

Article

A Potent (R)-alpha-bis-lipoyl Derivative Containing 8-Hydroxyquinoline Scaffold: Synthesis and Biological Evaluation of Its Neuroprotective Capabilities in SH-SY5Y Human Neuroblastoma Cells

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Abstract: A novel bis-lipoyl derivative containing 8-hydroxyquinoline scaffold (LA-HQ-LA, **5**) was synthesized as a new multifunctional drug candidate with antioxidant, chelant, and neuroprotective properties for the treatment of neurodegenerative diseases. We have investigated the potential effectiveness of LA-HQ-LA against the cytotoxicity induced by 6-OHDA and H₂O₂ on human neuroblastoma SH-SY5Y cell line. Our outcomes showed that LA-HQ-LA resulted in significant neuroprotective and antioxidant effects against H₂O₂- and 6-OHDA-induced neurotoxicity in human neuroblastoma SH-SY5Y cells, as assessed by MTT assay. In particular, it showed potent neuroprotective effects against 6-OHDA in RA/PMA differentiated cells at all the tested concentrations.

Keywords: (R)-alpha-lipoic acid; 8-hydroxyquinoline; neuroprotective agent; antioxidants; neurodegenerative diseases

1. Introduction

Neurodegenerative diseases comprise a condition in which nerve cells from brain and spinal cord are lost leading to either functional loss (ataxia) or sensory dysfunction (dementia). Mitochondrial dysfunctions, excitotoxicity, and, finally, apoptosis have been reported as pathological causes for aging and neurodegenerative diseases such as Parkinson's disease (PD), Alzheimer's disease (AD), multiple sclerosis (MS), and amyotrophic lateral sclerosis (ALS) [1]. Although any disease mentioned above is characterized by its own molecular mechanism and clinical onset, common pathways can be identified as: 1) oxidative stress and free radicals formation; 2) metal dyshomeostasis; 3) protein misfolding and aggregation; and 4) mitochondrial dysfunctions [2]. Oxidative stress plays a key role in neurodegenerative diseases: it arises due to an imbalance between pro-oxidant/antioxidant homeostasis that further takes part in generation of ROS and free radicals potentially toxic for neuronal cells [3]. Furthermore, in all forms of ROS generation molecular oxygen needs to be activated by a range of metallo-enzymes, thus facilitating ROS generation upon interaction of redox metals with O₂ using various catalytic pathways. Since free radicals are toxic to cells, under normal circumstances, cells have an efficient regulating system for O₂ and metal ion interaction leading to free radicals and ROS generation [4]. In fact, during the Fenton reaction, hydroxyl radicals are produced from hydrogen peroxide in the presence of a metal in a low oxidation state:



Fenton chemistry may occur in neurons of the nervous tissue where levels of both hydrogen peroxide and cerebral biometals—such as Fe³⁺, Cu²⁺, and Zn²⁺—are found. Several studies have indicated that cerebral biometal dyshomeostasis and oxidative stress are intimately associated [5]. An *in vitro* AD model has shown that both Aβ-40 and Aβ-42 deposits are formed after incubation of immobilized β-amyloid oligomers with Cu²⁺, Zn²⁺, or Fe³⁺. In these conditions, Fe³⁺ promoted the deposition of fibrillar amyloid plaques, while Cu²⁺ and Zn²⁺ only induced the formation of amorphous aggregates [5]. In an *in vitro* PD model, it has been found that Fe³⁺ enhanced intracellular aggregation of α-synuclein and led to the formation of advanced glycation end products. The accumulation of these factors strongly contributed to the progression of the neurodegenerative process [6].

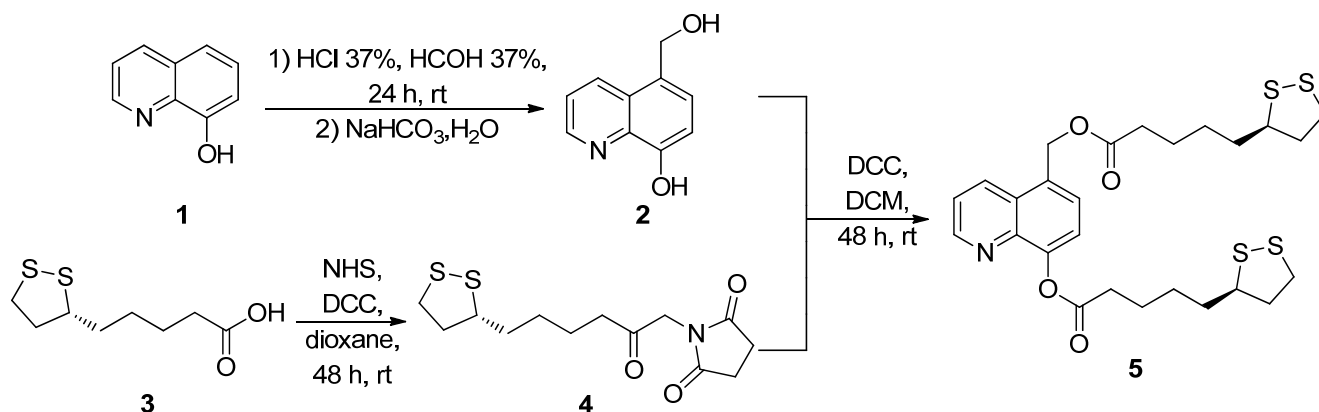
Increasingly medicinal chemistry approaches are currently under study to discover new drugs able to remove excess of specific metals [7,8] and to prevent or block the oxidative process that characterizes PD and AD [9–12]. Taking into account that drugs with two or more useful biological activities for the same pathology may represent an important pharmacological advance, we are currently interested in multifunctional drugs that combine potent antioxidant, chelant, and neuroprotective properties in a single molecule for the treatment of PD and AD [13–16]. For this purpose, to design a novel class of compounds with a multimodal mechanism of action, we selected the hydroxyquinoline (HQ) scaffold as a privileged structure since it is a clinically relevant bioactive metal chelator. Recently, 8-hydroxyquinoline (8-HQ, **1**) derivatives have found application in PD and AD drug discovery [17] since 8-HQ: 1) is able to cross the blood–brain barrier (BBB) [18]; 2) is a strong iron chelator with antioxidant property [19–21]; and 3) is able to protect against the precipitation of β-amyloid plaques in presence of Cu²⁺, Fe³⁺, Zn²⁺—compared to clioquinol—due to its ability to chelate these metals [22].

