Tricin Isolated from Enzyme-Treated *Zizania latifolia* Extract Inhibits IgE-Mediated Allergic Reactions in RBL-2H3 Cells by Targeting the Lyn/Syk Pathway

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**Abstract:** Tricin, a flavone present in rice bran, is confirmed as the major efficacious compound present in the enzyme-treated *Zizania latifolia* extract (ETZL), which protects against UVB-induced skin-aging. However, the suppressive mechanism of tricin on allergic responses remains unknown. The present study, therefore, aimed to determine the mechanisms of tricin and ETZL on mast cell degranulation in IgE-activated rat basophilic leukemia cell line (RBL-2H3) cells. We investigated the regulatory effects of tricin and ETZL on degranulation, production of cytokines and lipid mediators, and signaling proteins involved in the IgE-bound high-affinity IgE receptor activation, mitogen-activated protein kinase, arachidonic acid and Syk. The production of β-hexosaminidase, tumor necrosis factor-α, interleukin-4, leukotrienes (LT) B<sub>4</sub>, LTC<sub>4</sub> and prostaglandin E<sub>2</sub> in IgE-stimulated RBL-2H3 cells were significantly inhibited by exposure to tricin or ETZL. Moreover, tricin and ETZL inhibit the phosphorylation of cytosolic phospholipase A<sub>2</sub>, 5-lipoxygenase and cyclooxygenase-2. Furthermore, the phosphorylation of Akt, ERK, p38, JNK, protein kinase Cδ and phospholipase Cγ<sub>1</sub> were effectively suppressed by both samples. Exposure to tricin or ETZL also significantly decreases the phosphorylation of Lyn and Syk, but has minimal effect on Fyn. Taken together, our data indicate that tricin and ETZL are potential anti-allergic materials that could be applied for the prevention of allergy-related diseases.

**Keywords:** tricin; enzyme-treated *Zizania latifolia* extract; anti-allergic activity; FcεRI signaling pathway; MAPK signaling pathway; arachidonic acid signaling pathway

1. Introduction

*Zizania latifolia* (Gramineae) is the only member of the wild rice genus *Zizania* native to Asia, including China, Korea and Japan. *Z. latifolia* Turcz differs botanically from the other wild rice species, including *Z. aquatica*, *Z. palustris* and *Z. texana*. The *Z. latifolia* grains impart numerous pharmacological effects, including suppression of hyperlipidemia [1], reduction of oxidative stress in cells [2], decreasing blood glucose levels [3], improvement of insulin resistance [3] and anti-obesity [1]. Recently, it has been reported that the aerial portion of *Z. latifolia* exhibits a variety of inhibitory activities, such as H<sub>2</sub>O<sub>2</sub>-induced apoptosis in Neuro2A cells [4], angiotensin-converting enzyme and oxidative stress [5] and ultraviolet (UV) B-induced wrinkle formation in SKH-1 hairless mice and in human dermal fibroblasts [6,7]. The methanol extract of the aerial portion suppresses allergic responses in rat basophilic leukemia cell line (RBL-2H3) through inhibition of compound 48/80-induced degranulation, antigen-induced β-hexosaminidase release and phorbol 12-myristate 13-acetate plus A23187-induced tumor necrosis factor (TNF)-α production [8].
In our previous study, we identified five derivatives of tricin (4',5,7-trihydroxy-3', 5'-dimethoxyflavone) (Figure 1) in the methanol extract prepared using the aerial portion of Z. latifolia [8]. The tricin derivatives were determined to prevent release of β-hexosaminidase in RBL-2H3 cells sensitized with anti-dinitrophenyl (DNP) immunoglobulin E (IgE). Exposure to tricin is known to suppress cyclooxygenase (COX) activity in mice, and prostaglandin E2 (PGE2) production in colon cells and murine plasma [9]. Tricin was found to be the quantitatively major component obtained in 70% ethanol extract of Z. latifolia. Treatment with an enzyme mixture significantly increased the tricin content in a hydrolyzed ethanol extract [7].

