
Inverse Agonist of Estrogen-Related Receptor γ Enhances Sodium Iodide Symporter Function Through Mitogen-Activated Protein Kinase Signaling in Anaplastic Thyroid Cancer Cells

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Anaplastic thyroid cancer (ATC), a rare thyroid cancer with poor prognosis, is associated with insufficient function of the sodium iodide symporter (NIS). Estrogen-related receptor γ (ERR γ) is a member of the orphan nuclear receptors with important functions in cell development and homeostasis. However, there are no reports that demonstrate whether ERR γ is related to NIS function. Here, we evaluated the role of ERR γ in the regulation of NIS function in ATC cells using GSK5182, an inverse agonist of ERR γ . **Methods:** Two ATC cell lines, BHT-101 and CAL62, were incubated with GSK5182 at various time points and doses. The NIS function in the ATC cells was serially assessed by their uptake of radioiodine. The effects of GSK5182 on ERR γ and the mitogen-activated protein (MAP) kinase pathway, as well as on NIS protein, were evaluated by immunoblot assay. To examine whether the GSK5182-induced NIS functional activity can be affected by inhibition of the MAP kinase pathway, the MAP kinase activity and levels of radioiodine uptake were determined after application of a mitogen-activated protein kinase kinase (MEK) inhibitor to GSK5182-treated cells. Finally, the cytotoxic effect of ¹³¹I was determined by clonogenic assay. **Results:** Treatment with GSK5182 resulted in dose- and time-dependent increases in iodide uptake in ATC cells, which were accompanied by both the downregulation of ERR γ protein and the activation of extracellular signal-regulated kinase (ERK) 1/2. Both the increased radioiodine uptake and ERK1/2 activation of ATC cells were completely inhibited by the specific MEK inhibitor. GSK5182 treatment enhanced the membrane localization of NIS in both ATC cell lines. Accordingly, preexposure to GSK5182 enhanced the cytotoxic effects of ¹³¹I treatment in ATC cells. **Conclusion:** These findings suggest that the inverse agonist of ERR γ enhances the responsiveness of radioiodine therapy by modulating NIS function in ATC cells via the regulation of ERR γ and the MAP kinase signaling pathway.

Key Words: sodium iodide symporter (NIS); anaplastic thyroid cancer; estrogen-related receptor gamma (ERR γ); radioiodide therapy; MAP kinase signaling

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Anaplastic thyroid cancer (ATC) is one of the most aggressive and lethal cancers known to affect humans (1,2). ATC rapidly metastasizes from the thyroid glands to the lungs, bone, regional lymph nodes, and brain (3). This is in contrast to the benign nature of well-differentiated thyroid cancer, which accounts for most thyroid cancers. Treatment of ATC with surgery, radiotherapy, and chemotherapy alone or in combination has had little to no effect on patient survival (4–7). Consequently, the development of novel therapeutic approaches is urgently required.

The sodium iodide symporter (NIS) is a plasma membrane glycoprotein that mediates the active influx of iodide into cells (8). In thyroid cancer, endogenous NIS accommodates the widespread clinical application of radioiodine therapy, which has been shown over the years to be an effective method of eliminating malignant cells, with minimal adverse effects. Poorly differentiated cancer cells, including ATC cells, tend to exhibit a progressive dedifferentiation that leads to a decrease in levels of NIS (9,10). This results in the inability of ATC cells to accumulate a high iodine concentration and hence their resistance to radionuclide therapy, which finally leads to poor prognosis (11). Thus, many attempts have been made to restore NIS function in ATC cells using methods such as gene delivery (12,13) and epigenetic modulation with epigenome-modifying drugs (14), but few satisfactory outcomes have been achieved.

Estrogen-related receptors (ERR α , ERR β , and ERR γ) are constitutively active nuclear receptors that bear high levels of sequence identity to estrogen receptors (15). The ERR isoforms are primarily expressed in several organs, such as the heart, brain, kidney, pancreas, and liver (16,17). Recently, studies have shown that ERR γ is involved in metabolic diseases such as type 2 diabetes mellitus, alcohol-induced oxidative stress, liver injury, and microbial infection

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through impaired hepatic gluconeogenesis (18,19), hepatic insulin signaling (20), and iron metabolism (21).

