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Through scaffold modification to 3,5-diaryl-4,5-dihydroisoxazoles: new potent and selective inhibitors of monoamine oxidase B

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ABSTRACT

3,5-Diaryl-4,5-dihydroisoxazoles were synthesized and evaluated as monoamine oxidase (MAO) enzyme inhibitors and iron chelators. All compounds exhibited selective inhibitory activity towards the B isoform of MAO in the nanomolar concentration range. The best performing compound was preliminarily evaluated for its ability to bind iron II and III cations, indicating that neither iron II nor iron III is coordinated. The best compounds racemic mixtures were separated and single enantiomers inhibitory activity evaluated. Furthermore, none of the synthesised compounds exhibited activity towards MAO A. Overall, these data support our hypothesis that 3,5-diaryl-4,5-dihydroisoxazoles are promising scaffolds for the design of neuroprotective agents.

ARTICLE HISTORY

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KEYWORDS

3,5-diaryl-dihydroisoxazoles; MAO B selective inhibitors; neuroprotective agents

Introduction

Amine oxidase is a widely diffused family of enzymes responsible for the oxidative deamination of both endogenous and exogenous monoamines, diamines and polyamines. These enzymes are divided into two classes, flavin adenine dinucleotide amino oxidase (FAD-AO) (EC 1.4.3.4)¹ and copper topaquinone amino oxidase (Cu/TPQ-AO)². The former, named monoamine oxidase (MAO), contains flavin adenin dinucleotide (FAD) as a cofactor while the latter contains 2,4,5-trihydroxyphenylalanine quinone as cofactor (TPQ-Cu) and are inhibited by semicarbazide. Regarding FAD-AO, two isoforms, MAO-A and MAO-B, have been isolated, differing from central nervous system (CNS) location, substrate specificity and sensitivity to inhibitors^{3,4}. MAO enzymes⁵ play a key role in the metabolism of monoamine neurotransmitters, and have been therefore studied in a number of psychiatric and neurological diseases⁶. Moreover, the MAO catalysed oxidative deamination of amines leads to the formation of hydrogen peroxide which is co-responsible for neurodegeneration⁷. Not surprisingly a number of MAO inhibitors have been investigated as neuroprotective agents in the treatment of Parkinson and Alzheimer diseases^{3,4,6,8–10}. We have previously identified highly potent and selective MAO B inhibitors characterised by a 3,5-diaryl-4,5-dihydro-(1H)-pyrazole scaffold¹¹ and investigated on the structural requirements for the enzyme inhibition (Figure 1). In particular, the presence of two aryl substituents on the dihydro-(1H)-pyrazole nucleus appeared as essential for MAO B selectivity and inhibition potency activity 12-15. Hence, we designed and studied 2-thiazolylhydrazones derivatives, ideally originated by removing C4 and C5 from the dihydropyrazole ring. These derivatives exhibited high potency and selectivity towards MAO B¹⁶⁻¹⁹. More recently, through the isosteric replacement of the carbon 4 of the previously reported dihydropyrazole derivatives¹¹ with an

oxygen atom, we have designed and synthesised two series of 2,3-dihydro-oxadiazoles^{20–22} that exhibited high selectivity and activity towards MAO B. Moreover, we observed that the introduction of a 3,4-dichlorophenyl substituent in the position 2 of the dihydro-oxadiazole core is crucial for both selectivity and potency²¹. According to these structural information and with the aim of setting new hit structures for MAO B inhibition, we have devised a synthetic pathway to obtain 3,5-diaryl-4,5-dihydroisoxazoles as bio-isosters of both the previously reported 3,5-diaryl-4,5-dihydro-(1*H*)-pyrazoles and 2,5-diaryl-2,3-dihydro-1,3,4-oxadiazoles.

Methods

Materials and apparatus

Starting materials and reagents were obtained from commercial suppliers and were used without purification. All melting points were determined on a Stuart SMP11 melting points apparatus and are uncorrected (East Yorkshire, UK). Electron ionization mass spectra were obtained by a Fisons QMD 1000 mass spectrometer (70 eV, 200 mA, ion source temperature 200 °C) (Loughborough, UK). Samples were directly introduced into the ion source. Found mass values are in agreement with theoretical ones. Melting points, yield of reactions and the analytical and descriptive data of derivatives are reported in Table 1.

