

A

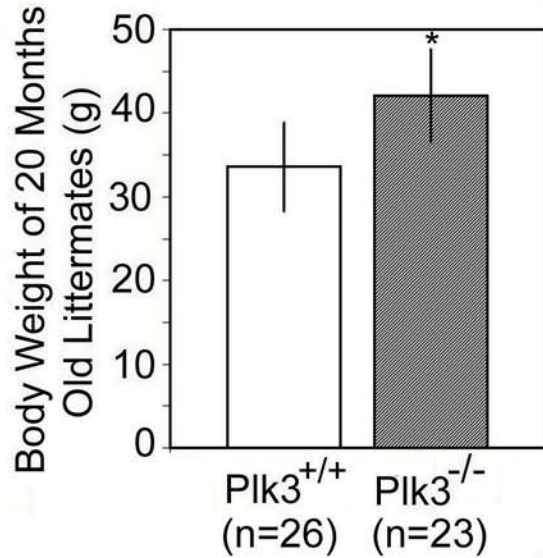


Table 1: Tumor development in *Plk3*^{+/+} vs *Plk3*^{-/-} littermates

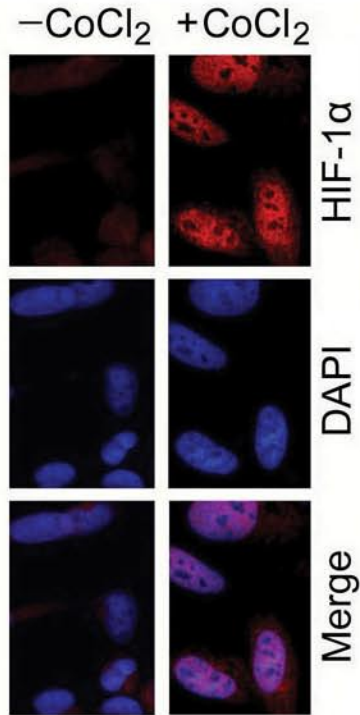
Tumor Type	Lung	Liver	Kidney	Intestine	Uterus	Total # of tumors
<i>Plk3</i> ^{+/+} (n=21)	2 (9.5%)	0	0	0	0	2 (9.5%)
<i>Plk3</i> ^{-/-} (n=26)	8(30.8%)	5 (19.2%)	2 (7.6%)	1 (3.8%)	2 (7.6%)	18 (69.2%)

Table 2: Ectopic expression of *Plk3*-KD suppresses HIF-1 α induction

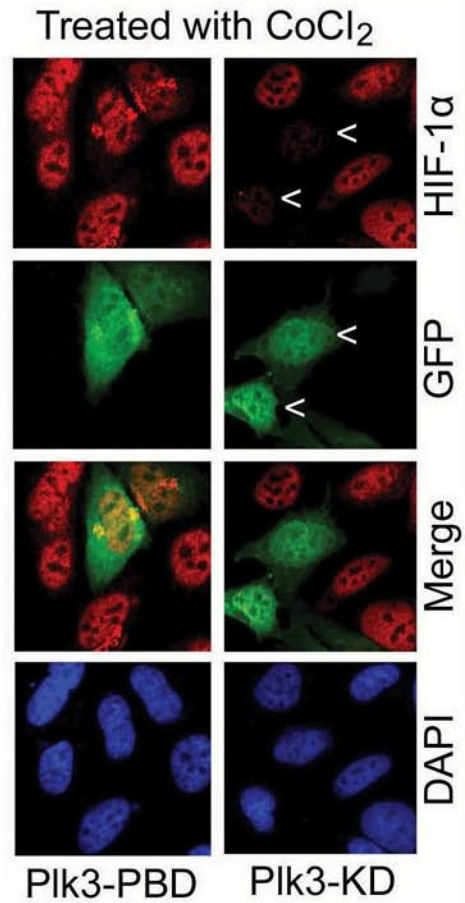
Constructs	GFP	<i>Plk3</i> -FL	<i>Plk3</i> -PBD	<i>Plk3</i> -KD	<i>Plk3</i> -KD ^{T219D}
HIF-1 α Signals	1.0	0.7	1.3	0.5	1.2

Approximately equal amounts of proteins from cells transfected with various expression constructs and treated with nickel ions, as shown in Figure 6C, were blotted for HIF-1 α . Specific HIF-1 α signals were quantitated by densitometry. After normalizing with β -actin signals, HIF-1 α signals are presented (arbitrary unit). The data were summarized from two independent experiments.

A



B



A. HeLa cells treated with CoCl₂ for 4 h were stained with the HIF-1 α antibody (red). DNA was stained with DAPI (blue). Representative cells are shown.
 B. HeLa cells transfected with various expression constructs for 24 h and then treated with CoCl₂ for 4 h were stained with antibodies to HIF-1 α (red) and GFP (green). DNA was stained with DAPI (blue). Representative cells are shown. Arrowheads indicate PIk3-KD-expressing cells with suppressed accumulation of nuclear HIF-1 α after CoCl₂ treatment.