Crystal structure studies have demonstrated that ERRs are constitutively active without their ligand, and several small-molecule ligands can either activate or repress the functional activity of the ERRs (22). Among them, GSK5182 (a 4-hydroxy tamoxifen analog) is a selective inverse agonist of ERR γ that does not interact with another nuclear receptor. Several lines of evidence have revealed that GSK5182 not only alleviates diabetes symptoms through the inhibition of hepatic gluconeogenesis in a PGC-1 α (peroxisome proliferator-activated receptor γ coactivator-1 α)-dependent manner (19,23) but also shows antimicrobial effects by the reduction of ERR γ -mediated hepcidin messenger RNA expression (21). Although the biologic effects of ERR γ have been extensively investigated in various disease models, the role of ERR γ in NIS function in ATC remains to be clearly determined.

Among various nuclear receptors the retinoic acid receptor, through its activator, retinoic acid, has been proven effective in restimulating iodide uptake in nonthyroidal cancer and thyroid cancer in vitro and in vivo (24,25). We speculated that the ERR γ of the nuclear receptor family might also be involved in controlling NIS function in thyroid cancer, similar to the biologic roles of RA and RAR on NIS function.

To explore the effects of ERR γ on NIS function in ATC cells, we adopted GSK5182 as a specific inverse agonist of ERR γ and investigated the changes in endogenous ERR γ protein and MAP kinase signaling by this agent. The effects of GSK5182 on the functional activity and expression level of NIS protein in ATC cells were also assessed. Finally, the enhanced cytotoxic effects of ¹³¹I from GSK5182 treatment were evaluated in ATC cells.

MATERIALS AND METHODS

Cells

Two ATC cell lines, BHT-101 and CAL-62, were purchased from Deutsche Sammlung von Mikroorganismen und Zellkulturen. Both cell lines were maintained in high-glucose Dulbecco modified Eagle medium supplemented with 10% fetal bovine serum and a 1% antibiotic-antimycotic (HyClone; GE Healthcare) at 37°C in a 5% CO₂ atmosphere.

¹²⁵I Uptake Assay

Cells were plated in 24-well plates for 24 h and then treated with GSK5182 synthesized by Daegu-Gyeongbuk Medical Innovation Foundation and prepared as a 100 mM stock solution in dimethyl sulfoxide and stored at -80°C for 24 h. After aspiration of drug-containing medium, the cells were washed with 1 mL of Hanks balanced salt solution (HBSS) and incubated with 500 μ L of HBSS containing 0.5% bovine serum albumin (bHBSS), 3.7 kBq of carrier-free ¹²⁵I (Perkin-Elmer), and a 10 μ mol/L solution of sodium iodide (specific activity of 740 MBq/mmol) at 37°C for 30 min. The cells were then washed twice with ice-cold bHBSS and were lysed with 500 μ L of 2% sodium dodecyl sulfate. The radioactivity was measured using a γ counter (Cobra II; Canberra Packard, Packard Bioscience). The radioactivity of the cells was normalized using total protein concentrations determined by a bicinchoninic acid protein assay kit (Pierce Protein Biology). The cells were preincubated with 300 μ M KClO₄ (as a specific inhibitor for NIS) for 30 min to inhibit iodide uptake, followed by a ¹²⁵I uptake test as described above.

Clonogenic Assay

The cells were plated into 6-well plates and left for 24 h. After treatment with 25 μ M GSK5182 for 24 h, drug-containing medium

was discarded and the cells were washed twice with phosphate-buffered saline (PBS). The medium was then replaced with Dulbecco modified Eagle medium in the presence or absence of 1,850 kBq (50 μ Ci) of ¹³¹I (Korea Institute of Radiological and Medical Sciences) for 6 h. The cells were washed with cold bHBSS and left in regular culture medium for the time corresponding to 6 doublings. Finally, the cells were fixed in 4% paraformaldehyde solution and stained with 0.05% crystal violet. Control and ¹³¹I treated colonies with more than 50 cells were counted.

