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## **Ki-67 supports global gene expression programmes driving tumourigenesis 2**

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1 **Ki-67 supports global gene expression programmes driving tumourigenesis**

2

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## 22 **Abstract**

23 Ki-67 is a nuclear protein universally expressed in proliferating vertebrate cells (1), a  
24 characteristic that underlies its widespread use as a cancer biomarker. It organises  
25 heterochromatin (2,3) and the mitotic chromosome periphery (2,4–6), but it is dispensable for  
26 cell proliferation *in vivo*<sup>2</sup>. It thus remains unclear whether Ki-67 confers any advantages to  
27 cancer cells. Here, using mouse models of breast and intestinal cancer, we show that Ki-67  
28 ensures gene expression programmes required for both carcinogenesis and tumour  
29 immunogenicity. We find that germline disruption of Ki-67 protects mice against intestinal  
30 tumourigenesis, while in mammary carcinoma cells, ablation of Ki-67 slows tumour growth  
31 and inhibits metastasis, despite the failure of cells lacking Ki-67 to induce an effective anti-  
32 tumour immune response. Mechanistically, Ki-67 loss leads to widespread deregulation of  
33 gene expression, including genes involved in the epithelial-mesenchymal transition (EMT),  
34 immune responses and drug metabolism. This results in loss of mesenchymal and stem cell  
35 characteristics, downregulation of MHC class I expression and sensitisation to various drug  
36 classes. Our results suggest that, by structuring chromatin to allow global transcriptome  
37 changes, Ki-67 allows cancer cells to adapt to their environment. This is required for all stages  
38 of tumourigenesis and drug resistance, but, by maintaining tumour immunogenicity, it also  
39 confers a targetable vulnerability on cancer cells.

40

## 41 **Results and discussion**

42 Although Ki-67 is universally expressed in dividing cancer cells, cell proliferation is not  
43 affected by Ki-67 gene disruption in any cell type yet analysed *in vitro*, nor during mouse  
44 development<sup>2,5–7</sup>. However, Ki-67 might still be important in carcinogenesis since it  
45 contributes to organisation of chromatin<sup>2</sup>, which controls gene expression and thus cellular  
46 plasticity, a characteristic of cancer cells<sup>8,9</sup>. We first tested the involvement of Ki-67 in tumour  
47 initiation using an *in vitro* model. We transduced our previously generated *Mki67*<sup>+/+</sup> or TALEN-  
48 mutated *Mki67*<sup>-/-</sup> 3T3 fibroblasts<sup>10</sup> with oncogenic mutant H-Ras (G12V), and assessed colony  
49 formation as an indicator of transformation. While H-Ras<sup>G12V</sup>-transduced control 3T3 cells  
50 efficiently formed colonies, *Mki67*<sup>-/-</sup> 3T3 cells did not, indicating that Ki-67 expression is  
51 required for transformation (Extended Data Fig. 1). We next determined whether Ki-67  
52 knockout affects tumour initiation *in vivo*. We used a mouse model of colon carcinogenesis  
53 induced chemically by azoxymethane / dextran sodium sulphate (AOM-DSS) treatment<sup>11</sup>, and  
54 tested the effect of a germline TALEN-disruption of Ki-67 (*Mki67*<sup>2ntΔ/2ntΔ</sup>) that we generated<sup>10</sup>.  
55 As expected, AOM-DSS efficiently induced colon tumours within 16 weeks in wild-type and  
56 *Mki67*<sup>2ntΔ/+</sup> mice; however, no macroscopic lesions were observed in *Mki67*<sup>2ntΔ/2ntΔ</sup> mice

57 (Fig.1a,b). We confirmed that this was not due to an inability to induce inflammation, since  
58 DSS alone triggered similar inflammatory responses in mice of all genotypes (Extended Data  
59 Fig. 2). Furthermore, immune cell proliferation and differentiation in *Mki67*<sup>2ntΔ/2ntΔ</sup> mice  
60 appeared normal in response to haemolytic anaemia triggered by phenylhydrazine treatment  
61 (Extended Data Fig. 3). This suggests that Ki-67 is specifically required for initiation of  
62 tumourigenesis. To confirm this, we used a genetic model of intestinal tumourigenesis. We  
63 crossed *Mki67*<sup>2ntΔ/2ntΔ</sup> mice with *Apc*<sup>Δ14/+</sup> mice, which rapidly develop tumours in the intestine  
64 due to loss of the second allele of the *Apc* tumour suppressor gene<sup>12</sup>. While *Apc*<sup>Δ14/+</sup>  
65 *Mki67*<sup>2ntΔ/+</sup> mice formed multiple intestinal tumours as expected, tumour burden was strongly  
66 reduced in *Apc*<sup>Δ14/+</sup> *Mki67*<sup>2ntΔ/2ntΔ</sup> mice (Fig. 1c,d). These results indicate that Ki-67 is required  
67 for efficient initiation of tumourigenesis induced either by an oncogene, chemical mutagenesis  
68 or loss of a tumour suppressor.

69 We next asked whether Ki-67 is required for tumour growth and metastasis from  
70 already transformed cells. As a first approach, we knocked down Ki-67 by stable shRNA in a  
71 commonly used aggressive human cervical cancer cell line, HeLa S3, and grafted the  
72 resulting cells and an shRNA control line into opposing flanks of athymic nude mice. Ki-67  
73 knockdown was maintained *in vivo* (Extended Data Fig. 4a,b) and, as expected, this did not  
74 affect cell proliferation (as shown by unchanged PCNA and mitotic indices; Extended data  
75 Fig. 4c,d). However, tumour growth was severely slowed (Fig. 2a) and histopathological  
76 analysis showed vast areas of necrosis and apoptosis accompanied by disorganised vascular  
77 and lymphatic networks, indicating disrupted angiogenesis (Fig. 2b,c). To ensure that  
78 phenotypes of Ki-67 downregulation were not specific to the tumour type or knockdown  
79 approach, we generated a biallelic *MKI67* gene knockout (KO) by CRISPR-Cas9 in human  
80 MDA-MB-231 triple negative breast cancer cells (Extended Data Fig. 4e,f). As expected,  
81 *MKI67*<sup>-/-</sup> cells proliferated normally *in vitro* (Extended Data Fig. 4g,h) but tumours from  
82 xenografts in nude mice grew much slower than controls, again showing normal cell  
83 proliferation but defective angiogenesis (Fig. 2d-f). Thus, in xenograft experiments, Ki-67  
84 sustains tumour growth but not by controlling cell proliferation.

85 To determine effects of Ki-67 loss in a mouse model that more closely mimics human  
86 breast cancer, we used 4T1 mammary carcinoma cells. This cell line is derived from BALB/c  
87 mice, is highly tumourigenic, spontaneously metastasises to distant organs<sup>13,14</sup>, and, when  
88 grafted in a syngeneic background, a biphasic tumour progression profile is observed due to  
89 an induced immune response<sup>15</sup>. As expected, CRISPR-Cas9-mediated *Mki67* knockout did  
90 not affect 4T1 cell proliferation rates (Extended Data Fig. 5a-d). We engrafted WT and Ki-67  
91 knockout 4T1 cells orthotopically into mammary fat pads in both immune-proficient syngeneic



92 BALB/c mice and immunodeficient (athymic-nude and NOD/SCID) mice, the comparison of  
93 which should allow identification of the effects of the immune system. In immunodeficient  
94 mice, tumours from *Mki67*<sup>-/-</sup> 4T1 cells grew significantly slower than from Ki-67-expressing  
95 4T1 cells (Fig. 3a-c). In immune-proficient BALB/c mice, control 4T1 tumours established  
96 quickly and, as expected, initially regressed before regrowing (Fig. 3d). However, in these  
97 mice, *Mki67*<sup>-/-</sup> 4T1 cells formed primary tumours as efficiently as controls, but no initial  
98 regression occurred, overriding the slower growth of knockout tumours and resulting in higher  
99 final tumour burden than controls (Fig. 3d). This suggested a defective immune response in  
100 mice bearing Ki-67 knockout tumours. We analysed this by immunophenotyping of spleens  
101 and lymph nodes. The fraction of dendritic cells, B-cells and T-cells (except for T-reg) was  
102 reduced in *Mki67*<sup>-/-</sup> 4T1 tumour-bearing mice, as a result of a dramatic increase in cells  
103 positive for CD11b (a common myeloid marker, expressed by monocytes, neutrophils, NK  
104 cells, granulocytes, macrophages and myeloid-derived suppressor cells (MDSC)) (Extended  
105 Data Fig. 6). In agreement, immunohistological analysis revealed increased infiltration of  
106 MDSC in knockout tumours (using the MDSC marker, GR1; Fig. 3e), possibly indicating an  
107 immunosuppressive environment that protects the tumours against immune-mediated  
108 cytotoxicity.

109 In immune-proficient mice, despite the impaired immune-response to *Mki67*<sup>-/-</sup> 4T1 cells  
110 and a consequently higher tumour burden, there were noticeably less metastases than from  
111 control 4T1 cells, although this failed to reach statistical significance ( $p = 0.13$ ) (Fig 3f).  
112 However, the reduction in visible metastasis from *Mki67*<sup>-/-</sup> 4T1 cells was highly significant in  
113 immunodeficient mice (Fig. 3g) or in immune-proficient mice injected in the tail vein to  
114 circumvent the immune-targeting of tumour cells at the primary-site (Fig. 3h). We quantified  
115 micrometastases in nude mice by dissociating lung tissue after 2 weeks and growing cells in  
116 the presence of 6-thioguanine, to which 4T1 cells are resistant<sup>14</sup>. The number of metastatic  
117 cells from Ki-67 knockouts was reduced nearly 100-fold (Extended Data Fig. 7). These results  
118 highlight a trade-off for cancer cells, whereby Ki-67 expression sustains tumour growth and  
119 promotes metastasis, yet also confers efficient targeting by the immune system.

120 Since we previously found that Ki-67 knockout in mouse NIH3T3 cells or knockdown  
121 in human cancer cells does not alter cell proliferation nor induce cell death, but disrupts  
122 chromatin organisation<sup>2</sup> and alters gene expression, we hypothesised that the latter might  
123 explain the requirements for Ki-67 in tumourigenesis. To explore this idea we first compared  
124 transcriptomes of WT and *Mki67*<sup>-/-</sup> NIH3T3 cells by RNA-seq. Consistent with roles for Ki-67  
125 in maintaining global chromatin organisation, this revealed widespread changes in gene  
126 expression, with 2558 genes significantly deregulated in independent clones of *Mki67*<sup>-/-</sup> cells

127 (q < 0.05) (Extended Data Fig. 8a; Supplementary Table 1). Next, we investigated how  
128 transcriptomes are remodelled in 4T1 cells lacking Ki-67. In these cancer cells, *Mki67*  
129 knockout caused genome-scale gene expression alterations: 4979 genes were deregulated,  
130 of which 1239 and 585 genes were >2-fold down-regulated and up-regulated, respectively  
131 (Fig. 4a; Extended Data Fig. 8b; Supplementary Table 2). There was little overlap in the  
132 deregulated genes between *Mki67*<sup>-/-</sup> (epithelial) and NIH3T3 (mesenchymal) cells (Extended  
133 Data Fig. 8b; Supplementary Tables 1,2) suggesting that Ki-67 ensures gene expression  
134 programmes in different cell types by maintaining chromatin organisation rather than directly  
135 regulating specific genes. RNA-seq of early stage tumours from WT and Ki-67 mutant 4T1  
136 cells grafted into nude mice showed that Ki-67-dependent transcriptome changes were  
137 preserved *in vivo* (Fig. 4b).

138 We performed pathway analysis of genes highly deregulated in Ki-67 knockout 4T1  
139 cells to better understand the links between Ki-67-mediated gene expression control and  
140 tumourigenesis. We found upregulation of Notch targets and downregulation of genes  
141 involved in the inflammatory response, Wnt pathway and the EMT (Fig. 4c) which we  
142 validated by qRT-PCR (Extended Data Fig. 8c). Importantly, although the Notch pathway is  
143 oncogenic in T cell acute lymphoid leukaemia, it can act as a tumour suppressor in specific  
144 cellular contexts<sup>16</sup>, can block Wnt signaling<sup>17,18</sup> (a driver of tumourigenesis, cell stemness and  
145 the EMT) and induce drug resistance<sup>19</sup>. We confirmed Notch pathway upregulation at the  
146 protein level (Fig. 4d). We further verified that Ki-67 knockout 4T1 cells had lost expression  
147 of the mesenchymal marker vimentin but upregulated E-cadherin, and had a more epithelial  
148 morphology (Fig. 4e). Downregulated genes were enriched in targets of nuclear factor  
149 erythroid 2-related factor 2 (NFE2L2), which upregulates genes in response to oxidative  
150 stress; interferon regulatory factor 8 (IRF8); polycomb-repression complex 2 (PRC2), which  
151 mediates H3K27me3 and is a well-characterised regulator of the EMT<sup>20,21</sup>; and the  
152 pluripotency factors Nanog and Sox2 (Extended Data Fig. 8d). We focused on PRC2 as we  
153 previously found<sup>10</sup> that in human osteosarcoma U2OS cells, Ki-67 binds to the essential  
154 PRC2 component SUZ12. To test whether the repression of the EMT in Ki-67 knockout  
155 depends on PRC2, we disrupted *Suz12* or *Ezh2* in WT and *Mki67*<sup>-/-</sup> 4T1 cells using CRISPR-  
156 Cas9. This partially rescued expression of vimentin (Extended Data Fig. 8e,f). However, it did  
157 not restore tumour growth rates nor metastasis *in vivo* from *Mki67*<sup>-/-</sup> 4T1 cells, nor affect  
158 tumourigenicity of control cells, in nude mice (Extended Data Fig. 8g). Thus, the PRC2  
159 complex activity is restrained by Ki-67, but does not in itself have either oncogenic or tumour  
160 suppressive activity in this system. As such, the reported PRC2 roles in cancer<sup>22,23</sup> might not

161 apply generally, and a single chromatin regulation pathway cannot account for Ki-67 roles in  
162 supporting the EMT and tumourigenesis.

