Thymic Involution in Viable Motheaten (*me*<sup>+</sup>) Mice is Associated with a Loss of Intrathymic Precursor Activity

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Mice homozygous for the viable motheaten (*me*) allele manifest abnormalities in thymocytopoiesis, are severely immunodeficient, and develop autoimmune disorders early in life. Premature thymic involution occurs in *me/me*<sup>-</sup> mice, and their bone marrow prothymocytes are unable to repopulate the thymus of adoptive recipients following intravenous (i.v.) transfer. However, analysis of thymocytopoiesis following intrathymic (i.t.) adoptive transfer of bone marrow from *me/me*<sup>-</sup> mice demonstrates the presence of normal numbers of prothymocytes. To investigate intrathymic development in *me/me*<sup>-</sup> mice, we determined intrathymic precursor cell number and activity. Dual labeling analyses showed that an involuted *me/me*<sup>-</sup> thymus is relatively enriched (fivefold) in CD<sup>4</sup>CD<sup>8</sup>- thymocytes (intrathymic precursor phenotype) compared with wild-type (+/+) thymus. However, thymocytes from *me/me*<sup>-</sup> mice were deficient in precursor activity when adoptively transferred i.t. into irradiated recipients. Thymocytes recovered from the involuted thymus of aged or steroid-treated normal mice also displayed reduced precursor activity. However, the phenotypic profile of thymocyte subsets from steroid-treated mice was enriched in single positive cells (mature phenotype) and was distinctly different from the subset distribution of thymocytes in *me/me*<sup>-</sup> and aged mice. These results suggest that intrathymic precursor activity in *me/me*<sup>-</sup> mice is decreased, and may be reflective of decreased prothymocyte seeding to the thymus in *vivo*. In addition, the results suggest that the thymic involution in *me/me*<sup>-</sup> mice is not due solely to effects of corticosteroids.

KEYWORDS: Viable motheaten mice, thymocytopoiesis, involution, aging, steroids, thymus.

INTRODUCTION

Mice homozygous for the motheaten (*me*) or viable motheaten (*me*<sup>+</sup>) alleles display numerous defects in lymphopoiesis. These defects include a paucity of surface immunoglobulin-positive (Ig<sup>+</sup>) B cells and abnormal peripheral T-cell functions (Green and Shultz, 1975; Shultz et al., 1984; Greiner et al., 1986; Shultz and Sidman, 1987; Shultz, 1988). Furthermore, a number of autoimmune disorders develop that include hyper-gamma-globulinemia, autoantibody production, and immune complex glomerulonephritis (Kincade, 1981; Shultz and Sidman, 1987; Shultz, 1988). The usual cause of death is a characteristic hemorrhagic macrophagic pneumonitis that is believed to be autoimmune in origin (Shultz and Sidman, 1987; Shultz, 1988). This autoimmune syndrome can be transferred to irradiated recipients using *me/me*<sup>-</sup> bone marrow stem cells (Kincade, 1981; Shultz, 1988).

Developmental abnormalities are evident in cells that express the enzyme terminal deoxynucleotidyl transferase (TdT; presumptive T- and B-lymphoid stem cells) in *me/me* and *me/me*<sup>-</sup> bone marrow and thymus (Landreth et al., 1981; Greiner et al., 1986). These bone marrow and thymus TdT<sup>-</sup>-cell developmental defects appear to be associated with the defects in thymocytopoiesis and may have a causal relationship to the peripheral T- and B-lymphocyte abnormalities. Moreover, at least some of the TdT<sup>-</sup>- (Landreth et al., 1981; Greiner et al., 1986) and B-cell progenitor cell (McCoy et al., 1985; Greiner et al., 1986; Kincade, 1987) developmental abnormalities,
when analyzed in selective in vitro assay systems (Medlock et al., 1987; Hayashi et al., 1988), appear to be secondary to bone marrow micro-environmental defects.

Studies on thymopoiesis in me/me and me'/me" mice have demonstrated a premature thymic involution, commencing as early as 4 weeks of age (Greiner et al., 1986; Shultz and Sidman 1987; Shultz, 1988). Furthermore, we have shown that both me/me and me'/me" bone marrow prothymocytes fail to repopulate the thymus of irradiated recipients after intravenous (i.v.) transfer, but generate normal numbers of thymocytes following intrathymic (i.t.) adoptive transfer (Greiner et al., 1986). Mixing of me'/me" bone marrow with normal bone marrow restores the ability of me'/me" prothymocytes to generate thymocytes after i.v. injection (Komschlies et al., 1987), again suggesting that a marrow micro-environmental defect may underlie certain of the early developmental defects observed in lymphopoiesis in me/me and me'/me" mice.

The apparent prothymocyte homing defect in me/me and me'/me" bone marrow cells may have in vivo consequences, since postnatal maintenance of normal thymopoiesis is dependent upon a low, but continual, seeding of the thymus by bone marrow hemopoietic precursors (Scollay and Shortman, 1984; Scollay et al., 1986). Furthermore, the ability of bone marrow precursors to generate thymocytes following entry into the thymus appears to be self-limiting, and will cease within 4 to 5 weeks following thymic entry (Goldschneider et al., 1986; Scollay et al., 1988). Because me/me and me'/me" bone marrow prothymocytes appear to be unable to repopulate the thymus of irradiated recipients following i.v. transfer, we have postulated that this defect may result in an early loss of intrathymic precursors in me/me and me'/me" mice, leading to premature thymic involution (Greiner et al., 1986; Komschlies et al., 1987).