¹H-NMR (Table 2) were registered on a Bruker AMX 300 MHz spectrometer (Billerica, MA). All samples were measured in CDCl₃. Chemical shifts are reported referenced to the solvent in which they were measured. Coupling constants *J* are expressed in Hz. Elemental analyses were obtained on a Perkin-Elmer 240 B microanalyser (Waltham, MA). Analytical data of the synthesised compounds are in agreement within ±0.4% of the theoretical values.

Potent and selective MAO B inhibitors

Figure 1. Scaffold evolution of MAO B inhibitors: from dihydro-pyrazoles to dihydro-isoxazoles.

Table 1. Chemical, analytical, and physical data of derivatives EMAC II (i-m).

			C_H_N				
Compound	R	Cryst. solvent	Calc.	Found	M.P. (°C)	Yield %	E.I. 70 eV mass spectra m/z (rel. ab.)
EMAC II i	4-Cl	Ethanol	70.72, 5.19, 5.15	70.67, 5.17, 5.12	134-5	42	271 (100); 240 (78); 139 (17)
EMAC II j	4-CH ₃	Ethanol	81.24, 6.82, 5.57	81.27, 6.83, 5.54	103-4	46	251 (100); 220 (83); 119 (13)
EMAC II k	4-F	Ethanol	75.28, 5.53, 5.49	75.31, 5.54, 5.43	132-4	41	255 (100); 224 (75); 123 (19)
EMAC II I	3,4-Cl	Water/ethanol	62.76, 4.28, 4.57	62.72, 4.31, 4.54	134-5	65	305 (100); 274 (81); 173 (22)
EMAC II m	4-OCH₃	ethanol	76.38, 6.41, 5.24	76.33, 6.40, 5.22	131-3	44	267 (100); 236 (71); 135 (9)

Table 2. ¹H NMR data and main fragments in E.I. mass spectrometry of derivatives EMAC II (i-m).

Compound	1 H NMR δ (ppm)		
EMAC II i	¹ H-NMR: (300 MHz, CDCl ₃) δH 2.38 (s, 3H, CH ₃), 3.23 (dd, 1H, Jab 16.5, Jax 7.8, CH ₂ , isoxazole), 3.76 (dd, 1H, Jab 16.5, Jbx 10.1, CH ₂ , isoxazole), 5.59 (dd, 1H, Jbx 10.8, Jax 8.1, CH, isoxazole), 7.21 (d, 2H, J 7.8, CH, arom.), 7.33 (m, 4H, CH, arom.), 7.57 (2H, d, J 8.1, CH, arom.).		
EMAC II j	¹ H-NMR: (300 MHz, CDCl ₃) &H 2.37 (s, 3H, CH ₃), 2.43 (s, 3H, CH ₃), 3.30 (dd, 1H, Jab 16.5, Jax 8.3, CH ₂ , isoxazole), 3.72 (dd, 1H, Jab 16.5, Jbx 10.8, CH ₂ , isoxazole), 5.66 (dd, 1H, Jbx 10.8, Jax 8.3, CH, isoxazole), 7.16–7.29 (m, 6H, CH, arom.), 7.57 (d, 2H, J 8.3, CH, arom.).		
EMAC II k	¹ H-NMR: (300 MHz, CDCl ₃) δH 2.35 (s, 3H, CH ₃), 3.27 (dd, 1H, Jab 16.7, Jax 8.0, CH ₂ , isoxazole), 3.74 (dd, 1H, Jab 16.5, Jbx 10.7, CH ₂ , isoxazole), 5.69 (dd, 1H, Jbx 10.7, Jax 8.3, CH, isoxazole), 7.04 (t, 2H, J 8.3, CH, arom.), 7.21 (d, 2H, J 8.1, CH, 4-CH ₃ -phenyl), 7.36 (dd, 2H, CH, J 8.3/5.3, arom.), 7.56 (d, 2H, J 8.2, CH, 4-CH ₃ -phenyl).		
EMAC II I	¹ H-NMR: (300 MHz, CDCl ₃) δH 2.38 (s, 3H, CH ₃), 3.27 (dd, 1H, Jab 16.7, Jax 7.8, CH ₂ , isoxazole), 3.78 (dd, 1H, Jab 16.4, Jbx 10.8, CH ₂ , isoxazole), 5.67 (dd, 1H, Jbx 10.8, Jax 7.8, CH, isoxazole), 7.21 (m, 3H, CH, arom.), 7.44 (d, 1H, J 8.2, CH, arom.), m, 7.49 (d, 1H, J 1.2, CH, arom.), 7.56 (d, 2H, J 8.1, CH, arom.).		
EMAC II m	¹ H-NMR: (300 MHz, CDCl ₃) δH 2.38 (s, 3H, CH ₃), 3.30 (dd, 1H, Jab 16.7, Jax 7.8, CH ₂ , isoxazole), 3.71 (dd, 1H, Jab 16.5, Jbx 10.6, CH ₂ , isoxazole), 3.80 (s, 3H, OCH ₃), 5.66 (dd, 1H, Jbx 10.5, Jax 8.6, CH, isoxazole), 6.89 (d, 2H, J 8.3, CH, arom.), 7.21 (d, 2H, J 8.2, CH, 4-CH ₃ -phenyl), 7.31 (d, 2H, J 8.7, CH, arom.), 7.58 (d, 2H, J 8.2, CH, 4-CH ₃ -phenyl).		

