Olanzapine Increases RGS7 Protein Expression via Stimulation of the Janus Tyrosine Kinase-Signal Transducer and Activator of Transcription Signaling Cascade

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ABSTRACT

Atypical antipsychotics such as olanzapine have high affinity for multiple monoamine neurotransmitter receptors and are the mainstay of pharmacological therapy for treatment of schizophrenia. In addition to blocking monoamine receptors, these drugs also affect intracellular signaling cascades. We now report that 24-h treatment with 300 nM olanzapine causes desensitization of serotonin (5-HT)2A receptors in A1A1v cells, a rat cortical cell line, as indicated by a reduction in inositol phosphate accumulation following stimulation with a 5-HT2A/2C receptor agonist (−)−1-(2,5-dimethoxy-4-lodophenyl)-2-aminopropane HCl. Olanzapine treatment for 24 h increased the levels of 5-HT2A receptors in both cytosol (254 ± 34% of control levels) and membrane fractions (206 ± 14% of control levels) and RGS7 proteins in both cytosol (193 ± 32% of control levels) and membrane fractions (160 ± 18% of control levels) as measured on Western blots. Increased phosphorylation of Janus tyrosine kinase (JAK) 2 and increased phosphorylation and nuclear translocation of signal transducer and activator of transcription (STAT) 3 with 24-h olanzapine treatment demonstrate activation of the JAK-STAT signaling cascade. Pretreatment with a JAK inhibitor, AG490 [α-cyano-(3,4-dihydroxy)-N-benzylcinnamamide], prevented the olanzapine-induced increase in membrane RGS7 protein levels; AG490 alone had no effect on RGS7 protein levels. We verified that treatment with AG490 reduced phosphorylation of JAK2 and inhibited the nuclear localization of phospho-STAT3. Interestingly, treatment with the JAK inhibitor had no effect on 5-HT2A receptor protein levels. These data suggest that olanzapine-induced activation of the JAK-STAT signaling cascade causes increased expression of RGS7 protein, which in turn could mediate desensitization of 5-HT2A receptor signal transduction caused by olanzapine because RGS7 binds to Goq protein and accelerates GTP hydrolysis.

Atypical antipsychotics are widely prescribed for the treatment of schizophrenia. They are classified as atypical because of their ability to achieve antipsychotic effects with lower rates of extrapyramidal side effects compared with first generation antipsychotics such as haloperidol. In addition, selected atypical antipsychotics also improve certain aspects of cognitive function in schizophrenic patients, whereas typical antipsychotics may worsen cognition (Meltzer et al., 1999). A typical antipsychotics improved side efficacies and efficacy has been attributed to the high-affinity interaction with 5-HT2A receptors (Kasper et al., 1999). Atypical antipsychotics have also been shown to block other 5-HT3 receptor subtypes, mainly 5-HT3A and 5-HT3C (Lucaites et al., 1996). However, only 5-HT3C receptor antagonism is suggested in contributing to the atypical antipsychotic effects (Herrick-Davis et al., 2000; Rauser et al., 2001).

The 5-HT2A receptor subtype has been implicated in various psychiatric disorders including depression, anxiety, and schizophrenia (Glennon et al., 1984). Olanzapine is an atypical antipsychotic, approved for the treatment of schizophrenia and bipolar disorder. Olanzapine has been also studied in the treatment of disorders such as substance abuse, aggression/violence, borderline personality disorder, and obsessive-compulsive disorder (Littrell et al., 2006).

Atypical antipsychotics as well as a specific 5-HT2A receptor antagonist, MDL 100,907, desensitize 5-HT2A-mediated responses (Willins et al., 1999). However, the molecular
mechanisms involved in antagonist-induced desensitization of 5-HT₂A receptor signaling are not well understood. By understanding the molecular mechanisms underlying the effects of olanzapine and other atypical antipsychotics, we hope to gain insight into targets for therapeutic treatment of psychiatric disorders. It has been recently reported that olanzapine increases extracellular receptor kinase (ERK) 1/2 phosphorylation in rat prefrontal cortex (Fumagalli et al., 2006). Furthermore, changes in mRNA levels of both 5-HT₂A receptors with 5-HT₂A receptor antagonists after both chronic and short-term treatments have been reported (Buckland et al., 1997). These studies suggest that atypical antipsychotics may target intracellular pathways shutting information from the receptor to the nucleus.

