

SUPPLEMENTARY MATERIAL

Table S1. Primers used in real-time PCR analyses.

Gene	Forward primer 5'-3'	Reverse primer 5'-3'
<i>Actb</i>	ATTACTGCTCTGGCTCCTA	ATCTGCTGGAAGGTGGAC
<i>Ccr2</i>	AGAGAGCTGCAGCAAAAGG	GGAAAGAGGCAGTTGCAAAG
<i>Cxcl1</i>	CCGCTCGCTTCTCTGTGC	CTCTGGATGTTCTTGAGGTGAATC
<i>Cyp1a1</i>	ACAGTGATTGGCAGAGATCG	GAAGGGGACGAAGGATGAAT
<i>Cyp1b1</i>	TTCTCCAGCTTTTTCCTGT	TAATGAAGCCGTCCTTGTC
<i>Gapdh</i>	GCCTTCCGTGTTCTACCC	CAGTGGGCCCTCAGATGC
<i>Ido1</i>	CCCACACTGAGCACGGACGG	GCCCTTGTCGCAGTCCCCAC
<i>Il1b</i>	TGA CGG ACC CCA AAA GAT GAA GG	CCA CGG GAA AGA CAC AGG TAG C
<i>Il1ra</i>	TTGTGCCAAGTCTGGAGATG	CAGCTGACTCAAAGCTGGTG
<i>Il6</i>	CCGGAGAGGAGACTTCACAG	TCCACGATTTCAGAGAAC
<i>Il10</i>	GAGAAGCATGGCCCAGAAATCAAG	ATCACTCTTCACCTGCTCCACTGC
<i>Il17a</i>	GACTACCTCAACCGTCCAC	CCTCCGATTGACACAGC
<i>Il22</i>	CTGCCTGCTTCTCATTGCCCTGTG	GATGTACGGCTGCTGGAAGTTGG
<i>Mmp2</i>	AACGGTCGGAATACAGCAG	ATGGGTGGATCTTCATGGGG
<i>Mmp9</i>	TAGCACAACAGCTGACTACG	ATCCTGGTCATAGTTGGCTG
<i>S100a8</i>	TCGTGACAATGCCGTCTGAACTG	TGCTACTCCTTGTGGCTGTCTTTG
<i>S100a9</i>	CGCAGCATAACCACCATCATC	GCCATCAGCATCATACACTCC
<i>Snai1</i>	CTTGTGTCTGCACGACCTGT	CATCCGAGTGGGTTTGGAGG
<i>Twist1</i>	CTGCCCTCGGACAAGCTGAG	CTAGTGGGACGCGGACATGG
<i>Vegfa</i>	TATTCAGCGGACTCACCAGC	AACCAACCTCCTCAAACCGT
<i>Vegfb</i>	TGACGATGGCCTGGAATGTG	GAGGATCCTGGGGCTGTCT
<i>Vegfc</i>	GCTGATGTCTGTCTGTACCC	AGAAGGTGTTGTGGCTGCT
<i>Vegfr2</i>	TCCACATGGGCGAATCACTC	GCAATTCTGTCACCCAGGGA
<i>Zeb1</i>	GGAGAGGTGACTGGTTGTGG	GCCACATCAGCAATAGCAGC

