

Supplementary Information

Dichloroacetate radiosensitizes hypoxic breast cancer cells
de Mey et al.

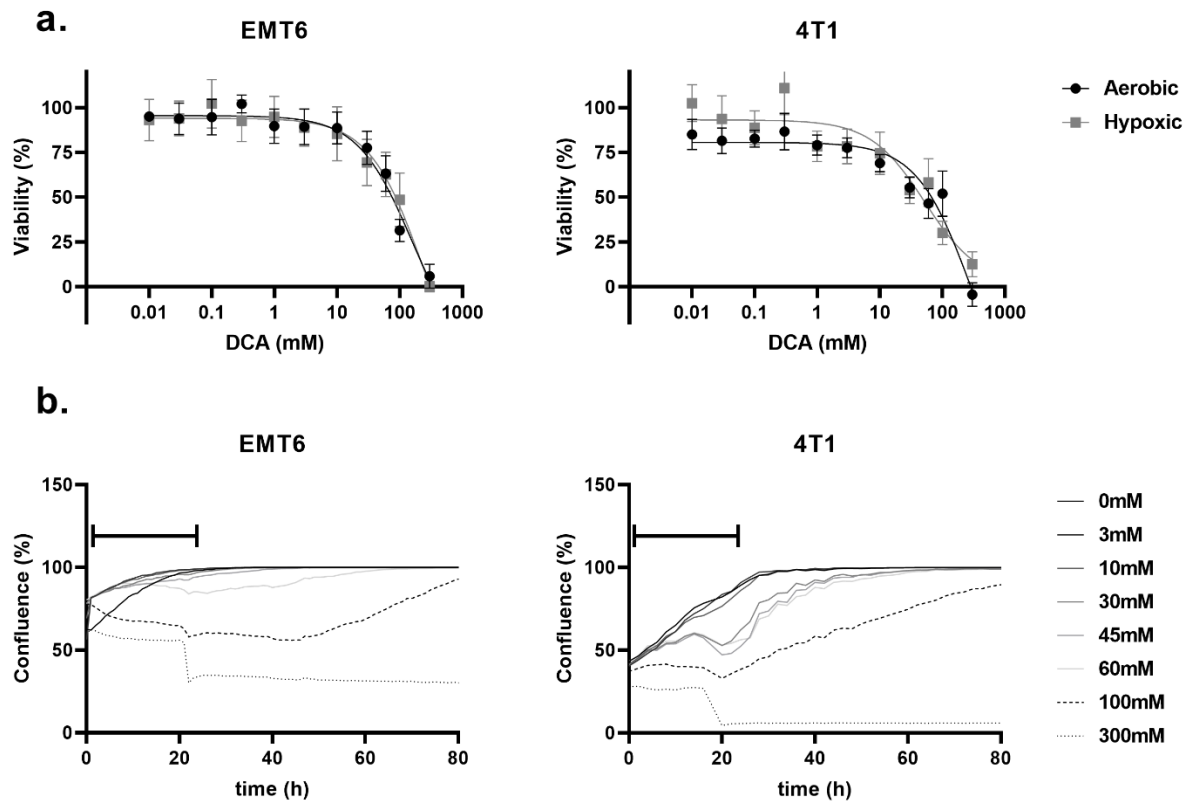


Figure S1: Growth of EMT6 and 4T1 breast cancer cells after treatment with DCA at indicated conditions. (a) Cell viability was assessed by MTT assay after treatment with DCA, and IC_{50} was calculated accordingly. (b) Cell proliferation was evaluated by live-confluence measurement with the Incucyte system. Treatment time of DCA is indicated with the capped line.

