER α levels in ER+ xenografts and in rat uteri
- Currently evaluating oral PROTACs in a variety of clinically-relevant ER+ BC models

Targeted and Selective Degradation of ER α by PROTACs

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