Hybrid Azine Derivatives

Subjects: Chemistry, Medicinal

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Infectious diseases caused by microorganisms are a major threat to human health, mostly because of drug resistance, multi-drug resistance and extensive-drug-resistance phenomena to microbial pathogens. During the past, obtaining hybrid azaheterocyclic drugs represents a powerful and attractive approach in modern antimicrobial therapy with very promising results including overcoming microbial drug resistance.

hybrid compounds antimicrobial pyridine quinoline

1. Introduction

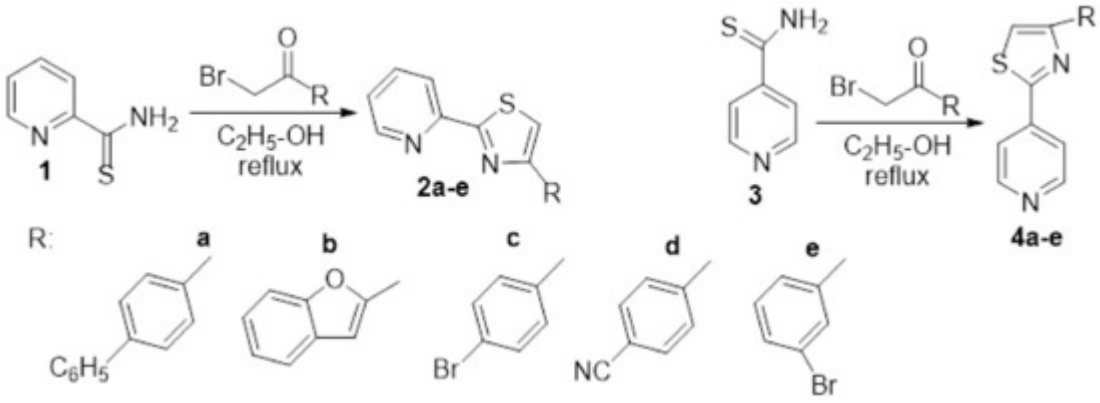
According to the WHO, infectious diseases caused by microorganisms represent a major threat that affects society and human health, exerting great pressure on health systems, individuals and communities [1]. In particular, overconsumption and widespread use and misuse of antimicrobial agents have resulted in the emergence of drug resistance, multi-drug resistance and extensive-drug-resistance phenomena to microbial pathogens and many other drawbacks (toxicity and non specificity of drugs, high prices, etc.). So far, searching for new chemical entities with improved antimicrobial properties remains a very challenging and important task in medicinal chemistry.

During the last few years, molecular hybridization represents a powerful tool in drug design, by merging two or more drug pharmacophores in a single hybrid multi-functional molecule. Usually, the resulting hybrid entity has superior properties compared with conventional classic drugs, with dual or multiple target mechanisms, better biological activity and specificity, less side effects and toxicity, less drug–drug interactions, etc. [2][3]. As a result of this approach, important advances have been achieved in antimicrobial therapy, some of the present drugs from the market have a hybrid structure and some hybrid structures are in different clinical trials [2][3][4][5][6][7][8].

A literature survey revealed that azines are privileged scaffolds in current medicinal chemistry and drug discovery, possessing a large variety of biological activities, such as: antibacterial, antifungal, antiplasmodial and antimalarial, anthelmintic, antitubercular, antiviral, anticancer, anti-inflammatory, antihypertensive, diuretic, antithrombic, anticoagulant, antidepressant, anxiolytic, anticonvulsant, analgesic, antiulcer, antidiabetic, antihistaminic, etc. [4][5][6][7][8]. As a matter of fact, the greatest majority of the existing drugs from the market contain in their structure a nitrogen heterocycle, some of them being a hybrid structure, which justifies the demand of the pharmaceutical industry for such drugs with nitrogen heterocycle skeleton.

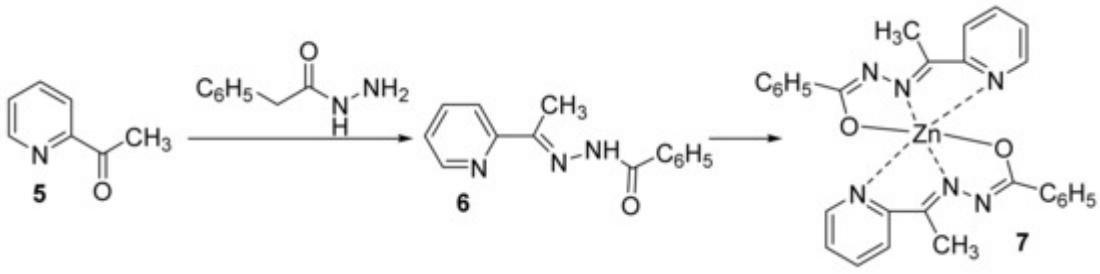
2. Six-Member Ring Azaheterocycles with One Nitrogen Atom. Hybrid Pyridine

In their attempt to identify new antimicrobial compounds, Eryılmaz et al. [9] designed and synthesized different hybrid pyridine derivatives bearing in the 2- and 4-position of the ring of a thiazole moiety. The synthesis was straight and efficient, involving a Hantzsch cyclocondensation of pyridine-2- and 4- carbothioamide **1** and **3** with acetophenone derivatives, when the desired hybrid 4-(*R*-2-yl)-2-(pyridin-2-yl)thiazole **2a**–**e** and 4-(*R*-2-yl)-2-(pyridin-4-yl)thiazole **4a**–**e** are obtained, Scheme 1. The synthesized compounds were tested for their antibacterial activity [four strains, *Gram-positive* (*Bacillus cereus*, *Staphylococcus aureus*) and *Gram-negative* (*Escherichia coli*, *Pseudomonas aeruginosa*)] and antifungal activity (one strain, *Candida albicans*) *via* minimal inhibitory concentration (MIC) method and DNA cleavage activity studies. The researchers established interesting correlation structure-biological activity (SAR), the most relevant finding being that 4-pyridine thiazole hybrid compounds **4a**–**e** showed more potent activity than **2a**–**e**. The most promising compound was found to be **4c** (MIC values 0.01 mM) exhibited on the bacterial strains *Staphylococcus aureus* and *Bacillus cereus*.



Scheme 1. Reaction pathway to obtain hybrid thiazole-pyridine **2a**–**e** and **4a**–**e**.