![Figure 1. Chemical structure of tricin.](image)

Mast cells play a major role in IgE-mediated allergic responses (hypersensitivity) by producing inflammatory mediators such as histamine, leukotrienes, prostaglandins and cytokines upon binding of an antigen to the IgE-bound high-affinity IgE receptor (FccRI) [10]. Since histamine released from cytosolic granules in mast cells induces acute allergic responses, antihistamines are commonly used to treat allergic diseases. Arachidonic acid metabolites (such as leukotrienes) and cytokines (including TNF-α and interleukin (IL)-4) are also critical in prompting allergic reactions [10]. Leukotrienes (LT) such as LTB4 and LTC4 attract neutrophils and stimulate the late-phase allergic reaction [11]. Cytokines released by a T-helper type 2 (Th2) response (such as IL-4, IL-6 and IL-13) activate B cells to produce IgE, thereby activating mast cells and inducing allergic inflammation [12].

In addition, FccRI stimulation results in the immediate activation of two Src family kinases, Lyn and Fyn, and a tyrosine kinase spleen tyrosine kinase (Syk), to initiate mast cell degranulation, with subsequent activation of the phospholipase Cγ (PLCγ) and protein kinase B (Akt) signaling pathways [10,13]. The underlying pathophysiologic mechanisms of allergic response are well-documented by previous in vitro and in vivo studies [14]. A mitogen-activated protein kinase (MAPK) cascade is an important signaling pathway that regulates the differentiation, activation, proliferation, degranulation and migration of immune cells, including mast cells. The p38 MAP kinase activates IL-4 in bone marrow mast cells, and the activation of c-Jun N terminal kinase (JNK) is also responsible for the production of pro-inflammatory cytokines in mast cells, including IL-2, IL-6 and TNF-α [15]. The phosphorylated Akt regulates the transcription of IL-2 and TNF-α promoters [12]. Moreover, the overexpression of Syk activates extracellular signal-regulated kinase 1/2 (ERK) phosphorylation, and hence the arachidonic acid signaling pathway, including activation of the expressions of cytosolic phospholipase A2 (cPLA2), 5-lipoxygenase (5-LO) and COX-2, and secretions of leukotrienes and prostaglandins [11,12]. Phosphorylation of ERK1/2 plays a role in signaling TNF-α, IL-3, IL-5 and IL-13 secretion in mast cells [12,15]. Therefore, decreasing secretions of cytokines (TNF-α, IL-4), leukotrienes and prostaglandins, is crucial in anti-allergic reactions.

Antihistamines and mast cell stabilizers are widely used as anti-allergy agents for ameliorating allergic inflammatory symptoms, despite some side-effects such as drowsiness. Recently, efficacious edible plant materials devoid of any side effects are gaining attention as safer anti-allergic agents. RBL-2H3 cells express FccRI on the cell membrane, and the IgE-bound FccRI can be activated by antigens; thus, these cells are widely used for in vitro investigations on the mechanism of mast cell activation [16,17].
Mechanisms associated with the anti-allergic action of tricin isolated from enzyme-treated *Z. latifolia* extract (ETZL) remain unclear. Therefore, to understand the mechanism by which tricin or ETZL alleviates the allergic response in IgE-mediated allergic reactions in RBL-2H3 cells, this study investigated the regulatory effects on signaling cascades, such as FcεRI, arachidonic acid and MAPKs.

2. Results and Discussion

2.1. Effects of the Non-Enzyme Treated Ethanol Extract of *Z. latifolia* (NEZL), ETZL and Tricin on Anti-DNP IgE-Mediated Degranulation in RBL-2H3 Cells

As reported in our previous studies, the methanol extract from aerial *Z. latifolia* exerts excellent anti-degranulation activity in RBL-2H3 cells, and contains tricin and its derivatives as the major components for anti-allergic activity. The identified derivatives include tricin-7-O-β-D-glucopyranose, tricin-4’-O-(threo-β-guaiacylglyceryl) ether 7-O-β-D-glucopyranose, tricin-4’-O-(erythro-β-guaiacylglyceryl) ether 7-O-β-D-glucopyranose and tricin-4’-O-(erythro-β-guaiacylglyceryl) ether 7”-O-β-D-glucopyranose [8,18]. The same derivatives were also identified for the ethanol extract of *Z. latifolia* (data not published). Additionally, an enzymatic treatment applied to augment the amount of tricin increased the content by 1.27-fold in ETZL as compared to non-enzyme treated ethanol extract (NEZL), giving 25.0 mg/100 g dried *Z. latifolia* with extraction yield of 17.45%, estimating 0.14% tricin in ETZL [6]. Our previous quantitative study identifying the tricin derivatives in ETZL confirmed the simultaneous increase of four derivatives along with tricin levels, as compared to the contents obtained in NEZL (manuscript in preparation). Based on these data, we concluded that the commercially available major compound tricin, and not its derivatives, is the active component and is also the most appropriate and potent compound to be applied as a marker for industrial quality control of the extract of *Z. latifolia* (ETZL).

