Epidermal growth factor-like domain 8 inhibits the survival and proliferation of mouse thymocytes

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Abstract. Thymic epithelial cells (TECs) play a critical role in T-cell development through their intercellular interactions and by producing various soluble proteins, such as growth factors, cytokines and chemokines. In this study, we report a new role for epidermal growth factor-like domain 8 (EGFL8) in the regulation of the survival and proliferation of mouse thymocytes. Mouse recombinant EGFL8 (rEGFL8) protein was produced using an E. coli system and its biological role in mouse thymocytes was determined. The injection of rEGFL8 in mice in vivo resulted in a decrease in the weight of the thymus, as well as in the number of total thymocytes; rEGFL8 also inhibited thymocyte proliferation and induced thymocyte apoptosis. Furthermore, rEGFL8 suppressed the expression of the Notch downstream targets, Hes1 and Hey1, in mouse thymocytes and TECs, indicating that EGFL8 negatively regulates the Notch signaling pathway in these cells. The identification of the role of EGFL8 in thymocytes may aid in the determination of the fate of thymocytes during T-cell development.

Introduction

Epidermal growth factor-like domain (EGFL), an evolutionarily conserved protein domain, is found in many vertebrate proteins that are involved in several essential cellular activities, such as blood coagulation, fibrinolysis, cell adhesion and development (1). EGFL8, a newly identified member of the EGFL family, was originally identified as a paralog of EGFL7 by a BLAST search of the mouse genome (2). EGFL7 has recently emerged as a secreted angiogenic signaling molecule, which enhances vasculogenesis and angiogenesis by promoting endothelial cell adhesion, proliferation, chemotraction, migration, sprouting and invasion (3-7). Furthermore, EGFL7 stimulates embryonic stem cell proliferation, but inhibits the proliferation of adult neuronal stem cells, demonstrating its differential mechanisms of action during cell proliferation between different stem cell types (7,8).

However, little is known about the characterization and the biological role of EGFL8. A structural analysis of the EGFL8 protein predicted that it is a secretory protein (9). EGF-like repeats, located in the extracellular domains of EGFL8 and Notch receptors, play a central role in controlling the Notch signaling pathway (10). As recently demonstrated, EGFL8 expression was significantly decreased in patients with colorectal and gastric cancer, suggesting that EGFL8 may have a distinct expression pattern and mechanism of action in cancer progression (11,12).

We have previously demonstrated that the mouse EGFL8 gene plays a functional role in T-cell development in a gain-of-function and loss-of-function study with an EGFL8 gene overexpressing vector and EGFL8 siRNA (13). Based on these previous findings, in the present study, we investigated the functional role of the EGFL8 protein in mouse thymocytes and thymic epithelial cells (TECs), which are pivotal for both T-cell development and T-cell repertoire selection (14). To identify the potential importance of the EGFL8 protein, in this study, we designed and optimized a protocol to produce and purify large amounts of mouse recombinant EGFL8 (rEGFL8) protein. Using high-purity mouse rEGFL8, we demonstrate that EGFL8 inhibits the survival and proliferation of thymocytes during T-cell development. In addition, the mechanistic regulatory role of EGFL8 in mouse thymocytes and TECs was determined.

Materials and methods

Cell line and cell culture. The mouse thymic cortical epithelial reticular cells (1308.1) were kindly provided by Dr Barbara B. Knowles (The Jackson Laboratory) (15). The cells were cultured in Dulbecco's modified Eagle's medium (DMEM), containing 10% (v/v) fetal bovine serum (FBS), 100 U/ml penicillin and 100 µg/ml streptomycin (all from Gibco Life Technologies, Grand Island, NY, USA) at 37°C in 5% CO₂ incubator.

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Cloning of mouse EGFL8 gene. The coding sequences of mouse EGFL8 total cDNA were isolated and amplified by PCR and cloned into pCDNA3.1 (Life Technologies, Carlsbad, CA, USA). They were amplified by PCR using the following oligonucleotide primers: EGFL8 forward, 5'-TTT CAA AGA GAG TTT GGG AGT G-3' and reverse, 5'-CAC CAC GTG TT CTG TGG TA-3' to create the NcoI and XhoI restriction sites at the start and stop codon sites. The PCR product was cloned into the pET28α vector, which carries a C-terminal HisTag/thrombin/T7Tag configuration plus an optional C-terminal His-Tag sequence.

