Off-Target Serine/Threonine Kinase 10 Inhibition by Erlotinib Enhances Lymphocytic Activity Leading to Severe Skin Disorders

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ABSTRACT

Skin disorders are among the most common adverse events related to treatment with epidermal growth factor receptor (EGFR) kinase inhibitors, and of these, erlotinib is known to cause more frequent and severe skin disease than other agents in this class. Although previous reports have shown that cutaneous manifestations are triggered by the inhibition of multiple EGFR-related homeostatic functions of the skin, this mechanism alone cannot explain the differences in frequency and severity of skin disorders caused by different kinase inhibitors. In this study, we focused on the relationship between the off-target kinase inhibition and aggravation of skin disorders. Based on calculations using reported $K_d$ values and plasma drug concentrations, serine/threonine kinase 10 (STK10) and Ste20-like kinase (SLK) were selected as candidates preferentially inhibited by erlotinib over gefitinib. In vitro experiments confirmed that STK10 and SLK kinase activity are inhibited by erlotinib at clinical concentrations, whereas only STK10 is slightly inhibited by gefitinib. It was also shown that erlotinib up-regulated lymphocytic responses such as interleukin (IL)-2 secretion and cell migration at clinical concentrations, whereas gefitinib did not affect lymphocyte activity. Moreover, small interfering RNA experiments revealed that STK10 plays a major role in up-regulation of the lymphocytic responses induced by erlotinib treatment. Finally, the role of erlotinib-induced lymphocyte activation was assessed in vivo using irritant hypersensitivity models. The results indicated that erlotinib aggravates cutaneous inflammatory reactions through the activation of lymphocytic responses such as IL-2 secretion and cell migration. These results demonstrated that off-target inhibition of STK10 by erlotinib enhances lymphocytic responses, which lead to the aggravation of skin inflammation.

Introduction

Erlotinib and gefitinib are low-molecular-weight tyrosine kinase inhibitors. Their primary target is epidermal growth factor receptor (EGFR), and they have been used to treat patients with non–small-cell lung cancer (Kris et al., 2003; Shepherd et al., 2005). In addition to erlotinib and gefitinib, other kinase inhibitors and monoclonal antibodies that target EGFR, including lapatinib, cetuximab, and panitumumab, have also been developed and used for the treatment of various types of cancer (Bonner et al., 2006; Geyer et al., 2005). Skin disorders, including rash, pruritus, dry skin, and acne, are among the most common adverse events observed in patients treated with these drugs (Albanell et al., 2002; Segaert and Van Cutsem, 2005). After a week of treatment with erlotinib or gefitinib, acniform eruptions with severe pain and itching appear primarily on the face and upper trunk (Hidalgo et al., 2001; Pallis et al., 2003). These symptoms can worsen a patient’s quality of life and can lead to the dose reduction or discontinuation of treatment with EGFR inhibitory agents (Shepherd et al., 2005).

Many studies have examined the relationship between EGFR inhibition and the development of cutaneous side effects (Albanell et al., 2002; Guttmann-Yassky et al., 2010). Inhibition of EGFR-mediated signaling pathways induces multiple effects in basal keratinocytes, including growth arrest, decreased migration, increased cell attachment, aberrations of cell cycle, expression of transforming growth factor-β, and recruitment of inflammatory cells (Albanell et al., 2002; Guttman-Yassky et al., 2010).
mal differentiation, apoptosis, and stimulation of inflammatory systems, all of which result in distinctive cutaneous manifestations (Kari et al., 2003; Mascia et al., 2003; Woodworth et al., 2005). However, the prevalence and severity of drug-related skin disorders vary significantly among EGFR kinase inhibitors. In Japanese patients, the frequency of rash was higher in patients receiving erlotinib (97%) (Tarceva, Japanese package insert, 2010; Chugai Pharmaceutical Corp, Tokyo, Japan) than in those on gefitinib (63%) (Iressa, Japanese package insert, 2010; AstraZeneca, Osaka, Japan). Lapatinib, a dual EGFR and human epidermal growth factor receptor 2 tyrosine kinase inhibitor used for the treatment of human epidermal growth factor receptor 2-positive breast cancer, was reported to induce rash in 49.8% of patients receiving monotherapy (Toi et al., 2009). In addition, rash severity also differs in patients taking erlotinib versus gefitinib. In phase I/II studies of erlotinib in the Japanese population, the frequency of rash graded using the National Cancer Institute Common Terminology Criteria for Adverse Events levels G2 and G3 was 67 and 4%, respectively (Tarceva Japanese package insert), whereas in phase II trial of gefitinib in patients composed of Japanese and non-Japanese patients, the frequency of rash graded G2 and G3 was 19 and 1%, respectively (Fukuoka et al., 2003). Moreover, the IC50 values for erlotinib, gefitinib, and lapatinib were reported to be 2, 23, and 11 nM, respectively (Moyer et al., 1997; Rusnak et al., 2001; Albanell et al., 2002). Considering the steady-state unbound plasma concentrations of each drug under clinical conditions, it is estimated that EGFR is almost completely inhibited in each case. This suggests that factors other than EGFR inhibition also contribute to the severe cutaneous side effects of erlotinib.