Thus, to investigate the mechanistic action of tricin in an anti-allergic reaction, we first evaluated the effect of NEZL, ETZL and tricin isolated from ETZL on allergic responses in the rat mast cell line RBL-2H3. In the present study, 0.1% tricin was used for each concentration of ETZL to check the efficacy of ETZL derived from tricin. Anti-DNP IgE-sensitized RBL-2H3 cells were exposed to varying concentrations of tricin and ETZL, and degranulation was triggered by adding dinitrophenyl-human serum albumin (DNP-HSA) as antigen. The effects of NEZL, ETZL and tricin were determined on the cell viability by the MTT assay. As shown in Figure 2A, exposure of an equal number of viable cells to NEZL, ETZL and tricin exhibits no cytotoxicity at any of the tested concentrations (p > 0.05). Effects on the IgE-mediated degranulation were then evaluated by determining the activity of β-hexosaminidase, a biomarker of histamine release, in IgE-activated RBL-2H3 cells. As expected, NEZL exhibit lower inhibitory activity against the release of β-hexosaminidase (50% inhibitory concentration (IC50), 106.79 µg/mL), as compared to ETZL (IC50, 70.79 µg/mL) (p < 0.001) (Figure 2B). In addition, tricin inhibited the release of β-hexosaminidase in a concentration-dependent manner with an IC50 value of 184.84 ng/mL (Figure 2B). These data indicate that the inhibitory effects are not the result of the cytotoxicity, and the degranulation-inhibitory activity of ETZL is mainly exerted by the major component, tricin.

Monitoring of mast cell degranulation using the β-hexosaminidase assay in RBL-2H3 cells is widely used to determine the effectiveness on anti-allergic responses, and is a convenient in vitro cell model for studying new molecules that inhibit mast cell activation or degranulation [19,20]. Histamine, secreted with β-hexosaminidase from mast cells, induces allergic inflammation, resulting in chronic allergic symptoms such as atopy, asthma, etc. Recent studies have reported the suppressive effects of flavonoids [21,22] and plant extracts [17] on mast cell degranulation. Among the various bioactive plant materials, *Actinidia arguta* extract has been recognized as a functional ingredient in facilitating the prevention of atopy, food allergy and asthma [23–25].
2.2. Effects of Tricin and ETZL on the Secretion of Cytokines and Lipid Mediators

Following the immediate histamine release subsequent to antigen-mediated FcεRI aggregation, degranulation of mast cells results in the secretion of inflammatory mediators, including TNF-α, IL-4, leukotrienes and prostaglandins, which play key roles during the late phase of inflammatory reactions [16,17]. Inhibiting these mediators from activated mast cells is considered a promising strategy to ameliorate allergic responses. Thus, we examined the effects of tricin and ETZL on the release of inflammatory cytokines (TNF-α and IL-4) by ELISA in anti-DNP-IgE-stimulated RBL-2H3 cells. As presented in Figure 2, the ETZL, containing a 1.27-fold higher tricin content, exhibited a 1.51-fold higher inhibitory activity than NEZL. We therefore decided to perform the subsequent experiments using tricin and ETZL, and not NEZL. Both TNF-α and IL-4 secretions were increased by antigen stimulation, to 55.3, 47.2 and 43.0 pg/mL at 10, 50 and 100 ng/mL, respectively, and ETZL (48.8, 31.1 and 10.8 pg/mL at 10, 50 and 100 µg/mL, respectively) in dose-dependent manners (Figure 3A). Similarly, IL-4 secretion was also suppressed by tricin (21.4, 17.4, 13.4 pg/mL at 10, 50 and 100 ng/mL, respectively) and ETZL (20.5, 13.9 and 6.9 pg/mL at 10, 50 and 100 µg/mL, respectively) (Figure 3B). Similar to the result of Figure 2, the release-suppressive effect of ETZL (p < 0.001) was higher than tricin (p < 0.01) at 100 µg/mL, indicating that apart from tricin, ETZL may have a 1.51-fold higher inhibitory activity than tricin.