The aim of this work was to combine the antioxidant and neuroprotective properties of (R)-alpha-lipoic acid (LA, **3**) and the chelant activities of 8-HQ (**1**) [23] to obtain a novel multi-target ligand, LA-HQ-LA (**5**) with multifunctional neuroprotective profile. LA-HQ-LA was obtained by linking via two ester bonds the 8-HQ derivative (5-hydroxymethyl-8-hydroxyquinoline, **2**) to LA, thus increasing the lipophilicity of this molecule. LA-HQ-LA can cross plasma membranes and release HQ and two molecules of LA, thus triggering a significant decrease in oxidative stress from human SH-SY5Y neuroblastoma cells. In addition, due to the different chemical nature of the ester bonds, the derivative **5** could gradually provide a continuative and time-controlled release of LA—an elevator of GSH levels that are lower in some cerebral areas of patients affected by neurodegenerative diseases [24]—and HQ directly to specific groups of neurons characterized by cellular stress and metals accumulation.

2. Results and Discussion

Starting from 8-HQ (**1**), the required starting material 5-hydroxymethyl-8-hydroxyquinoline (**2**) was obtained in good yield, using a known procedure [25]. The new multi-target ligand LA-HQ-LA (**5**) was synthesized by direct condensation of 5-hydroxymethyl-8-hydroxyquinoline (**2**) and LA-NHS (**4**), previously prepared as reported by Nefkens *et al.* [26] (Scheme 1). The chemical structure of LA-HQ-LA was confirmed by ^1H -, ^{13}C -NMR, IR, and MS spectra data.

Scheme 1. Synthesis of LA-HQ-LA (**5**).

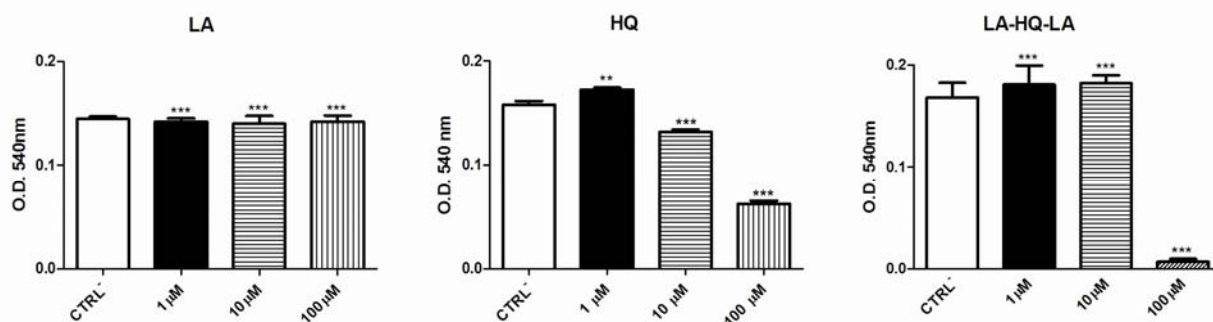


The neuroprotective and antioxidant capacities of LA-HQ-LA against oxidative stress were assayed by using the human SH-SY5Y neuroblastoma cell line, which is a reliable model for studying the neurotoxic effect of agents such as H₂O₂, 6-OHDA, and frequently used for elucidating the mechanisms of neurodegenerative diseases [27].

First of all, to define the suitable concentration range, the effects on cell proliferation of LA, HQ, and LA-HQ-LA were determined by colorimetric MTT assay (Figure 1). Thus, we performed dose-response experiments (with compound concentrations of 1, 10, and 100 μM) to verify if, 24 h after the treatment, the compounds added to the cells had any effect on the cell proliferative capacity. The compound concentrations of 1 and 10 μM did not show significant differences compared to the control, while at 100 μM , an antiproliferative activity was observed (Figures 1–2). In particular, at 100 μM , LA still retained a proliferative activity, while LA-HQ-LA resulted in an antiproliferative activity,

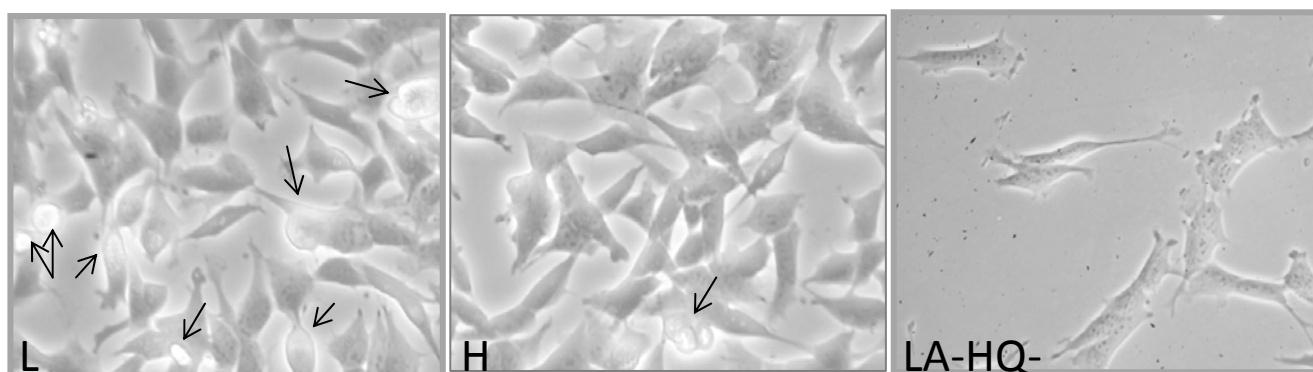
since there are no visible proliferating cells (Figure 2). Based on the results obtained, the utilized compound concentration in all the experiments reported in this study was 1 μ M.

Figure 1. Dose-response effects of LA, HQ, and LA-HQ-LA in undifferentiated SH-SY5Y human neuroblastoma cells^a.



^aMTT reduction assay in undifferentiated SH-SY5Y human neuroblastoma cells in the presence of LA, HQ, and LA-HQ-LA. The cells were incubated for 24 h with increasing concentrations (1, 10, and 100 μ M) of the compounds. After this period, cell viability was quantified by measuring MTT reduction. CTRL⁻: negative control w/o toxic. The means $\pm$ SEM derived from 3 different experiments (each with n=16; *** $p < 0.0001$, ** $0.0001 < p < 0.001$, * $0.001 < p < 0.05$, n.s. $p > 0.05$). * refers to SH-SY5Y human neuroblastoma cells treated only with the neuroprotective compound.

