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Recent developments in the synthesis and biological activity of acridine/acridone analogues

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Many people in the world struggle with cancer or bacterial, parasitic, viral, Alzheimer's and other diseases. Therefore, many scientists seek new, more effective, more selective and less toxic drugs. Acridine/acridone derivatives constitute a class of compounds with a broad spectrum of biological activity and are of great interest to scientists. To date, many acridine/acridone analogues have been obtained, which, *inter alia*, exhibit antitumour (e.g., (1–5)), antimicrobial (e.g., (59)), and antiviral (e.g., (61)) activities and are applicable in the treatment of Alzheimer's disease (e.g., (26)). However, in many cases, their clinical application is limited and excluded because of side effects. In this survey, we describe acridine and acridone derivatives reported since 2013, methods of their synthesis and their potential clinical applications.

Introduction

Acridine derivatives form an important class of heterocycles containing nitrogen due to their broad range of pharmaceutical properties. Acridine derivatives are characterized by unique physical and chemical properties, biological activities, and industrial applications. In the nineteenth century, acridine derivatives were already used industrially as pigments and dyes.¹ More critical to the pharmaceutical industry, acridine derivatives have exhibited bioactivities such as anti-inflammatory,^{2,3} anticancer,⁴ antimicrobial,⁵ antitubercular,^{6,7} antiparasitic,⁸ antimalarial,^{9–11} antiviral^{12,13} and fungicidal activities.¹⁴ Acridine derivatives have been shown to be effective as inhibitors of acetylcholinesterase.¹⁵ Furthermore, acridines are used as dyes, fluorescent materials for visualization of biomolecules, and in laser technologies.¹⁶ These properties of acridines are attributed to their semi-planar heterocyclic structure, which appreciably interacts with different biomolecular targets. Acridine/acridone derivatives are found in natural plants and various marine organisms.^{17,18} Notably, the anticancer activity of acridine/acridone derivatives has attracted increasing interest. To date, many derivatives of acridine have been synthesized and tested for antitumour activity. The unique planar ring structure allows acridone derivatives to act as DNA intercalators^{19,20} and to inhibit topoisomerase or telomerase enzymes.^{21–24} A variety of acridine/acridone derivatives have been synthesized; analogues such as *N*-(2-(dimethylamino)ethyl)acridine-4-carboxamide (DACA) (1),^{25–27} triazoloacridone (C-1305) (2)²⁸ and amsacrine (*m*-AMSA) (3)²⁹ (Fig. 1) have entered clinical studies. Among them, *m*-AMSA (3) was the first synthetic drug exhibiting clinical efficacy as a topoisomerase inhibitor. Many *m*-AMSA derivatives (AHMA (4),

D3CLP (5)) (Fig. 1) have been developed for stronger anti-cancer properties and removal of many harmful side effects.^{27,30}

Intermolecular interactions in acridine and acridinium derivatives determine their biological and physical properties including their chemiluminogenic abilities. Therefore, hydrogen bonding and π - π interactions within the Hirshfeld surface have been studied. Recently, Wera and co-workers reported the synthesis and structural investigations of some new acridine and acridinium derivatives.³¹

Naturally occurring acridine/acridone

Acridine/acridone alkaloids

Alkaloids are naturally occurring chemical compounds, which have a wide range of pharmacological activities (e.g., antimalarial, anticancer, and antibacterial).³² To date, the literature describes a number of acridine/acridone alkaloids, which have been tested as anticancer, antibacterial and antimalarial agents and against Alzheimer's disease.¹⁸ Examples of these compounds are cystodytin A (6) (isolated from various marine organisms) and acronycine (7) (isolated from bark of Australian scrub ash tree) (Fig. 2).^{33–35}

Arai *et al.* describe the bioassay-guided fractionation and Ngn2 promoter activity of acridine alkaloids (8–10) from an extract of a culture of *Streptomyces* sp. IFM 11440.¹⁷ Inubosin B (9) bearing a hydroxy group at the 4 position showed potent Ngn2 promoter activity, which was dose-dependent. Moreover, compound (9) demonstrated more activity than the positive control baicalin (11), while Insubosin A (8) and Insubosin C (10), which have a hydroxy group at the 5 position, did not show significant activity (Fig. 3).¹⁷

The Wouatsa group isolated acridone alkaloids (12–21) (Fig. 4) from the MeOH extract of the fruits of *Zanthoxylum zanthoxyloides* and *Zanthoxylum lepreurii*, of which six were new

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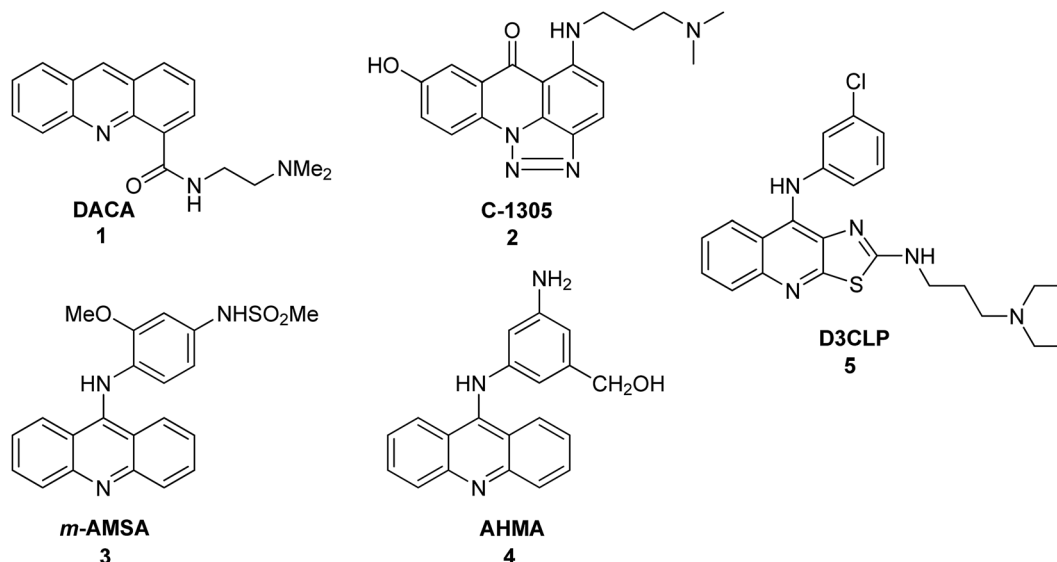


Fig. 1 Structures of DACA (1), C-1305 (2), *m*-AMSA (3), AHMA (4), and D3CLP (5).^{27–30}



Fig. 2 Structure of cystodytin A (6) and acronycine (7).³³

Fig. 4 Structures of acridones (12–21).³⁶

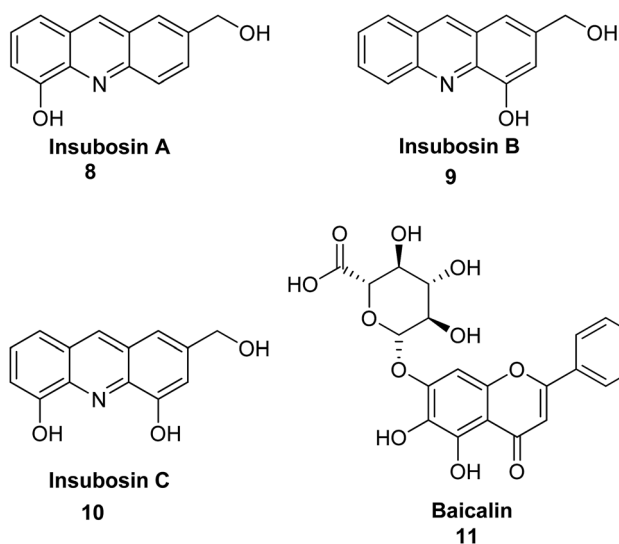


Fig. 3 Structure of acridine alkaloids (8–10) and baicalin (11).¹⁷

(12, 13, 14, 19, 20 and 21). Acridone molecules (19–21) that have a tetracyclic acridone structure as their carbon skeleton have been named as zanthacridone.³⁶

All newly discovered acridones (12, 13, 14, 19, 20 and 21) were tested for antibacterial activity against *Bacillus subtilis* (MTCC 121), *Micrococcus luteus* (MTCC 2470), *Staphylococcus aureus* (MTCC 96), and *Pseudomonas aeruginosa* (MTCC 741). Compounds (12, 14, 19, and 21) exhibited activity against MTCC 741 with MIC values equal to 500, 125, 250, and 250 $\mu\text{g mL}^{-1}$, respectively. Furthermore, analogue (14) inhibited MTCC 2470 (MIC 125 $\mu\text{g mL}^{-1}$), while molecules (15 and 18) failed to show appreciable activity against MTCC 741 but showed good activity in the inhibition of MTCC 96 (MIC 250 and 500 $\mu\text{g mL}^{-1}$, respectively), MTCC 121 (MIC 250 and 250 $\mu\text{g mL}^{-1}$, respectively) and MTCC 2470 (MIC 125 and 500 $\mu\text{g mL}^{-1}$, respectively).³⁶

Acridones (12, 15, 18, and 19) were also tested for *in vitro* cytotoxicity against liver (WRL-68), colon (Caco2), breast (MCF-7) and prostate (PC-3) cancer cell lines by MTT assay. Studies indicated that compound (12) showed moderate cytotoxic activity against WRL-68 with an IC₅₀ value equal to 86 μM .³⁵ Moreover, quantitative SAR and molecular modelling studies



showed that molecules (**12**, **20**, and **21**) are inhibitors of aromatase and glycosyltransferase.³⁶

Further acridone alkaloids chlorospermine A (**22**), chlorospermine B (**23**) and the known atalaphyllidine (**24**) and acrifoline (**25**) (Fig. 5) were isolated from the stem bark of *Glycosmis chlorosperma* by Beniddir *et al.*³⁷ They used bioassay-guided isolation for this purpose.

Acridone molecules (**22–25**) were tested for DYRK1A, CLK1, CDK1, CDK5, GSK3, and CK1 activity. Acrifoline (**25**) showed inhibition activities of DYRK1A, CLK1, GSK3, CDK1, and CDK5 with IC₅₀ values of 0.075, 0.17, 2, 5.3 and 9 μM, respectively. Chlorospermine B (**23**) and atalaphyllidine (**24**) exhibited activity against DYRK1A with IC₅₀ values of 5.7 and 2.2 μM, respectively. Furthermore, compound (**23**) demonstrated inhibition of CLK1 (IC₅₀ = 7 μM), while chlorospermine A (**22**) failed to show any activity against various kinases. Thus, studies performed suggest that acridone may constitute a molecular skeleton in search of new inhibitors of DYRK1A.³⁷ Moreover, a molecular docking study of (**22–35**) showed that interactions of the ligand with conserved Lys188 and backbone atoms in the hinge region (Glu239 and/or Leu241) are necessary for good biological activity. The most active compound (**25**) forms hydrogen bonds between the hydroxy group at C-6 and Glu203 and the conserved Lys188 and interacts with backbone atoms of Glu239 and Leu241 through hydrogen bonds involving the C-1 hydroxy group. Atalaphyllidine (**24**) is proposed to interact with backbone atoms of Glu239, Leu241, and Ile165 through hydrogen bonds involving the C-9, C-1, and C-5 oxygen atoms, respectively. In turn, derivative (**23**) showed hydrogen bonds between the hydroxy group in position C-5 and the side chains of Glu203 and Lys188, while inactive chlorospermine A (**22**) forms hydrogen bonds between its oxygen atoms at C-5, C-9, and C-2' and backbone NH of Leu241 and side chains of Asn244 and Asp307 (from the DFG motif), respectively.³⁷

Citrusinine-I (**26**) (Fig. 6) is an example acridone alkaloid isolated from *Swinglea glutinosa*. These analogues were described as herbicide models, and their inhibition of photosynthesis was investigated against both photosystem I and II.

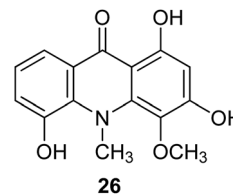


Fig. 6 Structure of citrusinine-I (**26**).³⁸

Measurements *in vitro* demonstrated significant influence on various stages of reactions in the course of photosynthesis; however, extracts provided only several milligram amounts of tested compounds, and design of multi-stage synthesis would be necessary to accomplish *in vivo* examinations.³⁸

Teow presented the methods for treating various liver diseases and certain cancers by used compositions of acridone, xanthone, thioxanthone, tocotrienol and oleanolic triterpenoid compounds.³⁹ He isolated hepatic therapeutic actives from the dried stem, bark and roots of *Cratoxylum cochinchinense* plant. The acridone molecules are at least one of the anti-viral components in the hepatic pharmaceutical active which probably function by inhibiting the growth and multiplication of the chronic hepatitis HBV and chronic hepatitis HCV in patients afflicted with chronic hepatitis B or chronic hepatitis C.

Synthetic derivatives of acridine/ acridone

Acridine/acridone as inhibitors of acetylcholinesterase

Alzheimer's disease (AD) is the most common form of dementia, an incurable and neurodegenerative disease, which manifests *via* progressive decline in intellectual abilities, caused by the disappearance of nerve cells.^{38,40} Because AD is currently one of the largest global medical problems, it has attracted the attention of medicinal chemists, who have studied new drugs for this disease.⁴¹ The drugs used in AD include cholinesterase inhibitors, which enhance neurotransmission of cholinergic synapses in the brain and thus enhance intellectual activity.⁴² The acridine derivative used in medicine as an anti-AD drug is tacrine (**27**) (Fig. 7).⁴³

Acridines can be also used to diagnosis of neurodegenerative disorders, such as AD. Fuchigami *et al.* developed ¹²⁵I-labeled acridines possessing high affinities for Aβ aggregates involved in these pathologies.⁴⁵ During *in vivo* experiments on Aβ plaques in the brain slices of Tg2576 mice, the most promising results were obtained in case of 6-iodo-2-methoxy-9-methylaminoacridine and 2,9-dimethoxy-6-iodoacridine as potential imaging agents for amyloid in living brain.

