MESOIONIC SYDNONE: A REVIEW IN THEIR CHEMICAL AND BIOLOGICAL PROPERTIES

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ABSTRACT

Various literature sources have documented sydnones as important molecules with exclusive chemical properties and a wide spectrum of bioactivities. Sydnone can be defined as a five-membered pseudo-aromatic heterocyclic molecule. Classically, 1,2,3-oxadiazole forms the main skeleton of sydnone. The molecule has delocalized balanced positive and negative charges. The five annular atoms share the positive charge and the enolate-like exocyclic oxygen atom bears the negative charge. The hydrogen atom at the position C4 was proved to have acidic and nucleophilic functionalities making the sydnone ring reactive towards electrophilic reagents. These unique chemical features enable sydnones to interact with biomolecules resulting in important therapeutic effects like anticancer, anti-diabetic, antimicrobial, antioxidant and anti-inflammatory. Consequently, we aim from the current article to review the available chemical and pharmacological information on sydnone and its derivatives.

Keywords: Sydnone, Mesoionic, Heterocycles, Anticancer, Antimicrobial, Anti-inflammatory

INTRODUCTION

Sydnones are the most studied compounds amongst the mesoionic family due to their interesting structures, chemical properties, synthetic utility and biological activities. Many reports stated that a covalent structure is not sufficient to represent the sydnone molecule satisfactorily [1, 2]. However, 1,2,3-oxadiazolidine bearing a carbonyl function has recently been the major representative of sydnones because FTIR spectroscopy showed a carbonyl stretch frequency attached to C5 of the ring like in 4-acetyl-3-tolybsydnone which exhibited a strong band at 1763 cm⁻¹ [3]. X-ray analysis revealed a bond length of 1.196 Å which corresponds to an exocyclic C=O double bond [4]. Classically, the sydnone ring can be prepared from the cyclization of N-nitroso amino acids with acetic anhydride [5]. Later, many attempts were employed to improve the yield of cyclization by using a stronger dehydrating agent such as trifluoroacetic acid anhydride or thionyl chloride [6]. Since their first preparation, sydnones attracted the attention of medicinal chemists and pharmacologists to investigate their biological applications. Their distinguished chemical structure enables them to bind and deactivate a variety of biomolecules like DNA and enzymes. A vast range of therapeutic properties has been demonstrated including antimicrobial, anti-inflammatory, anti-cancer, antioxidant and anti-diabetic [7]. The present review demonstrates the important chemical and biological data on sydnones starting from their early discovery in 1935 until today.

Chemistry of sydnone

Definition of sydnone

The word sydnone was originated from the phrase “University of Sydney” where this class of compounds was first prepared by Earl and Mackney in 1935. They suggested the formation of fused three- and four-membered ring product (I) from the action of acetic anhydride on N-nitrosophenylglycine [5] which was later considered wrong by other chemists.

Firstly, a fused ring system is unlikely to be formed by a simple intramolecular rearrangement and would be a highly strained unstable structure due to the existence of a β-propiolactone group. Therefore, Baker and his collaborator omitted the bridge bond and recommended a partially aromatic five-membered ring (II and III) which was a hybrid of many zwitterionic forms [8]. Secondly, acid hydrolysis decomposes sydnone into hydrazine, carboxylic acids and carbon dioxide while hot aqueous sodium hydroxide can revert the sydnone into the starting N-nitroso compound. These two facts indicate that the bicyclic system proposed by Earl is improbable [9]. Thirdly, other researchers proved that acetic anhydride can convert the dextro-rotary N-nitroso-N-phenylalanine into the optically inactive N-phenyl-C-methylsydnone (IV). The loss of optical activity implies either racemization or a change in the hybridization of C4 from a chiral sp³ state into an achiral sp². The oxygen atom attached to C5 was proved to be in an enolate form due to the rapid formation of a mono-bromo derivative (V) in glacial acetic acid and bromine [10].