Figure 2. Morphological analysis of undifferentiated SH-SY5Y human neuroblastoma cells in the presence of LA, HQ, and LA-HQ-LA.^a

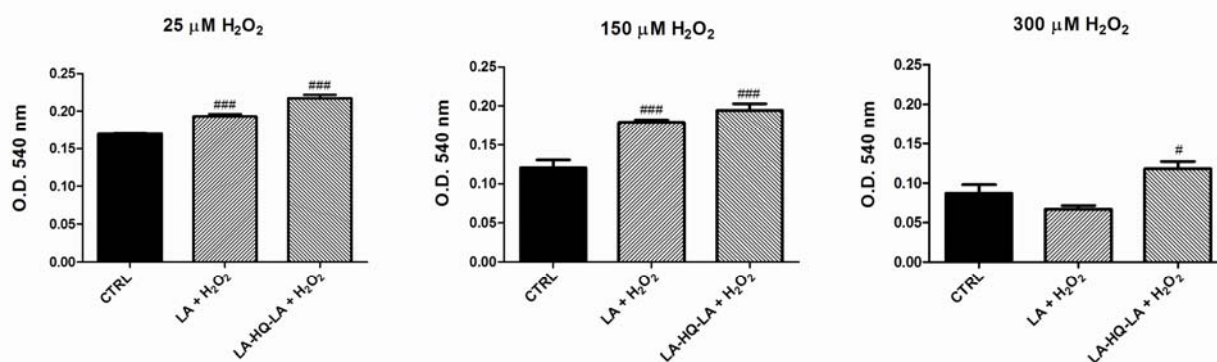


^aMorphological analysis by phase-contrast microscopy of undifferentiated SH-SY5Y human neuroblastoma cells. The cells were incubated with LA, HQ, and LA-HQ-LA (100 μ M) for 24 h. The arrows indicate the cell in proliferation. (Magnification 40X)

Because the sensitivity of cells to toxic agents may be dependent on the state of their neuronal differentiation, we used undifferentiated and RA-differentiated SH-SY5Y cells to study the neuroprotective effect of our compounds following H_2O_2 insult [28-29]. Thus, undifferentiated SH-SY5Y cells were pre-treated with LA, HQ, and LA-HQ-LA for 24 h and then exposed to H_2O_2 (25-150-300 μ M) for other 24 h (cell viability was detected using MTT assay as shown in Figure 3).

Both compounds (LA-HQ-LA and LA) showed a significant protective effect against H_2O_2 at 25 and 150 μM H_2O_2 . Moreover, the neuroprotective effect of LA-HQ-LA was appreciably higher in respect to LA and the control at 300 μM . In Figure 3, we did not insert the results obtained for HQ since, as evidenced by the morphological evaluation (Figure 4), at 1 μM and in presence of 25 μM of H_2O_2 it was demonstrated as toxic for the undifferentiated SH-SY5Y cells. This deleterious effect on SH-SY5Y cells could be due to a synergic action between HQ and H_2O_2 . HQ, being a metal chelator of iron, copper, and zinc (transition metals that react easily with ROS), avoids the Haber-Weiss reaction between the metal and the superoxide anion; in these oxidative stress conditions, superoxide anion is overproduced, thereby damaging cells. On the other hand, H_2O_2 directly produces a high quantity of ROS further damaging the cells. These two combined actions determine a deleterious synergic effect on cells.

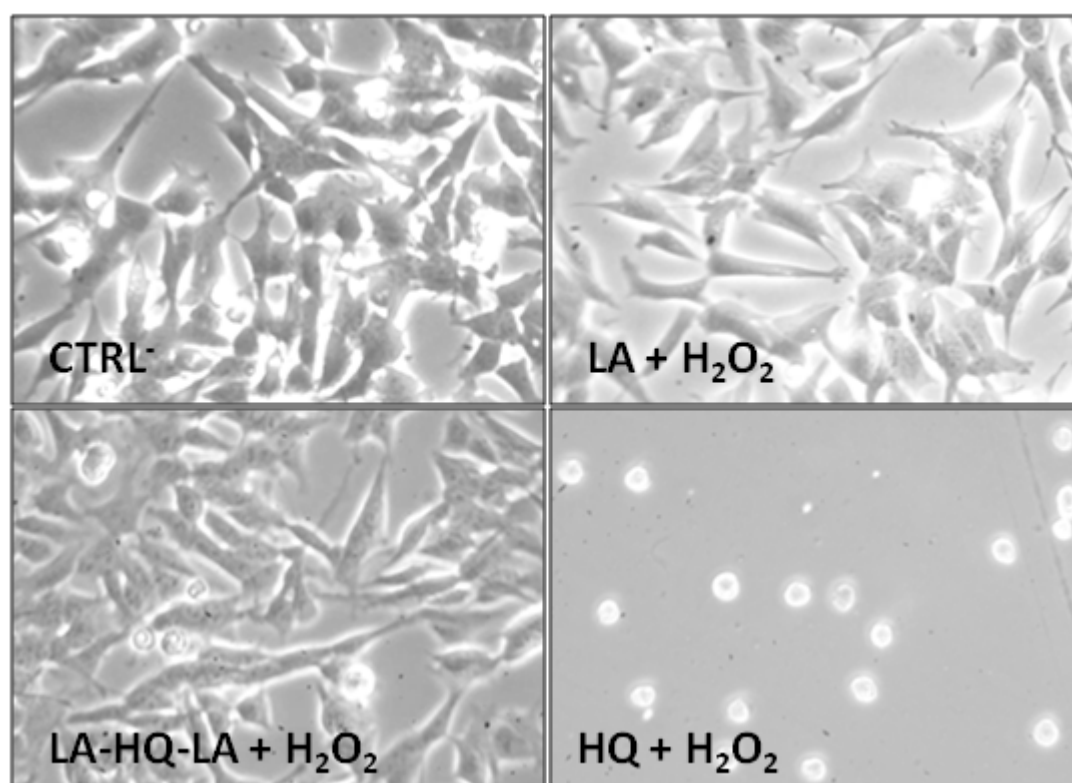
Figure 3. Neuroprotective effects of LA and LA-HQ-LA in undifferentiated and H_2O_2 -lesioned SH-SY5Y human neuroblastoma cells.^a



^a MTT reduction assay in undifferentiated and H_2O_2 -lesioned SH-SY5Y human neuroblastoma cells in the presence of LA and LA-HQ-LA. The cells were incubated with the compounds (1 μM) 24 h before and during a 24-h incubation period with increasing concentrations (25, 150, or 300 μM) of H_2O_2 . After this period, cell viability was quantified by measuring MTT reduction. CTRL: control with H_2O_2 . The means $\pm$ SEM derived from 3 different experiments (each with $n=16$; ### $p < 0.0001$, ## $0.0001 < p < 0.001$, # $0.001 < p < 0.05$, n.s. $p > 0.05$). # is refers to SH-SY5Y human neuroblastoma cells treated with a combination of neuroprotective compound and the toxin.