The Hamulakova group reported the potential of novel tacrine/acridines (**28–30**, **31a–c**) (Fig. 7) as human acetylcholinesterase (hAChE) and human butyrylcholinesterase (hBChE) inhibitors.⁴⁶ *In vitro* studies have shown that the compounds (**29**, **30**, **31b** and **31c**) proved to be especially potent against hAChE with IC₅₀ values in the range of 2 to 8 nM, and analogues (**28**, **29** and **31a–c**) were the most effective inhibitors against hBChE with IC₅₀ values 0.4–20 nM; however, tacrine (**27**) as

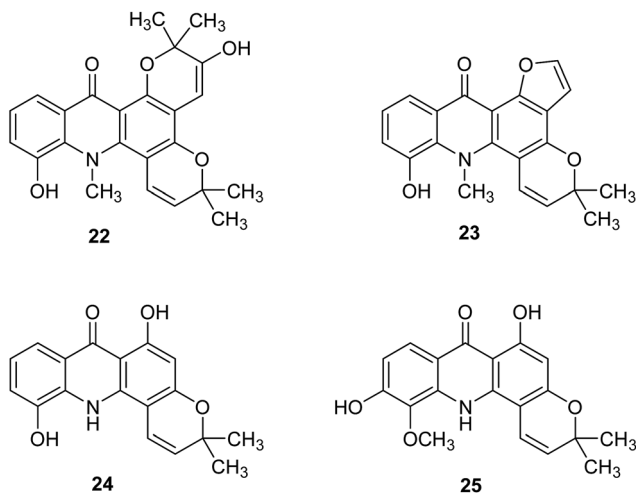


Fig. 5 Structure of acridone alkaloids (**22–25**).³⁷



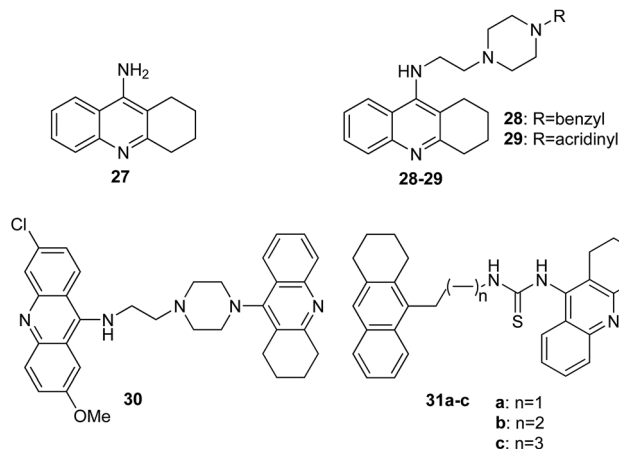


Fig. 7 Structure of tacrine (27) and tacrine/acridine derivatives (28–30, 31a–c).⁴⁴

a reference compound showed less activity against hAChE ($IC_{50} = 500$ nM) and hBChE ($IC_{50} = 23$ nM).⁴⁴ Docking models of the binding of compound (29) into hAChE showed cation- π interactions between Tyr341 and Trp286 residues of the peripheral anionic site and the positively charged acridine ring and showed cation- π interactions between Trp86 and the protonated tacrine ring. Additionally, stabilization of compound (29) may be due to a hydrogen bond between the amino group of tacrine and the oxygen atom of Tyr337.⁴⁴

Furthermore, Janockova *et al.* studied the interaction of derivative (30) with ctDNA.⁴⁷ The results of UV-Vis absorption spectroscopy showed that the binding of compound (30) occurred with a binding constant value $K = 2.0 \times 10^3$ M⁻¹. Thus, the results are indicative of a strong interaction between analogue (30) and ctDNA. Additionally, they demonstrated that

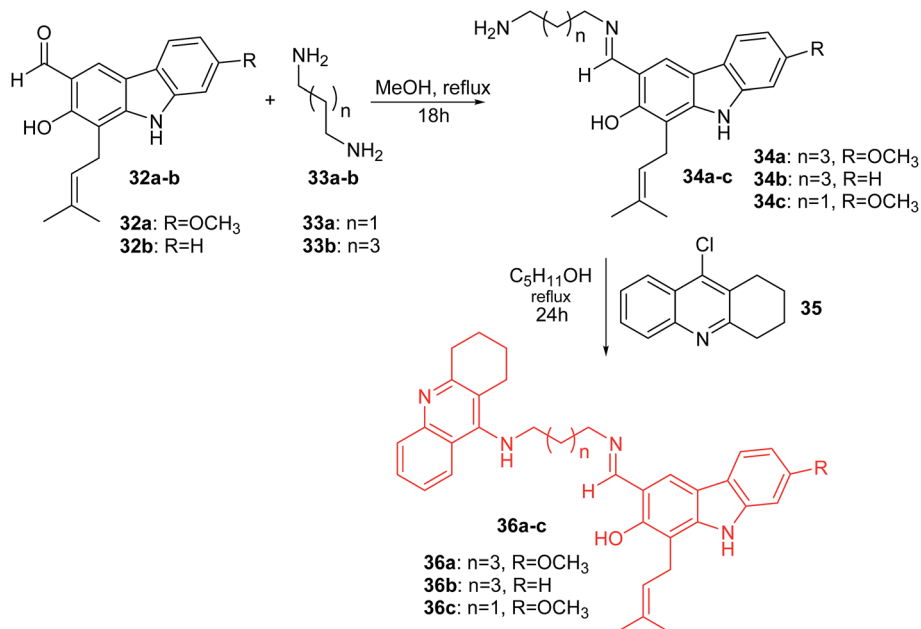
(30) is a dual topo inhibitor, acting both as a topo I inhibitor and as a topo II catalytic inhibitor.⁴⁶

Thiratmatrakul and co-workers synthesized the new tacrine-carbazole hybrids (36a–c) as potential multifunctional agents for the treatment of AD.⁴⁸ The intermediate imines (34a–c) were prepared by reaction of 7-methoxyheptaphylline (32a) or heptaphylline (32b) with 1,3-diaminopropane (33a) or 1,5-diaminopentane (33b) in MeOH. The reaction of the corresponding imines (34a–c) with 9-chlorotacrine (35) in the presence of pentanol obtained compounds (36a–c) (Scheme 1).⁴⁷

Derivative (36a) revealed the most potent activity against AChE inhibitory and antioxidant action. Additionally, the molecular modelling study showed that synthesized analogues demonstrated binding with the catalytic active site (CAS) and the peripheral anionic site (PAS) of AChE. Furthermore, tacrine-carbazole hybrids exhibited a neuroprotective effect against oxidative stress induced by H₂O₂ and A β 1-42 toxicity. Moreover, analogue (36a) exhibited an ability to improve short-term and long-term memory deficit in mice induced by scopolamine.⁴⁷

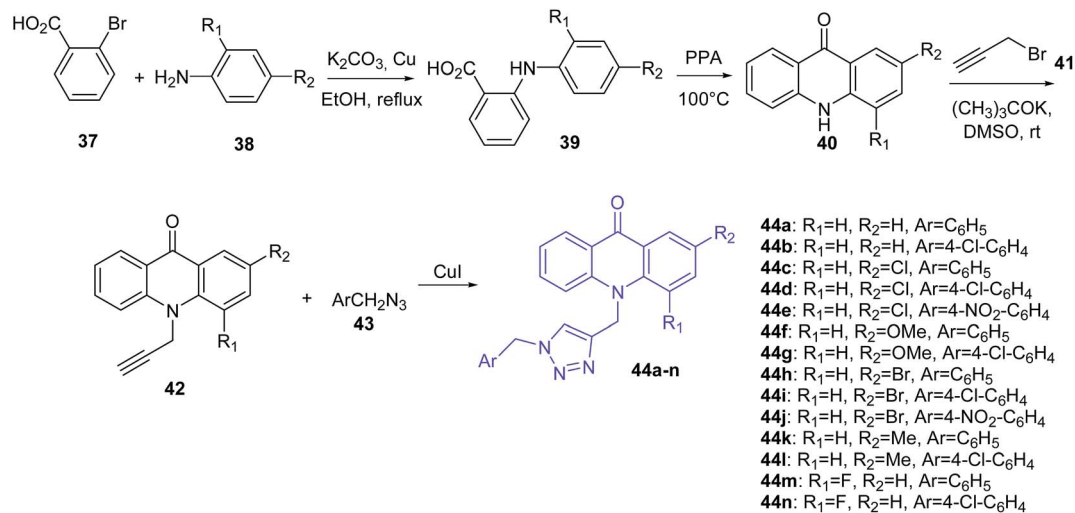
Mohammadi-Khanaposhtani *et al.* presented the synthesis of triazole derivatives of acridone (44a–n) (Scheme 2).⁴⁹ In the first step, 2-arylamino benzoic acids (39) were obtained by the Ullmann condensation reaction of 2-bromobenzoic acid (37) and various aniline derivatives (38) in the presence of potassium carbonate and copper in EtOH at reflux. Subsequent cyclization of compound (39) by PPA at 100 °C obtained acridone derivatives (40), which by reaction with propargyl bromide (41) using potassium *tert*-butoxide in DMSO at room temperature yielded 10-(prop-2-yn-1-yl)acridin-9-one derivatives (42). Then, compounds (42) were converted to the target triazole analogues (44a–n) using CuI and freshly prepared azide derivative (43).⁴⁸

Obtained derivatives of acridone (44a–n) were tested *in vitro* for inhibition of acetylcholinesterase (AChE) and butyrylcholinesterase (BChE). The best ability to inhibit AChE was



Scheme 1 Synthesis of tacrine-carbazole hybrids (36a–c).⁴⁸



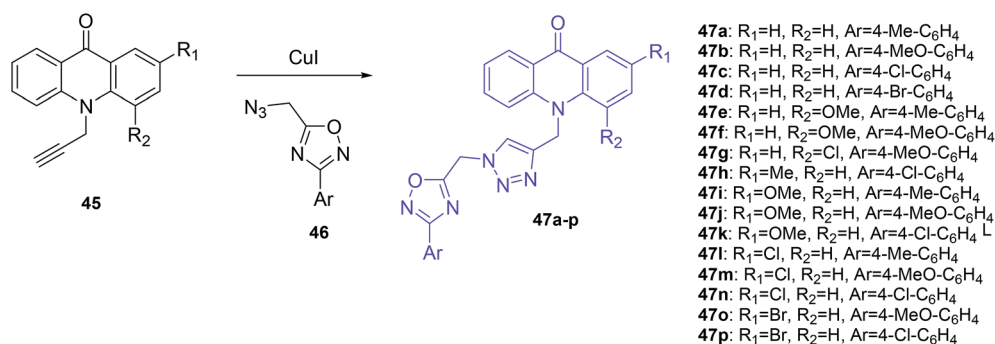
Scheme 2 Synthesis of triazole derivatives of acridone (**44a–n**).⁴⁹

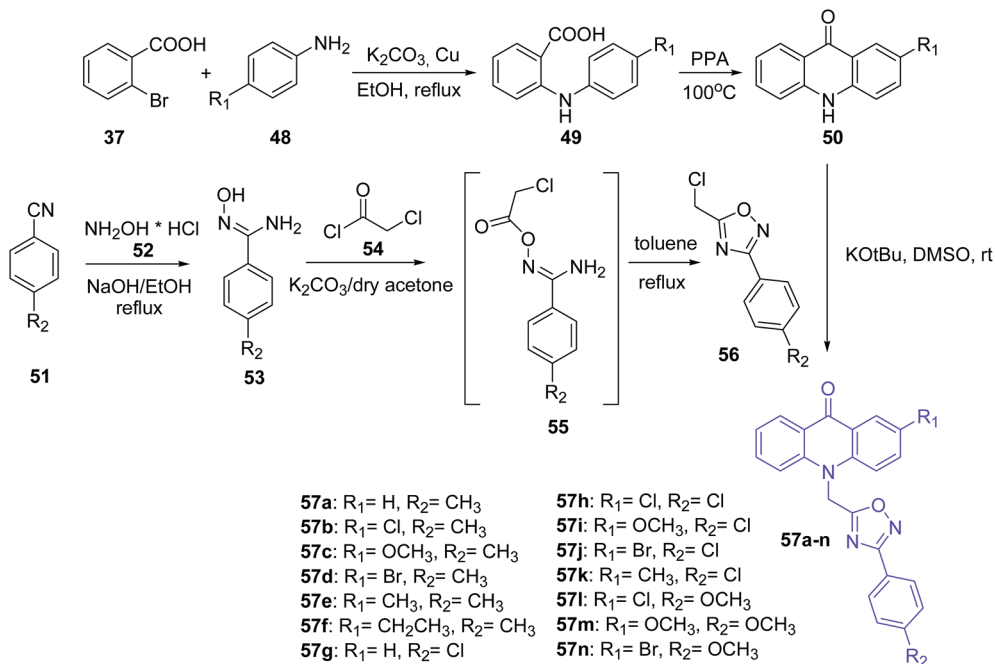
shown by compound (**44g**) (IC₅₀ = 7.31 μM), with a 4-substituted chlorine on the pendant benzyl group.⁴⁸ In turn, derivatives (**44e** and **44j**) with strong electron-withdrawing nitro groups into the pendant benzyl group showed decreased activity. Additionally, the presence of groups in the order of OMe > Me ≥ Cl > Br instead of hydrogen on the 2-substituted acridone moieties provided an increase in inhibitory activity. Thus, the presence of methoxy and chlorine groups contributes to good interactions with the active sites of enzymes and plays an important role in the inhibitory activity. Docking studies of compound (**44g**) showed π–π interactions between Phe330 and the 1,2,3-triazole moiety and between Trp84 and the acridone moiety. Additionally, derivative (**44g**) forms hydrogen bonds between the oxygen of OMe and Ser122. Furthermore, a weak interaction between chlorine and the backbone carbonyl group of amino acid is beneficial.⁴⁸

The Mohammadi-Khanaposhtani group also demonstrated acridone-1,2,4-oxadiazole-1,2,3-triazole hybrids as inhibitors of AChE and BChE.⁵⁰ In this research, a series of new acridone-1,2,4-oxadiazole-1,2,3-triazoles (**47a–p**) were obtained according to the synthetic pathway given in Scheme 3. The target compounds (**47a–p**) were synthesized by the reaction of 10-

(prop-2-yn-1-yl)acridin-9-one derivatives (**45**) in the presence of CuI and the freshly prepared azide derivative (**46**).⁵⁰

Among these compounds, (**47a–d**, **47k** and **47n**) showed an inhibitory effect against AChE with IC₅₀ values ranging between 11.55 and 77.79 μM. The most activity was shown by compound (**47b**) containing an unsubstituted acridone ring and 4-methoxyphenyl-1,2,4-oxadiazole moieties, while derivatives (**47c–j**, **47l–m**, and **47o–p**) containing acridone moieties 2 or 4-substituted by methoxy, chlorine or bromine groups demonstrated no observable activity. Additionally, it may be that a methoxy group into the pendant benzyl group imparted a higher activity than chlorine, methyl, and bromine substituents. Docking studies showed that the orientation of analogue (**47b**) in the active site of AChE was the same as donepezil. Namely, compound (**47b**) forms π–π interaction between the acridone moiety and Phe331 and Trp84 in the catalytic anionic site (CAS). Additionally, compound (**47b**) is proposed to interact with the hydroxyl group of Ser200 in the catalytic triad site through hydrogen bonds involving the carbonyl group of acridone. Further, the 1,2,4-oxadiazole moiety exhibited π–π interaction with a phenyl group of Tyr121 in the peripheral anionic site (PAS).⁵⁰

Scheme 3 Synthesis of acridone-1,2,4-oxadiazole-1,2,3-triazole derivatives (**47a–p**).⁵⁰

Scheme 4 Synthesis of compounds (57a–n).⁵¹

Mohammadi-Khanaposhtani *et al.* presented the synthesis of novel acridone-based 1,2,4-oxadiazoles (57a–n) as potential anticonvulsant agents. In the first step, 2-arylamino benzoic acids (49) were prepared by the Ullmann condensation reaction of 2-bromobenzoic acid (37) and different anilines (48).⁵¹ Then, the cyclization of compounds (49) by PPA yielded acridone derivatives (50). The second intermediate 3-aryl-5-(chloromethyl)-1,2,4-oxadiazole derivatives (56) were obtained starting from appropriate benzonitriles (51) and hydroxylamine hydrochloride (52) in the presence of sodium hydroxide in EtOH at reflux. Derivatives (53) and chloroacetyl chloride (54) reacted in dry acetone using potassium carbonate to yield *N*-(2-chloroacetoxy)-4-substituted benzimidamides (55). Subsequent cyclization of compounds (55) in refluxing toluene yielded 3-aryl-5-(chloromethyl)-1,2,4-oxadiazole derivatives (56). The desired final products (57a–n) were synthesized by reaction of acridone derivatives (50) with 1,2,4-oxadiazole derivatives (56) in the presence of potassium *tert*-butoxide in DMSO at room temperature (Scheme 4).⁵¹

