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Noncanonical K27-Linked Polyubiquitination of TIEG1 Regulates Foxp3 Expression and Tumor Growth

Dong-Jun Peng,*†‡ Minghui Zeng,*†‡ Ryuta Muromoto,§ Tadashi Matsuda,§ Kazuya Shimoda,¶ Malayannan Subramaniam,‖ Thomas C. Spelsberg,‖ Wei-Zen Wei,*†‡ and K. Venuprasad*†,‡

Earlier, we demonstrated the essential role of Kruppel-like transcription factor, TIEG1, in TGF-β-induced regulatory T cell (Treg) development. In this article, we demonstrate that IL-6, which promotes Th17 development, abrogated TIEG1 nuclear translocation and inhibited TGF-β–induced Treg development. Tyrosine kinase Tyk2-mediated phosphorylation of TIEG1 at Tyr179 promoted noncanonical K-27–linked polyubiquitination, which inhibited TIEG1 nuclear translocation. To test the role of TIEG1-regulated Treg/Th17 development in antitumor immunity, we analyzed TRAMP-C2 tumor growth in TIEG1−/− mice. The defective Treg development and elevated Th17 response resulted in enhanced immune reactivity in the tumor and inhibition of TRAMP-C2 tumor growth in TIEG1−/− mice. Thus, our results uncovered a novel regulatory mechanism that modulates Tregs and may regulate tumor progression. The Journal of Immunology, 2011, 186: 5638–5647.

Natural regulatory T cells (Tregs) develop from single-positive thymocytes during their maturation in the thymus and bear a diverse TCR repertoire against a broad range of self-Ags (1). In addition to thymus-derived natural Tregs, Foxp3+ Tregs are generated extrathyemically by the conversion of naïve CD4+ T cells by the action of TGF-β, known as induced regulatory T cells (iTregs) (2, 3). Because most tumor-associated Ags are aberrantly expressed self-Ags, Tregs play a critical role in suppressing antitumor immune response (4, 5). Furthermore, most malignant cells secrete large amounts of TGF-β (6–10), which was shown to convert effector T cells into tumor Ag-specific Tregs by inducing Foxp3 expression (6, 11–13). Such tumor-induced Tregs suppress the priming and effector functions of antitumor effector cells and form a broad self-amplifying immunosuppressive network (14). Therefore, overcoming TGF-β–induced expansion and de novo generation of Tregs is critical for the design of effective immunotherapeutic strategies for successful cancer treatment.

Although TGF-β is a potent anti-inflammatory cytokine that induces Foxp3 expression and Treg differentiation, in the presence of the proinflammatory cytokines IL-6, IL-21, and IL-23, CD4+ T cells differentiate into Th17 cells that can promote antitumor immunity (15–19). The intracellular signaling pathways that link TGF-β signaling to these diverse, and even opposing, T cell functions remain largely unclear. Whether Ag-stimulated CD4+ T cells differentiate into Foxp3+ Tregs or Th17 cells depends upon the cytokine-regulated balance between Foxp3 and RORγt. STAT3 is a crucial component of IL-6–mediated regulation of Th17 cells (20). Beyond these observations, the downstream mechanisms are not known.

We previously reported that Itch-mediated monoubiquitination is essential for TIEG1 nuclear translocation and Foxp3 expression. Interestingly, Itch also targets TIEG1 for polyubiquitination in the transient overexpression system (21). However, the physiological relevance of TIEG1 polyubiquitination is not known. Ubiquitin contains seven lysine residues, and its linkage to a substrate generally occurs via K48 or K63. K48-linked polyubiquitin predominantly targets proteins for proteasomal degradation, whereas K63-linked poly- and monoubiquitination regulate subcellular localization, protein function, or protein–protein interactions (22, 23). E3 ligases often target the same substrates differently for mono- or polyubiquitination under different physiological conditions. Such a phenomenon was reported for p53 and PTEN ubiquitination by MDM2 and Nedd4, respectively (24–26). However, the molecular signals and precise mechanisms that determine mono- versus polyubiquitination are not completely understood. In this article, we demonstrate that TGF-β and IL-6 differentially promote mono- and polyubiquitination of TIEG1 and regulate Treg/Th17 differentiation. The impact of TIEG1 on tumor immunity was investigated in TIEG1−/− mice. Growth of TRAMP-C2 tumor was reduced in TIEG1−/− mice and was accompanied by reduced Foxp3+ Tregs and an elevated Th17 response, suggesting that TIEG1 deficiency tilted the balance from suppressive toward effector immunity.

Materials and Methods

Mice

TIEG1−/− mice were described previously (27). Rag1−/− mice were purchased from The Jackson Laboratory (Bar Harbor, ME); C57BL/6 mice were purchased from Charles River Laboratory. All mice were housed in...
microisolator cages in the barrier facility of Karmanos Cancer Institute. All experiments were performed in accordance with the guidelines of the Institutional Animal Care and Use Committee of Wayne State University. Tyk2−/− mice were described previously (28) and were kept under specific pathogen-free conditions. All experiments were performed according to the guidelines of the Institutional Animal Care and Use Committee of Hokkaido University.