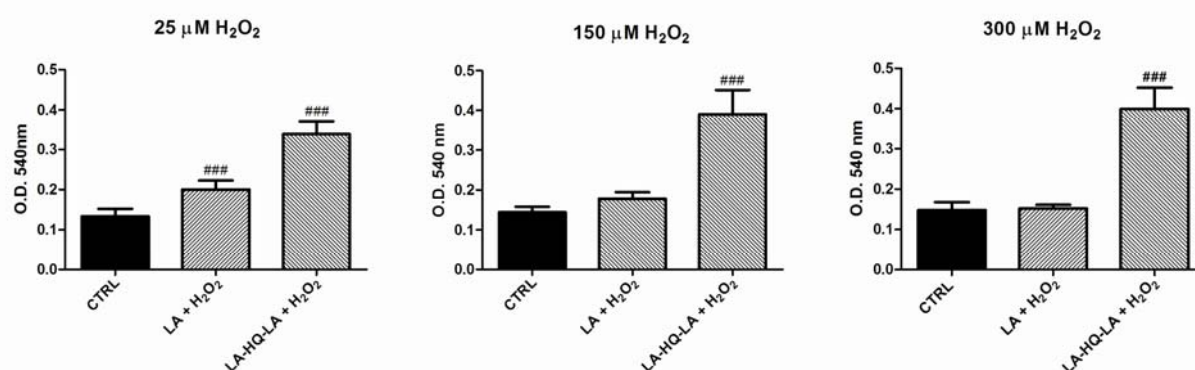
Many evidences indicated that the neuronal differentiation of SH-SY5Y neuroblastoma cells seems appropriate for studying neurotoxicity since the proliferation rate is limited and cells morphologically resemble the neuronal phenotype [30]. Thus, we differentiated our cells with RA to increase the cholinergic properties of SH-SY5Y cell line and we lesioned them with increasing concentrations (25–150–300 μM) of H_2O_2 (Figure 5). Results confirmed a significant neuroprotective effect of LA-HQ-LA, at all doses of H_2O_2 , against LA and control. These results confirmed several previous reports [28, 31–32] which showed that the differentiation of SH-SY5Y cells by RA enhanced their resistance to the action of neurotoxic agents. In fact, comparing Figures 3 and 5, indicating cellular viability, we observed that at high concentrations of H_2O_2 (300 μM), the optical density O.D. values were reduced more in the undifferentiated than differentiated SH-SY5Y cells. Thus, the undifferentiated cells were much more vulnerable to H_2O_2 than RA-treated ones.

Figure 4. Morphological analysis of undifferentiated and H_2O_2 -lesioned SH-SY5Y human neuroblastoma cells in the presence of LA, LA-HQ-LA, and HQ.^a



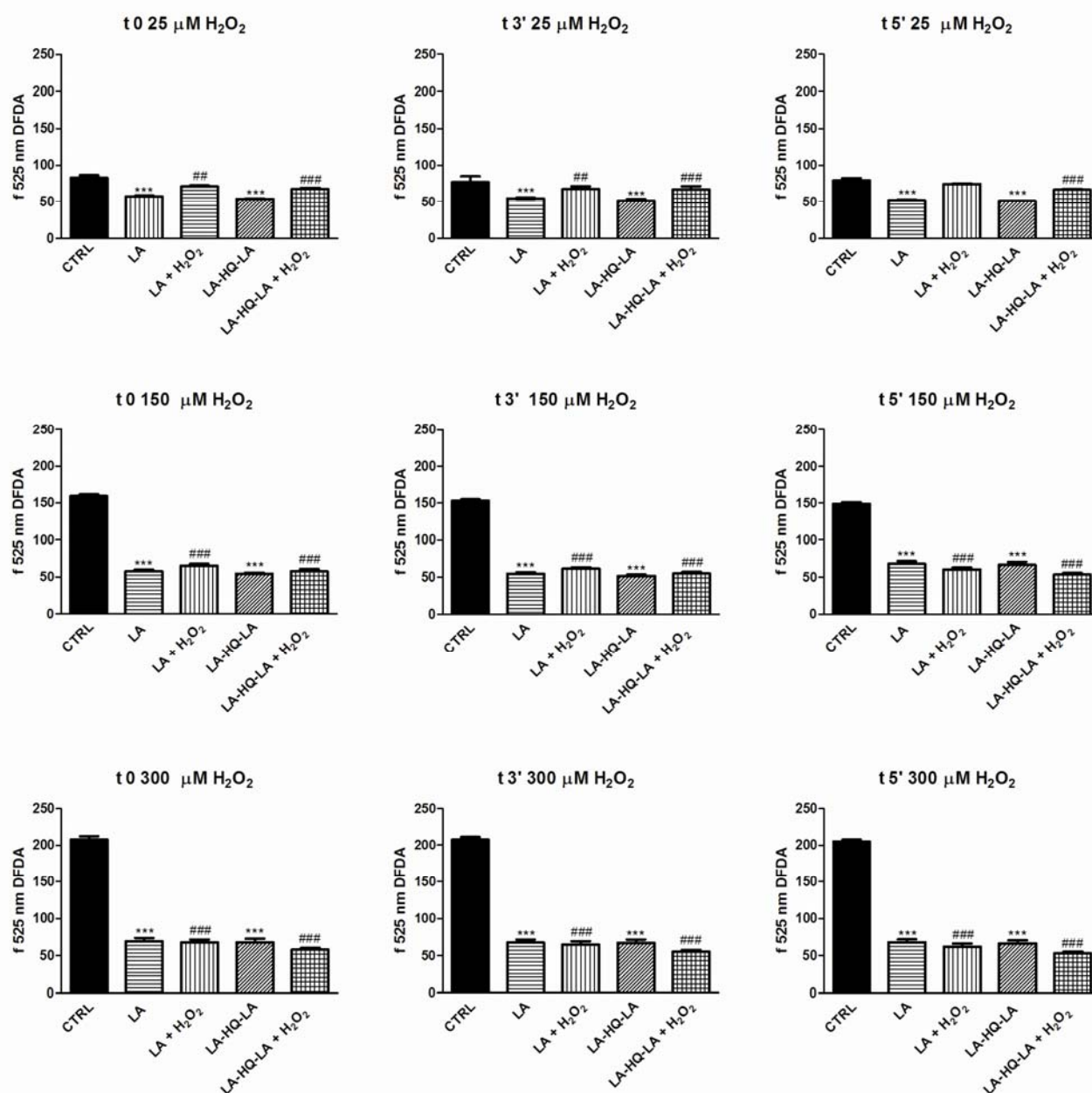
^a Morphological analysis by phase-contrast microscopy of undifferentiated and H_2O_2 -lesioned SH-SY5Y human neuroblastoma cells. The cells were incubated with LA, LA-HQ-LA, and HQ (1 μ M) 24 h before and during a 24-h incubation period with 25 μ M of H_2O_2 . CTRL⁻: negative control w/o H_2O_2 . (Magnification 40X)