Western Blot

The cells were treated with or without GSK5182 for 24 h and washed twice with cold PBS and lysed with radioimmunoprecipitation assay buffer containing complete protease inhibitor cocktail (Roche). In case of plasma membrane protein for NIS, samples were prepared with a protein biotinylation kit (EZ-Link sulfo-NHS-biotin; Thermo Scientific) according to the manufacturer's instructions. Briefly, either untreated or treated cells were washed twice with ice-cold PBS/CM (PBS containing 0.1 mM calcium chloride and 1 mM magnesium chloride, pH 7.3) and incubated with EZ link NHS-sulfo-SS-biotin (1 mg/mL) in PBS/CM for 30 min at 4°C. The reaction was quenched by 2 washes with cold 100 mM glycine in PBS/CM and further incubation with 100 mM glycine in PBS/CM at 4°C for 20 min. The cells were then quickly washed 2 times with PBS/CM before lysis with radioimmunoprecipitation assay buffer containing a protease inhibitor cocktail and phosphatase inhibitors (Roche) for 1 h at 4°C with constant shaking. Lysates were centrifuged at 16,000g for 30 min at 4°C. A portion of the supernatant was used for total cell protein immunoblots. The remaining sample was used to obtain membrane protein by incubation with 100 μ L of streptavidin beads (Thermo Scientific) for 1 h at room temperature. The beads were washed 3 times with radioimmunoprecipitation assay buffer, and bound proteins were eluted with 50 μ L of Laemmli buffer (62.5 M Tris, pH 6.8; 20% glycerol; 2% sodium dodecyl sulfate; 5% b-mercaptoethanol; and 0.01% bromphenol blue) for 30 min at room temperature. Equal amounts of total and biotinylated plasma membrane protein were loaded in each lane and resolved by 4%–12% gradient Bis-Tris gel (Invitrogen). Proteins were transferred to 0.2- μ m polyvinylidene fluoride membrane (Invitrogen). Membranes were incubated overnight at 4°C with primary mouse monoclonal human NIS-specific antibody (dilution, 1:1,000, Thermo Scientific, catalog no. MS-1653-P1, clone: FP5A), followed by incubation with horseradish peroxidase-conjugated secondary antibody at room temperature. ECL-Plus (Amersham Pharmacia) was used to detect peroxidase activity according to the manufacturer's protocol.

Similarly, for other proteins also, equal amounts of protein were loaded in each lane and resolved by 4%–12% gradient Bis-Tris gel (Invitrogen). Proteins were transferred to a 0.2- μ m polyvinylidene fluoride membrane (Invitrogen). The membranes were incubated overnight at 4°C with primary antibodies and then with the appropriate horseradish peroxidase-conjugated secondary antibody at room temperature. ECL-Plus was used to detect peroxidase activity according to the manufacturer's protocol. Band densities were determined by ImageJ software.

Statistical Analysis

All data are expressed as mean \pm SD, and statistical significance was determined using an unpaired Student test in Prism 5 (GraphPad). *P* values of less than 0.05 were considered statistically significant.

RESULTS

Enhanced Uptake in ATC Cells by Inverse Agonist of ERR γ

Treatment with GSK5182 significantly increased radioiodine uptake in both ATC cell lines in a dose-dependent fashion (Fig. 1A). The maximum relative increase in iodide uptake was 2.3- and 2.9-fold in the BHT-101 and CAL-62 cells, respectively, compared with the

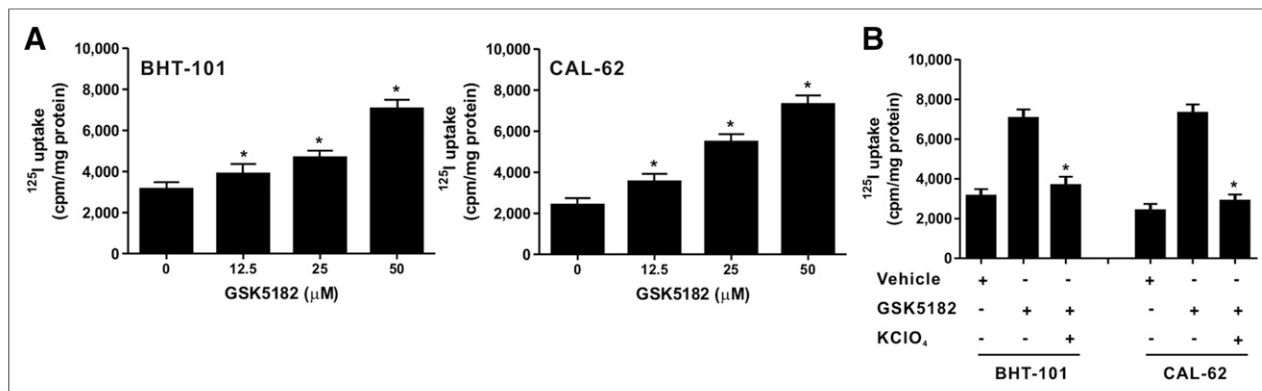


FIGURE 1. Effects of GSK5182 on kinetics of iodide uptake in the 2 ATC cell lines. (A) Each cell line was treated with various doses of GSK5182 for 24 h, and dose-dependent increased iodide uptake was confirmed in both. (B) ¹²⁵I uptake was inhibited in both cell lines by treatment with 300 μM KClO₄ for 30 min. **P* < 0.05 when compared with GSK5182. Data are mean ± SD of 3 samples per group.

vehicle group, which was obtained at a concentration of 50 μM GSK5182. Accordingly, mainly 50 μM GSK5182 was used for the further iodide uptake tests.