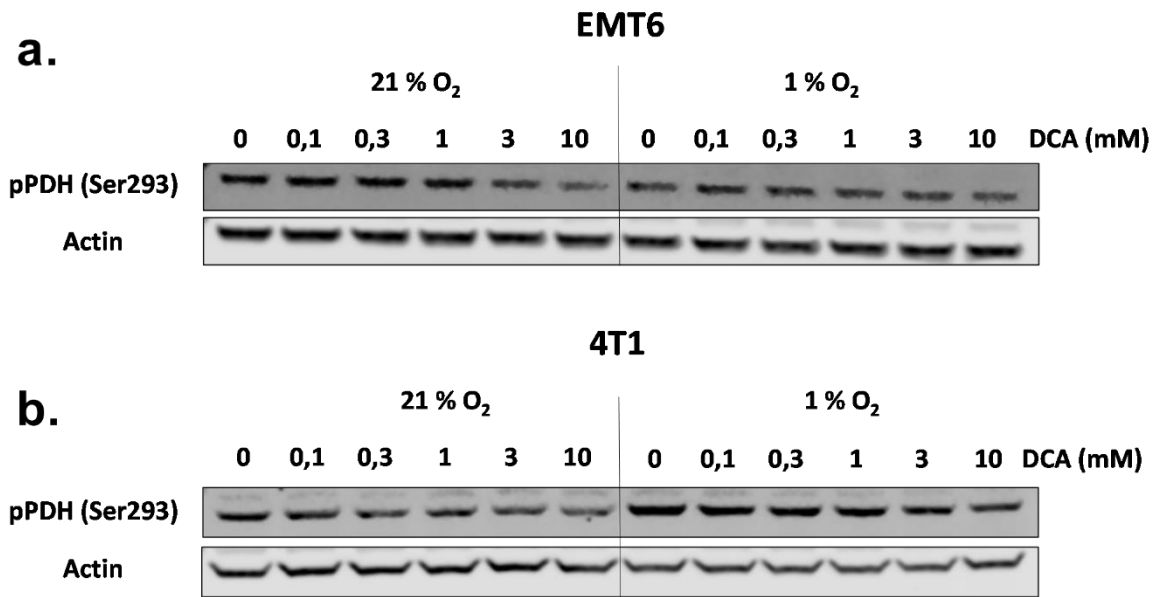


Figure S2: DCA decreases phosphorylated PDH of TNBC cells. Representative western blot of p-PDH (Ser293) in EMT6 (a) and 4T1 (b) cells, after treatment with DCA (0.1mM, 0.3mM, 1mM, 3mM and 10mM) under aerobic and hypoxic conditions.