Figure 5. Neuroprotective effects of LA and LA-HQ-LA in RA-differentiated and H_2O_2 -lesioned SH-SY5Y human neuroblastoma cells.^a



^a MTT reduction assay in RA-differentiated and H_2O_2 -lesioned SH-SY5Y human neuroblastoma cells in the presence of LA and LA-HQ-LA. The cells were incubated with the compounds (1 μ M) 24 h before and during a 24-h incubation period with increasing concentrations (25, 150, or 300 μ M) of H_2O_2 . After this period, cell viability was quantified by measuring MTT reduction. CTRL: control with H_2O_2 . The means $\pm$ SEM derived from 3 different experiments (each with $n=16$; ### $p < 0.0001$, ## $0.0001 < p < 0.001$, # $0.001 < p < 0.05$, n.s. $p > 0.05$). # is refers to SH-SY5Y human neuroblastoma cells treated with a combination of neuroprotective compound and the toxin.

Figure 6. Antioxidant capabilities of LA and LA-HQ-LA in RA-differentiated and H₂O₂-lesioned SH-SY5Y human neuroblastoma cells ^a.



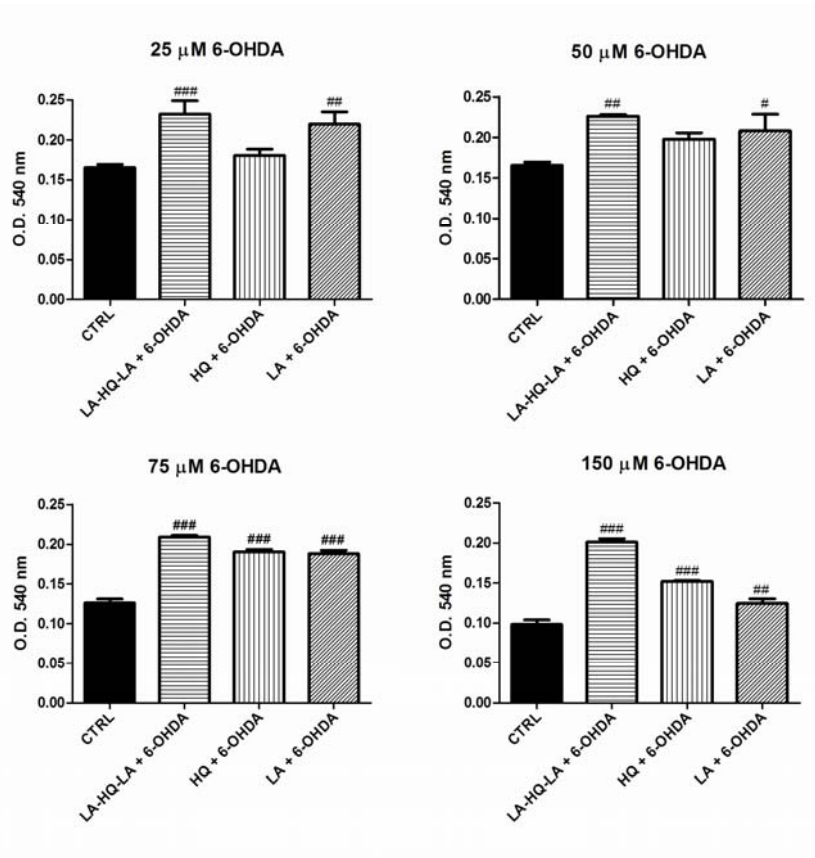
^a Quantitative analyses of ROS expression. ROS measurement by H₂DCF-DA fluorescence in RA-differentiated SH-SY5Y human neuroblastoma cells exposed to LA-HQ-LA or LA 24 h before and during a 5-minute incubation period with increasing concentrations (25, 150, or 300 μM) of H₂O₂. In this figure the fluorescence at three points (t₀, t₃ and t₅) during the time-course are reported. CTRL: control with H₂O₂. The means ± SEM derived from two different experiments (each with n=10; ### or *** p < 0.0001, ## or ** 0.0001 < p < 0.001, # or * 0.001 < p < 0.05, n.s. p > 0.05). * refers to SH-SY5Y human neuroblastoma cells treated only with the neuroprotective compound. # refers to SH-SY5Y human neuroblastoma cells treated with a combination of the neuroprotective compound and the toxin.

In the above reported experiments, exposing SH-SY5Y neuroblastoma cells to increasing concentrations of H₂O₂, we induced reactive oxygen species (ROS) generation. An overproduction of

ROS and a lower antioxidant capability of the cells result in oxidative stress that characterizes several neurodegenerative pathologies. To accurately measure ROS and the cell capability to counteract this insult, we used cell permeable fluorescent and chemiluminescent probes. 2'-7'-Dichlorodihydrofluorescein diacetate (H₂DCF-DA) is one of the most widely used techniques for directly measuring the redox state of a cell [33]. Our results showed that, at t_0 - t_5 and at all the concentrations of H₂O₂, LA-HQ-LA and LA exerted a powerful antioxidant effect returning ROS levels similar to the control (Fig. 6). These data confirmed that our compound possesses good antioxidant and neuroprotective capabilities.