Expression of mouse EGFL8 gene. The E. coli bacteria were cultured for 2 h, stimulated by the addition of 0.2 mM IPTG, and then cultured for an additional 4 h. The bacteria were harvested immediately by centrifugation at 6,000 rpm for 8-10 min, and the pellet was either frozen at -80°C until purification or directly resuspended in 60 ml lysis buffer [50 mM Tris-HCl (pH 8.0), 100 mM NaCl and 5 mM EDTA]. Subsequently, 0.5% Triton X-100, 0.1 mM phenylmethylsulfonyl fluoride (PMSF) and 1 mM dithiothreitol (DTT) were added to this pellet. Subsequently, the pellet was sonicated and centrifuged at 12,000 rpm for 15 min. The supernatant was discarded and the pellet was resuspended in lysis buffer. The following steps were repeated as above except for the addition of 10 mM MgCl₂, 0.01 mg/ml DNase and 0.1 mg/ml lysozyme to the Triton X-100, PMSF and DTT mixture. The pellet mixture was incubated for 20 min at room temperature, sonicated and centrifuged as described above. The supernatant was discarded and the pellet was resuspended in 60 ml lysis buffer. Subsequently, only PMSF and DTT were added, sonicated and centrifuged as described above. The supernatant was discarded and the pellet was resuspended in 40 ml of 8 M urea [100 mM Tris-HCl (pH 8.0), 50 mM glycine]. The pellet was then slowly shaken for 1 h at room temperature to completely solubilize the proteins. After a short centrifugation to remove non-solubilized proteins, the protein concentration was determined by a NanoDrop 2000 spectrophotometer (Thermo Scientific, Wilmington, DE, USA). The proteins (1-2 mg/ml) were dialyzed through a dialysis bag. Two liters of dialysis buffers [20 mM Tris-HCl (pH 8.0), 150 mM NaCl and 0.1 mM DTT] were used at 4°C for 12 to 15 h and the process was repeated. The third dialysis buffer was used under the same conditions except for the absence of DTT.

Ni-NTA column purification. Purification of His-tagged proteins for the recombinant protein was then performed. The column was washed with distilled water and then with 6X Ni-NTA washing buffer. The protein mixture was added to the Ni-NTA column and slowly passed through the column. The bound EGFL8 protein was eluted with 6X Ni-NTA elution buffer. The eluted protein was subjected to sodium dodecyl sulfate-polyacrylamide gel electrophoresis (SDS-PAGE) after determining its concentration.

Mass spectrometry. The rEGFL8 protein was subjected to 10% SDS-PAGE, stained with Coomassie blue, and the protein bands were excised from the gel. The excised gel pieces were transferred to microcentrifuge tubes containing 0.5 ml of distilled water. The protein was then digested with trypsin. The single band was used for matrix-assisted laser desorption/ionization time-of-flight mass spectrometry (MALDI-TOF-MS/MS).

Experimental animals and treatment with rEGFL8 in vivo. C57BL/6 mice (Dae Han Bio Link, Chungbuk, Korea) were intravenously (i.v.) injected with rEGFL8-8 (100 µg) and were sacrificed 12 h, 1, 2 and 3 days after injection. Animal care and all experimental procedures were conducted in accordance with the ‘Guide for Animal Experiments’ published by the Korean Academy of Medical Sciences.

Proliferation and apoptosis assay. The mice were injected intraperitoneally (i.p.) with BrdU (BD Biosciences Pharmingen, San Diego, CA, USA) or with PBS. The thymus was isolated by dissection 2 h after BrdU injection. The isolated cells were stained for CD4, CD8, CD25, CD44 and BrdU according to the BrdU Flow kit manual (BD Biosciences Pharmingen). For the apoptosis assay, the isolated cells were stained for CD4, CD8, CD25, CD44 and Annexin V according to the Annexin V detection kit protocol (BD Biosciences Pharmingen).