![Figure 2](image-url)

Figure 2. Effects of tricin and enzyme-treated Zizania latifolia extract (ETZL) on cell viability and degranulation in IgE-stimulated rat basophilic leukemia (RBL-2H3) cells. (A) Cell viability; (B) β-hexosaminidase. RBL-2H3 cells were seeded in a 24-well plate (1 × 10^5 cells/well) overnight at 37 °C, and further incubated with anti-dinitrophenyl (DNP)-IgE (0.05 µg/mL) for 24 h. The IgE-sensitized cells were preincubated with varying concentrations of tricin (10–500 ng/mL) or ETZL (10–500 µg/mL) for 1 h, and subsequently stimulated with dinitrophenyl-human serum albumin (DNP-HSA) (0.1 µg/mL) for 4 h. All values are the mean ± S.E.M. of three independent experiments. Values of ** p < 0.01 and *** p < 0.001 were considered significantly different to the anti-DNP IgE plus DNP-HSA, unpaired Student’s t-test. # p < 0.001 compared to the control.
higher than tricin (p < 0.01) at 100 μg/mL, indicating that apart from tricin, ETZL may have other bioactive compounds imparting the anti-allergic inflammation, including derivatives of tricin and other trace amounts, as shown in our previous reports [8].

The histamine released from mast cells increases the expression level of FccRI on mast cells, thereby stimulating the cells for further antigen-induced activation, resulting in the production of inflammatory cytokines and lipid mediators. Recent studies have determined that apart from early-phase allergic responses, mast cells also contribute to the late-phase reactions by modulation of cytokines and mediators [17]. TNF-α, a potent inflammatory cytokine, is produced in activated macrophages and T cells, as well as antigen-stimulated mast cells [11]. IL-4 secreted from mast cells induces the production of IgE in B cells and Th2 cell differentiation, which subsequently causes anaphylaxis [11]. Several active flavonoids, such as kaempferol [26], quercetin [26], luteolin [26], chalcone [27] and curcumin [28], inhibit the antigen-induced degranulation, and the production of TNF-α and IL-4 in IgE-sensitized RBL-2H3 cells; these molecules have therefore been suggested as effective applications for the prevention of immediate-phase as well as late-phase reactions.

Conversely, the activation of arachidonate cascade in IgE-stimulated mast cells induces the production of pro-inflammatory lipid mediators such as LTB4, LTC4 and PGE2, which initiate chronic inflammation in allergy-related diseases [17]. We therefore evaluated the effects of tricin and ETZL on the production of LTB4, LTC4 and PGE2. Preincubation with varying concentrations of tricin or ETZL

![Figure 3. Effect of tricin and ETZL on pro-inflammatory cytokines in IgE-mediated RBL-2H3 cells. (A) Tumor necrosis factor (TNF)-α secretion; (B) IL-4 secretion. IgE-sensitized RBL-2H3 cells were pre-incubated with tricin or ETZL for 1 h before antigen treatment for 4 h. All values are the mean ± S.E.M. of three independent experiments. Values of * p < 0.05, ** p < 0.01 and *** p < 0.001 were considered significantly different to the anti-DNP IgE plus DNP-HSA, unpaired Student’s t-test. # p < 0.001 compared to the control.](image-url)
prior to antigen stimulation markedly suppresses the production of LTB₄ (30.4 pg/mL at 100 ng/mL tricin, 17.2 pg/mL at 100 µg/mL of ETZL), LTC₄ (200.0 pg/mL at 100 ng/mL of tricin, 86.3 pg/mL at 100 µg/mL of ETZL) and PGE₂ (273.0 pg/mL at 100 ng/mL of tricin, 237.2 pg/mL at 100 µg/mL of ETZL) (p < 0.001) (Figure 4A–C). These results indicate that tricin as well as the tricin-containing ETZL inhibit leukotriene as well as prostaglandins synthesis, thereby preventing allergic inflammation induced by LTB₄, LTC₄ or PGE₂.

