



Figure S3. pLe^X-ω1 increases adipose tissue type 2 immune cells and improves whole-body insulin sensitivity in a dose-dependent manner. Mice were fed a LFD (white bars) or a HFD for 12 weeks, and next received biweekly intraperitoneal injections of PBS (black bars) or 10 μg, 25 μg or 50 μg pLe^X-ω1 (green bars) during 4 weeks (A). At the end of the experiment, eWAT was collected, processed and analyzed as described in the legend of Figure 1. The numbers of CD4 T cells and ILCs per gram tissue (B), and the frequencies of IL-13+ CD4 T cells (C) and ILCs (D) were determined. Numbers of eosinophils (E) and macrophages (F) per gram tissue, and percentages of CD11c⁺YM1⁻ and CD11c⁻YM1⁺ macrophages (G) were determined. Body weight (H-I) and body composition (J) were determined after 4 weeks. Food intake was monitored throughout the treatment period (K). Fasting blood glucose (L) and plasma insulin levels (M) were determined at week 4, and HOMA-IR (N) was calculated. An i.p. insulin tolerance test (O-P) was performed at week 3. Results are expressed as means ± SEM. * $P < 0.05$ vs HFD or as indicated (n = 3-4 mice per group).