Altered transcription factor trafficking in oxidatively-stressed neuronal cells

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ABSTRACT

Age-related neurodegenerative diseases are associated with alterations in gene expression in affected neurons. One of the mechanisms that could account for this is altered subcellular localization of transcription factors, which has been observed in human post-mortem brains of each of the major neurodegenerative diseases, including Parkinson's disease (PD). The specific mechanisms are yet to be elucidated; however a potential mechanism involves alterations in nuclear transport. In this study, we examined the nucleocytoplasmic trafficking of select transcription factors in response to a PD-relevant oxidative injury, 6-hydroxydopamine (6OHDA). Utilizing a well-established model of ligand-regulated nucleocytoplasmic shuttling, the glucocorticoid receptor, we found that 6OHDA selectively impaired nuclear import through an oxidative mechanism without affecting nuclear export or nuclear retention. Interestingly, impaired nuclear import was selective as Nrf2 (nuclear factor E2-related factor 2) nuclear localization remained intact in 6OHDA-treated cells. Thus, oxidative stress specifically impacts the subcellular localization of some but not all transcription factors, which is consistent with observations in post-mortem PD brains. Our data further implicate a role for altered microtubule dependent trafficking in the differential effects of 6OHDA on transcription factor import. Oxidative disruption of microtubule-dependent nuclear transport may contribute to selective declines in transcriptional responses of aging or diseased dopaminergic cells.

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1. Introduction

Age-related neurodegenerative diseases, such as Parkinson's disease (PD), Alzheimer's disease (AD), amyotrophic lateral sclerosis (ALS), and Huntington's disease (HD), are associated with alterations in expression of specific genes in affected neurons [1–3]. Transcription factors play a key role in the regulation of gene expression and the activity of many transcription factors is limited by their localization within the cytoplasm. A number of mechanisms are utilized to direct sequestered transcription

factors into the nucleus in response to specific signals [4–11]. Since nucleocytoplasmic protein trafficking can be bidirectional [12], transcription factors exhibiting predominant nuclear localization may transit through the cytoplasm and also function in nongenomic signaling networks [13–16]. Interestingly, alterations in the subcellular localization of a number of transcription factors are documented in human post-mortem brains of each of the major neurodegenerative diseases [5]. Examples include cytoplasmic aggregation of phosphorylated (p)CREB (cAMP response element binding protein) and lack of nuclear pCREB in PD [17], reduced nuclear localization of Nrf2 (nuclear factor E2-related factor 2) in AD but not in PD [18], increased cytoplasmic ATF2 (activating transcription factor 2) levels in AD [19], and increased cytoplasmic:nuclear ratios of TDP-43 (TAR DNA-binding protein 43) in frontotemporal lobar dementias and ALS [20].

Extensive evidence of oxidative damage to proteins, lipids, and DNA is found upon analysis of post-mortem brains of various neurodegenerative diseases, including PD [21–31]. Furthermore, genetic and environmental models of various neurodegenerative diseases support the notion that oxidative stress plays a primary role in their pathogenesis [24–26]. In the case of PD, mutations or deficiency of proteins such as Parkin, PINK1 (PTEN-induced putative kinase 1), α-synuclein, DJ-1, and LRRK2 (leucine-rich repeat kinase 2), have been shown to increase susceptibility to oxidative stress-mediated cell death [32–34]. Parkinsonian mimetics such as, 6-hydroxydopamine (6OHDA), 1-methyl-4-phenyl-1,2,3,6-tetrahydropyridine (MPTP), rotenone, and paraquat, generate reactive oxygen species (ROS) and cause neuronal death

[35]. These studies suggest a key role for oxidative stress in the pathogenesis of these diseases.

6OHDA has been widely used to study neuronal injury responses in PD. It is an analog of catecholamine neurotransmitters that is taken up into cells by dopamine and norepinephrine reuptake transporters. It is commonly used to model PD-relevant oxidative stress as it demonstrates an early autoxidation phase of ROS production and also has a delayed phase of mitochondrial ROS production [36]. Although used as an exogenous neurotoxin, there is evidence of spontaneous ring hydroxylation of dopamine in vivo, with elevated body fluid levels of 6OHDA detected in patients treated with L-Dopa [37,38].

6OHDA treatment leads to increased cytoplasmic accumulation of (p) ERK (extracellular signal-regulated kinase) [39] and decreased nuclear levels of GFP (green fluorescent protein)-ERK2 [40], as well as decreasing the nuclear to cytoplasmic ratios of pCREB in SH-SY5Y cells and in primary midbrain neurons [17]. These findings are consistent with alterations observed in dopaminergic neurons in post-mortem PD brains [5]. CREB-regulated gene transcription, which is important for axon growth, mitochondrial biogenesis, and neuronal survival, was also repressed in 6OHDA treated cells. Given the dynamic shuttling of transcription factors between the nuclear and cytoplasmic compartments, there are a number of mechanisms that could result in altered distribution of transcription factors in degenerating neurons, including compartmentalization, altered expression levels, sequestration in protein aggregates, and nuclear transport deficits [5]. Interestingly, 6OHDA impairment of CREB-mediated transcription (e.g. Bcl2 [B-cell lymphoma 2] and BDNF [brain-derived neurotrophic factor]) was reversed with cAMP treatment, which activates CREB through mechanisms not requiring active nuclear import [17,41]— suggesting a possible impairment in the nucleocytoplasmic trafficking of CREB. The observed steady state alterations in the distribution of transcription factors could result from altered nuclear import/export with significant consequences on neuronal survival. The main goal of this study was to directly examine, in a neuronal cell line, the impact of 6OHDA-induced oxidative stress on distinct steps in the regulated nucleocytoplasmic trafficking of select transcription factors.

2. Materials and methods

2.1. Cell culture

SH-SY5Y neuroblastoma cells (ATCC, Manassas, VA) were grown on 10 cm cell culture dishes containing Dulbecco's Modified Eagle's Medium (DMEM) with 10% heat-inactivated fetal calf serum, 2 mM L-glutamine, and 10 mM HEPES. Cells were maintained at 37 °C in a humidified 5% CO_2 incubator.