We have investigated this hypothesis in the present study using the i.t. adoptive transfer assay to quantitate directly the intrathymic precursor activity prior to and during thymic involution in me'/me" mice. Our results indicate that the pool of intrathymic precursors in me'/me" mice decreases early in life, in association with the onset of thymic involution. Furthermore, the thymic involution observed in me'/me" mice does not appear to be similar to that induced following administration of corticosteroids.

RESULTS

Phenotypic Characteristics of me'/me" Thymocytes During Thymic Involution

Investigations of murine thymocyte subpopulations have revealed a large diversity of cell subsets (Scollay et al. 1988; Wilson et al., 1988). Four major phenotypic populations have been extensively characterized, and are identified as CD4-CD8- (double negative, DN), CD4+CD8+ (double positive, DP), and CD4+CD8-(or CD4+CD8-) (single positive, SP). Because the thymus of me'/me" mice has been shown to undergo premature thymic involution, commencing as

| Genotype | Thymus cell number (<10^6) | Animal age (weeks) | Thymocyte subpopulations | CD4+CD8- (%) | CD4+CD8+ (%) | CD4+CD8+ (%) | CD4+CD8-(%) |
|----------|----------------------------|-------------------|--------------------------|--------------|-------------|-------------|-------------|
| me'/me"  | 1.6                        | 7                 |                           | 31.3         | 33.1        | 7.5         | 27.5        |
|          | 3.8                        | 6                 |                           | 15.8         | 48.7        | 15.8        | 19.7        |
|          | 9.0                        | 4                 |                           | 24.4         | 43.2        | 21.8        | 10.8        |
|          | 13.0                       | 4                 |                           | 14.0         | 47.8        | 28.3        | 9.9         |
|          | 58.0                       | 5                 |                           | 5.5          | 69.4        | 15.3        | 15.3        |
|          | 72.5                       | 4                 |                           | 11.6         | 65.0        | 15.7        | 7.6         |
|          | 135.0                      | 5                 |                           | 8.7          | 71.7        | 13.0        | 6.9         |
|          | 150.0                      | 4                 |                           | 4.0          | 88.7        | 5.0         | 2.2         |
|          | 270.0                      | 4                 |                           | 4.1          | 85.1        | 8.1         | 2.8         |
| Wild-type (+/+)| 285.0±37.7                 | 4                 |                           | 3.9±0.1      | 80.7±1.3    | 9.4±0.4     | 6.0±0.9     |

*Thymocytes from various aged C57BL/6j-me'/me" mice were analyzed for expression of CD4 and CD8 antigens using two-color immunofluorescence staining (see Materials and Methods). Data represent individual me'/me" thymuses except for the C57BL/6j+/+ data that represent the mean percentage±standard deviation for three thymuses from 4-week-old mice.
early as 4 weeks of age (Greiner et al., 1986; Shultz and Sidman, 1987; Shultz, 1988), we examined the phenotypic characteristics of the thymocytes prior to and during this involution process. As shown in Table 1 and Figure 1B, the loss of cellularity in the me/me thymus is evident as early as 4 weeks of age compared with untreated control B6+/+ mice. The loss of cellularity, variable between 4 and 7 weeks of age, is accompanied by a loss in the absolute numbers of DN, DP, and SP thymocytes compared with weanling B6 wild-type mice, with losses of 95%, 99.5%, and 98.4%, respectively, in the most severely involuted thymuses. The relative proportions of each unique thymocyte subset, however, were not as severely altered (Fig. 1B). There was a relative increase in the proportion of DN thymocytes in the most severely involuted thymus, approaching 30% of the total thymocyte population as compared to that of approximately 5% DN cells in the thymus of control B6+/+ mice. In the severely involuted thymuses (less than 13x10^6 total thymocytes), the DP and SP populations were approximately equal, each comprising 33-49% of the remaining cells present, whereas in normal B6+/+ mice, the DP and SP populations normally comprise approximately 80% and 15%, respectively, of a thymocyte population (Scollay et al., 1988). However, during the involution process, as thymus cellularity decreased, but prior to the most severely involuted state, the relative proportions of each

![Figure 1](image-url)
of the major thymocyte subsets approximated that observed in normal mice (Table 1).

**Intrathymic Precursor Activity of me^meme^m**

**Thymocytes During Thymic Involution**

The relative proportion of DN thymocytes was increased in the involuted me^meme^m thymus (Table 1 and Figure 1B), although the absolute number of DN thymocytes was decreased (Table 1). The increased percentage of DN thymocytes may be reflective of a compensatory enrichment in intrathymic precursors (i.e., DN thymocytes that possess the ability to repopulate the thymus of irradiated recipients upon adoptive transfer). Alternatively, the DN thymocytes remaining in the involuted thymuses of me^meme^m mice may be CD3^+^ DN thymocytes with no precursor activity upon adoptive transfer (Scollay et al., 1988). To investigate these various possibilities, we used the i.t. adoptive transfer assay (Goldschneider et al., 1986; Scollay et al., 1988; Shortman et al., 1988) to quantitate the intrathymic precursor activity of thymocytes in me^meme^m mice. This assay system was used because intrathymic precursors poorly repopulate the thymus following i.v. adoptive transfer (Kadish and Basch, 1977; Fowlkes et al., 1985; Scollay et al., 1988), and intrathymic precursor activity is readily quantified using the i.t. adoptive transfer assay (Goldschneider et al., 1986; Scollay et al., 1988).