Figure 4. Inhibitory effect of tricin and ETZL on formation of leukotrienes (LT)B₄, LTC₄ and prostaglandin E₂ (PGE₂) in RBL-2H3 cells stimulated by an IgE–antigen complex. (A) LTB₄ secretion; (B) LTC₄ secretion; (C) PGE₂ secretion. RBL-2H3 cells were seeded in a 24-well plate and cultured overnight at 37 °C; cells were subsequently washed and further incubated with anti-DNP-IgE (0.05 µg/mL) for 24 h. The cells were incubated with tricin (10–500 ng/mL) or ETZL (10–500 µg/mL) for 1 h, followed by stimulation with DNP-HSA (0.1 µg/mL) for 4 h. All values are the mean ± S.E.M. of three independent experiments. Values of * p < 0.05, ** p < 0.01 and *** p < 0.001 were considered significantly different to the anti-DNP IgE plus DNP-HSA, unpaired Student’s t-test. # p < 0.001 compared to the control.
We further investigated the effects of tricin and ETZL on the activation of enzymes involved in the synthesis of eicosanoids, including cPLA₂, 5-LO and COX-2, in IgE-activated RBL-2H3 cells. Both samples inhibit the phosphorylation of cPLA₂, 5-LO and COX-2, as seen in Figure 5A,B, suggesting that tricin and ETZL suppress the formation of arachidonate metabolites through inhibition of 5-LO (an early enzyme in leukotriene synthesis) and COX-2 expression (the rate-limiting enzyme in prostaglandin synthesis).

**Figure 5.** Effect of (A) tricin or (B) ETZL on phosphorylation of rate limiting enzymes in the arachidonate cascade. The expressions of p-cytosolic phospholipase A2 (cPLA₂), 5-lipoxygenase (5-LO), COX-2 and β-actin were determined as described in the Western blot method. All values are the mean ± S.E.M. of three independent experiments. Values of *p < 0.05, **p < 0.01 and ***p < 0.001 were considered significantly different to the anti-DNP IgE plus DNP-HSA, unpaired Student’s t-test. #p < 0.001 compared to the control.

Phosphorylation of cPLA₂ generates arachidonic acid (a potent metabolic precursor) from phospholipid; subsequently, the arachidonic acid metabolizing enzymes (COX-2 and 5-LO) produce PGE₂ and LTs, respectively, from arachidonic acid. Numerous studies have reported that various compounds (including flavonoids, alkaloids, terpenoids and saponins) are natural inhibitors of arachidonic acid metabolite formation and the metabolite-mediated allergic inflammation [29]. Tricin from *Oryza sativa* L. is also reported as an anti-inflammatory molecule for COX-2-mediated PGE₂ production in LPS-stimulated human peripheral blood mononuclear cells [30] as well as in human colon-derived human colon epithelial cell or HCA-7 cells (in vitro) and in *ApcMin* mice (in vivo) [9]. Our data also indicate that tricin isolated from ETZL is a natural inhibitor of allergic inflammatory LTs and PGs.

### 2.3. Effects of Tricin and ETZL on the MAPK Signaling Pathway

Since activation of MAPKs plays major roles in the formation of inflammatory cytokines, we next examined the effects of tricin and ETZL on the Akt and MAPK signaling pathways in IgE-stimulated RBL-2H3 cells [10]. The phosphorylation of Akt, p38 and JNK induces the expression of proinflammatory cytokines such as TNF-α and IL-4, and the phosphorylation of ERK activates cPLA₂ to synthesize arachidonic acid metabolites by 5-LO or COX-2, inducing the secretions of LTB₄, LTC₄ or PGE₂. Akt is a downstream effector of Syk, and the phosphorylation of Akt is involved in mast cell degranulation [14].
Proteins involved in the MAPK signaling cascade were phosphorylated upon IgE-mediated mast cell activation. However, exposure to tricin and ETZL inhibited the phosphorylation of Akt and MAPKs (such as ERK, p38 and JNK) that regulate the cytokine expression (Figure 6A,B). This indicates that tricin and ETZL directly block the activation of Syk-dependent signaling, thereby preventing the secretions of cytokines (TNF-α and IL-4) as well as arachidonate metabolites (LTB₄, LTC₄ and PGE₂).

**Figure 6.** Effect of (A) tricin and (B) ETZL on phosphorylation of proteins in the mitogen-activated protein kinase (MAPK) signaling cascade, in IgE-activated RBL-2H3 cells. IgE-sensitized RBL-2H3 cells were exposed to tricin or ETZL for 1 h, and subsequently stimulated by antigen for 10 min. All values are the mean ± S.E.M. of three independent experiments. Values of * p < 0.05, ** p < 0.01 and *** p < 0.001 were considered significantly different to the anti-DNP IgE plus DNP-HSA, unpaired Student’s t-test. † p < 0.001 compared to the control.