To put an end to the previous debate, Baker and Ollis extensively reviewed all proposed structures of sydnone and suggested more clear description as follows:

1. A single covalent structure from the preceding suggestions does not fairly describe sydnone. In other words, the sydnone molecule should be considered as a hybrid of bipolar and tetrapolar forms (VI-XI) whose contribution to the hybrid ring is not equal. Consequently, sydnones are described as mesoionic compounds.

2. Compared to cyclopentadienyl anion (XIIa, XIIb), tropylium cation (XIIIa, XIIIb), furan, pyrrole and pyridine, the sydnone ring...
has all requirements to develop aromatic properties. The exocyclic oxygen atom provides an electron to the ring to complete the sextet of $^2$ electrons. Moderately satisfied, they came up with the formula XIV to represent the sydnone structure.

![Diagram of the sydnone structure XIV](image)

After all, they defined sydnone as a mesionic compound consisting of a five-membered aromatic heterocyclic ring mainly 1,2,3-oxadiazole or a six-membered ring in some cases. The molecule is neutral and has a positive charge in common between the annular atoms balanced by a negative charge borne on an exocyclic atom(s). Even though there is no single polar or covalent structure for sydnone, the structure XIV is being used as a representative in the majority of literature up to date [1, 2].

**Synthesis of sydnone**

Primarily, sydnone was prepared by Earl and his colleague by the cyclodehydration effect of acetic anhydride on the N-nitroso derivatives of amino acids. They reported that the dissolution of N-nitroso-N-arylglucine in excess acetic anhydride at room temperature resulted, after 24 h, in a nitroso-free, crystalline and stable heterocyclic product which was later referred to as sydnone. The preparation of the N-nitroso intermediate was accomplished by the conventional nitrosation of the amino group of N-phenylglucine by the nitrous acid generated from the reaction of sodium nitrite and hydrochloric acid [5]. The N-nitrosation of N-phenylglucine in neutral conditions was described later by Applegate and Turnbull using isosamyl nitrite (IAN) in dimethoxymethane (DME) at room temperature (Scheme 1). They claimed that IAN was successfully used to prepare the N-nitroso derivative of N-(2-acetylphenyl)glucine with high yield compared to the acid-based method which led to the formation of C-nitrosoglycine [11].

![Scheme 1: Preparation of N-nitroso analogues in neutral conditions](image)

Later, Baker et al. deduced the mechanism of cyclization of the N-nitroso starting material by losing a water molecule which involves four steps as presented in Scheme 2. Firstly, a mixed anhydride intermediate XV will be formed from the effect of acetic anhydride on the free nitroso acid whose carbonyl group will evolve strong cationic intermediate XVII. Secondly, a nucleophilic attack of the nitroso oxygen on the acid carbonyl group will lead to ring closure (XVI). Thirdly, an acetate group is lost, and a double bond between the two nitrogen atoms is formed (XVII). Lastly, loss of proton and formation of enolic oxygen will produce the final sydnone product XVIII [6].

![Scheme 2: Mechanism of ring closure and sydnone formation](image)

However, Eade and Earl found that the preparation of some sydnone analogues such as nitro-containing sydnone took a long time up to 7-30 d at room temperature with low to moderate yields. They claimed that heating accelerated the formation of sydnone ring, even though it reduced the yield due to rapid hydrolysis of the product by the hot acidic reaction medium [9]. Therefore, Baker and his colleagues reported for the first time an instant and complete separation of N-arylsydnones in 90% yield when they used trifluoroacetic anhydride (TFGAA) as a dehydrating agent [6]. The later synthetic route had been successfully utilized to prepare some complicated sydnones such as N,N-polyaliphatic bis-sydnone at a yield of 70-80% [12]. Moreover, heat-labile sydnones such as 3-(2-methoxy carbonylphenyl) sydnone was prepared in a considerable yield of 75% within one hour using TFGA in dichloromethane at 5°C [13]. Many alternative reagents were also employed to prepare the sydnone system. In 1950, Baker et al. used thionyl chloride. They reported that the conversion of N-nitroso amino acids into sydnone took place within a few minutes using thionyl chloride in dry ether at room temperature giving a low yield of 28%. On the other hand, using thionyl chloride in a mixture of cold diene and pyridine resulted in an improved yield (75%) within 25 min [6].