Reagents and Abs

Cytokines TGF-β1 and IL-6 were purchased from PeproTech. The following Abs were used for immunoblotting and immunoprecipitation: anti-KLF10/TIEG1 (ab73537; Abcam), anti-phospho-tyrosine (PY99; Santa Cruz), anti-His (GE Healthcare), rabbit anti-Tyk2 (C20; Santa Cruz), mouse anti-GST (B14; Santa Cruz), rabbit anti-hemagglutinin (HA) (Y-11; Santa Cruz), anti–c-Myc (9E10; Santa Cruz), anti-actin (AC-15; Sigma), anti-histone H3 (Biolegend), and anti-Xpress (Invitrogen). Ni+-NTA beads were purchased from Qiagen.

Isolation of CD4+CD25+ T cells

CD4+CD25+ T cells from mouse spleen and lymph node were isolated using the CD4+CD25+ T Cell Isolation Kit (Miltenyi Biotec), according to manufacturer’s protocol. T cells were cultured in RPMI 1640 medium supplemented with 10% FBS and stimulated with 2 μg/ml plate-bound anti-CD3 (eBioscience) and 2 μg/ml soluble anti-CD28 (Biolegend).

Flow cytometry

Surface and intracellular staining were performed using the Mouse Regu- latory T Cell Staining Kit (88-8111; eBioscience), according to the manufacturer’s protocol. Briefly, spleen and lymph node cells were sus- pended in Flow Staining Buffer with 0.5 μl anti–CD4-FITC and incubated at 4°C for 30 min. Cells were fixed, permeabilized, subjected to in- tracellular staining with anti–Foxp3-PE or negative control Abs by in- cubation at 4°C for 30 min. Cells were fixed, permeabilized, subjected to in- 3

Quantitative RT-PCR

Total RNA was extracted using an RNeasy Mini kit (Qiagen), according to the manufacturer’s instructions, and reverse transcribed into cDNA using a Verso cDNA Synthesis kit (Thermo Scientific). Real-time PCR was performed using a Mastercycler reagentplex system (Eppendorf) and Light- Cycler 480 SYBR Green I Master kit (Roche). The abundance of mRNA was normalized to that of actin mRNA. The following primers were used: Foxp3 primers, 5′-CCATCCCAGAGGTCTTGG-3′ and 5′-ACCATGAC- TAGGGGACCTGA-3′; actin primers, 5′-GAAATCTGCCTGACATCCC- AAAG-3′ and 5′-TGTGGTTTCATGATGCACAG-3′; IFN-γ primers, 5′-GAAGCTGCCAAGAGGTGTA-3′ and 5′-TGGTGTTTGCACCTCAAA-3′ and IL-17 primers, 5′-TTTACCTCCTTGCGGCAA-3′ and 5′-CTTCTCCCTCCATGACAC-3′.

Plasmid construction and cell transfection

Mouse TIEG1 cDNA (Open Biosystems) was cloned into pEFe4-His vector (Invitrogen) to generate plasmid-encoding Hisx4-Xpress-tagged TIEG1. The truncated TIEG1 was amplified by PCR and then cloned into EcoRI/XbaI sites of pEFe4-His B vector (Invitrogen). The tyrosine residue mutants of TIEG1 were generated using a site-directed mutagenesis kit (Stratagene). Myc-Ich plasmid was a kind gift from Dr. Francesca Bernassola (Department of Experimental Medicine and Biocbchemical Sciences, University of Rome). Flag-Ty2 and Flag-Tyk2/2930I–encoding plasmids were constructed by subcloning the Tyk2 or Tyk2/2930I fragment into the EcoRI site of pCMV-Tag 2A vector. Plasmids encoding HA-tagged ubiquitin and ubiqui- tinyl lysine mutants were obtained from Addgene (29), with the exception of ubiquitin mutants of K6, K11, K27, K29, and K27R, which were generated using a site-directed mutagenesis kit. 293T cells were cultured in DMEM medium supplemented with 10% FBS, 100 μ/ml penicillin, and 100 μg/ml streptomycin. Cell transfection was performed using Lipo- fectamine 2000 (Invitrogen), according to the manufacturer’s instruction.

Immunoprecipitation and Western blotting

Cell lysates were prepared in Nonidet P-40 (NP-40) lysis buffer (150 mM NaCl, 50 mM Tris-Cl [pH 8] and 1% NP-40 with the addition of protease inhibitor mixture [Roche] and 1 mM vanadate). Cell lysates were incubated with 1–2 μg Ab at 4°C for ~3 h, followed by the addition of 25 μl protein A/G plus agarose (Santa Cruz Biotechnology) for an additional hour at 4°C. The immunoprecipitates were washed five times with NP-40 lysis buffer and analyzed by Western blotting with appropri- ate Abs. Briefly, proteins were electrophoresed by 10–12% denaturing polyacrylamide gels and transferred to polyvinylidene difluoro- mem- brane (Millipore). The membranes were probed with primary Abs, washed, and incubated with appropriate HRP-conjugated secondary Abs. The membranes were visualized by an ECL reagent (Amsersham Pharmacia Biotech). Membranes were stripped by incubation for 30 min at 65°C in stripping buffer and then probed with other Abs (21). In some experiments, cytoplasmic and nuclear fractions were prepared from total-cell lysates using Paris Protein and a RNA Isolation kit (Ambion).

