Isoxazole derivatives as new nitric oxide elicitors in plants

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Abstract

Several 3,5-disubstituted isoxazoles were obtained in good yields by regiospecific 1,3-dipolar cycloaddition reactions between aromatic nitrile oxides, generated in situ from the corresponding hydroxyimidoyl chlorides, with non-symmetrical activated alkynes in the presence of catalytic amounts of copper(I) iodide. Effects of 3,5-disubstituted isoxazoles on nitric oxide and reactive oxygen species generation in Arabidopsis tissues was studied using specific diaminofluoresceine dyes as fluorescence indicators.

Introduction

Isoxazoles are an interesting class of N-heterocyclic compounds intensely studied mainly due to their wide range of biological activity [1,2]. Isoxazole compounds show antiviral [3,4], antithrombotic [5-9], analgesic [9], COX-2 inhibitory [10,11], anti-inflammatory [9,11], antinociceptive [12] and anticancer [13] activities. Several isoxazole derivatives have GABA_A antagonist [14] and T-type Ca²⁺ channel blocking activities [15]. Commercial drugs featuring an isoxazole moiety include the COX-2 inhibitor Valdecoxib and the β-lactam antibiotics Cloxacillin andDicloxacillin. An isoxazole derivative, namely
3,5-difluorophenyl-[3-methyl-4-(methylsulfonyl)]isoxazol-5-yl)methanone, was recently reported as an inducer of nitric oxide producing elicitor in plants [16,17]. Nitric oxide (NO), which has been demonstrated to be a major gasotransmitter in mammals, is also involved in the orchestration of various plant physiological responses, playing an important role in the regulation of interactions between plant and microorganisms and in plant defense mechanisms against stresses [18,19]. Consequently, there is interest in the biological evaluation of further isoxazole derivatives.

Many synthetic approaches towards the isoxazole core include the reactions of hydroxylamine with aryl-[β]-diketones [20], α,β-unsaturated carbonyl compounds [21], or α,β-unsaturated nitriles [22], and 1,3-dipolar cycloaddition reactions between alkenes or alkynes and nitrile oxides [23-25].

Nitrile oxides are known as reactive 1,3-dipoles involved in 1,3-dipolar cycloaddition reactions with various dipolarophiles generating five-membered heterocyclic compounds, such as isoxazoles, isoxazolines, oxadiazoles, oxadiazolines, dioxazolidines etc. [23-25]. Intermediate nitrile oxides are usually generated in situ by the oxidative dehydrogenation of aldoximes in the presence of various oxidants [26-29], or by the dehydrohalogenation of hydroxymoyl halides promoted by organic or inorganic bases [30-32]. A less used synthetic procedure involves the oxidative dehydration of primary nitro compounds with isocyanates in the presence of tertiary alkylamines [33].

Generally, the cycloaddition reactions of nitrile oxides to alkenes yield isoxazolines or a mixture of isoxalines and isoxazoles. Cycloaddition reactions of nitrile oxides to alkynes yield isoxazoles directly, without a catalyst, but the yields of isoxazole products are quite low because of side reactions and both regioisomers are generally obtained [23-25]. The one-pot 1,3-dipolar cycloaddition reaction of a nitrile oxide, generated in situ from the corresponding hydroxymoyl chloride, with an in situ brominated electron-deficient alkeno led to the intermediate bromoisoxazoline from which, by loss of HBr, a 3,5-disubstituted isoxazol derivative is formed as major regioisomer [34]. Based on the copper(I)-catalyzed click reactions of organic azides with terminal acetylenes [35], different copper(I)-catalyzed synthetic procedures towards isoxazole derivatives were developed [36,37].