Some special structures of sydnones were reported as unexpected products of the cyclodehydration of the N-nitroso derivatives of amidino dicarbonylic acids. For example, 4', 4'-methylenedioxy sydnone was prepared in a yield of 80-88% using the previous reagents compared to 5-30% by the classic Earl’s method [9, 16, 17].

![Diagram of sydnone structures](image)

Physicochemical properties of sydnone

The electronic structure of sydnone ring

Using a new method of molecular orbital calculation named as $\alpha$-technique modified from Hückel framework, Kier and Roche calculated...
The properties of the ultraviolet spectra of sydnones were well studied due to conjugation as in 3-phenylsydnone and 3-(1-naphthyl) sydnone which absorb at the lower wavelength (<300 nm). For example, 3-acetylsydnone absorbs at a longer wavelength such as 4-acetylphenylsydnone whose UV maxima was at 317 nm. Similarly, 4-acylated sydnone absorbs at a longer wavelength such as 4-acetylphenylsydnone whose UV maxima was at 317 nm. Conjugation between sydnone ring and the aromatic system substituted at N3 was not essential for the stability of sydnones because the resonance between the two systems was minor [21].

However, more recent studies by Fan and his lab-mates gave a better insight into bond length and nature, atomic charges and electron density distribution for thirteen different sydnone compounds. Ring aromaticity is gained from the unevenly delocalized e-electrons with higher density on both N2-N3 and N3-C4. The calculated bond order of the N-O was 1.22 Å which is close to 1.14 Å of the double bond N=N, the X-ray confirmed its single bond nature [19, 20]. Interestingly, the high electron density of C4 in structures XXV and XXVI was later supported by the finding of Greco and O’Reilly who reported a strong acidity of 3-phenylsydnone with pKa value of 18-20. Consequently, sydnone is an electron donating reagent and the electrostatic substitution at this position is possible [21].

Sydnones spectral studies

Ultraviolet (UV) spectroscopy

The properties of the ultraviolet spectra of sydnones were well reviewed by Stewart [23] and Kier and Roche [24]. Briefly, absorption maxima in the range 290-340 nm was considered as a proof of the presence of the aromatic ring of sydnone. Alkyl sydnone absorbs at the lower wavelength (<300 nm). For example, 3-methylsydnone, 3-n-butyl sydnone and 3-cyclohexylsydnone showed their UV absorption maxima at 290, 289.5 and 292 nm, respectively. A bathochromic shift was observed for 3-arylsydnone due to conjugation as in 3-phenylsydnone and 3-(1-naphthyl) sydnone which absorb at 310 and 315, respectively. Many factors can remarkably affect the UV spectra of sydnones:

1. Conjugation: An aromatic system substituted at C4 of the sydnone ring has a stronger bathochromic effect such as 3-methyl-4-phenylsydnone whose UV maxima was at 317 nm. Similarly, 4-acetylsydnone absorbs at a longer wavelength such as 4-acetyl-3-phenylsydnone and 3-phenylsydnone absorb at 324 and 310 nm.

2. Steric factors can retard the conjugation due to the disturbance of the planarity of the molecule. The UV maxima of 3-(2, 6-dimethylphenyl) sydnone was found to be at 255 nm even shorter than that of 3-alkylsydnone which lacks conjugation.

3. Electrostatic interaction in bis-sydnone system makes the coplanarity system more rigid and therefore the UV absorption wavelength is unusually high like in XXVII, XXVIII, XXIX and XXXI whose maximum absorptions were at 292, 350, 292 and 303, respectively.