A few reports have described the unanticipated effects caused by the inhibition of off-target kinases. Although imatinib was designed to specifically target BCR-Abl, a gene product that causes chronic myeloid leukemia, it has been suggested that imatinib has direct effects on bone-resorbing osteoclasts and bone-forming osteoblasts through the off-target inhibition of c-fms, c-kit, carbonic anhydrase II, and platelet-derived growth factor receptor (Vandyke et al., 2010). In another example, sunitinib, a multitarget kinase inhibitor used to treat renal cell carcinoma and gastrointestinal stromal tumor, inhibits a number of growth factor receptors regulating both tumor cell proliferation and tumor angiogenesis (Gutman-Yassky et al., 2010). It has been suggested that the off-target inhibition of 5'-AMP-activated protein kinase plays a central role in sunitinib toxicity in cardiomyocytes (Kerkela et al., 2009).

Karaman et al. (2008) have reported comprehensive measurement of selectivity of kinase inhibitors. That analysis revealed that both gefitinib and erlotinib interact with several off-target kinases and with their primary target, EGFR. However, it is still unclear whether off-target inhibition by erlotinib or gefitinib plays a role in cutaneous side effects of these drugs. Here, to test that possibility, we compared off-target inhibition profiles of gefitinib and erlotinib at the steady-state unbound plasma concentrations found under clinical conditions. Our findings suggest that the off-target serine/threonine kinase 10 (STK10) is inhibited much more potently by erlotinib than by gefitinib under clinical conditions. Furthermore, in vitro experiments showed that erlotinib enhances lymphocytic responses such as cell migration and interleukin-2 (IL-2) secretion via STK10 inhibition. Our in vivo observations were consistent with these in vitro results. These results suggest that erlotinib exacerbates skin disorders through off-target kinase inhibition.

Materials and Methods

Cell Lines. 293FT cells were obtained from Invitrogen (Carlsbad, CA) and cultured in Dulbecco’s modified Eagle’s medium (Nacalai Tesque, Kyoto, Japan) supplemented with 10% fetal bovine serum, 2 mM l-glutamine, and 100 µg/ml penicillin and streptomycin (Invitrogen). Jurkat E6-1 cells were obtained from the American Type Culture Collection (Manassas, VA) and cultured in RPMI 1640 medium (Nacalai Tesque) supplemented with 10% fetal bovine serum and 100 µg/ml penicillin and streptomycin.

Animals. Male ddY mice were purchased from Japan SLC Inc. (Hamamatsu, Japan). Mice were allowed to acclimate to housing conditions for at least 7 days before handling and were analyzed at 6 weeks of age. All animal procedures were approved by the Institutional Animal Care and Use Committee of Graduate School of Medicine, the University of Tokyo.

Calculation of Occupancy Rates. Mean occupancy rate (R) of the kinase by the drug under clinical conditions was calculated according to the Michaelis-Menten-type equation \( R = \frac{C_{p,a,u}}{K_d + C_{p,a,u}} \), where \( C_{p,a,u} \) and \( K_d \) are the steady-state unbound plasma concentration of the drug obtained from Japanese package insert and the dissociation constant of the drug from the kinase obtained from a previous report (Karaman et al., 2008), respectively.

Preparation of Recombinant Kinase. Genes encoding human STK10 and Ste20-like kinase (SLK) were cloned using cDNA from MCF-7 cells and Jurkat E6-1 cells, respectively. Genes encoding mouse STK10 and SLK were cloned using cDNA from mouse spleens and mouse brains, respectively. Each gene was fused with a histidine tag at the N terminus and subcloned into the pcDNA3.3 vector (Invitrogen) to produce 6-His STK10/pcDNA3.3 and 6-His SLK/pcDNA3.3. 293FT cells were transfected with 6-His STK10/pcDNA3.3 or 6-His SLK/pcDNA3.3 using Lipofectamine 2000 (Invitrogen) according to the manufacturer’s protocol. Forty-eight hours after transfection, cells were lysed in a lysis buffer (phosphate-buffered saline, pH 8.0) containing 1% Nonidet P-40 (Nacalai Tesque, Kyoto, Japan) and a protease inhibitor cocktail (Roche Diagnostics, Indianapolis, IN). The recombinant proteins were purified using Profinity IMAC Ni-Charged Resin (Bio-Rad Laboratories, Tokyo, Japan).

In Vitro Kinase Assay. To measure the kinase activity of STK10 and SLK, 100 ng of each recombinant protein was incubated with 1 µg of myelin basic protein (Millipore Biosciences Research Reagents, Temecula, CA) and 1 mM ATP in kinase buffer (20 mM HEPES, 10 mM MgCl2, 3 mM MnCl2, and 0.1 mM bovine serum albumin, pH 7.6) for 1 h at 37°C. The remaining ATP concentration was determined using the Kinase-Glo Plus Luminescent Kinase Assay (Promega, Madison, WI) according to the manufacturer’s protocol. To determine the IC50 values for gefitinib and erlotinib for each kinase, each compound was added to the reaction mixture at the indicated concentrations. ATP concentrations were fitted to the standard IC50 model, and IC50 values were determined based on Powell’s nonlinear least square method with a uniform weighting factor using Scientist software (MicroMath, Salt Lake City, UT).