TLC chromatography was performed using silica gel plates (Merck F 254, Kenilworth, NJ), spots were visualised by UV light.

General procedure for the synthesis of compound derivatives

In a typical experiment, 0.0038 mol of the appropriate α , β -unsaturated ketone is dissolved in 30 ml of boiling absolute ethanol. On this solution, 10 ml of a freshly prepared

hydroxylamine hydrochloride (0.0057 mol) and potassium hydroxide (0.013 mol) water solution are added dropwise. The solution becomes progressively orange and the reaction is monitored by thin layer chromatography (TLC) (CH₂Cl₂/n-hexane: 20/1). By cooling the reaction, the formation of a precipitate is observed which was filtered and crystallised from ethanol.

By this procedure (Scheme 1), compounds EMAC II (i-m) were synthesised.

Scheme 1. Synthesis of EMAC II (i-m) compounds. Reagents and conditions: (i) ethanol, NaOH; (ii) ethanol, hydroxylamine hydrochloride, potassium hydroxide, reflux.

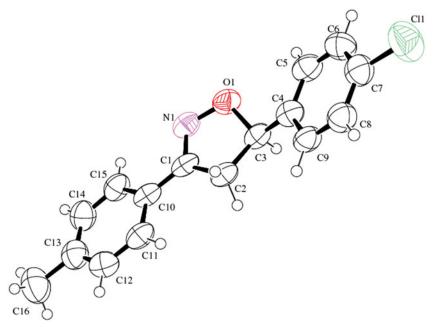


Figure 2. An ORTEP view of the molecular structure of (R)-(-)-EMAC II i enantiomer.

HPLC enantioseparations and stereochemical characterisation

Direct separation of the enantiomers of **EMAC II** (i-m) was achieved by HPLC using pure dichloromethane as the eluent. The resolving power of the immobilized-type was sufficiently high to achieve a baseline enantioseparation of all compounds in a short-time analysis. The optimized analytical conditions were easily scaled up to semipreparative level employing a 1 cm i.d. IA column. An amount of <20 mg of racemic samples was resolved for each chromatographic run, and both enantiomers were collected with high enantiomeric purity. The stereochemical characterization of compounds **EMAC II** (i-m) was performed by CD correlation comparing the maximum and minimum of ellitipticity of the CD spectra of the isolated enantiomers of compound **EMAC II** i of known stereochemistry (Figure 2), as previously described by some of us²³. Single enantiomers were resubmitted to biological assay.

Biological assay

The inhibitory activities of compounds **EMAC II (i-m)** on human recombinant MAO A and B isoforms, expressed in baculovirus infected BTI infected cells, as IC₅₀, are reported in Table 1. Tested compounds demonstrated no interference with the measurements,

since they were unable to directly react with the Amplex Red reagent. The kinetic parameters of hMAO-A and hMAO-B were evaluated in the presence of different tyramine concentrations. In our experiments, hMAO-A displayed a Michaelis constant (Km) of 514 ± 46.8 mM and a maximum reaction velocity (Vmax) of 301.4 ± 27.9 nmol/min/mg protein, whereas hMAO-B showed a Km of $104.7 \pm 16.3 \,\text{mM}$ and a Vmax of $28.9 \pm 6.3 \,\text{nmol/min/mg}$ protein (n = 5). Active compounds showed reversible behaviour according to the method proposed by Cer et al.²⁴ Hence, reported IC₅₀ is a useful tool to determine the relative activity of the compounds within the series as well as to determine substituent and key positions in the scaffold. All the tested compounds exhibited inhibitory activity towards hMAO-B at nM concentration. Single enantiomers activity was evaluated indicating that stereochemistry plays a relevant role in determining inhibitory concentrations. None of the tested compounds exhibited activity towards the A isoform of the enzyme at the tested concentration of 100 μM.