The Janus tyrosine kinase (JAK)-signal transducer and activator of transcription (STAT) signaling cascade has been reported to couple with 5-HT₂A receptors in skeletal muscles and vascular smooth muscle cells (Guillet-Deniaux et al., 1997; Banes et al., 2005). G protein-coupled receptor agonists, thrombin and angiotensin II, have previously been shown to activate the JAK-STAT signaling cascade (Bhat et al., 1994). JAK-STAT could be one of the possible signaling pathways involved in mediating olanzapine-induced receptor desensitization. JAKs are a small family of cytoplasmic tyrosine kinases initially identified as a mediator of cytokine receptor signaling (Ihle, 1995). Agonist stimulation of cytokine receptors causes phosphorylation of JAK, which in turn phosphorylates tyrosine residues on its target proteins, facilitating activation of specific STATs. Tyrosine phosphorylated STATs then undergoes dimerization and translocates to the nucleus, where it binds to target DNA sequences (Darnell, 1997).

 Alterations in proximal components of the 5-HT₂A receptor signaling system could mediate desensitization in response to increased activity of intracellular cascades such as JAK-STAT. 5-HT₂A receptors are classically linked to the Gα₁/G₁ protein family (Ivins and Molinoff, 1990). Activation of Gα₁/G₁ stimulates phospholipase C activity, which subsequently promotes the release of diacylglycerol and inositol triphosphate, which in turn stimulate protein kinase C activity and calcium release (Berg et al., 2001). It has been extensively reported that increased expression of regulator of G protein signaling (RGS) proteins cause desensitization of several G protein-activated receptor systems (Koelle and Horvitz, 1996). RGS proteins reduce the duration of signaling of many G protein-coupled receptors by their action as GTPases, accelerating the hydrolysis of GTP-bound Gα proteins or by blocking the interaction of Gα with its target proteins through a not well understood process known as effector antagonism (Roy et al., 2006).

Expression of RGS7 protein in rat frontal cortex is well documented (Zhang and Simonds, 2000; Krumins et al., 2004) and decreased 5-HT₂A receptor-mediated signaling via direct interaction of RGST protein with Gα₁, has been widely characterized in different systems (DiBello et al., 1998; Ghamvami et al., 2004). We hypothesize that the increased expression of RGS7 protein by the JAK-STAT signaling cascade contributes to olanzapine-induced desensitization of 5-HT₂A receptor signaling. In this study, we examined 5-HT₂A receptor and RGS7 protein levels in response to treatment with olanzapine and determined whether changes in these proteins are mediated by olanzapine-induced JAK-STAT signaling in A1A1v cells.

Materials and Methods

Cell Culture. A1A1v cells, a cortical cell line that expresses 5-HT₂A receptors, were used for all experiments and were generously donated by Dr. William Clarke and Kelly Berg (University of Texas Health Science Center, San Antonio, TX). Cells were grown in Dulbecco's modified Eagle's medium supplemented with 10% fetal bovine serum. Serotonin was removed from serum by filtration after treating with charcoal. Cells were grown in the serotonin-free serum media 24 h before treatment with olanzapine. Olanzapine was a generous gift from Eli Lilly & Co. (Indianapolis, IN). The treatment concentration (300 nM) was obtained by dissolving olanzapine in 20% acetic acid. The pH of the vehicle and olanzapine was adjusted to 6.5 with 10 N NaOH. A JAK inhibitor, AG490, was purchased from Calbiochem (San Diego, CA). It was reconstituted with dimethyl sulfoxide to obtain the desired concentration. (−)-1-(2,5-Dimethoxy-4-lodophenyl)-2-aminopropane HCl (DOI) was purchased from Sigma-Aldrich (St. Louis, MO). Hanks' balanced salt solution mix (1× Hanks' balanced salt solution, 20 mM LiCl₂, and 20 mM HEPES) was used to dissolve DOI. Cells were treated with either vehicle (20% acetic acid) or 300 nM olanzapine for 24 h for various experiments.

Cell Fractionation. Cells were separated into membrane and cytosol fractions using ultracentrifugation as described previously (Tucker, 2004). In brief, cells were collected in hypotonic buffer containing 0.25 M sucrose, 50 mM Tris-HCl, 5 mM EDTA, and protease inhibitor cocktail (Sigma-Aldrich). Cell homogenate was prepared by sonication three times for 10 s. The homogenate was spun at 100,000 g for 45 min at 4°C to produce a pellet, which is composed of membrane fraction and a supernatant, which is the cytosol fraction. The pellet was reconstituted with hypotonic buffer.