Infrared (IR) spectroscopy

A survey of the literature since their early preparation until today revealed two characteristic IR bands for sydnones. The stretch of sydnone carbonyl (C5=O) ranges from 1740 to 1770 cm⁻¹ while the absorption band of carbon-hydrogen (C4-H) was more than 3000 cm⁻¹. However, electrophilic substitution at C4 led to the loss of the carbon-hydrogen band and an increase in the wavenumber of the carbonyl up to 1780-1830 cm⁻¹. For example, acetylation of 3-(4-chlorophenyl) sydnone resulted in upshifting the CO band from 1750 cm⁻¹ to 1780 cm⁻¹. Nevertheless, more recent studies by Fan and his lab-mates gave a better insight into bond length and nature, atomic charges and electron density distribution for thirteen different sydnone compounds. Ring aromaticity is gained from the unevenly delocalized e-electrons with higher density on both N2-N3 and N3-C4. The calculated bond order of the N-O was 1.22 Å which is close to 1.14 Å of the double bond N=N, the X-ray confirmed its single bond nature [19, 20]. Interestingly, the high electron density of C4 in structures XXV and XXVI was later supported by the finding of Greco and O’Reilly who reported a strong acidity of 3-phenylsydnone with pKa value of 18-20. Consequently, sydnone is an electron donating reagent and the electrostatic substitution at this position is possible [21].

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refluxed with one molar equivalent of carboxylic acid and sydnone [29]. Later, other chemists speculated that the failure of Friedel-Crafts acylation was due to the coordination between Lewis acid and the exocyclic oxygen of the sydnone which eventually yielded a sydnone-containing fused ring compounds rather than the desired acylated product as shown in Scheme 3 [30]. More studies on the acylation of sydnone were later conducted employing a wide range of catalysts with a moderate to excellent yield as summarized in table 1.

Table 1: Acylation of 3-arylsydnone

| Catalyst | Experimental | Yield (%) | Reference |
|----------|--------------|-----------|-----------|
| Bis(triphenylphosphine)palladium(II) dichloride | Arylsydnone-4-yl copper, acyl chloride and the catalyst for 2 h at room temperature | 50-95 | [31] |
| PdCl₂ [(PH₃)₂]₂ | Reflux overnight in acetic anhydride | 25-86 | [32] |
| Montmorillonite K-10 | Reflux 7 h in acetic anhydride | 80-90 | [3] |
| 1,3-Dibromo-5,5-dimethylhydantoin (DBH) | Reflux 4 h in acetic acid at 110°C | 90-95 | [33] |
| N-Bromosuccinimide (NBS) | Refluxing an alkyl anhydride with the catalysts in acetonitrile at 95°C for 2-16 h | 48-93 | [34] |
| Bismuth(III) trifluoromethanesulfonate Bi(OTf)₃ and lithium perchlorate LiClO₄ | Overnight heating at 80°C in the presence of acetic anhydride in acetonitrile | 40-80 | [35] |
| Indium(III) trifluoromethanesulfonate In(OTf)₃ | Microwave irradiation for 15-45 min | 11-95 | [36] |

On the other hand, formylation of sydnone ring at C4 was conducted using the classic Vilsmeyer reaction with a yield up to 50%. Noteworthy, formylation occurred only at the sydnone ring, and no formyl group was found at the substituted phenyl ring [37]. Later, Yeh and his colleagues used the formylated sydnone to prepare 4-carbamoyl or 4-cyano sydnone from the reaction of 4-formyl-3-arylsydnone with concentrated or dilute sulfuric acid, respectively, in the existence of sodium azide [38]. Halogenation of sydnone ring

Many methods were developed for the introduction of halogens into carbon C4 of the sydnone ring Chloro, iso and bromo substituted sydrones were prepared successfully with a satisfactory yield as summarized in Scheme 4 [39-42]. To date, fluoro-containing derivatives at C4 have not been reported. However, Foster and his co-workers reported 4-trifluoromethyl-3-arylsydnone from 3,3,3-trifluorotridefluoro-2-(N-nitrosoarylamine)propanoic acid with 75-85% yield [28].