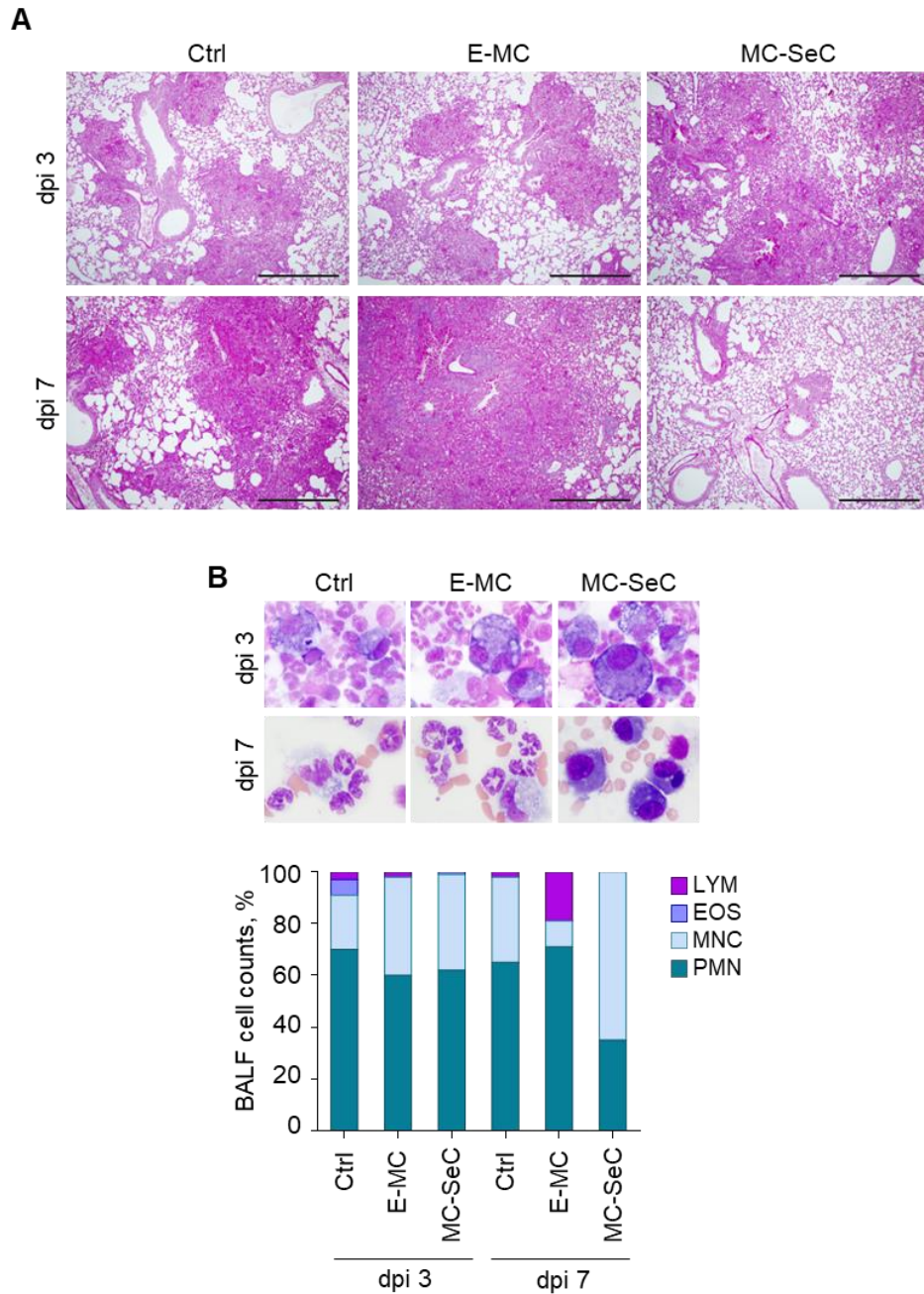


Figure S1. Grafted SeC protect from *A. fumigatus* intratracheal infection. (A,B) Control mice (Ctrl) and mice injected i.p. with MC-SeC (1.0×10^6 SeC/g body weight) or equivalent amount of E-MC were infected intratracheally with *A. fumigatus* conidia. Periodic acid-Schiff staining of lungs (A), and BALF morphometry analysis (B) were performed at 3 and 7 days post-infection (dpi). Reported are BALF differential cell counts. Scale bars (A), 1 mm; original magnification (B), 100X.