Comparing Figures 3, 5, and 6 we can observe that LA-HQ-LA showed a long-lasting neuroprotective activity respect to LA only in long-term experiments (Fig. 3, 5) probably due to a time-controlled release of its components.

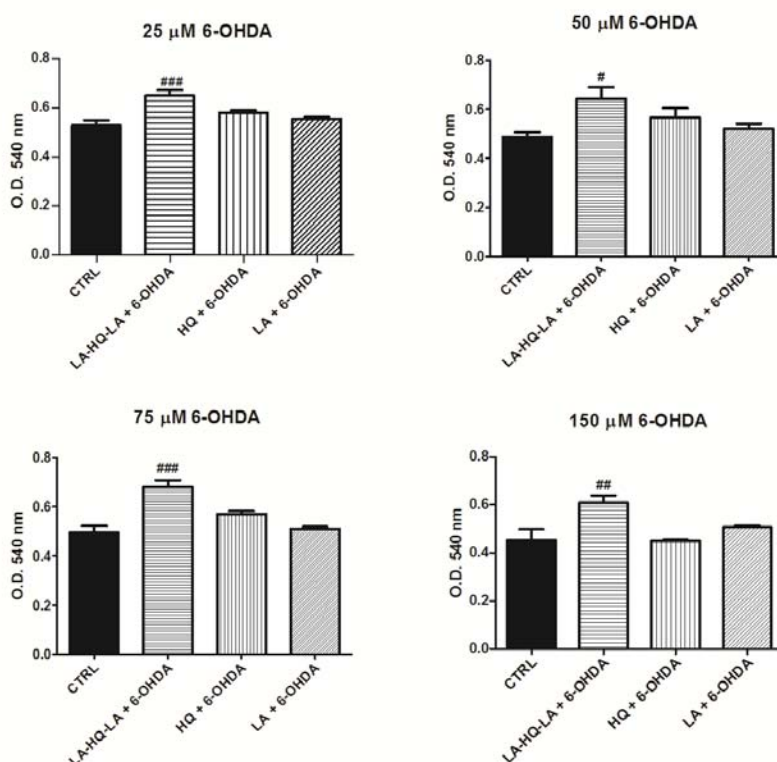
Figure 7. Neuroprotective effects of LA, HQ, and LA-HQ-LA in undifferentiated and 6-OHDA-lesioned SH-SY5Y human neuroblastoma cells ^a.



^a MTT reduction assay in undifferentiated and 6-OHDA-lesioned SH-SY5Y human neuroblastoma cells in the presence of LA, HQ, and LA-HQ-LA. The cells were incubated with the compounds (1 μM) 1 h before and during a 24 h incubation period with increasing concentrations (25, 50, 75, and 150 μM) of 6-OHDA. After this period, cell viability was quantified by measuring MTT reduction. CTRL: control with 6-OHDA. The means ± SEM derived from 3 different experiments (each with n=16; ### $p < 0.0001$, ## $0.0001 < p < 0.001$, # $0.001 < p < 0.05$, n.s. $p > 0.05$). # refers to SH-SY5Y human neuroblastoma cells treated with a combination of the neuroprotective compound and the toxin.

To further investigate the neuroprotective role in PD, it was necessary to differentiate SH-SY5Y neuroblastoma cells toward the DAergic phenotype using RA/PMA [34, 35]. We used the neurotoxin 6-OHDA as a toxic agent to lesionate the cells, as it is commonly used as a dopaminergic degeneration model for both *in vitro* and *in vivo* studies [36]. Like DA, 6-OHDA is quickly oxidized to form a variety of free radical species, and also induces ROS-dependent apoptosis in dopaminergic cells. In this second toxicity model, the neuroprotective effect of LA, HQ, and LA-HQ-LA against oxidative stress was evaluated by using the neurotoxin 6-OHDA in both the undifferentiated and RA/PMA-differentiated cells. The undifferentiated SH-SY5Y cells were treated with LA, HQ, and LA-HQ-LA for 1 h and then exposed to increasing concentrations of 6-OHDA (25–50–75–150 μM). After further 24 h of incubation, the cultures were assessed for viability by MTT assay (Figure 7). LA-HQ-LA was the most neuroprotective compound, among the ones investigated, mostly at the strongly neurotoxic 6-OHDA concentration of 150 μM . In RA/PMA-differentiated cells (Figure 8), LA-HQ-LA showed a strong neuroprotective effect at all the used concentrations of 6-OHDA. Also in these experiments, we observed a major susceptibility to toxic insult of undifferentiated respect to differentiated cells (Figures 7–8).

Figure 8. Neuroprotective effect of LA, HQ, and LA-HQ-LA in RA/PMA-differentiated and 6-OHDA-lesioned SH-SY5Y human neuroblastoma cells.^a



^a MTT reduction assay in RA/PMA-differentiated and 6-OHDA-lesioned SH-SY5Y human neuroblastoma cells in the presence of LA, HQ, and LA-HQ-LA. The cells were incubated with the compounds (1 μM) 1 h before and during a 24 h incubation period with increasing concentrations (25, 50, 75, and 150 μM) of 6-OHDA. After this period, cell viability was quantified by measuring MTT reduction. CTRL: control with 6-OHDA. The means $\pm$ SEM derived from 3 different experiments (each with $n=16$; ### $p < 0.0001$, ## $0.0001 < p < 0.001$, # $0.001 < p < 0.05$, n.s. $p > 0.05$). # refers to SH-SY5Y human neuroblastoma cells treated with combination of neuroprotective compounds and the toxin.

Our data showed that LA-HQ-LA has a significant neuroprotective effect against both H_2O_2 and 6-OHDA, higher than LA and HQ, up to 150 μM . However, LA-HQ-LA showed a different behavior in the response (mainly with 6-OHDA) in differentiated cells compared to the undifferentiated ones. This difference probably depends on the variable sensitivity of cells against neurotoxic agents. In fact, RA/PMA-differentiated cells exhibit 6-fold higher levels of dopamine D_2 and D_3 receptors in respect to undifferentiated or RA-differentiated cells [37]. Furthermore, it is also reasonable that LA-HQ-LA showed the greatest effect against neurotoxic agents, because LA itself is a neuroprotective agent for dopaminergic neurons [38]. The obtained results confirm that LA-HQ-LA is a valid molecule, potentially more effective in respect to LA and HQ. In particular, LA-HQ-LA—at low concentrations (1 μM)—might be a good therapeutic choice in diseases such as PD, where oxidative stress caused by oxidants plays a key role.

