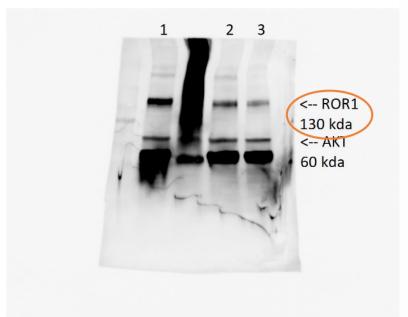
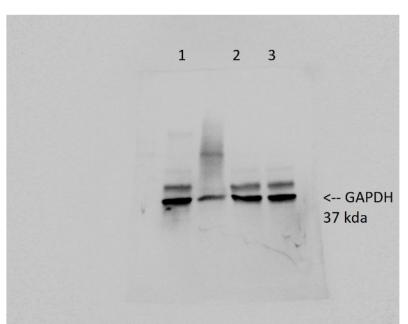
## Raw western blots (Figures 1-8)

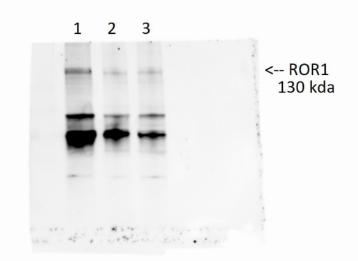
- All lanes used in the manuscript and used for densitometry analysis (N=3) are labeled.
  Unlabeled lanes were not used for analysis.
- 2. All proteins used in the manuscript and used for analysis are appropriately labeled. Unlabeled proteins were either not used for analysis or were labeled in subsequent slides.
- 3. All protein lysate was obtained from PC3 cells.
- 4. All densitometry analysis was normalized to housekeeping protein (GAPDH).
- 5. Blue lines indicates how the blot was spliced for final presentation.
- 6. Single blue line indicates where the blot was physically cut.
- 7. Red boxes indicate how the blot was cropped for publication (approximately).
- 8. Labeling:
  - Lanes
    - 1: Control
    - 2: Strc 250  $\mu$ M- 24h treatment
    - 3: Strc 250  $\mu$ M- 48h treatment
  - Proteins
    - pGSK3b = phospho-GSK3β (ser9)
    - pAKT = phospho-AKT (ser473)











1

2 3

Figure 1

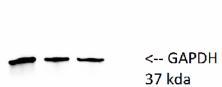
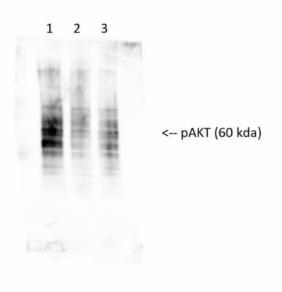
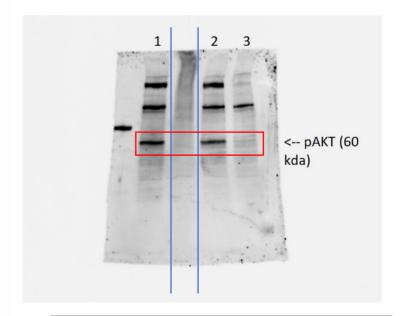
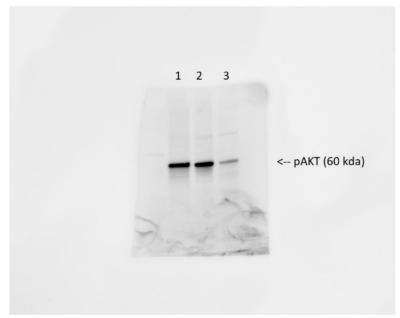