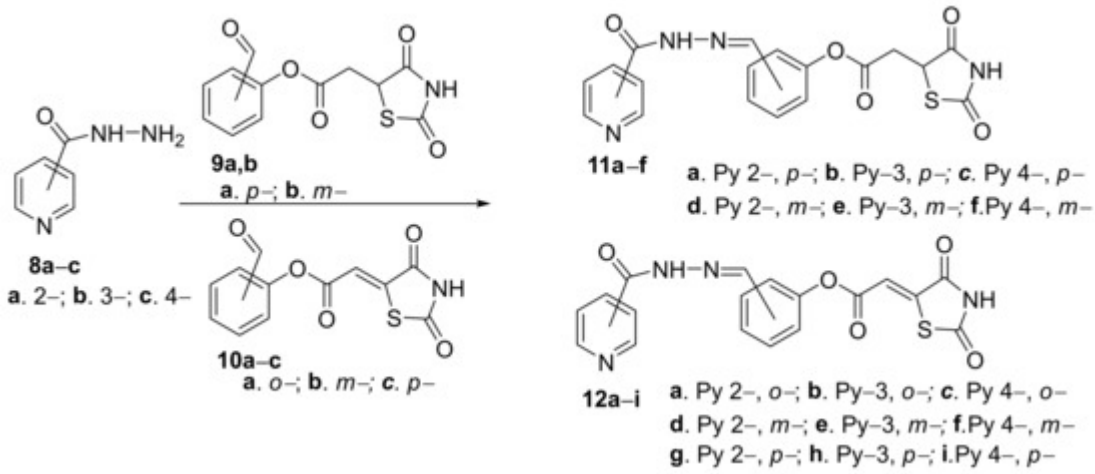
In a subsequent paper, some of the above researchers (Cinarli et al. [10]) synthesized different hybrid aroylhydrazone-pyridine-metal derivatives. The newly hybrid aroylhydrazone-pyridine metal derivatives [ZnL_2] **7** have been synthesized in two steps: an initial cyclocondensation of pyridine-2-acyl derivative **5** (with aroylhydrazone leading to pyridine-aroylhydrazone ligand **6**) is followed by complexation with M^{2+} metal (Zn^{2+}), Scheme 2.



Scheme 2. Reaction pathways to obtain hybrid metal-pyridine derivatives **7**.

The synthesized compounds were tested for their antibacterial activity (four strains, *Pseudomonas aeruginosa*, *Escherichia coli*, *Bacillus cereus* and *Staphylococcus aureus*) and antifungal (one strain, *Candida albicans*) activity *via* minimal inhibitory concentration method. The [ZnL_2] **7** has been found to be more active than pyridine-aroylhydrazone ligand **6** in all microorganisms (MIC = 11.71 µg/mL for bacteria and MIC = 23.43 µg/mL for *C. albicans*). The researchers claim that the synthesized new complex acts on microorganisms by disrupting the cell wall structure. The DNA binding interactions was also determined experimentally by spectrophotometric and electrochemical methods. The obtaining data indicate that ligand **6** and hybrid [ZnL_2] **7** interact the most with guanine base, and charge transfer is from DNA guanine bases to the molecular structures. Moreover, antioxidant activity was determined, and the hybrid [ZnL_2] **7** acted as a scavenger against peroxide radicals.

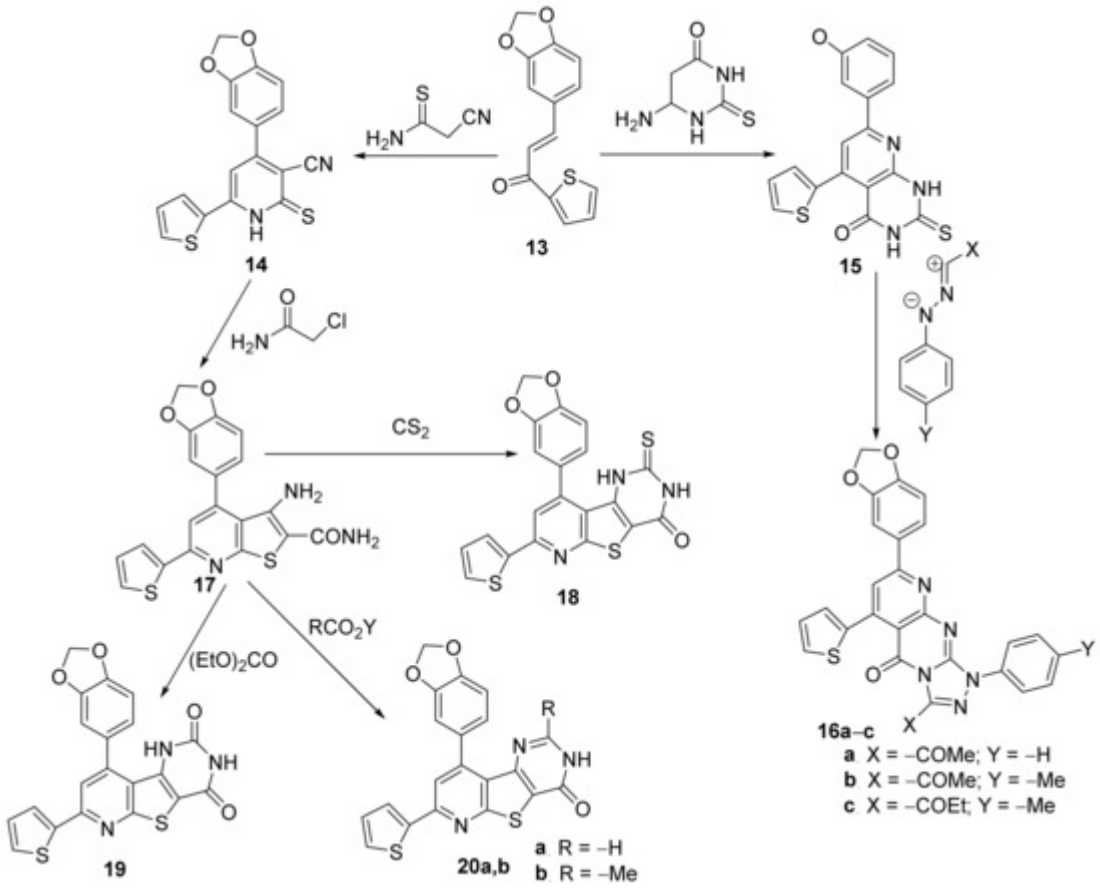
Trotsko et al. [11] designed and synthesized different hybrid pyridine derivatives bearing at the 2-, 3- or 4- position of the ring of a thiazolidine-2,4-dione moiety. The synthesis involve a condensation reaction of hydrazonyl-pyridine **8a**–**c** with the corresponding (2,4-dioxo-1,3-thiazolidin-5-yl/ylidene) **9a**,**b**/**10a**–**c**, which are leading to the desired hybrid pyridine-2,4-dioxo-1,3-thiazolidin-5-yl derivatives **11a**–**f** or pyridine-2,4-dioxo-1,3-thiazolidin-5-ylidene derivatives **12a**–**i**, Scheme 3.



