

# Supplementary Data: Figures and Tables

Sulistiyani et al. IJMS Mar 2021

Revised 17 March 2021

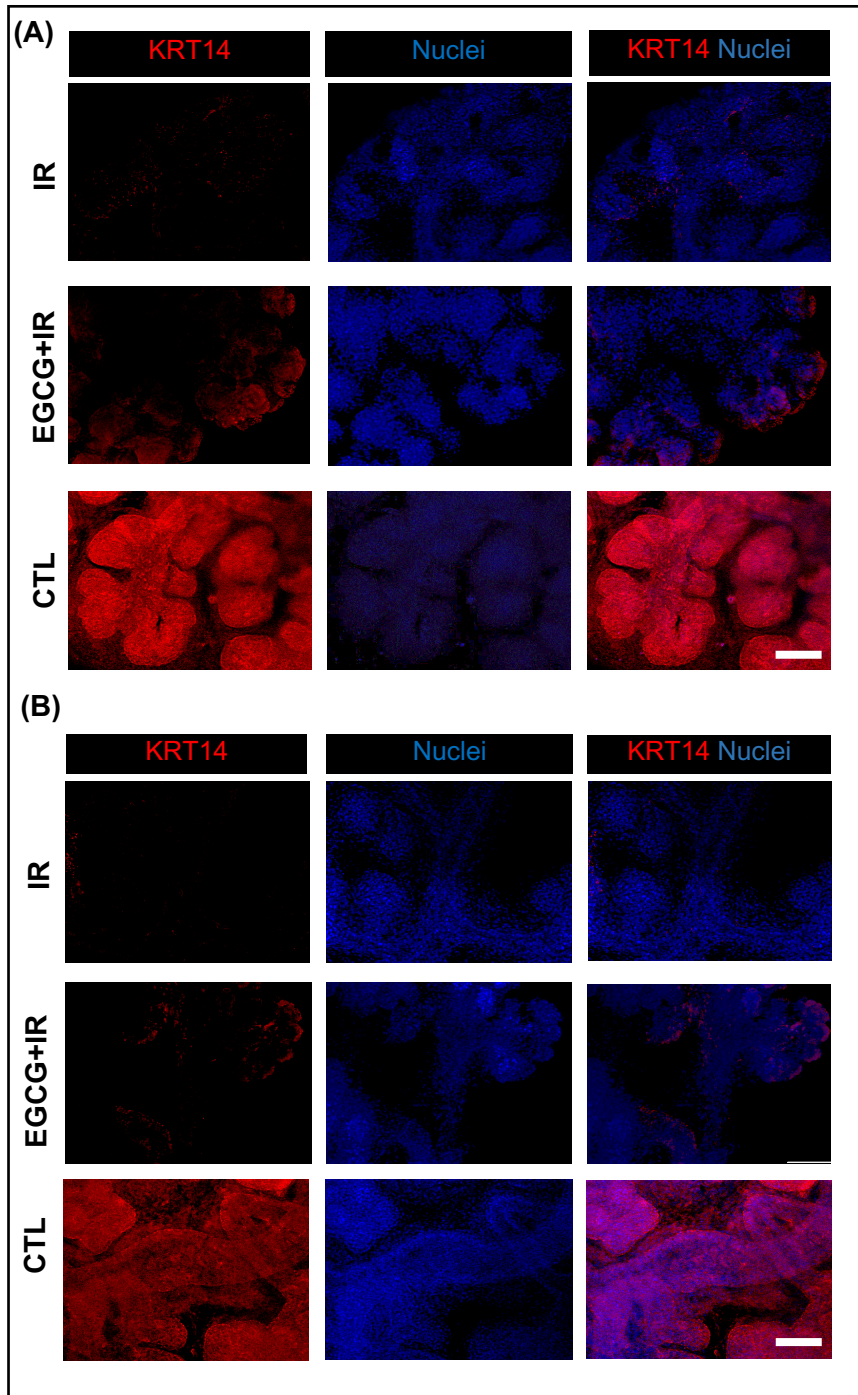
**Table S1. List of primary and secondary antibodies were used.**

Manufacturers: Life Technologies and Invitrogen are subsidiaries of Thermo Fisher Scientific (USA), Santa Cruz (USA), Abcam (UK), Cell Signaling Technologies (USA), and R&D systems (USA).