Thymocytes recovered from me^meme^m mice with involuted thymuses were relatively deficient, on a per cell basis, in their ability to repopulate the thymus of adoptive recipients as compared to thymocytes recovered from B6+/+ mice. The i.t. transfer of 1.2 to 1.9x10^6 thymocytes from me^meme^m mice with 2.1±1.0x10^6 cells/thymus resulted in the generation of <1x10^6 (nondetectable) (n=6) donor-origin cells on day 14, and the transfer of 1x10^6 or 2.5x10^6 thymocytes from B6+/+ mice resulted in the generation of 3.9±1.0x10^5 (n=4) and 8.0±4.3x10^5 (n=3) donor-origin cells, respectively, on day 14. In addition, it was of interest to compare the "total" precursor population in the thymus of me^meme^m mice with that of normal B6+/+ mice during the involusion process. To compare this directly, the relative intrathymic precursor activity of each thymus, which takes into account the total cellularity of the thymus, hence the total precursor pool within the thymus, was calculated (see Materials and Methods).

Thymocytes from 4-week-old me^meme^m mice clustered into two distinct groups when the relative intrathymic precursor activity was calculated (Fig. 2A). In one group, the relative intrathymic precursor activity was equivalent to that of age-matched B6+/− littermates. In the second group, the relative intrathymic precursor activity was decreased in relation to that of B6+/− littermates (Fig. 2A). In 5-, 6-, and 7-week-old me^meme^m mice, the relative intrathymic precursor activity was reduced in relation to that of age-matched B6+/− littermates, approaching nondetectable levels of precursor activity in 60% (3/5) of 6-week-old me^meme^m mice and 100% (3/3) of 7-week-old me^meme^m mice. No differences were observed in the relative intrathymic precursor activity of B6+/− littermates between 4 and 7 weeks of age (Fig. 2A), which, as observed in bone marrow prothymocyte activity (Komschlies et al., 1987), were comparable to that of wild-type B6+/+ mice (data not shown). Furthermore, in 3-week-old me^meme^m mice, we observed a decrease in intrathymic precursor activity prior to detectable thymic involution, suggesting that the loss of intrathymic precursor activity precedes thymic involution (data not shown).

Due to the variability in intrathymic precursor activity observed in involuting me^meme^m thymuses, especially in 4-week-old me^meme^m mice, we determined whether the extent of thymic involution in me^meme^m mice (loss in total cellularity) was related to the level of intrathymic precursor activity. As shown in Fig. 2B, there is a strong linear correlation (r^2=0.872) between the cellularity of the me^meme^m thymus and the relative intrathymic precursor activity detectable during involution; as thymic cellularity decreased, the amount of intrathymic precursor activity correspondingly decreased.

**Phenotypic Characteristics of Thymocytes in Aged B6 Wild-Type Mice During Thymic Involution**

Motheaten (me/me) and me^meme^m mice have expected lifespans of 22 and 61 days, respectively (Shultz and Sidman, 1987; Shultz, 1988), and we have previously suggested (Medlock et al., 1986) that these mice may be representative of an accelerated aging process in the immune system. Therefore, it was of interest to compare directly
Intrathymic Precursor Activity of Thymocytes in Aged Mice During Thymic Involution

Thymocytes from weanling (1-month-old) B6+/+ mice were comparable to that of weanling B6+/− littermate mice in intrathymic precursor activity (Figs. 2A and 3A). By 11 to 13 months of age, however, the intrathymic precursor activity of B6+/+ mice was decreased by 57 to 87% as compared to that of 1-month-old B6+/+ mice and continued to decrease with time to essentially nondetectable levels (98% decrease) of intrathymic precursor activity in 100% (4/4) of 16–17-month-old mice (Fig. 3A). As observed in involuting meV/meV thymuses, the decrease in thymus cellularity exhibited a strong linear correlation (r²=0.900) with the loss of intrathymic precursor activity with age (Fig. 3B).

Prothymocyte Activity in the Bone Marrow of Aged Mice

Viable motheaten (meV/meV) mice with severely involuted thymuses still contain normal numbers of bone marrow prothymocytes as quantitated by the i.t. adoptive transfer assay system, but no detectable intrathymic precursor activity (Greiner et al., 1986; Komschlies et al., 1987; Fig. 2). Therefore, we determined whether the loss of intrathymic precursor activity in B6+/+ aged...
mice is associated with a decrease in bone marrow prothymocyte activity. Bone marrow was recovered from 12-month-old B6+/+ mice (57 to 87% decrease in intrathymic precursor activity) and injected both i.v. and i.t. into young (4–6-week-old) irradiated adoptive recipients to determine their relative bone marrow prothymocyte activity. As shown in Table 3, in contrast to that observed in meV/meV mice and as shown previously by other investigators (Hirokawa et al., 1986), the bone marrow of aged mice is able to repopulate the thymus of young adoptive recipients following i.v., as well as i.t., adoptive transfer. Furthermore, on a per cell basis, bone marrow from 1-year-old donors does not differ significantly from that of weanling (1-month-old) bone marrow in relative prothymocyte activity. For the i.v. adoptive transfer system, the relative prothymocyte activity is 219±109 (mean±S.D.) units for young donors and 421±147 units for aged ones. Similar results are observed for the i.t. adoptive transfer system in which the relative prothymocyte activity is 10,410±7,563 units for young bone marrow and 16,245±8,189 units for aged marrow.