Ubiquitination assay

Forty-eight hours after transfection with the indicated plasmids, 293T cells were treated with 10 μM proteasome inhibitor MG132 (Calbiochem) for 6 h, harvested, and lysed in NP-40 lysis buffer. Hisx4-TIEG1 was isolated by the Ni-NTA affinity-purification method, according to a published protocol (30). Briefly, cell lysates were diluted in 1 ml guanidine buffer (6 M guanidine-HCl, 0.1 M NaHPO4/NaH2PO4 [pH 8]) with additions of 10 mM imidazole and 25 μl Ni-NTA agarose (Qiagen). The mixtures were incubated at 4°C for 3 h. The precipitated proteins were subsequently analyzed by immunoblotting with appropriate Abs.

GST pull-down assay

GST-TIEG1, GST-Ty2, and GST-Ty2/K930I fusion proteins were puri- fied from BL21 (DE3) by glutathione Sepharose beads (Amsersham Pharmac- ia Biotech), according to the manufacturer’s instructions. The purified GST, GST-TIEG1, GST-Ty2, and GST-Ty2/K930I proteins bound to Glutathione Sepharose beads were incubated with cell lysates at 4°C for 2–3 h, washed five times with PBS, and subjected to immunoblot analy- sis.

Retroviral construction and transduction

Flag-TIEG1 and Flag-TIEG1/Y179F cDNA amplified by PCR were cloned into retroviral vector MigR1. The retroviral particles were produced by Plat- E packaging cell line transiently transfected with retroviral constructs. Retroviral supernatants were collected 48 h after transfection. naïve CD4+CD25+ T cells from TIEG1−/− mice were stimulated with 2 μg/ml anti-CD3 (eBioscience) and 2 μg/ml soluble anti-CD28 (Biolegend) for 24 h and infected by adding 1:2 volume of retroviral supernatants supplemented with polynucleotide (6 μg/ml; Sigma), followed by centrifugation at 3000 rpm for 1 h at 32°C. Retroviral infection was repeated once after 24 h after the initial infection. T cells were maintained in complete RPMI 1640 medium with 100 μ/ml IL-2.

Conditioned medium

TRAMP-C2 cells were cultured in RPMI 1640 medium supplemented with 10% FBS, 5 μg/ml insulin (i6634; Sigma), 100 μ/ml penicillin, and 100 μg/ml streptomycin. The conditioned medium (CM) was collected from 3-d culture of TRAMP-C2 cells and prepared by passing through a 0.45-μm filter. Isolated CD4+CD25+ T cells were incubated with T cells medium (RPMI 1640 supplemented with 10% FBS) or CM in the presence of 2 μg/ml plate-bound anti-CD3 and soluble anti-CD28. T cells were collected and analyzed by quantitative PCR, as described above.

In vivo tumor growth

TRAMP-C2 cells were cultured in RPMI 1640 medium supplemented with 10% FBS, 5 μg/ml insulin (i6634; Sigma), 100 μ/ml penicillin, and 100 μg/ml streptomycin. LLC cells were cultured in DMEM medium supplemented with 10% FBS, 100 μ/ml penicillin, and 100 μg/ml streptomycin. A total of 0.5–5 × 106 TRAMP-C2 cells or 1 × 106 LLC cells were injected s.c. into 8–12-wk-old male TIEG1−/− mice or age-matched male C57BL/6 mice. Tumor growth was monitored thrice a week by measuring tumor length and width. The volume of the tumor was estimated by the formula (length × width)3/2.

Adaptive transfer of TIEG1−/− T cells into Rag1−/− mice

Naïve CD4+CD25− T cells (3 × 106) and CD8 T cells (1 × 106) isolated from TIEG1−/− and TIEG1−/− mice were adoptively transferred into Rag1−/− mice by i.v. injection 3 d after TRAMP-C2 tumor cell inoculation. Tumor growth was monitored thrice a week.
Results

IL-6 negatively regulates TIEG1 nuclear translocation via Tyk2-mediated phosphorylation