Sydnone lithiation and its application

The preparation of 3-aryl-4-lithiosydnone is straightforward and can be achieved by n-butyllithium in tetrahydrofuran (THF) within one hour. Fuchigami and others reported that the 4-sydnonyl anion generated from 4-lithiosydnone in THF could react easily and selectively with various chemicals to introduce heteroatom groups at the C4 position as a sole product with an excellent yield. They found that 4-sydnonyl anion had less nucleophilicity features than the ordinary aryl anion and therefore the reaction with phosphorus acid esters, tin (II) chloride and antimony trichloride was not successful as presented in Scheme 5 [43, 44].

Later, dilithium species (4-lithio-o-lithio arylsydnone) were reported for the first time by reacting 3-arylsydnone with 2.2 equivalents of n-butyllithium and tetramethylethlenediamine (TEMEDA) in THF at 78°C. The di-lithiated product was subsequently subjected to a variety of electrophiles to produce the disubstituted sydnone such as di-formyl, di-trimethylsilyl, di-halo, di-alkyl, di-thioether [45, 46] or fused sydnone rings such as 4-hydroxy-4-substituted sydno(3,4-a)(4H)indoles [47]. Also, the acylation of 3-arylsydnone at the ortho-ary1 position was accessed by adding Weinreb amides directly to the reaction pot of dilithium analogues as shown in Scheme 6 [48].
The biological activity of sydnone

The distinguished structure of sydnone having positive and negative charges along with its aromaticity and high lipophilicity enables it to react with biomolecules like DNA and enzymes. Consequently, sydnones exert a wide array of biological activities like anti-inflammatory, analgesic, anti-arthritis, cytotoxicity, anti-parasite (malaria and leishmaniasis), anti-diabetic, antioxidant, antimicrobial and nitric oxide donation [7]. In our present review, we will focus on the most studied and investigated biological activities.

Anti-inflammatory activity

The first report on the anti-inflammatory activity of sydnone-containing compounds was in 1974 by Wagner and Hill who reported that sydnones bearing 2-arylthioethyl or 2-arylsulfonylethyl at the position N3 were promising scaffolds for designing new anti-inflammatory drugs. Structure-activity relationship revealed that a small lipophilic group like methyl or hydrogen atom at C4 was essential for the activity. Aromatic ring attached to the sulfur increased the potency with a maximum activity when both ortho positions were substituted with an electronegative atom. They found that the inhibition of arthritic swelling by 4-methyl-3-[2-(phenylthio)ethyl] sydnone was equal to hydrocortisone and phenylbutazone while 4-methyl-3-[2-[2,4-dichlorophenylthio)ethyl] sydnone was six times stronger than hydrocortisone [49, 50].

Studies on the anti-inflammatory activity of more sydnone analogues were continued later by combining sydnone with other pharmacophores such as thiazole, pyrazole and styrly ketone. It was found that 3-substituted-4-(thiazol-4-yl) sydnone had a weak to moderate activity [51]. On the other hand, the presence of 5-arylpriazole at C4 of the sydnone ring resulted in a favourable anti-inflammatory activity as an anti-arthritis, anti-edema, and analgesic with less ulcerogenic side effects. For example, compound XXXV was found to be more effective than aspirin; ED₅₀ 28.3 vs. 81.4 mg/Kg, respectively [52]. Sydnones containing substituted styrlyketone were also investigated for their anti-inflammatory activity. Deshpande and his co-workers stated that some sydnonylstyrlyketone XXXVI showed significant analgesic activity especially when there was an electron withdrawing group attached to the styrly moiety such as furyl, 4-nitrophenyl, and 4-chlorophenyl. However, replacing the 4-methoxyphenyl group in XXXVI by 3-chloro-4-fluorophenyl enhanced the biological activity and decreased the ulcerogenicity compared to ibuprofen. Styrly-substituted sydnone exhibited a considerable analgesic activity in acetic acid-induced writhing but failed to show any significant activity in hot plate test suggesting that they act through peripheral rather than central effect [53, 54].

