SUPPLEMENTARY DATA

Supplementary Figure 1. Metabolic phenotype of male and female STAT4^{-/-}C57Bl6 mice and wild type controls on chow and high fat diets. Intraperitoneal glucose and insulin tolerance tests GTT and ITT) were performed on age-matched male and female mice after 16 weeks of high fat or chow diets. Data is expressed as area under curve (AUC) and represents average±SEM from n=6-8 females and n=6 males.



GTT males vs. females on chow diet

GTT males vs. females on high fat diet



ITT males vs. females on chow diet

ITT males vs. females on high fat diet



Figure justification: This supplemental information is important as some of the genetic alterations leading to improvement in insulin sensitivity may have a gender-specific bias. In this Figure we show that the improvement in insulin sensitivity was equally present in both the STAT4 deficient males and females compared to wild-type sex and age-matched controls.

SUPPLEMENTARY DATA

Supplementary Figure 2. Activation of insulin signaling pathway in isolated adipocytes Adipoctes separated by collagenase digestion from $STAT4^{-/-}$ and wildtype mice were treated in vitro with 5nM insulin for 10 min (n=3-7); activation of insulin receptor (IR) and protein kinase B (Akt) were measured by western blotting. Results are expressed as ratio of activated/basal phopho/total IR and Akt, respectively. Bars represent mean±SEM.



Figure justification: This supplemental information is important as it emphasizes that the improvement of insulin signaling in the STAT4-/- in vivo is a combination of STAT4 deficienty in adipocytes and the local pro-inflammatory milieu in adipose tissue.