Because TGF-β and IL-6 reciprocally regulate Treg and Th17 differentiation (16, 31–34), and TGF-β–induced TIEG1 nuclear translocation is essential for Foxp3 expression (21), we tested whether IL-6 negatively regulates TIEG1 to promote Th17 differentiation. Naive CD4+ T cells were stimulated with anti-CD3 + anti-CD28 in the presence of TGF-β alone or in combination with IL-6. To analyze the cellular distribution of TIEG1, we separated nuclear and cytoplasmic fractions and immunoblotted these fractions with anti-TIEG1 Ab. Consistent with our previous report (21), TGF-β treatment resulted in nuclear translocation of TIEG1; however, TIEG1 was predominantly located in the cytoplasm of the cells stimulated with IL-6 (Fig. 1A). This finding suggested that IL-6 inhibits nuclear localization of TIEG1 as a negative-regulatory mechanism to promote Th17. Because IL-6 stimulation induces tyrosine phosphorylation of several intracytoplasmic proteins, we tested whether TIEG1 is also tyrosine phosphorylated. As shown in Fig. 1B, TIEG1 was tyrosine phosphorylated in naive CD4+ T cells stimulated in the presence of TGF-β and IL-6 but not TGF-β alone, suggesting a critical role for phosphorylation-dependent cytoplasmic-to-nuclear shuttling of TIEG1 in Treg/Th17 differentiation. 

Next, we sought to identify the upstream kinase that induced TIEG1 phosphorylation. Tyk2, a nonreceptor tyrosine kinase belonging to the Jak family, is activated when IL-6 binds to its receptors (35, 36). A Tyk2 mutation is associated with hyper-IgE syndrome (37), and T cells from hyper-IgE syndrome patients failed to differentiate into Th17 cells (38). Defective IL-17 expression was also reported in γδ T cells from Tyk2−/− mice (39). In addition, a natural mutation in the pseudokinase domain of Tyk2 accounted for increased susceptibility to infection and resistance to collagen-induced arthritis in B10.Q/J mice (40, 41). All of these suggest a possible link between the IL-6–Tyk2 pathway in Treg and Th17 differentiation. We hypothesized that, upon binding of IL-6 to its receptor, activated Tyk2 phosphorylates TIEG1. To test this hypothesis, we transiently transfected 293T cells with plasmids encoding wild-type (WT) Tyk2 or kinase-deficient mutant (K930I) Tyk2 and TIEG1. After 24 h, cells were treated with a phosphatase inhibitor (pervanadate) for 20 min. Cell extracts were immunoprecipitated with anti-His and immunoblotted with antiphosphotyrosine to analyze TIEG1 tyrosine phosphorylation. As shown in Fig. 1C, WT Tyk2 induced TIEG1 phosphorylation but not the kinase mutant. To further confirm the specificity of Tyk2 as the kinase, we stimulated naive CD4 T cells from Tyk2+/+ and Tyk2−/− mice (28) with anti-CD3 and anti-CD28 in the presence of TGF-β alone or in combination with IL-6. TIEG1 was immunoprecipitated using anti-TIEG1 Ab and immunoblotted with anti–p-tyrosine Ab. As shown in Fig. 1D, stimulation of Tyk2+/+ CD4 T cells with IL-6 resulted in TIEG1 phosphorylation but not in Tyk2−/− cells. These data strongly supported that Tyk2 phosphorylates TIEG1.

To understand the molecular mechanism underlying Tyk2-mediated TIEG1 tyrosine phosphorylation, we examined whether Tyk2 physically associates with TIEG1. TIEG1 was coimmunoprecipitated with anti-Tyk2 Ab, or vice versa, in transiently transfected 293T cells (Supplemental Fig. 1A). Further, we generated

**FIGURE 1.** IL-6–Tyk2 pathway-mediated phosphorylation regulates TIEG1 nuclear localization and Treg/Th17 differentiation. A, Naive CD4+CD25− T cells were stimulated with anti-CD3 and anti-CD28 in the presence of TGF-β (5 ng/ml) alone or TGF-β (5 ng/ml) plus IL-6 (20 ng/ml). The nuclear and cytoplasmic fractions were immunoblotted with anti-TIEG1 Ab. The membranes were reprobed with anti-histone H3 and anti-actin Ab. C, cytoplasmic; N, nuclear. B, CD4+CD25− T cells were treated as in A. Endogenous TIEG1 from total-cell lysates was immunoprecipitated with anti-TIEG1 Ab and immunoblotted with anti–phosphotyrosine Ab. Blots were reprobed using anti-TIEG1 Ab, p-TIEG1, or tyrosine-phosphorylated TIEG1. As shown in Fig. 1C, WT Tyk2 induced TIEG1 phosphorylation but not TIEG1 tyrosine phosphorylation. Because TGF-β induces TIEG1 phosphorylation in naive CD4+ T cells, we stimulated naive CD4 T cells from Tyk2+/+ and Tyk2−/− mice (28) and TIEG1 was immunoprecipitated with anti-His Ab and immunoblotted with antiphosphotyrosine to analyze TIEG1 tyrosine phosphorylation. As shown in Fig. 1C, WT Tyk2 induced TIEG1 phosphorylation but not the kinase mutant. To further confirm the specificity of Tyk2 as the kinase, we stimulated naive CD4 T cells from Tyk2+/+ and Tyk2−/− mice (28) with anti-CD3 and anti-CD28 in the presence of TGF-β alone or in combination with IL-6. TIEG1 was immunoprecipitated using anti-TIEG1 Ab and immunoblotted with anti–p-tyrosine Ab. As shown in Fig. 1D, stimulation of Tyk2+/+ CD4 T cells with IL-6 resulted in TIEG1 phosphorylation but not in Tyk2−/− cells. These data strongly supported that Tyk2 phosphorylates TIEG1.