163 Like Ki-67, the EMT is required for metastasis in some mouse cancer models<sup>24,25</sup>. The  
164 EMT is also closely associated with a stem-like state<sup>26–28</sup>, as well as resistance to  
165 chemotherapeutic drugs<sup>29</sup> and induction of immune responses<sup>30</sup>. We therefore next  
166 investigated the importance of Ki-67 for these three EMT-linked characteristics.

167 First, we analysed two indicators of tumour cell stemness: the ability to form spheroids  
168 in the absence of adhesion to a surface<sup>26,31</sup>, and aldehyde dehydrogenase activity<sup>32,33</sup>. Both  
169 parameters were strongly reduced in Ki-67 knockout 4T1 cells (Fig. 4f,g).

170 Second, we noticed that 26 genes involved in drug metabolism were downregulated in  
171 Ki-67 knockout 4T1 cells, while only one was upregulated, suggesting that Ki-67 expression  
172 might determine sensitivity to chemotherapeutic drugs (Extended Data Fig. 9a,b). To test this,  
173 we performed an automated gene-drug screen using the Prestwick chemical library,  
174 composed of 1283 FDA-approved small molecules, as well as salinomycin, a positive control  
175 found to target cancer stem cells<sup>34</sup>, and 6-thioguanine, which was originally used to isolate  
176 4T1 cells<sup>13</sup>. Control 4T1 cells were sensitive to 102 drugs at 10 $\mu$ M concentration, while the  
177 two *Mki67*<sup>-/-</sup> clones were sensitive to 99 and 98 respectively, with 82 hits common to the three  
178 cell lines (Extended Data Fig. 9c; Supplementary Data). This suggests that Ki-67 loss does  
179 not qualitatively alter the drug-sensitivity profiles. We next determined the IC<sub>50</sub> of 10 common  
180 hits used in cancer therapy. Importantly, *Mki67*<sup>-/-</sup> cells were markedly more sensitive to all the  
181 molecules tested (Fig. 4h; Extended Data Fig. 9d).

182 Third, genes directing mouse major histocompatibility complex (MHC) class I-mediated  
183 antigen presentation *Tap2*, *Psmb8* and *Psmb9* were downregulated in *Mki67*<sup>-/-</sup> 4T1 cells  
184 (Extended Data Fig. 8c; Supplementary Table 2). Flow cytometry confirmed lower expression  
185 of the mouse MHC class I molecules H2D and H2K (Fig. 4i), providing an explanation for how  
186 Ki-67 knockout affects tumour recognition by the immune system. We obtained similar results  
187 in human *MKI67*<sup>-/-</sup> MDA-MB-231 cells, implying that roles of Ki-67 in maintaining MHC  
188 expression are conserved across species (Extended Data Fig. 10).

189 The above results show that Ki-67 is not required for cancer cell proliferation *in vivo*,  
190 but supports coherent transcriptional programmes that are involved in tumour cell plasticity,  
191 drug resistance and induction of immune responses. Together, these likely account for the  
192 roles of Ki-67 in tumourigenesis revealed in this study (Fig. 4j). Clues to its biochemical  
193 mechanism of action are suggested by the fact that Ki-67 is an intrinsically disordered protein  
194 (IDP) and behaves as a surfactant for mitotic chromatin<sup>5</sup>. Since we previously found that  
195 heterochromatin is more dispersed in cells lacking Ki-67 (ref. 2), we propose that these

196 properties may also allow Ki-67 to regulate interphase chromatin. Due to their conformational  
197 flexibility, IDPs can act as hubs to bind many protein partners<sup>35</sup>, and we previously identified  
198 many chromatin regulators interacting with Ki-67 (ref. 2). However, disrupting one such  
199 interactor, the PRC2 complex, does not restore tumourigenic capacity to Ki-67 mutant cancer  
200 cells, consistent with the idea that Ki-67 may act more globally.

201 While it has long been thought that global changes in nuclear structure observed in  
202 cancer cells might contribute functionally to carcinogenesis<sup>36</sup>, mechanistic insight into this  
203 question has been lacking. We speculate that the alterations of nuclear organisation seen in  
204 cancer cells may reflect their increased cellular plasticity. By binding multiple chromatin  
205 regulators and organising the transcriptome at a genome scale, Ki-67 might contribute to both  
206 nuclear structuration and cellular plasticity, thereby conferring on cancer cells the ability to  
207 adapt to their environment.

208

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299

300 **Competing interests**

301 The authors declare that they have no competing interests.

302

303 **Author Contributions**

304 D.F. conceived and supervised the project. K.M., L.K., V.Z., M.V. and D.F. designed  
305 experiments and interpreted the data. K.M., M.S., P.S., E.A., A.A., S.P., C.V., N.P., F.B., B.B.,  
306 M.P., C.H-K. and V.D. performed experiments. G.D. performed data analysis. K.M., L.K. and  
307 D.F. wrote the manuscript.

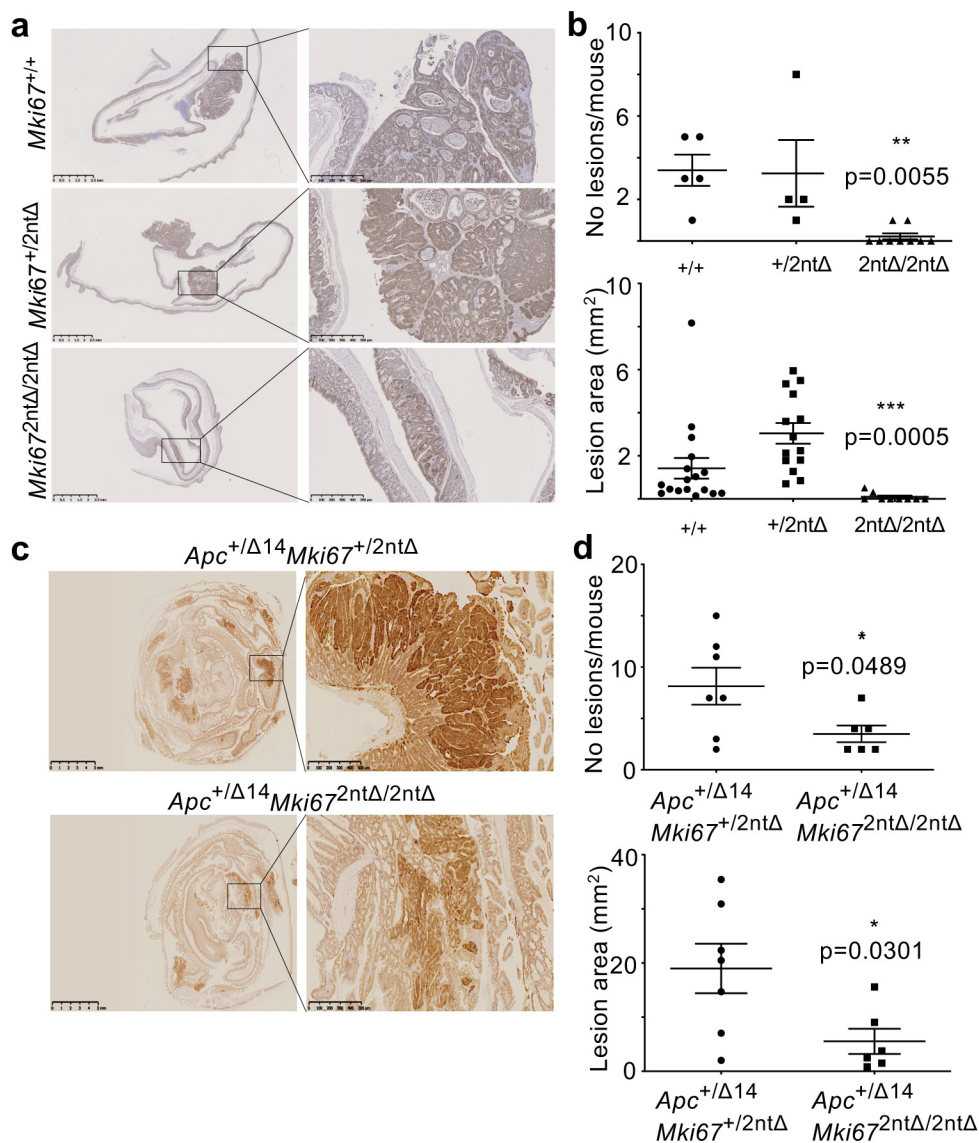
308

309 **Data availability**

310 All RNA-seq raw data will be deposited in GEO and will be publicly available prior to  
311 publication of the manuscript.

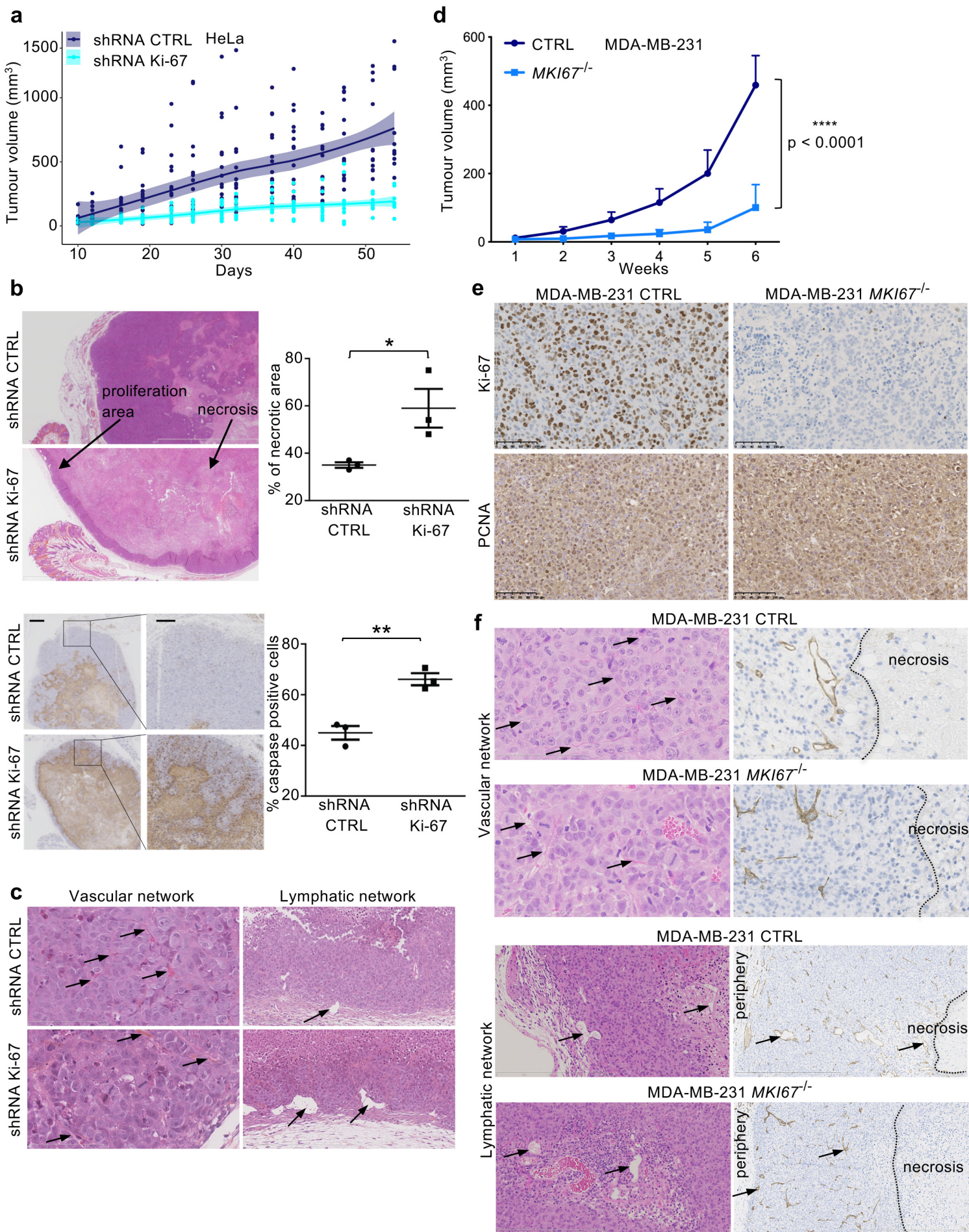
312



**Fig.1****Figure 1: Germline disruption of Ki-67 protects mice against intestinal tumourigenesis.**

**a.** IHC analysis of  $\beta$ -catenin expression in whole intestines recovered from each mouse group (*Mki67*<sup>+/+</sup>, *Mki67*<sup>+/2ntΔ</sup> and *Mki67*<sup>2ntΔ/2ntΔ</sup>) at week-16 post AOM-DSS treatment. Scale bar, 2.5mm. Insets show accumulation of  $\beta$ -catenin in nuclei. Scale bar, 500 $\mu$ m. **b.** Quantification of the number (top) and area (bottom) of neoplastic lesions. Error bar, SEM (n= 5 *Mki67*<sup>+/+</sup> mice; n= 4 *Mki67*<sup>+/2ntΔ</sup> mice; n= 9 *Mki67*<sup>2ntΔ/2ntΔ</sup> mice). **c.** IHC staining of  $\beta$ -catenin in whole intestines from 6-7-month old *Apc*<sup>+/Δ14</sup>*Mki67*<sup>+/2ntΔ</sup> and *Apc*<sup>+/Δ14</sup>*Mki67*<sup>2ntΔ/2ntΔ</sup> mice. Scale bar, 5mm. Insets show accumulation of  $\beta$ -catenin in nuclei. Scale bar, 500 $\mu$ m. **d.** Quantification of the number (top) and total area (bottom) of neoplastic lesions. Error bars, SEM (n= 7 *Apc*<sup>+/Δ14</sup>*Mki67*<sup>+/2ntΔ</sup> mice; n= 6 *Apc*<sup>+/Δ14</sup>*Mki67*<sup>2ntΔ/2ntΔ</sup> mice).