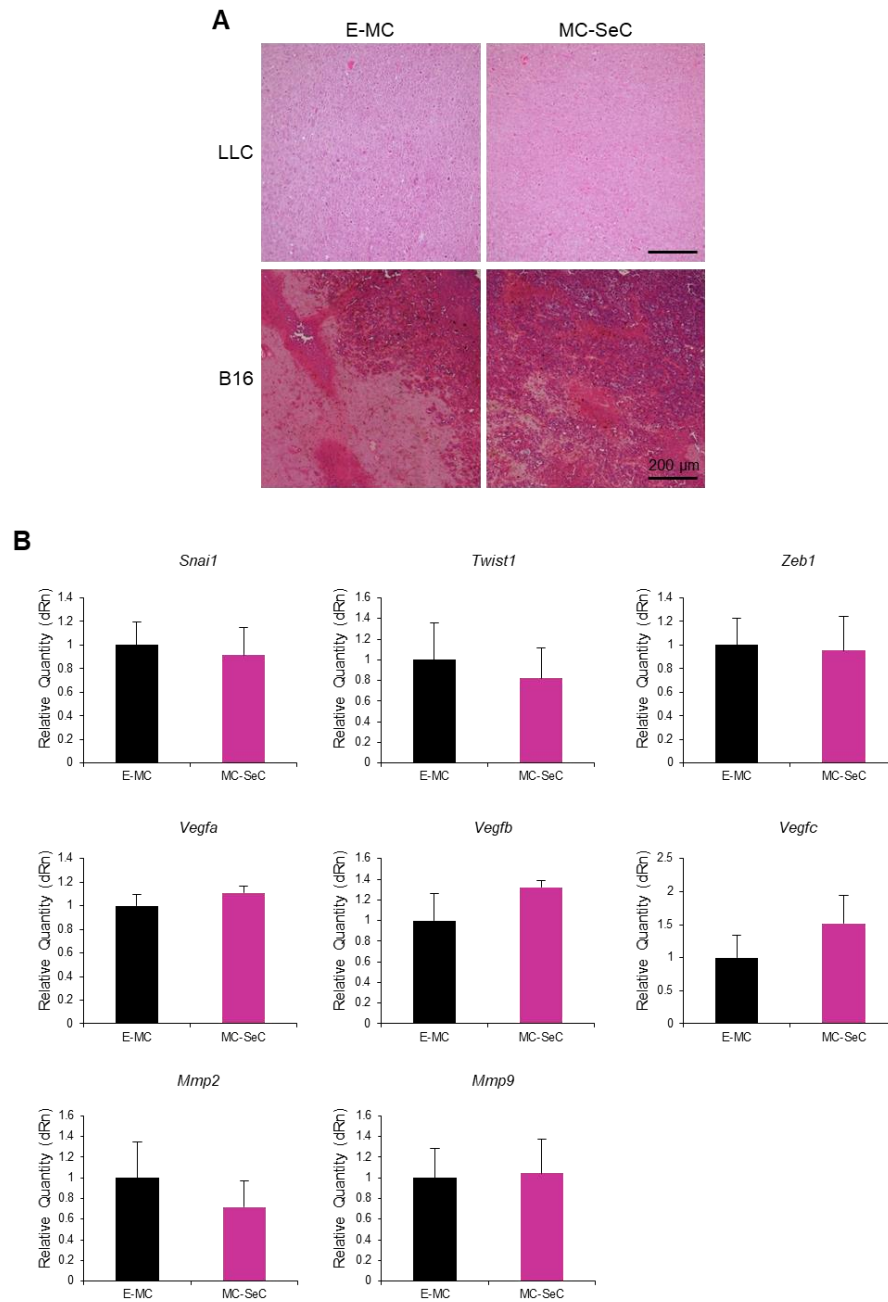


Figure S2. Grafted SeC do not affect primary tumor histology. (A) Primary LLC and B16 tumor masses developed in mice injected i.p. with MC-SeC (1.0×10^6 SeC/g body weight) or equivalent amount of E-MC were histologically evaluated after hematoxylin-eosin staining. Shown are representative images. Scale bars, 200 μ m. **(B)** LLC tumor masses were evaluated for the expression of epithelial-mesenchymal transition markers (*Snai1*, *Twist1*, and *Zeb1*), matrix metalloproteases (*Mmp2* and *Mmp9*), and angiogenesis markers (*Vegfa*, *Vegfb*, and *Vegfc*) by real-time PCR.