Most prepared compounds showed good anticonvulsant activities. Derivatives (57a–c and 57f–n) had potent anticonvulsant activity in the PTZ-induced seizure test (ED_{50} = 2.08–18.93 mg kg⁻¹). The most potent anticonvulsant activity against PTZ-induced seizures were demonstrated by 4-methoxyphenyl-oxadiazole derivatives (57l and 57m) bearing a 2-Cl or 2-MeO on the acridone ring, with ED_{50} values of 2.08 and 2.12 mg kg⁻¹, respectively. Additionally, studies on anticonvulsant activity against MES-induced seizures showed that all prepared compounds (57a–n) had potent anticonvulsant activity (ED_{50} s = 2.24–8.71 mg kg⁻¹). The most potent anticonvulsant activity against MES-induced seizures was shown by compound (57b). Moreover, products (57a–n) were screened for their neurotoxicity in mice using the rotarod test, and these derivatives

displayed good anticonvulsant activity with a high safety profile. Furthermore, a docking study of analogue (57l) in the BZD-binding site of the GABA_A receptor confirms possible binding of derivative (57l) with BZD receptors.⁵¹

Inhibitors of human carbonic anhydrase isozymes based on acridines and acridones analogs

Carbonic anhydrases (CA, EC 4.2.1.1) are Zn(II) ion-containing metalloenzymes that catalyse the interconversion of carbon dioxide (CO₂) and bicarbonate (HCO₃⁻) and the corresponding dehydration of bicarbonate in acidic medium with regeneration of CO₂.^{52,53} Furthermore, they participate in crucial physiological processes associated with, among others, pH homeostasis, electrolyte secretion, respiration, CO₂ and ion transport, some biosynthesis reactions (*e.g.*, gluconeogenesis, lipogenesis), and tumourigenicity.^{54–56} To date, sixteen different α -carbonic anhydrase isozymes have been described in mammals.⁵⁷ CA isoforms are present in a variety of tissues and are involved in several important biologic processes (*e.g.*, acid–base balance, respiration, carbon dioxide and ion transport, bone resorption, ureagenesis, gluconeogenesis, lipogenesis, and electrolyte secretion). CA II, which is the most active as a catalyst for carbon dioxide hydration, is found primarily in red blood cells and in other secretory tissues of the gastrointestinal tract, kidneys, lungs, eye, and CNS. Therefore, α -carbonic anhydrase isozymes that participate in these processes are important therapeutic targets with the potential to be inhibited/activated for the treatment of a range of disorders such as oedema, glaucoma, obesity, cancer, and epilepsy.⁵⁸

The Yesildag group presented a one-step method for the synthesis of acridine sulfonamide derivatives (61a–n).⁵⁹ Compounds (61a–n) were prepared by the reaction of dimedone



(58), 4-amino-*N*-[amino(imino)methyl]-benzenesulfonamide (59) and 4-cyano-benzaldehyde (60a-n) in the presence of sulfuric acid as a catalyst in water at room temperature (Scheme 5). This method has high yields and purities of the compounds obtained. In addition, it has advantages of using H₂SO₄ as an environmentally friendly catalyst and H₂O as a green solvent.⁵⁹

All synthesized compounds (61a-n) were tested *in vitro* with respect to inhibition of hydratase and esterase (hCA I and hCA II) isozymes. Analogues (61a-d and 61f-j) showed moderate esterase activities of hCA I and hCA II. Compound (61i), containing a methyl group on the *para* position of the benzyl group, was the most powerful inhibitor against hCA I (IC₅₀ = 47.2 μM) and hCA II (IC₅₀ = 50.1 μM). In turn, the derivatives containing larger functional groups (-OCH₃ and -SCH₃) were not active.

This may be related as the steric effects of these groups may block the enzyme-inhibitor interaction.⁵⁹

Other derivatives of acridine bis-sulfonamides (66a-l) were presented by Esirden *et al.*⁶⁰ The intermediate nitro-acridine sulfonamides (64a-b) were prepared by a coupling reaction of dimedone (58), 4-aminobenzesulfonamide (62) and *meta*- or *para*-substituted nitrobenzaldehydes (63a or 63b) in the molar ratio of 2 : 1 : 1 in the presence of DBSA as a phase transfer catalyst in H₂O. Then, the reduction of the nitro group was achieved by using excessive reduction-reactive aqueous sodium polysulfide. The final products (66a-l) were synthesized from appropriate sulfonyl chlorides and amino acridines (65a and 65b) in THF in the presence of triethylamine (TEA) (Scheme 6).⁶⁰ Derivatives (64a-b, 65a-b, and 66a-l) were tested for *in vitro* inhibition activity against the cytosolic human isoforms (hCA I,



Scheme 5 Synthesis of acridine sulfonamide derivatives (61a-n).⁵⁹



Scheme 6 Synthesis of acridine bis-sulfonamide derivatives (66a-l).⁶⁰

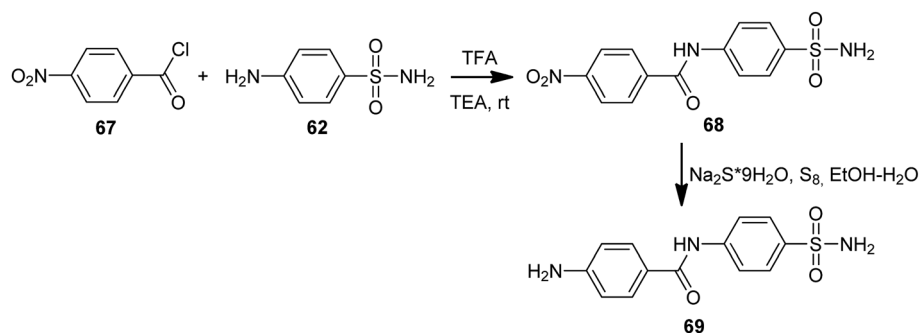


II, IX and XII). Compounds (**64a–b**, **65a–b**, and **66a–l**) were rather ineffective inhibitors against hCA I (0.69–8.91 μM). These analogues have low activity, probably due to the bulkiness of the tails that they incorporate and that most likely extend out of the active site.⁵⁹ In the case of hCA II, derivatives (**64a–b**, **65a–b**, and **66a–l**) were also rather ineffective inhibitors (0.10–0.96 μM), and no important effects of tail bulkiness were observed. This could be because hCA II has a wide entrance to the active site, and presumably the hydrophobic and hydrophilic interactions of these tails with the active site were sparingly effective. The transmembrane isoforms hCA IX and hCA XII were inhibited better than the cytosolic ones. However, compounds (**64a–b**, **65a–b**, and **66a–l**) showed more activity against hCA XII (0.09–1.12 μM) than against hCA IX (0.03–1.02 μM). The low inhibition of these analogues against the transmembrane isoforms (hCA IX and XII) may be because the variations in the structure are distant parts of the tail from the primary sulfonamide, which may result in these parts of the molecules being outside of the active site and causing only small interactions with amino acid residues crucial for the binding of inhibitors.⁶⁰

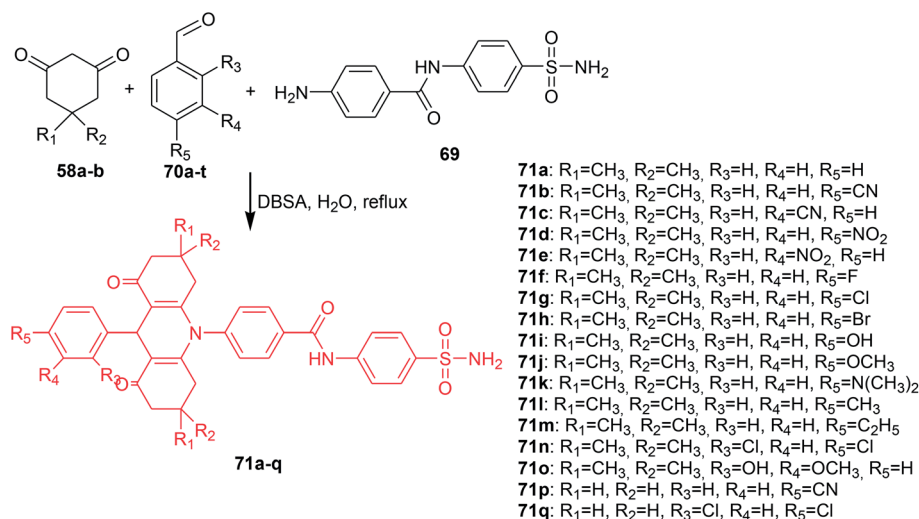
Ulus *et al.* presented the synthesis of acridine and bis acridine sulfonamides (**71a–r** and **73–74**) as cytosolic carbonic anhydrase isoforms I, II and VII inhibitors.⁶¹ The intermediate

(**69**) were obtained in two steps. Compound (**68**) was synthesized by reaction of 4-aminobenzenesulfonamide (**62**) with *p*-nitrobenzoyl chloride (**67**) in THF at room temperature in the presence of TEA. The nitro group of (**68**) was then reduced by aqueous sodium polysulfide in ethanol–water to obtain compound (**69**) (Scheme 7). Derivatives (**71a–r**) were prepared by coupling reactions of 4-amino-*N*-(4-sulfamoylphenyl)benzamide (**69**) with cyclic-1,3-diketones (**58a–b**) and aromatic aldehydes (**70a–t**) in the molar range of 1 : 2 : 1 in the presence of *p*-dodecylbenzenesulfonic acid (DBSA) as a phase transfer catalyst in H₂O (Scheme 8). Additionally, bis acridine sulfonamides (**73** and **74**) were synthesized in the molar range of 1 : 2 : 1 ((**69**) : 1,3-diketone (**58a**) : 1,3-benzenedialdehyde (**72a**) or 1,4-benzenedialdehyde (**72b**)) (Scheme 9).⁶¹

The prepared compounds (**71a–r** and **73–74**) exhibited good activity results against all three tested isoforms (hCA I, II and IV). Analogue (**71a**), containing only hydrogen, and derivative (**71o**), possessing OH and methyl moieties in the aromatic part of the molecule, were found to have the strongest activity against hCA I with K_{S} values 0.16 and 0.26 μM , respectively. Derivatives (**71a**, **71c**, **71e**, **71g**, and **71n**) showed effective hCA II inhibition with K_{S} values in the range of 15–23 μM . The best hCA IV inhibitors (K_{S} of 0.004, 0.005, and 0.022 μM) were compounds (**83a**, **83o**,



Scheme 7 Synthesis of 4-amino-*N*-(4-sulfamoylphenyl)benzamide (**69**).⁶¹



Scheme 8 Synthesis of acridine sulfonamides (**71a–q**).⁶¹



Scheme 9 Synthesis of bis acridine sulfonamides (73–74).⁶¹

and 71p), respectively, which were much more effective compared to acetazolamide (a reference compound).⁶¹

Acridine/acridone derivatives and their antibacterial and antiparasitic activity

To date, a great number of acridine compounds have been synthesized and clinically used as anti-bacterial drugs (*e.g.*, ethacridine (75)),¹⁸ anti-malarial agents (*e.g.*, quinacrine (76)) and anti-viral drugs (*e.g.* acranil (77)) (Fig. 8).^{62,63} The ability of acridine to bind to DNA has been used to inactivate genomes of viral and bacterial pathogens present in the cytoplasm and red RBC preparations used for clinical trials and transfusion. An example of such a compound is S-303 (78) (Fig. 8) containing the nitrogen mustard moiety. The ester bond is rapidly hydrolysed, which ensures a short half-life of the compound.⁶⁴

Acriflavine (ACF), a mixture of 3,6-diamino-10-methylacridinum chloride (trypaflavine) and 3,6-diaminoacridine (proflavine) showed

apart from antibacterial and anticancer, also potent antimalarial *in vitro* and *in vivo* properties in the nanomolar range.⁶⁵

In turn, Chiang proved that ACF can be an effective pharmaceutical for therapy in pathogenic protein diseases.⁶⁶ Additionally, ACF exhibited potential anticancer activity in mice and was approved by FDA for clinical trials.⁶⁷

The Joubert group presented the synthesis of artemisinin-acridine hybrids (82a–e).⁶⁸ The intermediate 9-aminoacridines (80a–e) were prepared by reaction of 6,9-dichloro-2-methoxyacridine (79) with the appropriate amine in the presence of K_2CO_3 in DMF. The desired final products (82a–e) were obtained using a microwave-assisted radiation method (150 W, 45 °C) by reacting compounds (80a–e) with 2-bromo-(10 β -dihydroartemisininoxy)ethane (81) in ACN in the presence of $NaHCO_3$ and KI as catalyst (Scheme 10).⁶⁸

Compounds (80a–e and 82a–e) were assayed *in vitro* for antimalarial activity against NF54 and Dd2 strains of *Plasmodium falciparum*. All prepared analogues (80a–e and 82a–e) were active and showed more antimalarial activity against NF54 than Dd2 inhibition. Derivative (82a), containing an ethylenediamine linker, exhibited the most potent activity with IC_{50} values equal to 2.6 nM against NF54 and 35.3 nM against Dd2. Interestingly, (82a) had 7-fold higher antimalarial potency than chloroquine (CQ) against both the NF54 and Dd2 strains, with high selective action towards the parasitic cells.⁶⁸

Furthermore, cytotoxic and anticancer effects of analogues (80a–e and 82a–e) on the CHO (Chinese hamster ovarian) and HeLa cell lines, respectively, were assessed by MTT assay. Derivatives (80a–e and 82a–e) have been found to be less cytotoxic compared to reference compound emetine. Hybrids (82a,d) showed stronger anticancer activity against the HeLa cells than the reference compounds (CQ and melphalan).⁶⁸

Bharathi *et al.* presented the synthesis, solvatochromic behaviour and larvicidal activity of substituted ethyl 10-chloro-4-(3,4-dimethoxyphenyl)-2-hydroxy-12-phenyl-1,4,5,6-tetrahydro-benzo[*a*]-acridine-3-carboxylates (85a–e).⁶⁹ Derivatives (85a–e) were obtained



Fig. 8 Structure of ethacridine (75), quinacrine (76), acranil (77) and S-303 (78).^{18,62–64}



Scheme 10 Synthesis of compounds (82a–e).⁶⁸

by reaction of (*E*)-7-chloro-2-(3,4-dimethoxybenzylidene)-9-phenyl-3,4-dihydroacridin-1(2*H*)-ones (83a–e) with ethyl acetoacetate (84) in the presence of 10% ethanolic NaOH (Scheme 11).⁶⁹

Derivatives (85b and 85e) showed good larvicidal activities against *A. stephensi* (LC₅₀ 126.3 and 108.17 mg L⁻¹, respectively) and *H. maculata* (LC₅₀ 111.57 and 241.05 mg L⁻¹, respectively). Furthermore, analogues (85b and 85e) were tested against non-target aquatic species (*S. annulatum* Fabricius, *Z. petiolatum* Rambur). The results showed that the prepared compounds are non-toxic. Derivatives (85b and 85e) were active against *S. annulatum* Fabricius and *Z. petiolatum* Rambur (with %

mortality values equal to 0.4 and 0.8 mg L⁻¹, % mortality values equal to 0.3 and 0.7 mg L⁻¹, respectively).⁶⁹

Medapi *et al.* presented the synthesis of *p*-phenylenediamine-linked acridine derivatives.⁷⁰ Acridine derivatives (91a–l and 92a–x) were synthesized according to Scheme 12. The synthetic pathway started from the Ullmann condensation reaction of 2-chlorobenzoic acid (86) and different *para*-substituted anilines (87a–c) to afford 2-(*p*-tolyl/4-chlorophenyl/phenylamino)benzoic acid derivatives (88a–c).