Scheme 3. Reaction pathway to obtain hybrid thiazolidine-pyridine **11a**–**f** and **12a**–**i**.

The in vitro antimycobacterial assay (*Mycobacterium tuberculosis*) of the newly obtained compounds reveals strong activity in the concentration range of 1–512 µg/mL and low cytotoxicity. Interesting SAR correlations have been performed, and the highest antimycobacterial activity (MIC = 1 µg/mL) was demonstrated for the hybrid pyridine derivatives bearing the thiazolidine-2,4-dione moiety at the 4-position of the pyridine ring (hybrids **11a**–**c** and **12g**–**i**).

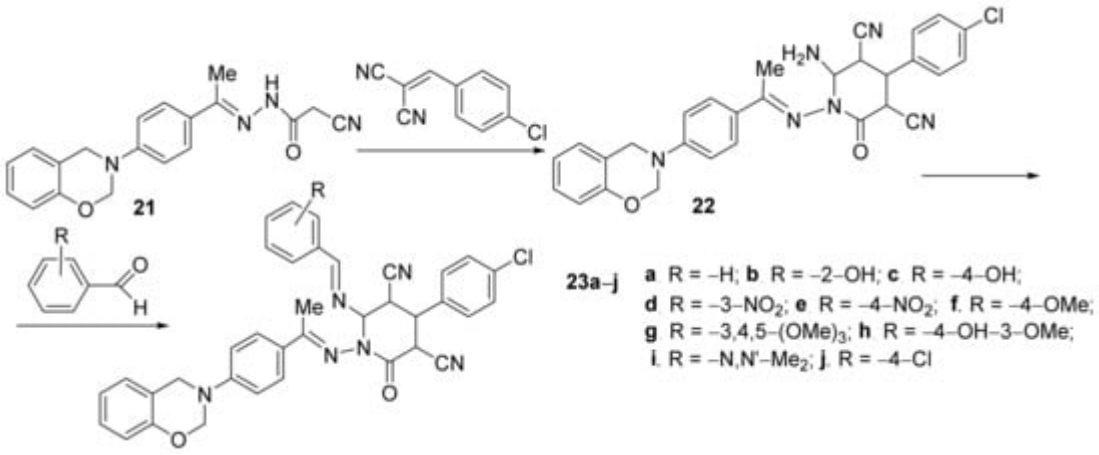
Sanad et al. [12] have performed an interesting study concerning the in vitro antimicrobial activity of some newly hybrid thieno-pyrimidin-pyridine derivatives. The synthesized compounds belonged to different classes of substituted pyridine: thiophen-dihydropyridine **14**, thiophen-pyrido-pyrimidin-4(1*H*)-one **15**, and fused pyridine: pyrido-thiophen-triazolo-pyrimidine **16a**–**c**, thiophen-pyrido-thieno derivative **17**, thiophen-pyrido-thieno-pyrimidin-4-one **18**, thiophen-pyrido-thieno-pyrimidin-2,4-dione **19**, thiophen-pyrido-thieno-pyrimidin-2-*R*-4-one **20**, Scheme 4.



Scheme 4. Reaction pathway to obtain hybrid thiophen-pyrimidin-pyridine **14**–**20**.

The synthetic approach is straight and efficient, involving typical organic chemistry reactions, mostly cyclocondensations. The synthesized compounds were tested in vitro for their antibacterial activity against *Escherichia coli* and *Klebsiella pneumoniae* as *Gram-negative* bacterial strains as well as against *Staphylococcus aureus* and *Streptococcus mutans* as *Gram-positive* bacterial strains. The obtained results (expressed as the diameter of inhibition zones (DIZ) and MIC) reveal that the thiophen-pyrido-thieno-pyrimidin-2-*R*-4-one **20a**,**b** exhibit the strongest antibacterial activities against all the tested bacteria, in the range of 40–60 mm for inhibition zones, respectively, 4–16 µg/mL for MIC values.

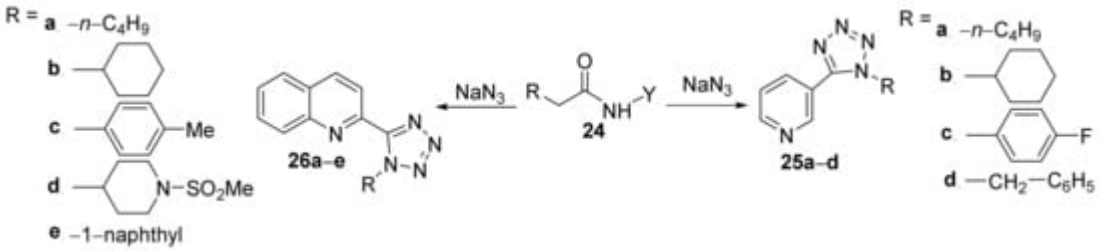
Desai et al. [13] have studied the in vitro antimicrobial activities of some newly hybrid oxazino-pyridine derivatives. The desired compounds, oxazin-3(4*H*)-yl)phenyl)ethyildene)amino)-6-((arylidene)amino)-4-(4-chlorophenyl)-2-oxo-1,2-dihydropyridine **23a**–**j**, were synthesized in two steps, by cyclocondensation of oxazine **21** followed by condensation of the intermediate **22**, Scheme 5.



Scheme 5. Reaction pathway to obtain hybrid oxazino-pyridine **23a**–**j**.

The synthesized hybrid compounds were tested for their in vitro antibacterial activity against various bacteria (*Escherichia coli*, *Pseudomonas aeruginosa*, *Staphylococcus aureus*, *Streptococcus pyogenes*) and fungus (*Candida albicans*, *Aspergillus niger*, *Aspergillus clavatus*) *via* the MIC method. Some compounds have proved to have a very powerful activity against bacteria *E. coli* (**23h**, MIC = 25 µg/mL) and against fungus *C. albicans* (**23f**, MIC = 50 µg/mL), respectively, *A. clavatus* (**23h**, MIC = 25 µg/mL).