Small Interfering RNA. siRNA was designed against human STK10 and SLK using BLOCK-it RNAi Designer (Invitrogen). Sequences were 5’-GCC UGU CUA CCC UGC AGA A-3’ for STK10 (siSTK10) and 5’-GCC AUA ACC AGA ACC UGA A-3’ for SLK (siSLK). siRNA duplexes containing a di-deoxynucleotide overhang at the 3’ terminus were synthesized (Sigma-Aldrich, St. Louis, MO). Each siRNA duplex was introduced into Jurkat E6-1 cells by electroporation using a Gene Pulser Xcell (Bio-Rad Laboratories) at a setting of 140 V, 1000 µF, and using a 0.1-cm gap cuvette according
to the manufacturer’s protocol. As a negative control (siNeg), we used siPerfect Negative Control (Sigma-Aldrich) that contains at least three miss-matches against human, mouse, and rat genes.

**Quantification of mRNA Expression by Real-Time PCR.** Total RNA was extracted from Jurkat E6-1 cells using RNAiso (Omega Bio-Tek, Lilburn, GA) according to the manufacturer’s protocol, and the prepared RNA was reverse-transcribed with ReverTra Ace (Toyobo Engineering, Osaka, Japan). Quantitative real-time PCR was performed using SYBR GreenER qPCR SuperMix Universal (Invitrogen), a Chromo4 (Bio-Rad Laboratories), and the associated software. Primers used for the quantification of gene expression were as follows: human STK10: forward, 5′-ATC CTT CGC CTC TCT ACC TT-3′; reverse, 5′-GCC TTG TAA ACC TTG CCG AA-3′; human IL-2: forward, 5′-TCTG TCT GTC GTG CTC TAC GAA-3′; reverse, 5′-GAC TGT CAG GGT AGT GGG ATT C-3′. The relative expression level of each gene was normalized to each β-actin.

**Western Blotting.** Forty-eight hours after siRNA introduction into Jurkat E6-1 cells, whole cell lysates were prepared with phosphate-buffered saline containing 1% NP-40 (lysis buffer). Proteins (20 μg/lane) were separated on 7.5% of SDS-polyacrylamide gels and transferred onto Immobilon membranes (Millipore Corporation, Billerica, MA). After a blocking treatment with Tris-buffered saline containing 5% skimmed milk, membranes were incubated with anti-human STK10 antibody or anti-human β-actin (BETHYL, Montgomery, TX) as a secondary antibody and anti-rabbit IgG antibody labeled with horseradish peroxidase (GE Healthcare, Chalfont St. Giles, Buckinghamshire, UK) was used. ECL Plus (GE Healthcare) was used for detection according to the manufacturer’s protocol and analyzed using a Chemidoc XRS (Bio-Rad Laboratories).

**Measurement of IL-2 by Enzyme-Linked Immunosorben Assay.** Jurkat E6-1 cells were seeded on a 96-well plate at a density of 5 × 10^4 cells/well for stimulation with Dynabeads Human T-Activator CD3/CD28 (Veritas, Tokyo, Japan) at a ratio of three beads to one cell. Lymph node cells isolated from ddY mice were seeded on a 96-well plate at a density of 5 × 10^5 cells/well for stimulation with Dynabeads Mouse T-Activator CD3/CD28 (Veritas) at a ratio of one bead to one cell. For costimulation experiments with PMA (20 ng/ml) and ionomycin (1 μM), both Jurkat cells and lymph node cells were seeded at a density of 2 × 10^5 cells/well. The indicated concentrations of gefitinib, erlotinib, and/or 2′-amino-2′-[2-(4-octylphenyl)ethyl]-1,3-propanediol, hydrochloride (FTY720; Cayman Chemical, Ann Arbor, MI) were orally administered on days –2, –1, and 0. The degree of ear flap swelling was measured 24 h after irritant treatment. To evaluate the effect of lymphocyte suppression, 3 mg/kg 2-amino-2′-[2-(4-octylphenyl)ethyl]-1,3-propanediol, hydrochloride (FTY720; Cayman Chemical, Ann Arbor, MI) was orally administered on days –2, –1, and 0. The degree of ear flap swelling was measured 24 h after irritant application. Infiltration of lymphocytes to the earflaps was also quantitated. For this experiment, treatment with erlotinib or gefitinib and irritant application were performed as described above, and mice were sacrificed 6 h after irritant challenge. The earflaps were excised and fixed in phosphate-buffered saline containing 4% paraformaldehyde. The tissues were embedded in paraffin, and then longitudinal thin sections were prepared for staining with hematoxylin-eosin as described previously. In one-third, one-half, and two-thirds distances from the base to the tip of earflap sections along with long axis, areas with 200 μm width were selected, and infiltrated lymphocytes were counted.