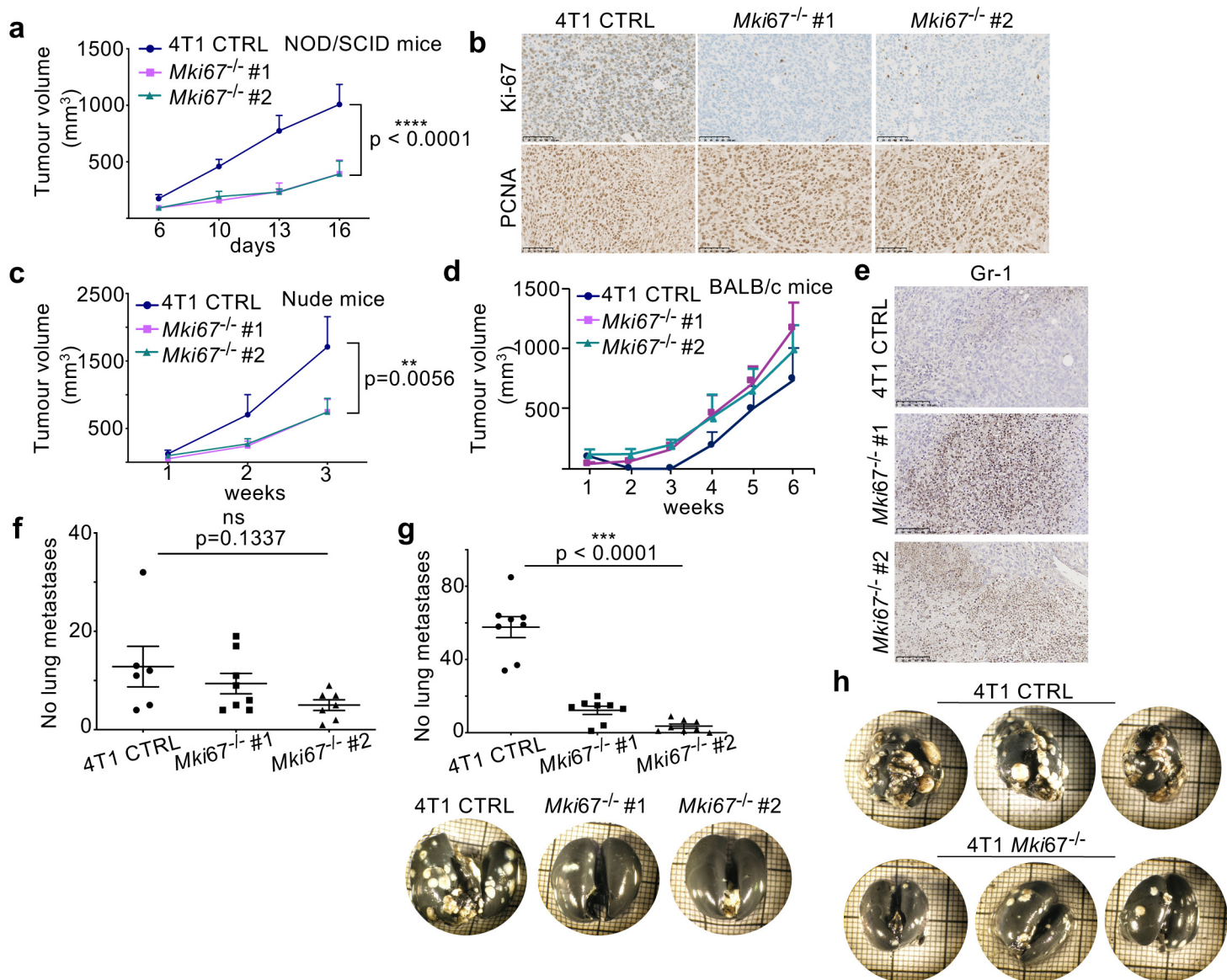


**Fig.2**

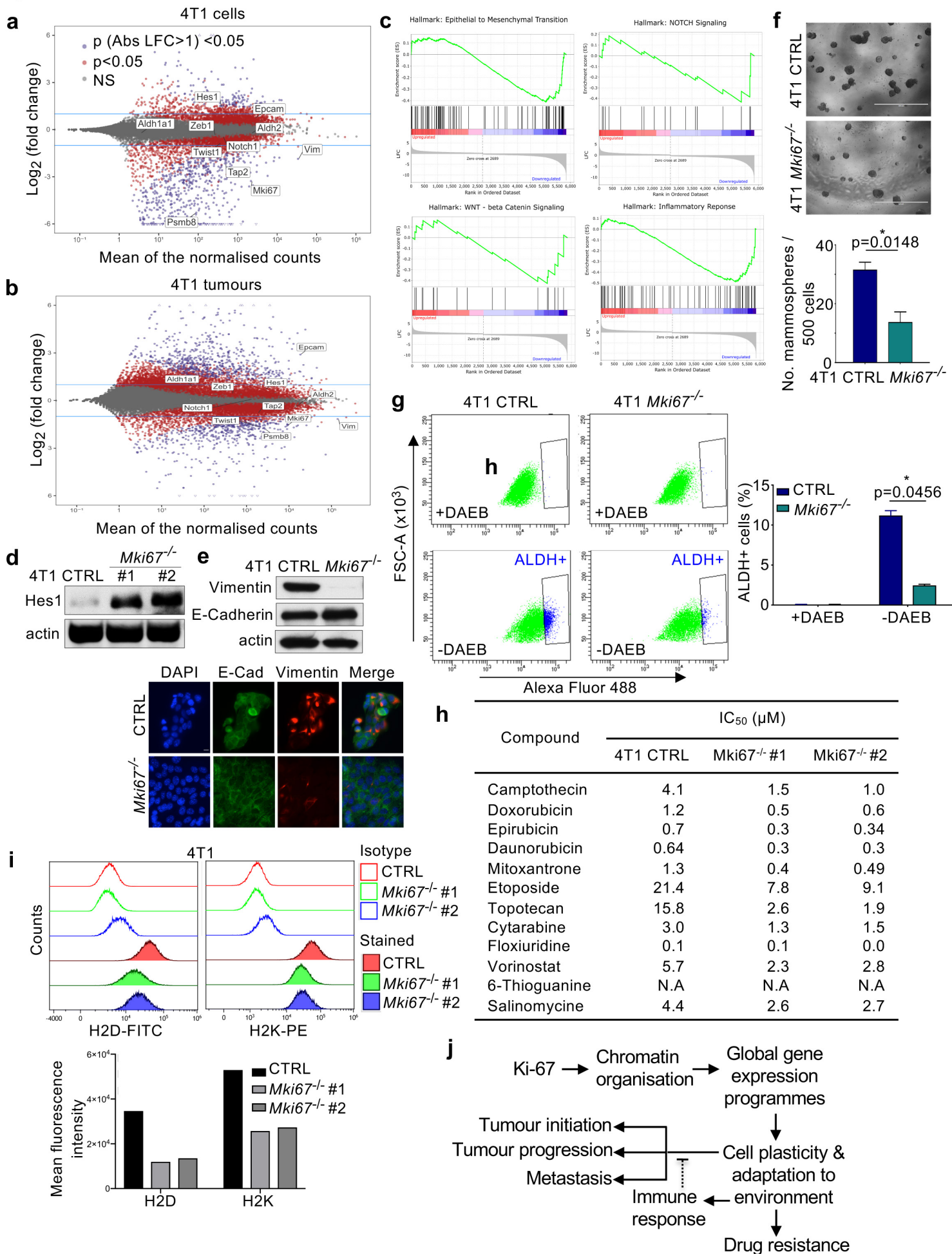
**Figure 2: Ki-67 sustains tumour growth independently of cell proliferation.**

**a-c.** HeLa shRNA CTRL or shRNA Ki-67 were injected subcutaneously into 6-8-week old nude mice. **a.** Tumour burden (n=19 mice). **b.** (Top left) Hematoxylin and eosin (H&E)-stained tumour sections highlighting proliferation and necrotic areas. Scale bar, 3mm. (Top right) Percentage of necrotic area for each of 3 separate tumours from different mice per cell line, mean  $\pm$  SEM. (Bottom left) Immunohistochemistry (IHC) staining for cleaved caspase 3. Scale bar, 1mm (250 $\mu$ m for insets). (Bottom right) Percentage of cleaved caspase 3-positive cells for tumours from different mice, mean  $\pm$  SEM. **c.** H&E-stained tumour sections showing vascular (arrows; scale bar, 100 $\mu$ m) and lymphatic networks (arrows; scale bar, 400 $\mu$ m). While in CTRL tumours fibrovascular network is thin and regular, in shRNA Ki-67 tumours, the network is disorganised, thick and fibrosed. The lymph vessels in shRNA Ki-67 tumours are dilated, characteristic for oedema and indicative of slow fluid flow. **d-f.** Xenografts of MDA-MB-231 CTRL or Ki-67 KO cells in mammary fat pads of nude mice. **d.** Tumour burden. Error bars, SEM (n=8 mice). **e.** IHC images of tumours stained for Ki-67 and PCNA. Scale bar, 100 $\mu$ m. **f.** IHC analysis of vascular and lymphatic networks (arrows). Left, H&E; scale bar, 400 $\mu$ m. Right, CD31 staining; scale bar, 800 $\mu$ m. Areas of necrosis are indicated. The defects in Ki-67 KO tumours are similar as those observed in shRNA Ki-67 tumours.



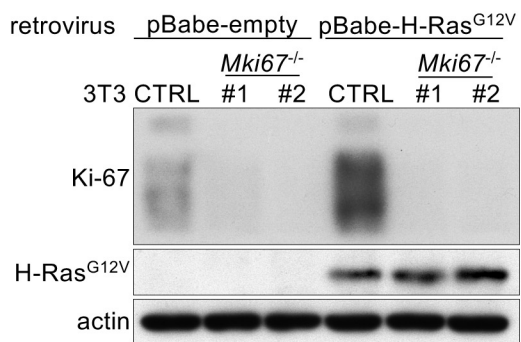
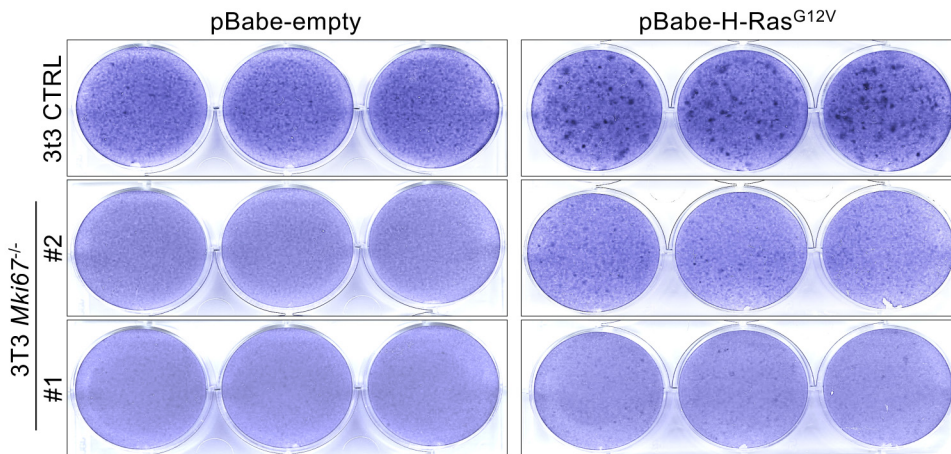
**Fig. 3****Figure 3: Metastasis and anti-tumour immune responses require Ki-67.**

**a.** 4T1 CTRL or *Mki67*<sup>-/-</sup> orthotopic xenografts in NOD/SCID mice. Tumour growth was monitored for 3 weeks (left). Error bars, SEM (n=6 mice). **b.** IHC staining for Ki-67 and PCNA in 4T1 CTRL or *Mki67*<sup>-/-</sup> tumours. Scale bar, 100µm. **c.** 4T1 CTRL or *Mki67*<sup>-/-</sup> orthotopic xenografts in athymic nude mice. Error bars, SEM (n=8 mice). **d.** Tumour growth of 4T1 CTRL or *Mki67*<sup>-/-</sup> orthotopic xenografts in immunocompetent BALB/c mice. Error bars, SEM (n=8 mice). **e.** IHC analysis of 4T1 CTRL or *Mki67*<sup>-/-</sup> tumours (week-4 post-transplantation) stained for Gr-1, a myeloid-derived suppressor cell (MDSC) marker. Scale bar, 100µm. **f.** Quantification of lung metastases in nude mice in **c**. Error bars, SEM. Bottom, representative images of lungs stained to visualise metastases (white nodules); scale bar, 10mm. **g.** Quantification of lung metastases in BALB/c mice in **d**. Error bars, SEM. **h.** 4T1 CTRL or *Mki67*<sup>-/-</sup> cells were injected via tail vein into immune-competent BALB/c mice. Representative images of stained lungs (day 21 post-injection), metastases are white. Scale bar, 10mm.