To investigate further the possible mechanisms that account for thymic involution in aged mice, especially since bone marrow precursor activity in older mice is comparable to that of young mice (Table 3), we determined the ability of young and older (12-month-old) bone marrow to repopulate the thymuses of older (10-month-old) irradiated recipients. Surprisingly, there were no significant differences (p>0.1 for the i.v. assay; p>0.5 for the i.t. assay) in the number of thymocytes generated by either young or old marrow on a per cell basis in older recipients as compared to that of young recipients. The relative prothymocyte activities calculated for both adoptive transfer assays are 477±127 units (i.v.) and 15,912±3,917 units (i.t.) for young bone marrow and 372±58 units (i.v.) and 14,070±9,478 units (i.t.) for aged marrow.

**Phenotypic Characteristics of Thymocytes in Corticosteroid-Treated Mice Following Thymic Involution**

To investigate whether the thymic involution observed in meV/meV mice resembled that of corticosteroid-induced thymic involution, normal B6+/+ mice were injected with dexamethasone (0.5 mg), and the thymus analyzed 48 h later. Steroid treatment resulted in a 97% decrease in cellularity, and a decrease in the absolute numbers of each of the four major thymocyte subsets (Table 4). The proportions of each of the thymus subsets that were present, however, were different (Fig. 1D) from that observed in an meV/meV involuted thymus (Fig. 1B). The proportion of DN thymocytes increased to 19±3.1%, similar to that of meV/meV thymocytes (range 5.5–32%), but the percentage of DP
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A

B

FIGURE 3 Thymocytes from 1-, 11-13-, and 16-17-month-old normal B6+/+ mice (Ly5.2) were tested for precursor activity following intrathymic adoptive transfer into irradiated B6-Ly5.1 congenic recipients. Relative intrathymic precursor activity was calculated as described (see Materials and Methods). Values represent data from individual thymuses. (A) Relative intrathymic precursor activity as a function of age normalized to that of control 4-week-old B6+/+ mice. (B) Relative intrathymic precursor activity as a function of thymus cellularity ($r^2=0.900$).

| TABLE 3 |
| Effect of Age on Bone Marrow Prothymocyte Activity in C57BL/6J+/+ Mice* |
| --- | --- | --- | --- |
| Age of donor (months) | Age of recipient (months) | Route of injection | Donor-origin thymocytes Percentage Cells/thymus ($\times 10^6$) |
| 1 | 1 | i.t. | 67.8±18.7 90.5±65.8 |
| 12 | 1 | i.t. | 85.2±11.0 81.2±40.9 |
| 1 | 10 | i.t. | 86.4±4.6 146.3±36.0 |
| 12 | 10 | i.t. | 78.4±12.3 112.6±75.8 |
| 1 | 1 | i.v. | 73.2±10.6 95.5±47.4 |
| 12 | 1 | i.v. | 82.1±5.2 105.3±36.9 |
| 1 | 10 | i.v. | 94.7±0.9 216.3±54.1 |
| 12 | 10 | i.v. | 89.5±6.2 148.9±23.2 |

*Bone marrow cells from young (1-month-old) and aged (12-month-old) mice were injected i.t. (0.2x10⁶) or i.v. (10x10⁶) into irradiated young (1-month-old) or aged (12-month-old) B6+/+ recipients. Twenty days post injection, thymuses were recovered and analyzed for donor-origin thymocytes using the indirect immunofluorescence staining technique (see Materials and Methods). Data represent mean±standard deviation for three or more animals per group.

Intrathymic Precursor Activity of Thymocytes in Steroid-Treated Mice

Thymocytes recovered from steroid-treated B6+/+ mice were tested for their intrathymic precursor activity in the i.t. adoptive transfer assay. The thymocyte repopulating activity of steroid-treated thymocytes (1.1±0.1 units) was decreased more than 99% when compared to the relative thymocyte repopulating activity of age-matched untreated B6+/+ (392.2±185.8 units) mice.

Prothymocyte Activity in the Bone Marrow of Steroid-Treated Mice

The high sensitivity of intrathymic precursors to steroids was surprising in light of previously published results detailing the relative steroid resistance of bone marrow prothymocytes (Greiner et al., 1982). To confirm this result, bone marrow cells were recovered from the same steroid-treated mice that were used as donors of thymocytes (see Table 4), and were injected both i.t. and i.v. into adoptive recipients. As expected, on a per cell basis and as previously reported

thymocytes (10.4±3.4%) was decreased relative to involuted me/me' thymocytes (range 33–72%), and SP thymocytes were the predominant thymocyte subset (65.8±5.4%) in steroid-treated mice (Table 4 and Fig. 1D).
TABLE 4

| Dexamethasone Treatment | Thymus Cell Number (x10^6) | Thymocyte Subpopulations |
|-------------------------|----------------------------|--------------------------|
|                         |                            | CD4^+CD8^- (%)           | CD4^+CD8^+ (%)           | CD4^-CD8^+ (%)           | CD4^-CD8^- (%)           |
| +                       | 7.5±4.1                    | 19.3±3.1b                | 10.4±2.4                 | 48.1±4.6                 | 17.8±3.8                |
| -                       | 173.3±10.4                 | 3.0±0.6                  | 82.4±5.9                 | 10.1±2.1                 | 4.5±1.2                 |