**Statistical Analyses.** All data are expressed as means ± S.D. from at least three independent experiments. Statistical analysis was performed using a Student’s t test or an analysis of variance followed by Bonferroni’s test where applicable.
Results

Comparison of Off-Target Kinase Occupancy Rates in Erlotinib and Gefitinib at Clinical Levels. To compare the off-target interaction profiles of erlotinib and gefitinib, we calculated their occupancy rates for 317 types of human kinases based on a previous report of high-throughput measurements of the $K_d$ values of various kinase inhibitors for human kinases (Karaman et al., 2008), taking into account the steady-state mean unbound plasma concentration of these agents under clinical conditions (Nakagawa et al., 2003; Yamamoto et al., 2008) (Fig. 1). It was estimated that both erlotinib and gefitinib occupy EGFR almost completely (100% by erlotinib and 99% by gefitinib). Cyclin G-associated kinase (GAK) was also estimated to be highly occupied by both agents (99% by erlotinib and 84% by gefitinib). In contrast, the estimated occupancy rates for STK10 and SLK were much higher for erlotinib than for gefitinib (92% by erlotinib and 12% by gefitinib for STK10, and 89% by erlotinib and 7% by gefitinib for SLK). It has been reported that both STK10 and SLK are members of the Ste20 family of serine/threonine protein kinases, and STK10 is reported to be expressed predominantly in lymphocytes (Walter et al., 2003) and to negatively regulate lymphocytic responses such as IL-2 expression and cell migration (Tao et al., 2002; Belkina et al., 2009). Given that the up-regulation of lymphocytic activity is likely to lead to the aggravation of inflammatory reactions, including skin disorders, we focused on STK10 and SLK in subsequent experiments.

Determination of IC$_{50}$ Values for STK10 and SLK by Erlotinib and Gefitinib. To confirm that STK10 and SLK kinase activity was actually inhibited by erlotinib under clinical conditions, we determined the IC$_{50}$ values of erlotinib and gefitinib for human STK10, SLK, and their mouse orthologs using an in vitro kinase assay system (Fig. 2). The results indicated that erlotinib inhibited STK10 and SLK more potently than gefitinib under clinical conditions. The IC$_{50}$ values of erlotinib and gefitinib for human STK10 were 160 and 1300 nM, respectively, and those for mouse STK10 were 350 and 1900 nM, respectively (Fig. 2, A and B). The IC$_{50}$ values of erlotinib and gefitinib for human SLK were 830 and 5200 nM, respectively, and those for mouse SLK were 480 and 1600 nM, respectively (Fig. 2, C and D). Based on these results, it was estimated that the inhibition rate for STK10 is approximately 60% for patients treated with erlotinib, but only 4% for patients treated with gefitinib. The inhibitory effects on SLK were estimated to be somewhat lower than those on STK10, and it was estimated that the inhibition rate was approximately 25% for patients treated with erlotinib and minimal for those treated with gefitinib.

Erlotinib Up-Regulates IL-2 Secretion and Cell Migration Activity in Lymphocytes. As mentioned above, it has been suggested that STK10 negatively regulates lymphocytic activity (Tao et al., 2002; Belkina et al., 2009). Therefore, we speculated that the possibility that the preferential inhibition of STK10 or SLK, which is closely related to
STK10, by erlotinib might enhance lymphocyte action. To compare the effects of erlotinib and gefitinib on lymphocytic IL-2 secretion, we measured IL-2 in the media by enzyme-linked immunosorbent assay 48 h after the activation of Jurkat E6-1 cells, which were derived from a human lymphocytic cell line. These cells were activated with CD3/CD28-coated beads, which mimic the role of antigen-presenting cells during in vivo T-cell activation (Fig. 3A). Addition of erlotinib to the media resulted in a dose-dependent increase in IL-2 secretion, whereas gefitinib had little or no effect. Unfortunately, we could not determine the EC50 values, because the viability of the cells tended to decrease over the 48-h period in which higher concentrations of erlotinib or gefitinib were added to the media. However, these results suggest that erlotinib up-regulates secretion of IL-2 from lymphocytes at clinical concentrations.

Next, we examined the effects of erlotinib and gefitinib on lymphocytic cell migration using a Transwell chemotaxis assay system (Fig. 3B). The addition of both erlotinib and gefitinib to the media led to an increase in the number of cells attracted by chemokines; however, erlotinib activated cell migration more potently than gefitinib. In this assay, cells were incubated for 2 h, and EC50 values were determined.

The EC50 values for lymphocytic cell migration were 470 nM for erlotinib and 1400 nM for gefitinib. The relative Emax values were 1.9 for erlotinib and 1.7 for gefitinib.