**Fig. 4**

**Figure 4: Ki-67 ablation deregulates gene expression programmes involved in cancer cell stemness, drug resistance and immunogenicity.**

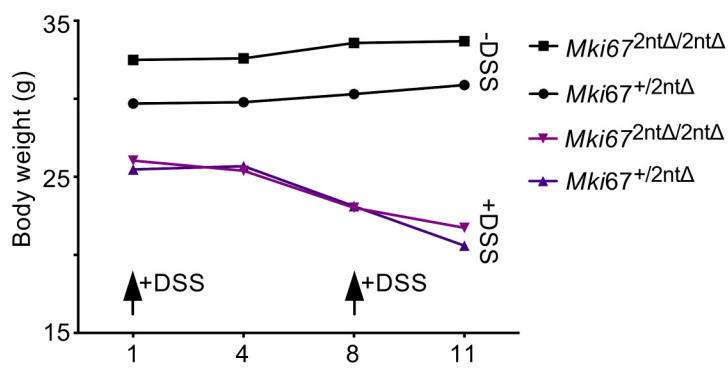
**a,b.** Dot plot analysis of differentially expressed genes (DEGs) in 4T1 *Mki67<sup>-/-</sup>* cells (**a**) and 4T1 *Mki67<sup>-/-</sup>*-derived tumours (**b**). Red dots: DEGs with adjusted p-value < 0.05; purple dots: genes whose log<sub>2</sub> fold change, LFC >1 or <-1, is statistically significant (adjusted p-value < 0.05); grey dots: not significant, NS. **c.** Gene Set Enrichment Analysis (GSEA) of highly down-regulated genes in 4T1 *Mki67<sup>-/-</sup>* cells. **d.** Immunoblot analysis of Hes1 expression in 4T1 CTRL and *Mki67<sup>-/-</sup>* cells. **e.** Immunoblot (top) and immunofluorescence (bottom; scale bar, 10µm) analysis of vimentin and E-cadherin expression in 4T1 CTRL and *Mki67<sup>-/-</sup>* cells. **f.** Mammosphere formation assay of 4T1 CTRL or *Mki67<sup>-/-</sup>* cells. Representative images (top; scale bar, 400 µm) and quantification (bottom; error bars, SEM, n=10) after 7 days. **g.** Aldehyde dehydrogenase 1 activity measured using a flow-cytometry assay in 4T1 CTRL and *Mki67<sup>-/-</sup>* cells. DAEB, inhibitor of ALDH was used as a negative control. Left, representative FACS images. Right, quantification of ALDH+ cells. Error bars, SEM (n=2 independent analyses). **h.** IC<sub>50</sub> of 4T1 CTRL or *Mki67<sup>-/-</sup>* cells treated with indicated compounds. **i.** Flow-cytometry analysis of MHC class I expression. 4T1 CTRL and *Mki67<sup>-/-</sup>* cells were stained with anti-H2D<sup>d</sup>-FITC and anti-H2K<sup>d</sup>-PE, or control isotypes. Top, representative images. Bottom, quantification (n=1 of each clone). **j.** Model of Ki-67 mechanism in tumour progression and cancer cell dissemination (see text).

**a****b****Extended Data Fig. 1: Ki-67 is required for cell transformation.**

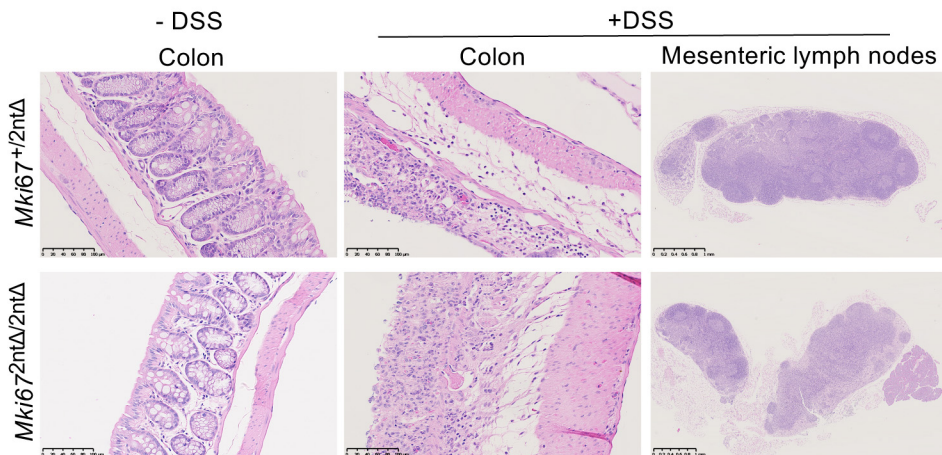
CTRL or TALEN-mutated *Mki67*<sup>-/-</sup> 3T3 fibroblasts were transduced with either empty or H-Ras<sup>G12V</sup>-expressing retroviruses. **a.** Immunoblot analysis of the expression of the indicated proteins. Actin serves as loading control. **b.** Crystal violet staining of the colonies formed.



a



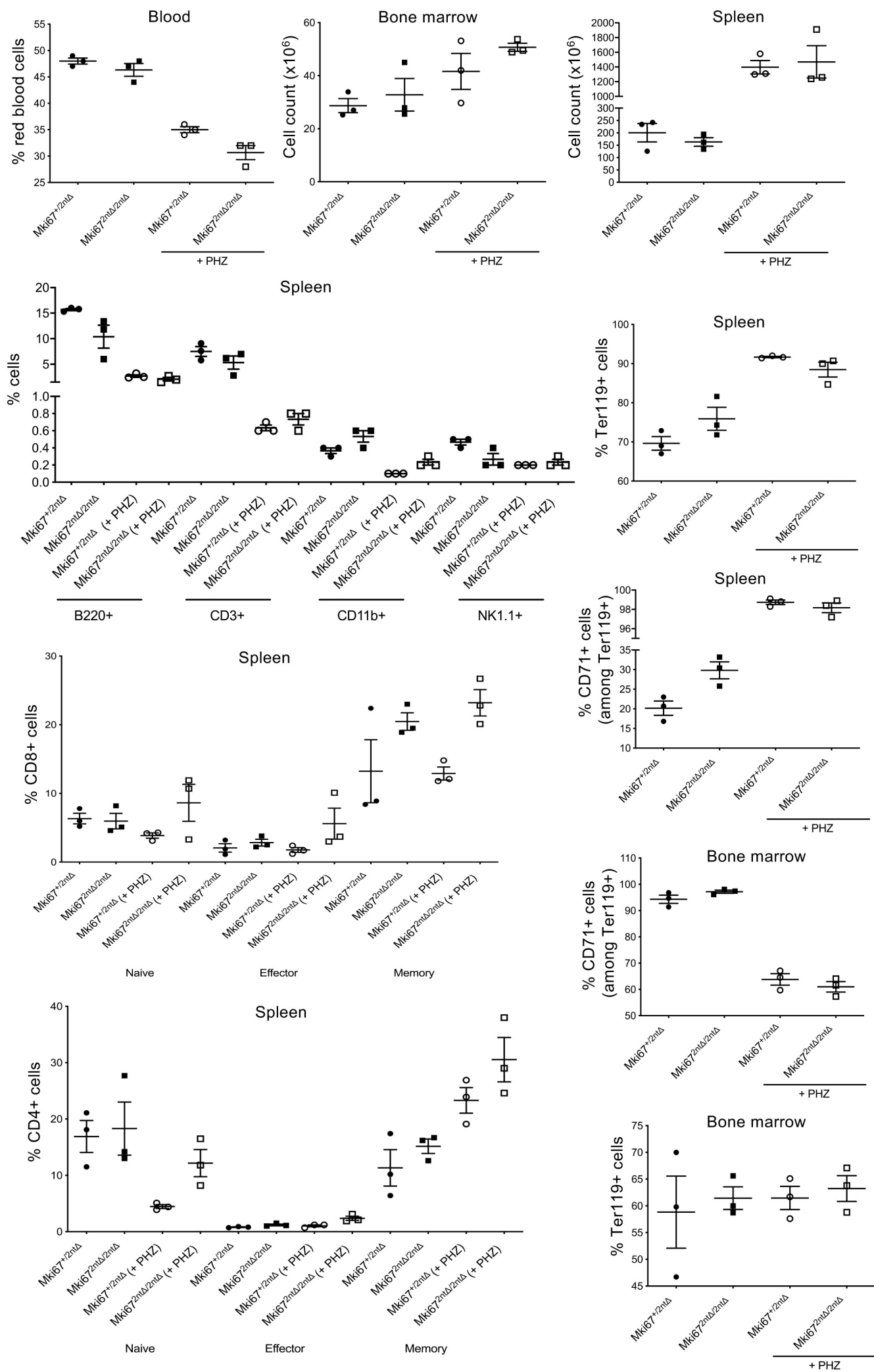
b



**Extended Data Fig. 2: *Mki67*<sup>mt/mt</sup> mice have normal inflammatory response.**

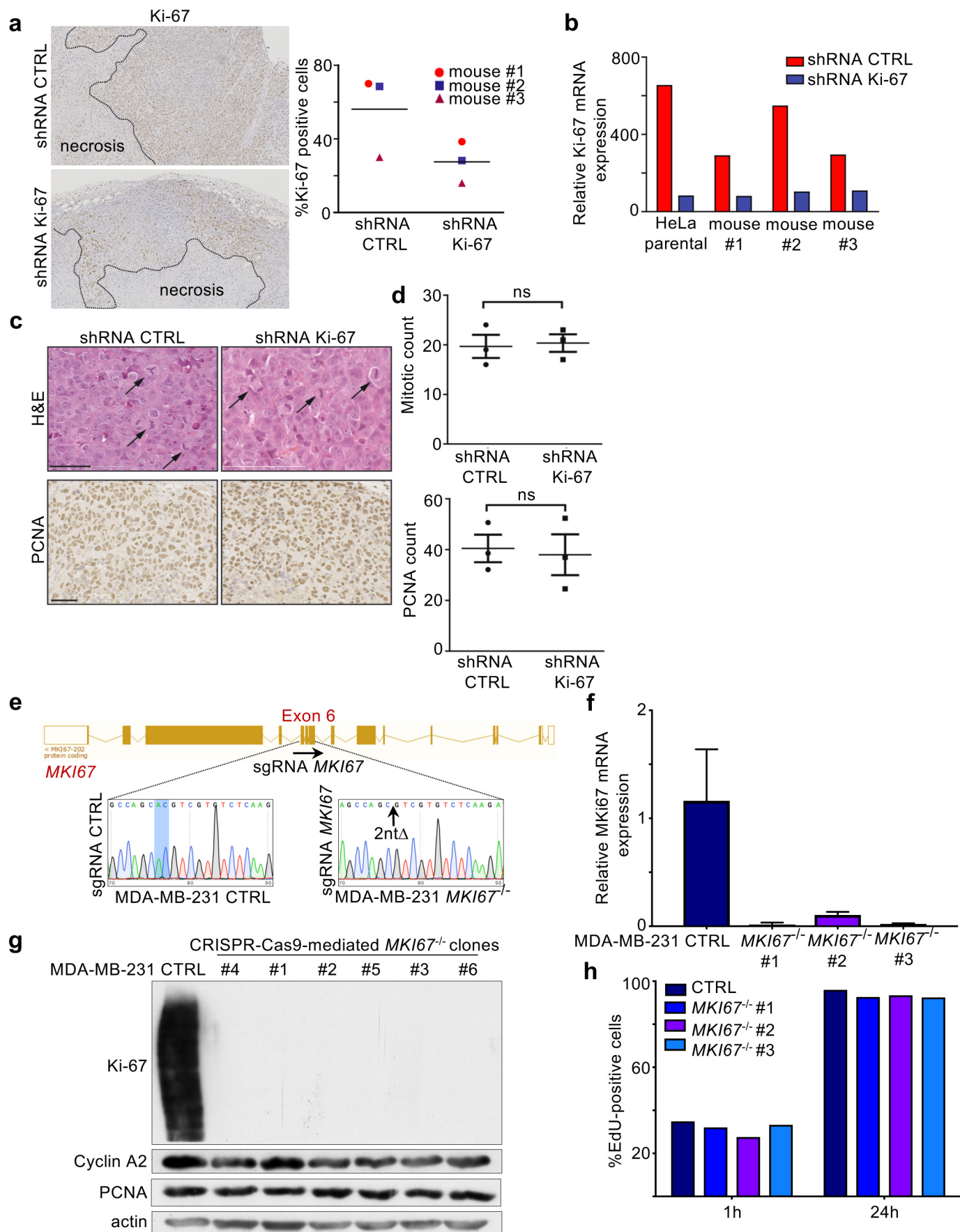
**a.** Body weight (g) measurement in *Mki67*<sup>+/2ntΔ</sup> and *Mki67*<sup>2ntΔ/2ntΔ</sup> mice during the AOM-DSS experiment. **b.**

Assessment of inflammatory response in colon and mesenteric lymph nodes in *Mki67*<sup>+/2ntΔ</sup> and *Mki67*<sup>2ntΔ/2ntΔ</sup> mice, treated or not with DSS (H&E staining; scale bar, 1mm).



