Modulation of Thymic Selection by Expression of an Immediate-early Gene, Early Growth Response 1 (Egr-1)

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Summary

The potential involvement of early growth response (Egr)-1, a zinc-finger transcription factor belonging to the immediate-early genes, in positive/negative selection of thymocytes has been implicated by its expression in the population of CD4+CD8+ double positive (DP) cells undergoing selection. To further investigate this possibility, transgenic mice overexpressing Egr-1 in thymocytes were bred with a transgenic mouse line expressing a T cell receptor (TCR) recognizing the H-Y male antigen in the context of H-2b class I major histocompatibility complex (MHC) molecules. In Egr-1/TCR H-Y double-transgenic mice, efficient positive selection of H-Y CD8+ T cells occurred, even in mice on either a nonselecting H-2d background or a β2-microglobulin (β2m)-deficient background in which the expression of class I MHC heavy chains is extremely low; no positive selection was observed on a Kb2/Db2/β2m2/2 background where class I MHC expression is entirely absent. Similarly, when the Egr-1 transgene was introduced into a class II MHC–restricted TCR transgenic mouse line, Egr-1/TCR double-transgenic mice revealed increased numbers of CD4+ T cells selected by class II MHC, as well as significant numbers of CD8+ T cells selected by class I MHC (for which the transgenic TCR might have weak affinity). Thus, Egr-1 overexpression allows positive selection of thymocytes via TCR–MHC interactions of unusually low avidity, possibly by lowering the threshold of avidity required for positive selection. Supporting this possibility, increased numbers of alloreactive T cells were positively selected in Egr-1 transgenic mice, resulting in a strikingly enhanced response against allo-MHC. These results suggest that expression of Egr-1 and/or its target gene(s) may directly influence the thresholds required for thymocyte selection.

Key words: Egr-1 • positive selection • T cell • avidity • transgenic mouse

1 Abbreviations used in this paper: DP, double positive; Egr-1, early growth response 1; Egr/H-Y, Egr-1 transgene–positive H-Y transgenic; I0, class I MHC–deficient; II0, class II MHC–deficient; ISP, immature SP; NL/H-Y, Egr-1 transgene-negative H-Y transgenic; PNAR, peanut agglutinin receptor; RAG, recombination-activating gene; SP, single positive; TG, transgenic.

Positive/negative selection of thymocytes at the CD4+CD8+ double positive (DP) stage is the key checkpoint for thymocyte maturation, at which their fate—whether they develop further to CD4+ or CD8+ single positive (SP) cells or die by clonal deletion—is decided (1, 2). Accumulating evidence indicates that the avidity between T cell receptors (TCR) expressed on thymocytes and MHC/antigen-peptide complexes displayed on the surface of APCs appears to define the type of intracellular signals generated, which promote thymocytes to be either positively selected (further developing to the SP stage) or negatively selected (clonally deleted); avidity is dependent on TCR–MHC affinity and the expression levels of both complexes (3, 4). It is believed that there are thresholds in the strength of TCR–MHC avidity that determine the nature and consequences of subsequent TCR signaling (3, 4). When thymocytes and APCs interact with high avidity, cells are negatively selected through the apoptotic pathway, whereas thymocytes are promoted to the SP cell stage when the avidity is moderate but sufficient. If the avidity is too low, cells cannot undergo either type of selection, resulting in their death as “neglected cells.” However, the molecular mechanism that defines these thresholds for selection events is unclear.

Signals generated during positive/negative selection must be differentially controlled; hence, rapidly responding transcriptional regulators able to elicit a cascade of changes in gene expression should be important. Immediate-early genes, expression of which is rapidly induced after cell-surface receptor ligation without de novo protein synthesis, are strong candidates for such a rapid response mediator (5).
Shao et al. reported recently that the expression of one of these immediate-early genes, early growth response (Egr)-1, a zinc-finger transcription factor rapidly induced by TCR ligation, appears to correlate with selection events, as it is expressed at much lower levels in DP cells from mutant mice deficient for both MHC class I and class II molecules than in DP cells from wild-type mice. In addition, Egr-1 expression in the mutant DP cells was able to be upregulated by anti-CD3 mAb ligation (6).