As part of our continued efforts to develop simple synthetic routes towards bioactive heterocyclic compounds [38-43], we report here the synthesis of several 3,5-isoxazole derivatives, bearing benzo[1,3]dioxole and thiophene scaffolds respectively, as well as their inductor effect on the generation of nitric oxide and reactive oxygen species in plant tissues. The benzo[1,3]dioxole framework is a constituent of some fragrances and flavors [44], and several bioactive compounds with a broad spectrum of applications [45-48]. The thiophene is a core system of a large number of bioactive molecules such as antineoplastic agents [49], non-steroidal anti-inflammatory drugs [50] or compounds with antibacterial activities against several Gram-positive strains [51]. Thus, 3,5-disubstituted isoxazole derivatives were obtained by regioselective 1,3-dipolar cycloaddition reactions of aromatic nitrile oxides to non-symmetrically activated alkyne derivatives in the presence of catalytic amounts of copper(I) iodide.

Results and Discussion
Synthesis of 3,5-disubstituted isoxazole derivatives
The regioselective copper(I)-catalyzed 1,3-dipolar cycloaddition reactions of aromatic nitrile oxides, generated in situ from the corresponding crude imidoyl chlorides, and non-symmetrically activated alkyne led to 3,5-disubstituted isoxazole derivatives. For this, aromatic aldehydes 1 are first converted to the corresponding aldoximes 2, via reactions with hydroxylamine, and the crude reaction products are transformed to the corresponding imidoyl chlorides 3 which are directly used in the next step without purifications. Catalytic amounts of copper(I) iodide and a base, such as KHCO₃, are added to an aqueous solution containing the crude imidoyl chlorides 3 and the non-symmetrically activated alkyne 4. The in situ generated aromatic nitrile oxides 5 undergo an addition to copper(I) acetylides formed in situ to give the 3,5-disubstituted isoxazoles 6-11 as single isomers, in moderate to good yields (Scheme 1).

All reactions between crude imidoyl chlorides 3 and the non-symmetrical activated alkynes 4 are carried out in aqueous solutions at room temperature. The final 3,5-disubstituted isoxazoles are easily separated by filtration, washed with water and recrystallised. The synthesized 3,5-disubstituted isoxazoles are presented in Table 1. 3,5-Disubstituted isoxazoles 6, 7, 9 and 10 (Table 1) have been previously prepared by different synthetic procedures, but compounds 6 [52], 7 [49] and 10 [53] have been only partly characterized, while for compound 9 [54] no characterization data have been reported.

3,5-Disubstituted isoxazole structures were unambiguously assigned on the basis of chemical and spectral analysis (IR, 1H, 13C and 15N NMR spectra). NMR spectra clearly indicated the presence of only one regioisomer for all synthesized 3,5-disubstituted isoxazoles. Signals in 1H and 13C NMR spectra were fully assigned based on H,C-HSQC and H,C-HMBC experiments. The 3,5-disubstitution was also experimentally proven.
by NOE experiments which indicated through space interaction between the H-4 proton and the protons from both 3- and 5-substituents.

**Biological activity**

We investigated the inductor effects of 3,5-disubstituted isoxazoles 6–11 on NO and reactive oxygen species (ROS) production in plant tissues. Usually, NO and ROS, such as O$_2^-$, OH· and H$_2$O$_2$, together are required to induce the activation of various defense-related enzymes in plants [55]. Plant cells contain oxygen radical detoxifying enzymes and nonenzymatic antioxidants which have an essential role in protection of plant cells from oxidative damages at the sites of ROS and NO generation [56,57]. Measuring the ROS and NO levels in plant tissues is often difficult due to high reactivity and extremely short physiological half-life of these free radicals [58,59].