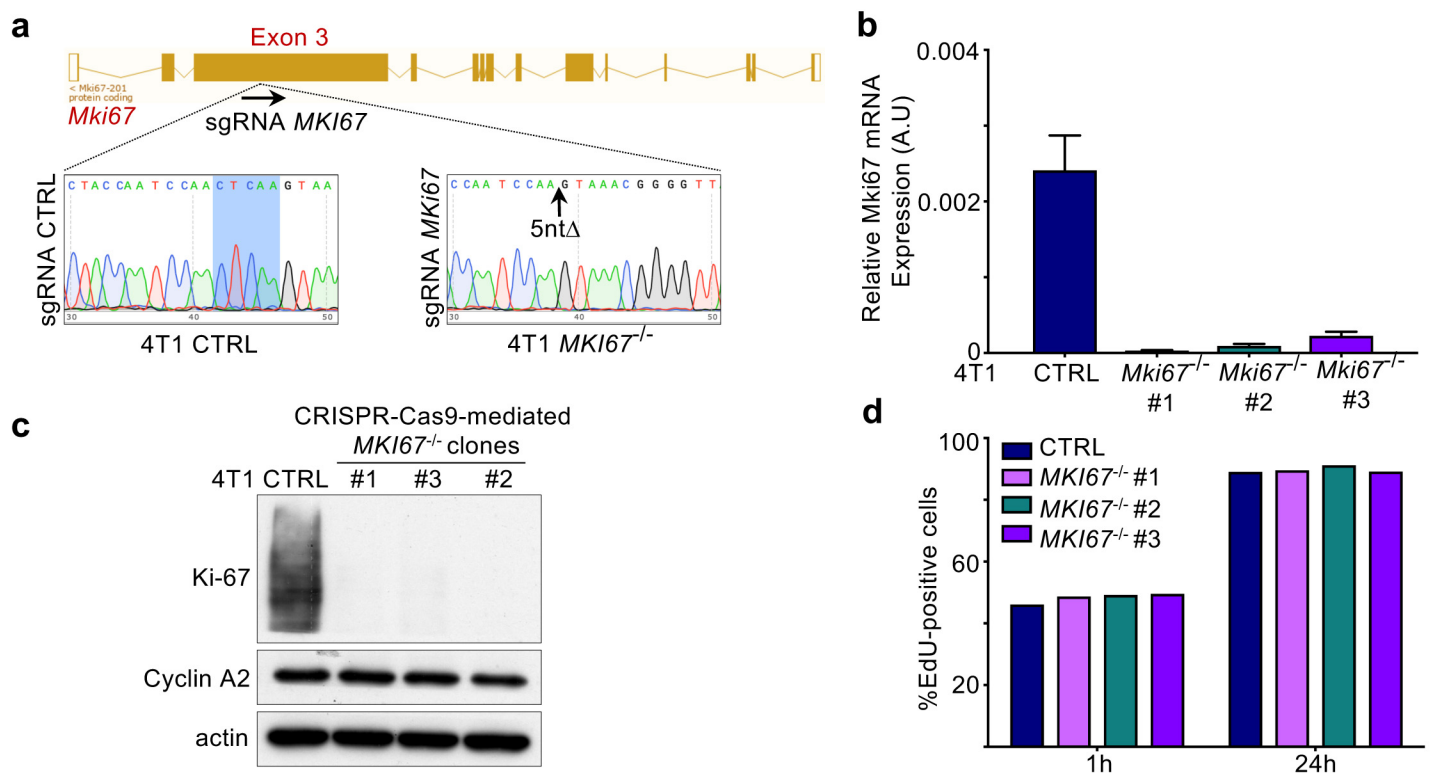
**Extended Data Fig. 3: *Mki67*<sup>2ntΔ/2ntΔ</sup> mice have normal response to haemolytic anaemia triggered by phenylhydrazine (PHZ) treatment.**

Analysis of number and percentage of indicated populations of blood, bone marrow and spleen cells in *Mki67*<sup>+ /2ntΔ</sup> and *Mki67*<sup>2ntΔ/2ntΔ</sup> mouse, treated or not with PHZ.



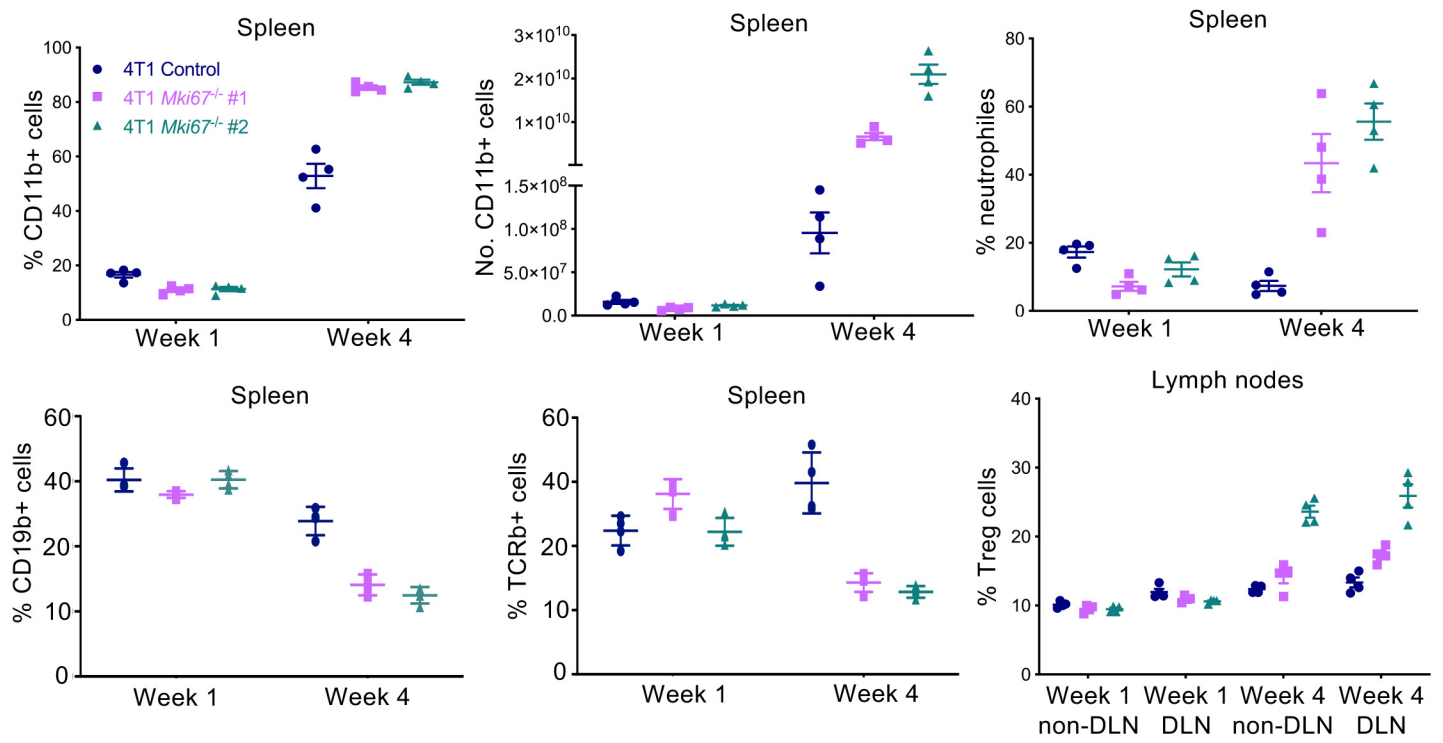
#### Extended Data Fig. 4: Ablation of Ki-67 does not affect cell proliferation.

**a-d.** Characterisation of HeLa shRNA CTRL and shRNA Ki-67 xenografts. **a.** Left, IHC staining for Ki-67; necrotic areas are highlighted. Right, quantification of Ki-67-positive cells in tumours ( $n=3$  mice). **b.** Relative Ki-67 mRNA expression levels in HeLa shRNA CTRL and shRNA Ki-67 parental cells and xenografts. **c.** H&E (top; arrows indicate mitotic cells) and PCNA (bottom) staining of HeLa shRNA CTRL and shRNA Ki-67 xenografts. **d.** Quantification of mitotic indices and PCNA-positive cells. **e-h.** Characterisation of MDA-MB-231 CTRL and *MKI67*<sup>-/-</sup> cells. **e.** Scheme of CRISPR-Cas9-mediated disruption of *MKI67* gene (targeting exon 6 and resulting in a 2nt-deletion). **f.** Relative Ki-67 mRNA expression in parental MDA-MB-231 cells and three *MKI67*<sup>-/-</sup> clones. **g.** Immunoblotting for the indicated proteins of parental MDA-MB-231 cells and six *MKI67*<sup>-/-</sup> clones. **h.** Quantification of the number of CTRL and *MKI67*<sup>-/-</sup> MDA-MB-231 EdU-positive cells, pulsed with EdU for either 1h or 24h.



### Extended Data Fig. 5: Knockout of *Mki67* in 4T1 cells.

**a.** Scheme of CRISPR-Cas9-mediated disruption of *Mki67* gene in 4T1 cells (targeting exon 3 and resulting in a 5nt-deletion). **b.** Relative Ki-67 mRNA expression in parental 4T1 cells and three *Mki67*<sup>-/-</sup> clones. **c.** Immunoblotting for the indicated proteins of parental 4T1 cells and *Mki67*<sup>-/-</sup> clones. **d.** Quantification of the number of CTRL and *Mki67*<sup>-/-</sup> EdU-positive cells, pulsed with EdU for either 1h or 24h.

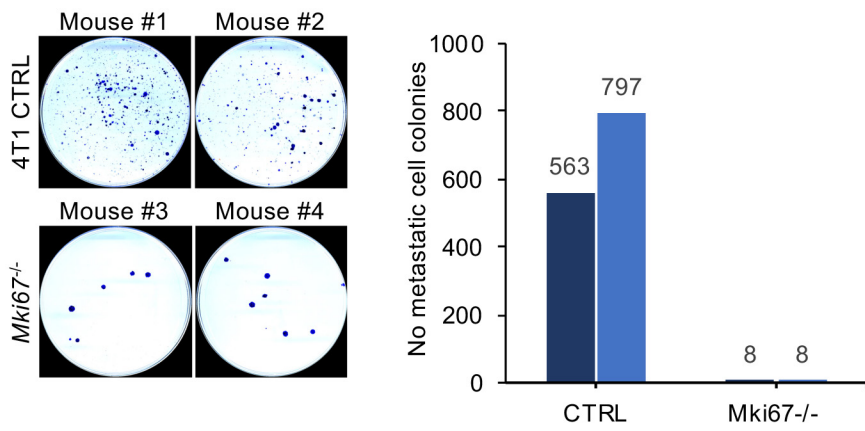


### Extended Data Fig. 6: Ki-67 knockout induces an immunosuppressive tumour environment.

Immunohistological analysis of spleens and lymph nodes (draining, DLN; and non-draining, non-DLN) performed in week 1 and 4 in mice xenografted with 4T1 CTRL and *Mki67*<sup>-/-</sup> cells (n=4).