*Four-week-old C57BL/6+/+ mice injected with dexamethasone (0.5 mg) and the thymuses recovered for analysis 48 h later. Thymocytes from these mice were analyzed for expression of CD4 and CD8 antigens using two-color immunofluorescence staining (see Materials and Methods). Data represent the mean percentage ± standard deviation for four mice in the dexamethasone-treated group and for three mice in the untreated group.

bPercentage for CD4^-CD8^- thymocytes excludes surface Ig^+ cells in the thymus.

...DN cells approximates that observed in normal thymuses (approximately 50%).

DISCUSSION

In the present study, we have demonstrated that me^+/me^+ invovled thymuses display a loss in all four of their major phenotypic thymocyte subsets (DN, DP, SP) by 4 to 7 weeks of age. Prior to this loss of thymocytes, a decrease in the intrathymic precursor activity was observed. This decrease was evident as early as 3 weeks of age in me^+/me^+ mice whose thymuses have not lost cellularity. At 4 weeks of age, approximately half of the me^+/me^+ mice examined had reduced intrathymic precursor activity, and the majority of the animals studied at 6 to 7 weeks of age demonstrated reduced activity. Although a relative increase in the percentage of DN thymocytes was observed...
in the involuting thymus of me"/me" mice, there
was a decrease in the absolute number of DN
cells relative to that of normal age-matched
B6+/+ mice. Because only DN thymocytes are
able to generate the other three phenotypic
thymocyte subsets (DP and SP) in vivo in adoptive
recipients (Scollay et al., 1988; Shortman et al.,
1990), our observations would suggest that
the intrathymic precursors, although reduced in
absolute numbers, may be relatively enriched in
involuting me"/me" thymuses. However, the DN
thymocytes that were present in the severely
involuted me"/me" thymuses were deficient in
their ability to generate thymocytes upon trans-
fer to irradiated adoptive recipients.

It has been calculated that as few as 1×10⁵ puri-
fied DN thymocytes can generate detectable
levels of donor-origin thymocytes in an
irradiated recipient's thymus, while up to 6×10⁷
DN thymocytes (2×10⁷ unfractionated thymo-
cytes) are required for saturation repopulation
of the thymus of an irradiated recipient (Scollay et
al., 1988). We injected in our experiments at least
5×10⁵ thymocytes from each involuted thymus
into the irradiated recipient's thymus. Because,
in the severely involuted thymus, at least 20% of
the thymocytes were DN (Table 1), at least
100,000 DN cells were injected into the thymus of
each of the adoptive recipients. Our failure to
find evidence of intrathymic precursor activity
following injection of approximately 100-fold
more DN thymocytes than should have been
necessary to generate thymocytes in adoptive
recipients (and approximately fourfold more DN
thymocytes on a per cell basis than that present
in normal thymus that gave high levels of
repopulation in adoptive recipients) suggests
that at a maximum, less than 1% of the DN
thymocytes in the severely involuted me"/me" thym-
uses could have had precursor activity.

The DN thymocytes that are present in normal
mice, however, are composed of at least 11 differ-
ent subsets, each with different functional capa-
bilities and kinetics of thymocyte repopulation in
adoptive recipients (Scollay et al., 1988). Also
contained within these various thymocyte sub-
sets are DN thymocytes that express either CD3
or surface Ig and are unable to function as intra-
thymic precursors in adoptive transfer assay sys-
tems (Scollay et al., 1988). In addition, variations
in the DN CD3⁺ population with respect to
mouse strains and to the class of T-cell receptor
expressed occurs (Shortman et al., 1990). Conse-
quently, we analyzed the DN thymocyte popu-
lation in the involuted thymuses to determine
whether there was an increase in the percentage
of DN cells expressing CD3 using the 2C11
monoclonal antibody that detects a portion of the
CD3 complex expressed by both alpha/beta- and
gamma/delta-expressing T cells. An increase in

| Animal age (months) | Thymus cell number (x10⁶) | CD4⁺CD8⁺ thymocytes (%) | Percentage of CD4⁺CD8⁺ thymocytes that are: |
|---------------------|--------------------------|-------------------------|--------------------------------------------|
|                     |                          |                          | CD3⁺ or sIg⁺ | CD3⁺ sIg⁺ |
| Untreated (1 month) | 173.3±10.4               | 3.9±0.5                 | 51 | 49 |
| Untreated (12 months) | 73.3±33.9               | 4.3±0.5                 | 51 | 49 |
| Untreated (18 months) | 53.0⁺†                  | 5.2                     | 67 | 33 |
|                       | 42.0                    | 3.5                     | 69 | 31 |
|                       | 23.0                    | 24.3                    | 81 | 19 |
| Steroid-treated (1 month) | 7.5±4.1                | 23.8±2.2               | 41 | 59 |
| me"/me" (1 month)  | 270.0⁺†                 | 4.1                     | 30 | 70 |
|                     | 150.0                   | 4.5                     | 20 | 80 |
|                     | 13.0                    | 14.0                    | 49 | 51 |

*Thymocytes from normal and involuted thymuses were analyzed for expression of CD4, CD8, CD3, and surface Ig using two-color immunofluorescence staining (see Materials and Methods). Data, unless otherwise marked, represent means±standard deviations of three or more thymuses.