We recently generated transgenic mouse lines over-expressing Egr-1 in thymocytes under the control of the lck-proximal promoter (7), and showed that in Egr-1 transgenic mice on a recombination-activating gene (RAG)-deficient background, thymocytes bypassed the block at the CD25−CD44−DN stage and matured to the immature single positive (ISP) cell stage. Here, the effect of Egr-1 expression on positive/negative selection is extensively analyzed by breeding Egr-1 transgenic mice with various TCR transgenic mice, as well as by evaluating positive selection of alloreactive T cells in Egr-1 transgenic mice. This report provides interesting insight into what is actually required for positive selection and what transcriptional pathways may be involved in the selection events.

Materials and Methods

Mice. All mice used here were maintained in the specific pathogen-free facility of the Basel Institute for Immunology. Mice were of mixed C57BL/6 backgrounds and were used for all experiments. Senologial reagents and Flow Cytometry. Reagents used for staining T cells and subsets thereof were as described (7–9, and references therein). Thymus, lymph node, and spleen cells were stained with saturating levels of mAbs and analyzed using a FACS Calibur 

Cell Survival Assay. Purified CD8+ T cells were sorted after staining thymocytes for CD4, CD8, T3,70, and peanut agglutinin receptor (PNA). Stained cells from each type of mice were cultured in triplicate in 96-well flat-bottomed microculture plates in complete DMEM culture medium including 10% FCS. 24, 48, 72, 96, and 120 h after starting the culture, the number of living cells was determined by a modification of a similar experiment described elsewhere (10). In brief, purified CD8+ T cells including 2 × 10^5 H-Y CD8+ T cells calculated by the percentage of T3,70+ population in the CD8+ cells after fluorocytometric analysis, were suspended in 200 μl of PBS, and injected intravenously into 6-8-wk-old male and female C58-d-deficient mice. In one experiment, cells from each type of donor mouse were injected into two pairs of male and female recipient mice. 4 d after injection, spleen cell suspension from each recipient was stained for CD8 and T3,70, and analyzed by FACScan (Becton Dickinson). Absolute number of H-Y CD8+ T cells was calculated from spleen cell number and percentage of CD8+ T cells in each type of mouse, including various numbers of H-Y CD8+ T cells were stimulated by 8 × 10^6 well of 3,000-rad irradiated spleen cells from male or female B6 mice in complete DMEM in 96-well microculture plates. Cultures were performed for 5 d, and proliferation was assessed by [3H]thymidine incorporation in the last 16 h of culture.

Bone Marrow Chimeras. To construct bone marrow chimeras, recipient mice were 10–19 g (from a cobalt source) irradiated. 24 h later, they were injected intravenously with 10^9 bone marrow cells from which mature T and B cells had been depleted by treatment with anti-CD4 (R1L72) and anti-CD8 (31M) mAbs plus Tox complement (Cedarlane Labs Ltd.). After grafting, mice were rested for 5 wk to allow reconstitution before analysis.

Results

Efficient Positive Selection of H-Y CD8+ T Cells by Low Avidity TCR-MHC Interactions in the Presence of Egr-1 Over-expression. As described in our previous report, Egr-1 transgenic mice (Egr-1-TG) exhibit transgene expression at levels far exceeding endogenous Egr-1, when analyzed by Northern blotting using thymic RNA from transgenic and negative littermate mice (7). As observed for many other transgenes driven by the lck-proximal promoter (11), Egr-1 transgene expression was detected in a wide range of developmental stages of thymocytes of Egr-1-TG, including DP cells, when analyzed by the reverse transcription PCR method (12) using RNA from sorted cells from each population (data not shown). As observed in our previous report, Egr-1 transgenic mice do not undergo either positive or negative selection in female or male mice on an H-2d background (8, 13). Fig. 1 shows CD4+/CD8+ profiles of mature thymocytes bearing the H-Y

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To assess the possible effect of Egr-1 expression on thymocyte selection, Egr-1-TG were cross-bred with transgenic mice expressing a TCR recognizing the H-Y male antigen in the context of H-2b class I, D3 molecules. It is well-established that thymocytes expressing H-Y-specific αβ TCRs are positively selected in female H-2b mice and negatively selected in male H-2b mice, but do not undergo either positive or negative selection in female or male mice on an H-2d background (8, 13). Fig. 1 shows CD4+/CD8+ profiles of mature thymocytes bearing the H-Y