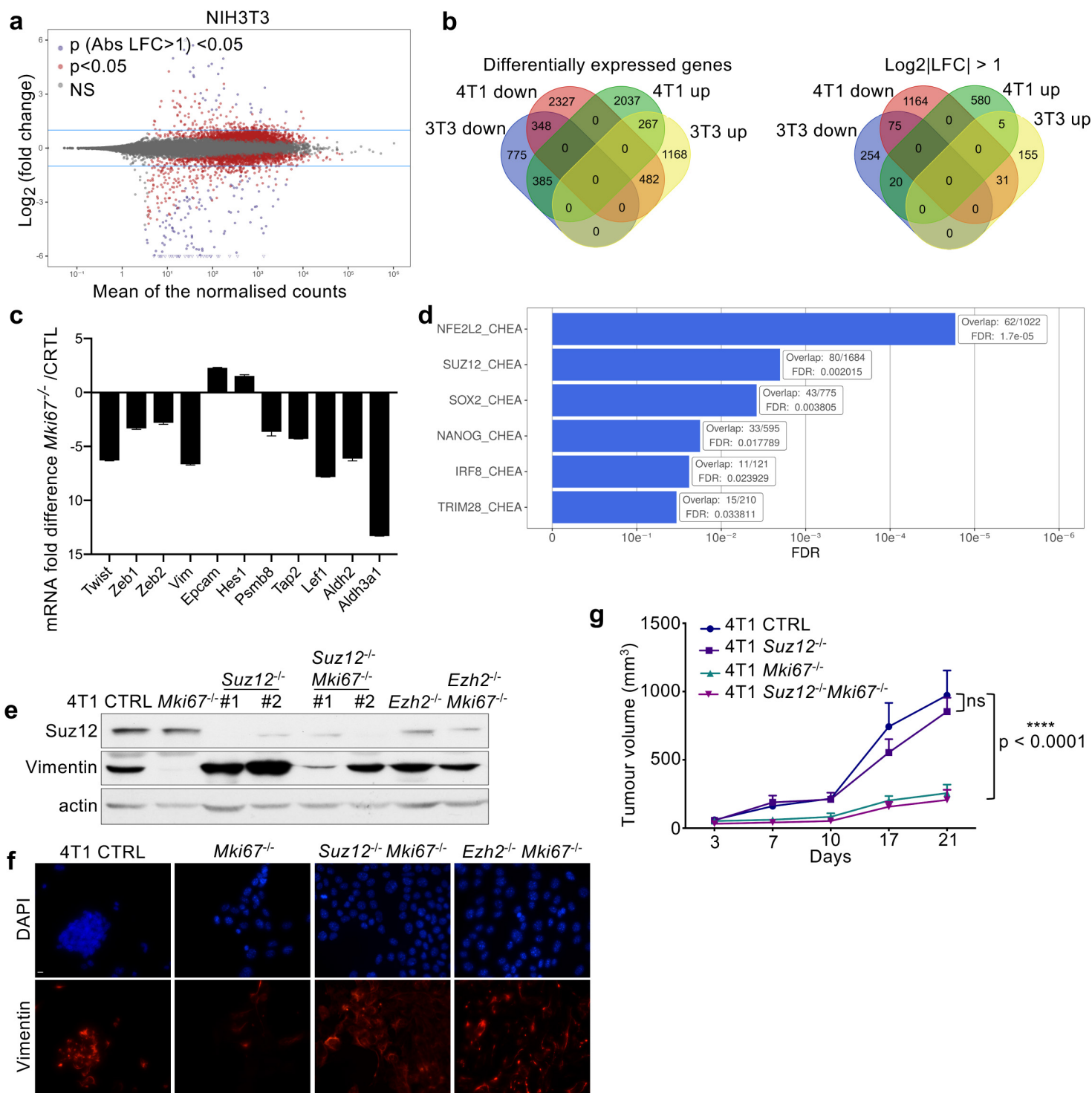
## Ext. Fig.7



### Extended Data Fig. 7: Ki-67 knockout reduces metastatic potential.

Lung tissue from mice injected via tail vein with 4T1 CTRL or *Mki67*<sup>-/-</sup> cells (2 per condition) was dissociated and resulting cells were maintained in the presence of 6-thioguanine, to select for 4T1 cells. Left, crystal violet staining of resulting colonies. Right, quantification of the number of 4T1 CTRL and *Mki67*<sup>-/-</sup> colonies.

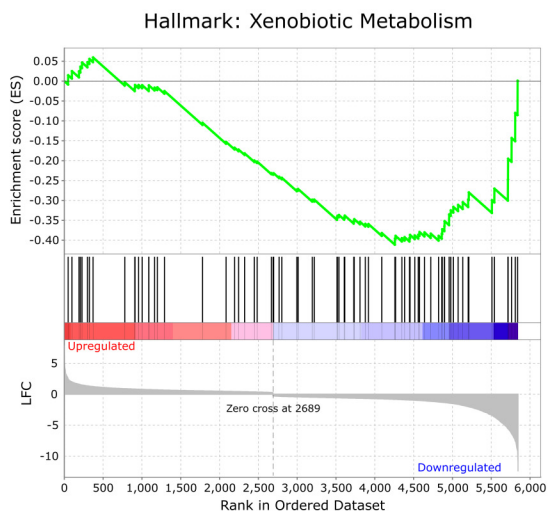




### Extended Data Fig. 8: Ki-67 controls gene expression, in part through PRC2.

**a.** Dot plot analysis of differentially expressed genes (DEGs) in 3T3 *Mki67*<sup>-/-</sup> cells. Red dots: DEGs with adjusted p-value < 0.05; purple dots: genes whose log<sub>2</sub> fold change, LFC > 1 or < -1, is statistically significant (adjusted p-value < 0.05); grey dots: not significant, NS. **b.** Venn diagrams of DEGs in 3T3 and 4T1 *Mki67*<sup>-/-</sup> cells under condition of p-value < 0.05 (left), and p-value (LFC > 1 or < -1) < 0.05 (right). **c.** Quantitative RT-PCR analysis of differentially expressed mesenchymal and epithelial markers in 4T1 *Mki67*<sup>-/-</sup> cells; fold change in expression  $\pm$  SD is shown. **d.** Gene set enrichment performed over the ENCODE and ChEA Consensus transcription factors from ChIP-X gene sets. FDR, false discovery rate adjusted p-values. **e.** Western-blot analysis of Suz12 and Vimentin levels in 4T1 cells, CTRL, and single or double knockouts for *Mki67*, *Suz12* and *Ezh2*. Actin serves as loading control. **f.** Immunofluorescence staining of Vimentin in 4T1 CTRL, *Mki67*<sup>-/-</sup>, *Mki67*<sup>-/-</sup>;*Suz12*<sup>-/-</sup> and *Mki67*<sup>-/-</sup>;*Ezh2*<sup>-/-</sup> cells. **g.** Tumour growth of 4T1 CTRL, *Suz12*<sup>-/-</sup>, *Mki67*<sup>-/-</sup>, and *Mki67*<sup>-/-</sup>*Suz12*<sup>-/-</sup> xenografts over 3 weeks (n=3).

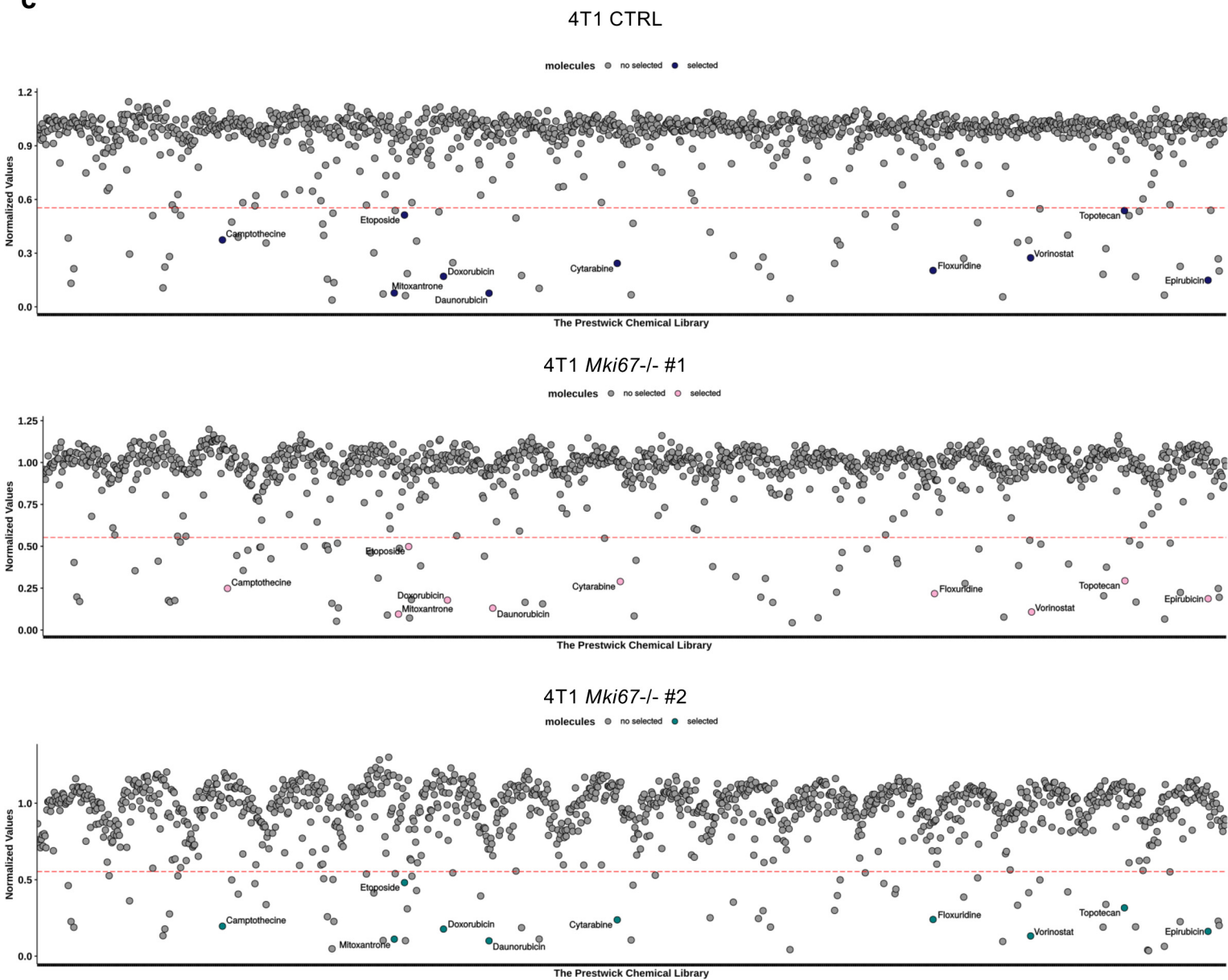
**a**



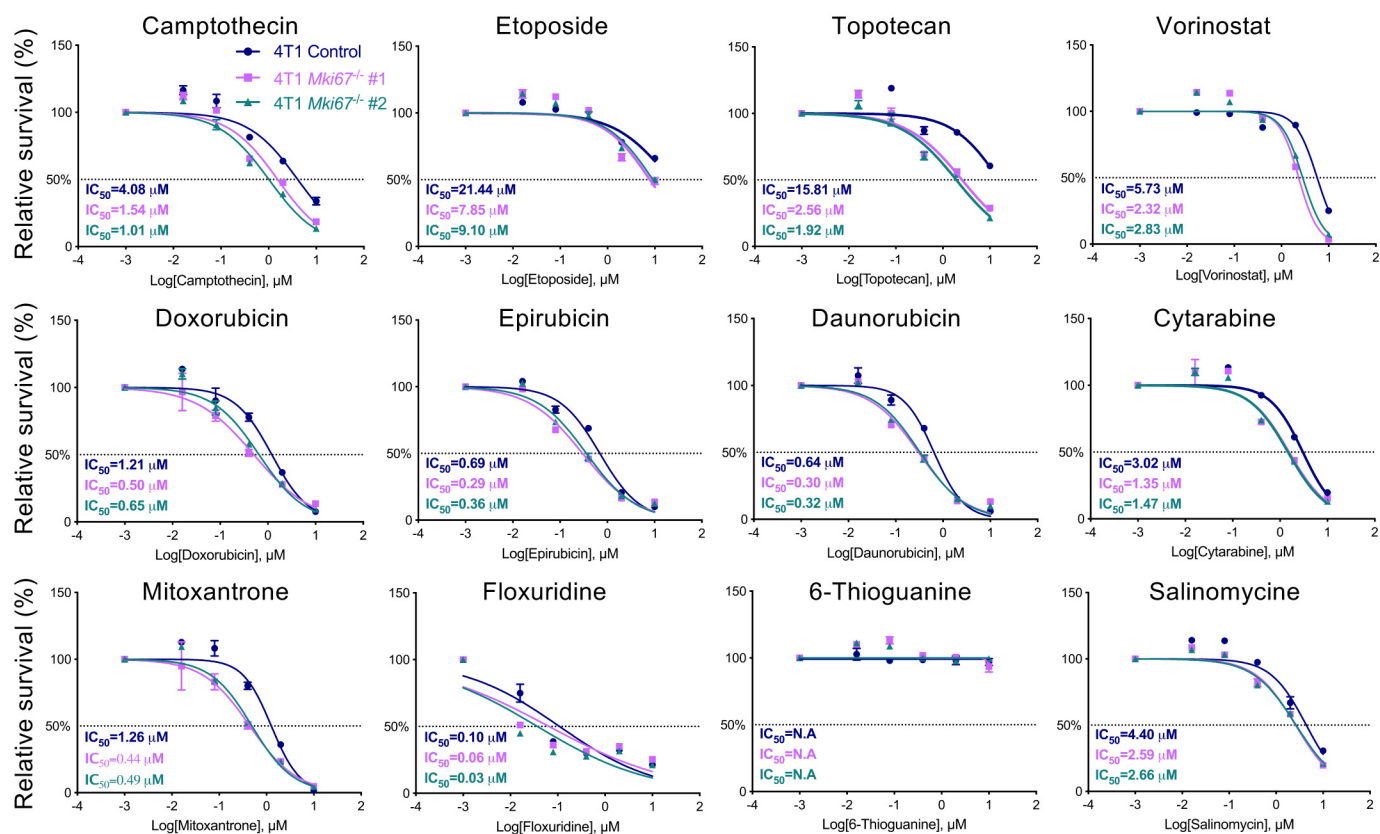
**b**

Upregulated	Downregulated
Phase I drug metabolising enzymes	
PTGS1	ALDH1I2, 3a1, 1b1 CYP1b1, 2c55, 3a13 PTGS2
Phase II drug metabolising enzymes	
	AOX1 BLVRB EPHX1 GSTA1, 2, 3 GSTM1, 2, 5 GSTT2, 3 MAOA MGST1, 2 NAT2, 6, 9 NQO1 PON3