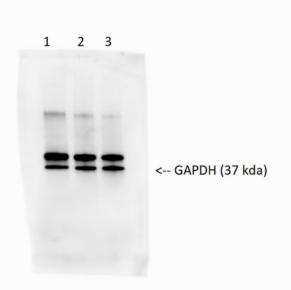
Figure 2

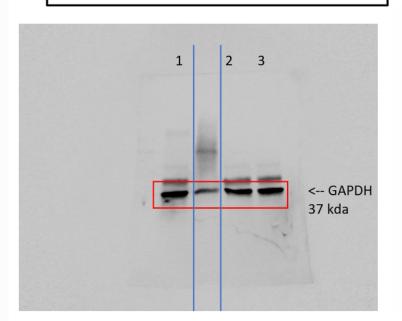


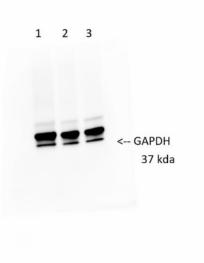




This replicate was used in Figure 2c of the manuscript







Vertical blue lines indicate how the blot was spliced for final presentation

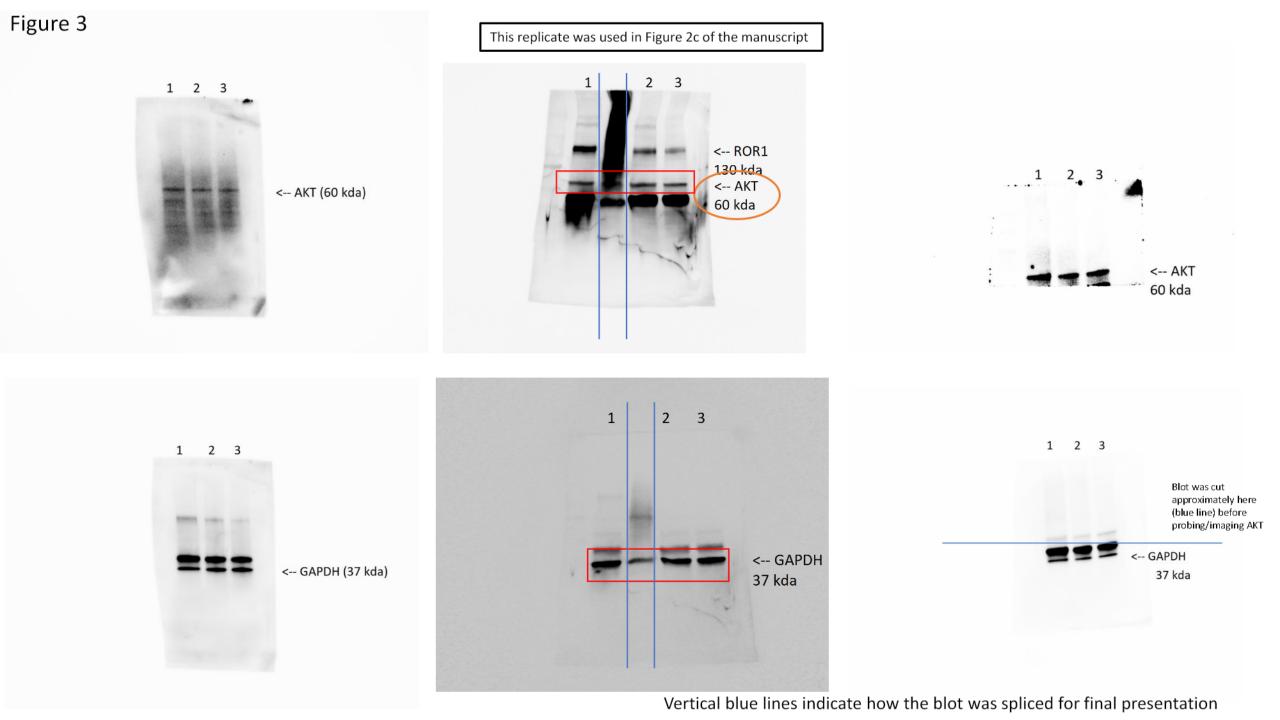
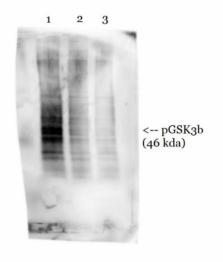
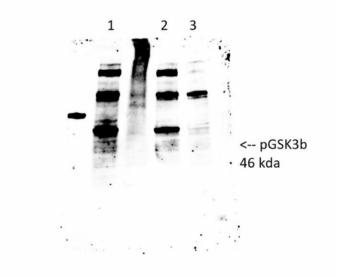
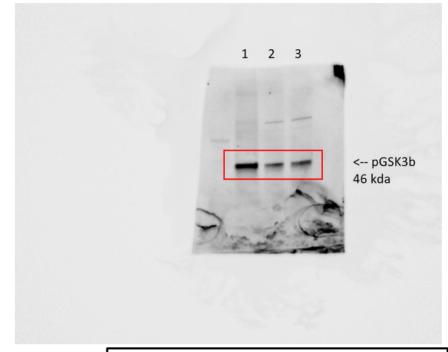