The protein amount was assessed with bicinchoninic acid protein assay kit (Pierce Chemical, Rockford, IL). Cytosolic and nuclear fractions were prepared as described previously (Andrews and Fuller, 1991) with some modification. In brief, cells were washed and scrunched into ice-cold PBS containing phosphatase inhibitors. The pellet was collected after centrifugation at 1000 g for 1 min and resuspended into buffer A (10 mM HEPES-KOH, pH 7.9, 1.5 mM MgCl₂, 10 mM KCl, 0.5 mM dithiothreitol, and protease inhibitor cocktail) and incubated for 10 min on ice followed by vortex mixing for 10 s. The supernatant contains the cytosolic fraction. The pellet was resuspended in buffer B (10 mM HEPES-KOH, pH 7.9, 25% glycerol, 420 mM NaCl, 0.2 mM EDTA, 1.5 mM MgCl₂, 10 mM KCl, 0.5 mM dithiothreitol, and protease inhibitor cocktail) and incubated on ice for 20 min. The nuclear fraction was obtained as the supernatant after centrifugation at 18,000 g for 2 min.

Western Analyses. Equal amounts of protein were separated and transferred to nitrocellulose membrane as described previously (Shi et al., 2007). Membranes were blocked either in Tris-buffered saline or PBS containing 5% (w/v) nonfat dry milk with 0.1% Tween 20. The following primary antibodies were used: anti-RGS7 (1:1000, polyclonal antibody; Upstate Biotechnology, Inc., Lake Placid, NY), anti-phospho-JAK2 (1:1000, polyclonal antibody; Affinity Bioreagents, Golden, CO), and anti-Jak2 (1:2000, polyclonal antibody; Upstate Biotechnology, Inc.). The anti-phospho-STAT1, STAT3, and STAT5 (1:1000, polyclonal antibodies) and anti-STAT1, STAT3, and STAT5 (1:1000, polyclonal antibody) were purchased from Cell Signaling (Danvers, MA). A monoclonal anti-actin antibody was from MP Biomedicals (1:10,000; Aurora, OH). Before incubation with a second primary antibody, blots were stripped with Restore Western blot stripping buffer (Pierce Chemical) by incubating at 37°C for 25 min. After incubation blots were removed from stripping buffer, washed three times for 10 min each with Tris-buffered saline or PBS containing 0.1% Tween 20, and blocked again. Protein bands were analyzed densitometrically using Scion Image software (Scion Corporation, Frederick, MD). The integrated optical density for the film
background was subtracted from the integrated optical density for each band. Each sample was measured in triplicate. RGS7 protein and 5-HT\textsubscript{2A} receptor protein levels were normalized to actin protein used as a loading control, and phosphoproteins were normalized to the corresponding total protein levels. Protein levels from olanzapine-treated cells were normalized to vehicle-treated cells.

**5-HT\textsubscript{2A} Receptor Antibody Production and Characterization.** A peptide corresponding to amino acids 22 to 41 of the rat 5-HT\textsubscript{2A} receptor (NH\textsubscript{2}-GG PRLYHINDPNSRDIANTSE-OH) was synthesized and used to produce antibodies by Biosynthesis, Inc. (Lewisville, TX). This sequence is 85% identical to the mouse 5-HT\textsubscript{2A} receptor sequence and 65% identical to the human 5-HT\textsubscript{2A} receptor sequence as determined using the NCBI Sequence Viewer. The same peptide sequence was used previously by Garlow et al. (1993) to produce antibodies against the 5-HT\textsubscript{2A} receptor. The antibodies produced were characterized using Western blotting and enzyme-linked immunosorbent assays. The antibody titer reported by Biosynthesis, Inc. was up to 1:25,600. A1A1v cells were used to verify the specificity of the 5-HT\textsubscript{2A} receptor antibodies. Cells were transfected with the human 5-HT\textsubscript{2A} receptor in pcDNA3.1+ (Guthrie DNA Resource Center, Sayre, PA) using 3 µg of DNA for every dish and the Lipo-fectamine Plus Reagent (Invitrogen, Frederick, MD).