3. Experimental Section

3.1. Chemistry

HQ (**1**) and LA (**3**) were purchased from Sigma Chemical Co. (St Louis MO, USA). All other chemicals used were of the highest commercially available purity. Chromatographic purifications were performed on silica gel using column chromatography (Merck 60, 70–230 mesh ASTM silica gel), and compounds were detected with UV light ($\lambda = 254$ nm). Optical rotations were taken at 20 °C with a Perkin-Elmer 241 polarimeter (Santa Clara, CA, USA). IR spectra were recorded with a Varian 1000 FT-IR spectrometer. NMR spectra were recorded with a Varian VXR-300 spectrometer (Varian Medical Systems, Inc., Palo Alto, CA, USA). Chemical shifts are reported in parts per million (δ) downfield from the internal standard tetramethylsilane (Me_4Si). Mass spectra were obtained by electrospray ionization (ESI) in positive mode using a LCQ (Thermo Finnigan) ion trap mass spectrometer (San Jose, CA, USA) equipped with an electrospray ionization (ESI) source. The capillary temperature was set at 300 °C and the spray voltage at 4.25 kV. The fluid was nebulized using nitrogen (N_2) as both the sheath gas and the auxiliary gas.

The identity of LA-HQ-LA was confirmed by NMR, IR, and MS spectra; the homogeneity was confirmed by TLC on silica gel Merck 60 F254. Before performing biological studies, the purity of LA-HQ-LA was checked by HPLC analysis; it was determined by analytical HPLC using a Waters 600 HPLC (Waters Co., Milford, MA, USA) equipped with an X-Bridge BEH130 C-18, 5 μm , 4.6 $\times$ 250 mm column with Waters 2996 PDA detector, and an isocratic elution using water (0.1% TFA) at flow rate of 1 mL/min. LA-HQ-LA was obtained with purity greater than 98%, determined by analytical HPLC at 280 nm.

3.1.1. Synthesis of 5-Hydroxymethyl-8-hydroxyquinoline (**2**)

A mixture of **1** (2 g, 13.77 mmol), concentrated hydrochloric acid (4.4 mL), and 37% formaldehyde (2.2 mL) was treated with hydrogen chloride gas and stirred overnight. The reaction mixture was filtered and the precipitate was dried giving the 5-hydroxymethyl-8-hydroxyquinoline hydrochloride as a yellow solid (yield: 93%). The product was then solubilized with water and added with $NaHCO_3$ (3.9 g, 46.48 mmol); the precipitate was dissolved and extracted with $CHCl_3/NaCl$, dried over Na_2SO_4 , and

the solvent was removed under vacuum obtaining **2** as a white solid (yield: 68%). $^1\text{H-NMR}$ (300 MHz, DMSO- d_6) δ : 4.78–4.83 (2H, m, CH₂, HQ), 5.16 (1H, s, OH, HQ), 6.96–7.57 (3H, m, Ar, HQ), 8.47–8.83 (2H, m, Ar, HQ), 9.70 (1H, s, br, OH); $^{13}\text{C-NMR}$ (300 MHz, DMSO- d_6): 111.35–153.07 (9 $\times$ CH, HQ).

3.1.2. Synthesis of 1-(-5-[1,2]dithiolan-3-yl-pentanoyl)-pyrrolidine-2,5-dione (LA-NHS, **4**)

A solution of **3** (3.69 g, 19.2 mmol) in dry dioxane (15.4 mL) was slowly added to a mixture of NHS (2.3 g, 20 mmol) and DCC (4.12 g, 20 mmol) in dry dioxane (37 mL) and stirred for 48 h at room temperature. After filtration of the precipitate, the solvent was evaporated, the residue was taken up in 2-propanol, and left overnight at 4 °C. The resulting white precipitate **4** was then filtered and dried (yield: 60%). $^1\text{H-NMR}$ (300 MHz, DMSO- d_6) δ : 1.25 (2H, m, CH₂, LA), 1.37–1.62 (4H, m, 2 $\times$ CH₂, LA), 1.84 (2H, m, CH₂, LA), 2.29 (2H, t, CH₂, LA), 2.42–2.51 (3H, m, CH₂ and CH, LA), 2.78 (4H, t, 2 $\times$ CH₂, NHS); $^{13}\text{C-NMR}$ (300 MHz, DMSO- d_6): 28.1 (CH₂, LA), 29.3 (CH₂, LA), 33.0 (2 $\times$ CH₂, NHS), 33.5 (CH₂, LA), 34.8 (CH₂, LA), 37.9 (CH₂, LA), 42.9 (CH₂, LA), 55.7 (CH, LA), 107.4 (CH, NHS), 172.3 (CO, LA), 205.2 (2 $\times$ CO, NHS).