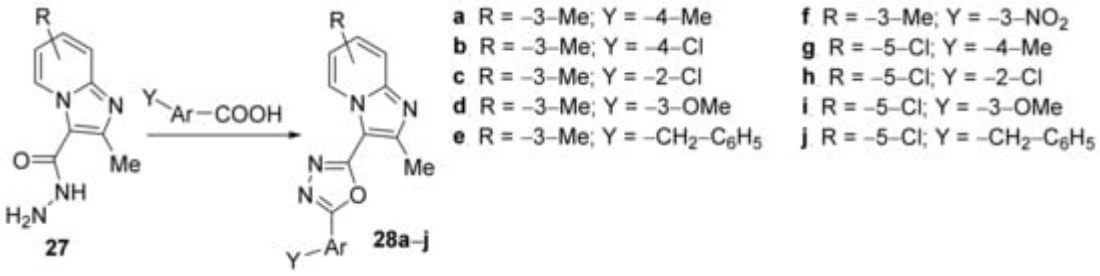
Sribalan et al. [14] have studied thein vitroantimicrobial activity of some tetrazole-heterocycle hybrid derivatives. The synthesis supposes a cyclocondensation reaction of amide precursors **24** with sodium azide, when the corresponding tetrazolo-pyridine **25a**–**d** and tetrazolo-quinoline **26a**–**e** hybrids are obtained, Scheme 6.



Scheme 6. Reaction pathway to obtain the tetrazolo-pyridine **25a**–**d** and tetrazolo-quinoline **26a**–**e** hybrids.

The synthesized tetrazolo-pyridine **25a**–**d** and tetrazolo-quinoline **26a**–**e** hybrids were tested for their in vitro antibacterial activity against various bacteria (*Klebsiella pneumoniae*, *Pseudomonas aeruginosa*, *Staphylococcus aureus*, *Streptococcus pyogenes*) and the fungus *Candida albicans*. An interesting SAR correlation has been performed. The compound **25a** (the pyridyl ring is decorated with *n*-butyl) proved to be the most active from the tetrazolo-pyridine series against all bacteria (DIZ in the range of 4–15 mm), having a superior inhibition to the standard drug (amikacin). The compound **26d** (the quinoline ring is decorated with a piperidyl-sulfonamide moiety) proved to be the most active from the tetrazolo-quinoline series against all bacteria (DIZ in the range of 4–10 mm), having a comparable inhibition to the standard. The antifungal activity was negligible.

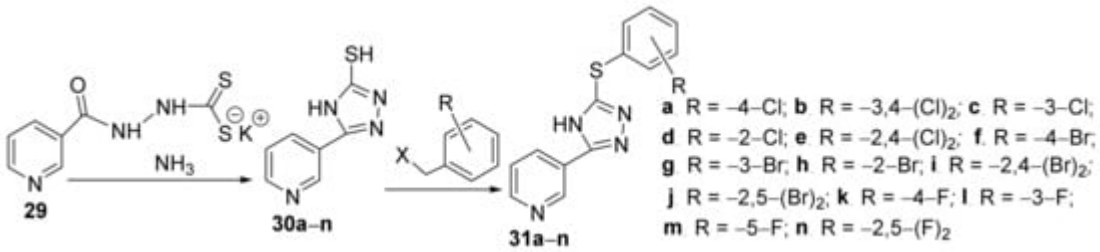
Kuthyala et al. [15] have studied the in vitro antimicrobial activity of some oxadiazolo-imidazopyridine hybrid derivatives. The synthesis was straight, involving a cyclocondensation reaction of hydrazonyl-imidazopyridine **27** with different benzoic acids, when the corresponding oxadiazolo-imidazopyridine hybrids **28a**–**j** were obtained, Scheme 7.



Scheme 7. Reaction pathway to obtain oxadiazolo-imidazopyridine hybrids **28a**–**j**.

The synthesized oxadiazolo-imidazopyridine hybrids **28a**–**j** were tested for their in vitro antibacterial activity against various human bacterial pathogens (*Escherichia coli*, *Klebsiella pneumoniae*, *Staphylococcus aureus*, *Bacillus subtilis*) and the fungus *Candida albicans* and *Aspergillus niger*. An interesting SAR correlation has been performed. The compounds **28f** and **28g** have high activity against *Gram-positive* bacteria *S. aureus* (MIC = 3.12 µg/mL), while compound **28f** proved to have high activity against fungus *C. albicans* (MIC = 12.5 µg/mL).

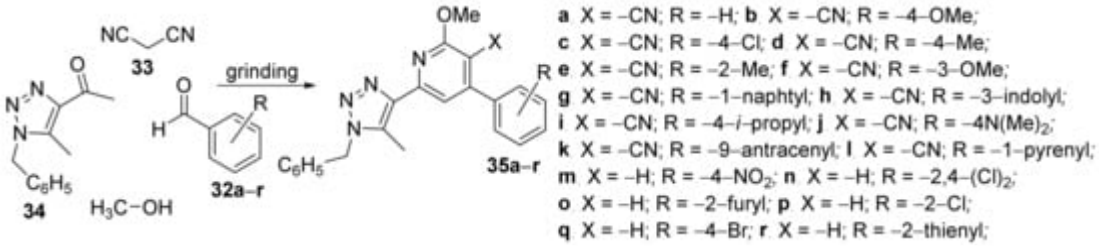
Ahirwaret al. [16] synthesized two new series of some 1,3,4-triazolo-pyridine hybrid derivatives and studied their antimicrobial activities. The synthesis was conducted in two steps: a cyclocondensation reaction of dithiocarbazate **29** with ammonia leading to the first class of hybrids triazolo-pyridine **30a**–**n**, then an alkylation reaction of **30a**–**n** with benzyl halide takes place leading to the second class of hybrids triazolo-pyridine **31a**–**n**, Scheme 8.



Scheme 8. Reaction pathway to obtain 1,3,4-triazolo-pyridine hybryds **30a**–**m** and **31a**–**m**.

The synthesized triazolo-pyridine hybrids **30a**–**n** and **31a**–**n** were evaluated for their in vitro antibacterial activity against *Gram-positive* bacteria (three strains: *Staphylococcus aureus*, *Streptococcus pyogenes*, *Enterococcus faecalis*) and *Gram-negative* bacteria (three strains: *Escherichia coli*, *Pseudomonas aeruginosa*, *Acinetbacter baumannii*) by MIC assay. From the tested compounds, two of them, **31h** and **31i**, have excellent activity against all strains (MIC in the range of 0.91–11 µg/mL).