**Inositol Phosphate Accumulation Assay.** Assays were performed as described previously (Berg et al., 1994; Shi et al., 2007). In brief, cells were seeded in 24-well plates at the density of 40,000/well. Cells were treated with vehicle or 300 nM olanzapine and labeled with 0.5 µCi [\textsuperscript{3}H]myoinositol/well for 24 h in serum-free Dulbecco’s modified Eagle’s medium. Cells were washed with Hank’s balanced salt solution containing 20 mM LiCl\textsubscript{2} and 20 mM HEPES, pH 7.4. Phosphoinositol hydrolysis was initiated with the addition of DOI at 37°C. Reaction was stopped after 30 min with ice-cold 10 mM formic acid. The accumulation of [\textsuperscript{3}H]-labeled inositol phosphate (IP) was determined by ion exchange chromatography.

**Statistics.** All statistical analyses were performed using the GB-STAT School Pack (Dynamic Microsystems, Silver Spring, MD). Data are expressed as means ± S.E.M. For Western blots and the IP accumulation assay, data were analyzed using a Student’s t test for equal variances.

**Results**

**IP Accumulation Assay.** Agonist-stimulated IP accumulation can be used to monitor desensitization of 5-HT\textsubscript{2A} receptor mediated signaling (Hanley and Hensler, 2002). Treatment with 300 nM olanzapine for 24 h significantly decreased (p < 0.05) DOI (10\textsuperscript{−4} M)-stimulated IP accumulation by 28 ± 1.9% compared with vehicle (20% acetic acid)-treated cells (Fig. 1). This decrease suggests a desensitization of 5-HT\textsubscript{2A}-mediated receptor signaling by olanzapine. The DOI concentration was chosen based on the previous dose-response experiments conducted in our laboratory (Shi et al., 2007).

**Olanzapine Induced Phosphorylation of JAK Kinase.** Guillet-Deniau et al. (1997) have shown that serotonin stimulation of 5-HT\textsubscript{2A} receptors causes phosphorylation of JAK2 kinase and association of the receptor with JAK2. To investigate whether olanzapine causes JAK2 phosphorylation, lysates of cytosol and membrane fractions prepared from vehicle (20% acetic acid)- and olanzapine (300 nM)-treated cells were examined by Western blot with anti-phospho-JAK2 antibody, then stripped and reprobed with anti-JAK2 antibody (Fig. 2A). Tyrosine phosphorylation of JAK2 was significantly increased (p < 0.05) to more than 232 ± 15% of the control levels in the membrane fraction of olanzapine-treated cells, whereas total JAK2 protein levels did not show any appreciable change.

**Phosphorylation and Nuclear Translocation of STAT Proteins.** Phosphorylated JAK2 facilitates activation of various STAT proteins. Tyrosine-phosphorylated STATs then undergo dimerization, translocate to the nucleus, and bind to the target DNA sequences. However, different STAT proteins mediate signaling cascades stimulated by different agonists. To determine which STAT protein(s) are activated in response to olanzapine, lysates of cytosolic and nuclear fractions from control (20% acetic acid)- and olanzapine (300 nM)-treated cells were analyzed by Western blot with anti-phospho-STAT1, STAT3, and STAT5 antibodies, then stripped and reprobed with corresponding anti-STAT antibodies. We found that phosphorylation and nuclear translocation of STAT3 was significantly increased to 171 ± 25% in the olanzapine-treated cells compared with vehicle-treated cells (Fig. 2B), whereas phosphorylation and nuclear translocation of STAT1 and STAT5 did not show any change (data not shown).

**Olanzapine Increases RGS7 Protein Levels.** To monitor the changes in RGS7 protein levels, lysates from cytosol and membrane fractions of vehicle (20% acetic acid)- and olanzapine (300 nM)-treated cells were analyzed by Western blot with anti-RGS7 antibody (Fig. 3A). We found that RGS7 protein levels were increased in the membrane fraction to 160 ± 18% of control levels and significantly increased in cytoplasmic fraction by 193 ± 32% of control levels in olanzapine-treated cells compared with vehicle-treated control cells.