Flow cytometric analysis. The following fluorochrome-conjugated monoclonal antibodies (mAbs) were purchased from BD Biosciences Pharmingen: Pacific Blue-conjugated anti-CD4 (RM4-5), aliphycocyanin (APC)-Cy7-conjugated anti-CD8 (53-6,7), PE-labeled anti-CD25 (PC61) and APC anti-CD44 (IM7). Flow cytometric analysis was performed using a FACScanto IIflow cytometer (BD Biosciences Pharmingen), and the acquired data were analyzed using the FlowJo software (Tree Star, Ashland, OR, USA).

Electron microscopy. For electron microscopic investigation, all specimens were processed according to a standard procedure. The ultra-thin sections were examined under a JEOL-1200 EXII transmission electron microscope.

Western blot analysis. The TECs were incubated with 100 ng/ml rEGFL8 for 12 h. The thymocytes were isolated from the mice injected with 100 µg of rEGFL8 i.v., after 12 h, 1 and 2 days. Following treatment with rEGFL8, the thymocytes and TECs were washed with cold PBS and the total proteins were then extracted from the cultured cells using a protein extraction solution (iNIRON Biotechnology, Seongnam, Korea) supplemented with a protease inhibitor mixture (Sigma-Aldrich, St. Louis, MO, USA). Protein concentrations were measured using the Bradford protein assay kit (Bio-Rad, Hercules, CA, USA). Anti-Hes-1 (sc-166378), anti-Hey-1 (sc-28746) (both from Santa Cruz Biotechnology, Santa Cruz, CA, USA) and anti-β-actin (Abcam, Cambridge, UK) antibodies were used for immunoblot analysis. The goat anti-mouse IgG-HRP (7076) and goat anti-rabbit IgG-HRP (7074) (both from Cell Signaling Technology, Danvers, MA, USA) were used as the secondary antibodies. Immunoreactivity was detected and quantified using a LAS-3000 imaging system (Fujifilm, Tokyo, Japan).

Statistical analysis. Data are expressed as the means ± SD. Statistical analyses were performed using the Student's t-test. A value of P<0.05 was considered to indicate a statistically significant difference.
Results

Expression and purification of mouse rEGFL8 protein. rEGFL8 was expressed in E. coli and purified. The harvested rEGFL8 protein was stained with Coomassie blue in 10% SDS-PAGE (Fig. 1A). We produced high-purity proteins after the third lysis step, suggesting that most of the secreted fusion proteins and most of the other bacterial proteins were removed by the lysis procedure. The dialysis and refolding steps were performed in the presence of urea, mainly as EGFL8 is prone to precipitation at high concentrations and is also prone to protein interactions with the exchange groups on the columns. However, to recover the biological activity of rEGFL8, urea needs to be removed and the protein should be refolded properly. The dialysis protein was analyzed on SDS-PAGE in comparison with cell lysates and supernatants to confirm the successful recovery of the recombinant protein after solubilization with 8 M urea. Fig. 1A illustrates an exact size of the desired protein for this study with no contaminated product from the E. coli cells. This result also verified that most or all of the inclusion bodies of rEGFL8 protein were solubilized in 8 M urea. The Ni-NTA-purified protein was loaded onto 10% SDS-PAGE for total protein analysis (Fig. 1B). The total Ni-NTA column-purified protein showed a single protein band with a specific size, which confirmed the successful rEGFL8 protein purification through the Ni-NTA column. The Ni-NTA-purified protein was also analyzed by western blot analysis for further confirmation (Fig. 1B). The western blot analysis result also showed a single and specific band of rEGFL8 protein. For further characterization, the rEGFL8 protein was run on 10% SDS-PAGE and the band was cut and trypsinized before loading onto the spectrometer (Fig. 2). All liquid chromatography (LC)-MS/MS data were submitted to a Mascot search; the peaks from the purified rEGFL8 protein sample are shown in Fig. 3, with the masses indicated on the top of each peak. The identified molecular weight of purified rEGFL8 measured by MS was 30.09 kDa, which was almost identical to the theoretical value.