	<b>Antibody</b>	<b>Dilution</b>	<b>Manufacturer / Catalog number</b>
<b>Secondary Antibodies</b>	Alexa Fluor 488	1:200	Life Technologies / A21200
	Alexa Fluor 594	1:200	Life Technologies / A21442
	Alexa Fluor 594	1:200	Life Technologies / A21442
	Alexa Fluor 488	1:200	Life Technologies / A11055
<b>Primary Antibodies</b>	Sox2	1:100	Santa Cruz / SC17320
	Ki67	1:200	Invitrogen / PA5-19462
	KRT14	1:500	Abcam / 181595
	AQP5	1:100	Abcam / AB92320
	Cleaved Caspase 3	1:200	Cell Signaling Technologies / 9664S
	β-3 tubulin (Tuj-1)	1:100	R&D systems / MAB1195

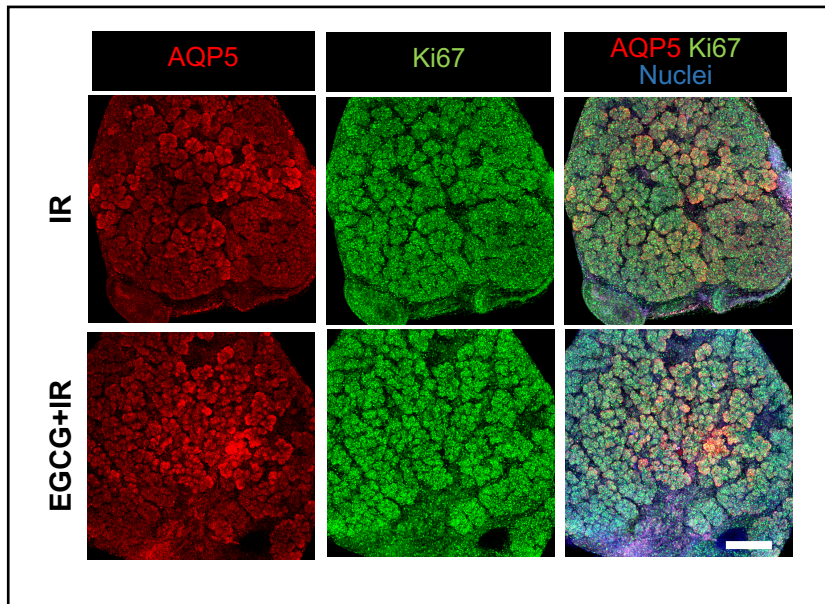
Table S2. List of oligonucleotide primer sequences

Gene	Forward sequence	Reverse sequence
<i>Ki67</i>	CATACCTGAGCCCATCACCA	GCTTTGCTGCATTCCGAGTA
<i>Sox 2</i>	CAGCATGTCCTACTCGCAGCAG	TGGAGTGGGAGGAAGAGGTAACC
<i>Sox10</i>	ATCAGCCACGAGGTAATGTCCAAC	ACTGCCCAGCCCGTAGCC
<i>Krt14</i>	CAGCCCCTACTTCAAGACCA	GTCGATCTGCAGGAGGACAT
<i>Aqp5</i>	TCTACTTCTACTTGCTTTTCCCCTCCTC	CGATGGTCTTCTTCCGCTCCTCTC
<i>Krt5</i>	TCCTGTTGAACGCCGCTGAC	CGGAAGGACACACTGGACTGG
<i>Acta2</i>	GGAGAAGCCCAGCCAGTCGC	AGCCGGCCTTACAGAGCCCA
<i>Krt19</i>	CCTCCCGAGATTACAACCACT	GGCGAGCATTGTCAATCTGT
<i>Mist1</i>	GCTGACCGCCACCATACTTAC	TGTGTAGAGTAGCGTTGCAGG
<i>Nkcc1</i>	TTCCGCGTGAAGTTCGTGG	TTGGTGTGGGTGTCATAGTAGT
<i>Pecam1</i>	TCCAACAGAGCCAGCAGTATGAGG	TCCAATGACAACCACCGCAATGAG
<i>Tubb3</i>	CCAGAGCCATCTAGCTACTGACACTG	AGAGCCAAGTGGACTCACATGGAG
<i>Rsp29</i>	GGAGTCACCCACGGAAGTTCGG	GGAAGCACTGGCGGCACATG