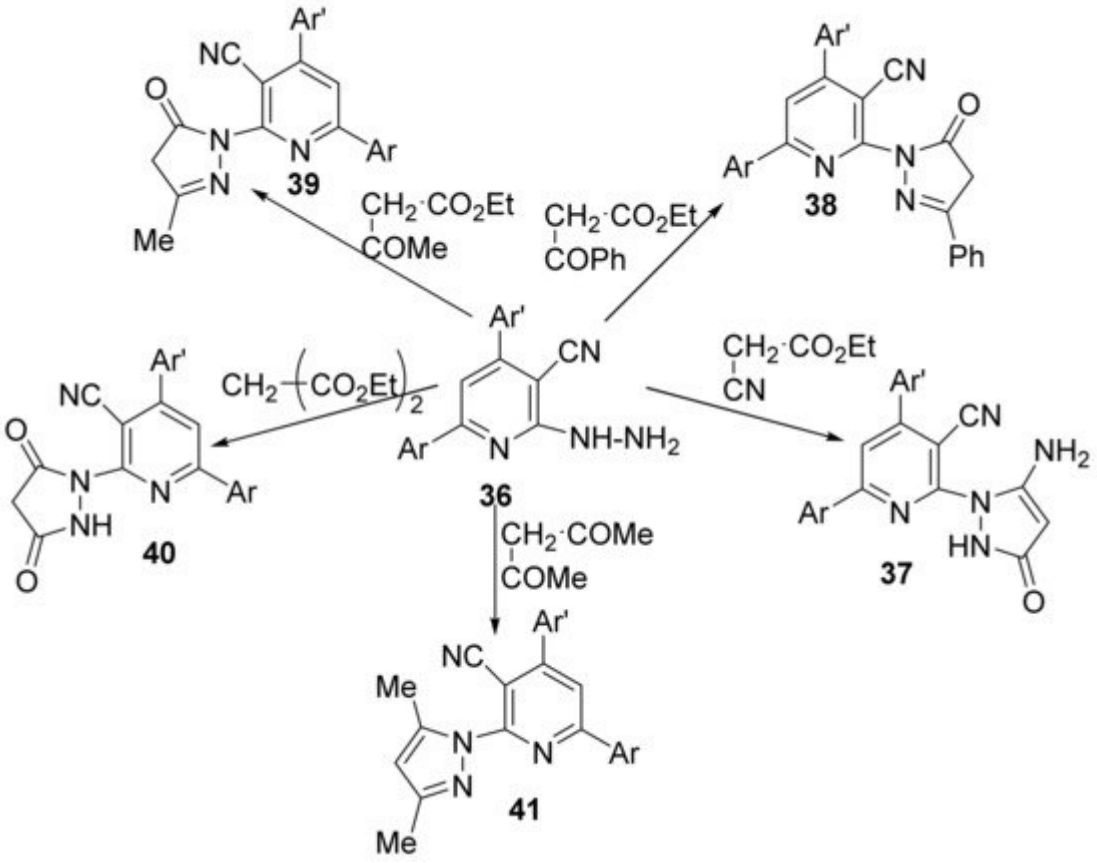
Jaabil et al. [17] have studied the in vitro antimicrobial activities of some newly hybrid 1,2,3-triazolo-pyridine derivatives. The synthesis was green and efficient, under grinding strategy at room temperature, involving *one-pot* sequential multicomponent reactions of aryl aldehydes **32a**–**r**, malonitrile **33**, methanol and 1,2,3-triazolyl ketone **34**, when the corresponding 1,2,3-triazolyl-pyridine/cyanopyridine hybrids **35a**–**r** were obtained, Scheme 9.



Scheme 9. Reaction pathway to obtain 1,2,3-triazolo-pyridine hybrids **35a**–**r**.

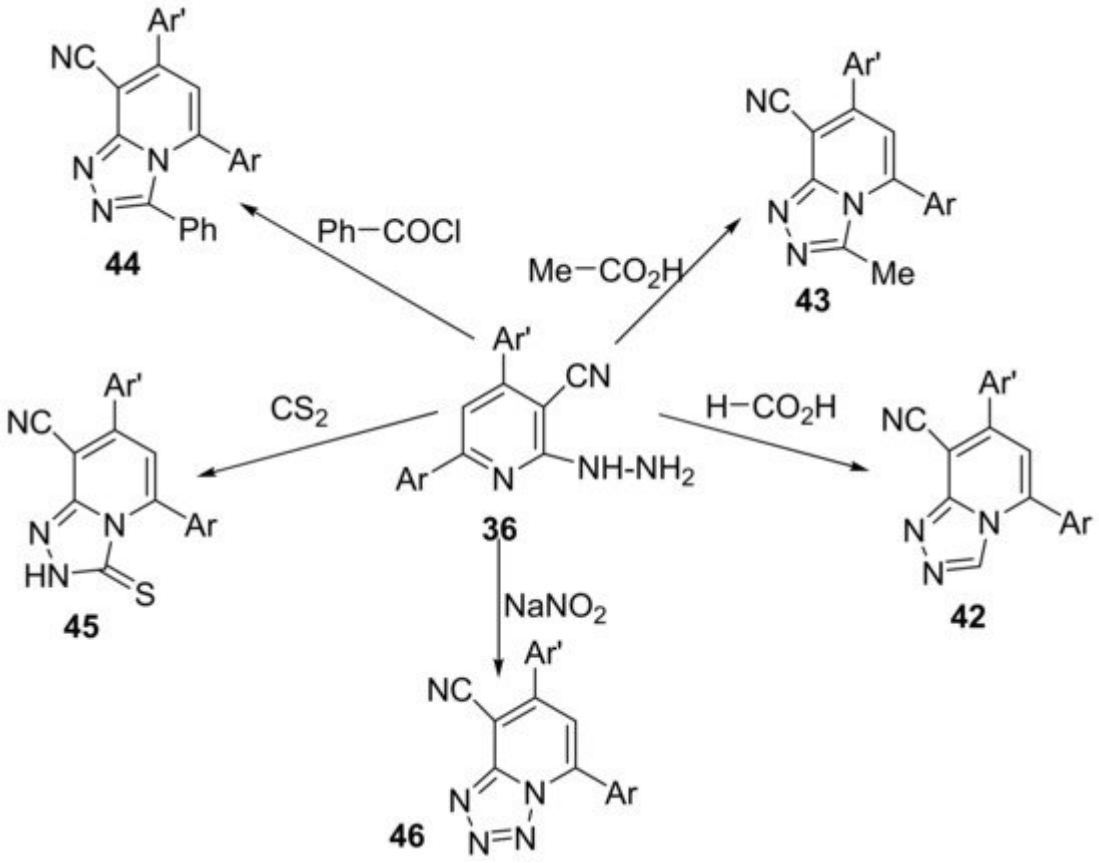
The synthesized 1,2,3-triazolo-pyridine hybrids **35a**–**r** were screened for their in vitro antibacterial activity against three human bacterial strains, *Staphylococcus aureus*, *Salmonella typhi* and *Escherichia coli*, using the MIC method. Some of the 1,2,3-triazolyl cyanopyridine hybrids displayed a remarkable activity against the tested germs, better than tetracycline (standard drug), according to the R-substituent from the phenyl ring. The most active compounds were **35c** (with R = −4-chloro-; MIC in the range of 50–90 µg/mL), **35e** (with R = −2-methyl-; MIC in the range of 40–90 µg/mL) and **35r** (with R = −2-thienyl; MIC in the range of 70–120 µg/mL). The hybrid 1,2,3-triazolo-pyridine compounds were also tested for their antioxidant activity in the assay by 2,2-diphenyl-1-picrylhydrazyl (DPPH) method, showing promising results.