Compounds (88a–c) easily participated in the cyclization reaction in the presence of POCl₃ at reflux and afforded the

Scheme 11 Synthesis of acridine derivatives (85a–e).⁶⁹

Scheme 12 Synthesis of substituted *N*-[4-(substituted-9-acridinyl)amino]phenyl analogues (**91a-l**, **92a-x**).⁷⁰

corresponding 9-chloro-2-substituted acridine derivatives (**89a-c**). Then, *N*¹-(2-methyl/2-chloro/simple acridin-9-yl)benzene-1,4-diamine derivatives (**90a-c**) were received in the microwave-assisted reaction of derivatives (**89a-c**) with *p*-phenylenediamine in the presence of PTSA (*p*-toluenesulphonic acid) as the catalyst in methanol. Finally, the acridine derivatives (**90a-c**) were reacted with different substituted aryl sulfonylchlorides and aryl isothio/isocyanates to afford sulphonamide (**91a-l**) and thio/urea (**92a-l**, **92m-x**) derivatives.⁷⁰

These derivatives are structurally similar to *m*-amsacrine (**3**), which acts as a eukaryotic topoisomerase II poison and as an MTB topoisomerase I inhibitor with an IC₅₀ of 10 μM. An MTB topoisomerase I relaxation assay indicated that molecules (**91a-l** and **92a-x**) showed inhibition at 50 μM. Thus, a methoxyl group present in the phenyl ring linker of *m*-amsacrine is of importance. All synthesized compounds (**91a-l** and **92a-x**) were tested for *Mycobacterium tuberculosis* (MTB) DNA gyrase activity. Derivatives (**91a-l** and **92a-x**) showed good activity with IC₅₀ values in the range of 5.2 to 33.9 μM. The most active were sulphonamide (**91a-l**) followed by thiourea (**92a-l**) derivatives, and the least active were urea (**92m-x**) derivatives. The most potent was derivative (**91b**) with a CH₃ group at the 2-position of the acridine moiety and an unsubstituted phenyl ring. Furthermore, analogue (**91b**) was shown to be significantly potent with an MTB MIC (IC₅₀ = 6.59 μM) and 11.78% inhibition at 50 μM against mouse macrophage cell line RAW 264.7.⁷⁰

Muscia *et al.* described the synthesis of acridine derivatives (**95a-b** and **96a-b**) as potential anti-tuberculosis agents.⁷¹ The method is based on the use of the microwave-assisted Friedländer reaction. Compounds (**95a-b**) were prepared by reaction of cyclohexanone **94** with 2-amino-5-chlorobenzophenone (**93a**) or 2-amino-5-nitro-benzophenone (**93b**) in the presence of TFA as a catalyst in an MW oven at constant power (400 W) (Scheme 13). Analogues (**96a-b**) were obtained by microwave-assisted reaction of 5,5-dimethylcyclohexane-1,3-dione (**58**) with (**93a**) or (**93b**) (Scheme 13).⁷¹

Scheme 13 Synthesis of acridine derivatives (**95a-b**, **96a-b**).⁷¹

Synthesized compounds (**95a–b**) and, synthesized compounds (**95a–b** and **96a–b**) were assayed for inhibition of *Mycobacterium tuberculosis* H₃7Rv (Mtb). Derivative (**95b**) revealed the most potent activity with IC₅₀ values equal to 44.35 μM.⁷¹

A different method of obtaining derivatives of 1,8-dioxoacridine (**100a–l**) was presented by Kaya's group.⁷² In the first step of the synthesis, the condensation of cyclohexane-1,3-dione (**97**) or dimedone (**58**) with *p*-methoxybenzaldehyde (**98**) afforded the tetraketone intermediates (**99a** or **99b**). Reaction of tetraketone (**99a–b**) with primary aromatic amines using the Dean-Stark apparatus in the presence of acetic acid–sodium acetate in toluene gave acridine derivatives (**100a–l**) (Scheme 14).⁷²

All the synthesized compounds (**100a–l**) were tested for antibacterial and antifungal *in vitro* activity. Most of the analogues showed significant activity against *E. coli*, *P. aeruginosa*, *S. enteritidis* and *S. aureus*. Furthermore, compound (**100k**) showed more effective activity against *S. enteritidis* than the conventional reference antibiotic erythromycin. Additionally, all derivatives had moderate antifungal activity against *C. albicans* and *C. glabrata*.⁷²

Markovich *et al.* presented the synthesis of 2-(4-methyl-1,3-thiazol-5-yl)ethyl esters of acridone carboxylic acids (**103a–g**) by the transesterification of the corresponding butyl esters of

acridone carboxylic acids (**101a–g**) with 5-(2-hydroxyethyl)-4-methylthiazole (**102**) in the presence of sodium methoxide as the catalyst (Scheme 15).⁷³

Analogues of acridone that have a carboxyl group directly bonded to the aromatic ring of acridone also exhibit biological activity.^{74–76} Therefore, 2-(4-methyl-1,3-thiazol-5-yl)ethyl esters of 2(4)-carboxyacridone (**106–107**) were prepared.⁷³ The reaction was carried out at 70 °C for 1 h using 2- or 4-carboxyacridone acid chloride (**104–105**) and 5-(2-hydroxyethyl)-4-methylthiazole (**102**) (Scheme 16).⁷³

Compounds (**103a–b** and **107**) were tested for antibacterial activity against strains of microorganisms (*E. coli*, *Ps. aeruginosa*, *Pr. vulgaris*, *S. aureus*, *B. subtilis*, *Candida albicans*). The prepared analogues were active and showed inhibition against all microorganisms. Moreover, derivative (**103b**) with a CH₃ group at the 2-position of the acridone moiety showed slightly higher activity against *Candida albicans* in comparison to Rivanol.⁷³

Bhardwaj *et al.* synthesized new acridone derivatives. Acridone was coupled *via* a linker group to an anti-inflammatory (AI) drug such as ibuprofen (**109**), (*S*)-naproxen (**110**), acetyl salicylic acid (**108**) or chlororofecoxib analogue (**114**).⁷⁷ The intermediate *N*¹⁰-(2-hydroxyethyl)acridone (**112**) was prepared



Scheme 14 Synthesis of derivatives of 1,8-dioxoacridine (**100a–l**).⁷²



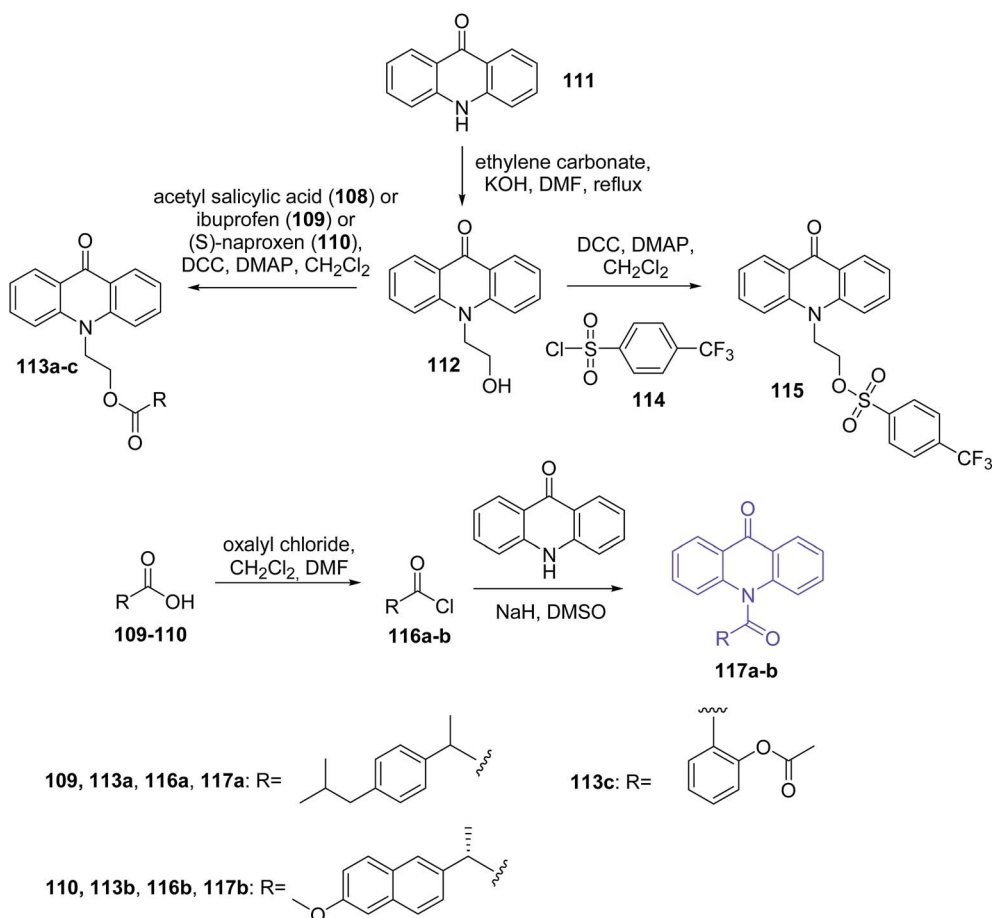
Scheme 15 Synthesis of acridone derivatives (**103a–g**).⁷³



Scheme 16 Synthesis of acridone derivatives (106–107).⁷³

by reaction of acridone (**111**) with ethylene carbonate in the presence of KOH. Acridone-ibuprofen (**113a**), acridone-(*S*)-naproxen (**113b**), acridone-acetyl salicylic acid (**113c**) or acridone-chlororofecoxib (**115**) derivatives were prepared by the condensation of compound (**112**) with ibuprofen (**109**), (*S*)-naproxen (**110**), acetyl salicylic acid (**108**) or 4-(trifluoromethyl) benzenesulphonyl chloride (**114**), respectively, in the presence of *N,N'*-dicyclohexylcarbodiimide (DCC) with 4-dimethylaminopyridine (DMAP) as catalyst in dry dichloromethane (DCM).

Additionally, the Bhardwaj group synthesized analogues (**117a–b**) by the condensation of acridone (**111**) with acid chloride derivatives of ibuprofen or (*S*)-naproxen (**116a–b**) in the presence of NaH in dry DMSO (Scheme 17).⁷⁷ Conjugates (**113a–c**, **115**, and **117a–b**) were assayed *in vitro* as COX-1 and COX-2 inhibitors. All prepared analogues (**113a–c**, **115**, and **117a–b**) showed more potent inhibition of COX-2 (IC₅₀ = 0.59–21.2 μM range) than COX-1 (IC₅₀ = 18.0 to 88.1 μM range). Furthermore, derivatives (**113a–c**, **115**, and **117a–b**) showed a higher COX-2

Scheme 17 Synthesis of acridone derivatives (113a–c, 115, 117a–b).⁷⁷

selectivity index (SI) than AI drugs (**108–110** and **114**). This may be caused by the fact that the derivatives (**113a–c**, **115**, and **117a–b**) have larger molecular volumes (353–412 Å³ range) than the smaller molecular volumes of the AI drugs (155–214 Å³ range). Thus, it is credible that the larger size of the conjugates of acridone may hinder their entry into the smaller COX-1 binding site (316 Å³), thus enabling selective entry into the larger COX-2 binding site (394 Å³). The most potent and selective COX-2 inhibitor was compound (**113a**) (IC₅₀ = 0.67 μM; SI = 110.6). Additionally, studies have shown that fluorescence emissions of conjugates (**113a–c**, **115**, and **117a–b**) (α_{em} = 415–445 nm) were not suitable for fluorescent imaging of cancer cells that over-express the COX-2 isozyme.⁷⁷

Anticancer activity of acridine/acridone analogues

The effectiveness of heterocyclic, tricyclic and planar arrays to intercalate between base pairs in the double-stranded DNA structure,^{78,79} and the ability of acceptor/donor nitrogen are two major considerations for the use of acridines/acridones as potential anti-cancer compounds.⁸⁰ Additionally, acridine/acridone derivatives prevent proper functioning of tumour cells, including by dysfunction relevant to the enzymes that control the topology of DNA function: topoisomerase, telomerase, and cyclin-dependent kinases.⁸¹ There are also reported ¹²⁵I-radiolabeled acridine derivatives investigated both *in vitro* and *in vivo* against melanoma.⁸²

In this survey, we describe the latest acridine and acridone derivatives as antitumour agents.

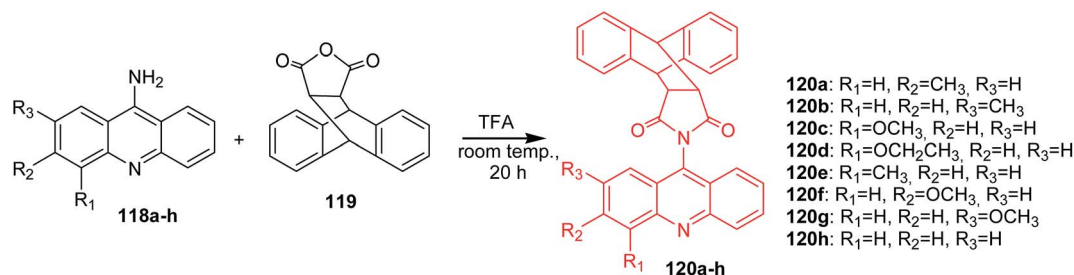
Stark and Dermawan reported a method for treating subjects having cancer that has developed resistance to tyrosine kinase inhibitors by using tyrosine kinase inhibitor (TKI) and NF-κB inhibitor.⁸³ NF-κB activation seems to be involved in the development of resistance to TKIs and other chemotherapeutic agents, and accordingly this resistance can be overcome by inhibiting NF-κB activation. The combination of the TKI and the NF-κB inhibitor provides a synergistic anticancer effect that overcomes the TKI resistance that has developed. For example, researchers administered to a subject a therapeutically effective amount of erlotinib (TKI inhibitor) and quinacrine (**76**) (NF-κB inhibitor) against erlotinib-resistant NSCLC cell lines (A549, H1975, and H1993). Results showed that the addition of quinacrine (**76**) to erlotinib treatment inhibits colony formation. Moreover, addition of quinacrine (**76**) to erlotinib treatment induced significant apoptosis in A549 and H1975 cells.

