

Human thymus contains IFN- α -producing CD11c⁻, myeloid CD11c⁺, and mature interdigitating dendritic cells

Nathalie Bendriss-Vermare, ... , Giorgio Trinchieri, Francine Brière

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Corrigendum

J. Clin. Invest. 107:835–844 (2001) During the preparation of this manuscript for publication, errors were introduced in Figure 3. The correct version, accompanied by the legend, appears below.³ Figure 3 Immunophenotype of isolated thymic DC subsets analyzed by flow cytometry. Thymic DCs were sorted into Lin⁻ (PE-Cy5) HLA-DR^{int} (FITC) and Lin⁻ HLA-DR^{hi} subsets. Anti-CD13-PE-Cy5 labeling of HLA-DR^{int} cells clearly resolved two distinct populations. CD13⁺ HLA-DR^{int}, CD13⁻ HLA-DR^{int}, and CD13⁺ HLA-DR^{hi} DCs were analyzed using PE-conjugated mAb's for the expression of a number of lymphoid, myeloid, costimulatory, and adhesion markers. Data shown are representative of three experiments. Ag, antigen.

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Nathalie Bendriss-Vermare, Clarisse Barthélémy, Isabelle Durand, Corine Bruand, Colette Dezutter-Dambuyant, Nathalie Moulian, Sonia Berrih-Aknin, Christophe Caux, Giorgio Trinchieri, and Francine Brière

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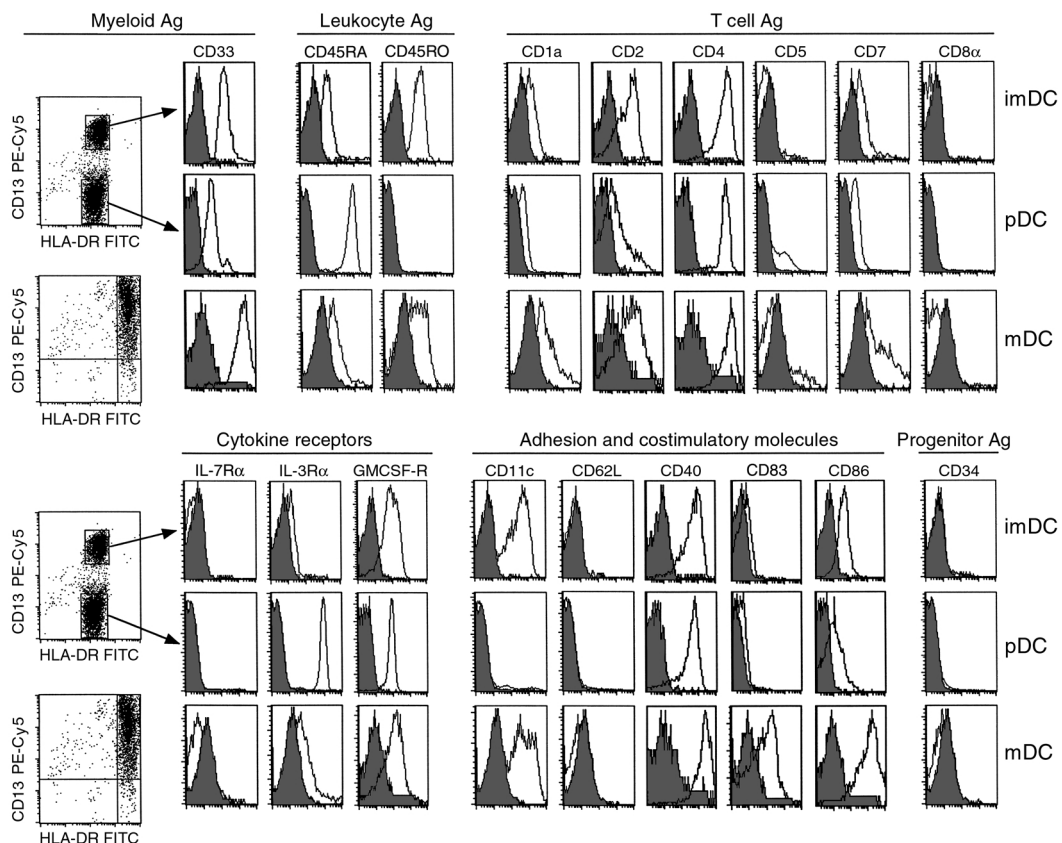


Figure 3

Immunophenotype of isolated thymic DC subsets analyzed by flow cytometry. Thymic DCs were sorted into Lin⁻ (PE-Cy5) HLA-DR^{int} (FITC) and Lin⁻ HLA-DR^{hi} subsets. Anti-CD13-PE-Cy5 labeling of HLA-DR^{int} cells clearly resolved two distinct populations. CD13⁺ HLA-DR^{int}, CD13⁻ HLA-DR^{int}, and CD13⁺ HLA-DR^{hi} DCs were analyzed using PE-conjugated mAb's for the expression of a number of lymphoid, myeloid, costimulatory, and adhesion markers. Data shown are representative of three experiments. Ag, antigen.