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TCR defined as T3.70 (mAb against H-Y TCR α chain) high, PNAr low. As shown in the upper panels, in H-2b/b females, both Egr-1-TG with an H-Y TCR transgene (Egr/H-Y) and Egr-1–TG-negative littermate mice with an H-Y TCR transgene (NL/H-Y) harbored mature H-Y CD81 SP thymocytes. The CD81 SP cells in these profiles do not contain immature CD8 SPs (ISPs), which exhibit high levels of PNAr (7). However, in H-Y mice, since expression levels of transgenic H-Y TCR in DN cells (which are PNAr low) are as high as those in mature SP cells (8), a significant proportion of DN cells as well as mature SP cells are plotted in these profiles. The numbers of total thymocytes were slightly larger in Egr/H-Y mice than in NL/H-Y mice (Table 1). However, the absolute numbers of mature H-Y CD81 SP cells were ~1.7 times higher in Egr/H-Y than in N L/H-Y mice (Table 1). In line with this, the percentages of T cells bearing the transgenic α chain (T3.70) in the DP and the total mature CD81 populations were larger in Egr/H-Y than in N L/H-Y mice (Table 1). Thus, H-Y CD81 T cells appear to be more efficiently positively selected in the presence of Egr-1 overexpression. More surprisingly, significant numbers of mature H-Y CD81 SP thymocytes were also observed in female Egr/H-Y mice on either a nonselecting H-2d/d background or a β2m-deficient (β2m0) H-2b/b background (14, 15), in which class I MHC heavy chains are expressed at extremely low levels (Fig. 1a, middle and bottom; absolute numbers of both total thymocytes and mature H-Y CD81 SP cells are shown in Table 1). Very few (<5 × 105) mature H-Y CD81 SP cells were detected in N L/H-Y mice on both H-2d/d and β2m0 backgrounds.

H-Y CD81 T cells were observed in peripheral lymphoid tissues in Egr/H-Y mice on H-2b/b, H-2d/d, and β2m0 backgrounds as well as in H-2b/b N L/H-Y female mice (numbers shown in Table 1). As observed in the thymus, the percentage of T cells bearing the transgenic α chain (T3.70) in splenic CD81 cells was larger in H-2b/b Egr/H-Y mice than in H-2d/d N L/H-Y mice (Table 1).

Figure 1. (a) Thymocyte suspensions from female Egr/H-Y and N L/H-Y mice on H-2b/b, H-2d/d, or β2m0 background were stained for CD4, CD8, H-Y transgenic α chain (T3.70), and PNAr, and analyzed by flow cytometry. CD4/CD8 plots of T3.70high, PNArlow cells are shown. The averages of the absolute numbers of mature H-Y CD81 SP cells (T3.70high, PNArlow) of each type of mice are indicated in each panel. Five pairs of female Egr/TG and N L/H-Y mice of each background were examined. (b) CD4/CD8 plots of T3.70high, PNArlow thymocytes from female β2m0 recipient mice into which female β2m0 Egr/H-Y bone marrow cells (BM) or female β2m0 N L/H-Y bone marrow cells were transplanted (left panels). Right, The same profile of female β2m0 recipient mice into which female β2m0 Egr/H-Y bone marrow cells were grafted. The averages of the absolute numbers of mature H-Y CD81 SP cells (T3.70high, PNArlow) from two sets of transfer experiments are indicated in each panel. (c) CD4/CD8 plots of T3.70high, PNArlow thymocytes from β2m0 or β2m0II0 Egr/H-Y mice. The averages of the absolute numbers of mature H-Y CD81 SP cells (T3.70high, PNArlow) of each type of mouse are indicated in each panel. Three pairs of female β2m0 and β2m0II0 Egr/H-Y mice were examined.
Peripheral H-Y CD8⁺ T cells revealed similar expression patterns of several activation and memory markers such as CD25, CD44, and Mel-14 in all types of mice (data not shown).