Felefel et al. [18] synthesized three new series of some pyridine hybrid derivatives (namely pyrazole-pyridine **37**–**41**, triazolo-pyridine **42**–**45** and triazino-pyridine **46**) and studied their antimicrobial activities. The synthesis is using as starting material 6-(3,4-dimethylphenyl)-2-hydrazinyl-4-(thiophen-2-yl)-pyridine-3-carbonitrile **36** which react with different compounds with methylene active group (namely acetyl acetone, diethylmalonate, ethyl cyanoacetate, ethyl benzoylacetate and/or ethyl acetoacetate) to produce the desired pyrazole-pyridine hybrid derivatives **37**–**41**, Scheme 10.



Scheme 10. Reaction pathway to obtain pyrazole-pyridine hybrids **37**–**41**.

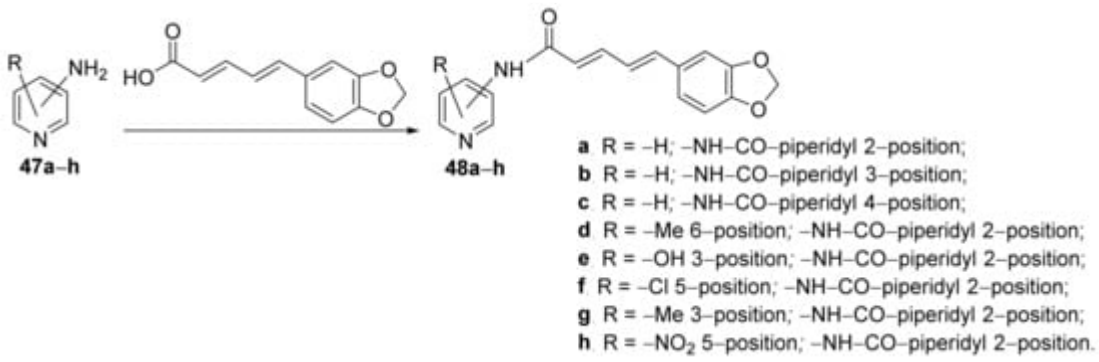
The synthesis of triazolo-pyridines **42**–**45** and tetrazolo-pyridines **46** use as starting material the same intermediate, the 6-(3,4-dimethylphenyl)-2-hydrazinyl-4-(thiophen-2-yl)-pyridine-3-carbonitrile **36**, which react with the appropriate formic acid, acetic acid, benzoyl chloride, carbon disulfide, respectively, sodium nitrite, to produce the desired hybrid derivatives **42**–**45** and **46**, Scheme 11.



Scheme 11. Reaction pathway to obtain triazolo-pyridines **42**–**45** and tetrazolo-pyridine **46** hybrids.

The synthesized pyridine hybrids **37**–**46** were screened for their in vitro antibacterial activity against *Gram-positive* bacteria (*Staphylococcus aureus* and *Bacillus subtilis*), *Gram-negative* bacteria (*Salmonella typhi* and *Escherichia coli*) and fungus (*Aspergillus flavus* and *Candida albicans*) using the disk diffusion agar technique. Some of the hybrids have significant antimicrobial activity, the most active compounds being **37** with a DIZ in the range of 10–17 mm. The antioxidant activity was also tested.

Amperayani et al. [19] synthesized a library of piperine-pyridine hybrid derivatives and studied their antimicrobial activities. The reaction pathway is straight, in one step, involving an acylation reaction of various amino-pyridine derivatives **47a**–**h**, when the corresponding hybrids piperine-pyridine derivatives **48a**–**h** are obtained, Scheme 12.

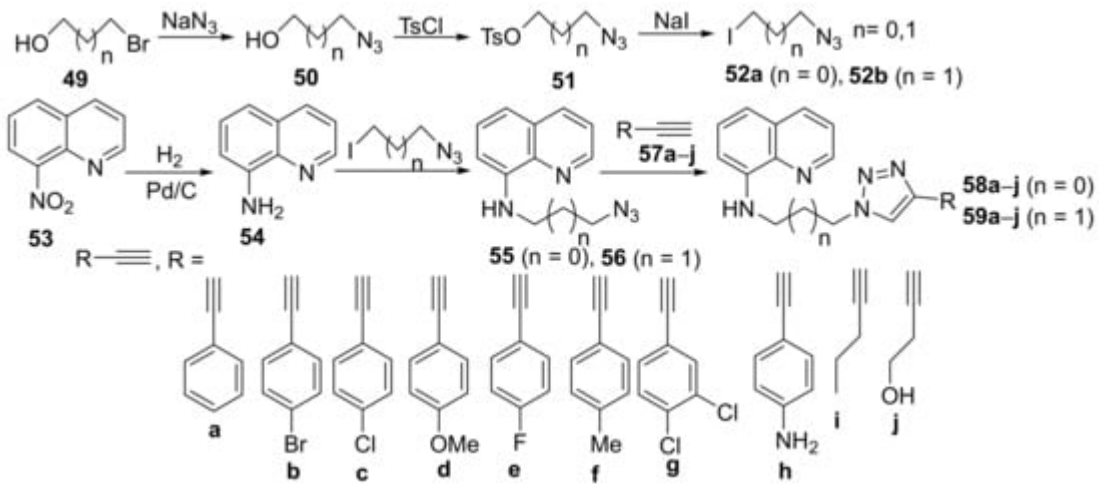


Scheme 12. Reaction pathway to obtain piperine-pyridine hybrids **48a**–**h**.

The synthesized piperine-pyridine hybrid derivatives **48a**–**h** were tested for their in vitro antibacterial activity against some *Gram-positive* and *Gram-negative* bacterial strains (*Bacillus subtilis*, *Streptobacillus*, *Staphylococcus aureus*, *Escherichia coli*, *Klebsiella pneumoniae*, *Pseudomonas aeruginosa*, *Enterococcus faecalis* and *Salmonella typhi*) and fungus strains (*Aspergillus niger*, *Aspergillus flavus*, *Aspergillus fumigatus* and *Candida albicans*) using the disk diffusion agar technique. The piperine-pyridine hybrids **48a**, **48d** and **48h** have very good activity against the *Gram-negative* strains *E. coli*, *K. pneumoniae*, *E. faecalis* and *P. aeruginosa*, having a DIZ in the range of 22–26 mm, superior to control standard drug). The antifungal activity of hybrids was moderate.

