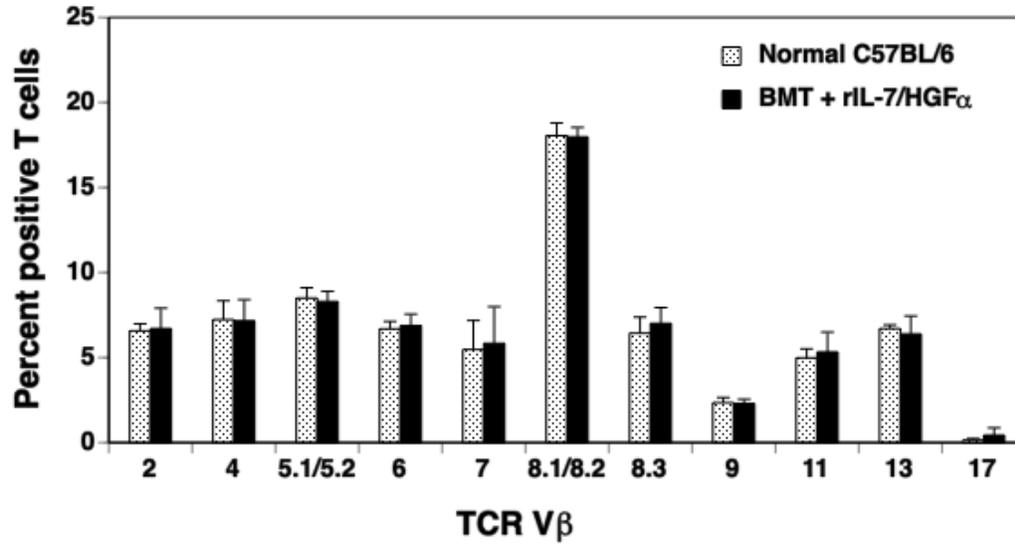


Supplemental Figure 1:

hrIL-7/HGF α -treated mice maintained increased numbers of total thymocytes and their subsets through day 90 post-BMT. Lethally irradiated BMT mice were treated with a panel of cytokines as in Figure 2C. Ninety days after BMT, the numbers of total thymocytes and their subsets were quantified by flow cytometry. Data shown are mean \pm SD from 4-5 mice per group. * P<0.05 compared with PBS-treated BMT mice. ** P<0.05 compared with hrIL-7+hrHGF α -treated BMT mice.



Supplemental Figure 2: Donor-origin peripheral T cells from hrIL-7/HGF α -treated mice have a diverse TCR repertoire. Lethally irradiated BMT mice were treated with hrIL-7/HGF α as in Figure 2C. One month after BMT, the expression of TCR V β families by donor-derived CD3⁺ cells in the peripheral blood was analyzed by flow cytometry. The results were compared with those of CD3⁺ cells from normal control mice. Mean percentages \pm SD from groups of 4-6 mice. No significant differences were observed between the experimental and control values for each TCR V β .

Supplemental Table 1. Primers used for cloning of human IL-7/HGF α (NK1), IL-7 and HGF α (NK1) genes into expression vectors

Primers	Primer Sequence (5'-3')
A	GTCAACATGTTCCATGTTTCT
B	CGACCCACCACCGCCCGAGCCACCGCCTCCGTGTTCTTTAGTGCCCAT
C	GGAGGCGGTGGCTCGGGCGGTGGTGGGTCGCAAAGGAAAAGAAGAAT
D	GCCTCACTATTCAACTTCTGAACACTG
E	GCCCTATCAGTGTTCTTTAGTGCCCAT
F	GCGAAC ATGTGGGTGACCAAACCTC
G	GCCTCACTAATGGTGATGGTGATGATGTTCAACTTCTGAACACTG