In other studies, sydnone ring hybrid with Mannich and Schiff bases XXXVII showed good anti-inflammatory and analgesic activities. Results showed that derivatives containing piperidine or morpholine had the highest anti-inflammatory activity in carrageenan-induced edemain rats with an analgesic activity comparable to the standard drug pentazocine [55]. A deeper insight into the mechanism of action of sydnones was provided by Kamble and his co-workers who postulated that benzophenone oxime compounds appended with sydnone directly inhibited phospholipase A2 (PLA2) by competing with the substrate at the binding cavity of the enzyme. Studies proved that PLA2 inhibitors can be as potent as steroidal anti-inflammatory drugs due to the reduction of lipid mediators which were usually secreted as a response to tissue injury [56]. Others found that 3-(4-chloro-3-nitro) sydnone mimicked the anti-inflammatory and immunosuppressive agents by reducing phagocytic activity, increasing superoxide anion production, inhibiting the production of nitric oxide, and declining interleukin-6 (IL-6) levels in peritoneal macrophages [57].

Cytotoxic and anticancer activity

In 1992, Grynberg and his co-researchers reported for the first time a successful usage of some sydnone derivatives as antitumor agents in vivo. They found that 3-(4-chloro-3-nitrophenyl) sydnone and 3-(4-pyridyl-2-thione-3-nitrophenyl) sydnone exhibited a significant cytotoxic activity against sarcoma 180, Ehrlich carcinoma and B10MCH fibrous histiocytoma. Remarkably, only the first one showed a growth inhibitory activity against L1210 leukaemia ascites tumours. They argued that the cytotoxic effect of sydnones might be due to inhibition of thymidine uptake by the cancerous cells [58]. Furthermore, sydnones were also linked to other pharmacologically active molecules to produce more potent cytotoxic agents. In this stream, sydnone-substituted chalcones were successfully synthesized and significantly inhibited the growth of Ehrlich ascites cells and Dalton’s lymphoma ascites cells. Noteworthy, the existence of a methyl group on the chalcone moiety enhanced the survival of the experimental tumour-bearing animals while chloride atom produced a toxic compound [59]. A few years later, other 3-(halogen-substituted phenyl) sydnones were synthesized and tested against many cancer cell lines in vitro. It was found that a fluoride atom at the para position of the phenyl ring resulted in a sound antiproliferative activity against breast cancer MCF7, lung cancer NCI-H460 and central nervous system cancer SF-268. In contrast, replacing the halogen atom by other heterocyclic rings such as indole and isoindole was detrimental to the cytotoxic activity [60].

In addition, new stilbene-sydnone hybrids were prepared and found to decrease the viability and proliferation of cervical carcinoma (HeLa), breast carcinoma (MCF7), colon carcinoma (SW620), pancreatic carcinoma (MiaPaCa2) and lung carcinoma (H460) cell lines in vitro. The most potent agents were found to have a chloride or methyl substituent on the stilbene moiety with a phenyl or methyl group at C4 of the sydnone ring like in structures XXXVIII and XXXIX [61]. Others reported a significant anticancer activity for new sydnones derivatized with imidazo [2,1-b][1,3,4]thiadiazole and coumarin at C4 of the sydnone ring (XL) against HT-29 human colorectal adenocarcinoma cell line. They found that the hydrophobicity of R was crucial for the cytotoxic activity and the existence of a chlorine atom on the coumarin ring (R') sharply raised the activity to be comparable to that of cisplatin [62].
Coordination complexes containing a metallic coordination center, sydnones, and other ligands were also synthesized and reported as new efficacious drugs such as the tridentate palladium Pd (II) complexes with thiosemicarbazone and phenyl sydnone (XLII). Sydnone and thiosemicarbazone bonded to the Pd (II) atom via the sydnone exocyclic oxygen, azomethine nitrogen and the sulfur atom afforded a yield of 40-60%. The complexes exhibited a sound antibacterial activity against human hepatocellular carcinoma (HepG2) and human cervical epithelial carcinoma (HeLa) cells. The IC_{50} of Pd-sydnone complexes were 0.77-2.25 μM against HepG2 and 0.36-1.30 μM against HeLa whereas the IC_{50} of the standard drug 5-fluorouracil were 6.94 and 0.71 μM, respectively [63].