To examine whether the increased radioiodine uptake is related to modulation of NIS function by GSK5182, KClO₄, a specific inhibitor of NIS, was coincubated with GSK5182-treated BHT-101 and CAL-62 cells and the change in radioiodine uptake levels was determined. The level of radioiodine uptake between vehicle-treated cells and combined vehicle and KClO₄-treated cells did not differ between the two ATC cell lines, indicating that these lines have negligible NIS-mediated iodide uptake (Supplemental Fig. 1; supplemental materials are available at <http://jnm.snmjournals.org>). KClO₄ completely blocked the enhanced radioiodine uptake in both types of GSK5182-treated cells (Fig. 1B), suggesting that the augmentation of iodide uptake is directly involved in the improved functional activity of NIS mediated by GSK5182.

We assessed the radioiodine uptake level in GSK5182-treated BHT-101 and CAL-62 cells according to different time points. Radioiodine uptake increased in both treated cell lines within as early as 2 h after treatment and peaked at 24 h (Supplemental Fig. 2).

Downregulation of Endogenous ERRγ by GSK5182 in ATC Cells

To determine the effect of GSK5182 on ERRγ protein levels in ATC cells, an immunoblotting analysis was conducted using an ERRγ-specific antibody. Both BHT-101 cells and CAL-62 cells showed endogenous expression of ERRγ protein. Treatment with GSK5182 markedly reduced ERRγ protein in both cells, with a relative decrease of 3.6-fold and 2.4-fold in the BHT-101 and CAL-62 cells, respectively (Fig. 2).

Increase in Membrane-Localized NIS in ATC cells Through Activation of MAP Kinase Signaling

A significant increase in phosphorylated MAP kinase levels, such as p44 and p42 extracellular signal-regulated kinase (ERK), was found in both ATC cell lines treated with GSK5182 (Fig. 3A). The relative increase in the phosphorylated forms of ERK1 and ERK2 was 1.6- and 2.1-fold, respectively, in BHT-101 cells. GSK5182 treatment resulted in 5.8- and 2.2-fold increases in phosphorylated p44 and p42 ERK, respectively, in CAL-62 cells and was completely inhibited by the selective mitogen-activated protein kinase kinase (MEK) inhibitor, PD98059 (Fig. 3B).

To determine the effect of GSK5182 on NIS protein status, we examined the change in levels of membranous and total NIS

proteins collected from GSK5182-treated ATC cells using the plasma membrane biotinylation kit, followed by immunoblotting with NIS-specific antibody. As illustrated in Figure 4A, GSK5182 induced drastic increases in plasma membrane-localized NIS

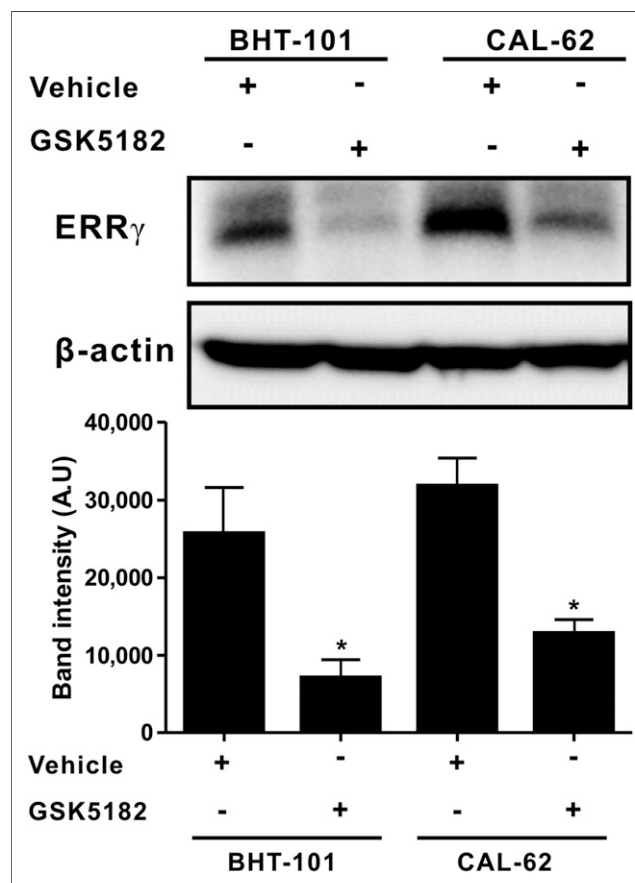


FIGURE 2. Effects of GSK5182 on endogenous ERRγ protein in ATC cells. Endogenously expressed ERRγ protein levels in ATC cells were decreased by GSK5182. Each cell line was treated with 50 μM GSK5182 for 24 h, and immunoblotting was done with ERRγ-specific antibody. Graphs depicts results of quantitative analysis of ERRγ protein levels by calibrated densitometer. **P* < 0.05 when compared with untreated cells. Data are mean ± SD of 3 samples per group, expressed in arbitrary units.