3. Six-Member Ring Azaheterocycles with One Nitrogen Atom. Hybrid Quinoline and Isoquinoline

In their attempt to obtain new quinoline derivatives with antimicrobial activity, Albayrak et al. [20] synthesized a library of 20 new triazolo-quinoline hybrid derivatives and studied their antimicrobial activities. The reaction pathway involves several steps (Scheme 13), starting from 8-nitroquinoline **53**. The initial reduction reaction of **53** is leading to 8-aminoquinoline **54**, which is suffering a subsequent *N*-alkylation with azido-iodo-propane **52a**,**b** (generated from the corresponding bromo-alkyl alcohol) leading to alkyl-azide-quinolines **55** and **56**. Finally, the alkyl-azide-quinoline derivatives are treated with the corresponding alkyne **57a**–**j** leading to the desired products, the triazolo-quinoline hybrid derivatives **58a**–**j** and **59a**–**j**.

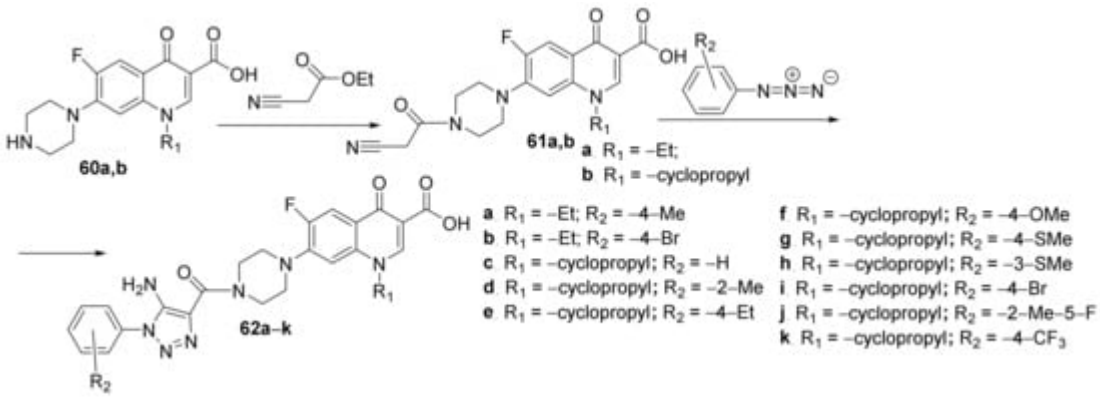


Scheme 13. Reaction pathway to obtain triazolo-quinoline hybrids **58a**–**j** and **59a**–**j**.

The synthesized triazolo-quinoline hybrid derivatives **58a**–**j** and **59a**–**j** were tested for their in vitro antibacterial activity against some *Gram-positive* and *Gram-negative* bacterial strains (*Bacillus subtilis*, *Streptococcus pneumoniae*, *Staphylococcus aureus*, *Escherichia coli*, *Klebsiella pneumoniae*, *Pseudomonas aeruginosa* and *Enterococcus faecalis*) and fungus strains (*Candida parapsilosis* and *Candida albicans*) using the disk diffusion agar technique. The triazolo-quinoline hybrid derivatives **58a**–**j** and **59a**–**j** manifest good activity against the tested strains. The most active compound was **58a**, having excellent activity against *E. coli*, *P. aeruginosa*, *K. pneumoniae*, *E. faecalis*, *S. aureus*, *S. pneumoniae*, *B. subtilis*, *C. albicans* and *C. parapsilosis*. In some cases, the

activity was several orders of magnitude superior to control drugs (DIZ of **58a** was in the range of 35–250 mm; control, ampicillin, respectively, fluconazole have had a DIZ of 35 mm).

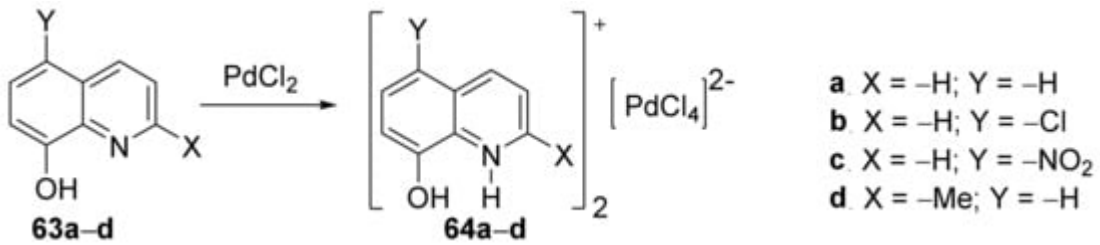
Hryhoriv et al. [21][22] synthesized two new classes of hybrid derivatives analogous to fluoroquinolones, namely piperidino-quinoline **61a**,**b** and 1,2,3-triazolo-piperidino- quinoline **62a**–**k**, and studied their antimicrobial activities. The first class of hybrids was obtainedviaan *N*-alkylation reaction of piperidino-quinoline **60a**,**b,** when the *N*-substituted-piperidino-quinoline hybrids **61a**,**b** are obtained. A click cyclocondensation reaction of **61a**,**b** occurs to the second class of hybrids, the 1,2,3-triazolo-piperidino-quinoline **62a**–**k**, Scheme 14.



Scheme 14. Reaction pathway to obtain *N*-substituted-piperidino-quinoline hybrids **61a**,**b** and1,2,3-triazolo-piperidino-quinoline hybrids **62a**–**k**.

The synthesized hybrid derivatives piperidino-quinoline **61a**,**b** and 1,2,3-triazolo-piperidino-quinoline **62a**–**k** were tested for their in vitro antibacterial activity against standard bacterial strains *Staphylococcus aureus* and *Escherichia coli*, respectively, and the fungus *Candida albicans* using the disk diffusion agar technique. The antimicrobial assay was also made by some clinical bacterial strains *S. aureus* and *E. coli*, respectively, and fungus *C. albicans* using the same method. The hybrid, 1,2,3-triazolo-piperidino-quinoline **62c** have a very good activity against the tested standard strains (DIZ in the range of 25–35 mm), having a superior inhibition zone to control (DIZ = 25 mm). Against clinical microbial strains, the activity was negligible.

