Synthesis of 3(2)-phosphonylated thiazolo[3,2-α]oxopyrimidines

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Abstract
A series of 3(2)-phosphonylated thiazolo[3,2-α]oxopyrimidines was synthesized for the first time by the reactions of chloroethynylphosphonates with unsubstituted and 5(6)-substituted 2-thiouracils. The reaction of chloroethynylphosphonates with 6-substituted 2-thiouracils bearing electron-donor groups (CH₃, Ph) proceeded with high regioselectivity involving the cyclization through the N₃-nitrogen atom to form new 3-phosphonylated thiazolo[3,2-α]-5-oxopyrimidines with good yield. In the case of unsubstituted and 5-methyl-2-thiouracils, cyclization occurred predominantly through the N₁ atom and partially via the N₃-nitrogen atom to form a mixture of the corresponding thiazolo[3,2-α]-7- and 5-oxopyrimidines. A dramatic change in the reaction regioselectivity was observed in the case of 6-trifluoromethyl-2-thiouracil that afforded 2- and 3-phosphonylated 5-oxothiazolopyrimidines in a 1:1 ratio.

Introduction
Thiazolopyrimidines, whose molecules includes both thiazole and pyrimidine rings, have a structural analogy with the antipsychotic drugs ritanserin and setoperone (Figure 1) [1-3]. To date, a wide spectrum of biological activity of thiazolopyrimidines has been determined: anticancer [4,5], antimicrobial [6,7], anti-inflammatory [8,9], and antiviral [10,11].

The best known methods for the preparation of thiazolopyrimidines are based on condensation reactions. The most commonly used synthesis is the three-component condensation of 2-aminothiazoline, aromatic aldehyde, and ethyl cyanoacetate, which leads to the formation of 5- and 7-oxothiazolopyrimidine-6-carbonitriles (Scheme 1) [12,13].
The synthesis of thiazolopyrimidines through the reaction of 2-aminothiazoles with 1,3-ketoesters in the presence of acids, bases or condensing agents (Scheme 2) has been studied fairly well [6,8,14-16].

The most accessible approach to the synthesis of 5H-thiazolo[3,2-a]pyrimidine-5(7)-ones is the reaction of 2-thiouracil derivatives with α-halo ketones and α-halo acids, involving successive alkylation and condensation steps (Scheme 3) [17-21].

A convenient one-step synthesis of thiazolopyrimidine-5-ones by reacting 6-methyl-2-thiouracils with bromoethynylketones has been reported by Shishkin and co-workers (Scheme 4) [20]. The authors for the first time proved the structure of the 5-oxo isomer by single crystal X-ray diffraction analysis.

The Pd-catalyzed Sonogashira coupling reaction between 2-thiouracil and propargyl bromide yielded 5H-thiazolo[3,2-a]pyrimidine-5-one (Scheme 5) [20-24].

Despite the wide variety of thiazolopyrimidines reported to date, phosphorylated analogues of compounds of this series are unknown. Of special interest is the design of molecules containing practically significant heteroaromatic rings and a biologically active and hydrolysis-resistant phosphonate group, as it has

Figure 1: Structure of ritanserin and setoperone drugs.
Herein, we report the synthesis of a new series of phosphonylated thiazolopyrimidines. In our studies, chloroethynylphosphonate was used as the phosphonylating agent, which allowed the formation of a thiazole ring with simultaneous phosphonylation of the latter. As the second reaction component, available 2-thiouracils were chosen as the most studied objects used for creating thiazolopyrimidine systems. The main objective of the study was to determine the regioselectivity of the reaction.

Literature data mainly report the formation of 5-oxopyrimidines by cyclization through the N₃ atom of the starting 2-thiouracil [14-24]. The formation of 7-oxopyrimidines by cyclization through the N₁ atom has been noted only in a few reports [29-32]. However, reliable data for the identification of the 5- and 7-oxo isomers are not available to date and the determination of the structures of 5- and 7-oxo isomers were mainly based on ¹H NMR spectroscopy. The most comprehensive and convincing evidence for the formation of a 5-oxothiazolopyrimidine was provided by Shishkin and co-workers [20], who performed single crystal X-ray diffraction analysis along with ¹H and ¹³C NMR spectral studies. Unfortunately, the majority of reports on the synthesis of thiazolopyrimidines relied on ¹H NMR data to prove the structure of the obtained compounds [18,33-35]. There are no systematic data on ¹³C NMR spectroscopy of thiazolopyrimidines, which could be used as an additional approach to estimate the regioselectivity of the reaction.