In this work, generation of both NO and ROS was proven by fluorescence microscopy using specific fluorescence indicators that help to exactly define the sites of generation. We used *Arabidopsis thaliana* wild type seeds, cultivated for six weeks in laboratory in Arasystem [60]. *Arabidopsis thaliana* was selected as model organisms for NO inductors because this is the flowering plant with the largest amount of knowledge on cellular and molecular biology and it has a relatively short life cycle. The *Arabidopsis* leaves were sprayed with fine suspensions of isoxazole inductors 6–11 at two different concentrations (10 μg/mL and 50 μg/mL, respectively) and collected after 24 h. Collected leaves were washed with distilled water and incubated with the specific fluorescence indicator for histochemical analysis of ROS and NO by fluorescence microscopy. *Arabidopsis* leaves untreated with inductor suspensions have been used as negative controls. As positive control, we used chitosan (β-1,4 linked glucosamine) with average molecular weight, a fungal elicitor with known effect as NO and ROS inductor on *A. thaliana* [61].

Intracellular ROS was visualized using 2′,7′-dichlorodihydrofluorescein diacetate (H$_2$DCF DA) as fluorescence indicator. The method is based on the oxidation of the non-fluorescent probe of 2′,7′-dichlorodihydrofluorescein diacetate to the highly fluorescent 2′,7′-dichlorofluorescein diacetate [62,63]. Intracellular NO was visualized using 4-amino-5-methylamino-2′,7′-dichlorodihydrofluorescein diacetate (DAF-FM diacetate), a non-fluorescent compound, which reacts with NO to form a
fluorescent benzotriazole and does not react with any ROS [64-67].

Fluorescence microscopy images of all 3,5-disubstituted isoxazoles-treated Arabidopsis leaves showed a pronounced presence of ROS at both concentrations of inductors (10 μg/mL and 50 μg/mL). Strong fluorescence densities were observed especially at higher concentration (50 μg/mL) of 3,5-disubstituted isoxazoles. Intensities of fluorescence revealed differences between tested compounds. These compounds can be listed in the increasing order of ROS generation efficacy as follows: 11 > 10 > 9 > 7 > 8 > 6 (Figure S2 in Supporting Information File 1).

Similarly, images of fluorescence microscopy revealed the presence of NO in all 3,5-disubstituted isoxazoles-treated Arabidopsis leaves, at both concentrations (10 μg/mL and 50 μg/mL), especially at higher concentration of compounds (50 μg/mL). The NO releasing capacity of newly synthesized 50 μg/mL, especially at higher concentration of compounds μg/mL and leaves, at both concentrations (10 mg/mL). The NO releasing capacity of newly synthesized 50 μg/mL, especially at higher concentration of compounds μg/mL). Strong fluorescence densities were observed especially at higher concentration (50 μg/mL) of 3,5-disubstituted isoxazoles followed the series: 9 > 11 > 10 > 8 > 7 = 6 (Figure S3 in Supporting Information File 1).

Fluorescence data indicate that the 3,5-disubstituted isoxazoles 6–11, particularly 9–11, are involved in NO and ROS production in Arabidopsis treated leaves.

Conclusion
Several 3,5-disubstituted isoxazoles were obtained by the convenient, regiospecific 1,3-dipolar cycloaddition reactions of aromatic nitrile oxides, generated in situ from the crude imidoyl chlorides, with non-symmetrical activated alkynes in the presence of catalytic amounts of copper(I) iodide. The effect of 3,5-disubstituted isoxazoles in generation of ROS and NO in plant tissues was investigated by fluorescence microscopy. The obtained data indicate that some of these compounds are chemical elicitors that induce NO and ROS generation in plant tissues and could activate various defense mechanisms in plants. Further research is in progress to assess the in planta mechanism of NO generation by these compounds.

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References

Supporting Information
Supporting Information File 1
Experimental procedures, characterization data, IR, 1H, 13C and 15N NMR data for all new compounds and fluorescence microscopy images of NO and ROS generation for all 3,5-disubstituted isoxazoles-treated Arabidopsis leaves.
[http://www.beilstein-journals.org/bjoc/content/supplementary/1860-5397-13-65-S1.pdf]


63. Chen, Y.; Mo, H.-Z.; Hu, L.-B.; Li, Y.-Q.; Chen, J.; Yang, L.-F. 

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