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GST-TIEG1 and GST-Tyk2 fusion protein and performed pull-down assays. GST-TIEG1 precipitated Tyk2 from the cellular lysate of 293T cells transiently transfected with Tyk2; similarly, GST-Tyk2 precipitated TIEG1 (Supplemental Fig. 1B). To test whether the endogenously expressed TIEG1 and Tyk2 interact in naïve CD4 T cells, we performed coimmunoprecipitation experiments using anti-TIEG1 and anti-Tyk2 Abs. As expected, anti-TIEG1 Ab coprecipitated Tyk2, and vice versa, in cells stimulated with IL-6 but not TGF-β alone (Fig. 1E).

TIEG1 phosphorylation at Y179 inhibits TIEG1 and Foxp3 expression

To identify the tyrosine residue that undergoes phosphorylation, we generated deletion mutants of TIEG1, as shown in Fig. 2A. Cotransfection with Tyk2 and the constructs encoding full-length TIEG1 or deletion mutants (1–220, and 1–370) resulted in TIEG1 phosphorylation but not deletion mutants 1–90, 1–160, 231–479, and 181–479 of TIEG1 (Fig. 2B). This result indicated that only tyrosine 179 is phosphorylated by Tyk2. To further confirm this result, we generated Y179F mutant TIEG1 (in which tyrosine 179 was mutated to phenylalanine). Cotransfection of TIEG1-Y179F with Tyk2 in 293T cells almost completely abolished phosphorylation (Fig. 2C). These results showed that Tyk2 phosphorylates TIEG1 at tyrosine 179. To understand the physiological relevance of these findings, we cloned TIEG1-Y179F mutant into MigR1-GFP retroviral vectors. We transduced TIEG1−/− naïve CD4+ CD25− cells with WT or TIEG1-Y179F mutant by retroviral transduction. Nuclear/cytoplasmic localization of TIEG1 was analyzed in FACS-sorted GFP+ cells following treatment with TGF-β1+ IL-6 (Th17-inducing) conditions for 7 d. The effect of TIEG1-Y179F on Foxp3 versus IL-17 expression was analyzed by real-time PCR. The results shown are from one representative experiment that was conducted three times.

FIGURE 3. Tyk2-mediated phosphorylation regulates TIEG1 nuclear localization and Treg/Th17 differentiation. A, Naïve TIEG1−/− CD4+ CD25− cells were reconstituted with WT or TIEG1-Y179F mutant by retroviral transduction. Nuclear/cytoplasmic localization of TIEG1 was analyzed in FAC-sorted GFP+ cells following treatment with TGF-β or TGF-β+ IL-6. B, GFP+ cells were stimulated with TGF-β (Treg-inducing) and TGF-β+ IL-6 (Th17-inducing) conditions for 7 d. The effect of Y179F mutation on Foxp3 versus IL-17 expression was analyzed by real-time PCR. The results shown are from one representative experiment that was conducted three times.
ubiquitination, other forms of polyubiquitination are more common than originally thought (43–46). Therefore, we used UbK6, UbK11, UbK27, UbK29, and UbK33 to identify the topology of TIEG1 polyubiquitination. Interestingly, coexpression of UbK27 resulted in TIEG1 polyubiquitination. We consistently observed moderate polyubiquitination when UbK29 was coexpressed but not with UbK6, UbK11, UbK33, UbK48, or UbK63 (Fig. 4D).

This suggested that Tyk2 promotes predominantly K27-linked polyubiquitination of TIEG1. To further confirm K27-linked polyubiquitination of TIEG1, we generated UbK27R (in which lysine residue at K27 of Ub was mutated to arginine). Coexpression of UbK27R with Itch, Tyk2, and TIEG1 resulted in markedly reduced TIEG1 polyubiquitination (Fig. 4E). To confirm that Tyk2-mediated phosphorylation is essential for K27-linked polyubiquitination, we coexpressed WT and Y179F mutant of TIEG1 with Itch, Tyk2, and UbK27. As shown in Fig. 4F, there was a marked defect in the polyubiquitination of Y179F mutant compared with WT TIEG1. These results collectively suggested that Tyk2-mediated phosphorylation of TIEG1 promotes K27-linked polyubiquitination of TIEG1. Based on our results, we propose that when the naive CD4 T cells are stimulated in the presence of TGF-β, Itch targets TIEG1 for monoubiquitination. Monoubiquitinated TIEG1 translocates to the nucleus and binds to Foxp3 promoter to induce Foxp3 expression (21), whereas in the presence of IL-6, Tyk2 phosphorylates TIEG1, and Itch targets phosphorylated TIEG1 for polyubiquitination and prevent its nuclear translocation. This may divert Foxp3 transcription and promote Th17 differentiation (Supplemental Fig. 2).