### 2.4. Effects of Tricin and ETZL on the Activation of the FcεRI Signaling Pathway

Upon binding of an antigen to the IgE-bound FcεRI, the signaling pathways involving PI3K/PLCγ/protein kinase C (PKC) and MAPKs are activated. The phosphorylation of PLCγ1 and PKCδ stimulates the secretion of histamine along with β-hexosaminidase and inflammatory cytokines, resulting in allergic reactions [8,31]. The phosphorylation of Lyn, Fyn and Syk is activated immediately upon FcεRI stimulation, and are known to play essential roles in the initiation of mast cell degranulation and activation [10,13,17]. Thus, down-regulating the phosphorylation of Syk and PLCγ blocks the FcεRI-mediated mast cell activation [16]. PLCγ also plays a key role in the generation of inositol-1,4,5-triphosphate to induce intracellular Ca²⁺ mobilization, which is a crucial step for mast cell degranulation [32,33]. Thus, we performed by Western blot to evaluate the effects of tricin and ETZL on activation of the IgE-FcεRI pathway in the mast cell line. Both tricin and ETZL significantly suppressed the phosphorylation of PLCγ1 and PKCδ (Figure 7A,B), thereby inhibiting the release of β-hexosaminidase and inflammatory cytokines in activated RBL-2H3 cells (Figure 2B).
Molecules 2020, 25, x 10 of 14

been reported to inhibit allergic reactions by suppressing the phosphorylation of Akt, ERK, p38 and JNK, as well as inhibiting Syk and Lyn in antigen-stimulated RBL-2H3 cells, suggesting possible therapeutic applications in allergy related diseases [17,34].

Figure 7. Inhibitory effect of tricin and ETZL on phosphorylation of proteins in the FcεRI signaling cascade in RBL-2H3 cells stimulated by an IgE–antigen complex. RBL-2H3 cells were seeded overnight in a 24-well plate at 37 °C; cultured cells were washed and further incubated with anti-DNP-IgE for 24 h. The cells were then incubated with varying concentrations of (A,C) tricin (10–500 ng/mL) or (B,D) ETZL (10–500 µg/mL) for 1 h, and subsequently stimulated by DNP-HSA for 4 h. All values are the mean ± S.E.M. of three independent experiments. Values of * p < 0.05, ** p < 0.01 and *** p < 0.001 were considered significantly different to the anti-DNP IgE plus DNP-HSA, unpaired Student’s t-test. # p < 0.001 compared to the control.

In conclusion, the present study demonstrates that ETZL exerts anti-allergic effects in IgE-stimulated mast cells, and tricin is responsible for its preventive action against the allergic reactions. The suppressive mechanism of tricin involved in the signaling cascade necessary for IgE-mediated

To investigate the regulatory effects of tricin and ETZL on the early stage of FcεRI signaling pathway, we next examined the phosphorylation levels of Fyn, Lyn and Syk in IgE-activated RBL-2H3 cells. The phosphorylation levels of Lyn and Syk were up-regulated by IgE-stimulation (2.37- and 2.56-fold, respectively, vs. untreated control), but Fyn levels were minimally altered (Figure 7C,D). Tricin and ETZL also down-regulated the enhanced expressions of Lyn (0.35- and 0.40-fold of IgE-treated control, respectively) and Syk (0.57- and 0.84-fold of IgE-treated control, respectively), but not Fyn (1.00- and
0.97-fold of IgE-treated control), suggesting that tricin and tricin-containing ETZL effectively prevent the IgE-mediated allergic reactions through inhibition of Syk-Lyn, which subsequently suppresses the activation of PLCγ and MAPK signaling in RBL-2H3 cells.

Among the numerous natural resources known to inhibit Syk and MAPK activation in mast cells, saponins obtained from roots of Platycodon grandiflorum and Loranthus parasiticus extract have been reported to inhibit allergic reactions by suppressing the phosphorylation of Akt, ERK, p38 and JNK, as well as inhibiting Syk and Lyn in antigen-stimulated RBL-2H3 cells, suggesting possible therapeutic applications in allergy related diseases [17,34].