2.2. Plasmids and transfections

SH-SY5Y cells were transfected with Lipofectamine™ 2000 reagent (Invitrogen, Carlsbad, CA) utilizing Opti-MEM® I reduced serum media (Invitrogen, Carlsbad, CA) according to the manufacturer's protocol at a final Lipofectamine concentration of 0.1%. Cells were grown for at least 48 h before transfection and then allowed to express the protein of interest for another 24 h–48 h before treatment. GR-GFP and Nrf2-GFP plasmids were kindly provided by Dr. Ian Macara (University of Virginia) and Dr. Manabu Furukawa (University of Nebraska), respectively.

2.3. Drug treatments

6OHDA (Sigma, St. Louis, MO) was prepared in cold sterile water immediately before use. Dexamethasone (Sigma), a synthetic glucocorticoid hormone, was prepared at a stock concentration of 10^{-3} M in 95% EtOH and used at a working concentration of 10^{-6} M unless otherwise stated. Cortisol (Sigma) was prepared at a stock concentration of 10^{-3} M to 10^{-4} M in 100% EtOH and used at a working concentration of 10^{-6} M to 10^{-7} M unless otherwise stated. For antioxidant treatment, a metalloporphyrin antioxidant, MnTBAP (A.G. Scientific, San Diego, CA), was utilized at a final concentration of 150 μM. Colchicine (Sigma) and Paclitaxel (Sigma) were prepared in dH_2O and DMSO, respectively, and used at the concentrations indicated in the text.

2.4. Monomeric tubulin extraction

Cells treated with either vehicle or the drug of interest were washed in 37 °C DPBS (Dulbecco's Phosphate-Buffered Saline) and then incubated with tubulin extraction buffer (10 mM PIPES, pH 6.8, 50 mM KCl, 2 mM EGTA, 1 mM $MgCl_2$, 2 M glycerol, 0.5% Triton X-100, 1 mM Na_3VO_4, and protease inhibitor cocktail [Sigma]) for 15 min at room temperature. The resulting lysate was then centrifuged at 37 °C at $16,000 \times g$ for 2 min to pellet any polymerized microtubules. The resulting supernatant, containing soluble (monomeric) tubulin, was then subjected to Western blotting. The protocol was modified from [42].

2.5. Immunoblotting

Cell extracts were collected and immunoblots were performed as previously described [43] with the following exception: BioRad (Hercules, CA) gel running system with a 10% gel was used to separate proteins. Primary antibodies used: anti-α-tubulin (1:5000; Sigma) and anti-GAPDH (1:10,000; Abcam). ImageJ (NIH) was utilized for densitometry (from $N \geq 3$ experiments).

2.6. Cell toxicity assay

Cells were plated in 96-well plates and treated with 150 μM 6OHDA for varying amounts of time. LDH (lactate dehydrogenase) release assay was carried out according to the protocol provided by the manufacturer (Promega, Madison, WI). Spectramax M2 plate reader was utilized to read the fluorescent signal (Molecular Devices, Sunnyvale, CA). Percent cytotoxicity was calculated by the following formula: $100 \times [(\text{Experimental} - \text{Medium Background}) / (\text{Maximum LDH release} - \text{Medium Background})]$. Maximum LDH release was determined by treatment of cells with lysis buffer as per manufacturer's protocol.

2.7. Quantitative image analysis and statistics

An inverted epifluorescent microscope (Olympus IX71) was used for imaging and ImageJ was utilized for quantification. Nuclear:cytoplasmic ratios were calculated from at least 50 randomly imaged cells per experiment (and pooled from at least three independent experiments) by measuring the fluorescent intensity within a randomly distributed region of interest of fixed size within the nuclear and the cytoplasmic compartments. Raw fluorescent intensity is shown in the figures. Nrf2 distribution was also scored into two categories: (1) cytoplasmic and (2) cytoplasmic and nuclear. Example images of these categories are shown in the Results section. Student's t test was used to compare means between two groups. One-way analysis of variance (ANOVA) followed by posthoc Tukey HSD test was used for multiple comparisons. A p-value<0.05 was considered statistically significant.

3. Results

3.1. Subcellular localization of GR in response to hormone

The glucocorticoid receptor (GR) was used to examine the effects of 6OHDA-induced oxidative stress on bidirectional nucleocytoplasmic trafficking. Transport of GR from the cytoplasm to the nucleus is hormone-dependent and bidirectional as hormone withdrawal triggers its export from the nucleus [12,44–46]. Therefore, both steps in the nucleocytoplasmic trafficking of this transcription factor are regulated
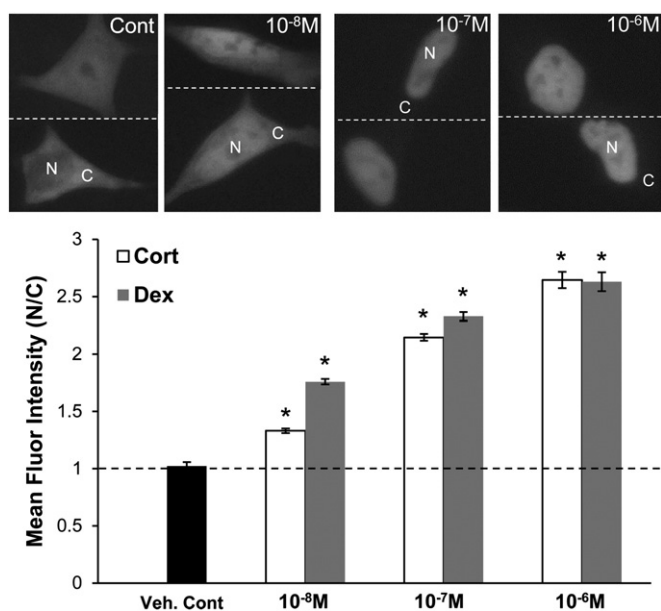
Fig. 1. Dexamethasone and cortisol induce GR nuclear import. SH-SY5Y cells expressing GR-GFP were treated for 1 h with Dex or Cort at the indicated concentrations. In the absence of either ligand, GR-GFP is cytoplasmic in the majority of the cells (~60%) with the remainder showing some nuclear localization as well. The raw mean fluorescent intensity within equal size nuclear and cytoplasmic regions (N/C ratio) for untreated cells is approximately 1. After hormone treatment, GR-GFP is primarily localized within the nucleus. Nuclear:cytoplasmic GR-GFP ratios were determined as noted in the Materials and methods section. Representative images from Dex treatment are shown. Mean +/− SEM, *$p < 0.05$ vs. Veh. Cont. Compiled from 3 independent experiments.