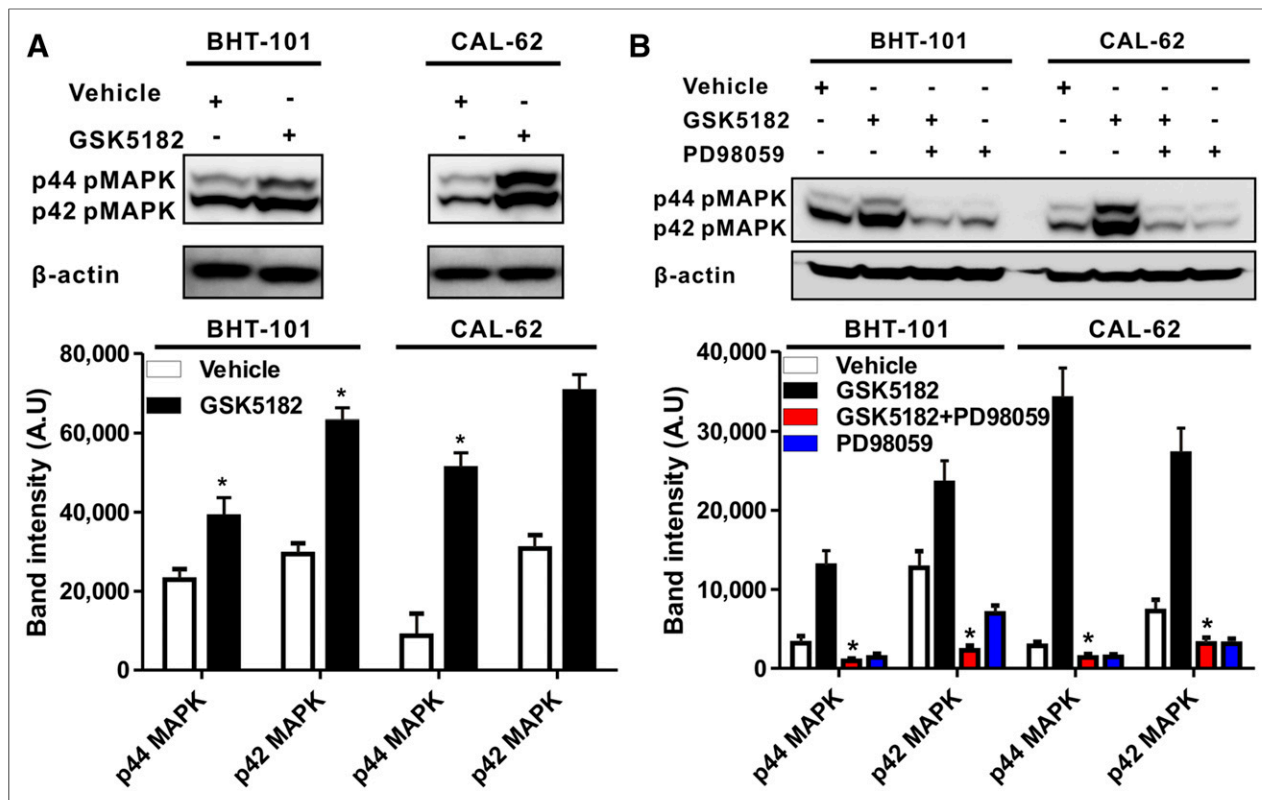


FIGURE 3. GSK5182-induced MAP kinase activation in ATC cells. (A) Activation of MAP kinase signaling in ATC cells by GSK5182. Each cell line was incubated with 50 μ M GSK5182 for 24 h, and phosphorylated ERK1/2 levels were determined with phospho-p44/42 specific antibody. Graphs depict quantitative analysis of phosphorylated p44/42-ERK levels by scanning densitometry. * $P < 0.05$ when compared with untreated cells. Data are mean \pm SD of 3 samples per group, expressed in arbitrary units. (B) Reversal of activated MAK kinase signaling by PD98059. Cells were cotreated with both 20 μ M PD98059 and GSK5182, and phosphorylated ERK1/2 levels were determined with immunoblotting. Graphs depict quantitative analysis of phosphorylated p44/42-ERK levels by scanning densitometry after cotreatment with both 20 μ M PD98059 and GSK5182. * $P < 0.05$ when compared with GSK5182 cells. Data are mean \pm SD of 3 samples per group, expressed in arbitrary units.