Thus, in the presence of Egr-1 overexpression, H-Y thymocytes appear to be efficiently positively selected even by H-2b class I molecules and by the extremely low levels of class II MHC (16, 17). As shown in Fig. 1, H-Y CD8⁺ T cells were detected in female recipients. These results suggest that H-Y CD8⁺ T cells from Egr/H-Y mice on all three backgrounds responded to both anti-CD3 mAb and Con A to a comparable degree as those from H-2b/b female NL/H-Y mice.

Next, to determine whether the H-Y CD8⁺ T cells specifically recognize the H-Y male antigen, equal numbers of H-Y CD8⁺ T cells from each type of mouse were injected intravenously into either male or female H-2b/b CD2⁺/⁺ recipients into which either Egr/H-Y or NL/H-Y T cells were transplanted. In contrast, no obvious H-Y CD8⁺ T cells were detected in female recipients. These results suggest that H-Y CD8⁺ T cells from all types of mice recognized and specifically responded to the male antigen; expansion of the H-Y CD8⁺ T cells resulted in accumulation of a significant population of those T cells in recipient spleens. Interestingly, the number of H-Y CD8⁺ T cells in male recipients into which cells from Egr/H-Y mice on all three backgrounds responded to both anti-CD3 mAb and Con A to a comparable degree as those from H-2b/b female NL/H-Y mice.

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### Table 1. Positive Selection of H-Y CD8⁺ T cells

<table>
<thead>
<tr>
<th>Mice</th>
<th>Absolute number (× 10⁶)</th>
<th>% of T3.70⁺ cells in:</th>
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<tr>
<td></td>
<td>Total</td>
<td>H-Y CD8⁺</td>
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<td></td>
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<td>DP</td>
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<tr>
<td>Thymus</td>
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<tr>
<td>N/L/H-Y (b/b)</td>
<td>7.1 ± 0.2</td>
<td>1.4 ± 0.3</td>
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<tr>
<td>Egr/H-Y (b/b)</td>
<td>8.0 ± 0.2</td>
<td>2.6 ± 0.2</td>
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<tr>
<td>Egr/H-Y (d/d)</td>
<td>7.0 ± 0.4</td>
<td>1.0 ± 0.1</td>
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<tr>
<td>Egr/H-Y (p2m⁰)</td>
<td>7.2 ± 0.1</td>
<td>0.9 ± 0.2</td>
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<td>Spleen</td>
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<tr>
<td>N/L/H-Y (b/b)</td>
<td>0.9 ± 0.1</td>
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<tr>
<td>Egr/H-Y (b/b)</td>
<td>1.5 ± 0.3</td>
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<tr>
<td>Egr/H-Y (d/d)</td>
<td>1.6 ± 0.4</td>
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<tr>
<td>Egr/H-Y (p2m⁰)</td>
<td>1.4 ± 0.2</td>
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Absolute numbers of total thymocytes and mature H-Y CD8⁺ T cells are shown. Percentages of T3.70⁺ population in DP or total mature CD8⁺ cells in the thymus and spleen are also presented. Each value represents the mean of data compiled from five mice ± SD.
in N.L/H-Y female H-2b/b mice in the absence of the Egr-1 transgene.

More Efficient Negative Selection in Egr-1 Transgenic Mice. In male H-2b/b thymi, H-Y thymocytes are negatively selected and die through the apoptotic pathway (8). As shown in Fig. 3, both Egr/H-Y and N.L/H-Y thymocytes have undergone clonal deletion. Interestingly, even more efficient negative selection appears to occur in Egr/H-Y mice, judging from the lower number of thymocytes in Egr/H-Y mice. However, male H-2d/d and β2m0 Egr/H-Y mice exhibited positive selection of H-Y CD8+ SP cells, and the numbers of mature H-Y CD8+ SP cells were comparable to those of females (data not shown).