Arya *et al.* presented the synthesis of acridine derivatives (**120a–h**, **121a–d**, **122a–d**, and **123a–d**).¹⁵ Compounds (**120a–h**) were prepared by condensation of appropriate 9-aminoacridine derivatives (**118a–h**) with 9,10-dihydroanthracene-9,10- α,β -succinic anhydride (**119**) in tetrahydrofuran at room temperature (Scheme 18). To synthesize analogues (**121a–d**, **122a–d**, and **123a–d**), the authors used microwave-assisted condensation of 9-aminoacridine derivatives (**118a–d**) with phthalic anhydride (**124**) or *cis*-1,2,3,6-tetrahydrophthalimide (**125**) or 2,5-pyrroledione (**126**) (Scheme 19).¹⁵

All the obtained compounds (**120a–h**, **121a–d**, **122a–d**, and **123a–d**) were tested for *in vitro* anticancer activity against human cancer cell lines (T47D, NCI H-522, HCT-15, PA-1, HepG-2). The most active was derivative (**120a**) against T47D (IC₅₀ = 5.4 μM), (**120g**) against NCI H-522 (IC₅₀ = 4.2 μM), (**121a**) against HepG-2 (IC₅₀ = 4.5 μM) and (**123b**) against HCT-15 (IC₅₀ = 2.4 μM). The structure–activity analysis showed that compounds containing a 7- or 8-substituted acridine moiety by an electron-donating group were more active. This may be due to more effective interaction of these molecules with the DNA compared to other derivatives.¹⁵

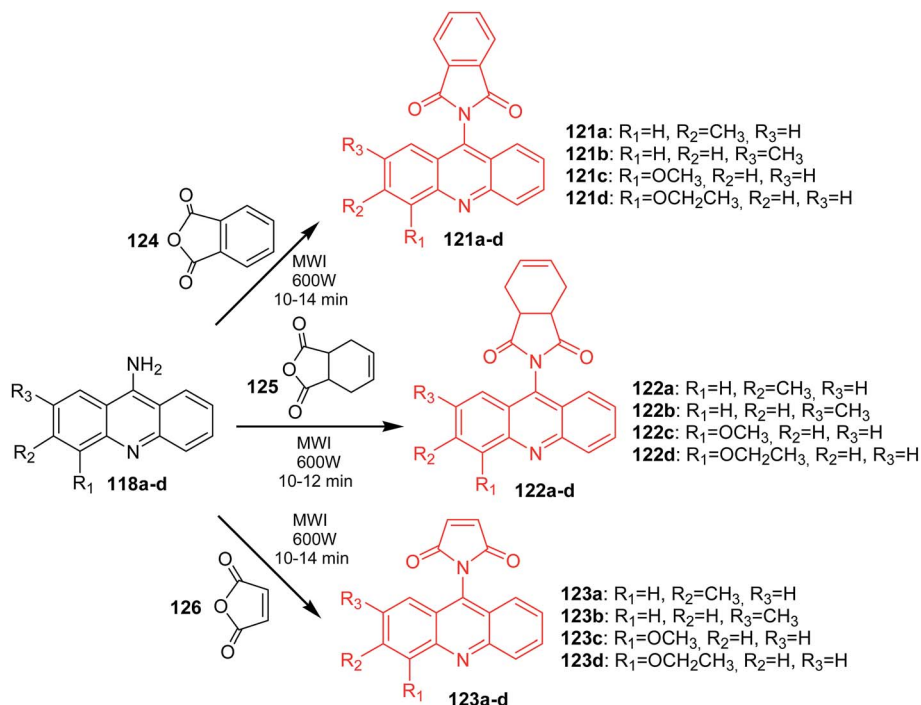
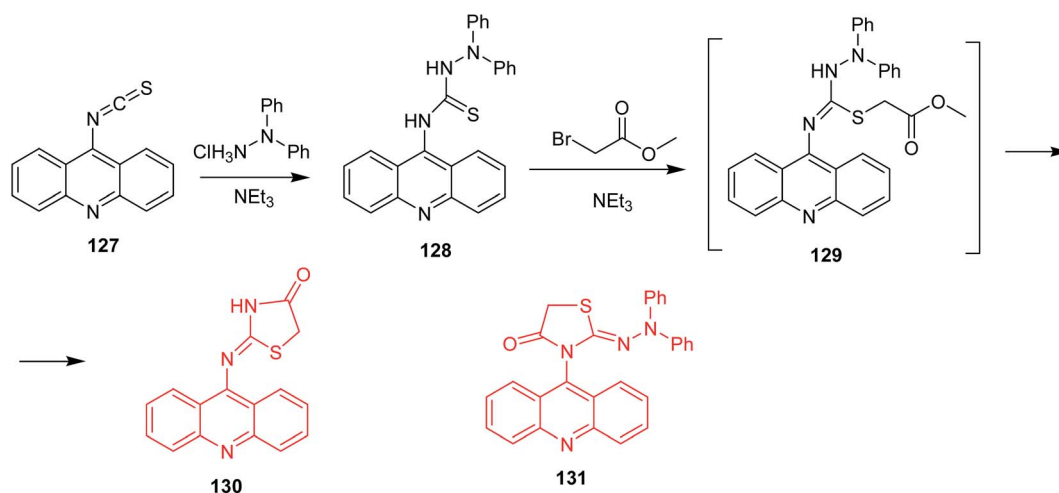
Salem *et al.* described thiazolidinone-acridines as new potential anticancer agents.⁸⁴ These compounds exhibited DNA-binding abilities and topoisomerase I and II inhibition. The most active derivative was 2-(acridin-9-yl)imino-3-diphenylamino-1,3-thiazolidin-4-one (**130**) (Scheme 20), which inhibited topoisomerase II at the concentration of 5 μM. Its synthesis included addition of 1,1-diphenylhydrazine to acridin-9-yl isothiocyanate (**127**) to form the respective thiosemicarbazide (**128**). In the next stage, the thiocarbonyl derivative underwent *S*-alkylation with methyl bromoacetate to intermediate (**129**) followed by cyclization to the final product (**130**). Of note, replacement of methyl bromoacetate by bromoacetyl bromide proceeded unselectively, and 3-(acridin-9-yl)-2-(2,2-diphenylhydrazono)-1,3-thiazolidin-4-one (**131**) was also isolated. The regioisomeric product (**131**) demonstrated lower activity against HL-60 cancer cells compared to (**130**).⁸⁴

Gao *et al.* reported the synthesis of benzimidazole acridine derivatives (**136a–p**).⁸⁵ 9-Chloroacridine derivatives (**134a–l**) were obtained by cyclization of the anthranilic acids (**133a–l**), which were synthesized by the Ullmann condensation. Then, the reaction of 2-aminomethyl benzimidazole (**135**) or compounds (**137a–b**) with the appropriate derivatives of 9-chloroacridine (**134a–l**) gave the desired final products (**136a–p**) (Scheme 21).⁸⁵



Scheme 18 Synthesis of acridine derivatives (**120a–h**).¹⁵

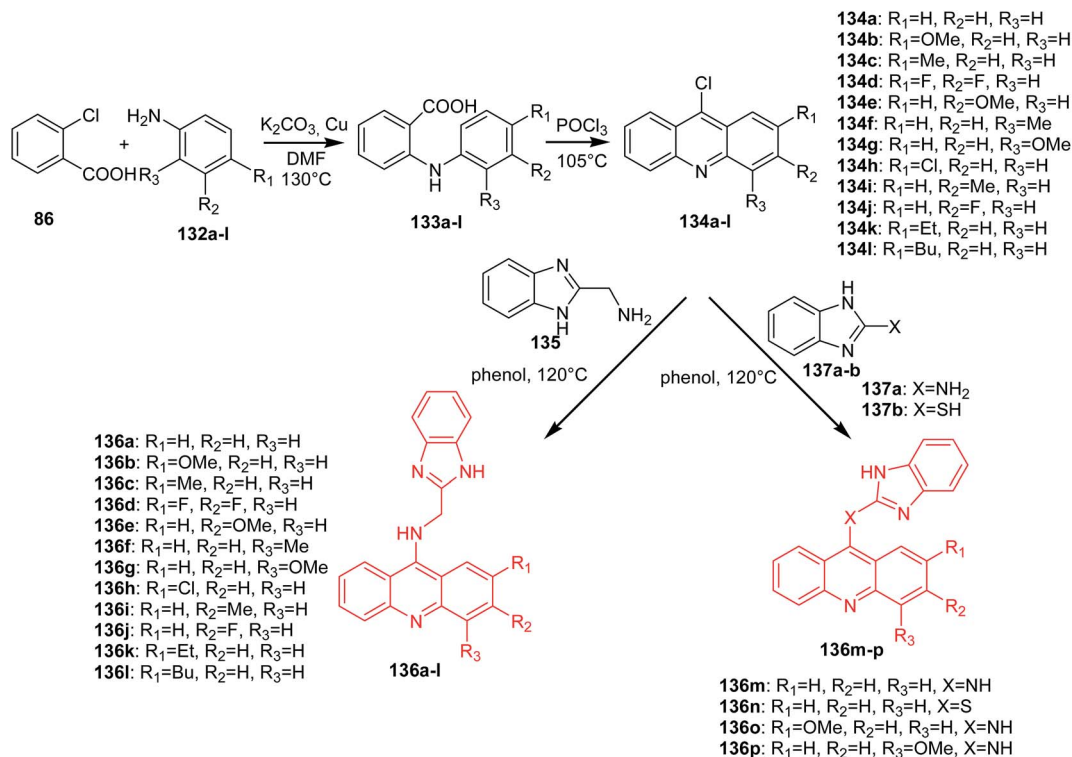


Scheme 19 Synthesis of acridine derivatives (121a-d, 122a-d, 123a-d).¹⁵Scheme 20 Synthesis of 2-(acridin-9-yl)imino-3-difenyamino-1,3-thiazolidin-4-one (130).⁸⁴

Analogues (**136a-p**) were subjected to MTT assay, which showed that most of them had good antiproliferative activity. Compound (**136l**) had the highest activity against K562 (IC₅₀ = 2.68 μM) and HepG-2 (IC₅₀ = 8.11 μM) cells. Moreover, the derivative (**136l**) inhibited topo I activity, as well as inducing apoptosis in K562 cell lines through a mitochondrial pathway.⁸¹ The structure-activity analysis showed that the substitution of the acridine moiety and the linker between the benzimidazole ring and acridine moiety had a great effect on the antitumour activity. A look at the structures of these molecules shows that substitution at position 2 of the acridine moiety with a butyl group makes this molecule (**136l**) more active than compounds

containing a 2-substituted acridine moiety by ethyl (**136k**) or methyl (**136c**) groups. In turn, analogue (**136a**), with no substituents, displayed similar activity to (**136c**), with a methyl substituent. Thus, both the size and length of substituents greatly influenced the cytotoxicity. Additionally, the electronegativity of the substituents also significantly affected the anticancer activity. For example, molecule (**136b**), with a 2-methoxy substituent, was approximately 3-fold more potent than a compound containing a 2-methyl group (**136c**) or a 2-chloro group (**136h**). Moreover, the linker between the benzimidazole ring and acridine moiety is also important in the cytotoxicity, which can be seen from the IC₅₀ value of analogues



Scheme 21 Synthesis of acridine derivatives (136a–p).⁸⁵

(136a,m,n, and o), among which derivative (136o), containing an NH-benzene-linker, showed the best anticancer activity.⁸⁵

Lang *et al.* synthesized the acridine derivatives (141a–h) as potent DNA-binding antitumour agents.³⁰ First, an Ullmann coupling of benzoic acids (86, 138) with anilines (139a–d) in DMF using Cu as the catalyst produced compounds (140a–f), which were refluxed in POCl₃ to afford the 9-chloroacridine derivatives (141a–f). The reaction of (141a–f) with substituted

benzylamines (142a–c) in the presence of KI and K₂CO₃ in absolute ethanol gave compounds (143a–h) (Scheme 22).³⁰

All synthesized compounds were assayed for inhibition of K562 leukaemia cells and hepatoma HepG-2 cells. These derivatives also exhibited good results in activity against K562 and HepG-2, among which analogue (143e) showed the best antitumour activity against both cell lines. Furthermore, fluorescence spectroscopy studies revealed that most of the

Scheme 22 Synthesis of acridine derivatives (143a–h).³⁰

derivatives can interact with DNA. Compound (**143e**) displayed the highest DNA-binding capability and inhibited topoisomerase I activity.³⁰

In turn, Gellerman described the synthesis of *N*-substituted 9-aminoacridine and bis-acridine derivatives containing electron-withdrawing groups (EWG) or electron-donating groups (EDG), including amino acid residues as potential anticancer agents.⁸⁶ These derivatives displayed good anti-tumour activity against the cancer cell lines MDM-MD-A31, MCF-7, HT29, OVCAR8, NCI-ADR, MCF-7mito and HI 299, and comparison with commercial 9-aminoacridine and *m*-amsacrine (**3**) drugs. Compound 2-(acridin-9-ylamino)-3-ethoxy-5,8-dihydroxynaphthoquinone revealed the most potent cytotoxic activity against all above mentioned cell lines than the anti-cancer drug *m*-amsacrine (**3**), probably due to enhancement of biologically important chelating properties, possibly leading to formation of more powerful DNA damaging reactive species.

Li *et al.* developed a procedure for the preparation of novel analogues of 9-amino-acridine (**146a-d**).⁸⁷ In a first step, *N*-phenylanthranilic acid (**133a**) was prepared by Ullmann reaction. Compound (**133a**) was stirred in POCl₃ to afford the 9-chloro-acridine (**134a**). Then, acridine (**134a**) by reaction with phenol was converted to 9-phenoxyacridine (**144**), which was reacted with the corresponding amines (**145**) to give derivatives (**146a-d**) (Scheme 23).⁸⁷

The synthesized compounds (**146a-d**) were tested for their antiproliferative activity against K562 and HepG-2 cells. The best activity against both cell lines was shown by derivative (**146c**) through binding with DNA and inhibiting topoisomerase I activity.⁸⁷

Roe *et al.* presented the synthesis of hybrid acridine-HSP90 ligand conjugates as telomerase inhibitors.⁸⁸ The intermediate G4-azide bearing fragments (**149a-d**) were prepared by nucleophilic coupling of compound (**147**) with the amino azides (**148a-d**) in CHCl₃ at reflux. A second intermediate GA-alkyne (**151**) was synthesized by reacting geldanamycin (**150**) with propargyl amine in DMF at room temperature. The desired final products (**153a-d**) were obtained by reaction of the corresponding G4-azide bearing fragments (**149a-d**) with

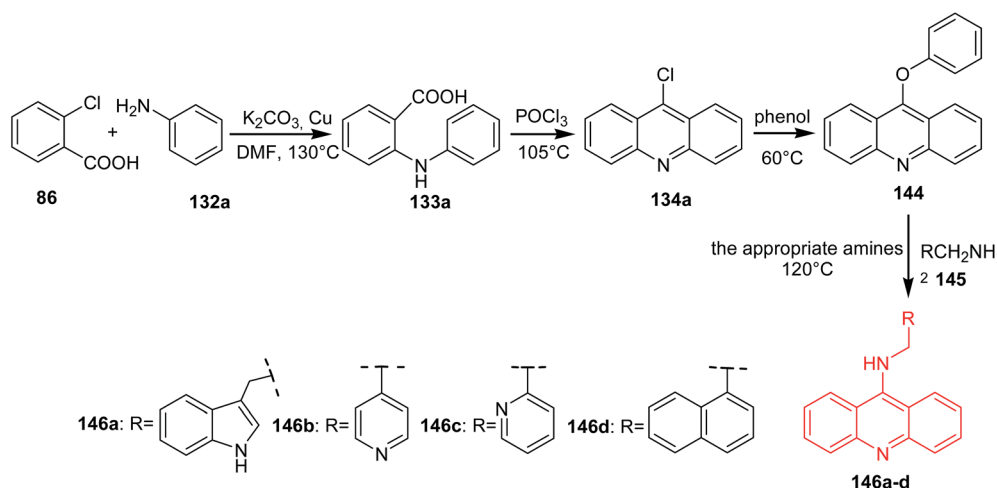
compound (**151**) using 1 : 1 *tert*-butanol : water, in the presence of CuSO₄·5H₂O as a catalyst and sodium ascorbate (**152**) (Scheme 24).⁸⁸

All synthesized compounds (**153a-d**) were assayed for inhibition of tumour cell growth including MCF7, A549, GIST48, and WI38. Derivatives (**153a-d**) showed the best activity against A549 (IC₅₀ = 0.1–0.5 μM range). Compounds (**153c-d**) containing the longest linkers (*n* = 9, and 10, respectively) revealed better potent cytotoxic activity than analogue (**153a**), with the shortest linker (*n* = 5).⁸⁸

Acridine derivatives are widely considered as potent photosensitizers for photodynamic therapy in tumour treatment. Recently, acridin-3,6-dialkylthioureas (**155**) were developed (Scheme 25), where the propyl analogue (R = *n*-Pr) revealed the highest photocytotoxicity against the L1221 cell line (IC₅₀ = 0.48 μM).⁸⁹ The mechanism of action of these compounds was also investigated, and the results strongly suggested that lysosomes of cancer cells were significant targets. Acridin-3,6-dialkylthioureas (**155**) were synthesized from 3,6-diaminoacridine (**154**) through relevant isothiocyanates.⁸⁹