†Due to variability among a single group, data are given for individual thymuses.
the proportion of DN CD3-expressing cells would indicate an increase in the percentage of CD4$^+$CD8$^-$ thymocytes that would be unable to generate thymocytes in adoptive recipients. We have demonstrated that the involution process in the me$^e$/me$^e$ mouse does not affect the relative proportion of these cells in the DN subset (Table 6).

In light of our findings, it appears that the thymic involution in me$^e$/me$^e$ mice may in part be due to the CD4$^+$CD8$^-$CD3$^-$sIg$^-$ subset being unable to generate thymocytes, as demonstrated by their transfer to adoptive recipients. To determine whether these precursors are arrested in development, it will be of interest in future studies to examine additional time points following adoptive transfer of the donor thymocytes to confirm the absence in involuted me$^e$/me$^e$ thymuses of each of the developmentally distinct populations that have been described. However, based on our inability to demonstrate any intrathymic precursor activity at day 14 following transfer to irradiated recipients, and because of the large excess of DN cells injected, it seems highly unlikely that we have missed a significant population of DN thymocytes in involuting me$^e$/me$^e$ thymus that possesses precursor activity.

Alternatively, we have previously suggested (Komschlies et al., 1987) that the bone marrow of me$^e$/me$^e$ mice is deficient in an accessory cell (or factor) that permits repopulation of the thymus of irradiated recipients following i.v. adoptive transfer. Addition of normal bone marrow to me$^e$/me$^e$ bone marrow restores its i.v. thymus repopulating capacity. Viable motheaten (me$^e$/me$^e$) bone marrow (as are normal thymocytes), however, is able to repopulate the thymus of irradiated recipients following i.t. injection without additional accessory cell (or factor) requirements. It has been shown previously that a normal thymus contains a population of bone marrow derived "accessory" dendritic cells of myeloid origin (Fowlkes and Pardoll, 1989; Robey et al., 1990; Spangrude and Scollay, 1990; von Boehmer, 1990) that are important in the clonal selection, survival, and differentiation of thymocytes (Robey et al., 1990; von Boehmer, 1990). Codevelopment of such myeloid stem cells following bone marrow i.t. injection appears to be requisite for the subsequent development of intrathymic precursors and donor-origin thymocytes (Spangrude and Scollay, 1990). Because numerous defects in myeloid cells have been described in me$^e$/me$^e$ mice in vivo and in vitro (McCoy et al., 1982, 1984; Medlock et al., 1987; Hayashi et al., 1988; Shultz, 1988; Van Zant and Shultz, 1989), the lack of these bone marrow derived dendritic cells in an me$^e$/me$^e$ thymus may result in the absence, death, and/or functional inactivity of me$^e$/me$^e$ intrathymic precursors. However, cell mixture experiments of me$^e$/me$^e$ thymocytes with normal bone marrow cells failed to "rescue" or provide evidence for suppression of the thymus repopulating capacity of me$^e$/me$^e$ DN thymocytes following i.t. adoptive transfer into irradiated recipients (unpublished observations). These results suggest that me$^e$/me$^e$ DN thymocytes, if they in fact are intrathymic precursors, may be anergic to such regulatory signals. It does not rule out, however, that these positive signals may be required only at specific, perhaps early, stages of precursor development. The setup of our assay system, in which "accessory" cells are not added until thymic involution (hence loss of intrathymic precursor activity) has already occurred, may be too late to rescue the intrathymic precursor population.

It was also of interest in these studies to compare the phenotypic and functional characteristics of thymocytes in an involuting me$^e$/me$^e$ thymus with other models of thymic involution in an attempt to gain insight into the possible mechanisms involved. Thymic involution has been observed in pregnancy, infection, surgery, drugs (including cyclosporin A), steroids, malnutrition, malignancy, and aging (Clarke and Kendall, 1989). Of special interest in our studies was the involution in me$^e$/me$^e$ thymus with that of thymuses in aged mice, because we have suggested previously (Medlock et al., 1986) that the me$^e$/me$^e$ mouse may represent an accelerated model of aging in the lymphopoietic system. Thymic involution in normal mice begins soon after puberty, while the animal is still increasing in body weight (Santisteban, 1960). There is a decrease in thymic weight with age, a loss of cortical thymocytes, an increase of thymic adipose tissue, and a general decrease in thymic function. In addition, the thymic atrophy is reflected in the peripheral tissues by a decrease in T-cell functional responses, and an increase in a CD4$^+$ Fgp-1$^+$ population reflective of memory T cells (Nagelkerken et al., 1991).
As predicted, by 12 to 18 months of age, the thymus cellularity in B6+/+ mice is reduced by approximately 90% in comparison to that of young (4-week-old) B6 mice. Our phenotypic studies have demonstrated that in an involuting aged thymus, essentially normal ratios of the four phenotypic subsets of thymocytes are present. Therefore, in the in vivo studies, equivalent numbers of young and aged DN thymocytes were injected into the recipient thymus. Even though this is the case, the 17-month-old thymus is severely deficient in intrathymic precursor activity, and displays essentially no ability to repopulate the thymus of irradiated recipients at the cell doses examined (3–5×10⁶ cells injected). One 18-month-old thymus, however, did exhibit a variation in the proportions of the major thymocyte subsets (24.3% DN, 43.6% DP, 32.1% SP). When the DN thymocyte population was also stained for CD3 or slg, it was observed that in this involuted thymus, 80% of the CD4⁺CD8⁺ thymocytes expressed either CD3 or Ig.