Coordination studies

The interaction between **EMAC II I** and the Fe²⁺ or Fe³⁺ ions was evaluated by recording the UV/VIS absorbance spectra of **EMAC II** I (Figure 3) in absence and in presence either of FeCl₃ or FeSO₄. **EMAC II I** was solubilised in DMSO then diluted to 1 mM in

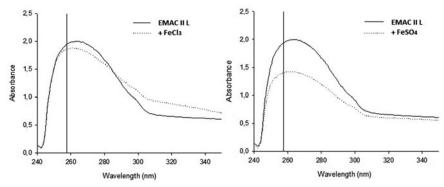


Figure 3. Effect of Fe^{2+} or Fe^{3+} ions on the spectrum of absorbance of EMAC II I. UV/vis spectrum was measured with 100 μ M of compound alone (unbroken line) or in the presence of 10 mM FeCl₃ or FeSO₄ (dotted line).

Table 3 Inhibitory activities towards hMAO-A and hMAO-B of FMAC II (i-m) derivatives

Compound	Structure	MAO-A (IC ₅₀)	MAO-B (IC ₅₀)	Ratio ^e
EMAC II i	H ₃ C CI	b	104.04 ± 3.69 nM	>961 ^d
EMAC II j	N-O H ₃ C CH ₃	b	41.05 ± 1.52 nM	>2436 ^d
EMAC II k	Ñ−ó H₃C F	b	320.22 ± 13.61 nM	>312 ^d
EMAC II I	N-O H₃C CI	b	$11.97 \pm 0.37 \text{nM}$	>8354 ^d
EMAC II m	H ₃ C OCH ₃	b	449.57 ± 18.02 nM	>222 ^d
Clorgiline L-Deprenyl Iproniazide	N=0	4.46 ± 0.32 nM ^c 67.25 ± 1.02 μM ^c 6.56 ± 0.76 μM	61.35 ± 1.13 μM 19.60 ± 0.86 nM 7.54 ± 0.36 μM	0.000073 3431.12 0.87
Moclobemide		361.38 ± 19.37 μM	7.54 ± 0.50 μW	<0.36 ^e

^aInactive at 1 mM (highest concentration tested).

50%DMSO solution. The UV/vis spectrum of the solution was recorded before and after addition of 10 mM final concentration either of FeCl₃ or FeSO₄. The spectra were recorded with a Ultrospec 2100 pro (Amersham Biosciences, Little Chalfont, UK) and analysed with SWIFT II-METHOD software (GE Healthcare, Little Chalfont, UK).

Molecular modelling studies

Protein preparation. Three-dimensional coordinates of the receptor²⁵ were obtained from the Protein Data Bank (PDB).²⁶ The protein was processed and the internal hydrogen bonding network of the receptor was optimized using the algorithm implemented in Protein Preparation wizard.²⁷

Ligand preparation. The ligands were built using Maestro GUI²⁷. The lowest energy conformer was considered for the subsequent computational studies. This was obtained with MacroModel version 7.2²⁸, considering MMFFs²⁹ as force field and solvent effects by adopting the implicit solvation model Generalized Born/Surface Area (GB/SA) water³⁰. The simulation was performed allowing 1000 steps Monte Carlo analysis with Polak-Ribier Conjugate

Gradient (PRCG) method and a convergence criterion of 0.05 kcal/ (mol Å).

Docking experiments. Previous reported and validated QMPL docking protocol was applied using default settings^{21,31}. In order to better take into account the induced fit phenomena, the most energy favoured generated complexes were fully optimized with the AMBER* united atoms force field³² in GB/SA implicit water,³⁰ setting 5000 steps iterations analysis with PRCG method and a convergence criterion of 0.1 kcal/(mol Å). Analysis of the results of the complex minimization was carried out taking into account the state equations (free energy of complex formation) computed at 300 K, applying molecular mechanics and continuum solvation models with the molecular mechanics generalized Born³³. The resulting complexes were considered for the binding modes graphical analysis with Pymol³⁴ and Maestro²⁷.