Defective iTreg development and elevated Th17 response attenuate TRAMP-C2 tumor growth in TIEG1−/− mice

Tyk2/Treg was shown to play a critical role in tumor immunity and immunotherapy (47). In hosts with large advanced tumors, Treg induction and expansion become dominant and result in poor effector T cell activation and tumor rejection (48–50). Phenotypic analysis of prostate-infiltrating lymphocytes revealed Th17/Treg skewing and an inverse correlation between Th17 cells and tumor progression (51). Moreover, vaccination with hsp70 induced IL-6 production in the prostate tissue, which triggered a Th17 response, resulting in rejection of established prostate tumors (52). All of these findings suggested an intimate connection between Treg/Th17 balance and immune surveillance. We previously demonstrated that TIEG1−/− CD4+ T cells were resistant to TGF-β-
mediated suppression and defective in Foxp3 expression upon TGF-β treatment in vitro (21). Another study reported enhanced IL-17 and IFN-γ production by TIEG1−/− T cells (53). Therefore, we hypothesized that a reduced Treg and an elevated Th17 response in TIEG1−/− mice would render enhanced antitumor immunity. This hypothesis was tested with prostate cancer TRAMP-C2 cells, which secrete large amounts of TGF-β. First, we tested whether TIEG1−/− CD4+CD25− cells are refractory to TRAMP-C2-derived TGF-β. Naive CD4+CD25− cells from TIEG1+/+ and TIEG1−/− mice were cultured with TRAMP-C2 CM. The cells were stimulated with anti-CD3 and anti-CD28 for 3–5 d (Fig. 5A). The effect of TRAMP-C2–derived TGF-β on the conversion of CD4+CD25− cells into Foxp3+ cells was compared by real-time PCR, using the total RNA isolated from these cells. As expected, TIEG1+/+ cells expressed Foxp3, but the level of Foxp3 induction was markedly lower in TIEG1−/− cells (Fig. 5B). To confirm that TGF-β derived from the TRAMP-C2 cells induced Foxp3 expression, we neutralized TGF-β in the supernatant using anti-TGF-β Ab. As expected, anti–TGF-β Ab inhibited Foxp3 expression in naive CD4 T cells, but the control isotype Ab did not (Fig. 5C). When IL-6 was added to TRAMP-C2 CM, TIEG1−/− T cells produced significantly higher levels of IL-17 compared with TIEG1+/+ cells (Fig. 5D).

To test the impact of TIEG1 deficiency on tumor growth, we inoculated (0.5 × 106–5 × 106) TRAMP-C2 cells s.c. in the flanks of TIEG1+/+ and TIEG1−/− mice. Tumor growth was monitored thrice a week for 60 d. As shown in Fig. 6A, a majority of the TIEG1−/− mice rejected TRAMP-C2 tumors; even when the tumors developed, their sizes were smaller (Fig. 6B). Because we demonstrated the essential role of TIEG1 in Treg development, we analyzed Foxp3+ Tregs in the spleen and draining lymph nodes of these mice by intracellular staining using anti-Foxp3 Ab. As shown in Fig. 6C and 6D, the percentage of CD4+Foxp3+ cells was markedly increased in the tumor-bearing TIEG1+/+ mice; however, in the TIEG1−/− mice, the percentage of Tregs remained similar to that in the naive mice. Similarly, we noticed markedly reduced Foxp3 expression in the tumor mass collected from TIEG1−/− mice compared with TIEG1+/+ mice. This suggested that the TIEG1 deficiency may affect the expansion and de novo generation of tumor-specific Tregs. Because tumor-specific Tregs actively suppress effector T cell responses, we tested whether defective Treg development in TIEG1−/− mice correlated with elevated immune reactivity in the tumor. As expected, we found elevated levels of IFN-γ and IL-17 in the tumor-infiltrated cells in TIEG1−/− mice (Fig. 6E) compared with TIEG1+/+ mice. Because we found elevated IL-17 in TIEG1−/− tumors, we analyzed the expression of IL-6 transcripts in the RNA isolated from the whole tumor tissue. We detected IL-6 mRNA in the tumors from TIEG1+/+ and TIEG1−/− mice (data not shown). This suggested that in the TRAMP-C2 tumor microenvironment there is an abundance of Treg-promoting TGF-β and Th17-promoting inflammatory IL-6. Because high levels of TGF-β produced by TRAMP-C2 cells induced Treg development and inhibited Th17 (54) in TIEG1+/+ tumor-bearing mice, Treg development may predominate, and TIEG1 deficiency may tilt the balance toward Th17 and antitumor immunity. To further test whether an enhanced antitumor immune response is unique to TRAMP-C2, we inoculated TIEG1+/+ and TIEG1−/− mice with LLC cells. As shown in Supplemental Fig. 3, reduced LLC tumor growth was observed in TIEG1−/− mice. These data collectively suggested that defective Foxp3 expression and elevated Th17 responses in TIEG1−/− mice result in robust antitumor immune response and tumor rejection.