It appears, however, that the loss of intrathymic precursor activity with age in mice is not due to a defect in the bone marrow prothymocyte population. We found that on a per cell basis, bone marrow from older mice is as efficient as young marrow in thymus repopulating capacity. Contradictory results exist in regards to the study of the aging process and the immune system. CFU-s activity and basal hemopoietic activity do not appear to be affected by age (Tyan, 1982; Williams et al., 1986). However, other investigators find prothymocyte and thymic activity to be impaired in aged mice (Kay, 1984; Hirokawa et al., 1986). We observed the opposite; the defect is not readily apparent in the thymus, as older irradiated recipients readily supported the generation of donor-origin thymocytes from both young and older marrow as well as did younger irradiated recipients. It must be cautioned that the irradiation model of repopulation may not directly represent the in vivo situation in regard to thymus function. Thus, irradiation may induce or reactivate "accessory cells" or factors that promote thymocyte regeneration that are quiescent in intact aged hosts. For example, an irradiated thymus may be induced to secrete increased levels of chemotactic factors (Harr et al., 1989) such as thymotaxin (Imhof et al., 1988; Deugnier et al., 1989). However, it is most likely not thymotaxin, because this compound has been found to be beta-2 microglobulin (Dargemont et al., 1989), and beta-2 microglobulin-deficient mutants have normal development of the thymus (Koller et al., 1990; Zijlstra et al., 1990). Alternatively other, as yet undescribed, factors may have a role in the irradiation induced ability of thymus in aged mice to support thymocyte production in the adoptive transfer system.

Similarly, we have provided evidence that the thymic involution in meᵡ/meᵡ mice is not due to increased levels of steroids. Although essentially all intrathymic precursor activity is eliminated by a single dose of dexamethasone, we observed significant differences in the phenotypic characteristics of the remaining thymocyte populations. Thus, in thymuses from steroid-treated mice, a large proportion of the remaining lymphoid cells were SP, and in thymuses from either meᵡ/meᵡ or aged mice, a much lower proportion of the lymphoid cells were SP. The characteristics of the thymocyte populations that were present following steroid treatment were similar to those previously reported by numerous investigators (Reichert et al., 1986; Vliet et al., 1986). Of interest in this study was the unexpected finding of the higher steroid sensitivity of intrathymic precursors compared to bone marrow prothymocytes.

In the bone marrow, the prothymocyte population is relatively resistant to steroids (Greiner et al., 1982). We confirmed these findings, and we were able to calculate that approximately 60% of bone marrow prothymocyte activity is lost following steroid treatment. This is in marked contrast to the loss of intrathymic precursors observed in the steroid-treated mice, in which greater than 99% of the precursor activity was lost. The loss of intrathymic precursor activity, however, is supported by phenotyping data. Approximately 40% of steroid-resistant DN thymocytes express CD3 or Ig and would be unable to generate thymocytes in adoptive recipients (Table 6). Furthermore, other investigators have shown that steroid treatment depletes CD5⁺ thymocytes (Scollay and Shortman, 1985), a population that contains intrathymic precursor activity (Fowlkes et al., 1985). Thus, although steroids resulting from the stress induced by the clinical symptoms of the autoimmune disease process may contribute to the early thymic involution observed in meᵡ/meᵡ mice, such contribution is most likely a small and insignificant part of the involution process.
MATERIALS AND METHODS

Animals
Inbred C57BL/6J (Ly 5.2) and Ly 5 congenic C57BL/6J-Ly 5.1 mice (hereafter referred to as B6 Ly 5.2 and B6 Ly 5.1, respectively) were purchased from Charles River (Boston, MA) and Harlan Sprague Dawley (Frederick, MD), respectively. C57BL/6J-meV/meV mice (hereafter referred to as meV/meV mice; Ly 5.2) were obtained from the colony maintained at the Jackson Laboratory by LDS. The inbred C57BL/6 mice are designated as Ly 5.2; the congenic is designated by Ly 5.1 (SJL/J phenotype). This nomenclature is consistent with that recently recommended by Morse et al. (1987), and is opposite from the designation of the monoclonal antibodies commercially available from Dupont/New England Nuclear. Unless otherwise noted, animals were placed into experiments when they were 4–6 weeks of age, and were maintained on commercial chow and acidified chlorinated water ad libitum.