and can be experimentally isolated. SH-SY5Y human neuroblastoma cells were transfected with an expression vector for GR-GFP and treated for 1 h with either Dexamethasone (Dex), a synthetic glucocorticoid, or Cortisol (Cort), the primary circulating glucocorticoid in humans. In the absence of Dex or Cort, GR-GFP is cytoplasmic in the majority of the cells (~60% on average) with the remainder of cells also showing some

nuclear localization. The mean fluorescent intensity within equal size nuclear and cytoplasmic regions (nuclear:cytoplasmic ratio) for untreated cells is on average around 1 (Fig. 1). A dose-dependent increase in nuclear transport of GR-GFP was observed with a significant nuclear localization attained at $\geq 10^{-7}$ M Dex or Cort (Fig. 1).

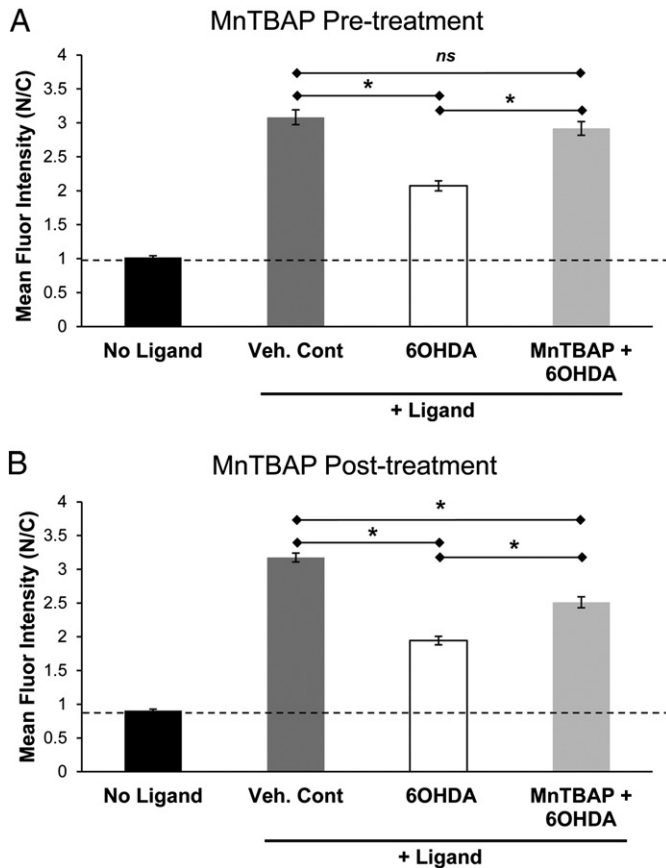### 3.2. 6OHDA treatment impairs nuclear import of GR

With 150 μM 6OHDA (dose where impaired CREB/pCREB localization is observed [17]), treatment for less than 6 h did not cause cell death (Fig. 2A). To determine if 6OHDA treatment causes an impairment in nuclear import, GR-GFP expressing SH-SY5Y cells were first treated with 150 μM 6OHDA for 1 h and 4 h and then treated with 1 μM Dex for 1 h to induce the nuclear localization of GR-GFP. As seen in Fig. 2B and C, pre-treatment with 6OHDA impaired the hormone-dependent nuclear import of GR-GFP. Significant reduction in the extent of nuclear import was also seen at doses varying from 90 μM to 300 μM and treatment times as early as 30 min (data not shown).

3.3. Antioxidant treatment reverses 6OHDA-induced impairment in nuclear import

6OHDA has the propensity to produce free radicals [36,47]. To determine whether the nuclear import impairment observed in 6OHDA-treated cells involves oxidative stress, SH-SY5Y cells expressing GR-GFP were treated with MnTBAP, a cell permeable SOD mimetic, 30 min before or 30 min after the start of 6OHDA treatment (150 μM; 4 h) (Fig. 3). When cells were treated with MnTBAP before being exposed to 6OHDA, a near complete rescue in GR-GFP nuclear import was observed (Fig. 3A). When cells were first treated with 6OHDA for 30 min before the addition of MnTBAP, a partial protection was observed (Fig. 3B). This is consistent with the observation that 6OHDA treatment as short as 30 min is enough to cause some impairment in GR nuclear import (data not shown). The rescue observed with MnTBAP treatment suggests that an oxidative mechanism is responsible at least in part for the 6OHDA-mediated reduction in GR nuclear import.
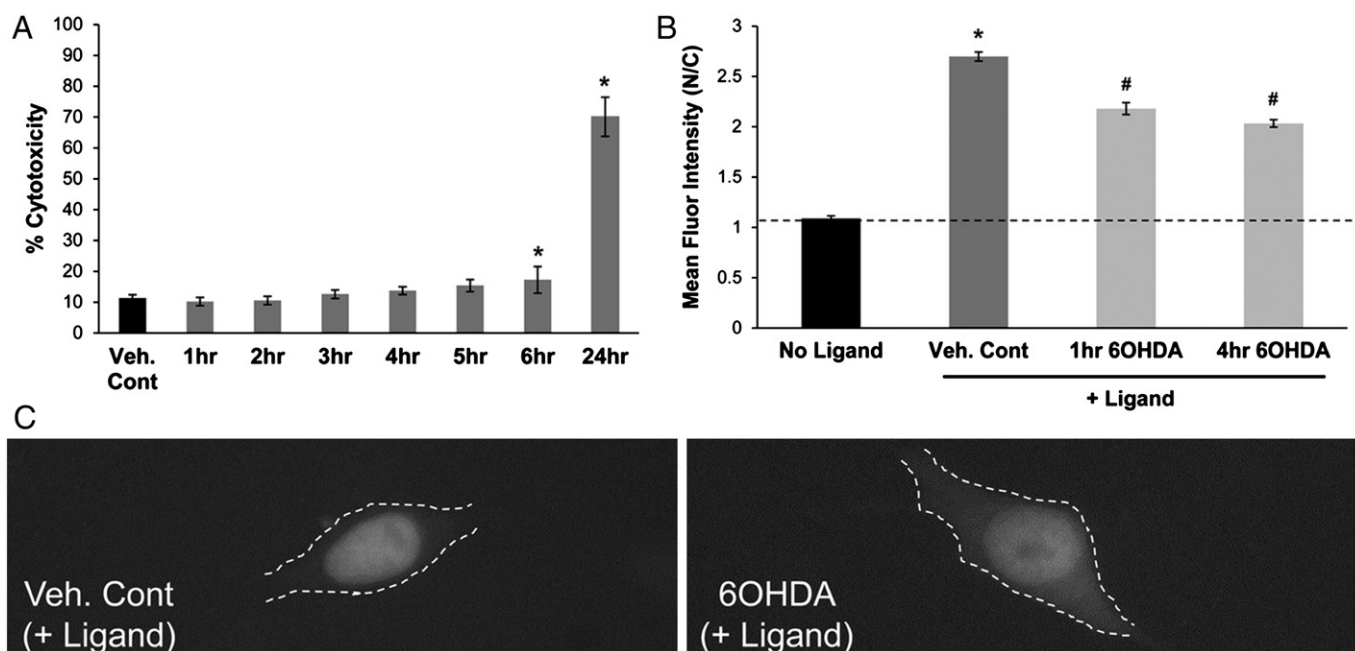
Fig. 2. 6OHDA reduces GR nuclear import. (A) LDH release assay showed no evidence of toxicity until 6 h of 6OHDA treatment (150 μM). Mean +/− SEM, *$p < 0.05$ vs. Cont. (B and C) GR-GFP expressing SH-SY5Y cells were exposed to 150 μM 6OHDA for 1 h and 4 h and then treated with 1 μM Dex for an additional 1 h. Significant reduction in GR import was observed in 6OHDA-treated cells. Mean +/− SEM, *$p < 0.05$ vs. No Ligand; #$p < 0.05$ vs. Veh. Cont. Compiled from 3 independent experiments.