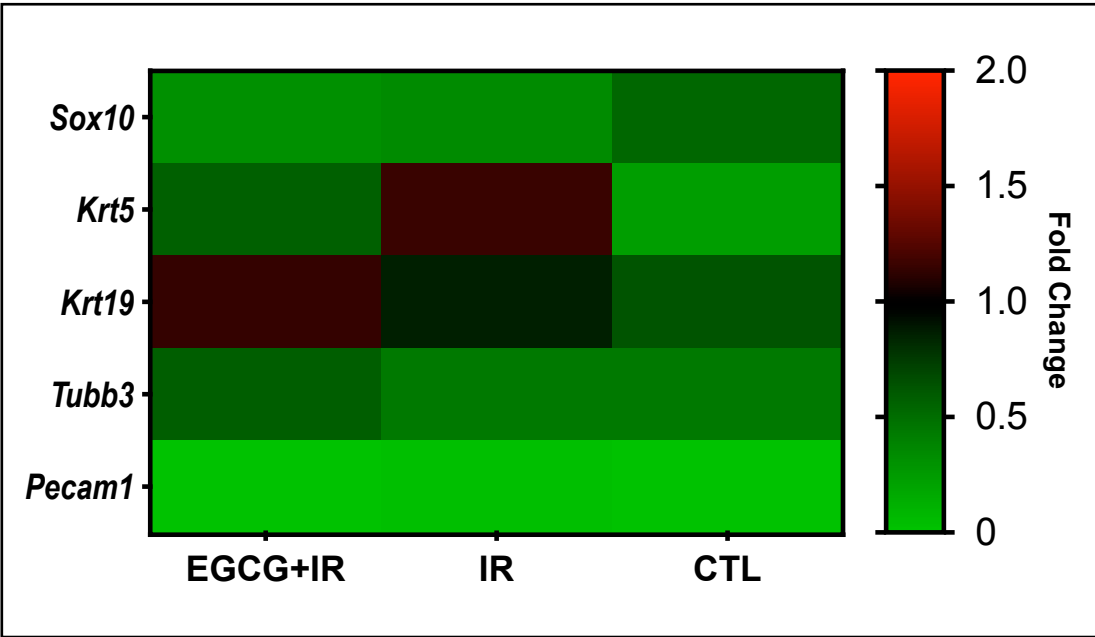


**Figure S1. Expression of epithelial progenitors in EGCG-treated glands after IR injury.** Expression of cytokeratin 14 (KRT14) progenitor markers in pro-acinar endbud compartments (A) and in ductal compartments (B) after immunofluorescence staining. Images shown are maximum intensity projections. Mag.: 20X. Scale bar: 100μm.

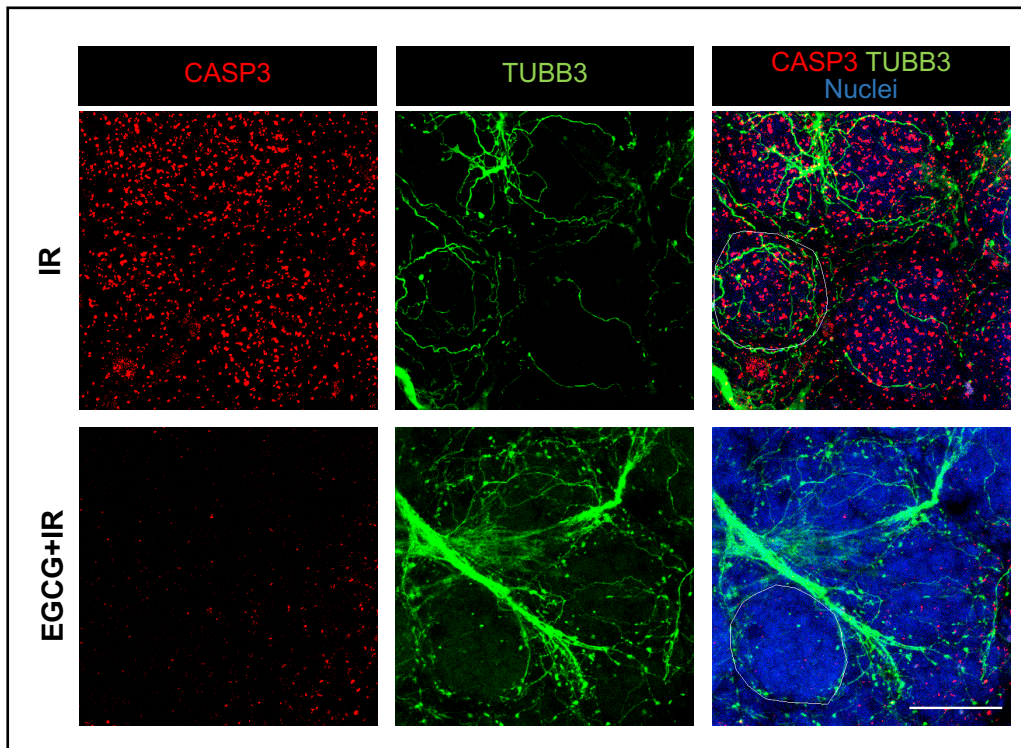




**Figure S2.** Expression of differentiated acinar epithelial and mitotic markers in EGCG pre-treated glands after IR injury. SG were immuno-stained with Aquaporin 5 (AQP5), a mitotic marker (Ki67) and incubated with a nuclear stain. Mag.: 10X. Scale bar: 200 $\mu$ m.



**Figure S3.** Heatmap with mean expression of other stem/progenitor, ductal epithelial, neuronal and vascular markers in the whole gland by qPCR. Data are presented as mean ( $n = 3$ ) of fold change relative to house keeping gene normalized to baseline from  $n = 3$ . Welch's Student *t*-test were performed between untreated and treated but not significant difference was observed. CTL represent non-irradiated untreated controls.



**Figure S4.** Expression of pro-apoptotic Caspase 3 marker in EGCG pre-treated glands after IR injury. SG were immuno-stained with cleaved-caspase 3 (CASP3),  $\beta$ -3 tubulin (TUBB3) to depict the boundaries of acinar buds where terminal neurons synapse. SG were also incubated with a nuclear stain. Mag.: 40X. Scale bar: 100 $\mu$ m.