**Characterization of 5-HT\textsubscript{2A} Receptor Antibody.** A1A1v cells were used to verify the specificity of the 5-HT\textsubscript{2A} receptor antibody. On Western blots prepared with A1A1v cell lysates, there was a prominent band with a molecular mass of approximately 42 kDa (Fig. 3B). This is very similar to the size of the band produced by in vitro transcription and translation of the human 5-HT\textsubscript{2A} receptor construct provided by the Guthrie cDNA Resource Center. Overexpression of the human 5-HT\textsubscript{2A} receptor construct in A1A1v cells resulted in
a more intense protein band detected on Western blots prepared with the 5-HT\textsubscript{2A} receptor antibody (Fig. 3B). Pread- sorption control experiments were also performed to verify the specificity of the antibody. Using homogenates from rat frontal cortex and lysates from A1A1v cells transfected with the human 5-HT\textsubscript{2A} receptor, the 42-kDa band 5-HT\textsubscript{2A} receptor band was no longer present in the Western blots prepared with the antibody preincubated with the peptide antigen (Fig. 3C). These experiments also demonstrate that the antibody produced, using a peptide based on the sequence for rat 5-HT\textsubscript{2A} receptor, cross-reacts with the human 5-HT\textsubscript{2A} receptor expressed in rat cells. Furthermore, the 5-HT\textsubscript{2A}
receptor antibody cross-reacts with the rat 5-HT2A receptor expressed in a rat cortical cell line.

**5-HT2A Receptor Protein Levels.** Cytosol and membrane fractions from vehicle (20% acetic acid)- and olanzapine (300 nM)-treated cells were analyzed by Western blot with the anti-5-HT2A antibody we generated. We found a significant increase ($p < 0.05$) in 5-HT2A receptor protein levels in both cytoplasmic (234 ± 32% of control level) and membrane fractions (206 ± 14% of control levels) of olanzapine-treated cells compared with vehicle-treated cells (Fig. 3D). We also assessed the purity of our membrane fraction after stripping and reprobing the same blot with an anti-NA^-K^-ATPase antibody. A band corresponding to NA^-K^-ATPase was mainly present in membrane fraction (data not shown).

**JAK Inhibition.** A JAK kinase inhibitor, AG490, was used to investigate whether inhibition of the JAK-STAT signaling cascade could reverse the increase of 5-HT2A receptor or RGS7 protein levels observed in response to olanzapine treatment. Cells were treated for 1 h with 0, 15, and 30 µM AG 490 before adding either vehicle or olanzapine. Twenty-four hours later, cells were lysed, and protein levels of phospho-JAK2, RGS7, and phospho-STAT3 were analyzed by Western blot. Olanzapine-induced phosphorylation of JAK2 was decreased with AG490 as shown in Fig. 4A. There was no change in the total JAK2 protein levels. With AG490 treatment, there was a similar decrease in olanzapine-induced STAT3 phosphorylation in the nuclear fraction as shown in Fig. 4B and no change in STAT levels, again confirming previous findings that activation of JAK2 causes phosphorylation and nuclear localization of STAT3.

If the JAK-STAT signaling cascade is mediating the olanzapine-induced increase in protein levels, then inhibiting this signaling cascade should prevent the increase in 5-HT2A receptor and RGS7 protein levels. To test this hypothesis, membrane fractions from vehicle- and olanzapine-treated cells pretreated with AG490 were analyzed by Western blot for 5-HT2A receptor and RGS7 protein levels. As in a previous experiment (Fig. 3A), olanzapine treatment for 24 h increased the levels of RGS7 protein (Fig. 5, A and B). Treatment with AG490 reduced the olanzapine-induced increase in RGS7 protein in the membrane fractions to the levels in the vehicle-treated cells (Fig. 5A). Treatment with AG490 alone had no effect on the levels of RGS7 protein. Although we observed a similar increase in levels of 5-HT2A receptor protein in olanzapine-treated cells compared with vehicle-treated cells as shown before in Fig. 3D, AG490 pretreatment did not alter protein levels of 5-HT2A receptor in cells treated with olanzapine (Fig. 5, C and D), suggesting that the increase in levels is not mediated by the JAK-STAT signaling cascade.

**Discussion**

This study demonstrates the involvement of an intracellular signaling cascade, the JAK-STAT pathway, in increasing the levels of RGS7 protein in response to treatment with olanzapine. The increased levels of RGS7 protein in turn could contribute to the desensitization of 5-HT2A receptor signaling induced by olanzapine in A1A1v cells by directly interacting with Gα11, and accelerating GTP hydrolysis. In contrast, we found an increase in 5-HT2A receptor protein levels with olanzapine treatment that was not associated with increased activation of the JAK-STAT signaling cascade. The increase in 5-HT2A receptor protein levels would not probably contribute to the desensitization response but could conceivably counter or moderate the desensitization response.