Positive Selection of AND-TCR T Cells in the Presence of Egr-1 Overexpression. Egr-1 transgenic mice were also bred with another type of transgenic mouse expressing a TCR recognizing the moth cytochrome C peptide in the context of class II MHC I-Ek (called AND mice; reference 9). The AND TCR has a weak affinity for I-A^d class II MHC molecules, and therefore, CD4+ SP thymocytes are positively selected in an H-2b/b background (9, 21). The upper panels of Fig. 4 show the CD4/CD8 profiles of mature thymocytes bearing the AND TCR (defined as transgenic TCR [Vb3]high, PNArlow) from Egr-1 transgene-positive AND mice (Egr/AND) and Egr-1 transgene-negative AND mice (N.L/AND), both of which are on an H-2b/b background. Egr/AND mice contained increased numbers of both total thymocytes (1.1 ± 0.1 × 10^8 in Egr/AND versus 0.8 ± 0.2 × 10^8 in N.L/AND; n = 4 each) and AND CD4+ SP thymocytes (3.5 ± 0.3 × 10^7 in Egr/AND versus 2.2 ± 0.3 × 10^7 in N.L/AND; n = 4 each). This is reminiscent of the increased numbers of H-Y CD8+ SP cells in H-2b/b Egr/H-Y mice and, again, suggests more efficient positive selection in the presence of the Egr-1 transgene. Interestingly, a significant number (8.0 ± 0.6 × 10^5; n = 4) of mature AND CD8+ SP cells were observed.

Figure 2. (a and b) CD8+ lymph node T cells were purified from female Egr/H-Y mice (Egr) on H-2b/b, H-2d/d, and β2m0 backgrounds and from female N.L/H-Y mice (N.L) on H-2b/b by depleting CD4+ cells and class II MHC-positive cells. Cells were stimulated by culture with (a) immobilized anti-CD3 mAb or (b) Con A. In both experiments, proliferation was assessed in duplicate samples by [3H]thymidine incorporation. Data are representative of four independent experiments. (c) Purified CD8+ T cells including 5 × 10^6 H-Y CD8+ cells in PBS were injected intravenously into male or female C3H/HeJ recipient mice. Four days after injection, the numbers of H-Y CD8+ cells in recipient spleens were evaluated and are represented by bars ± SD. Three sets of independent experiments were performed. Numbers of H-Y CD8+ cells in female recipients were identical or less than those of mice with no cells injected [donor (—)], which is indicative of background in the fluorocytometric analysis. Id) Purified CD8+ T cells including the indicated numbers of H-Y CD8+ cells were stimulated by 3,000-rad irradiated spleen cells from male or female B6 mice. Proliferation was assessed by duplicate [3H]thymidine incorporation. Results are representative of three sets of independent experiments; all experiments showed similar results.

Figure 3. CD4/CD8 profiles of thymocytes from male Egr/H-Y and N.L/H-Y mice on an H-2b/b background, gated on T370+ cells. Numbers in the profiles indicate the averages of the absolute cell numbers of positive cells within a quadrant, of all mice analyzed. Total thymocyte numbers for both types of mice are shown. Three pairs of male Egr/H-Y and N.L/H-Y mice were analyzed.
in Egr/AND mice. It is possible that the AND TCR also has a weak affinity for class I MHC (22) which is not normally sufficient to induce signal(s) for positive selection. Egr-1 overexpression might lower the threshold for positive selection, resulting in positive selection of AND cells by class I MHC, creating a population of AND CD8\(^+\) SP cells. This hypothesis is well-supported by analysis of Egr/AND mice on a \(\beta_2m^0\) background, where the number of AND CD8\(^+\) SP cells strikingly decreased (Fig. 4, middle). However, the difference between H-\(2^d\) CD8\(^+\) SP cells and AND CD8\(^+\) SP cells in \(\beta_2m^0\) Egr-1 transgenic mice is intriguing; very efficient positive selection of H-\(2^d\) CD8\(^+\) SP cells was observed, while almost no positive selection of AND CD8\(^+\) SP cells was achieved, in the \(\beta_2m^0\) background. This may be explained by a possible difference in the original affinity of H-Y TCR and AND TCR for class I MHC molecules. The former may have a relatively high affinity, which creates sufficient avidity for positive selection even with extremely low expression levels of class I MHC, when Egr-1 overexpression lowers the threshold. In contrast, the latter may have a low affinity, which contributes to sufficient avidity with normal expression levels of class I MHC in the presence of Egr-1 overexpression, but not with the extremely low levels of class I MHC in \(\beta_2m^0\) mice.