**STK10 Inhibition Plays a Major Role in the Up-Regulation of Lymphocytic Activation by Erlotinib.** We next examined the effects of RNAi gene suppression of STK10 and SLK to further establish whether the inhibition of STK10 or SLK by erlotinib mediates the up-regulation of lymphocyte action. Knockdown efficiency was confirmed by measuring mRNA and protein levels using quantitative real-time PCR and Western blotting, respectively, in Jurkat E6-1 cells after siRNA was introduced by electroporation (Fig. 4, A and B). Because it was reported previously that IL-2 production is negatively regulated by STK10 at the transcriptional level (Tao et al., 2002), we examined the effects of gene suppression on IL-2 mRNA expression. IL-2 mRNA expression was increased 2-fold after stimulation with CD3/CD28-coated beads in Jurkat E6-1 cells treated with siSTK10 relative to cells treated with siNeg, but no significant difference was observed in cells treated with siSLK versus siNeg (Fig. 4C). Moreover, the addition of erlotinib at clinical concentrations resulted in increased IL-2 mRNA expression under SLK knockdown and control conditions; however, it did not significantly affect IL-2 mRNA under STK10 knockdown conditions (Fig. 4C). Changes in IL-2 protein in the media were consistent with these changes in mRNA (Fig. 4D). These results suggest that up-regulation of IL-2 secretion in response to erlotinib is due to transcriptional up-regulation mediated by the inhibition of STK10 by erlotinib.

We next performed a transwell migration assay using Jurkat E6-1 cells under the various siRNA conditions (Fig. 4E). STK10 knockdown increased the number of cells attracted by chemokines compared with control and SLK knockdown conditions. As was the case for IL-2 production, the addition of erlotinib to the media at clinical concentrations resulted in increased chemokine-induced cell migration, whereas no significant change was observed in cells treated with siSLK versus siNeg (Fig. 4C). Moreover, the addition of erlotinib at clinical concentrations resulted in increased IL-2 mRNA expression under STK10 knockdown conditions (Fig. 4C). Changes in IL-2 protein in the media were consistent with these changes in mRNA (Fig. 4D). These results suggest that the up-regulation of lymphocytic action by erlotinib is largely mediated by STK10 inhibition.

**Erlotinib Does Not Affect the Mitogen-Activated Protein Kinase Extracellular Signal-Regulated Kinase Kinase-1 Pathway or Ca2+ Signaling Pathway.** It has been reported that mitogen-activated protein kinase extracellular signal-regulated kinase (ERK) kinase-1 (MEK1) increases the IL-2 gene transcription (Whitehurst and Geppert, 1996). To confirm whether the modulation of MEK1 is the cause of a high level of IL-2 secretion in response to erlotinib, the involvement of MEK1 in increased IL-2 secretion by erlotinib was investigated using the MEK1-selective inhibitor PD98059. In the case of Jurkat E6-1 cells stimulated with CD3/CD28 beads, PD98059 partially inhibited the secretion of IL-2. The resistant remainder is believed to be dependent on signaling pathways other than the MEK1/ERK pathway. Moreover, erlotinib stimulated IL-2 secretion significantly even in cells cotreated with PD98059 (Fig. 5A). These results suggest that the increased IL-2 secretion induced by erlotinib treatment is mediated by the signals other than MEK1 signals. In addition, we measured the IL-2 secretion in cells under PMA/ionomycin costimulation, which bypassed the T-cell receptor (TCR) signaling to activate Ras and calcineurin (Chatila et al., 1989; Franklin et al., 1994). Upon PMA/
on IL-2 secretion in vitro using lymph node cells from ddY mice. Upon stimulation with CD3/CD28 beads, the addition of erlotinib to the media resulted in a dose-dependent increase in IL-2 secretion, whereas gefitinib had no effect on lymph node cells at concentration used in clinical situations (Fig. 6D). In addition, erlotinib and gefitinib had no effect on the secretion of IL-2 in lymph node cells upon PMA/ionomycin costimulation (Fig. 6E). These results indicate that erlotinib exhibits similar lymphocytic activation properties in both Jurkat E6-1 cells and mouse lymph node cells. However, IL-2 secretion in response to PMA/ionomycin costimulation or CD3/CD28 bead stimulation was higher in lymph node cells than in Jurkat-E1 cells (Fig. 3). Several possibilities can be advanced to explain this point. Jurkat-E1 is an immortalized cell line that was cultured alone, whereas the lymph node cells were freshly isolated from mice as a mixture of several types of cells, including lymphocytes, macrophages, and dendritic cells. In general, physiological reactions of the immortalized cell line often proceed weakly compared with the case of primary cultured cells. Thus, we believe it is plausible that lymph node cells were more strongly activated than Jurkat cells, although Jurkat cells maintain their ability to produce IL-2 in response to stimulation. In addition, IL-2 production from T cells after stimulation might be up-regulated via cross-talk with other types of cells in the case of isolated lymph node cells.