Negative effects of rEGFL8 on the weight of the thymus and the number of thymocytes. To investigate the biological activity of rEGFL8 protein in mouse thymocytes, 100 µg of rEGFL8 protein were injected i.v. into the mice, and the weight of the thymus and the total number of thymocytes were determined (Fig. 4). The weight of the thymus and the number of thymocytes were significantly decreased after the rEGFL8 injection. The weight of the thymus and the number of thymocytes diminished from 1 day after rEGFL8 injection, and these effects reached a peak value at 3 days. The observation that the number of thymocytes markedly decreased after the administration of rEGFL8 raised the question of whether the subsets of thymocytes were differentially affected by treatment with rEGFL8. To address this issue, mouse thymocytes were freshly isolated 1, 2 and 3 days after rEGFL8 injection, and were stained with anti-CD4 and anti-CD8 mAbs (Fig. 5). Of note, there was a drastic decrease in both the percentage and number of CD4+CD8+ double-positive (DP) thymocytes 3 days after rEGFL8 injection (Figs. 5 and 6). Moreover, the CD4+ single-positive (SP) and CD8+ SP thymocyte subsets were also significantly reduced in number compared with the wild-type subset, although to a lesser degree than the CD4+CD8+ DP thymocytes on day 3 (Fig. 6).
Inhibitory effect of rEGFL8 on the survival and proliferation of thymocytes. The reduction in the number of thymocytes following treatment with rEGFL8 also raised the question of whether these changes occur due to the inhibition of cell proliferation or the promotion of apoptosis of thymocytes by rEGFL8. To shed light on this matter, the effect of rEGFL8 on...
cell proliferation was first investigated 12 h, 1 and 3 days after rEGFL8 injection. Freshly isolated thymocytes from the thymus 2 h after BrdU injection were stained with anti-CD4, anti-CD8, anti-CD25, anti-CD44 and anti-BrdU mAbs. The histogram shows BrdU+ thymocytes gated on the 4 major subsets of thymocytes (DN, DP, CD8 SP and CD4 SP) of rEGFL8-treated mice at the indicated time points. (B) The bar graph is a summary of the BrdU+ thymocyte frequency. All major thymocyte subsets, particularly at 12 h and 1 day after EGFL8 injection, exhibited a decrease in the percentage of BrdU+ cells compared with the control mice. Groups of mice were analyzed at each time point and data are presented as the means ± SD. *P<0.05. DN, double-negative; DP, double-positive; SP, single-positive.

To determine whether EGFL8 causes morphological changes in the mouse thymus at the ultrastructural level, transmission electron microscopy was performed after the injection of

Figure 8. (A) The effect of epidermal growth factor-like domain 8 (EGFL8) on cell survival was assessed by FACS analysis upon the administration of recombinant EGFL8 (rEGFL8) protein for 12 h, 1 and 3 days. Thymocytes from rEGFL8- or mock-treated mice were stained with the FITC-Annexin V and PI reagents. The histogram shows Annexin V+ thymocytes in the indicated thymocyte subpopulation. (B) The bar graph is a summary of the Annexin V+ thymocyte frequency. The percentage of apoptosis in the mouse thymocytes 12 h after EGFL8 injection significantly increased compared with the control thymocytes. Groups of mice were analyzed at each time point and data are presented as the means ± SD. *P<0.05.
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The thymus from the control mice showed normal ultrastructural features (Fig. 9A). However, electron micrographs of the thymus 12 h after the administration of rEGFL8 revealed apoptotic thymus cells, although their proportion was relatively small (Fig. 9B). Apoptotic cells (Fig. 9B and C) had typical morphological changes; the chromatins were condensed and aggregated into large dark, compact masses. Most of the apoptotic thymic cells appeared to be thymocytes, whereas the apoptosis of TECs and other thymic stromal cell types was rarely observed. In the thymus 1 day after the administration of rEGFL8, apoptotic thymocytes were observed to a much lesser extent than in at 12 h (Fig. 9C). Within 3 days after the rEGFL8 injection, apoptotic cells were rarely visible and the thymus exhibited an almost normal ultrastructural appearance (Fig. 9D).