**c**



d

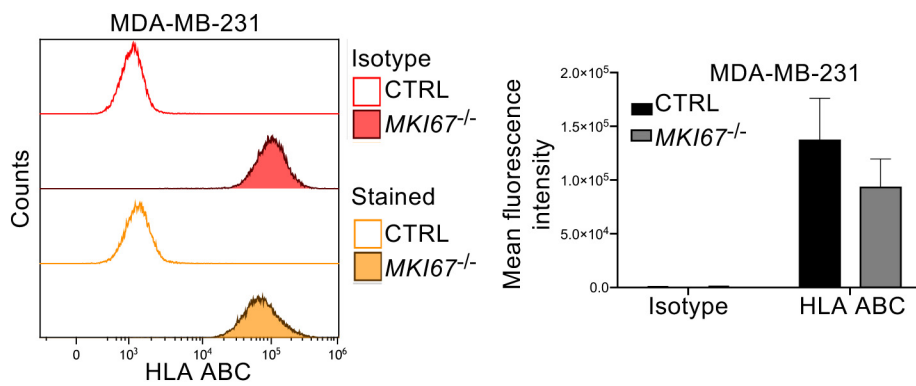


### Extended Data Fig. 9: Ki-67 knockout sensitises cells to various drugs.

a. Xenobiotic metabolism is one of hallmarks found in pathway analysis of genes downregulated in *Mki67*<sup>-/-</sup> cells. b. List of drug-metabolising enzymes with altered expression in *Mki67*<sup>-/-</sup> cells. c. Results of the automated gene-drug screen using the Prestwick chemical library. 'Hits' were identified applying the Z-score statistical method and are located below the dashed red line. Drugs chosen for subsequent dose-response experiments are highlighted in colour for CTRL and *Mki67*<sup>-/-</sup> clones. d. Dose-response curves of 4T1 CTRL or *Mki67*<sup>-/-</sup> cells treated with indicated compounds and their corresponding IC<sub>50</sub>.

**Ext. Fig.10**

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**Extended Data Fig. 10. MHC class I expression requires Ki-67 in MDA-MB-231 cells.** MDA-MB-231 CTRL and MKI67<sup>-/-</sup> cells were stained with anti-HLA-PE or control isotype. Top, representative images of FACS analysis. Bottom, quantification; median  $\pm$  SD (n=2 independent experiments).

## 1 **Methods**

2

### 3 **Cell lines and mice**

4 4T1 cells were provided by Robert Hipskind (IGMM, Montpellier); MDA-MB-231 cells were  
5 obtained from SIRIC, Montpellier. NIH 3T3, 4T1, MDA-MB-231 and S3 HeLa cells were  
6 grown in Dulbecco modified Eagle medium (DMEM - high glucose, pyruvate, GlutaMAX –  
7 Gibco® Life Technologies) supplemented with 10% foetal bovine serum (SIGMA or  
8 DUTSCHER). Cells were grown under standard conditions at 37°C in humidified incubator  
9 containing 5% CO<sub>2</sub>.

10 6-8 weeks-old female BALB/c (BALB/cOlaHsd), athymic nude (Hsd:Athymic  
11 Foxn1<sup>nu</sup>/Foxn1<sup>+</sup>), and NOD.SCID (NOD.CB17-Prkdc<sup>scid</sup>/NCrHsd) mice were purchased from  
12 Envigo. C57BL/6 Apc<sup>Δ14</sup> mice<sup>12</sup> were provided by Philippe Jay (IGF, Montpellier).

13

### 14 **Animal studies**

15 All animal experiments were performed in accordance with international ethics standards  
16 and were subjected to approval by the Animal Experimentation Ethics Committee of  
17 Languedoc Roussillon (n°APAFIS#8637-2017012311188975v3).

18

### 19 **Antibodies and plasmids**

20 Antibodies: Ki-67 (clone SP6; Abcam), cyclin A2 (6E6; Novocastra), PCNA (ab18197;  
21 Abcam), β-catenin (BD610154; BD-Bioscience), Ras (G12V Mutant Specific D2H12,  
22 #14412; CST), actin (A2066; Sigma), vimentin (D21H3, #5741; CST), E-Cadherin (24E10,  
23 #3195; CST), Suz12 (D39F6, #3737; CST), H3K27me3 (#39155, Active Motif), H2D<sup>d</sup>-FITC  
24 (#553579; BD Biosciences), H2K<sup>d</sup>-PE (SF1-1.1; BD Biosciences), HLA-PE (EMR8-5; BD  
25 Biosciences).

26 Lentiviral Vectors used: LentiCRISPRv2 (Addgene #52961), pMD2.G (Addgene #12259),  
27 psPAX2 (Addgene #12260). pHIV-Luc-ZsGreen (Addgene #39196) was used to generate  
28 bicistronic lentivirus that expresses both Luciferase and ZsGreen.

29 Retroviral vectors used: pBabe-puro (Addgene #1764), pBabe-puro H-Ras<sup>G12V</sup> (Addgene  
30 #1768). gag/pol (retroviral packaging) and Maloney (envelope) were gift from Leatitia  
31 Linares (IRCM, Montpellier).

32 pUC57-U6-sgRNA (#51132, Addgene); Cas9-VP12 (#72247, Addgene), modified by adding  
33 T2A-GFP at the C-terminal end (gift from Edouard Bertrand, IGMM).

34

### 35 **CRISPR/Cas9-mediated genome editing**



36 The sgRNAs targeting mouse *Mki67* exon 3 or human *MKI67* exon 6 and non-targeting  
37 control sequences were previously designed<sup>37</sup>, and cloning of the target sequence into the  
38 LentiCRISPRV2 lentiviral vector was conducted as described<sup>37</sup>. Lentiviruses encoding the  
39 sgRNA targeting sequences were produced. Transduced cells (4T1 and MDA-MB-231)  
40 expressing CRISPR/Cas9 were selected using puromycin. Resistant cells were isolated and  
41 seeded as single cell clones in 96 well-plates.

42 For Suz12 and Ezh2 knockout, sgRNA targeting sequences<sup>37</sup> were synthesised with Bpil  
43 sticky ends and cloned into the pUC57-U6-sgRNA vector. Cas9-VP12-T2A-GFP and sgRNA  
44 vectors were transfected into 4T1 WT and Ki-67<sup>-/-</sup> #2 clone using Lipofectamine 2000  
45 (ThermoFisher). 24h post-transfection, GFP-positive cells were sorted i96-well plates by  
46 flow cytometry (FACS Aria, BD). After 10-12 days of culture, colonies were picked and  
47 screened (Cellomics, Thermo), using Suz12 or H3K9me3 antibodies. Positive knockout  
48 clones were confirmed by PCR, IF and western blotting.

49

#### 50 **AOM-DSS-induced colon carcinogenesis**

51 Mice (*Mki67*<sup>+/+</sup>; *Mki67*<sup>+/<sup>2ntΔ</sup> & *Mki67*<sup>2ntΔ/<sup>2ntΔ</sup>) were given a single intraperitoneal injection of  
52 AOM (10 mg/kg in 0.9% saline; A5486, Sigma-Aldrich), and one week later, 2% Dextran  
53 Sodium Sulfate (DSS; MP Biomedicals) was administered in the drinking water for 7  
54 consecutive days. Mice were sacrificed at week 16-post AOM-DSS treatment and colon  
55 tissues were removed.</sup></sup>

56 Colons were flushed and fixed overnight in neutral buffered formalin (10%) before paraffin  
57 embedding. Briefly, 4μm thick sections were dewaxed in xylene and rehydrated in graded  
58 alcohol baths. Slides were incubated in 3% H<sub>2</sub>O<sub>2</sub> for 20 min and washed in PBS to quench  
59 endogenous peroxidase activity. Antigen retrieval was performed by boiling slides for 20 min  
60 in 10 mM sodium citrate buffer, pH 6.0. Nonspecific binding sites were blocked in blocking  
61 buffer (TBS, pH 7.4, 5% dried milk, 0.5% Triton X-100) for 60 min at RT. Sections were  
62 incubated with anti-β-catenin antibody diluted in blocking buffer overnight at 4°C. Envision+  
63 (Dako) was used as a secondary reagent. Signals were developed with DAB (Sigma-  
64 Aldrich). After dehydration, sections were mounted in Pertex (Histolab) and imaged using  
65 the Nanozoomer-XR Digital slide Scanner C12000-01 (Hamamatsu).

66

#### 67 **DNA replication assay, EdU labelling**

68 Cells were treated with 10 μM 5-ethynyl-2'-deoxyuridine (EdU; LifeTechnologies) for the  
69 indicated time, harvested, washed once with cold PBS, resuspended in 300μL cold PBS  
70 and fixed with 700μL ice-cold 100% ethanol. Click reaction was performed according to the

71 manufacturer instructions (Click-iT™ Plus EdU Alexa Fluor™ 647 Flow Cytometry Assay  
72 Kit; Invitrogen) and cells were analysed by flow-cytometry (BD FACSCanto II). FlowJo®  
73 software was used for analysis.

74

### 75 **Mammosphere assay**

76 The mammosphere formation assay was performed as previously described<sup>38</sup>. Briefly, 500  
77 cells were plated per well in a low-adherence 96-well plate coated with poly-2-hydroxyethyl-  
78 methacrylate (poly-Hema). After 7 days in culture (37°C, 5% CO<sub>2</sub>), images of formed  
79 mammospheres were acquired and counted using automated high-content microscopy  
80 analysis (Cellomics, Thermo).

81

### 82 **Aldehyde Dehydrogenase 1 (ALDH1) activity**

83 ALDH1 enzymatic activity was determined using the ALDEFLUOR kit (Stem Cell  
84 Technologies) according to the manufacturer instructions. For each sample, half the  
85 cell/substrate mixture was treated with diethylaminobenzaldehyde (DEAB; ALDH inhibitor).  
86 ALDEFLUOR/DEAB treated cells were used to define negative gates. FACS was performed  
87 with  $\geq 10^5$  cells.

88

### 89 **Xenografts**

90 Animals were housed in the animal facility of IGMM and were maintained in a pathogen-free  
91 environment and fed *ad libitum*.

92 To generate primary tumours,  $10^6$  cells (4T1) or  $3 \times 10^6$  cells (MDA-MB-231) of log-phase  
93 viable 'mouse pathogen-free' (Test: IMPACT1, Iddex) were implanted into the fourth inguinal  
94 mammary gland (in 50  $\mu$ l PBS (4T1) or 200  $\mu$ l PBS (MDA-MB-231)).

95 Primary tumour volume was measured every week by electronic calliper using the formula  
96 " $\pi/6 * S^2 * L$  (Smaller radius)\*L (Larger radius)".

97 At the end of the experiment, following sacrifice, primary tumours were excised and fixed  
98 over-night in neutral buffered formalin (10%) before paraffin embedding (see above). IHC  
99 analysis of Ki-67 expression of the different tumour tissue sections was conducted as  
100 described above.



## 101 **Visualisation of lung metastases**

102 Dissected lungs were stained with 15% India Ink diluted in distilled water, and subsequently  
103 washed with 5 ml Fekete's solution (40 ml glacial acid acetic, 80 ml formalin, 580 ml ethanol  
104 100%, 200 ml water). A binocular microscope connected to a digital camera was used to  
105 visualise and count the metastatic nodules.