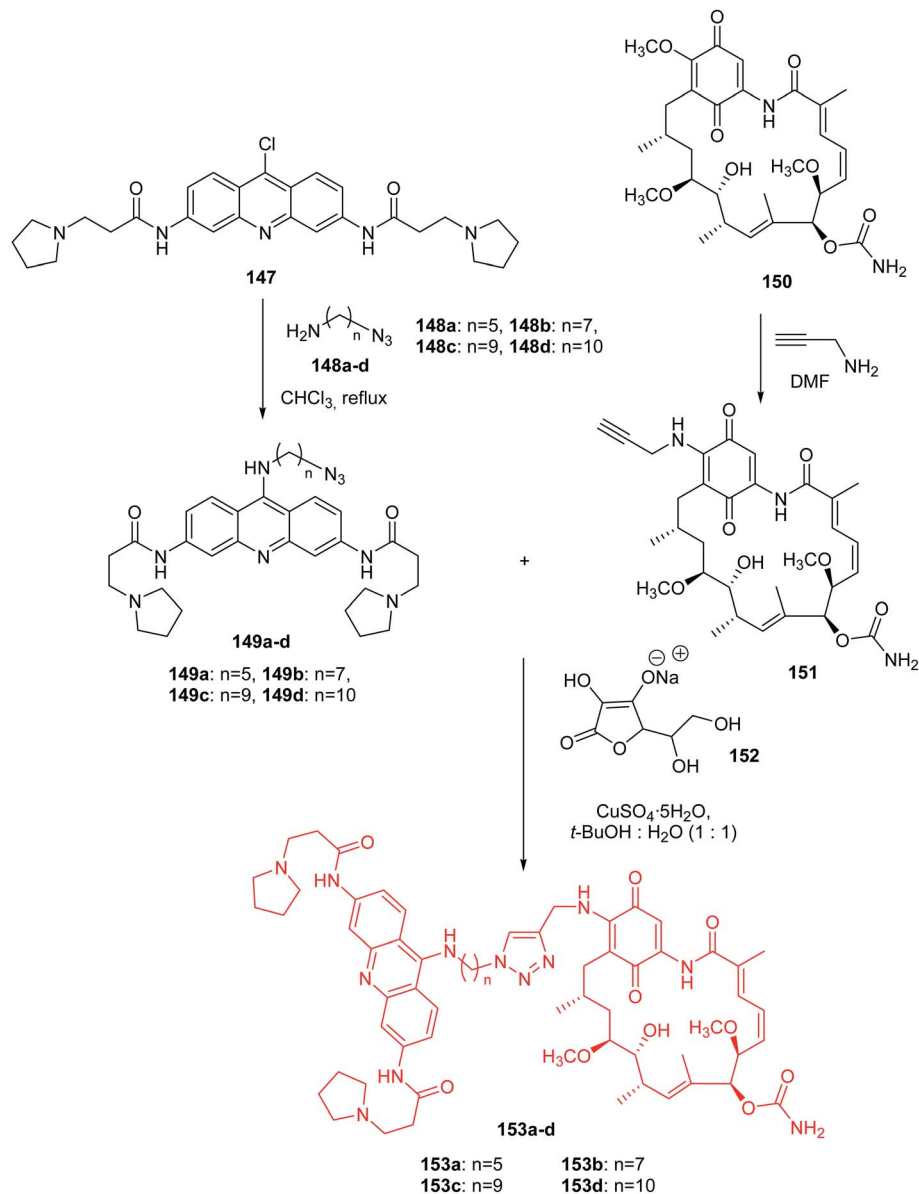
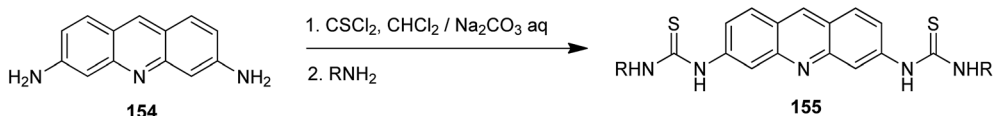
Similar 3,6-bis(3-alkylguanidino)acridines were synthesized by Plsikova and co-workers.⁹⁰ These derivatives exhibited DNA-binding abilities ($K = 1.25\text{--}5.26 \times 10^5 \text{ M}^{-1}$) and are potent DNA-intercalating antitumor agents. It was demonstrated that analogs with longer alkyl chains were able to penetrate cell membranes and suppress cell proliferation effectively. The compound with *n*-hexyl chains at nitrogen of guanidine moiety proved to be the most cytotoxic.

In turn, Huo *et al.* presented *N*-(benzo[*b*][1,10]phenanthrolin-7-yl)-2-(4-chlorobenzoyl)hydrazinecarbothioamide,⁹¹ 2-(4-methoxybenzoyl)-*N*-(2-methylacridin-9-yl)hydrazinecarbothioamide,⁹² 7-*p*-toluene-aminobenzo[*c*]acridine hydrochloride⁹³ and *N*-(4-methoxyphenyl)benzo[*c*]acridin-7-amine hydrochloride⁹⁴ as potential antitumor agents against a liver cancer, a lung cancer, a gastric cancer and various tumor cell strains. These compounds revealed strong cytotoxic activity against MGC-803 (IC₅₀ = 8.1, 11.2, 10.4 and 9.7 μg mL⁻¹, respectively), BEL-7404 (IC₅₀ = 24.2, 27.1, 23.1 and 20.3 μg mL⁻¹, respectively) and NCI-H460 (IC₅₀ = 35.3, 25.4,



Scheme 23 Synthesis of acridine derivatives (**146a-d**).⁸⁷



Scheme 24 Synthesis of acridine-HSP90 ligand conjugates (**153a-d**).⁸⁸Scheme 25 Preparation of acridin-3,6-dialkylthioureas (**155**).⁸⁹

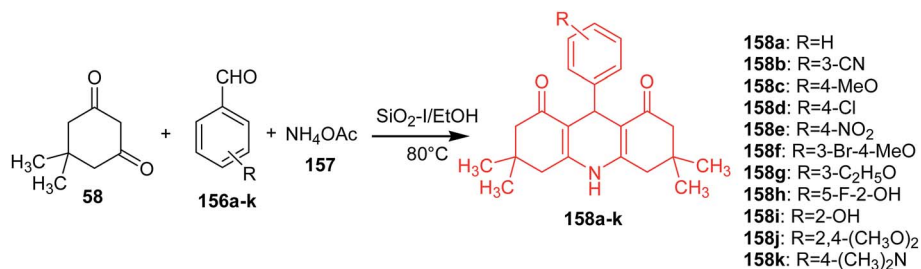
28.2 and 25.4 $\mu\text{g mL}^{-1}$, respectively). Moreover, measurements *in vivo* demonstrated that 7-*p*-toluene-aminobenzo[*c*]acridine hydrochloride and *N*-(4-methoxyphenyl)benzo[*c*]acridin-7-amine hydrochloride effectively inhibited the division of hepatocarcinoma cells and had a good inhibitory effect in mouse models of liver cancer.

Ramesh and Pasha described an efficient and economical synthesis of 9-aryl-hexahydro-acridine-1,8-diones (**158a-k**) by

cyclocondensation of dimedone (**58**), aromatic aldehydes (**156a-k**) and ammonium acetate (**157**) in the presence of a heterogeneous catalyst $\text{SiO}_2\text{-I}$ in ethanol at 80 °C (Scheme 26).⁹⁵

The obtained derivatives have shown promising antitumour activity against HepG-2 and MCF-7 cells. Compounds (**158b**, **158f** and **158j**) showed good activity against HepG-2 and MCF-7, analogue (**158i**) towards HepG-2, and the derivative (**158g**) inhibited MCF-7 cells.⁹⁵



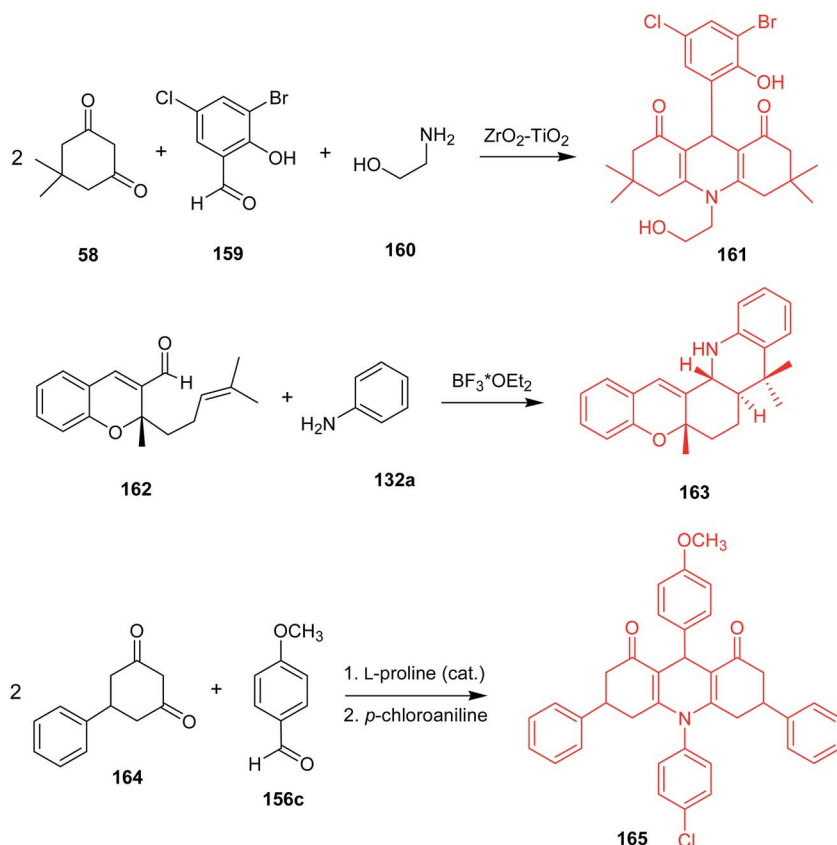


Scheme 26 Synthesis of 9-aryl-hexahydro-acridine-1,8-diones (**158a–k**).⁹⁵

Synthetic methods leading to desired acridine derivatives are still being developed and optimized. For instance, a ZrO₂-TiO₂ mixed oxide catalyst enabled a three-component, one-pot condensation of 5,5-dimethylcyclohexane-1,3-dione (**58**), 3-bromo-5-chloro-2-hydroxybenzaldehyde (**159**), and ethanolamine (**160**) to (**161**) (Scheme 27) in 99% yield.⁹⁶ Reddy *et al.* described a synthesis of trans-fused 5*H*-chromeno[2,3-*c*]acridine derivatives *via* intramolecular aza Diels–Alder reaction catalysed by Lewis acid, followed by 1,3-hydrogen shift,⁹⁷ where 5*a*,8,8-trimethyl-6,7,7*a*,8,13,13*a*-hexahydro-5*H*-chromeno[2,3-*c*]acridine (**163**) was obtained from (**162**) and aniline in the presence of BF₃·OEt₂ in 88% yield and 05 : 95 *cis/trans* ratio. Distribution of the products was in good agreement with theoretical studies.⁹⁸ Wang and co-workers reported 3,6-diphenyl-9-aryl-3,4,6,7,9,10-hexahydroacridine-1,8(2*H*,5*H*)-dione derivatives

received in the course of Knoevenagel, Michael, and cyclization reactions catalysed by *L*-proline.⁹⁹ For example, 10-(4-chlorophenyl)-9-(4-methoxyphenyl)-3,6-diphenyl-3,4,6,7,9,10-hexahydroacridine-1,8(2*H*,5*H*)-dione (**165**) was synthesized in the reaction of 5-phenylcyclohexane-1,3-dione (**164**), *p*-methoxybenzaldehyde (**156c**) and *p*-chloroaniline in an overall yield of 57%. The product (**165**) gave IC₅₀ = 82.98 μg mL⁻¹ against HepG-2 in the MTT assay.

Wang *et al.* synthesized the 12-(3,4,5-trimethoxyphenyl)benzo-*[b]*[1,3]dioxolo[4,5-*i*]acridine-6,11(5*H*,12*H*)-dione by cyclocondensation of 3,4-methylenedioxy aniline, 3,4,5-methoxybenzaldehyde, 2-hydroxy-1,4-naphthoquinone in the presence of *L*-proline in ethanol.¹⁰⁰ This molecule showed good activity against HepG-2 (IC₅₀ = 11.6 μM).



Scheme 27 Examples of newly obtained acridine derivatives **161**,⁹⁶ **163**,^{97,98} **165**.⁹⁹



Pickard *et al.* developed platinum–acridine anticancer hybrid agents, where replacement of acridine moiety by benz[*c*]acridine caused less toxicity *in vivo* in mice.¹⁰¹

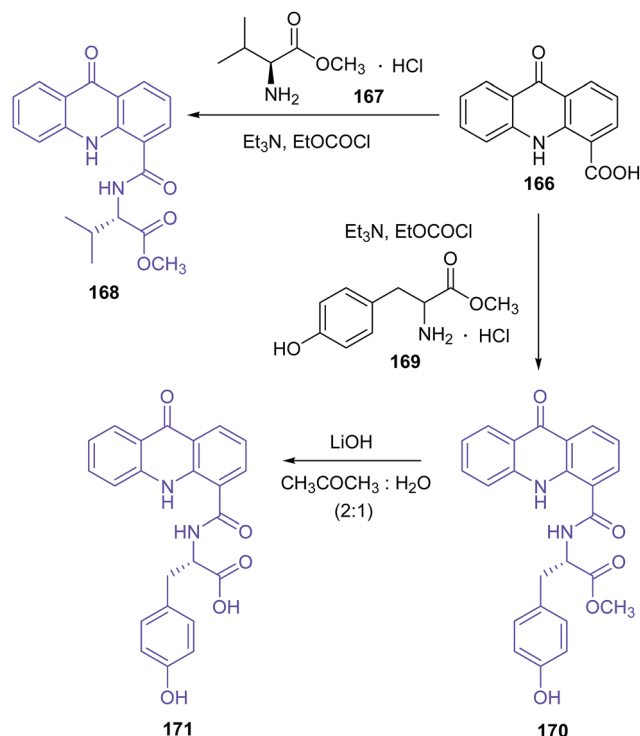
Cossio Mora *et al.* presented the synthesis of 1,2,3,4-tetrahydroacridine derivatives as histone deacetylase inhibitors (HDACs) and therapeutic agents for preventing or treating malignancies associated with aberrant histone acetylation such as cancer, hematological malignancies, proliferative diseases, neurological disorders, and immunological disorders.¹⁰² These compounds were tested *in vitro* with respect to inhibition of histone deacetylase: human isoforms HDAC1, 2, 3, 4, 5, 6, 7, 8, 9, 10 and 11 and HeLa cell line. *N*-Hydroxy-7-((1,2,3,4-tetrahydroacridin-9-yl)amino)-heptanamide and 7-((2,3-dihydro-1*H*-cyclopenta[*b*]quinolin-9-yl)amino)-*N*-hydroxy-heptanamide showed the best activity against all histone deacetylase isoforms (IC₅₀ = 0.01–1.57 and 0.007–1.28 μM, respectively) and against HeLa (IC₅₀ = 0.023 and 0.019 μM, respectively).

Belmont *et al.* described tetrahydrocyclopenta[*c*]acridine derivatives as kinase inhibitors in particular for treating cancer.¹⁰³ The 5-hydroxy-1-trimethylsilyl-3,3*a*,4,5-tetrahydro-2*H*-cyclopenta[*c*]acridin-2-one was the most effective inhibitor against CDK1 and CDK5 with IC₅₀ values of 0.56 and 1.6 μM, respectively. In turn, 5-hydroxy-9-methoxy-1-trimethylsilyl-3,3*a*,4,5-tetrahydro-2*H*-cyclopenta[*c*]acridin-2-one showed the best activity against HT 29 (IC₅₀ = 6.5 μM).

