

## Analysis of the adult thymus in reconstitution of T lymphocytes in HIV-1 infection

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### Erratum

*J. Clin. Invest.* 103:453–460 (1999). A production error resulted in the incorrect printing of Figure 1; the correct display appears below. We regret the error and have provided corrected reprints to the corresponding author: Barton F. Haynes, Box 3703, Duke Hospital, Durham, North Carolina 27710, USA. Phone: (919) 684-5384; Fax: (919) 681-8992; E-mail: Hayne002@mc.duke.edu

Figure 1 Immunohistological analysis of the thymus in HIV infection. (a–d) Thymus from HIV-1+ patient no. 1 with no thymopoiesis. (e–h) Thymus from HIV-1+ patient no. 2 with areas of active thymopoiesis. (a) Hematoxylin and eosin stain of patient no. 1's lymphoid thymus. ×13. (b) A similar area as in a, with thymic epithelium in immunohistological analysis reactive with antikeratin antibody (brown central areas). All keratin+ thymic epithelium (e) in the true thymus is collapsed (dark brown areas) and devoid of lymphocytes, with a surrounding infiltrate of blue mononuclear cells present in the thymic perivascular space (P). ×13. (c) Immunohistological stain of CD8+ T cells (brown cells; see arrows for examples) in the perivascular space (P) around a central empty thymic epithelial island (e). The dotted line surrounds thymic true epithelial thymus areas (e), and the short arrow points out a rare CD8+ T cell within the true epithelial thymus (e). ×66. (d) Many of the perivascular space (P) CD8+ cells are reactive with MAB TIA-1 (arrows) and [...]

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**Figure 1**

Immunohistological analysis of the thymus in HIV infection. (a–d) Thymus from HIV-1+ patient no. 1 with no thymopoiesis. (e–h) Thymus from HIV-1+ patient no. 2 with areas of active thymopoiesis. (a) Hematoxylin and eosin stain of patient no. 1's lymphoid thymus.  $\times 13$ . (b) A similar area as in a, with thymic epithelium in immunohistological analysis reactive with antikeratin antibody (brown central areas). All keratin<sup>+</sup> thymic epithelium (e) in the true thymus is collapsed (dark brown areas) and devoid of lymphocytes, with a surrounding infiltrate of blue mononuclear cells present in the thymic perivascular space (P).  $\times 13$ . (c) Immunohistological stain of CD8<sup>+</sup> T cells (brown cells; see arrows for examples) in the perivascular space (P) around a central empty thymic epithelial island (e). The dotted line surrounds thymic true epithelial thymus areas (e), and the short arrow points out a rare CD8<sup>+</sup> T cell within the true epithelial thymus (e).  $\times 66$ . (d) Many of the perivascular space (P) CD8<sup>+</sup> cells are reactive with MAB TIA-1 (arrows) and therefore are mature effector cytotoxic T cells.  $\times 66$ . (e–h) are from patient no. 2's thymus.  $\times 33$ . (e) Light microscopic view of patient no. 2's thymus (hematoxylin and eosin stain with a Hassall's body [h] in the thymus medulla). (f) Immunohistological analysis with antikeratin antibody, with areas of normal-appearing keratin<sup>+</sup> thymic epithelium (brown areas) filled with lymphocytes (blue areas) intermingled with thymic epithelium (arrows). Most developing thymocytes are CD3<sup>+</sup> T cells (arrows in g), many of which are normal CD1a<sup>+</sup> cortical thymocytes (brown cells, arrows in h). A subset of these CD1a<sup>+</sup>, CD3<sup>+</sup> immature thymocytes were actively dividing as determined by nuclear reactivity with MAB mib-1 (not shown). MAB, monoclonal antibody.