In a class II MHC-deficient (II\(^0\)) background, in which “leaky” expression of class II molecules is not detected (23), AND CD4\(^+\) cells virtually disappeared in both Egr/AND and NL/AND mice (Fig. 4, bottom). This is consistent with failed positive selection of H-Y CD8\(^+\) T cells on an I\(^0\) background, which entirely lacks class I MHC expression. Deficiency of positive selection in the complete absence of MHC molecules indicates a requirement of TCR-mediated signaling to promote DP cells to the SP stage, even in the presence of Egr-1 overexpression.

Increased Frequency of Alloreactive T Cells in Egr-1 Transgenic Mice. Alloreactive T cells which recognize allo-MHC complexes are believed to be positively selected by auto-MHC/antigen-peptide complexes for which the alloreactive TCRs have sufficient, but not too much, avidity (24). Therefore, whether Egr-1 expression might affect the frequency of alloreactive T cells was determined using “normal” (no TCR transgenes introduced) Egr-1 transgenic mice. Fig. 5 demonstrates allo-MLR responses of H-\(2^{b_0}\) Egr-1-TG and NL mice to \(\beta_2m^0\) mice with a mutated class II and BALB/c (H-\(2^d\)) spleen cells. Transgenic T cells exhibited enhanced responses to both \(\beta_2m^0\) and BALB/c spleen cells. Similar enhanced allo-responses of H-\(2^{b_0}\) transgenic T cells were elicited towards B6 (H-\(2^{b_0}\)) spleen cells (Fig. 5 b). These enhanced allo-responses of transgenic T cells appear to be due to an increased frequency of alloreactive T cells, but not to hyperproliferation of each T cell, because fewer transgenic T cells responded to the stimulators, as shown by the limiting dilution of responding cells in Fig. 5, a and b, and transgenic and negative littermate T cells exhibited comparable proliferative responses to Con A (Fig. 5 d). Overall, Egr-1 overexpression appears to allow positive selection of a larger number of alloreactive T cells, some of which originally might not have had sufficient affinity with the auto-MHC to be positively selected in normal mice.

Discussion

This is the first description of a transcription factor behaving as a modulator of positive selection; Egr-1 overexpression allows positive selection of thymocytes mediated by low avidity interactions with ligands that are usually not sufficient, perhaps by lowering the threshold of avidity required for positive selection. One might argue that Egr-1 overexpression induces survival effectors, such as antiproliferative factors (25–30) in thymocytes. This may result in accumulation of the few leaky mature SP cells, which are the positively selected by H-Y CD8\(^+\) SP cells that are detectable in both H-\(2^{d_0}\) and \(\beta_2m^0\) NL/H-Y mice, thus forming a seemingly positively selected cell population. However, this is unlikely for at least three reasons. First, purified H-Y CD8\(^+\) SP cells from H-\(2^{b_0}\), H-\(2^{b_0}\), or \(\beta_2m^0\) female Egr/H-Y mice and H-\(2^{b_0}\) female NL/H-Y mice showed comparable survival curves...
When cells were simply cultured in vitro (data not shown; see Materials and Methods). Second, 105b2m0 recipient mice into which p2mEgr/H-Y bone marrow cells were transplanted did not harbor the significant H-Y CD8+ SP cell population that was seen in H-2d and p2m female Egr/H-Y mice, although their thymi contained detectable leaky H-Y CD8+ SP cells. Third, both male H-2b Egr/H-Y and NL/H-Y thymocytes exhibited clonal deletion of H-Y cells, indicating no rescue from apoptosis of Egr/H-Y cells undergoing negative selection. On the contrary, even more efficient negative selection appears to occur in Egr/H-Y mice. By these criteria, Egr-1 overexpression does not appear normal both in responses to various stimuli and in specificity of antigen-peptide recognition, as demonstrated by the functional analysis of mature H-Y CD8+ T cells. This is supported convincingly by the analysis of 5 × 105 B cell-depleted lymph node cells from either Egr-1 transgenic (TG) or negative littermate (NL) mice, both with an H-2d background, stimulated with 3,000-rad irradiated 5 × 106 spleen cells from b12m12 (H-2d) mice. (A) MLR responders were titrated B cell-depleted lymph node cells from transgenic (TG) or negative littermate (NL) mice, both with an H-2d background, stimulated with 3,000-rad irradiated 5 × 106 spleen cells from b12m12 (H-2d) or b12m0 (H-2b) female Egr/H-Y mice. (B) MLR was performed as above, with H-2b responder T cells and b6 (H-2b) stimulator cells. The smallest number of responder cells which 'H' thymidine incorporation was detected in transgenic T cells (envelope) and in negative littermate T cells (ellipse) in each panel is indicated. All experiments were repeated with four pairs of Egr-1 transgenic and negative littermate mice. In all panels, each value represents the mean of all results ± SD. (a) 5 × 105 B cell-depleted lymph node cells from transgenic (TG) and negative littermate (NL) mice were stimulated by 1 μg/ml of Con A for 72 h. Proliferation was assessed by duplicate thymidine incorporation for the last 16 h. Bars represent the mean of all results of three independent experiments ± SD.