Fig. 3. MnTBAP rescues 6OHDA-induced impairment in GR nuclear import. SH-SY5Y cells expressing GR-GFP were treated with MnTBAP, a cell permeable SOD mimetic, 30 min before (A) and 30 min after (B) the start of 6OHDA treatment (150 μM for 4 h). A complete and a partial protection was observed with pre- and post-6OHDA treatments with MnTBAP, respectively. Mean +/− SEM, *$p < 0.05$. Compiled from 3 independent experiments.

3.4. Nuclear export of GR is not altered in 6OHDA-treated cells

In addition to impaired nuclear import, enhanced nuclear export could also explain the altered transcription factor distribution observed in neurodegenerative diseases. To study if nuclear export is affected by 6OHDA-induced oxidative stress, a hormone withdrawal study was performed in SH-SY5Y cells expressing GR-GFP. Cort, the naturally occurring ligand for GR, dissociates rapidly from GR and is therefore better suited for studying the effects of hormone withdrawal [46,48]. SH-SY5Y cells expressing GR-GFP were first treated with 10^{-7} M Cort for 1 h to induce the nuclear localization of GR-GFP. The cells were then withdrawn from the ligand for up to 4 h in the presence of either vehicle or 150 μM 6OHDA. No significant change in nuclear export of GR-GFP was observed in cells exposed to 6OHDA compared to vehicle control (Fig. 4).
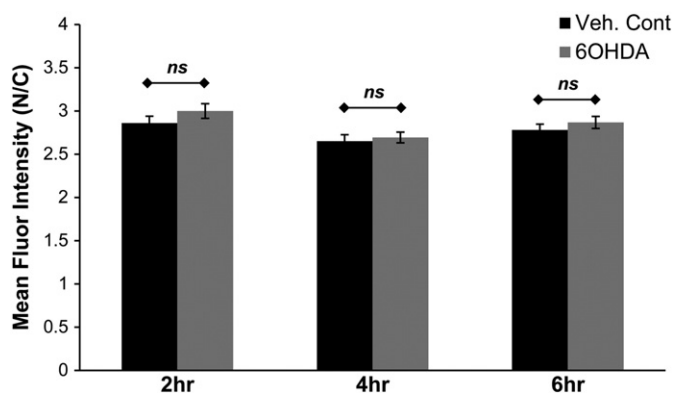
3.5. 6OHDA does not alter GR nuclear retention

6OHDA-induced oxidative stress could damage the nuclear envelope or chromatin binding sites and therefore alter nuclear retention of GR. To investigate this possibility, SH-SY5Y cells expressing GR-GFP were first treated with Dex for 1 h to fully induce the nuclear translocation of GR-GFP and then exposed to 150 μM of 6OHDA from 2 to 6 h in the presence of Dex (Fig. 5). If 6OHDA-induced oxidative stress damaged the nuclear envelope or reduced GR binding to high affinity chromatin binding sites, an increase in ligand-bound cytoplasmic GR-GFP could occur. However, no significant change in subcellular localization was observed—suggesting that 6OHDA does not reduce GR-GFP nuclear retention or trigger its leakage from a "damaged" nucleus. In combination with the results of GR nuclear import and export assays, our data suggest that 6OHDA-induced oxidative stress impairs the hormone-dependent nuclear import of GR.