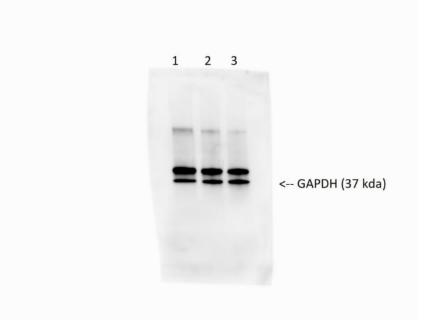
Figure 4

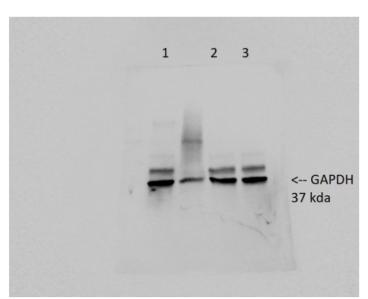


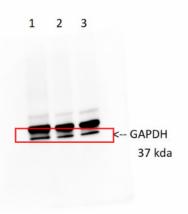




This replicate was used in Figure 2c of the manuscript







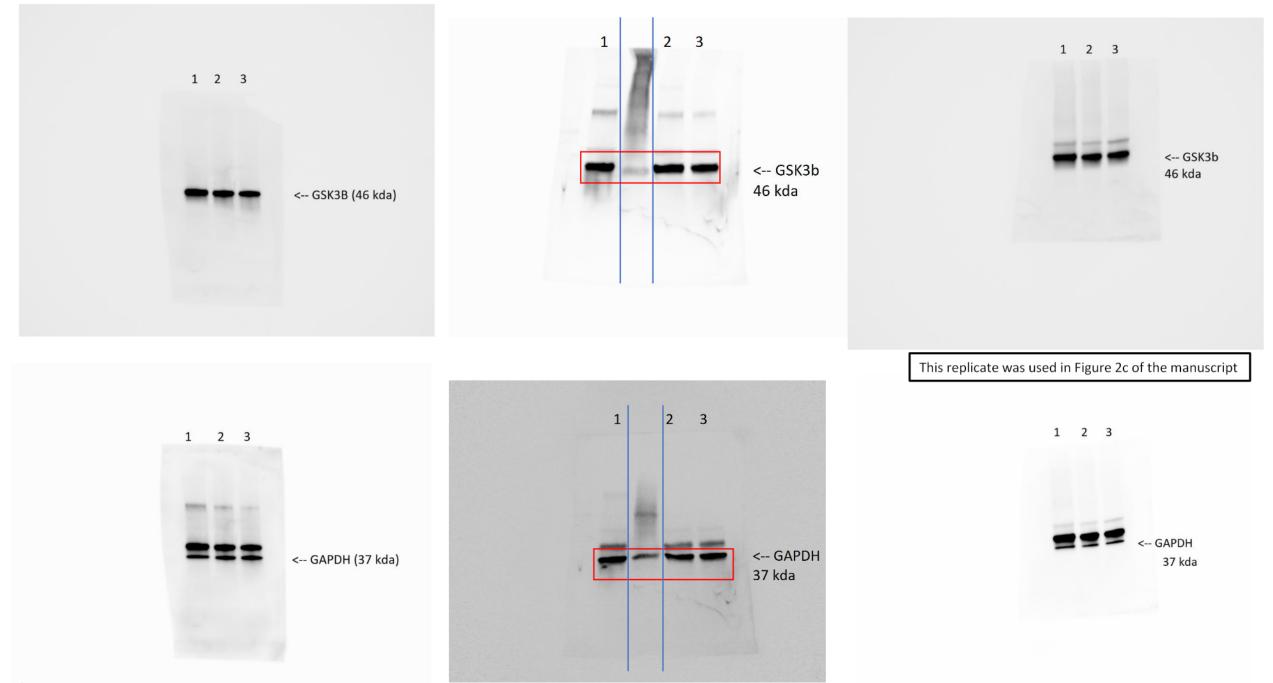
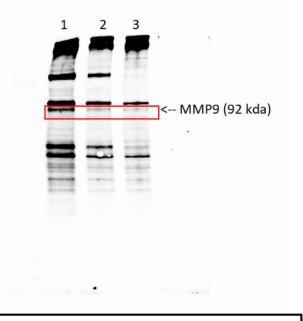
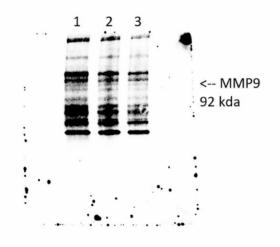