Activation of Lymphocytic Responses Mediates the Exacerbation of Skin Inflammation by Erlotinib Treatment. We also measured lymphocyte infiltration in earflaps by tissue section staining. Only erlotinib treatment increased lymphocyte infiltration in the inflamed area (Fig. 7A). To confirm our in vitro finding of the up-regulation of IL-2 secretion and lymphocyte migration in response to erlotinib treatment, we

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**Administration of Erlotinib Exacerbates Irritant-Induced Skin Inflammation.** Our in vitro experiments indicated that the preferential inhibition of STK10 by erlotinib enhances the lymphocytic response, leading to the aggravation of skin disorders. To confirm this, we performed an in vivo irritant hypersensitivity assay using male ddY mice. In these experiments, the swelling of earflaps 24 h after topical application of croton oil was used as an indicator of irritant hypersensitivity, and erlotinib or gefitinib was orally administered four times over 24 h. The dosage regimens for erlotinib and gefitinib were designed to maintain the inhibition rate for STK10 at a level similar to that of patients treated with these agents, based on previous reports of the mouse pharmacokinetic profiles (Marchetti et al., 2008; Wang et al., 2008) and the IC$_{50}$ values for human and mouse STK10 measured in the present study (Fig. 2). To confirm the STK10 inhibition profiles, we measured plasma concentrations of erlotinib and gefitinib at the indicated time points and compared the estimated STK10 inhibition profiles in mice and humans (Fig. 6, A and B). Earflap swelling was significantly exacerbated by erlotinib administration compared with controls, but gefitinib administration did not affect earflap swelling (Fig. 6C).

To confirm that mouse lymphocyte responses are actually affected by erlotinib as seen in human cell line Jurkat-E6-1 cells, we also determined the effect of erlotinib and gefitinib on IL-2 secretion, whereas erlotinib did not (Fig. 5B). This result suggests that IL-2 secretion triggered by PMA/ionomycin costimulation depends more strongly on MEK1 signaling than on CD3/CD28 stimulation and also suggests that erlotinib does not affect the MEK1 pathway or Ca$^{2+}$ signaling pathway.

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![Graphs and images illustrating the effects of erlotinib and gefitinib on IL-2 secretion and lymphocyte migration](https://example.com/graphs/471)

**Fig. 4.** STK10 inhibition plays a major role in the up-regulation of lymphocytic activity by erlotinib. A, changes of STK10 (left) and SLK (right) mRNA levels by RNAi gene suppression. Each data point represents the mean ± S.D. (n = 3). B, changes of STK10 (left) and SLK (right) protein levels by RNAi gene suppression. C, effects of RNAi gene suppression of STK10 or SLK on IL-2 mRNA expression in Jurkat E6-1 cells stimulated with beads under erlotinib or gefitinib treatment conditions. Each data point represents the mean ± S.D. (n = 3). D, effects of RNAi gene suppression of STK10 or SLK on IL-2 secretion from Jurkat E6-1 cells stimulated with beads under erlotinib or gefitinib treatment conditions. Each data point represents the mean ± S.D. (n = 3). E, effects of RNAi gene suppression of STK10 or SLK on the migration of Jurkat E6-1 cells induced by SDF-1 under erlotinib or gefitinib treatment conditions. Each data point represents the mean ± S.D. (n = 3). *p < 0.01.
administered anti-IL-2 antibody or FTY720, an immunosuppressant sequestering T lymphocytes within peripheral lymphoid tissues, to mice treated with the irritant. Treatment with anti-IL-2 antibody or FTY720 reduced ear flap swelling in erlotinib-treated animals (Fig. 7, B and C). These results indicate that the up-regulation of lymphocytic activation by erlotinib treatment contributes to the aggravation of skin inflammation.

**Discussion**

A limited number of reports have focused on the inhibition of off-target kinases by clinically used kinase inhibitors (Bain et al., 2007; Karaman et al., 2008). These reports have only addressed the magnitude of off-target and primary target inhibition rates, and the pharmacokinetic properties of these anticancer agents have not been considered. However, the off-target kinase inhibition rate is directly correlated with the development of adverse effects. The off-target inhibition rate for a kinase inhibitor that shows 100-fold selectivity is estimated to be 8.3% if the plasma concentration of the agent is adjusted to achieve 90.0% inhibition of the primary target. It appears that 100-fold selectivity is safe enough in this case. However, if the inhibition rate for the primary target is adjusted to 99.5%, the off-target kinase inhibition rate is estimated to increase to 66.6%, possibly resulting in adverse effects. This estimation also demonstrates that the off-target inhibition rate shows large variability depending on the plasma concentration of the drug. For cytotoxic anticancer agents, clinical doses are typically determined based on the maximum tolerated dose estimation (Sleijfer and Wiemer, 2008). However, this is not always the case for tyrosine kinase inhibitors, and the primary target inhibition rates achieved at clinical doses vary among kinase inhibitors. For imatinib, the inhibition rate for Bcr-Abl, the primary target of this agent, is estimated to be 90% based on the reported IC_{50} value and the steady-state unbound plasma concentration (Deininger et al., 2005). For erlotinib and gefitinib, which are the focus of this study, the EGFR inhibition rates are estimated to be 100 and 99%, respectively (Fig. 1). Therefore, it is necessary to convert K_d or IC_{50} values for off-target kinases into inhibition rates at clinical concentrations taking into consideration the pharmacokinetic properties of tyrosine kinase inhibitors.

