Figure S1. Increased CD11b cells in  $LT\beta^{-/-}$  mouse tracheas following infection with *M*. *pulmonis*. LYVE-1<sup>+</sup> positive lymphatic vessels (red) and CD11b<sup>+</sup> positive infiltrates (green) in the tracheas of mice infected with *M. pulmonis*. Scale bar is 200 µm.

**Figure S2. Absence of increased lymphangiogenesis in TNFR1**<sup>-/-</sup> **mouse skin following the induction of inflammation.** Fluorescent microscopy of the site of immunization with ovalbumin and CFA after nanoparticle injection reveals no lymphatic vessel networks in proximity to the immunization depot site in TNFR1<sup>-/-</sup> mice. Black blood vessels are apparent and even more obvious after immunization. Lymphatic vessels would be bright green (See Fig. 4). Representative images of 2 or 3 mice per time point from two separate experiments (13 mice total).

**Figure S3.** LTα , LTβ, and TNFα mRNAs at the site of immunization in the skin at day 7. WT mice express LTα, LTβ and TNFα mRNA in the skin, spleen and peripheral lymph node (PLN). LTα<sup>-/-</sup> mice had no detectable LTα mRNA in the skin or spleen as expected, but demonstrated LTβ and TNFα mRNA expression in these organs. LTβ<sup>-/-</sup> mice expressed TNFα and LTα mRNA in the skin and mesenteric lymph node (MLN). LTα mRNA is 485 bp; partially spliced is 709 bp; LTβ mRNA is 640bp; TNF mRNA is 700bp.

# Figure Supplement 1

# Pathogen-free Infected WT

### LTα-/-



# Figure Supplement 2.



#### TNFR1-/-

day 7

## day 14

# Figure Supplement 3



# - 650 bp - 500 bp - 400 bp

- 850 bp