A single example of the use of ¹³C NMR spectroscopy for unambiguous establishing the structure of thiazolo[3,2-a]pyrimidines obtained was given by Iranian researchers [36], but only for the 5-oxo isomers. In our opinion, the presence of a phosphorus fragment in a thiazolo[3,2-a]pyrimidine molecule significantly facilitates the determination of the structure by means of ¹³C, ¹H, and ³¹P NMR spectroscopy methods.

Results and Discussion

Aiming to synthesize a new series of phosphonylated thiazolopyrimidines, we performed reactions of unsubstituted and substituted 2-thiouracils 1a–e with chloroethynylphosphonates 2a–c. We found that the reaction with 6-substituted 2-thiouracil bearing either methyl or phenyl groups occurred regioselectively with a N₃ atom ring-closure to afford the 3-phosphonylated thiazolo[3,2-a]-5-oxopyrimidines 3a–f in good yields (Scheme 6). Likely, in this case the attack by the N₃ atom was more favorable than by the N₁ atom due to the steric effect of the substituent in position 6. The reactions proceeded under mild conditions within 3–5 hours. Anhydrous K₂CO₃ was used as a base to neutralize HCl formed during the reaction. The need for the use of anhydrous solvents and reagents is caused by the possibility of the formation of byproducts, if any, due to hydrolysis as we have noted earlier in the case of the reactions of chloroethynylphosphonates with nitrogen-containing nucleophiles [37-39].

The assignment of the reaction product to the 5-oxo isomer was made based on ¹³C NMR spectral analysis: the carbon atoms of
Scheme 6: Reactions of 6-substituted 2-thiouracils 1a, b with chloroethynylphosphonates 2a–c.

\[ \begin{align*}
\text{HN} & \text{N} \\
\text{S} & \text{R} \\
\text{O} & \text{R} \\
\end{align*} \]
\[ \text{Cl} \]
\[ \text{OR} \]
\[ \text{K}_2\text{CO}_3 \]
\[ \text{CH}_3\text{CN, rt} \]
\[ \begin{align*}
\text{HN} & \text{N} \\
\text{S} & \text{R} \\
\text{O} & \text{R} \\
\end{align*} \]
\[ \text{P} \]
\[ \text{OR} \]
\[ \text{OR} \]

Scheme 7: Reaction of 5-methyl-2-thiouracil (1c) with chloroethynylphosphonates 2a–c.

The CH=CR fragment (R=CH₃, Ph) are represented by a strong signal at δ_C 101–105 ppm (CH=) and a weak signal at δ_C 157–158 ppm (=CR). These data coincide with those for 3-phenylthiazolo[3,2-a]-5-oxopyrimidine [40]. In addition, the structures of the phosphorylated thiazolopyrimidines 3a and 3d were unambiguously confirmed by single crystal X-ray diffraction data.

The presence of the CH₃ group at the position 5 of the thiouracil ring changes the reaction regioselectivity, as the main direction is cyclization through the N¹ nitrogen atom with the formation of 3-phosphonylated thiazolo[3,2-a]-7-oxopyrimidines 4a–c and 5-oxo regioisomers 5a–c in a ≈1:0.1–0.3 ratio with yields of 87–91% (Scheme 7).

The structure of thiazolopyrimidines 4a–c and 5a–c is difficult to establish from the ¹H and ¹³C NMR spectral data. The signals of the vinyl proton of the uracil moiety are represented by quartets (J_HH = 1.4 Hz) in the δ_H 8.1 ppm region (quartet at δ_H 6 ppm for 6-methyl-5-oxo isomer). In the ¹³C NMR spectra,
the signals of ethylene carbons of the uracil ring are presented at δC 131 and 122 ppm (δC 101 and 158 ppm for 5-oxopyrimidines 5a–c). These data are in accordance with values of the chemical shifts of carbon atoms and protons of the pyrimidine ring in 3-methylthiazolo[3,2-a]pyrimidine-7-one [29], 3-phenylthiazolo[3,2-a]pyrimidine-7-one [40], and 5-phenylthiazolo[3,2-a]pyridine-7-one [16].

A similar reaction outcome was observed when chloroethynylphosphonates 2a–c were reacted with unsubstituted 2-thiouracil (1d). The cyclization reaction also proceeded predominantly through the N1 atom to form the corresponding 3-phosphonylated thiazolo[3,2-a]-7-oxopyrimidines 6a–c together with a small amount of 5-oxo isomers 7a–c (Scheme 8). In addition, it should be noted that the regioselectivity of the reaction was higher when using diisopropyl 2-chloroethynylphosphonate (2c).

The assignment of the major reaction product to the 7-oxo isomers was made by help of 13C NMR spectroscopy. In the 13C NMR spectra of the thiazolopyrimidines 6a–c, the O=C–CH=CH fragment is observed by signals of equal intensity at δC 112–113 and 135–136 ppm. However, the unambiguous proof of the structure of thiazolopyrimidine-7-one 6b was obtained by single crystal X-ray diffraction analysis.