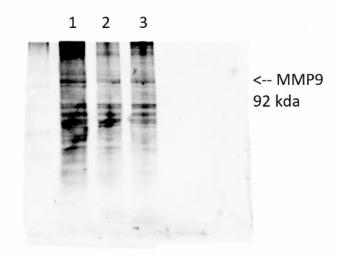
Figure 5

Vertical blue lines indicate how the blot was spliced for final presentation

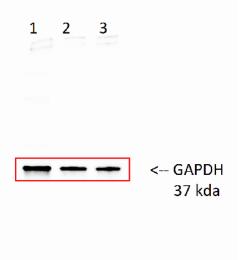
Figure 6

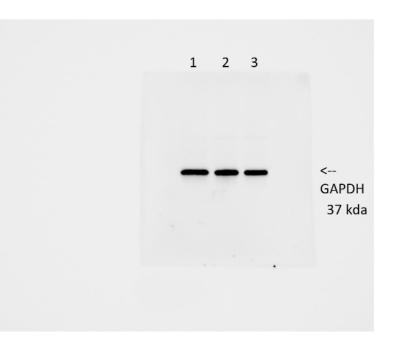






This replicate was used in Figure 4d of the manuscript





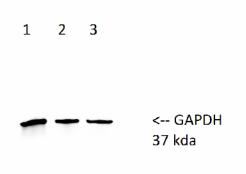
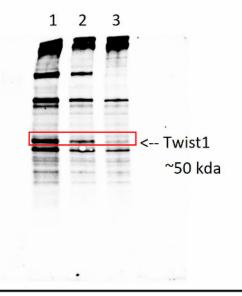
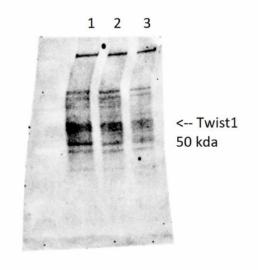


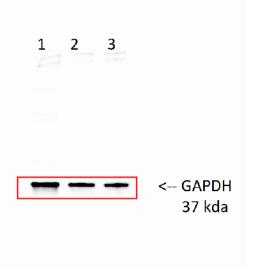
Figure 7

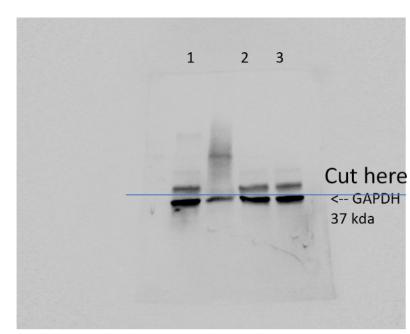


1 2 3 <--- Twist1 ~50 kda



This replicate was used in Figure 4d of the manuscript





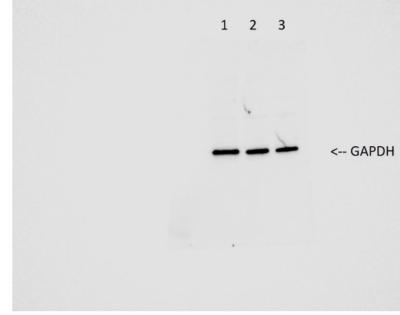
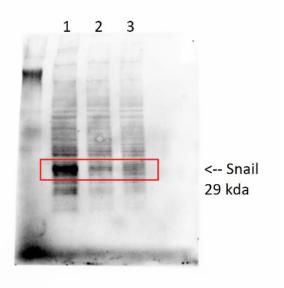
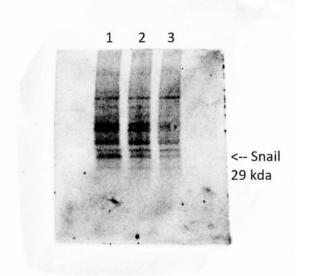
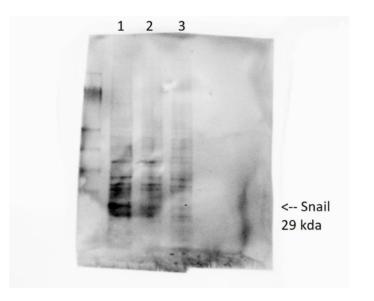