3.6. Nrf2 translocates into the nucleus in response to 6OHDA

One of the transcription factors that did not show reduced nuclear localization in post-mortem human PD brains is Nrf2, a key protein in the defense against cellular stresses. Under basal metabolic conditions, Nrf2 is sequestered in the cytoplasm and targeted for proteasomal degradation via its association with Keap1 (kelch-like ECH-associated protein 1). Upon exposure to ROS, electrophiles, or serum factors, Nrf2 dissociates from the Keap1 inhibitory complex and translocates into the nucleus, where it upregulates stress response genes and protects against neurotoxic insults [49–55]. For this reason, studies of Nrf2 nuclear translocation typically involve pre-incubation under serum-deprived conditions to shift the basal distribution of Nrf2 toward the cytoplasm [54,56]. SH-SY5Y cells expressing Nrf2-GFP exhibited reduced nuclear:cytoplasmic ratios when grown in low-serum media (1% FBS) for 24 h (Fig. 6A). Serum deprivation also caused an increase in the percentage of cells with cytoplasmic
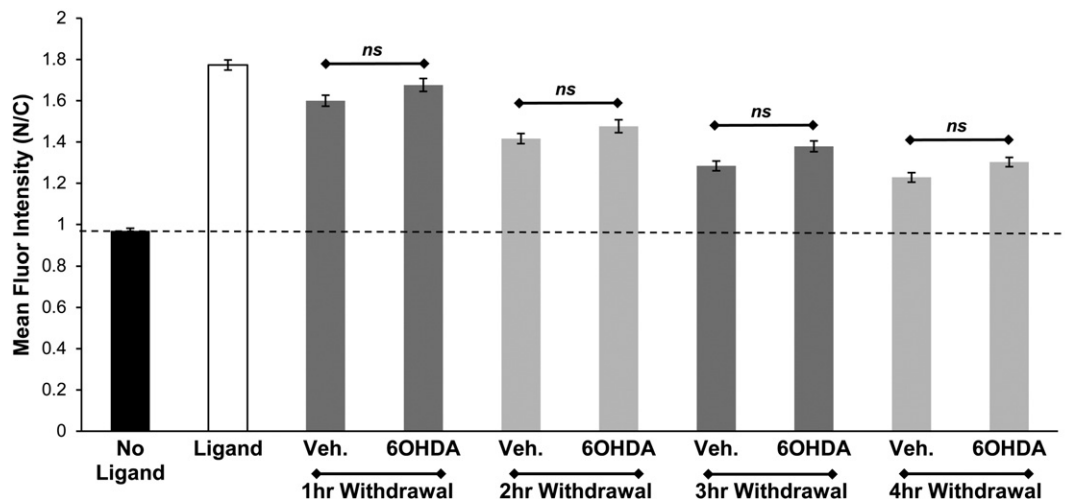
Fig. 4. 6OHDA does not alter GR nuclear export. SH-SY5Y cells expressing GR-GFP were treated with 10^{-7} M Cort for 1 h and then withdrawn from the ligand in the presence of vehicle or 6OHDA for up to 4 h. No significant change in nuclear export of GR-GFP was observed in 6OHDA treated cells. Mean +/− SEM, $p > 0.05$. Compiled from 3 independent experiments.

Fig. 5. 6OHDA does not alter GR nuclear retention. SH-SY5Y cells expressing GR-GFP were first treated with Dex for 1 h and then exposed to 150 μM of 6OHDA from 2 to 6 h in the presence of Dex. No significant change in nuclear retention of GR-GFP was observed. Mean $+/-$ SEM, $p > 0.05$. Compiled from 3 independent experiments.

localized Nrf2-GFP and a decrease in the percentage of cells showing both cytoplasmic and nuclear localization (Fig. 6B).
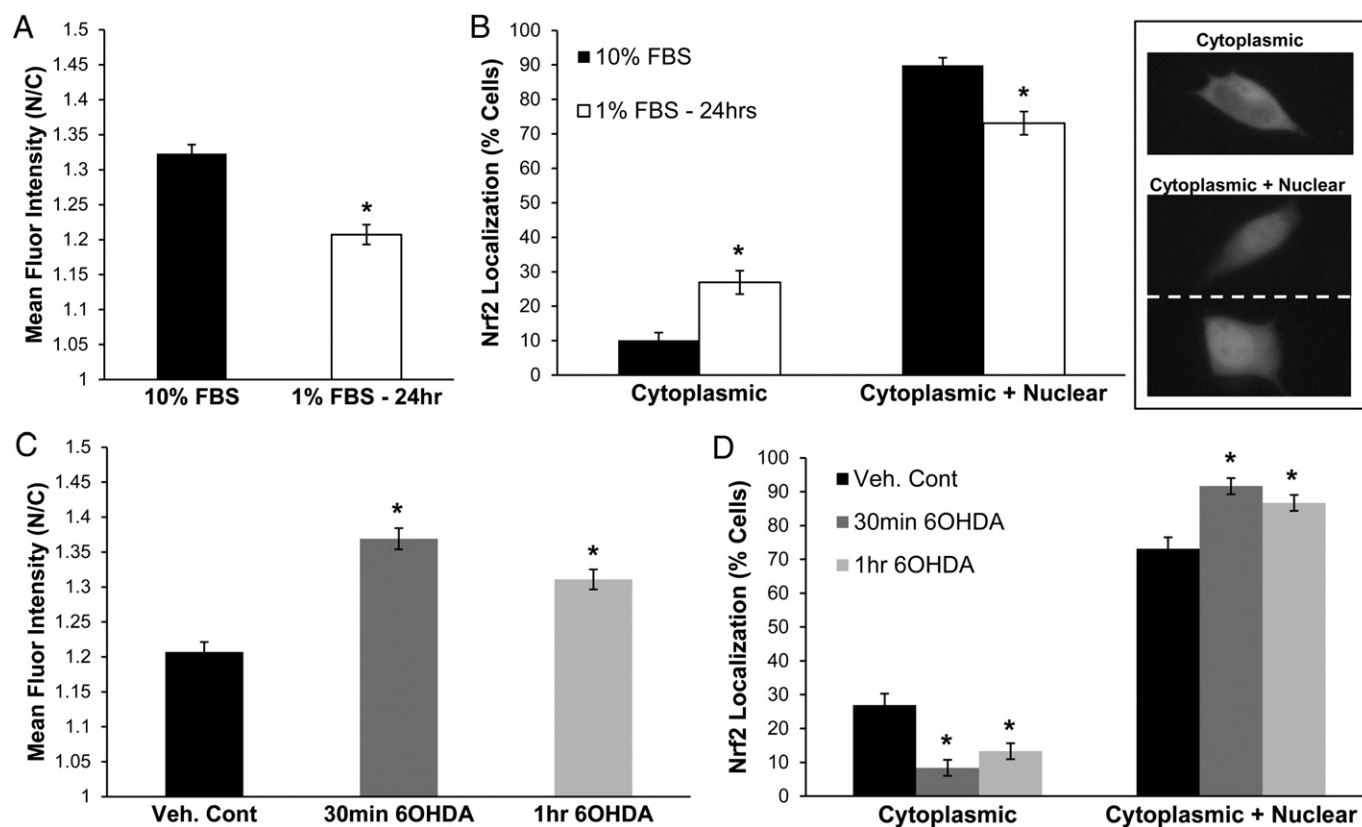
Nrf2 nuclear localization in response to 6OHDA-induced oxidative stress was then studied. Serum deprived SH-SY5Y cells expressing Nrf2-GFP were treated with increasing doses of 6OHDA (from 30 μM to 150 μM), which caused a progressive increase in the nuclear localization of Nrf2. The results from cells treated with 150 μM 6OHDA are shown in Fig. 6C. Quantitation of Nrf2-GFP localization revealed a decrease in the percentage of cells showing predominant cytoplasmic localization and an increase in the percentage of cells showing both cytoplasmic and nuclear Nrf2 localization(Fig. 6D). These results are