protein, with an immature form in both ATC cell lines, compared with control cells. Quantitative analysis of the band intensity revealed 3.8- and 6.0-fold increases in membrane NIS protein in the BHT-101 and CAL-62 cells, respectively (Supplemental Fig. 3). However, there was no significant change in the levels of NIS protein in total cell lysates of untreated and GSK5182-treated ATCs. A radioiodine uptake test with PD98059 treatment demonstrated that the GSK5182-induced iodide uptake reverted to the basal level in both ATC cell lines, whereas incubation of cells with PD98059 alone did not affect iodide uptake (Fig. 4B).

Amelioration of ^{131}I -Mediated Cytotoxicity by GSK5182 in ATCs

As shown in Figure 5 and Supplemental Figure 4, clonogenic assay with ^{131}I demonstrated minimal cytotoxic effects in the ATC cells treated with either GSK5182 or ^{131}I alone. The relative colony-forming ability in CAL-62 cells for the ^{131}I and GSK5182 groups was $92.9\% \pm 5.8\%$ and $94.5\% \pm 10.8\%$, respectively. Similar to CAL-62 cells, BHT-101 cells had a relative colony-forming ability of $95.2\% \pm 4.2\%$ and $93.2\% \pm 5.5\%$ in the ^{131}I and GSK5182 groups, respectively. However, the combination of GSK5182 with ^{131}I markedly reduced the colony-forming ability to approximately $58.5\% \pm 7.4\%$ and $72.8\% \pm 2.4\%$ in CAL-62 and BHT-101, respectively.

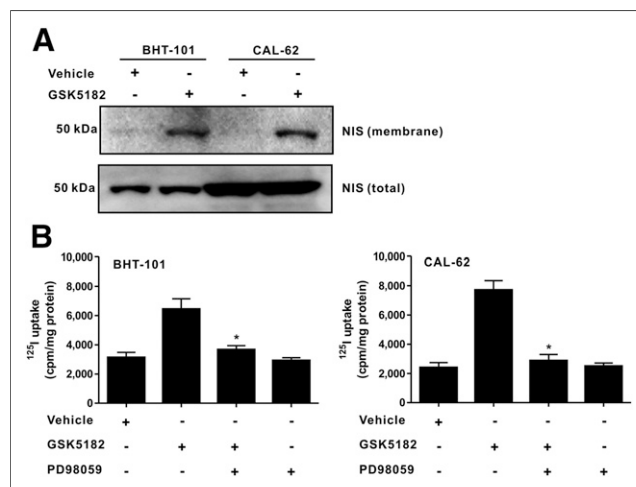


FIGURE 4. Enhancement of membrane-localized NIS protein level by GSK5182. (A) Detection of membrane and total NIS protein. Membrane fraction of cells treated with or without GSK5182 was collected and immunoblotting analysis was done with NIS-specific antibody. (B) Inhibition of increased iodide uptake by PD98059 in GSK5182-treated ATC cells. Cells were cotreated with both 20 μ M PD98059 and GSK5182, and iodide uptake levels were determined. * $P < 0.05$ when compared with GSK5182. Data are mean \pm SD of 3 samples per group.

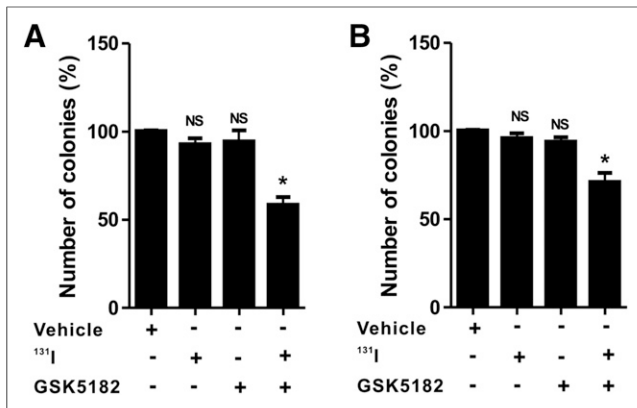


FIGURE 5. Increased cytotoxicity of ¹³¹I by GSK5182 against ATC cells. CAL-62 (A) and BHT-101 (B) cells were pretreated with or without GSK5182 before day 1. After being washed, they were further coincubated with 1,850 kBq (50 μCi) of ¹³¹I for 6 h. Graphs depict percentage of colonies after ¹³¹I treatment. **P* < 0.05. NS = not significant when compared with untreated cells. Data are mean ± SD of 3 samples per group.