An alternative argument is that Egr-1 overexpression itself might be sufficient to promote the DP to SP transition, with no interaction between thymocytes and ligands required. When the Egr-1 transgene was introduced into a RAG-deficient background, thymocytes did overcome the "maturation block at the CD25+CD44-DN stage and developed into immature CD8 single positive (ISP) cells without any signaling through the TCR (7). However, this was not the case for positive selection at the DP stage. As we demonstrated in a previous report, although irradiated Egr-1 transgenic mice on a RAG-deficient background developed DP cells expressing the Egr-1 transgene but lacking TCR expression, neither CD4+ nor CD8+ SP cells were observed (7). This suggests that DP cells do require some interaction with ligand(s) to proceed to the SP stage even in the presence of Egr-1 overexpression. This is supported convincingly by the analysis of 105b2m0 recipient mice into which p2mEgr/H-Y bone marrow cells were transplanted, in which no positive selection of H-Y CD8+ T cells occurred, as well as of Egr/AND mice on a class II MHC-deficient background, which harbored no AND CD4+ T cells. Hence, Egr-1 expression appears not to directly initiate transition of DP cells to the SP stage, but may influence the threshold of avidity required for positive selection.

Functions of T cells selected by unusually low avidity TCR-MHC interactions in the presence of Egr-1 overexpression appear normal both in responses to various stimuli and in specificity of antigen-peptide recognition, as demonstrated by the functional analysis of mature H-Y CD8+ T cells in H-2d and p2m Egr/H-Y mice. However, long-term survival of a naive population of peripheral H-Y CD8+ T cells in H-2d Egr/H-Y mice could be affected, as a recent report by Tanchot et al. (1996) implicated a requirement of the right MHC for survival of peripheral naive T cells. Therefore, a large proportion of peripheral H-Y CD8+ T cells in H-2d Egr/H-Y mice might be newly generated T cells. Tanchot et al. also demonstrated a requirement of only a nonspecific class II MHC for survival of peripheral memory T cells (16). This is consistent with the comparable expression patterns of memory markers such as Mel-14 in the peripheral H-Y CD8+ T cells in H-2d Egr/H-Y mice and H-2d NL/H-Y mice. Phenotype in Egr-1 transgenic mice indicates a physiological role of Egr-1 in determining the threshold required.
for positive selection. Perhaps related to this, Egr-1 overexpression may also affect negative selection. As described above, more efficient negative selection appeared to be achieved by Egr-1 overexpression in male H-2b β2m- H-Y mice. However, thymocytes express other members of Egr family genes, such as Egr-2, -3, and -4, all of which share a highly conserved DNA-binding domain (5, 6). These members might functionally compensate for each other, and hence no alteration of thymic selection is observed in Egr-1-deficient mice as discussed previously (7, 31). Therefore, a set of genes controlled by a common binding site for Egr genes, including Egr-1, might be critical in defining the overall thresholds required for thymic selection. Further characterization of Egr target genes should shed light on the precise molecular mechanism underlying the positive/negative selection of thymocytes.

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