similar to the studies examining Nrf2 nuclear translocation in response to H_2O_2 treatment [54]. Thus, inhibitory effects of 6OHDA on nuclear transport are selective and influence the active nuclear import of GR but not of Nrf2.

3.7. 6OHDA increases levels of unpolymerized tubulin

The nucleocytoplasmic trafficking of some transcription factors, including CREB [57] and GR [58,59], is dependent on microtubules (MTs) in some cells. Hence, alterations in MT function due to 6OHDA-induced oxidative stress could help explain the selective trafficking impairments. This is of particular relevance given the alterations in MT function observed in several age-related neurodegenerative diseases [60,61]. In the case of PD, both toxin (e.g. MPP+ and rotenone) and genetic (e.g. α-synuclein and LRRK2) models implicate impaired MT function as a common pathway of neuronal degeneration [62–65].

We investigated the effects of 6OHDA-induced oxidative stress on polymerization of tubulin. We first determined the effects of MT-altering agents, colchicine and paclitaxel, on the levels of unpolymerized tubulin (Fig. 7A). SH-SY5Y cells were treated with either vehicle, 1 μM colchicine (2 h), or 50 nM paclitaxel (1 h) and then incubated with tubulin extraction buffer to extract monomeric tubulin. Colchicine increased and paclitaxel reduced the levels of monomeric tubulin, respectively, while not significantly affecting the levels of total tubulin (Fig. 7A and B). These concentrations of MT-altering agents did not cause cell death (data not shown). We then examined the effects of 6OHDA on the levels of unpolymerized tubulin. SH-SY5Y cells were treated with 150 μM 6OHDA from 1 h to 4 h after which monomeric tubulin was extracted. Cells treated with 6OHDA showed increased levels of monomeric tubulin (Fig. 7C). Levels of total tubulin were not altered in 6OHDA-treated cells (Fig. 7D).

Fig. 6. 6OHDA induces Nrf2 nuclear localization. (A and B) Standard culture conditions elicit Nrf2 nuclear translocation. To desaturate the assay, cells were incubated in low serum media (1% FBS) for 24 h and nuclear localization was quantified by two methods as described in the Materials and methods section. Example images of cells scored as cytoplasmic Nrf2-GFP or cytoplasmic + nuclear Nrf2-GFP are shown. Mean $+/-$ SEM, $*p < 0.05$ vs. 10% FBS. (C and D) Treatment of serum deprived SH-SY5Y cells with 150 μM 6OHDA caused an increased nuclear and a decreased cytoplasmic localization of Nrf2-GFP. Mean $+/-$ SEM, $*p < 0.05$ vs. Veh. Cont. Compiled from 3 independent experiments.
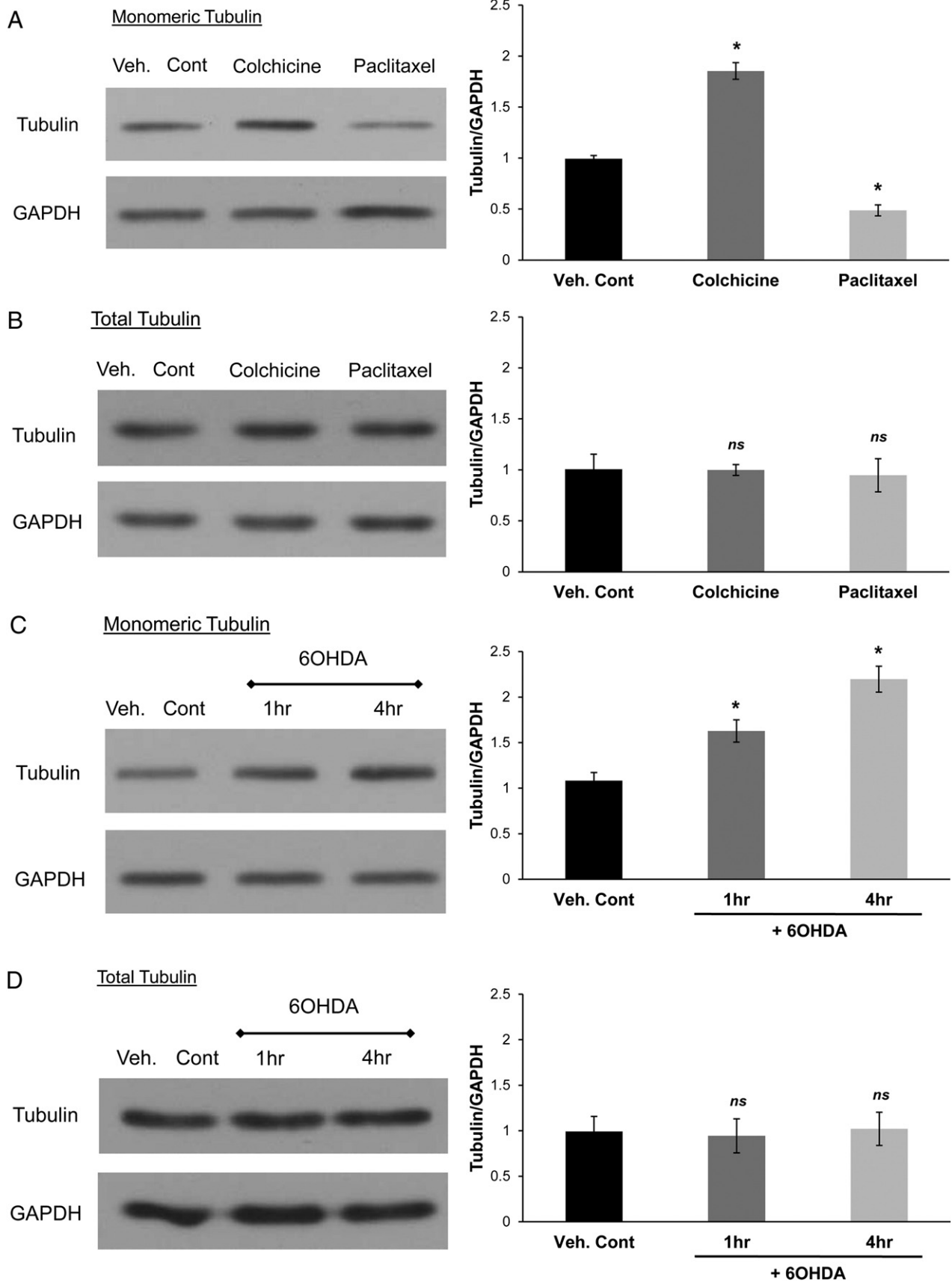
3.8. MTs regulate GR but not Nrf2 nuclear trafficking