Results and discussion

Pursuing in our research on the design and synthesis of new agents for the selective inhibition of MAO B, we have synthesised a series of 3,5-diaryl-4,5-dihydro-isoxazoles EMAC II (i-m). All the

 $^{^{}b}$ Inactive at 100 μM (highest concentration tested). At higher concentration, the compounds precipitate.

^cResults are mean \pm SEM from five experiments. Level of statistical significance: ^cP < 0.01 versus the corresponding IC₅₀ values obtained against MAO-B, as determined by ANOVA/Dunnett's.

 $^{^{}m d}$ Values obtained under the assumption that the corresponding IC₅₀ against MAO-A is the highest concentration tested (100 μ M).

^eSelectivity ratios [IC₅₀(MAO-A)]/[IC₅₀(MAO-B)].

compounds were submitted to biological evaluation to investigate their ability to inhibit both MAO isoforms. The structure and the MAO A and MAO B IC₅₀ values and selectivity ratios of the new derivatives are reported in Table 3. All the compounds inhibit MAO B isozyme with IC50 values ranging from higher to lower nanomolar range. Accordingly to what is observed in our previous report on similar derivatives²¹, compound EMAC II I, bearing a 3,4-dichlorophenyl moiety in the position 5 of the dihydroisoxazole ring resulted, the most active within all the tested compounds. Based on this observation we assumed that, in the formation of the enzyme-ligand complex, this portion of the inhibitor is, in all probability, oriented similarly with respect to our previously reported compounds. Interestingly, none of the new compounds exhibited activity towards the MAO A isoform of the

Table 4. Inhibitory activities towards hMAO-A and hMAO-B of EMAC II (i-m) sinale enantiomers.

Compound	MAO-A (IC ₅₀)	MAO-B (IC ₅₀)	Ratio
R-(—) EMAC II i	a	15.27 ± 0.95 nM	6549 ^b
S-(+) EMAC II i	a	a	
R-(-) EMAC II j	a	$2.55 \pm 0.11 \text{nM}$	39 216 ^b
S-(+) EMAC II j	a	$31.37 \pm 1.67 \mu\text{M}$	3188 ^b
R-(—) EMAC II k	a	299.86 ± 12.04 nM	333.5 ^b
S-(+) EMAC II k	a	$33.82 \pm 1.12 \mu\text{M}$	2956 ^b
R-(-) EMAC II I	a	1.89 ± 0.16 nM	52 910 ^b
S-(+) EMAC II I	a	$10.96 \pm 0.83 \mu M$	9124
R-(-) EMAC II m	a	126.16 ± 8.44 nM	792.6 ^b
S-(+) EMAC II m	a	291.82 ± 13.45 nM	342.6

 IC_{50} values are the mean \pm SEM from five experiments.

enzyme up to 100 µM concentration. Considering the presence of an asymmetric carbon at the position 5 of the dihydro-isoxazole nucleus we performed, semipreparative chromatographic enantioseparation of all the new derivatives EMAC II (i-m). The separated enantiomers were then submitted to biological evaluation (Table 4). The obtained data pointed out the importance of enantiomeric separation for both potency and selectivity. In particular, in the case of compound R-(-)-EMAC II I, the IC₅₀ value towards MAO B, drops from 11.97 nM to 1.89 nM, while the selectivity towards MAO B is increased by more than six folds. Moreover, considering the growing interest in the development of iron chelators for the treatment of neurodegenerative disorders^{35–37}, we have preliminarily investigated our best compound able to bind iron II and iron III cations following a previously reported procedure³⁸. According to our preliminary results (Figure 3), compound EMAC II I do not have the ability to bind neither iron II nor iron III cations.

In order to explore the nature of the ligand-receptor interactions, we have carried out docking experiments considering the MAO B active site by means of QMPL docking protocol^{31,39,40}. We have focused our attention on the most active compound R-(–)-EMAC II I, which shows the lowest IC_{50} value and the highest selectivity towards MAO-B.