To confirm that TIEG1 deficiency regulates de novo generation of Tregs/Th17 cells in the tumor microenvironment, we set up an adoptive-transfer model. In this model, Rag1−/− mice were injected with TRAMP-C2 cells, followed by infusion of CD25-depleted CD4 (3 × 105) and CD8 T cells (1 × 105) from naive TIEG1+/+ and TIEG1−/− donors. In the WT environment, tumors grew progressively; however, tumor growth was markedly reduced and delayed in Rag1−/− mice that received TIEG1−/− T cells (Fig. 7A, 7B). Analysis of Foxp3 expression in tumor-infiltrating

**FIGURE 5.** Defective TGF-β induced Treg development in TIEG1−/− naive CD4 T cells. A, Experimental procedures. TRAMP-C2 cells were cultured for 3 d without change of the medium. CM was collected and incubated with naive CD4+CD25− T cells isolated from spleen and lymph nodes of TIEG1+/+ and TIEG1−/− mice in the presence of anti-CD3 and anti-CD28. After 3–5 d of culture with CM, T cells were harvested for Foxp3 analysis. B, Real-time PCR analysis of Foxp3 expression. CD4+CD25− T cells were stimulated with or without CM as in A, and Foxp3 expression level was analyzed by real-time PCR. The increased folds of Foxp3 mRNA were normalized to that of actin mRNA. C, In a similar experiment to B, anti-TGF-β (15 μg/ml) or control isotype Ab was added to the cultures. Foxp3 mRNA was analyzed by real-time PCR. D, In a similar experiment to A, IL-6 was added to the T cell cultures. IL-17 mRNA was analyzed by real-time PCR.
lymphocytes by intracellular staining revealed that a significant number of TIEG1+/+ donor CD4+CD25+ cells was converted into Foxp3+ cells. However, we detected only a few Foxp3+ cells in Rag1−/− mice that received CD25-depleted TIEG1+/−/− T cells (Fig. 7C). As expected, we also observed markedly increased numbers of IFN-γ+ and IL-17+ cells in the Rag1−/− mice that received TIEG1+/−/− T cells (Fig. 7D). These results showed that iTregs generated in the tumor microenvironment played a significant role in the antitumor immune response.

Discussion

Although protein ubiquitination has been classically considered as a death signal, recent studies convincingly demonstrated that it also represents a means of protein modification, affecting protein–protein interaction, phosphorylation, and subcellular localization (55). K48-linked polyubiquitin chains are the principal signals for targeting substrates for degradation by the 26S proteasome, whereas K63-linked chains act in a range of processes, including protein trafficking, DNA repair, and inflammation. In addition, endogenous K6, K11, K27, K29, or K33-linked polyubiquitination of protein substrates has been reported; however, their function remains largely nebulous (44, 56). In this study, we demonstrated that Itch differentially targets TIEG1 for mono- and polyubiquitination and regulates its nuclear-to-cytoplasmic shuttling during Treg/Th17 differentiation. Interestingly, TIEG1 polyubiquitination was K27 linked, and whose function remains largely unknown. The E3 ligase TRAF6 was reported to promote K6, K27, and K29 ubiquitination of the Parkinson’s disease proteins DJ-1 and α-synuclein, resulting in their accumulation in the cytoplasmic aggregates (57). K27-linked polyubiquitination was also shown to promote lysosomal localization of Jun (58). Our initial analysis revealed a bipartite nuclear localization signal (59–61) within the second zinc finger domain of TIEG1 (K. Venuprasad, unpublished observations), and K27-linked ubiquitination may block this nuclear localization signal. Another important observation of our study was that Tyk2, which is activated by IL-6, phosphorylates TIEG1 and acts as a recognition signal for K27-linked polyubiquitination. Previous studies showed that phosphorylation can create recognition-signal phosphodegrons for binding of an E3 ligase to the substrate (42, 62). Phosphorylation can also result in the exposure of degrons by inducing conformational change, although the phosphate itself does not directly contribute to recognition on the degron. In addition, phosphorylation can regulate the access of an E3 ligase to its targets via...
phosphorylation-dependent transport of the substrate or the ligase between cellular compartments (42). It is not clear how TIEG1 phosphorylation results in K27-linked polyubiquitination, and it requires further detailed investigation.

Another important finding of our study was that inoculation of TRAMP-C2 cells into TIEG1−/− mice resulted in tumor rejection. Also, reduced Foxp3+ Tregs and elevated IFN-γ and IL-17 by tumor-infiltrated cells suggested a robust immune reactivity against TRAMP-C2 cells in TIEG1−/− mice. Again, our adoptive-transfer studies using Rag1−/− mice clearly suggested that TIEG1−/− T cells failed to convert into Foxp3+ Tregs, resulting in elevated Th17 responses and reduced tumor growth, which suggested a critical role for TIEG1 in immune surveillance. However, TIEG1 was shown to regulate transcription of TGF-β itself by binding to the consensus sequences on the TGF-β promoter (53). In addition to defective Treg development and elevated Th17 responses, TIEG1 may regulate additional mechanisms that contribute to the observed antitumor effects in TIEG1−/− mice.