Antibodies
The unconjugated monoclonal antibodies directed against the Ly 5.1 and Ly 5.2 alloantigens were obtained from Dupont/New England Nuclear (Boston, MA) and were developed for immunofluorescence analysis using an F(ab')2 fragment of FITC-conjugated goat antimouse IgG (heavy- and light-chain-specific) obtained from Organon Teknika-Cappel (Malvern, PA). Monoclonal antibodies for use in the dual labeling studies, FITC-conjugated antimouse Lyt-2 and Phycoerythrin (PE)-conjugated antimouse L3T4, were obtained from Becton Dickinson Immunocytochemistry Systems, Mountain View, CA.

Dexamethasone-Resistant Thymocytes and Bone Marrow Cells
Dexamethasone-resistant thymocytes and bone marrow cells were obtained 48 h after i.p. injection of 0.5 mg of dexamethasone sodium phosphate (LyphoMed, Inc., Melrose Park, IL) into 4–6-week-old mice (Vliet et al., 1986; Compton et al., 1987).

Cell Suspensions
Bone marrow cells were obtained by flushing tibias and femurs with cold medium (Hepes buffered RPMI 1640). After repeated gentle pipetting to disperse the cells, the marrow was centrifuged at 150 x g for 5 min and resuspended in cold medium for cell counting. Thymus cell suspensions were prepared by gently pressing the tissues through a 50-mesh cell sieve followed by washing in cold medium. Cell viability was determined by exclusion of 0.1% trypan blue and was greater than 95% in all cases (Greiner et al., 1986).

Immunofluorescence Analysis
Thymocytes were labeled for immunofluorescence analysis as previously described (Goldschneider, 1986). Briefly, a single-cell suspension of thymocytes (1 x 10^6/well) was added to individual wells of 96-well U-bottom microtiter plate and spun at 150 x g for 2 min. The supernatant was removed, the cells resuspended directly in 10 µl of a saturating concentration of the primary antibody (anti-Ly 5.1 or anti-Ly 5.2) or medium (control) and incubated for 20 min at 4 °C. The cells were spun at 150 x g for 2 min, the supernatant removed and the cells were washed twice in cold medium containing 0.13% sodium azide. After the last wash, the cells were resuspended directly in 10 µl of an appropriate dilution of the developing antibody and incubated for 20 min at 4 °C. After the second incubation, the cells were washed, fixed in buffered formalin, and analyzed by flow microfluorimetry for relative immunofluorescence on a FACS 4 (Becton-Dickinson Immunocytochemistry System, Sunnyvale, CA). All viable nucleated cells were gated for analysis, and dead cells and red blood cells were excluded electronically. In two-color analysis, thymocytes were incubated with FITC-anti-Lyt 2 alone, with PE-anti-L3T4 alone, or simultaneously with both. In some cases, to exclude the possible contribution of surface Ig’ B cells in the involuted thymuses in the calculation of double negative (CD4^-CD8^-) thymocytes, anti-CD4, anti-CD8, and FITC-conjugated goat antigoat mouse IgG were used to stain the thymocytes.

Irradiation
Recipients received the indicated level of whole body irradiation from a ^137Cs source (gamma cell 40 irradiator, Atomic Energy of Canada Limited,
Ottawa, Canada) at a dose rate of approximately 95 rads/min 2–6 h prior to bone marrow cell transfer (Greiner et al., 1986).

**Intrathymic Adoptive Transfer Assay System**

Details of the intrathymic (i.t.) adoptive transfer assay system for prothymocytes have been described previously (Goldschneider et al., 1986; Greiner et al., 1986; Komschlies et al., 1987; Scollay et al., 1988). Briefly, irradiated (700 rad) recipients received varying numbers of thymocytes or bone marrow cells in 20 μl of media distributed equally between each thymus lobe (10 μl/lobe). Thirteen to fifteen days after thymocyte transfer or 18 to 20 days after bone marrow cell transfer, the thymus was removed, and the percentage and number of donor- and host-origin cells were determined by immunofluorescence analysis of Ly 5.1 (host) and Ly 5.2 (donor) bearing thymocytes using flow microfluorimetry.

**Intravenous Adoptive Transfer Assay System**

This assay has been described in detail (Goldschneider et al., 1986; Greiner et al., 1986; Komschlies et al., 1987). Briefly, irradiated recipients (700 rad) were injected i.v. in the tail vein with varying numbers of bone marrow cells (see Results). Twenty to twenty-three days later, the percentage and number of donor- (Ly 5.2+) and host- (Ly 5.1+) origin thymocytes were determined by immunofluorescence analysis using flow microfluorimetry.

**Calculation of Relative Intrathymic Precursor or Bone Marrow Prothymocyte Activity**

To calculate the relative intrathymic or prothymocyte precursor activity of a donor thymus or bone marrow when using either the i.t. or i.v. adoptive transfer assay system, the following formula was used:

\[
\text{Number of thymocytes generated from donor-origin cells} \times \text{Cell dose} \times \text{donor cell number} = \text{precursor activity}
\]

For example, if 1x10^6 thymocytes injected from a donor thymus that had 3x10^6 cells generated 15 x10^6 donor-origin thymocytes in the adoptive recipient, the total intrathymic precursor activity of the donor thymus would be 450x10^6.

**Calculation of Normalized Intrathymic Precursor Activity**

Once relative intrathymic precursor activity was calculated for individual thymuses, the mean (±standard deviation) intrathymic precursor activity was calculated for the control thymuses. The experimental thymus intrathymic precursor activity was then normalized as a ratio of the mean control thymus value.

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