3.1.3. Synthesis of LA-HQ-LA (**5**)

5-Hydroxymethyl-8-hydroxyquinoline (**2**) (440 mg, 2.5 mmol) was dissolved in dry CH₂Cl₂ (72 mL) and then added to a stirring mixture of **4** (3.03 g, 10 mmol) and DCC (2.06 g, 10 mmol) in dry CH₂Cl₂ (72 mL). The reaction mixture was left at room temperature for 48 h under stirring, then the solution was filtered and the solvent was removed. The obtained residue was chromatographed using petroleum ether: AcOEt (6:4) as eluant to give LA-HQ-LA as a yellow oil (yield: 57%). R_f = 0.72 (petroleum ether:AcOEt (6:4)); $[\alpha]_D^{20} = +20.0$ ($c=1$ in MeOH); $^1\text{H-NMR}$ (300 MHz, CDCl₃) δ : 1.42 (2H, m, CH₂, LA), 1.8 (12H, m, 6 $\times$ CH₂, LA), 2.36 (2H, m, CH₂, LA), 2.47 (2H, m, CH₂, LA), 2.81 (2H, t, CH₂, LA), 3.12 (4H, m, 2 $\times$ CH₂, LA), 3.49 (1H, m, LA), 3.63 (1H, m, LA), 5.52 (2H, s, Ar-CH₂-O), 7.42 (1H, d, HQ), 7.50 (1H, m, HQ), 7.60 (1H, d, HQ), 8.38 (1H, d, HQ), 8.93 (1H, d, HQ). $^{13}\text{C-NMR}$ (300 MHz, CDCl₃) δ : 24.85 (CH₂, LA), 24.93 (CH₂, LA), 28.85 (OOCCH₂-CH₂, LA), 28.96 (OOCCH₂-CH₂, LA), 34.18 (OOC-CH₂, LA), 34.24 (OOC-CH₂, LA), 34.77 (CH₂, LA), 34.9 (CH₂, LA), 38.74 (2 $\times$ CH₂, LA), 40.41 (CH₂, LA), 40.48 (CH₂, LA), 56.49 (CH, LA), 56.65 (CH, LA), 63.48 (COO-CH₂-HQ), 121.07 (CH, HQ), 122.27 (CH, HQ), 128.25 (CH, HQ), 130.30 (C-OCO, HQ), 132.56 (CH, HQ), 148.34 (C-N, HQ), 150.56 (N-CH, HQ), 172.48 (COO, LA), 173.39 (COO, LA); IR (KBr): $\tilde{\nu} = 3409, 2932, 2283, 1733, 1461, 1376, 1259, 1182, \text{cm}^{-1}$; MS (ESI) m/z 552.12 (M+H)⁺.

3.2. Neuroprotective Studies

3.2.1. SH-SY5Y Cell Culture

Human SH-SY5Y neuroblastoma cells (EGACC, Sigma–Aldrich, UK) were grown at 37 °C, in a humid 5% CO₂, in Dulbecco's Modified Eagle's medium supplemented with 10% heat-inactivated fetal bovine serum, penicillin (100 U/mL), streptomycin (100 $\mu\text{g/mL}$), and 1% L-glutamine. SH-SY5Y cells were plated onto 96-well plates (2,700 cell/well). Undifferentiated cells (UN) were

grown for 24 h in normal medium and then incubated with the compounds (LA, HQ, and LA-HQ-LA). After 24 h of incubation, the cultures were assessed for viability by using a colorimetric assay based on the ability of living cells to reduce a tetrazolium-based compound to a blue formazan product (MTT assay). At the end of incubation, 20 μ L of MTT (5 mg/mL in PBS) were added to each well and the incubation continued for an additional 3 h at 37 °C. The plate was centrifuged at 2000 rpm for 15 min. The MTT solution was carefully decanted off and formazan crystals were dissolved with 200 μ L DMSO at 37 °C for 30 min. Finally, samples were read in a Titertek Multiscan Microeliza Reader (Flow Laboratories, Irvine, UT, USA) at 540 nm. All MTT assays were repeated nine times. The neuroprotective effects of HQ (1 μ M), LA (1 μ M), and LA-HQ-LA (1 μ M) in undifferentiated cells were tested for 24 h in the presence of two neurotoxic agents: H₂O₂ (25-150-300 μ M) and 6-OHDA (50-75-150 μ M).

To obtain a cholinergic phenotype, SH-SY5Y cells were grown in a medium containing retinoic acid (RA) (10 μ M) for 3 days; then the medium was removed and replaced with fresh RA (10 μ M) medium for other 3 days of differentiation. Alternatively, to have a dopaminergic phenotype, the cells were treated with RA/ phorbol 12-myristate 13-acetate (PMA) in a medium supplemented of RA (10 μ M) for 3 days; then the medium was removed and replaced with growth medium containing PMA (80 nM) for a subsequent 3 days. The neuroprotective effects of LA, HQ, and LA-HQ-LA at the concentration of 1 μ M were evaluated on differentiated cells exposed to 6-OHDA (25-50-75-150 μ M) and H₂O₂ (25-150-300 μ M).

3.2.2. Measurement of Intracellular ROS

Intracellular ROS were quantified by the 2',7'-dichlorodihydrofluorescein diacetate (H₂DCF-DA) assay using a Microplate Fluorometer SPECTRAmax Gemini XS (Molecular Devices, Sunnyvale, CA, U.S.A.) at excitation and emission wavelengths of 480 nm and 530 nm, respectively, and analyzed by SOFTmax Pro software (Version 5.0; Molecular Devices). DCFH-DA is transported across the cell membrane and deacetylated by esterases to form the non-fluorescent 2',7'-dichlorfluorescein (DCFH). This compound is trapped inside the cells. Next, DCFH is converted to DCF through the action of peroxide rated by the presence of peroxidase. SH-SY5Y cells were plated (2000 cells/well) into special-optics 96-well plates (Corning-Costar). 24 hours later, the cells were washed 3 times with imaging buffer (125 mM NaCl, 5 mM KCl, 1.2 mM MgSO₄, 5 mM glucose, 25 mM Hepes, 2 mM CaCl₂). Then 10 μ M DCFH-DA media solution were added and the plates were incubated at 37 °C for 30 minutes. After 2 washings with imaging buffer, cells were treated with 25, 150, or 300 μ M H₂O₂ for immediate fluorescence measurement. The fluorescence intensity is proportional to the ROS levels within the cell cytosol and in inverse proportion to the antioxidant capacity of the cells themselves. Plates, maintained at 25 °C, were read for kinetic analysis in increments from 0 to 5 minutes every 30 seconds [39].

Statistical analysis. The statistical analysis (unpaired *t* test) was performed with GraphPad Prism 5 software version 5.0. One-way ANOVA was computed for each level of treatment followed by Dunnett's *t*-test *post hoc*.

4. Conclusions

In conclusion, we synthesized a novel multi-target ligand—containing antioxidant and chelant groups—that might represent a step forward in the search for new molecules against neurodegenerative diseases. LA-HQ-LA was superior to the single molecules (LA and HQ), for the antioxidant and neuroprotective activities against 6-OHDA and H₂O₂. Particularly, it displayed a strong neuroprotective effect against all the used concentrations of 6-OHDA, reporting the values similar to those of the control. Clearly, proof of the concept will involve an investigation of its *in vivo* neuroprotective profile.

Although much needs to be understood in terms of pathogenesis of neurodegeneration, it is clear that the simultaneous modulation of oxidative stress and metal brain levels can be considered as a potential strategy to interrupt the vicious cycle that accelerates the progression of the neurodegenerative diseases, such as PD and AD.

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Conflict of Interest

The authors declare no conflict of interest.

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