The expansion and de novo generation of tumor-specific Tregs has emerged as a major obstacle in successful immunotherapy against tumors (11). Especially in hosts bearing large advanced tumors, Treg induction and expansion become dominant, resulting in poor effector T cell activation and tumor rejection (48–50). For example, immunizing tumor-bearing mice with vaccine carrying a tumor Ag elicited tumor Ag-specific effector T cells and Tregs (48). Similarly, clinical investigations with human papilloma virus E6/E7 vaccines in cervical cancer patients provide further support for the concomitant induction of tumor-specific Tregs (50). Using the TRAMP-C2 prostate cancer cell line, which secretes high levels of TGF-β (7), it was demonstrated that tumor-secreted TGF-β converted the tumor Ag-specific T cells into Foxp3+ Tregs (6). Such Tregs inhibit the priming and effector function of antitumor effector cells and form a self-amplifying immune-suppressive network through its interaction with various APCs in the tumor microenvironment (14). Therefore, developing a vaccination strategy to increase the frequency of tumor-specific effector T cells, as well as to prevent Treg induction, is essential for successful treatment. Depleting Tregs using anti-CD25 Ab was demonstrated to enable the rejection of several types of transplantable tumors. Because Tregs are required for maintaining self-tolerance, prolonged, global, and nonspecific depletion of Tregs may breach self-tolerance (11, 63). Therefore, strategies are needed to selectively block the development of Tregs in the tumor site. It will be advantageous to eliminate suppressors while redirecting the precursor T cells to differentiate into effector cells, such as Th1 or Th17 cells (23). We describe a phosphorylation-dependent regulation of Treg/Th17 development by TIEG1. TIEG1 deficiency resulted in elevated IL-17 expression and reduced TRAMP-C2 tumor growth. Because higher levels of TGF-β promote the development of Tregs and suppress IL-6–mediated Th17 cell differentiation (54, 64, 65), it is likely that, in the absence of TIEG1, the effect of high concentrations of TGF-β in the tumor microenvironment is nullified, resulting in elevated IL-17 expression. The role of IL-17 in tumor immunopathology has been controversial. Overexpression of IL-17 in some tumor cell lines promotes angiogenesis and tumor growth, suggesting a protumor activity (66). However, transgenic T cells polarized to a Th17 phenotype by treatment with TGF-β and IL-6 were shown to eradicate tumors (17–19). A significant inverse correlation between Th17 and tumor progression was reported in patients with prostate cancer (51). Therefore, Th17 cells provide protection against tumors in certain cases, as we observed in TIEG1-deficient mice. A better understanding of the molecular regulation of Treg/Th17 conversion could help to improve cancer immunotherapy strategies.
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Disclosures
The authors have no financial conflicts of interest.

References

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Supplementary Figure Legend

Supplementary Figure 1. TIEG1 physically interacts with Tyk2. A, 293T cells were transfected with plasmids encoding His-TIEG1, Flag-Tyk2, or their combination for 48 h. Cells were lysed and immunoprecipitated (IP) with anti-His or anti-Flag, and analyzed by immunoblotting with anti-Flag or anti-TIEG1. The amount of Flag-Tyk2 and His-TIEG1 in the whole cell lysates was shown in lower panel. B, GST, GST-Tyk2 or GST-TIEG1 were incubated for 2-3 h at 4°C with the lysates of 293T cells transfected with His-TIEG1 (left) or Tyk2 (right). The precipitants were analyzed by immunoblotting with anti-TIEG1 or anti-Tyk2 respectively.

Supplementary Figure 2. Proposed model of TIEG1 ubiquitination in Treg/Th17 differentiation. TGF-β promotes monoubiquitination and nuclear translocation of TIEG1 resulting in Foxp3 expression in naïve CD4 T cells (left). In the presence of IL-6, Tyk2-mediated phosphorylation results in K27-linked polyubiquitination of TIEG1. This negatively regulates TIEG1 and inhibits Foxp3 transcription which may promote Th17 development (right).

Supplementary Figure 3. Reduced LLC tumor growth in TIEG1-/- mice. A, Kinetics of LLC tumor growth in TIEG1+/+ mice (left; n=4) and TIEG-/- mice (right; n=4). \(1 \times 10^6\) LLC cells were injected s.c. into the flanks of mice, and tumor volume was monitored by measuring the length and width of the tumor three times per week. Day 0, the date of inoculation. B, Representative images of tumor mass from tumor bearing TIEG1+/+ and TIEG1-/- mice. Images were taken at day 25 after inoculation of LLC cells.
Supplementary Figure 1
TGF-β

Itch

Ub

Nucleus

Cytoplasm

IL-17

IL-6

Tyk2

P

STAT3/RORγT

Supplementary Figure 2