We then examined the effects of directly modulating MTs on the trafficking of GR and Nrf2. Colchicine and paclitaxel were utilized to disrupt or stabilize, respectively, the MT network in SH-SY5Y cells expressing GR-GFP and Nrf2-GFP (Fig. 8). SH-SY5Y cells expressing GR-GFP were exposed to these agents at doses and times indicated above and then treated with Cort to induce the nuclear translocation of GR-GFP. MT stabilization using paclitaxel did not alter the nuclear import of GR-GFP, however a decrease in nuclear import was observed in cells where the MT network was disrupted by colchicine treatment (Fig. 8A). Neither MT stabilization nor disruption significantly impaired Nrf2-GFP subcellular localization compared to vehicle in response to 6OHDA stress (Fig. 8B). Thus, the effect of 6OHDA on MT stability was similar to that of colchicine, and colchicine recapitulated the selective nuclear import impairment elicited by 6OHDA.

4. Discussion

In this study, we examined the effects of 6OHDA on the nucleocytoplasmic trafficking of two transcription factors that are sequestered in the cytoplasm, but utilize distinct mechanisms for release from cytoplasmic anchors and import into the nucleus. Previously published work has demonstrated adverse functional effects of altered pCREB or pERK2 subcellular localization on midbrain dopaminergic neuron survival [17,39], but the mechanism underlying such localization changes remained unclear. Utilizing a well-established model of regulated nuclear transport, the glucocorticoid receptor, we found that 6OHDA treatment elicited early impairments in nuclear import without causing reduced nuclear retention or enhanced nuclear export. Treatment with an antioxidant rescued the defect in GR nuclear import, supporting an oxidative mechanism for the 6OHDA-induced effects.

While relatively high doses of H_2O_2 and other oxidants can affect nuclear transport through global mechanisms in non-neuronal cells [10,66–72], the effects of sublethal/prelethal oxidative stress have not been previously studied in neuronal cells. Interestingly, the impaired nuclear import observed in our system did not result from global nuclear import failure as Nrf2 nuclear localization remained intact in 6OHDA-treated cells. The selective nature of these effects suggests a specific alteration that impacts the subcellular localization of some but not all transcription factors. Our in vitro results are consistent with observations that Nrf2 shows clear nuclear localization in post-mortem PD substantia nigra neurons [18], while kinases and transcription factors activated by trophic signals show abnormal cytoplasmic accumulation [5]. These findings suggest that the impaired neurotrophic response may outweigh the intact antioxidant response after oxidative injury, leading to neuronal degeneration.

Interestingly, GR has been linked to the regulation of the dopaminergic system, including through its interactions with the nuclear receptor related 1 protein (Nurr1)—which is important for dopaminergic differentiation and phenotype maintenance as well as being associated with PD [73–83]. Furthermore, glucocorticoids have been shown to play a protective role in models of PD, although the mechanisms may involve more than one cell type in the brain [74,77]. In any case, selective impairment of a subset of transcription factors in the aging or oxidatively stressed brain may account for downregulation of dopaminergic differentiation markers observed in PD models [84–90] and for alterations in neurotrophic gene expression observed in PD midbrain tissues [1,91–94].

The mechanism(s) underlying the selectivity in nuclear trafficking of proteins observed in this model of parkinsonian injury and in diseased PD brains may be of particular relevance to chronic neurodegeneration. Since not all transcription factors are affected, a global mechanism that would impair all nuclear transport, such as bioenergetic collapse, is unlikely. This is further supported by the early time points at which the trafficking impairment becomes apparent. The current data implicate a role for altered microtubule dependent trafficking. Some transcription factors utilize MTs for their transport. Indeed, a sequence within the parathyroid hormone related protein was recently found to efficiently transport fusion proteins into the nucleus via a MT facilitated "fast track" mechanism [95]. Oxidative stress (e.g. H_2O_2) has been linked to impaired MT function, such as decreased tubulin polymerization rate and MT growth rate [96–99]. In our model, we found that 6OHDA increased the levels of free monomeric tubulin similar to colchicine, suggesting enhanced depolymerization and/or reduced polymerization. Furthermore, colchicine-mediated disruption but not stabilization of MTs mimicked the effects of 6OHDA in reducing nuclear import of GR without affecting Nrf2 import. These results suggest that 6OHDA modulates transcription factor trafficking through effects on MT-dependent transport of transcription factors.

How could 6OHDA-induced oxidative stress affect the state of tubulin polymerization? A possible mechanism is through the direct oxidative modification of the tubulin heterodimer, which has numerous cysteine residues. Oxidation of cysteine residues has been shown to reduce tubulin polymerization rate and hence could explain the increased levels of unpolymerized tubulin after 6OHDA treatment [100–102]. Furthermore, rates of tubulin polymerization and depolymerization are also regulated by microtubule-associated proteins (MAPs) and the binding of these proteins to MTs could be directly regulated by oxidative modifications [98,103–105]. The binding of MAPs to MTs is also regulated by tubulin post-translational modifications (PTMs), such as acetylation, tyrosination/detyrosination, and polyglutamylation [106,107]. Modulation of tubulin PTMs by oxidative stress therefore represents another means by which 6OHDA could impact the state of tubulin polymerization.

