Table S1. Eggs with and without detectable tumour at day 14 embryonation in Chick embryo after tumour cell implantation

	Eggs with tumour /Total eggs	Eggs without tumour /Total eggs	p value
SW620shCon	24/42	18/42	
SW620shC1GT	8/42	34/42	0.001
SW620shCon+Gal3	8/24	16/24	
SW620shC1GT+Gal3	3/24	21/24	0.011

Table S2. Cell membrane proteins extracted by galectin-3 and MGL from both SW620shcon and SW620shC1GalT1 cells

Cell membrane proteins extracted by galectin-3 from both SW620shCon and SW620shC1GT cells	Cell membrane proteins extracted by MGL from both SW620shCon and SW620shC1GT cells	
14-3-3 protein	Annexin A1	
Annexin A1	Annexin A2	
Annexin A2	Annexin A3	
Annexin A3	CD98 antigen	
CD98 antigen	Clathrin heavy chain 1	
Ep-CAM	Complement component 1 Q subcomponent-binding protein	
Galectin-1	Dnj3	
Galectin-3-binding protein	eIF-2gX	
HMG-2	Extended synaptotagmin-1	
HMGA1	FAU ubiquitin-like and ribosomal protein S30	
LAMP-1	G25K GTP-binding protein	
LAMP-2	Galectin-1	
LCP-1	Galectin-3	
MCT 4	Galectin-3-binding protein p105	
Membrane-associated progesterone receptor component 1	Magnesium transporter protein 1	
Moesin	MCT 1	
p27RF-Rho	MCT 4	
p59-Fyn	Moesin	
Prohibitin-2	Multifunctional protein ADE2	
Protein S100-A10	MYH-1c	
Protein S100-A6	Parkinson disease protein 7	
Receptor of activated protein C kinase 1	Plastin-2	
SLC12A2	Protein S100-A4	
Spectrin alpha chain, non-erythrocytic 1	Receptor of activated protein C kinase 1	
Spectrin beta chain, non-erythrocytic 1	ZO-1	
Src substrate cortactin		
Vimentin		

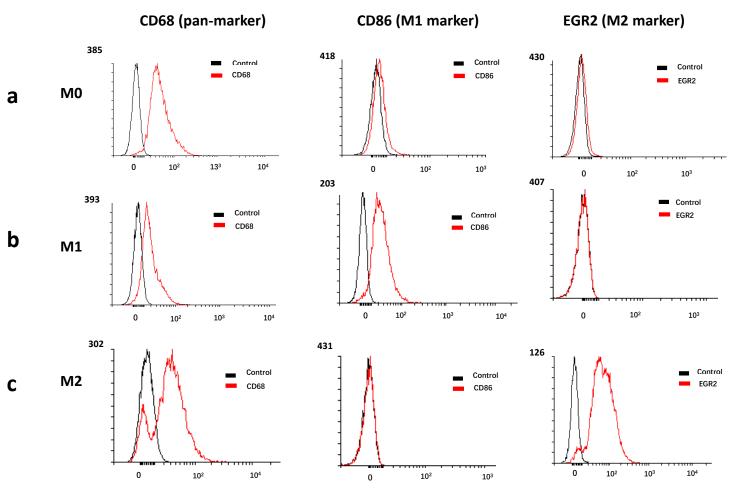
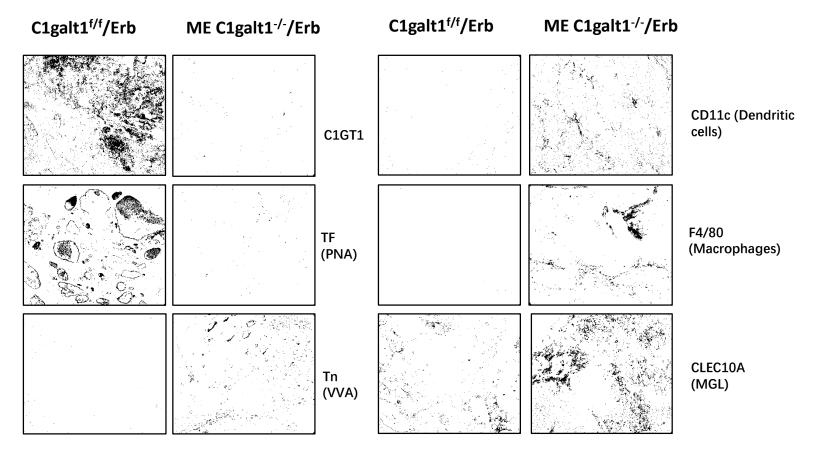


Figure S1. Confirmation of THP-1 macrophage differentiation. THP-1 cells were incubated with 10 ng/ml PMA for 48 h at 37°C. The cells were released with NECDS, washed once with PBS and fixed with 2% formaldehyde for 30 min. After two washes with PBS, the cells (5x10<sup>5</sup> cells/ml) were incubated with 1% BSA/PBS for 20 min before incubation with PE-conjugated antibodies against pan-macrophage marker CD68, M1 macrophage marker CD86 and M2 macrophage marker EGR2 and analysed by flow cytometry (a). THP-1 differentiated M0 macrophages were treated with 50 ng/ml IFN-γ and 100 pg/ml LPS for 24 h before analysed by flow cytometry for expression of macrophage markers (b). THP-1 differentiated M0 macrophages were treated with 60 ng/ml IL-4 for 72 h before analysed by flow cytometry for expression of macrophage markers (c). Representative flow cytometry histograms from three independent analysis are shown.



**Figure S2. Binary images in evaluation of macrophage and DC tumour infiltration.** Immunohistochemistry images of the mice sections were quantified using Fiji Image J. The final binary images of Fig 7 c (left panel) and d (right panel) were shown as example here.