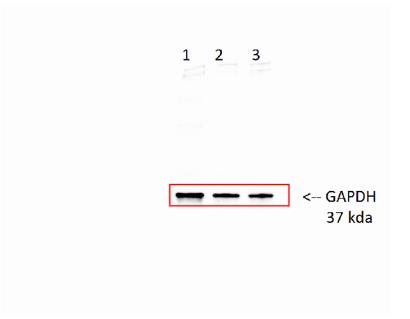
Figure 8



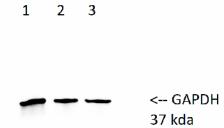




This replicate was used in Figure 4d of the manuscript



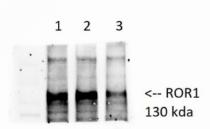


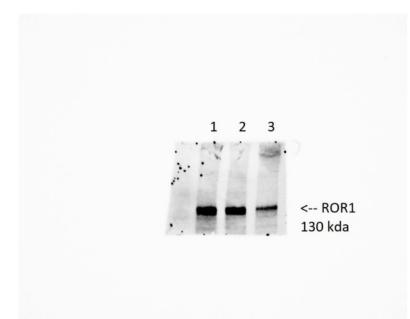


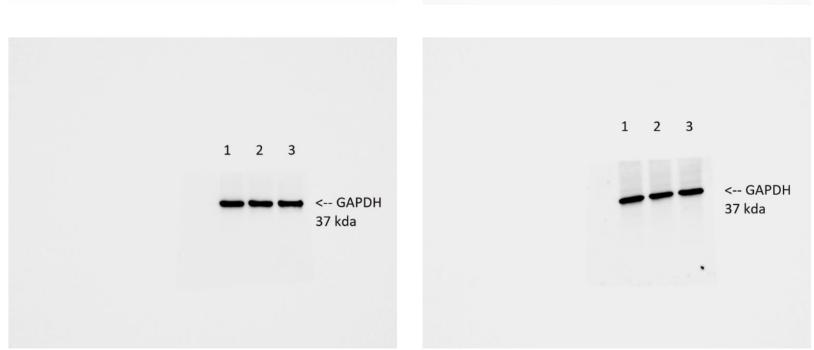
## Raw western blots (Figures 9-13)

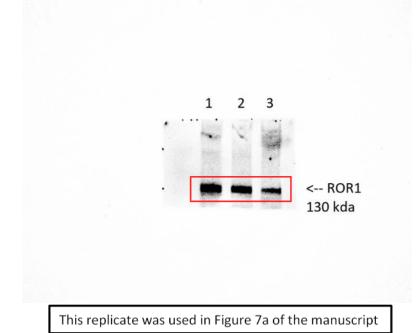
- 1. All lanes used in the manuscript and used for densitometry analysis (N=3) are labeled. Unlabeled lanes were not used for analysis.
- All proteins used in the manuscript and used for analysis are appropriately labeled. Unlabeled proteins were either not used for analysis or were labeled in subsequent slides.
- 3. All protein lysate was obtained from DU145 cells.
- 4. All densitometry analysis was normalized to housekeeping protein (GAPDH).
- 5. Red boxes indicate how the blot was cropped for publication (approximately).
- 6. Labeling:
  - Lanes
    - 1: Control
    - 2: Strc 250 μM- 24h treatment
    - 3: Strc 250 μM- 48h treatment
  - Proteins
    - pGSK3b = phospho-GSK3β (ser9)
    - pAKT = phospho-AKT (ser473)

Figure 9 Membranes were cut approximately at 60-70 kda region before probing/imaging ROR1 and GAPDH.