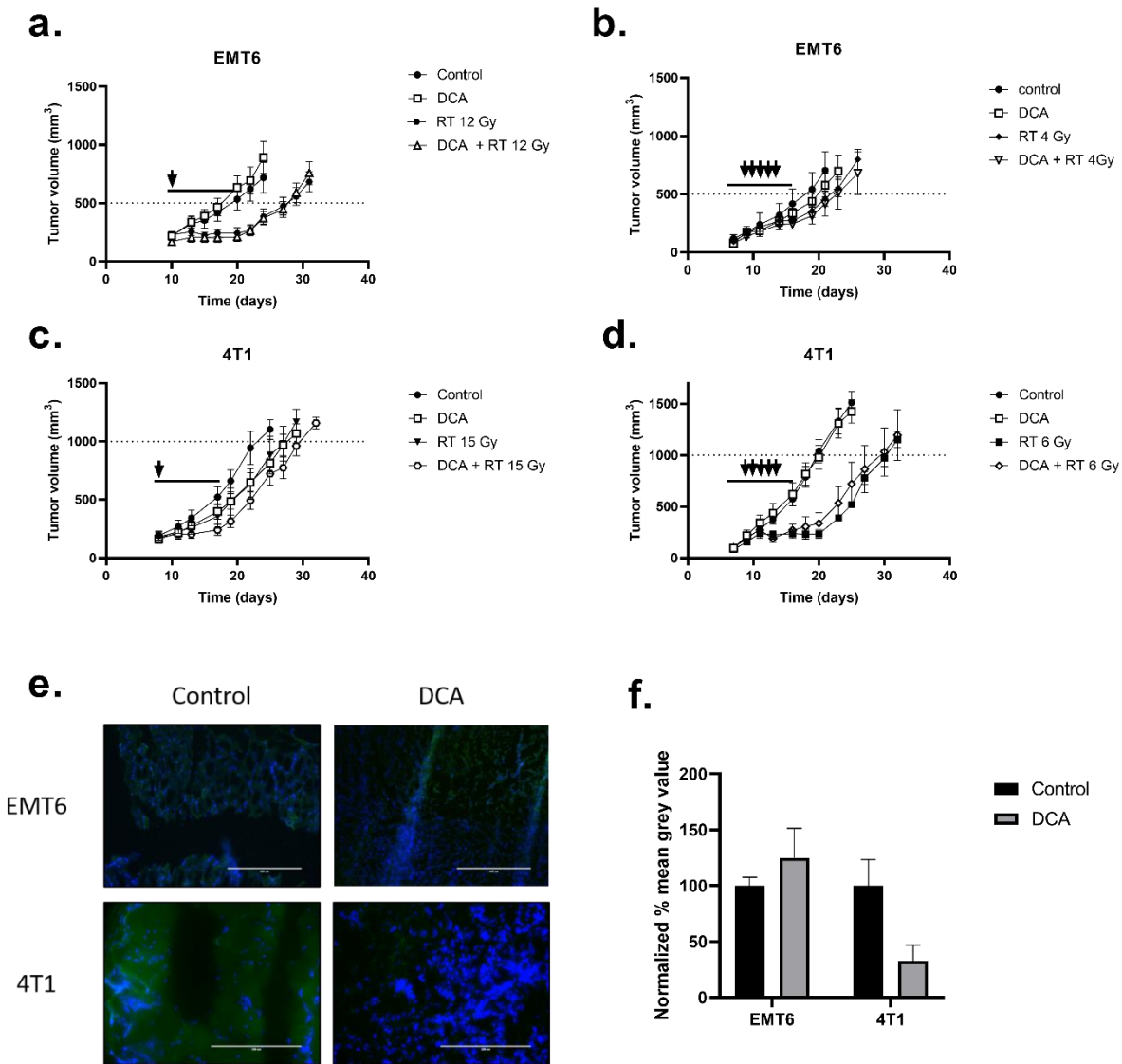


Figure S3: DCA fails to radiosensitize tumour cells *in vivo*. Tumour growth of EMT6 tumours in Balb/c mice administered with DCA 300mg/kg (ip. or it.) for 10 consecutive days and single dose radiation (12Gy) (a) or fractionated radiation (5*4Gy) (b). Tumour growth of 4T1 tumours in Balb/c mice daily for treated with DCA 300mg/kg (ip.) for 10 consecutive days and single fraction radiation of 15Gy (c), or treated with fractionated radiation of 5*6Gy (d). (e) Representative pictures and quantified data (f) of pimonidazole and dapi staining of EMT6 and 4T1 tumours.

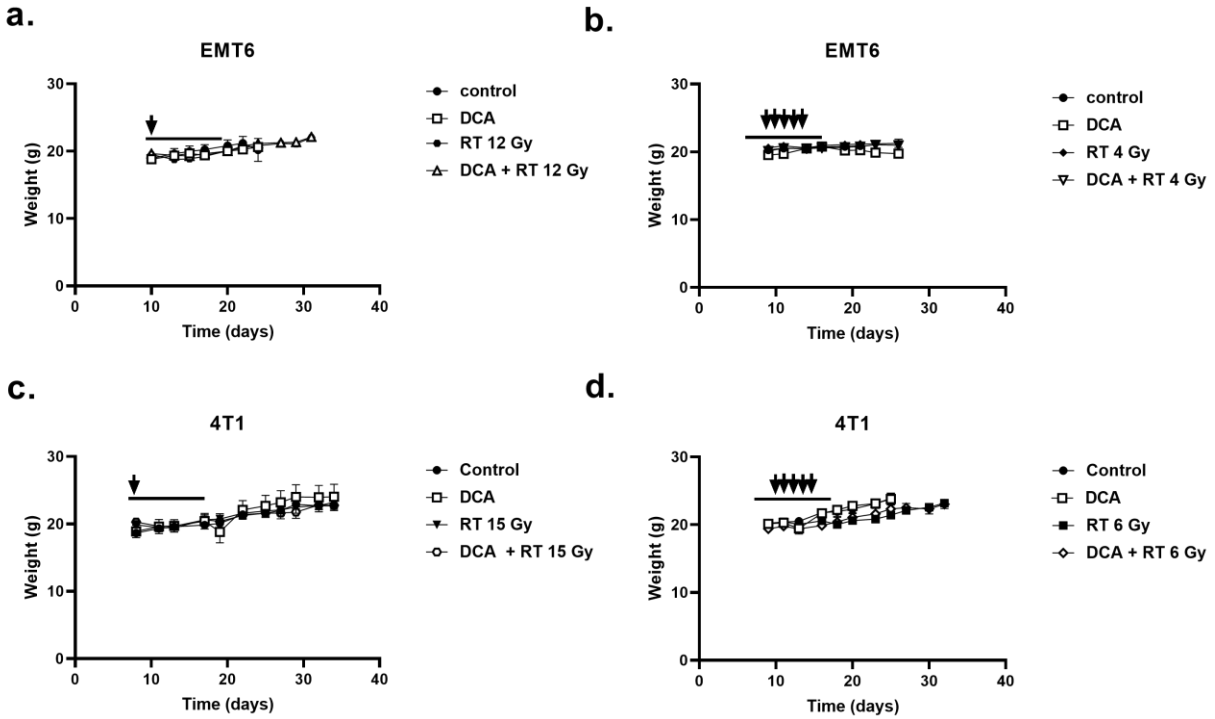


Figure S4: DCA and radiotherapy treatment does not cause toxicity in the mice. Weight of EMT6 inoculated Balb/c mice treated with DCA 300mg/kg (i.p. or i.t.) for 10 consecutive days and single dose radiation (12Gy) (a). Weight of EMT6 inoculated Balb/c mice treated with DCA 300mg/kg (i.p. or i.t.) for 10 consecutive days and fractionated radiation (5*4Gy) (b). Weight of 4T1 inoculated Balb/c mice treated with DCA 300mg/kg (i.p.) for 10 consecutive days and single fraction radiation of 15Gy (c). Weight of 4T1 inoculated Balb/c mice treated with DCA 300mg/kg (i.p.) for 10 consecutive days and treated with fractionated radiation of 5*6Gy (d).