## 106 **Automated drug library screen**

107 We performed an initial screen of all 1,280 FDA-approved drugs in the Prestwick Chemical  
108 Library (Prestwick Chemical, Illkirch-Graffenstaden, France) at 10  $\mu$ M in white clear-bottom  
109 96-well plates. Briefly, we prepared seeding solutions at a density of  $2 \times 10^5$  cells/mL in  
110 DMEM complete medium (10%FBS), and dispensed 100  $\mu$ L into 96-well plates using a  
111 Multidrop Combi Dispenser (Thermo Scientific™). Next, we prepared 1x treatment solutions  
112 of the Prestwick compounds in DMEM complete medium at a concentration of 10  $\mu$ M using  
113 a Tecan EVO200 robotic liquid handling system (Tecan Trading AG). We removed medium  
114 from the pre-seeded 96-well plates and we dispensed 100  $\mu$ L of 1x treatment solutions using  
115 the Tecan robot. We incubated plates at 37°C, 5% CO<sub>2</sub> for 48 h in an automated incubator  
116 (Cytomat, Thermo Scientific™) associated with the Tecan robot. After treatment, cell  
117 survival was determined using the CellTiter-Glo® Luminescent Cell Viability Assay  
118 (Promega Corporation) in accordance with the manufacturer's instructions. Luminescence  
119 signal was automatically measured with a Tecan Infinity F500 microplate reader (Tecan  
120 Trading AG). "Hits" were identified applying the Z-score statistical method (see statistical  
121 analysis section). The same method was used for subsequent dose-response experiments  
122 on selected "hits".

123

## 124 **Induction of anemia with PHZ**

125 Anemia was induced in adult *Mki67<sup>2ntΔ/2ntΔ</sup>* mice and littermate controls (*Mki67<sup>+/2ntΔ</sup>*) by  
126 injection of 40 mg/kg phenylhydrazine (PHZ) on days 0, 1, and 3. Animals were sacrificed  
127 on day 5 and blood, spleen and bone marrow were collected. Hematocrit was measured as  
128 previously described<sup>39</sup>. Cell suspensions were generated and immunophenotyping  
129 performed by flow cytometry.

## 130 **Immunophenotyping and flow cytometry analyses**

131 Cells isolated from peripheral lymph nodes tumor (draining and non-draining for tumor-  
132 bearing mice) as well as spleen and bone marrow, were stained with Sytox blue or Live/dead

133 fixable viability dye (Life Technologies and eBioscience, respectively) together with the  
 134 appropriate conjugated anti-CD3, CD45, CD62L, CD4, CD8, CD44, CD25, F4/80, CD19,  
 135 B220, NK1.1, Gr1, CD11b, CD11c, Ter119, CD71 and Foxp3 antibodies  
 136 (eBioscience/ThermoFisher or Becton Dickinson, San Diego, CA). Intracellular staining for  
 137 Foxp3 was performed using the eBioscience Fixation/Permeabilization kit.

138 Cells were analysed on a FACS Fortessa (BD Biosciences) flow cytometer. Data analyses  
 139 were performed using FlowJo Mac v.10 (Tree Star) or DIVA (Becton Dickinson) software.

140

141 Prior to intracellular cytokine stainings, cells were activated with PMA (Sigma-Aldrich; 100  
 142 ng/ml)/ Ionomycin (Sigma-Aldrich; 1ug/ml) in the presence of brefeldin A (Sigma-Aldrich;  
 143 10ug/ml) for 3.5-4h at 37°C. Cell surface staining was performed, after which cells were  
 144 fixed and permeabilised using a fixation/permeabilization kit (BD Biosciences or  
 145 eBioscience) followed by intracellular cytokine staining.

146

#### 147 **Antibodies used for immunophenotyping**

148

	clone	fluorochrome	reference
Murine anti-CD3	145-2C11	V421	562600
Murine anti-CD4	RM4-5	BV711	563726
Murine anti-CD8	53-6.7	FITC	553031
Murine anti-CD8	53-6.7	AF700	564983
Murine anti-CD62L	MEL-14	APC	553152
Murine anti-CD44	IM7	PE-CF594	562464
Murine anti-CD45	30-F11	AAF	47-0451-82
Murine anti-CD25	PC61	AAF	47-0251-82
Murine anti-CD19	1D3	BV650	563235
Murine anti-NK11	PK136	APC	550627
Murine anti-CD11b	M1/70	BV605	563015
Murine anti-CD11c	HL3	PE	557401
Murine anti-LY6G	RB6-8C5	PE-CF594	562710
Murine anti-LY6C	AL-21	V450	560594
Murine anti-F4/80	BM8	PEC7	25-4801-82
Murine anti-Foxp3	FJK-16S	PEC7	25-5773-82
Murine anti-TNF $\alpha$	MP6-XT22	PE	12-7321-82
Murine anti-IL-2	JES6-5H4	PEC7	560538
Murine anti-IFN $\gamma$	XMG1.2	APC	17-7311-82
Murine anti-IL17A	TC11-18H10	BV605	564169
Murine anti-Ter119	Ter-119 (hamster)	FITC	557915

Murine anti-CD71	C2	BV605	563013
Murine anti-B220	RA3-6B2	eF780	47-0452-82

149

## 150 **Cell extracts and Western-blotting**

151 Frozen cell pellets (harvested by trypsinization, washed with cold PBS) were lysed directly  
152 in Laemmli buffer at 95°C. Protein concentrations were determined by BCA protein assay  
153 (Pierce Biotechnology). Proteins were separated by SDS-polyacrylamide gel  
154 electrophoresis (SDS-PAGE) (7.5% and 12.5% gels) and transferred to Immobilon  
155 membranes (Milipore) at 1.15 mA/cm<sup>2</sup> for 120 min with a semi-dry blotting apparatus.  
156 Membranes were blocked in TBS-T pH 7.6 (20 mM Tris, 140mM NaCl, 0.1% Tween-20)  
157 containing non-fat dry milk (5%), incubated with the primary antibody for 2 hours at RT or  
158 over-night at 4°C, washed several times with TBS-T for a total of 45 minutes, incubated with  
159 secondary antibody at 1/5000 dilution for 1 hour at RT and washed several times in TBS-T.  
160 Signals were detected using Western Lightning Plus-ECL (PerkinElmer) and Amersham  
161 Hyperfilm™ (GE Healthcare).

162

## 163 **qRT-PCR**

164 For reverse transcription (cDNA synthesis), 500 ng of purified RNA in total volume of 10µl,  
165 extracted by RNeasy Mini Kit (Qiagen), were mixed with 1µl of 10mM dNTPs mix  
166 (LifeTechnologies) and 1µl of 50µM random hexaprimers (NEB). Samples were incubated  
167 at 65°C for 5 minutes, transferred to ice, 5µl 5x First Strand Buffer, 2µl 100mM DTT and 1µl  
168 RNasin RNase Inhibitor (Promega) were added and samples were incubated at 25°C for 10  
169 minutes, 42°C for 2 minutes. 1µl of SuperScript® III Reverse Transcriptase  
170 (LifeTechnologies) was added to each sample and incubated at 50°C for 60 minutes, 70°C  
171 for 15 minutes.

172 qPCR was performed using LightCycler® 480 SYBR Green I Master (Roche) and  
173 LightCycler® 480 qPCR machine. The reaction contained 5ng of cDNA, 2µl of 1µM qPCR  
174 primer pair, 5µl 2x Master Mix, and final volume made up to 10µl with DNase free water.  
175 Primers used for both mouse (Mki67) and human (MKI67) Ki-67 in addition to the  
176 housekeeping genes are similar to those used by Sobecki et al., 2016. qPCR was conducted  
177 at 95°C for 10 min, and then 40 cycles of 95°C 20s, 58°C 20s and 72°C 20s. The specificity  
178 of the reaction was verified by melting curve analysis.

179

## 180 **PCR primers used in this study.**

181

Gene name	Sequence	PrimerBank ID	Reference
Actb	Fw- GGCCCAGAGCAAGAGAGGTATCC Rv- ACGCACGATTTCCCTCTCAGC	-	40
Gapdh	Fw- CGTCCCGTAGACAAAATGGT Rv- TGA CTGTGCCGTTGAATTTG	-	41
B2m	Fw- ACGTAACACAGTTCCACCCG Rv- CAGTCTCAGTGGGGGTGAAT	-	42
Gusb	Fw- AACAAACACACTGACCCCTCA Rv- ACCACAGATCGATGCAGTCC	-	42
Zeb2	Fw- GGCAAGGCCTTCAAGTACAA Rv- AAGCGTTTCTTG CAGTTTGG	-	43
Twist1	Fw- AGCGGGTCATGGCTAACG Rv- GGACCTGGTACAGGAAGTCGA	-	44
Tap2	Fw- CTGGCGGACATGGCTTTACTT Rv- CTCCC ACTTTTAGCAGTCCCC	7549791a1	45
Psmb8	Fw- ATGGCGT TACTGGATCTGTGC Rv- CGCGGAGAACTGTAGTGTCC	6755208a1	45
Zeb1	Fw- CGCCATGAGAAGAACGAGGAC Rv- CTGTGAATCCGTAAGTGTCTTT	357197186c2	45
Vim	Fw- CTGCTTCAAGACTCGGTGGAC Rv- ATCTCCTCCTCGTACAGGTCCG	227430362c2	45
Epcam	Fw- GGAGTCCCTGTTCCATTCTTCT Rv- GCGATGACTGCTAATGACACCA	11229327c3	45
Hes1	Fw- GATAGCTCCCGGCATTCCAAG Rv- GCGCGGTATTTCCCAACA	31560817c2	45
Lef1	Fw- GCCACCGATGAGATGATCCC Rv- TTGATGTCCGGCTAAGTCGCC	118130133c1	45
Aldh1l2	Fw- TTTCTGAGGGGATCAAGGC Rv- GACCTCGAATCCAGTTATGCAA	283436217c3	45
Aldh3a1	Fw-AATATCAGTAGCATCGTGAACCG Rv- GGAGAGCCCCTTAATCGTGAAA	6680676a1	45

182

183

#### 184 **Colony formation assay**

185 3T3 wild-type (WT) and two Ki-67 TALEN-mutant clones were transduced with either empty  
 186 control or H-Ras<sup>G12V</sup> expressing retroviruses. After verification of Ras<sup>G12V</sup> expression, cells  
 187 were seeded at 10<sup>5</sup> cells/well (6 well-plate) in triplicate and allowed to grow for 2-weeks

188 (media were changed every 2 days). Cells were then fixed (4% formaldehyde) and stained  
189 with 0.5% (w/v) crystal violet to visualise the colonies formed.

190

### 191 **RNA sequencing library prep**

192 Total RNA was extracted using Trizol (Life Technologies) following manufacturer's  
193 instructions from wild type and two clones of 4T1 Ki-67 knockout cell lines. RNA integrity  
194 was analysed on Agilent 2100 bioanalyzer. For library preparation, cDNA synthesis was  
195 performed on rRNA-depleted samples using the TruSeq Stranded Total RNA Library  
196 Preparation (RS-122-2301). All sequencing libraries were prepared in two or three biological  
197 replicates. Indexed cDNA libraries were sequenced by MGX (Montpellier) on Illumina  
198 HiSeq2000 with a single 50 bp read and a 10 bp index read.

199

### 200 **Sequencing of cDNA libraries and data processing**

201 FastQC was used to perform quality control of the sequencing. Using the tool STAR  
202 2.6.0a(2), all the reads that passed the quality control were aligned to the mouse reference  
203 genome (GRCm38.p6) and the counts per gene were quantified. The release 93 of the  
204 Ensembl mouse genome annotations were used for establishing the coordinates of each  
205 gene and their transcripts. Differential expression analysis was performed in R using the  
206 DESeq2(2) library embedded in an in-house script. After normalization of the read counts  
207 per gene, a negative binomial generalised linear model was fitted considering single factor  
208 design for assessing the differential expression between Mki67 knockout and wild-type  
209 samples. Gene set enrichment analysis of the 4T1 *Mki67*<sup>-/-</sup> cell line vs WT cell line was  
210 performed using the javaGSEA(3) desktop application with a log-fold-change pre-ranked  
211 gene list.

212

### 213 **Gene set enrichment for transcription factors.**

214 The gene set enrichment was performed over the ENCODE and ChEA Consensus TFs from  
215 CHIP-X gene set, which contains consensus target genes for transcriptions factors present  
216 in ENCODE and ChEA databases. The p value was computed with the Fisher Exact test  
217 and then adjusted for multiple hypothesis testing in order to obtain the FDR (false discovery  
218 rate) adjusted p-value.

219

### 220 **MHC class I expression analysis**

221 The MHC class I expression analysis was performed as previously described<sup>46</sup>. 4T1 CTRL  
222 and *Mki67*<sup>-/-</sup> cells were incubated with anti-H2D<sup>d</sup>-FITC, anti-H2K<sup>d</sup>-PE or control isotypes, and



223 MDA-MB-231 CTRL and *Mki67*<sup>-/-</sup> cells were incubated with anti-HLA-PE or control isotype,  
224 in PBS/BSA 2% at 4°C for 20 min. After 2 washes in PBS/BSA 2%, cells were analysed by  
225 FACS.

226

## 227 **Statistical analysis**

228 Significant differences between the different experimental groups were tested using an  
229 unpaired two-tailed Student's *t*-test or ANOVA in Prism 5 (GraphPad). For all analyses, *p*  
230 values < 0,05 (\*), *p* values < 0,01 (\*\*), *p* values < 0,001 (\*\*\*) and *p* values < 0,0001 (\*\*\*\*)  
231 were considered to indicate a statistically significant result.

232

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