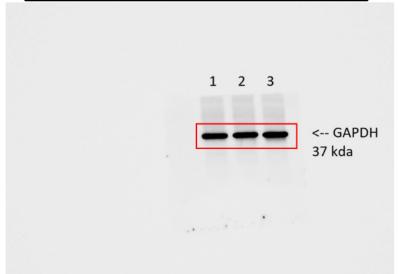


Figure 10

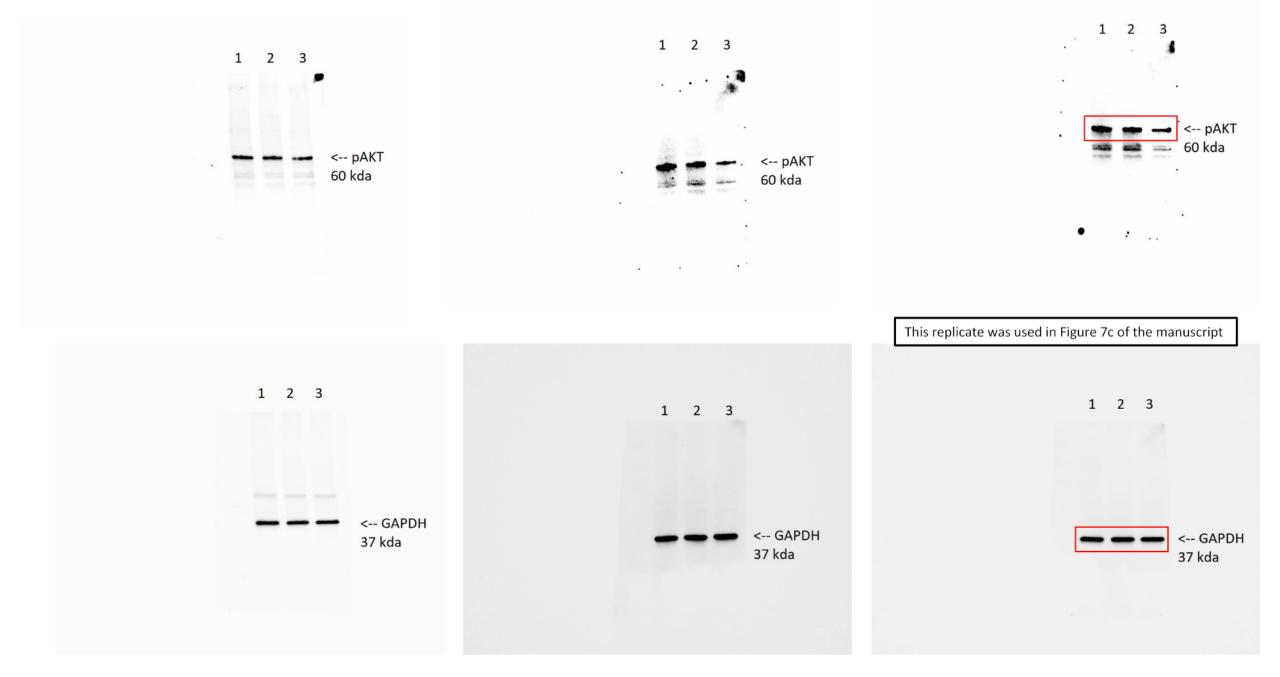


Figure 11

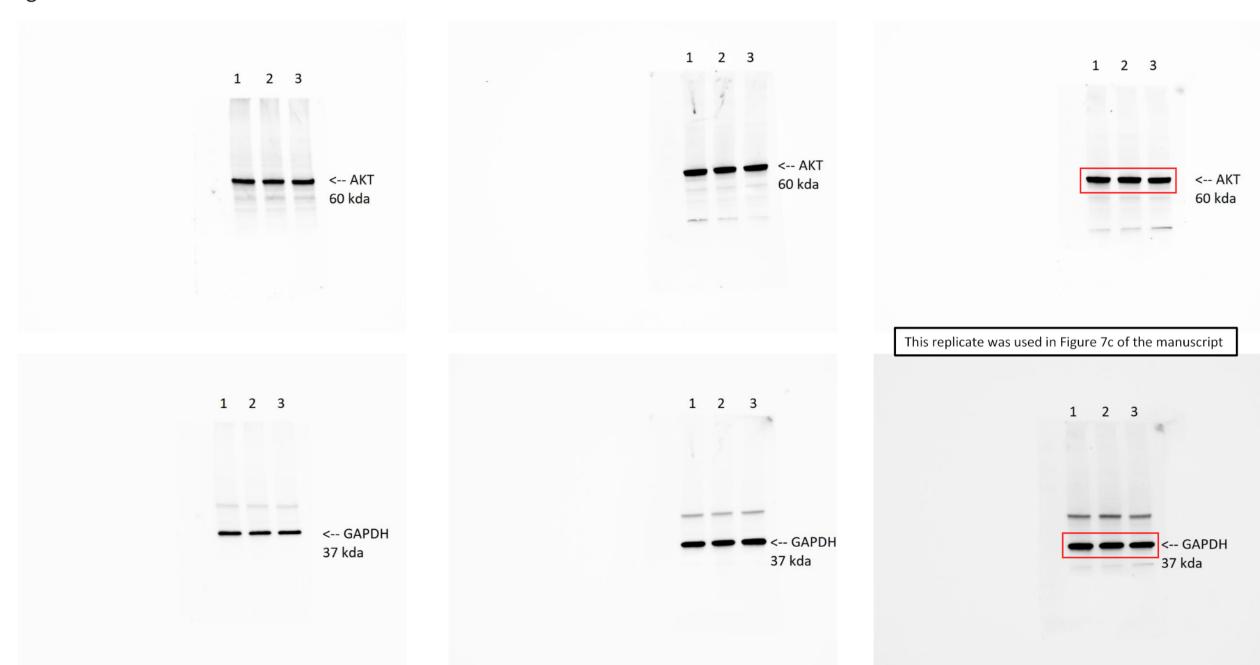
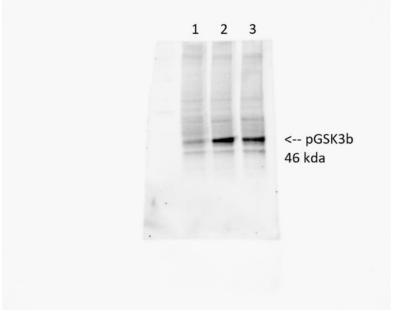
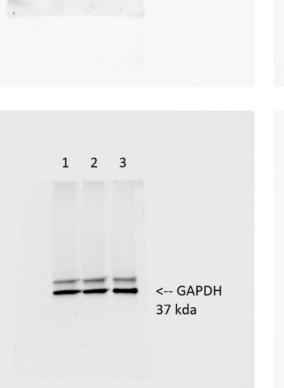