Inhibitory effects of rEGFL8 on the expression of Hes1 and Hey1. To determine the effect of rEGFL8 on the Notch signaling pathway in mouse TECs, TECs were treated with rEGFL8 protein and the expression of Hes1 and Hey1 was then assessed by western blot analysis. Remarkably, the expression of Hes1 and Hey1 was downregulated in the TECs (Fig. 10A). To determine the effect of EGFL8 on the Notch signaling pathway in mouse thymocytes, rEGFL8 protein was injected into the mice, and the expression of Hes1 and Hey1 was assessed by western blot analysis. The expression of Hes1 and Hey1 was significantly downregulated in the mouse thymocytes (Fig. 10B). The inhibitory effect of rEGFL8 on Hes1 expression was more pronounced 12 h after rEGFL8 injection. The decrease in Hey1 expression was most evident 1 day following treatment with EGFL8.

Discussion

In the present study, to our knowledge, we present for the first time an optimized protocol for the production of high-purity mouse rEGFL8 protein; we also demonstrate the biological activity of the EGFL8 protein in mouse thymocytes. The biological activity of rEGFL8 in mouse thymocytes and TECs
was investigated as the modulation of the \textit{EGFL8} gene in TECs provided evidence of its negative regulatory role on the activity of TECs and the development of mouse thymocytes in our previous study (13). The \textit{in vivo} experiment with rEGFL8 demonstrated its inhibitory effect on the development of mouse thymocytes in a time-dependent manner in the present study. rEGFL8 also induced a decrease in the weight of the thymus, as well as in the number of mouse thymocytes primarily by the suppression of thymocyte proliferation as assessed by BrdU cell proliferation assay and, to a lesser extent, by the induction of apoptosis in thymocytes as revealed by flow cytometric analysis and electron microscopy. In a previous study, a differential equation model of thymocyte dynamics was constructed, which showed that cell proliferation, differentiation, and cell death in the thymus may account for both the total number of thymic cells and the fraction of various types of immature and mature thymocytes (16). According to this model, a decrease in the proliferation rate or an increase in the rate of apoptosis may be some of the parameters that account for the reduction of thymic size and thymocyte number that occurs due to thymic involution.

As regards the reduction in the number of thymocytes, the suppressive effect of rEGFL8 on thymocytes had no subset specificity, although the administration of rEGFL8 induced a profound decrease in the number of DP thymocytes. The reason why there was a marked reduction in the number of DP thymocytes compared with the other subsets may be related with their proportion and the absolute number in the control mice.

As regards the molecular mechanisms underlying the inhibitory effects of EGFL8 on the proliferation of mouse thymocytes and its promotion of apoptosis in mouse thymocytes, the involvement of EGFL8 in the Notch signaling pathway contributes, at least in part, to this negative regulatory role of rEGFL8, since it inhibited the expression of the Notch downstream effectors, Hes1 and Hey1, in the mouse thymocytes and TECs. Notch signaling is directly involved in the regulation of thymic T-cell development (17). Notch signaling has also been found to play a central role in reconstitution after transplantation, immunomodulation, or the development and maturation of thymocytes (18). Taken together, the data from the present study suggest that EGFL8 acts as a negative regulatory factor in the critical steps of mouse T-cell development, such as thymocyte proliferation and survival, through the inhibition of Notch signaling in mouse thymocytes and TECs.

In conclusion, the pET-28a-EGFL8 vector was expressed in \textit{E. coli} (DE3) and a relatively large amount of mouse rEGFL8 was successfully produced and purified in this study. Treatment with rEGFL8 reduced the weight of the thymus and the number of thymocytes, suppressed thymocyte proliferation, induced thymocyte apoptosis and inhibited Notch signaling (downregulation of Hes1 and Hey1 expression) in mouse thymocytes and TECs.

Therefore, the data from the present study suggest that EGFL8 acts as a negative regulatory factor in the critical steps of mouse T-cell development. Further studies are required to fully elucidate the functional role of EGFL8 in diverse physiological and pathological processes.

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