DISCUSSION

We have demonstrated that GSK5182, an inverse agonist of ERRγ, reduces the endogenous ERRγ protein level in ATC cells and leads to the activation of MAP kinase signaling (Fig. 6). Importantly, treatment with GSK5182 enhances radioiodine uptake in ATC cells in a dose- and time-dependent fashion. Both the GSK5182-induced ERK1/2 activation and the increase in radioiodine uptake are completely inhibited by PD98059, a selective MEK inhibitor. Further examination showed that GSK5182 increases NIS plasma membrane protein without affecting levels of total NIS protein. Furthermore, the enhancement of NIS function modulated by GSK5182 leads to increased susceptibility of ATC cells to radioiodine therapy.

Recently, many studies have shown interesting therapeutic outcomes from GSK5182-mediated ERRγ regulation in various

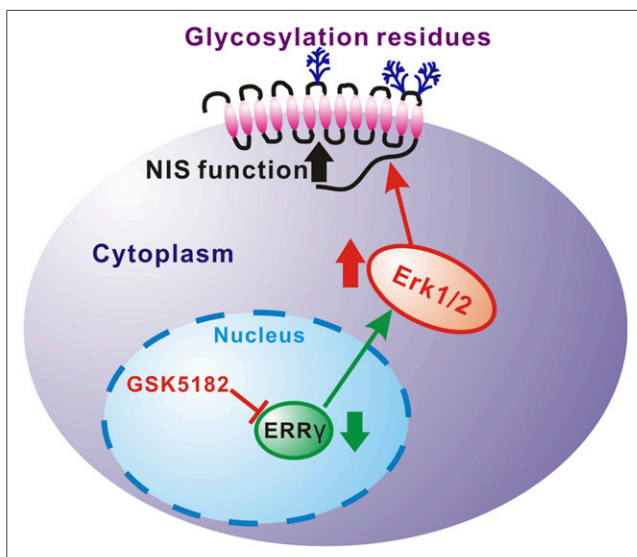


FIGURE 6. Proposed mechanism of GSK5182-induced modulation of NIS function in ATC cells through MAP kinase pathway.

metabolic and cardiac diseases (18,20,21,26), but no studies have unraveled the involvement of GSK5182-modified ERRγ in regulating NIS function in thyroid cancer, especially ATC. In an attempt to determine the effects of GSK5182 on radioiodine uptake in ATC cells, we selected 2 different ATC cells, BHT-101 and CAL-62, which have characteristic mutations of the K-RAS and BRAF genes, respectively. These cell lines have been used to evaluate the therapeutic efficacy of several drugs in inducing cytotoxic effects directly as well as modulating NIS function (14,27). Interestingly, the incubation of ATC cells with GSK5182 led to dose- and time-dependent increases in radioiodine uptake, but negligible uptake was shown in vehicle-treated cells. From these findings, we questioned whether the increased iodide uptake was associated with modulated NIS function. Thus, an iodide uptake test was further performed by introducing a specific inhibitor of NIS protein, potassium perchlorate (KClO₄), to GSK5182-treated cells. KClO₄ has been widely used to validate the functional activity of NIS protein (8). Iodide uptake analysis with KClO₄ clearly showed the inhibition of enhanced radioiodine uptake to basal level, revealing that increased iodide uptake is related to the GSK5182-induced modulation of NIS function. Next, we speculated that the enhanced iodide uptake might be involved in a change in endogenous ERRγ protein by GSK5182. We observed that both ATC cell lines exhibited a substantial and similar level of expression of ERRγ protein, and treatment with GSK5182 significantly reduced the ERRγ protein level. Taken together, these results suggest that the GSK5182-induced reduction of endogenous ERRγ protein may affect NIS function and finally lead to increased radioiodine uptake in ATC cells.