It is important to note, that the reaction of dimethyl 2-chloroethynylphosphonate (2a) with 2-thiouracil had some features. The reaction proceeded with the formation of a mixture of 7-oxo and 5-oxo isomers in a ≈ 1: 0.1–0.3 ratio. However, a decrease in the signal of the 7-oxo isomer was observed in the 31P NMR spectrum upon standing at room temperature for 24 hours. The analysis of the formed precipitate identified product 6aa, representing the monodealkylation product of the dimethylphosphonate group. A similar phenomenon has been reported earlier [41].

In the case of 6-trifluoromethyl-2-thiouracil (1e), a dramatic change in the reaction regioselectivity was observed. The cyclization took place with the formation of a mixture of the corre-
As in the case of the 6-substituted 2-thiouracils 1a and 1b, the presence of a trifluoromethyl group in position 6 favored an attack by the N₃-nitrogen atom, resulting in the formation of the 5-oxo isomer. The formation of the 2-phosphonylated thiazolopyrimidine-5-one could be explained by a primary attack of the electrophilic carbon atom bonded to the chlorine atom by the N₃ nitrogen atom of the uracil fragment followed by cyclization.

The mixture of phosphonylated 5-oxothiazolopyrimidines 8e and 9e was separated by column chromatography and the ¹H, ¹³C, and ³¹P NMR spectral data of the individual thiazolopyrimidine isomers did not allow convincing assignment. In the ¹H NMR spectra, the proton of the thiazole ring of both isomers is represented by a doublet signal in the low field region at δH 7.91 (JHP = 3.2 Hz) and 8.41 ppm (JHP = 7.5 Hz), whereas the corresponding proton of the uracil ring resonated as a singlet at δH 6.61 and 6.71 ppm, respectively. Note that in the case of the reported 3-(4-bromophenyl)-7-trifluoromethyl-5H-[1,3]thiazolo[3,2-a]pyrimidine-5-one [18] the signal of the uracil proton H⁶ appeared at δH 6.67 ppm.

The ¹³C NMR spectra of the isomers 8e and 9e differed only in the values of the carbon signals at the phosphorus atom, i.e., doublets at 119.78 (JHP = 209.5 Hz) and 130.24 ppm (JHP = 217.5 Hz). The remaining signals were completely identical. Finally, the X-ray diffraction analysis of the 2-phosphonylated thiazolopyrimidine 9e allowed us to uniquely determine its structure.

In our opinion, the unusual formation of the 2-phosphonylated thiazolopyrimidine can be explained by the electron-withdrawing effect of the trifluoromethyl group in the starting 2-thiouracil. In contrast to 6-methyl- or 6-phenyl-2-thiouracil, where the nucleophilicity is localized on the sulfur atom, the presence of the electron-withdrawing CF₃ group in 6-trifluoromethyl-2-thiouracil (1e) enhances the acidity of the N₃H hydrogen by direct conjugation to the carbonyl moiety. As a result, 2-thiouracil 1e acts as an ambident nucleophile. Thus, the attack of the carbon atom attached to the chlorine by the N₃ nitrogen atom is accompanied by the elimination of hydrogen chloride (Scheme 10). A further 5-endo-dig-type cyclization results in the formation of the 2-phosphonylated 5-oxopyrimidines 9a–c. The formation of the 3-phosphonylated 5-oxopyrimidines 8a–c is due to the implementation of the usual favorable direction with the attack of chloroethynylphosphonate by the sulfur atom [33,42-46].

**Conclusion**

In conclusion, a series of phosphorylated thiazolo[3,2-a]oxopyrimidines was first synthesized by reacting unsubstituted and substituted 2-thiouracils with chloroethynylphosphonates. The main regularities of this reaction were revealed. In the case of

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**Scheme 9:** Reaction of 6-trifluoromethyl-2-thiouracil (1e) with chloroethynylphosphonates 2a–c.
Scheme 10: A plausible mechanism of the reaction between 6-trifluoromethyl-2-thiouracil (1e) and chloroethynylphosphonates.

6-substituted 2-thiouracil the primary attack by the most favorable nucleophilic site C=S takes place with further cyclization through the N3 atom of 2-thiouracil to form 5-oxopyrimidines. When using unsubstituted and 5-substituted 2-thiouracils, cyclization occurs predominantly through the N1 atom of the uracil ring, leading to the formation of 7-oxopyrimidines.

Supporting Information
Supporting Information File 1
General experimental procedure, characterization data, and copies of NMR spectra.
[https://www.beilstein-journals.org/bjoc/content/supplementary/1860-5397-16-161-S1.pdf]

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