In conclusion, the present study demonstrates that ETZL exerts anti-allergic effects in IgE-stimulated mast cells, and tricin is responsible for its preventive action against the allergic reactions. The suppressive mechanism of tricin involved in the signaling cascade necessary for IgE-mediated allergic reactions, were investigated via the FcεRI signaling pathway related to mast cell degranulation and secretions of cytokines and chemo-attractants. Tricin inhibits the release of β-hexosaminidase, production of proinflammatory cytokines TNF-α and IL-4 through the inhibition of PLCγ1/PKCδ and MAPKs pathways, and also the synthesis of arachidonate metabolites LTB4, LTC4 and PGE2 by affecting the enzyme activities of 5-LO and COX-2 as well as ERK and cPLA. Furthermore, our data demonstrates that phosphorylation of Lyn and Syk proteins involved in the early stage of FcεRI signaling pathway are effectively suppressed by tricin. In the current study, the numerous reactions that inhibit allergic responses after exposure to tricin were investigated at various phosphorylation steps in the signaling pathways triggered by antigen bound IgE-FcεRI on mast cells. Further studies are required to determine the in vivo efficacy of tricin containing ETZL, using allergy animal models for asthma, allergic rhinitis and atopic dermatitis. Overall, our data confirms that tricin and ETZL possess potential anti-allergic activity, and tricin can be used as a phytochemical to inhibit the mast cell-derived mediator release, in the development of functional products related to allergic disorders.

3. Materials and Methods

3.1. Reagents

3-(4,5-dimethylthiazol-2-yl)2,5-diphenyltetrazolium bromide (MTT), dinitrophenyl-human serum albumin (DNP-HSA), anti-DNP-IgE and 4-nitrophenyl N-acetyl-β-d-glucosaminide (p-NAG) were purchased from Sigma Aldrich Co. (St. Louis, MO, USA). Antibodies against phospho-protein kinase B (Akt; #9271), phospho-cPLA2 (#2831), phospho-extracellular signal-regulated kinase 1/2 (ERK; #9101), phospho-c-Jun N terminal kinase 1/2 (JNK; #2821), phospho-Src family protein kinase (Lyn; #2731), phospho-p38 (#9211), phospho-PKCδ (#2055), phospho-PLCγ1 (#2821), phospho-spleen tyrosine kinase (Syk; #2710), horseradish peroxidase-linked anti-rabbit secondary antibody (#7074) and anti-mouse secondary antibody (#7076) were purchased from Cell Signaling Technology (Beverly, MA, USA). Specific antibodies against phospho-Fyn (orb128087) were obtained from Biorbyt Ltd. (Cambridge, UK). A specific antibody against 5-LO (#10007820) and EIA kits for LTC4, LTB4 and PGE2 were obtained from Cayman Chemical Co. (Ann Arbor, MI, USA). ELISA kits for IL-4 and TNF-α were purchased from R&D systems, Inc. (Minneapolis, MN, USA). Minimum Eagle’s medium (MEM) and fetal bovine serum (FBS) were from the American Type Culture Collection (Manassas, VA, USA). All other chemicals and reagents used were of analytical grade.

3.2. Plant Material and Preparation of ETZL and Tricin

The aerial parts of Z. latifolia were purchased from the Pureunsan Agricultural Association Corporation (Dongdaemun-gu, Seoul, Korea) and ETZL was provided by the BTC Corporation (Sangnok-gu, Ansan, Korea). Briefly, the dried leaves of Z. latifolia were incubated with mixed hydrolysis enzymes (cellulase, hemicellulase and pectinase) at 35 °C for 16 h in H2O, followed by heating to inactivate the enzymes. The extracted solution was filtered and acquired. After extraction of the residue with 70% ethanol (Duksan Science, Seoul, Korea) at 80 °C for 6 h, the filtered and extracted
solution was mixed with the first enzyme extract and the total enzyme extract was concentrated and
dried to produce ETZL. Tricin was isolated from ETZL as previously described [8,20]. The isolated
tricin was reanalyzed by HPLC and was >99% pure.

3.3. Cell Culture

RBL-2H3 cells, a mast cell line originating from rat basophilic leukemia, were cultured in MEM
medium supplemented with 10% (v/v) FBS, 100 U/mL penicillin and 100 µg/mL streptomycin, at 37 °C
in a humidified atmosphere of 5% CO₂. All experiments contain a control group as the vehicle control
group, comprising 0.1% DMSO.