Singh *et al.* presented the synthesis of amino acid derivatives of acridone as potential leads to anticancer drugs.¹⁰⁴ Compound (168) was prepared by the reaction of the 9-oxo-9,10-dihydroacridine-4-carboxylic acid (166) with *L*-valine methyl ester hydrochloride (167) in the presence of TEA and ethyl chloroformate. A similar reaction of acridone (184) with *L*-tyrosine methyl ester hydrochloride (169) procured derivative (170). For deprotection of the ester group, analogue (170) was treated with LiOH to give compound (171) (Scheme 28). Then, derivative (171) was reacted with *L*-proline methyl ester hydrochloride (172) to provide compound (173). Deprotection of the carboxyl function yielded analogue (174), which on treatment with glycine methyl ester hydrochloride (175) gave compound (176). The final product (177) was prepared by hydrolysis of derivative (176) using LiOH (Scheme 29). Similarly, the sequential incorporation of glycine (175) and proline (172) in compound (181) yielded derivative (199).¹⁰⁴

Derivatives (186, 188, 189, 195, 196 and 199) were tested for antiproliferative activity against the C6 glioblastoma cell line (IC₅₀ = 14–20 μM range). Further, cell cycle analysis showed that compounds (195, 196 and 199) arrested C6 cells in the G0/G1 phase of the cell cycle. Moreover, analogues (195, 196 and 199) were seen to upregulate mortalin and HSP70 expression thus indicating their senescence-inducing potential.¹⁰⁴

Zhang *et al.* presented the synthesis of novel pyridyl acridones as potent DNA-binding and apoptosis-inducing agents.¹⁰⁵ Acridone-4-carboxylic acid derivatives (184*a–e*) were obtained by cyclization of the anthranilic acids (183*a–e*), which were prepared from the Ullmann condensation of 2,4-dichlorobenzoic acid (138) with various anthranilic acid derivatives (182*a–e*) in DMF using Cu as the catalyst (Scheme 30). Then, the reaction of compounds (184*a–e*) with the corresponding

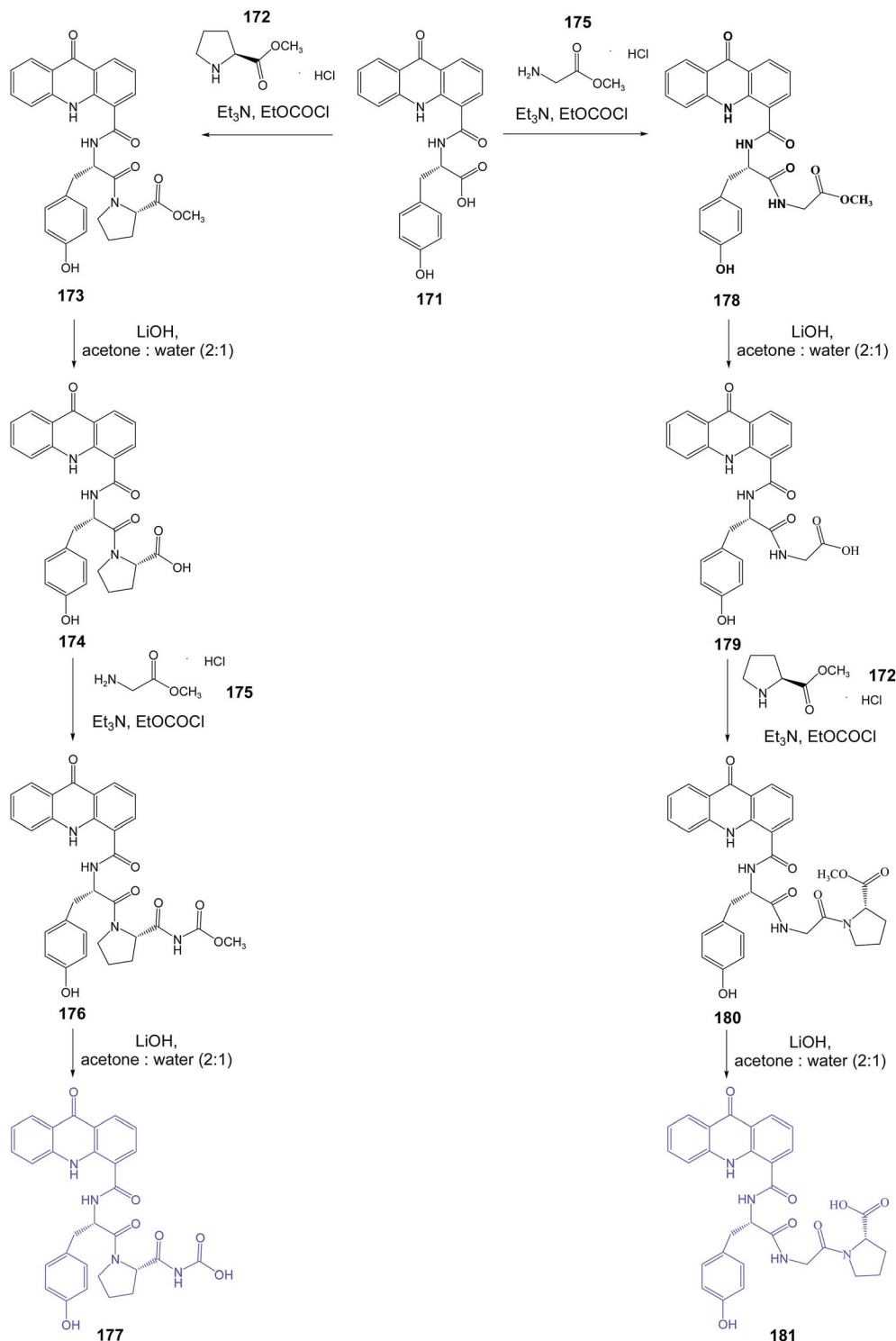


Scheme 28 Synthesis of the amino acid derivatives of acridone (168, 170, 171).¹⁰⁴

primary aliphatic amines in the presence of *N,N'*-carbonyldiimidazole (CDI) as the condensation agent gave the intermediate acridone-4-carboxamides (185*a–j*). Nucleophilic substitution of compounds (185*a–j*) with appropriate picolylamines or 4-pyridineethanamine obtained the final products (186*a–w* and 187*a–c*).¹⁰⁵

Most of the compounds (186*a–w* and 187*a–c*) exhibit good cytotoxicity against K562 cells. The best ability to inhibit K562 was shown by the compound containing a methyl group on the C5 position of acridone (186*d*) (IC₅₀ = 0.46 μM). The structure–activity analysis showed that compounds containing two methylene units between the *N,N*-dimethylamino group and 4-carboxamide group displayed better antiproliferative activity than derivatives containing three methylene units. Moreover, analogues with an *N,N*-dimethylamino group on the 4-carboxamide side chain displayed better antitumour activity than compounds with a methoxy group, which showed no cytotoxicity. To evaluate whether the number of pyridyl groups has an effect on antitumour activity (IC₅₀ = 11.2–13.9 μM range), the Zhang group introduced a 2,2'-bis pyridyl group to the acridone scaffold.¹⁰⁵ Moreover, compound (186*d*) was tested for *in vitro* anticancer activity against cancer cell lines including NCI-H520, U251, A375, A172, HeLa, CNE-2, U118-MG, HepG-2 and MCF-7 cells. Testing results showed that (186*d*) has broad antitumour activities. Derivative (186*d*) showed the best antitumour activity against A375 and HepG-2 cells with IC₅₀ values equal to 0.16 and 0.32 μM, respectively. Furthermore, spectrophotometric methods, viscosity measurements and topo I inhibition assays demonstrated that (186*d*) interacts with DNA, inhibits





Scheme 29 Synthesis of the dipeptide derivatives of acridone (177, 181).¹⁰⁴

topo I activity and induces apoptosis through the mitochondrial pathway.¹⁰⁵

The Gao group synthesized new derivatives of 10-(3,5-dimethoxy)benzyl-9(10*H*)-acridone (192a–g).¹⁰⁶ Starting from the commercially available acridone (111) by the nitration reaction gave 2,7-dinitro-9(10*H*)-acridone (188), which in the presence of 3,5-dimethoxybenzyl was converted to 2,7-dinitro-10-(3,5-

dimethoxy)benzyl-9(10*H*)-acridone (189). Then, the reduction of the nitro group by Na₂S·9H₂O in ethanol gave 2,7-amino-10-(3,5-dimethoxy)benzyl-9(10*H*)-acridone (190). Subsequently, the derivative of acridone (190) was acylated with 3-chloropropionyl chloride to obtain 2,7-bis(3-chloropropionamido)-10-(3,5-dimethoxybenzyl)-9,10-dihydroacridone (191). As a result of nucleophilic substitution of compound (191) with an



Scheme 30 Synthesis of pyridyl acridones (**186a-w**, **187a-c**).¹⁰⁵

appropriate amine, desirable acridone derivatives (**192a-g**) were synthesized (Scheme 31).¹⁰⁶

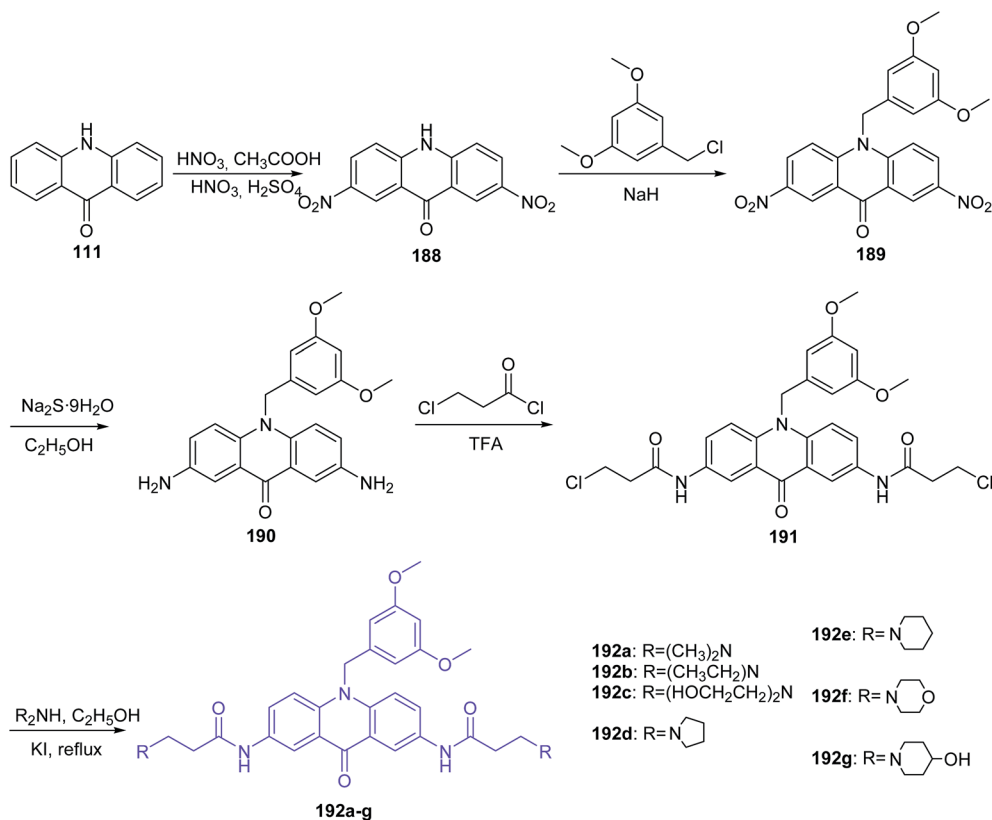
The final products (**192a-g**) were evaluated for anti-proliferative activity. The highest activity (IC₅₀ = 0.3 μM) against leukaemia CCRF-CEM cells was shown by compound (**192a**), with dimethylamine substituents at the termini of the chains. Compounds (**192b** and **192c**), containing diethylamine and diethanol amine groups, revealed a decreased inhibitory effect. Furthermore, (**219a**) demonstrated low toxicity against 293T cells (IC₅₀ > 100 μM).¹⁰⁶

In search of anticancer compounds, Gao and co-workers developed amino acid acridone analogues possessing a 3,5-dimethoxyphenyl moiety, and the most potent one was *N*-(10-(3,5-dimethoxybenzyl)-9,10-dihydro-9-oxoacridin-2-yl)-2-aminoacetamide hydrochloride (**194**) (Scheme 32).^{107,108} Analogue (**194**) exhibited significant activity against CCRF-CEM, A549, HepG-2, and MCF-7 cells together with low cytotoxicity to normal cells. The authors have more recently reported more detailed data on the mechanism of action of this compound against CCRF-CEM leukaemia cells, and 55 proteins were identified as involved in the proteomic processes.

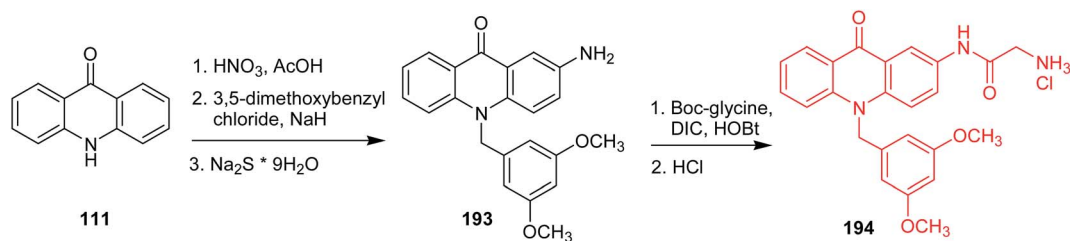
According to the results, cell death can be caused by multiple mechanisms, such as inhibition of aerobic glycolysis, mitochondrial oxidative phosphorylation, DNA damage, and oxidative stress.¹⁰⁸ The synthetic pathway included nitration of acridone (**111**) followed by *N*-alkylation with 3,5-dimethoxybenzyl chloride and reduction to the respective amine. Then, 2-amino-10-(3,5-dimethoxy)-benzyl-9(10*H*)-acridone (**193**) was coupled with *N*-(*tert*-butoxycarbonyl)glycine in the presence of *N,N'*-diisopropylcarbodiimide (DIC) and 1-hydroxybenzotriazole (HOBt). In the last stage, the target compound (**215**) was produced in the course of treatment of the respective Boc-protected derivative with HCl.¹⁰⁷

Prasad *et al.* presented the synthesis of nitric oxide-releasing acridone carboxamide derivatives as potential anticancer agents.¹⁰⁹ Acridone-2-carboxylic acid (**197**) was prepared by cyclization of the 2,4'-iminodibenzoic acid (**196**), which was synthesized by the Ullmann condensation reaction of 2-chlorobenzoic acid (**86**) with *p*-aminobenzoic acid (**195**). Then, acridone-2-carboxamides (**198a-f**) were obtained by the reaction of compound (**197**) with an appropriate amine. In turn, derivatives (**199a-d**) were synthesized by stirring acridone-2-carboxamides with the alkylating agent





Scheme 31 Synthesis of 10-(3,5-dimethoxybenzyl)-9(10H)-acridone derivatives (192a–g).¹⁰⁶



Scheme 32 Synthesis of *N*-(10-(3,5-dimethoxybenzyl)-9,10-dihydro-9-oxoacridin-2-yl)-2-aminoacetamide hydrochloride (194).¹⁰⁷

1-bromo-3-chloropropane or 1-bromo-4-chlorobutane in the presence of the strong base sodium hydride. Subsequently, compounds (199a–d) had introduced the nitric oxide-donating moiety by using silver nitrate in dry acetonitrile to afford the NO-donating acridone carboxamides (200a–d) (Scheme 33).¹⁰⁹

All synthesized compounds were assayed for inhibition of the human breast cancer cell line MCF7 and its two drug cross-resistant phenotype sublines (P-gp and BCRP), doxorubicin-resistant MCF7/Dx (P-gp expression) and mitoxantrone-resistant MCF7/Mr (BCRP expression) cell lines. Compounds fused with NO donors (200a–d) revealed the most potent cytotoxic activity against MCF7, MCF7/Mr and MCF7/Dx cells with IC_{50} values varying from 0.8 μM to 1.6 μM , 1.7 μM to 2.7 μM and 1.9 μM to 4.2 μM , respectively. Moreover, derivatives (200b–d) were tested for colorectal cancer cell line (SW1398, WiDr and LS174T) activity and showed potent, selective inhibition (IC_{50} =

