SUPPLEMENTARY DATA

Supplementary Table 1. % Donor- and host-type blood mononuclear cells in mixed chimeric NOD at different time points after MHC-mismatched HCT (Mean \pm SE, N=4)

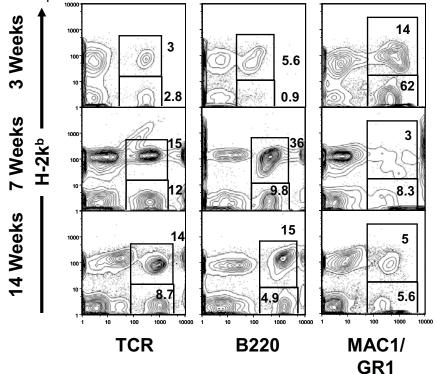
		TCRβ ⁺	B220 ⁺	MAC1/Gr1 ⁺
3 Weeks	Donor- type	26.6 ±9.1	599±12.1	12.7 ±3.0
	Host-type	73.4 ± 9.1	40.1 ±12.1	87.3 ± 3.0
7 Weeks	Donor- type	61.3 ±7.7	60.6 ± 17.6	22.3 ±7.3
	Host-type	38.7 ± 7.7	394 ± 17.6	77.7 ± 7.3
14 Weeks	Donor- type	594 ± 16.3	55.2 ± 16.8	23.8 ± 6.6
	Host-type	40.6 ± 16.3	44.8 ± 16.8	76.2 ± 6.6

NOTE: Donor-type cells were defined as H-2k^{b+} and host-type cells were defined as H-2k^{b-}; no significant difference was found with percentage of donor- or host-type cells between 7 and 14 weeks after HCT.

Supplementary Table 2. % $H-2k^{b+}$ or $H-2k^{b-}$ T, B, Macrophages/granulocyte, and dendritic cells in spleen of C57BL/6 (Mean \pm SE, N=4)

		$TCR\beta^+$	B220 ⁺	MAC1/Gr1 ⁺	CD11c ⁺
C57BL/6	H-2k ^{b+}	100 ±0	920 ±0.2	92.0 ±1.8	98.7 ±0.5
	H-2k ^{b-}	0 ±0	1.0 ± 0.2	8.0 ± 1.8	1.3 ± 0.5

Supplementary Figure 1. Mixed chimerism in NOD recipients conditioned with radiation free anti-CD3 based regimen. WT NOD mice were conditioned with anti-CD3/CD8 ($5\mu g$ each) on days -10 and -5. On day 0, the conditioned mice were transplanted with graded doses ($5-20x10^6$ each) of CD4⁺ T cell-depleted splenocytes and whole BM from WT C57BL/6 donors to induce chimerism. Chimerism was judged by analysis of donor- and host-type peripheral blood mononuclear cells at 3, 7, and 14 weeks after HCT. One representative of 4 examined mice is shown.

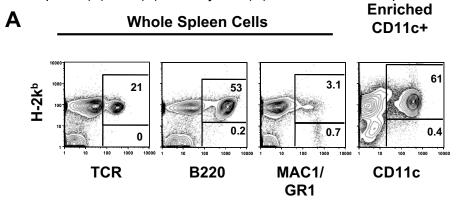


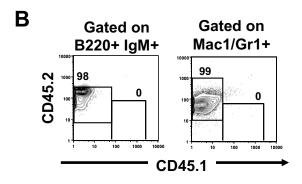
Supplementary Figure 2. Representative insulitis levels from Conditioning Alone mice, ~70 days after first round of conditioning. Brown – Insulin.

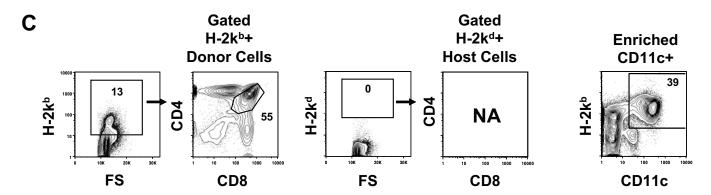
Insulitis

40x

Supplementary Figure 3. Donor-type control staining. Spleen, BM, and thymus cells of WT C57BL/6 donors were stained with anti-H-2k^b vs anti-TCR β , B220, MAC-1/Gr-1 or CD11c. Expression of H-2k^b by TCR β ⁺ T cells, B220⁺ B cells, MAC-1/Gr-1⁺ macrophages/granulocytes, and CD11c⁺ DCs from spleen (A), BM (B), and thymus (C) are shown. One representative of 4 examined mice is shown.







Supplementary Figure 4 - No expansion of T cells expressing transgenic TCR V β 4 coupled with endogenous V α s in mixed chimeric BDC2.5-NOD recipients. Transgenic BDC2.5 NOD were conditioned with anti-CD3 (5 μ g) on day -7. On day 0, the conditioned mice were transplanted with BM cells (50x10⁶) from C57BL/6 donors. 60 days after HCT, peripheral blood mononuclear cells were analyzed for V β 4⁺ CD4⁺ T cell expression of endogenous V α 2, V α 8.3 and V α 11. WT-NOD and BDC2.5-NOD without HCT were used as controls. Gated CD45.2⁻ V β 4⁺CD4⁺ or V β 4⁻CD4⁺ host-type T cells are shown in V β 4 vs V α 2, V α 8.3, V α 11. One representative staining pattern of 4 examined mice is shown. Mean±SE of V β 4⁺V α + or V β 4⁻V α + cells amongst gated CD4⁺T cells is shown.