3.4. Cytotoxicities of Tricin and ETZL in RBL-2H3 Cells

Briefly, the RBL-2H3 cells were seeded in a 24-well plate (1 × 10⁵ cells/well) in MEM medium
supplemented with 10% FBS, and cultured overnight at 37 °C. The cells were washed with 1 × PBS,
and subsequently incubated with anti-DNP-IgE (0.05 µg/mL) for 24 h. The IgE-sensitized cells were
incubated with varying concentrations of tricin or ETZL (0–500 ng/mL or 0–500 µg/mL, respectively) for
1 h. DNP-HSA (0.1 µg/mL) was then added to the wells, and the mixture was incubated for a further
4 h. To measure cell viability, the absorbance was measured at 550 nm using a well-plate multi-reader.

3.5. Regulation of β-Hexosaminidase Releases of ETZL and Tricin in IgE-Stimulated RBL-2H3 Cells

The RBL-2H3 cells were incubated in a 24-well plate (1 × 10⁵ cells/well), overnight at 37 °C.
Cultured cells were washed with 1 × PBS, and subsequently incubated with anti-DNP-IgE for 24 h.
IgE-sensitized cells were exposed to varying concentrations of tricin or ETZL (0–500 ng/mL or
0–500 µg/mL, respectively) for 1 h, spiked with DNP-HSA and then incubated for a further 4 h. To
measure the amount of β-hexosaminidase activity released from the cells, the culture medium was
transferred and centrifuged (13,000 × g for 10 min) at 4 °C. The supernatant (25 µL) was mixed with
50 µL p-NAG (10 mM) in 0.1 M sodium citrate buffer (pH 4.5) in a 96-well plate, and then incubated
for 1 h at 37 °C. The reaction was terminated by adding sodium carbonate buffer (0.1 M, pH 10.0). The
β-hexosaminidase activity was determined by measuring the difference in absorbance at 405 nm.

3.6. Effect of ETZL and Tricin on the Cytokine and Chemokine Secretions in IgE-Mediated RBL-2H3 Cells

To measure the cytokine levels in the culture medium, all culture media were centrifuged (13,000 × g
for 10 min) at 4 °C, and the samples were stored at −80 °C until use. The concentrations of IL-4
and TNF-α were determined using ELISA kits (R&D systems, Inc.), according to the manufacturer’s
instructions. To determine the levels of LTB₄, LTC₄ and PGE₂ all culture media were centrifuged
(13,000 × g for 10 min) at 4 °C, and the supernatant was stored at −80 °C until use. The concentrations
of LTB₄, LTC₄ and PGE₂ were determined using EIA kits (Cayman Chemical, Inc., Ann Arbor, MI,
USA), according to the manufacturer’s instructions.

3.7. Western Blot

RBL-2H3 mast cells were lysed in lysis buffer (Pro-Prep™, iNtRON, Seongnam, Korea) containing
protease inhibitor (P3100_001, GenDEPOT, Barker, TX, USA) and phosphatase inhibitor (P3200_001,
GenDEPOT, Baker, TX, USA), with subsequent incubation on ice for 30 min. Protein concentrations
in cell lysates were determined using a BCA protein assay (Thermo Fisher Scientific, Waltham, MA,
USA). Proteins (50 µg) were loaded and separated by 10% sodium dodecyl sulfate-polyacrylamide
gel electrophoresis, transferred to PVDF membranes (Whatman GmbH, Dassel, Germany), blocked
with 5% skim milk in Tris-buffered saline containing 0.1% Tween-20 for 1 h, and then incubated with
primary antibodies for 16 h at 4 °C. After three washes in Tris-buffered saline containing 0.1% Tween-20,
membranes were incubated with horseradish peroxidase-linked secondary antibodies (Cell Signaling
Technology, Beverly, MA, USA) for 1 h. Proteins were detected by enhanced chemiluminescence, and
visualized using image software (UVP Vision Works® LS Image Acquisition & Analysis Software, Upland, CA, USA).

3.8. Statistical Analysis

Data are presented as means ± SD values of three experiments. Differences among groups were examined using Student’s t-test, and a significant difference is considered at $p < 0.05$, $p < 0.01$ and $p < 0.001$.

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**Sample Availability:** Samples of the compounds are available from the BTC Corporation.

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