In addition to altered MT dynamics, there are other potential mechanisms that could contribute to selective impairments in nuclear transport. Oxidative injury, e.g. H_2O_2, has been shown to alter the distribution of the small G protein Ran, which is a key player in the classic import/export cycle [68,70,71,108]. Although such an alteration would be expected to affect nuclear transport of many proteins, it is possible that different transcription factors have differential sensitivity to the breakdown of the Ran gradient across the nucleus or utilize an alternative non-Ran dependent mechanism for their nuclear import [11,109]. Another possibility is the involvement of carrier proteins for nuclear import, importins. Different transcription factors utilize different importins for their nuclear import, as is the case for CREB, GR, and Nrf2—hence oxidative stress-induced alterations in different importins could also help explain the observed selectivity [110–112]. Evidence of altered levels or localization of specific importins has been described in various neurodegenerative diseases [10].

In summary, we have found that 6OHDA selectively impairs nuclear import through an oxidative mechanism that may involve alterations in MT-dependent trafficking of transcription factors. Given the importance of transcription factors for regulating gene expression, alterations in their nucleocytoplasmic transport could have significant consequences for neuronal differentiation, function, and survival. For chronic diseases such as PD, oxidative stress related to aging, the cellular handling of the oxidative catecholamine neurotransmitters, or mitochondrial pathology

Fig. 7. 6OHDA increases levels of unpolymerized tubulin. (A) SH-SY5Y cells were treated with either vehicle, 1 μM colchicine (2 h), or 50 nM paclitaxel (1 h) and then incubated with tubulin extraction buffer to extract monomeric tubulin and resolved by Western blotting. Colchicine increased and paclitaxel reduced the levels of monomeric tubulin, respectively. Mean $+/-$ SEM, $*p < 0.05$ vs. Veh. Cont. (B) Whole cell extracts of colchicine and paclitaxel treated cells were subjected to Western blotting, which showed no significant changes in total tubulin levels. Mean $+/-$ SEM, $p > 0.05$ vs. Veh. Cont. (C) SH-SY5Y cells were treated with 150 μM 6OHDA for 1 h and 4 h after which monomeric tubulin was extracted. 6OHDA-induced oxidative stress increased the levels of monomeric tubulin. Mean $+/-$ SEM, $*p < 0.05$ vs. Veh. Cont. (D) Whole cell extracts of 6OHDA-treated cells were subjected to Western blotting, which showed no changes in total tubulin levels. Mean $+/-$ SEM, $p > 0.05$ vs. Veh. Cont. Western blot results were quantified and compiled from 3 independent experiments.
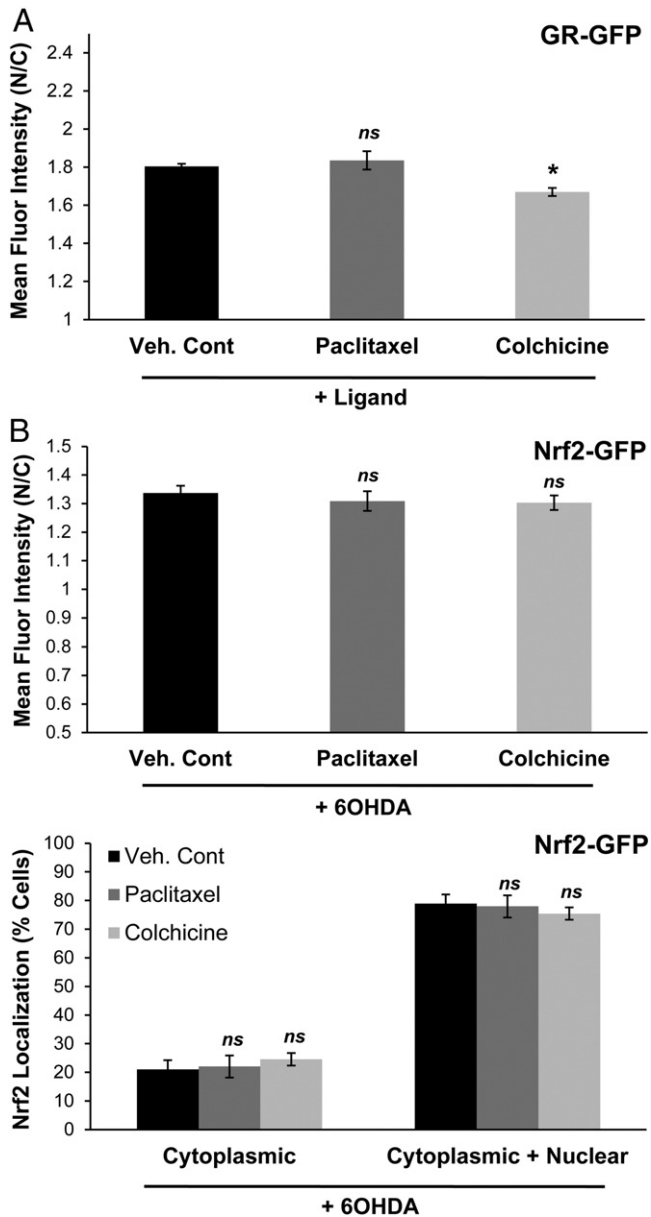
Fig. 8. MTs regulate GR but not Nrf2 nuclear trafficking. (A) GR-GFP expressing SH-SY5Y cells were treated with either paclitaxel or colchicine and then treated with Cort for 30 min to induce GR-GFP nuclear import. Reduced GR-GFP nuclear import was observed in colchicine treated cells but not in paclitaxel treated cells. Mean $+/-$ SEM, $*p < 0.05$ vs. Veh. Cont. (B) Nrf2-GFP expressing SH-SY5Y cells were treated with either paclitaxel or colchicine and then treated with 150 μM 6OHDA for 30 min to induce Nrf2-GFP nuclear localization. No significant impairment was observed with either MT-altering agent. Mean $+/-$ SEM, $p > 0.05$ vs. Veh. Cont. Compiled from 3 independent experiments.

could contribute to reduced trophic and reparative transcriptional responses through selective impairments in nuclear import.

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