MAP kinases are evolutionarily conserved enzymes that transfer signals from cell surface receptors to critical intracellular molecules (28). Among the MAP kinases, p38 MAP kinase has been shown to have a role in maintaining normal basal expression of serotonin transporters (29) and improving the activation of norepinephrine transporters (30). Knostman et al. reported that PI3K activation induces underglycosylated intracellular NIS protein expression in breast cancer (MCF-7) cells and leads to iodide uptake ability (31). Another report has shown that MEK inhibitor decreases NIS protein levels through lysosome-mediated NIS degradation in human breast cancer cells, suggesting that MEK activation has as an important role in maintaining NIS protein stability in human breast cancers (32). More recently, Lee et al. (12) reported that activation of protein kinase C signaling with phorbol 12-myristate 13-acetate down-regulated iodide uptake in NIS-expressing nonthyroidal cancer cells and that EGF-mediated MAP kinase activation enhanced radioiodine uptake. On the basis of these studies, it was presumed that the enhanced radioiodine uptake of ATC cells might be linked to upregulation of these kinase pathways. From an immunoblotting analysis with phospho-ERK1/2-specific antibody, we found that ERK signaling is constitutively active in both ATC cell lines and that treatment with GSK5182 increases the phosphorylated MAP kinase level. Conversely, PD98059, a selective inhibitor of MAP kinase, inhibits the increased phosphorylation of ERK1/2 MAP kinase as well as the iodide uptake induced by GSK5182. It would therefore be reasonable to postulate that the enhanced radioiodine uptake of ATC cells may be associated with the activation of MAP kinase signaling that results from the GSK5182-mediated reduction of ERRγ protein.

The total amount of NIS protein, as well as plasma-membrane-localized NIS protein, is crucial to the capacity of thyroid cells to transport iodide. Even though sufficient expression of NIS protein is

observed in thyroid cancer cells, intracellular retention of NIS prevents iodide uptake from being sufficient to generate an effective radioiodine therapy. The NIS transporter needs to be translocated to the plasma membrane to be fully functionalized in thyroid cells. Thyroid-stimulating hormone is needed to generate membrane retention of NIS, and its depletion leads to an acute decrease in iodide uptake in thyroid cells (33). Furthermore, breast cancer cells with sufficient endogenous expression of NIS had a poor ability to take up iodide, partially resulting from intracellular localization of NIS protein (24,34).

On the basis of these reports, we speculated that both the downregulation of ERR γ protein level and activation of MAP kinase by GSK5182 may affect total and plasma membrane NIS protein status. We could detect the endogenously expressed total NIS protein without an increase in membrane NIS protein in vehicle-treated cells. Although several reports have demonstrated that ATC cells do not express NIS protein (35), we could evidently detect NIS protein expression in both ATC cell lines in this study. The discrepancy may be due to the development of anti-NIS-specific antibody with high sensitivity and specificity as well as better experimental conditions. Surprisingly, biotinylation experiments with GSK5182-treated ATC cells revealed an increase in immature NIS protein at the plasma membrane. But there was no change in the NIS protein in total cell lysate protein. The exact mechanism responsible for the GSK5182-mediated increase in iodide transport is not yet clear. Because the general posttranscriptional mechanism of both the localization and the function of many transporters includes the glycosylation and phosphorylation processes (36), similar processes may be involved in NIS regulation.

Successful radioiodine therapy against thyroid cancer depends mainly on effective accumulation of radioiodine. It would be reasonable to postulate that the enhanced radioiodine uptake brought about by GSK5182 may induce cytotoxic effects against ATC cells. Notably, in our study GSK5182 treatment was sufficient to enhance the killing effect of therapeutic radioiodine on ATC cells, but treatment with ¹³¹I alone could not induce sufficient cytotoxic effects on ATC cells. These findings suggest that ERR γ -modulated NIS function can be a reasonable approach to achieving a radioiodine accumulation that allows for acceptable therapeutic outcomes with clinically used doses of ¹³¹I. Further preclinical investigations using nuclear medicine imaging instruments such as SPECT or PET are required to evaluate the effects of GSK5182-mediated MAP kinase activation on the efficacy of radioiodine accumulation in ATC cancer in living mice.

CONCLUSION

Our findings suggest that ERR γ -regulated MAP kinase pathways play an important role in regulating NIS function. The findings also suggest that it is realistically feasible to appropriately regulate these signaling pathways to enhance NIS function and enable radioiodine therapy, thus allowing new therapeutic strategies for ATC to be uncovered. In the future, the ERR γ inverse agonist can be introduced as an adjuvant in the development of new therapeutic protocols for patients with ATCs. Our current study guarantees further investigations for new drug candidates or GSK5182 derivatives to regulate ERR γ activity and mediate a more effective reinduction of radioiodine uptake in ATC cells. Because the lack of iodide organification limits effective radioiodine therapy in ATC cells, further study is needed to learn the effects of GSK5182 on iodide organification in ATC cells.

DISCLOSURE

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