1.7–11 μM against SW1398, 2.8–19.1 μM against WiDr and 3.1–18.2 μM against LS174T).¹⁰⁹

The Mohammadi-Khanaposhtani group presented the synthesis of new 9(10H)-acridinone-1,2,3-triazole derivatives (207a–n) against human breast cancer cell lines and apoptosis-inducing agents.¹¹⁰ The intermediate acridone derivatives (203) were prepared by cyclization of 2-arylaminobenzoic acids (202), which were obtained from the Ullmann condensation reaction of 2-bromobenzoic acid (37) with the corresponding aniline derivatives (201). Then, compounds (203) were converted to the 10-(prop-2-yn-1-yl)acridin-9-one derivatives (204) using propargyl bromide in the presence of potassium *tert*-butoxide in DMSO at room temperature. The final products (207a–n) were prepared by the reaction of compounds (204) with *in situ* prepared 1-(azidomethyl)-4-methoxybenzene derivatives (206) from chloride (205) (Scheme 34).¹¹⁰



All prepared compounds (**207a–n**) were assayed for inhibition of three human breast cancer cell lines, MCF-7, T-47D, and MDA-MB-231. Compounds (**207a–g**) with mono-methoxy substituents on the pendant benzyl group of the 1,2,3-triazole ring displayed better antitumour activity ($IC_{50} = 11.0–60.0 \mu\text{M}$) than derivatives (**207h–n**) with tri-methoxy groups, which showed no cytotoxicity ($IC_{50} > 100 \mu\text{M}$). Analogue (**207c**), with a methoxy group at the 2-position, showed the best activity against MCF-7 ($IC_{50} = 11.0 \mu\text{M}$), T-47D ($IC_{50} = 14.5 \mu\text{M}$) and MDA-MB-231 ($IC_{50} = 16.6 \mu\text{M}$). Moreover, acridine orange/ethidium bromide and Annexin V-FITC/propidium iodide (PI) double staining showed that derivative (**207c**) can induce apoptosis.¹¹⁰

Kumar and co-workers synthesized new N10-substituted acridone derivatives (**209a–n**, **210a–b**, **211a–b**, and **212a–b**).¹¹¹ Compounds (**209a–n**) were obtained by reaction of the appropriate alkyl halides in aqueous NaOH in the presence of TBAB and 2-butanone with acridin-9(10H)-one (**111**) or 2-fluoroacridin-9(10H)-one (**208**) (Scheme 35). Analogues (**210a–b**) were prepared by reaction of (**111**) or (**208**) with hydroxylamine hydrochloride and sodium acetate in water (Scheme 35), while derivatives (**211a–b**) were synthesized in two steps. Reacting (**111**) or (**208**) in DMF and NaH with ethyl bromoacetate produced compounds (**211a–b**). Further hydrolysis of (**211a**) or (**211b**) in dry ethanol, sodium hydroxide in water, and then by acidification HCl yielded derivatives (**212a–b**) (Scheme 35).¹¹¹

The cytotoxic effect of analogues (**209a–n**, **210a–b**, **211a–b**, and **212a–b**) on the four cell lines (MCF-7, A-549, HT-29, and

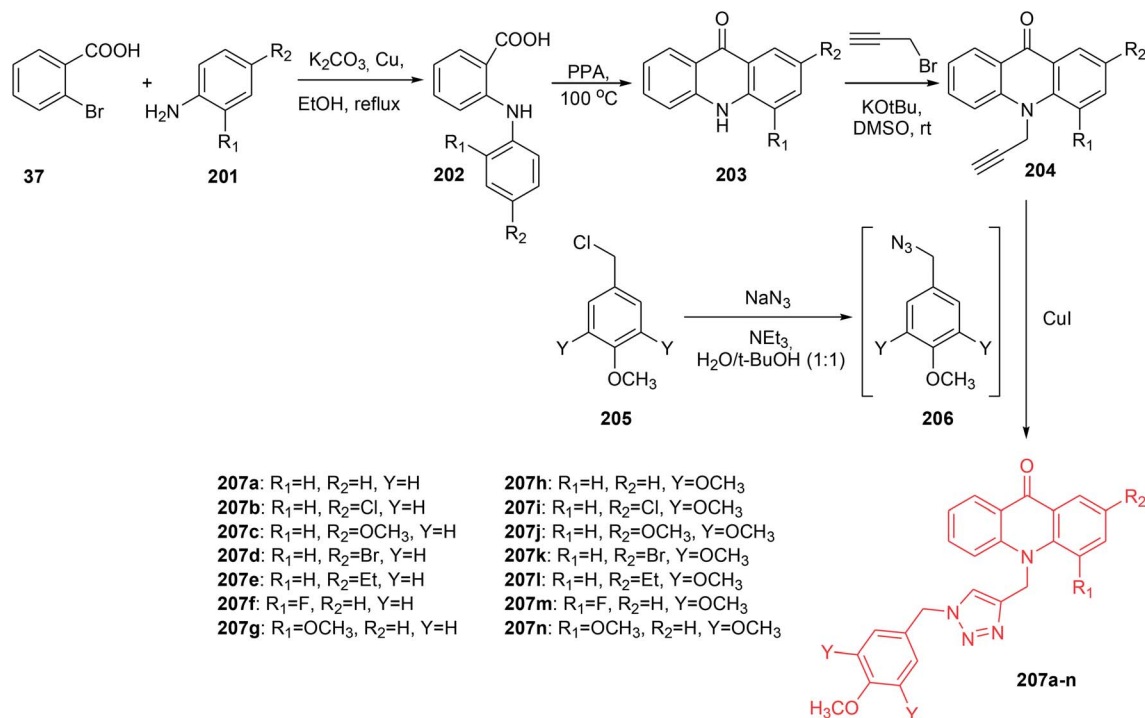
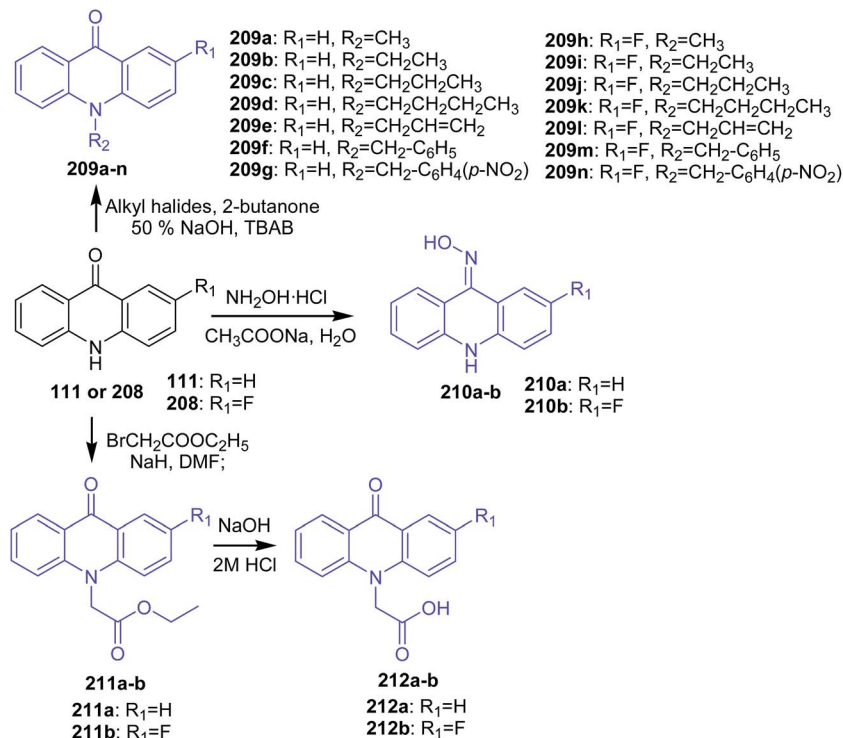
HeLa) were assessed by MTT assay. Compound (**209n**), containing a *p*-nitrobenzyl group at the N10 position and a fluoro group at the C2 position, revealed good cytotoxic activity against MCF-7, A-549, and HeLa with IC_{50} values equal to 57.4, 43.1, and 39.5 μM , respectively. In turn, derivative (**209h**), with a methyl group at the N10 position of 2-fluoroacridone, showed moderate inhibitory effect, and analogues containing higher alkyl, unsaturated alkyl, ester, and oxime groups were not active. Thus, the N10 position and the C2 position of the acridone ring had a significant influence on the antiproliferative activity of the compounds. Furthermore, the authors conducted *a silico* study to check the multi-drug resistance modulator (MDR) effect of synthesized derivatives using docking simulations in the ATP binding site of P-gp. Derivatives (**209g** and **209n**) showed similar interactions. These analogues (**209g** and **209n**) showed a hydrogen bond between the carbonyl oxygen at the 9th position of the acridone ring with Tyr1393 and a π - π stacking interaction between the benzyl ring substituted at the 10th position of the acridone moiety with Tyr2352. Furthermore, the nitro group of (**209g** and **209n**) also showed weak non-covalent interactions with positively charged magnesium ion (Mg^{2+}). Analogues (**209f**, **209g**, **209m**, **209n**, and **212a–b**) showed considerable interactions into the ATP binding site of P-gp and can therefore be considered to display MDR modulatory activity as well.¹¹¹

Kumar *et al.* presented the synthesis of 2-(9-oxoacridin-10(9H)-yl)-*N*-phenyl acetamides.¹¹² 2-(9-Oxoacridin-10(9H)-yl)-*N*-phenylacetamide derivatives (**215a–o**) were prepared by the



Scheme 33 Synthesis of nitric oxide releasing acridone carboxamide derivatives (**200a–d**).¹⁰⁹

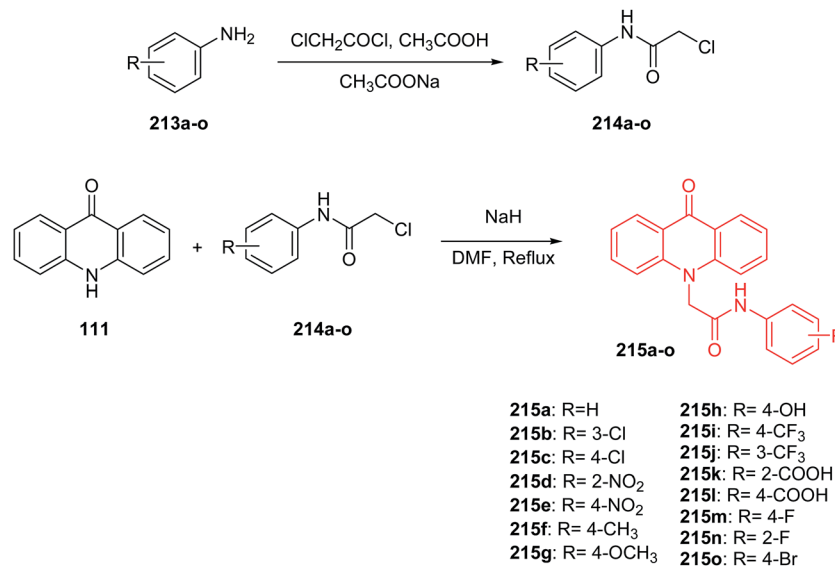


Scheme 34 Synthesis of compounds (207a–n).¹¹⁰Scheme 35 Synthesis of N10-substituted acridone derivatives (209a–n, 210a–b, 211a–b, 212a–b).¹¹¹

condensation reaction of acridone (**111**) with 2-chloro-*N*-phenylacetamides (**214a–o**) in the presence of sodium hydride in DMF at reflux (Scheme 36).¹¹²

Compounds (**215a–o**) were tested for *in vitro* anticancer activity against three human cancer cell lines (MCF-7, HeLa, and A-549). Derivatives a with carboxylic group at position 2 or 4





Scheme 36 Synthesis of intermediate compounds (214a-o) and final target compounds (215a-o).¹¹²

of the phenyl ring (215k and 215l) showed the greatest inhibition ability against MCF-7 ($\text{IC}_{50} = 6.07$ and $7.12 \mu\text{M}$, respectively), HeLa ($\text{IC}_{50} = 8.80$ and $10.66 \mu\text{M}$, respectively) and A-549 ($\text{IC}_{50} = 7.39$ and $8.80 \mu\text{M}$, respectively). Moreover, studies performed showed that analogues (215k, 215l, 215d, 215e and 215h), bearing electron-withdrawing groups ($-\text{COOH}$, $-\text{NO}_2$, $-\text{OH}$), exhibited better cytotoxic activity compared to unsubstituted (215a) or compound (215f) bearing electron-releasing groups ($-\text{CH}_3$). Furthermore, the computational analysis showed that all prepared products display significant interactions with the ATP binding site, and they may therefore serve as MDR modulators.¹¹²

Conclusion

Bacterial, parasitic and viral infections and above all neoplastic diseases are still a serious challenge for scientists. Acridine/acridone analogues constitute a group of compounds with antibacterial, antiparasitic, anti-malarial, anti-HIV and anti-tumour properties. However, the clinical usefulness of acridine/acridone derivatives is limited due to the risk of high toxicity and tumour resistance. Recent research has opened new directions for the design and development of new and more potent inhibitors of acetylcholinesterase, inhibitors of human carbonic anhydrase isozymes and antibacterial, antiparasitic and anticancer drugs.¹¹³⁻¹¹⁷

The described compounds showed potent activity against hAChE ($\text{IC}_{50} = 3-8 \text{ nM}$) and their derivatives against hBChE ($\text{IC}_{50} = 0.4-20 \text{ nM}$). These analogues were more active than the positive control tacrine,⁴⁶ while compound (44g) (Scheme 2) showed activity against AChE with an $\text{IC}_{50} = 7.31 \mu\text{M}$.⁴⁹

Synthesized acridine derivatives have exhibited good results in activity against isoforms (hCA I, II and IV).⁶¹ Also described are acridine derivatives as antimalarial agents.⁶⁸ A novel inhibitor of *M. tuberculosis* DNA gyrase was prepared that

showed good activity with IC_{50} values in the range of 5.2 to $33.9 \mu\text{M}$.⁷⁰ Derivative (100k, Scheme 14) exhibited more effective activity against *S. enteritidis* than erythromycin,⁷² while derivative (103b, Scheme 15) exhibited higher activity against *Candida albicans* than Rivanol.⁷³

Compounds (121a, 136l, 143e, 146c, 158b, 158f, 158i, 158j, 186d) showed promising antitumour activity against HepG-2 (ref. 15, 30, 85, 87, 95 and 105) Additionally, derivatives (136l, 143e, 146c, 186d) exhibited good activity results against K562 through binding with DNA and inhibiting topo I activity.^{30,85,87,105} Furthermore, analogue (192a) showed the highest activity ($\text{IC}_{50} = 0.3 \mu\text{M}$) against leukaemia CCRF-CEM cells.⁸⁹ Derivatives (153a-d, Scheme 24) showed the best activity against A549 ($\text{IC}_{50} = 0.1-0.5 \mu\text{M}$)⁸⁰ and (200a-d) (Scheme 33) revealed the most potent cytotoxic activity against MCF7 ($\text{IC}_{50} = 0.8-1.6 \mu\text{M}$).⁹²

PEGylated polyacridine peptides are considered as non-viral gene delivery systems.¹¹⁸ Khargharia *et al.* reported synthesis of DNA binding polyacridine peptides followed by their *in vivo* gene expression in mice. Length of polyethylene glycol (PEG) moiety influenced DNA polyplexes administration dramatically, whereas its location occurred to be not so crucial. Investigation of the relationship between structure of PEGylated polyacridine peptides and pharmacokinetics, biodistribution and metabolism can lead to novel compounds in gene therapy.

Thus, the acridine/acridone moiety can still serve as a lead structure and can be further modified to increase the specificity of new analogues towards tumour cells and bacterial, parasitic and viral infections.

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