Olanzapine and other antipsychotics have been recently reported to stimulate other signaling cascades, including increased phosphorylation of ERK1/2 in rat frontal cortex (Fumagalli et al., 2006) and increased phosphorylation of Akt/PKB and p38 in PC12 cells (Lu et al., 2004). The increased activation of signaling cascades induced by antipsychotics suggests that changes in gene expression regulated by these cascades could contribute to the positive therapeutic benefits seen in schizophrenic patients.

Several microarray studies have explored the effects of olanzapine and other antipsychotics on gene expression, reporting changes in expression of a host of genes including RGS proteins, and gene families linked with synaptic plas-
ticity and presynaptic neurotransmission (Fehér et al., 2005; Fatemi et al., 2006). Fatemi et al. (2006) reported an increase in expression of RGS19 mRNA and a decrease in expression of RGS2 mRNA in the frontal cortex of rats treated for 3 weeks with olanzapine. GAP activity of RGS19 is associated with $G_{i/o}$, a member of the $G$ protein family, whereas 5-HT$_2A$ receptors are coupled with $G_{q/11}$ proteins. Therefore, increased RGS19 is unlikely to affect olanzapine-induced desensitization of 5-HT$_2A$ receptor signaling. Although RGS2 protein associates with $G_{q/11}$ protein and could decrease the 5-HT$_2A$ receptor signaling, a decrease in expression of RGS2 protein could not cause the desensitization of 5-HT$_2A$ receptor signaling induced by olanzapine because a decrease in RGS protein expression would probably result in increased receptor signaling. Although RGS2 is expressed in the frontal cortex, it is not known if it colocalizes with 5-HT$_2A$ receptors in the frontal cortex.

None of these microarray studies identified alterations in RGS7 protein or 5-HT$_2A$ receptor mRNA levels. It is important to remember that changes in mRNA levels may or may not result in changes in protein levels, and changes in protein levels may not be due to changes in mRNA levels. It is also important to emphasize that these studies were conducted in different experimental settings with different time course and dose regimens than our study. Each study highlights important findings, further extending our understanding of the mechanism of action of antipsychotics. It is becoming evident that there could be numerous pathways and alterations in gene expression that lead to the development of psychosis (Ko et al., 2006) and its treatment with atypical antipsychotics.

Genetic risk factors for schizophrenia are beginning to be identified and include RGS4 protein with RGS4 protein expression being reduced in the frontal cortex in schizophrenics (Gu et al., 2007). Like RGS7 protein, RGS4 regulates 5-HT$_2A$ receptor signaling. Individualized treatment approaches based on a patient’s genetic profile are being pioneered in cancer treatment and could be developed for schizophrenia and other mood disorders. For example, olanzapine-induced increases in expression of RGS7 protein could mitigate reductions in RGS4 protein expression in patients with alterations in RGS4. Based on the concept of individualized treatment approaches, it would be important to determine which other atypical antipsychotics increase RGS7 and possibly RGS4 protein expression.

Direct association of JAK2 and STAT3 with 5-HT$_2A$ receptors and activation of the JAK-STAT signaling cascade by 5-HT$_2A$ receptor agonists has been reported previously (Guillet-Deniau et al., 1997). In the present study, we found JAK2 activation and increased phosphorylation and nuclear localization of phospho-STAT3 in olanzapine-treated cells. It is interesting to note that phosphorylation and nuclear translocation of other isoforms of STAT proteins, STAT1 and STAT5, did not show any change with olanzapine treatment. Activation of STAT proteins, which are transcription factors, could bolster the previous notion that antipsychotic agents affect expression of various genes.