It should be pointed out that MAO-B cavity is rather narrow, therefore, small changes are reflected in the biological activity. In this respect, the best compound shows remarkable difference of activity between the R and the S enantiomers. An analogous behaviour was observed in the previous series 11,20. According to docking and post docking best scored pose, the most stable complex configurations of both R and S enantiomers are depicted in Figure 4(a,b).

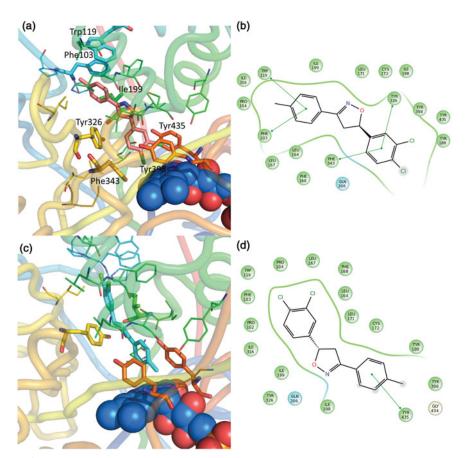


Figure 4. Putative binding mode of compound EMAC II I. (a) R-(-)-EMAC II I/MAO-B complex; (c) S-(+)-EMAC II I/MAO-B complex; (b, d) compound 2D representation and binding pocket interacting residues: green, hydrophobic; cyan, polar residues. Green arrows indicate $\pi-\pi$ interactions.

^alnactive at 100 μM (highest concentration tested).

^bValues obtained under the assumption that the corresponding IC_{50} against MAO-B is the highest concentration tested (100 μ M).

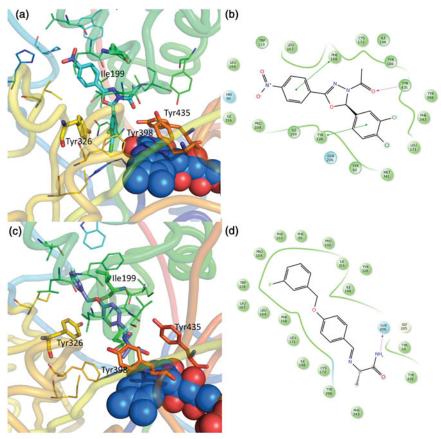


Figure 5. Binding mode of previously published compounds and approved inhibitors. (a) 4f-(R) MAO-B complex²¹; (c) safinamide MAO-B complex (pdb code 4v5z)²⁵; (b, d) compound 2D representation and binding pocket interacting residues: in green, hydrophobic; cyan, polar residues. Green arrows indicate $\pi-\pi$ interactions, magenta arrow indicates H bonds.

The *R* enantiomer is stabilized by $\pi - \pi$ interactions with Trp119, Phe103, Phe343, Tyr326 and some hydrophobic interactions with several residues in the binding pocket: Leu 164, Leu 167, Phe 168, lle 316, lle 199, lle198, Leu 171 and the Tyr of the aromatic cage 435, 298, 188.

Conversely, the S enantiomer is oriented in the opposite way: with the dichlorophenyl substituent in the entrance cavity and the 4-methylphenyl substituent in the catalytic cavity toward the FAD. The complex results in less stability and gives rise only to a single $\pi - \pi$ interaction with Tyr 435 and several hydrophobic interactions (Figure 4(c,d)).

Compared with previous series²¹, the absence of acetyl group allows a better accommodation in the cavity (Figures 4(a) and 5(a)). Once again, the importance in crossing both entrance and catalytic cavity, in order to achieve the MAO-B isoform selective activity is confirmed. This essential requisite has been reported by several studies^{25,41,42} and can be observed in co-crystallized selective MAO B inhibitors. A typical example is the inhibitor safinamide $(Figure 5(c,d))^{25,43}$

Conclusions

In summary, a small library of 3,5-diaryl-4,5-dihydroisoxazoles have been designed and synthesized on the basis of previously reported MAO inhibitors. Our finding highlighted the capability of 3,5-diaryl-4,5-dihydroisoxazoles to exert specific recognition and inhibition against the isoform B of MAO. The stereochemical characterization of R/S enantiomers confirms an enantioselective effect against MAO B isoform. This scaffold is definitively interesting for the development of novel selective neuroprotective agents.

Disclosure statement

The authors have no declarations of interest to report.

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