To further elucidate the mechanism of action of sydnone cytotoxicity, investigations were carried out on 3-(4-chloro-3-nitrophenyl) sydnones (SYD-1). It was reported that mitochondria were the main targets of SYD-1 by lowering cellular energy production. Lipophilicity of SYD-1 enables it to interact with mitochondrial membrane resulting in altering of the components' redox state and finally shrinking of the mitochondria. It was also found that sydnone can lower the activity of glutamate dehydrogenase (GLDH). Being a nitric oxide donor, SYD-1 can interfere with the mitochondrial respiratory chain such as cytochrome oxidase leading to less oxygen utilization. Collectively, sydnone-induced cell death can be considered as a consequence of increased apoptosis [66]. Moreover, SYD-1 effectively reduced the oxidative stress of rat isolated mitochondria because it inhibited oxoglutarate-induced lipoperoxidation, reduced the formation/opening of Ca^{2+}-mediated permeability transition pores, and suppressed nicotinamide adenine dinucleotide phosphate-oxidase (NADPH) [65]. Recently, Galuppo et al. evaluated the biological activity of SYD-1 in tumour-bearing Wistar rats. They reported a significant decrease in tumor volume and tumor weight after 12 d of treatment by SYD-1 at a dose of 75 mg/Kg. Histological examination of the tumor revealed coagulative necrosis and apoptotic spots. Therefore, they attributed sydnone activity to the activation of apoptosis pathways by decreasing the expression of anti-apoptotic enzyme Bcl-2 and increasing both apoptotic bodies and pro-apoptotic proteins (Bax and p53). However, splenomegaly was observed in the treated animals, and it was linked to sydnone-induced extravascular hemolysis [66].

**Antimicrobial activity**

It has been demonstrated by numerous studies that sydnone derivatives have antibacterial and antifungal activities. Penicillin 3-arylsyndone hybrids XLII were prepared by Naito and his colleagues from 3-arylsydnone-4-carboxylic acid and 6-aminopenicillanic acid. They were found to be active against penicillinase-producing bacteria strains. On the other hand, penicillin 3-alkylsyndrones were inactive against the same resistant strains. It was postulated that the existence of a phenyl group at N3 of the sydnone ring resulted in steric hindrance which protects the β-lactum carbonyl in a manner similar to that of oxacillin XLII [67].

Sydnone-chalcone hybrids were also prepared and showed high antibacterial activity against gram-positive bacteria (**Staphylococcus aureus**) and weak activity against gram-negative bacteria (**E. coli**).

However, they did not exert antifungal activity. The existence of nitro group at the chalcone moiety enhanced the antibacterial activity. Remarkably, bromination of the α, β-unsaturated ketone of the chalcone and the position C4 in the sydnone ring resulted in good bactericide molecules [68].

Recently, more sydnone-containing antimicrobial agents were also prepared with more potent and wide spectrum activity. To mention some examples, 4-aminotriazine linked to the carbon C4 of the sydnone ring via mercaptoacetyl (XLIV), triazole (XLV) and benzothiazole (XLVI) linked to the sydnone ring via Mannich base showed promising antibacterial and antifungal activities even better than some known agents like nitrofurazone and ciprofloxacin [69-71].
Antioxidant activity

Sydnones and its related compounds have been reported as antioxidant agents in various literature sources. In 1994, it was found that sydnone ring can enhance the antioxidant activity of chalcone by inhibiting lipid peroxidation and scavenging free radicals. Of interest, sydnone-substituted chalcones suppressed superoxide production by peritoneal macrophages in vitro in the presence of phorbol myristate acetate ester (PMA) which was linked to tumour growth [59]. Recently, 3-(halogen-substituted phenyl) sydnones combined with chalcone were reported as strong 2,2-diphenyl-1-picrylhydrazyl (DPPH) free radical scavengers. The existence of fluoride and chlorine atoms at the phenyl ring of the sydnone moiety increased the antioxidant activity by nine folds compared to the commonly used antioxidant agent butyl hydroxy anisole (BHA) [77].