In the present study, we focused on the relationship between off-target kinase inhibition and the aggravation of skin disorders, focusing on two kinase inhibitors that primarily target EGFR, erlotinib and gefitinib. Based on estimations of human kinase occupancy by erlotinib and gefitinib, we determined that erlotinib interacts strongly with GAK, STK10, SLK, and EGFR (Fig. 1). A number of studies have described the involvement of EGFR inhibition in the pathophysiological mechanisms of cutaneous toxicity observed in patients treated with EGFR kinase inhibitors (Lacouture, 2006). The primary effect is damage to the sebaceous glands and follicular epithelia, which leads to altered epidermal growth and differentiation. These cellular and structural changes subsequently trigger the release of chemokines, leading to the infiltration of mononuclear leukocytes including T lymphocytes, inflammatory dendritic cells, and macrophages to the lesion area (Guttman-Yassky et al., 2010). Histological findings show that the earliest change is the infiltration of T lymphocytes immunoreactive for CD45RO, a marker of T lymphocyte activation, which is followed by abundant neutrophil infiltration (Busam et al., 2001). T cells are believed to facilitate inflammation through the release of effector molecules (Deane and Hickey, 2009). A series of reactions, sometimes along with the bacterial infection, finally result in cutaneous injury (Guttman-Yassky et al., 2010). However, these proposed mechanisms do not fully explain the difference in the severity of skin disorders induced by erlotinib and gefitinib.

It seems unlikely that the inhibition of GAK is involved in the aggravation of skin disorder because gefitinib also interacts strongly with GAK (Fig. 1). In addition, the frequency of skin rash in patients taking lapatinib, a kinase inhibitor primarily targeting EGFR and Her2, is comparable with that in patients taking gefitinib, despite the fact that lapatinib does not interact with GAK (Karaman et al., 2008; Toi et al., 2009). In contrast, erlotinib shows a greater interaction with STK10 and SLK than with gefitinib (Fig. 1). STK10 and SLK are serine/threonine protein kinases that are closely mapped in the phylogenetic tree of the Ste20 family, and STK10 is predominantly expressed in lymphocytes, whereas SLK is ubiquitously expressed (Yamada et al., 2000; Walter et al., 2003). It has been reported that STK10 negatively regulates IL-2 expression in lymphocytes at the transcriptional level and also negatively regulates cell motility through the phosphorylation of ezrin-radixin-moesin proteins (Tao et al., 2002; Belkina et al., 2009). Although there are reports that SLK is required for breast cancer cell motility, disassembly of actin stress fibers, and radial microtubule organization, little is known about the function of SLK in lymphocytes (Burakov et al., 2002; Belkina et al., 2009).
et al., 2008; Roovers et al., 2009). Considering these points, we selected STK10 and SLK as candidates for off-target inhibition, leading to inflammatory responses associated with the development of skin disorders using in vitro and in vivo experimental systems.

An in vitro kinase assay showed that the inhibition rates for STK10 and SLK by erlotinib in the steady state are 59 and 21%, respectively, whereas gefitinib showed only a slight inhibition of STK10 at clinical concentrations (Fig. 2). The IC_{50} values, which we determined from our in vitro experiments, were somewhat higher than the reported K_{d} values (Moyer et al., 1997; Albanell et al., 2002). Differences in experimental conditions might explain this difference. We determined the IC_{50} values using enzyme kinetics data obtained in the presence of 1 μM ATP, and the reported K_{d} values were determined using a high-throughput competitive binding assay (Karaman et al., 2008). However, we confirmed that erlotinib significantly inhibits STK10 at clinical concentrations. Our cell-based assays showed that erlotinib enhances lymphocytic responses such as IL-2 secretion and cell migration (Fig. 3). The EC_{50} values of erlotinib and gefitinib for the enhancement of cell migration were comparable with the IC_{50} values for STK10 inhibition (Fig. 3). In addition, RNAi gene suppression experiments indicated that the up-regulation of lymphocytic activity by erlotinib or gefitinib is primarily mediated by STK10 inhibition (Fig. 4).

We obtained similar results by using kinase dead forms of STK10 and SLK in place of siRNA (data not shown). Considering these in vitro results, it can be hypothesized that the preferential inhibition of STK10 by erlotinib leads to enhanced lymphocyte migration to the lesion area and increased local IL-2 secretion.

T-cell activation is induced by a series of intracellular signaling cascade initiated by signals from the TCR/CD3 complex and other costimulatory molecules including CD28 (Clevers et al., 1988). CD28 provides an essential costimulatory signal that approximates T cell and antigen-presenting cells and augments the production of IL-2 (Lenschow et al., 1996). In our present study, we used CD3/CD28 beads stim-