Our experiments with a JAK kinase inhibitor, AG490, suggest that the increased levels of RGS7 protein are mediated by the JAK-STAT signaling cascade in response to olanzapine treatment. Interestingly, AG490 selectively targeted the membrane-localized RGS7 protein and had no effect on...
the levels in cytosol fraction. The overall effect of AG490 is a decrease in the total levels of RGS7 protein (i.e., membrane plus cytosol), suggesting that olanzapine-induced JAK-STAT signaling increases the levels of RGS7 protein rather than causing a redistribution of the protein. We speculate that transcriptional activity of phospho-STAT3 could increase RGS7 mRNA expression and be responsible for increased levels of RGS7 protein in the membrane fraction. The mechanism(s) involved in the selective decrease in the membrane fraction are unknown but could involve a chaperone protein. However, further studies are needed to test these hypotheses. In contrast, changes in 5-HT2A receptor protein levels seem to be regulated by a different mechanism because pretreatment with AG490 fails to reverse the olanzapine-induced increase in receptor protein levels.

Most atypical antipsychotics cause desensitization of 5-HT2A receptor signaling. The decrease in DOI-stimulated IP accumulation with olanzapine treatment in our study is consistent with these findings and supports the use of A1A1 cells as a model to study the actions of olanzapine on 5-HT2A receptor signaling. Previous experiments in A1A1 cells using the selective 5-HT2A receptor antagonist, MDL 100,907, demonstrated that IP accumulation stimulated by the 5-HT2A/2C-selective agonist DOI is not likely due to stimulation of 5-HT2C receptors (Shi et al., 2007). Numerous investigations have explored the mechanisms by which antipsychotics cause desensitization of 5-HT2A receptor signaling. This desensitization by antipsychotics could be mediated by receptor internalization, uncoupling of G proteins from receptor, and receptor down-regulation (Roth et al., 1995; Williams et al., 1999) in addition to increases in RGS7 protein.

Other mechanisms have been associated with desensitization of 5-HT2A receptor signaling as a result of treatment with atypical antipsychotics. Several studies specifically investigated changes in the transcript level of 5-HT2A receptors with both short-term and chronic antipsychotic treatments (Burnet et al., 1996; Doat-Meyerhoefer et al., 2005). Buckley et al. (1997) reported a significant decrease in receptor mRNA in hippocampus, brain stem, and midbrain, whereas no significant change was observed in other brain regions after 32 days of treatment with the atypical antipsychotic clozapine. However, in the same study, 4 days of clozapine treatment did not produce any significant change, but a trend for a decreased mRNA expression was observed in major brain areas. As noted previously, changes in mRNA levels do not necessarily result in corresponding changes in protein levels, so changes in 5-HT2A receptor mRNA levels are not necessarily inconsistent with the current findings. We found an increase in 5-HT2A receptor protein levels in both the cytosol and membrane fractions. Previous studies reported a sizable amount of 5-HT2A receptors in the cytosol in addition to that found in the membrane (Cornea-Hebert et al., 1999). Several reports suggest a decrease in $B_{max}$ with no change in $K_a$ after treatment with antipsychotics (Matsubara and Meltzer, 1989; Doat-Meyerhoefer et al., 2005). A decrease in the density of 5-HT2A receptors without any change in affinity is consistent with the receptor internalization previously reported (Roth et al., 1995; Williams et al., 1998). An increase in 5-HT2A receptor protein levels in the cytosol, as we found with Western blot analysis, could reflect internalized receptors. Furthermore, an increase in total 5-HT2A receptor protein levels is not necessarily inconsistent with a decrease in receptor density. 5-HT2A receptor proteins under go extensive post-translational modifications that could affect ligand binding. It is difficult to speculate at this point about the post-translational modifications that are required for or inhibit ligand binding. It is likely that multiple mechanisms contribute to desensitization of 5-HT2A receptor signaling and that the specific mechanisms involved are probably tissue-specific.

In summary, our in vitro data highlight a new role of JAK-STAT signaling in treatment with olanzapine. Increased activation of this pathway by olanzapine increases expression of RGS7 protein. Increased RGS7 protein could directly contribute to the desensitization of 5-HT2A receptor signaling by accelerating hydrolysis of GTP-bound $G_{q/11}$ protein. Unfortunately, the JAK inhibitor AG490 interferes with the IP accumulation assay and therefore precludes our ability to determine whether JAK-STAT signaling contributes to the desensitization of 5-HT2A receptors in our model system. We are further investigating whether increased RGS7 protein is the result of increased transcriptional activity and whether blocking membrane localization of RGS7 protein inhibits desensitization of 5-HT2A receptor signaling cascade. Further studies are also needed in animal models to confirm these findings in vivo. Overall, the results from this study provide a further understanding of possible involvement of intracellular pathways in mediating the effects of atypical antipsychotics.

References