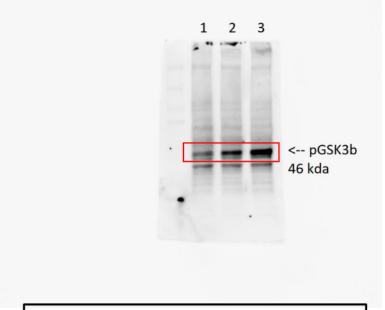
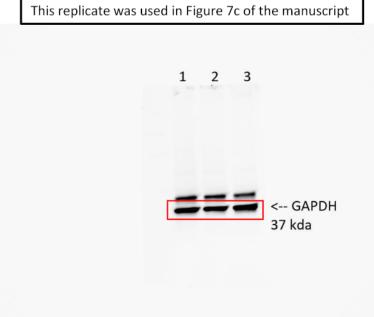


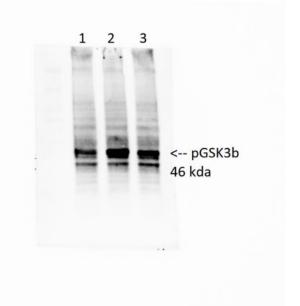
Figure 12











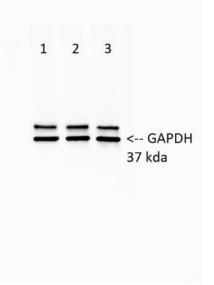


Figure 13