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**Fig. 6.** Erlotinib exacerbates the skin inflammatory response to croton oil. Inhibition ratios of STK10 in ddY mice orally administered erlotinib (A) or gefitinib (B). Each data point represents the mean ± S.D. (n = 5). The solid line represents the fitted inhibition rate profile of mSTK10 by erlotinib in the mouse model, and the dashed line represents the predicted inhibition rate profile of hSTK10 by erlotinib under human clinical conditions. C, effects of erlotinib or gefitinib administration on changes in earflap swelling. Each data point represents the mean ± S.D. (n = 5), *, P < 0.05; **, P < 0.01. D, effects of erlotinib and gefitinib on IL-2 secretion in mouse lymph node cells stimulated with Dynabeads Mouse T-Activator CD3/CD28. Each data point represents the mean ± S.D. (n = 4). E, effects of erlotinib and gefitinib on IL-2 secretion in mouse lymph node cells stimulated with PMA/ionomycin. Each data point represents the mean ± S.D. (n = 4).
ulation, which mimics natural TCR signaling. TCR signaling is further amplified by 70-kDa ζ-chain-associated protein kinase, which leads to the activation of multiple pathways, including ERK, c-Jun NH2-terminal kinase, nuclear factor-κB, p38, and nuclear factor of activated T cells, which ultimately induce IL-2 gene expression (Chu et al., 1998). Meanwhile, costimulation with PMA/ionomycin bypasses the TCR signaling and activates the signaling downstream of phospholipase Cγ, a downstream effector of 70-kDa ζ-chain-associated protein kinase (Chatila et al., 1989; Franklin et al., 1994). Because erlotinib could up-regulate the IL-2 secretion under the CD3/CD28 stimulation not upon PMA/ionomycin stimulation (Fig. 5, A and B), up-regulation of IL-2 secretion by erlotinib via STK10 inhibition could be dependent on the signals other than the downstream signals of phospholipase Cγ. These results are consistent with previous reports that STK10 down-regulates the IL-2 secretion upon Raji/SEE stimulation but not upon PMA/ionomycin cotreatment. The authors of this report considered that STK10 might act on early events in T-cell activation (Tao et al., 2002).

Our observations in a mouse contact hypersensitivity model were consistent with the in vitro observations (Figs. 6 and 7). Infiltration of lymphocytes was up-regulated in the earflaps of erlotinib-treated mice compared with gefitinib- or control-treated animals (Fig. 7). Treatment with anti-IL-2 antibody or FTY720 also resulted in a reduction in ear flap swelling induced by erlotinib treatment (Fig. 7). These observations are also consistent with the proposed mechanism of skin rash development in patients treated with erlotinib, in which inflammatory reactions are dominated by mononuclear leukocytes, including lymphocytes and neutrophils (Guttmann-Yassky et al., 2010).

It has been reported that there is significant interindividual variation in plasma concentrations of erlotinib and gefitinib (Li et al., 2006; Thomas et al., 2009). Preclinical studies have demonstrated that CYP3A4 is primarily involved in the metabolism of erlotinib and gefitinib, and therefore, pharmacokinetic interactions may take place with drugs that inhibit or induce CYP3A4 (McKillop et al., 2005; Li et al., 2007). Itraconazole is a potent inhibitor of CYP3A4 and the plasma area under the curve of gefitinib was reported to increase by 78% with the coadministration of itraconazole (Swaisland et al., 2005). Coadministration of ketoconazole, another strong inhibitor of CYP3A4, was reported to cause a 2-fold increase in plasma area under the curve and the maximum plasma concentration of erlotinib (Rahib et al., 2008). Based on the IC50 values obtained from our in vitro kinase assay and the mean unbound plasma concentrations of erlotinib and gefitinib at the steady state, it is estimated that the mean rate of STK10 inhibition was approximately 59% for erlotinib but only 4% for gefitinib (Fig. 2). If interindividual variations in CYP3A4 activity and/or drug-drug interactions were to result in a 2-fold higher unbound plasma concentration of erlotinib or gefitinib compared with the mean values, the inhibition rate for STK10 would increase to 74% for erlotinib but would remain at 8% for gefitinib. In contrast, the inhibition rate for EGFR, which is almost 100% for both erlotinib and gefitinib, would not be affected by these conditions. These estimations indicate that interindividual differences in the severity of skin rash among patients treated with erlotinib may be due to interindividual variability in the STK10 inhibition rate. This hypothesis is also supported by a previous report showing that variability in skin rash susceptibility was associated with trough erlotinib plasma concentrations (Rudin et al., 2008).

In conclusion, we propose a mechanism in which erlotinib exacerbates skin disorders through the enhancement of lymphocytic responses such as lymphocyte migration and increased IL-2 secretion via STK10 inhibition. Because it has been demonstrated that off-target inhibition by kinase inhibitors can result in the development of adverse effects under clinical conditions, it would be beneficial to determine the comprehensive inhibitory profiles of kinase inhibitors at the preclinical stage. In cases in which the inhibition rate for off-target kinases is predicted to be variable within the clinical concentration range for reasons such as interindividual variability in pharmacokinetics and drug-drug interactions, clinical dose determination studies should be performed carefully and strategically to maximize therapeutic effects and minimize side effects.

Fig. 7. IL-2 secretion and lymphocyte infiltration contribute to the aggravation of the inflammatory response by erlotinib. A, effects of erlotinib or gefitinib administration on ear flap lymphocyte infiltration. Each data point represents the mean ± S.D. (n = 3). B, effects of anti-IL-2 Ab on ear flap swelling aggravated by erlotinib treatment. Each data point represents the mean ± S.D. (n = 6). C, effects of FTY720 on the ear flap swelling aggravated by erlotinib treatment. Each data point represents the mean ± S.D. (n = 8). * P < 0.05.
Aggravation of Skin Rash via STK10 Inhibition by Erlotinib