Additionally, sydnones substituted at C4 with thiazolidinone and thiazole rings exhibited a moderate to potent DPPH free radical scavengers. The 2,3-dihydrothiazole ring linked to 3-phenylsydnone yielded powerful and rapid antioxidant compounds (LII) whose scavenging activities were comparable to that of α-tocopherol. On the other hand, sydnones bearing 4-oxothiazoline (LIII) were less active. The absence of N-H group rendered the latter to be a weak scavenger [78, 79].

\[
\begin{align*}
\text{LII} & & \text{LIII} \\
\begin{array}{c}
\text{O} \\
\text{N} \\
\text{H} \\
\text{N} \\
\text{O}
\end{array} & & \\
\begin{array}{c}
\text{O} \\
\text{N} \\
\text{H} \\
\text{N} \\
\text{O}
\end{array}
\end{align*}
\]

Antimalarial activity

In 1965, sydnone-based derivatives emerged as a new class of antimalarial agents. It was reported that 3-piperonylsydnone (LIV) and 3-phenylsydnone showed activity against Plasmodium berghei; the main parasite that causes malaria in mice. Nyber and Chen stated that 3-piperonylsydnone exhibited antimalarial activity when administered orally or subcutaneously at a dose of 10 mg/Kg with no toxic side effects even at a dose up to 500 mg/Kg [80]. Since 3-phenylsydnone was less toxic and higher toxicity, it was of interest for other researchers to conduct a structure-activity analysis on the antimalarial activity of sydnone and piperonyl compounds. It was found that the N-N bond is essential for the antimalarial activity either in the sydnone or in the piperonyl moiety as highlighted in structures LIV-LVII. However, 3-piperonylsydnone was still the most active molecule among all tested compounds [81]. On the other hand, 4,4-bis (acetamidophenyl) sulfone derivatives were very potent antimalarial agents, while sydnone rendered it less active or inactive when they were combined in one structure LVIII [82]. Unfortunately, studies on sydnone-containing antimalarial agents were discontinued.

\[
\begin{align*}
\text{LIV} & & \text{LV} \\
\begin{array}{c}
\text{O} \\
\text{N} \\
\text{H} \\
\text{N} \\
\text{O}
\end{array} & & \\
\begin{array}{c}
\text{O} \\
\text{N} \\
\text{H} \\
\text{N} \\
\text{O}
\end{array} & & \\
\begin{array}{c}
\text{OCH}_3 \\
\text{OCH}_3
\end{array}
\end{align*}
\]

\[
\begin{align*}
\text{LVII} & & \text{LV} \\
\begin{array}{c}
\text{O} \\
\text{N} \\
\text{H} \\
\text{N} \\
\text{O}
\end{array} & & \\
\begin{array}{c}
\text{O} \\
\text{N} \\
\text{H} \\
\text{N} \\
\text{O}
\end{array} & & \\
\begin{array}{c}
\text{N} \\
\text{H} \\
\text{N}
\end{array}
\end{align*}
\]

CONCLUSION

The wide spectrum of chemical and biological properties of sydnone and its derivatives reported in diverse literature sources makes it of paramount interest for chemists and pharmacologists. Thereupon, sydnones were considered chemically and medicinally versatile and robust molecules. They merit more exploration to furnish novel sydnone analogues linked to various substrates as potential scaffolds for the discovery of new drugs.

CONFLICTS OF INTERESTS

Declared none

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