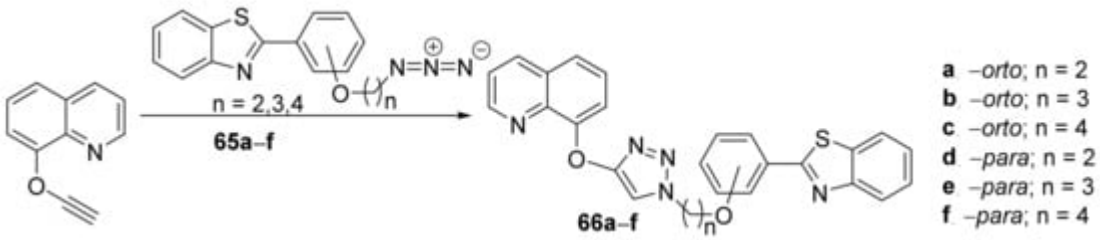
Drweesh et al. [23] synthesized hybrid organic-inorganic derivatives and studied their antimicrobial activities, antiproliferative activity, and radical scavenging properties. In order to synthesize the desired palladium-quinoline derivatives **64a**–**d**, they used organic cation modulation, doing a complexation reaction with $PdCl_2$ of the quinolines **63a**–**d**, Scheme 15.



Scheme 15. Reaction pathway to obtain metal-quinoline hybrids **64a**–**d**.

The synthesized palladium-quinoline derivatives hybrids **64a**–**d** and the free ligands **63a**–**d**, were tested for their in vitro antimicrobial activity against 14 standard microbial strains (*Gram-positive* and *Gram-negative* bacteria, fungus: *Bifidobacterium animalis*, *Lactobacillus plantarum*, *Bacillus subtilis*, *Staphylococcus aureus ATCC 663*, *Staphylococcus aureus ATCC 25923*, *Pseudomonas aeruginosa*, *Proteus mirabilis* , *Escherichia coli*, *Salmonella enterica*, *Candida albicans*, *Saccharomyces boulardii*, *Aspergillus flavus*, *Trichoderma viridae*, *Aspergillus niger*). All hybrid compounds **64a**–**d** have high antimicrobial activity against all tested strains, with minimum inhibitory concentration values ranging from 1.95 to 250 µg/mL. The results of DNA interaction studies indicate that the hybrids **64a**–**d** and the free ligands **63a**–**d**, interact with the DNAvia an intercalation mechanism (the aromatic chromophore intercalates the base pairs of DNA; compound **64a** has the highest binding affinity). The anticancer activity was also studied, with compounds **64a** and **64b** having selective and high cytotoxicity against human lung and breast cancer cells.

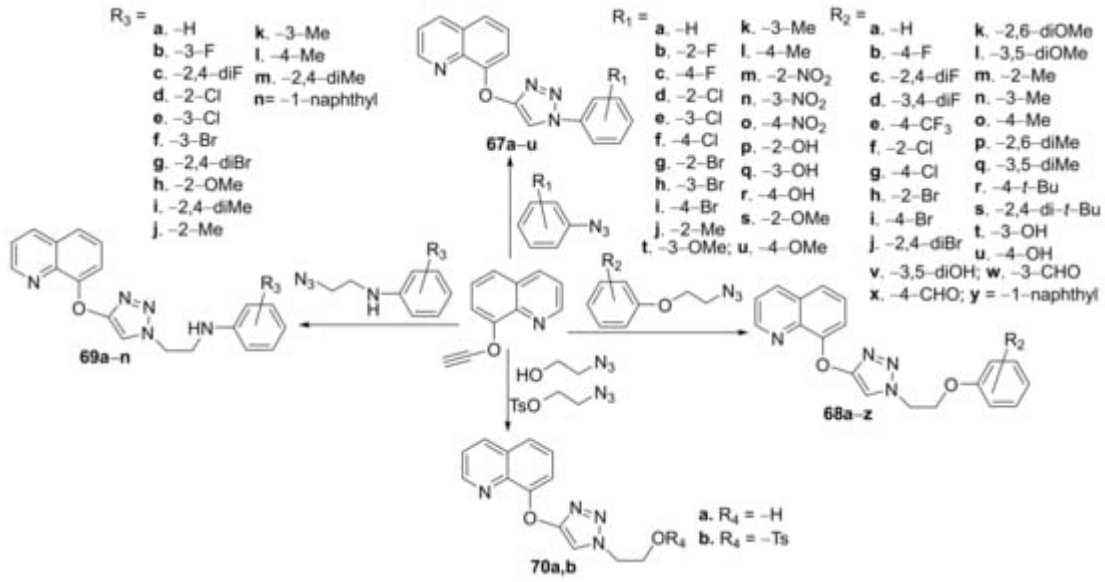
Nehra et al. [24] synthesized a series of triazole-benzothiazole-quinoline hybrids and studied their antimicrobial properties. The reaction pathway is straight and efficient (Scheme 16), involving a click cyclocondensation reaction of azido-alkyl-benzothiazole **65a**–**f** (generated in situ from the corresponding bromo-alkyl derivative) with the corresponding alkyne-quinoline, leading to the desired products, triazole- -benzothiazole-quinoline hybrids **66a**–**f**.



Scheme 16. Reaction pathway to obtain triazole-benzothiazole-quinoline hybrids **66a**–**f**.

The synthesized hybrids **66a**–**f** were evaluated for their in vitro antimicrobial activity against two *Gram-positive* strains (*Staphylococcus aureus* and *Bacillus subtilis*) and two *Gram-negative* strains (*Escherichia coli* and *Pseudomonas aeruginosa*) and two fungal strains (*Candida tropicalis* and *Aspergillus terreus*). The tested hybrids have good antimicrobial activity against both bacteria and fungus. The most promising compound was proved to be **66a**, with an antibacterial (DIZ in the range of 15–17 mm) and antifungal (DIZ in the range of 21–34 mm) activity superior to reference ciprofloxacin (DIZ = 22 mm) and fluconazole (DIZ = 20 mm), respectively. Interesting molecular docking studies were also performed.

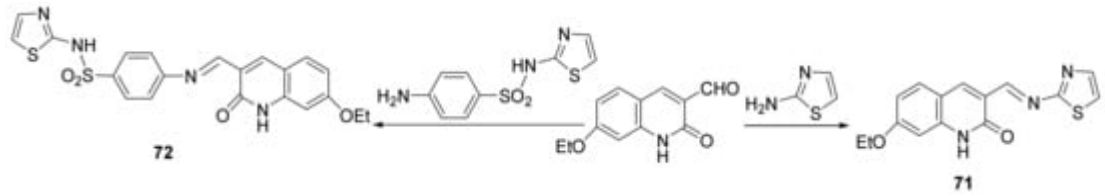
Awolade et al. [25] synthesized a library of triazole-quinoline hybrids and studied their antimicrobial properties. The reaction pathway is straight involving click chemistry of various azides with triple bond derivatives, *via* copper(I)-catalyzed azide-alkyne 3 + 2 dipolar cycloaddition reactions, Scheme 17.



Scheme 17. Reaction pathway to obtaintriazole-quinoline hybrids **67a**–**u**, **68a**–**z**, **69a**–**n** and **70a**,**b**.

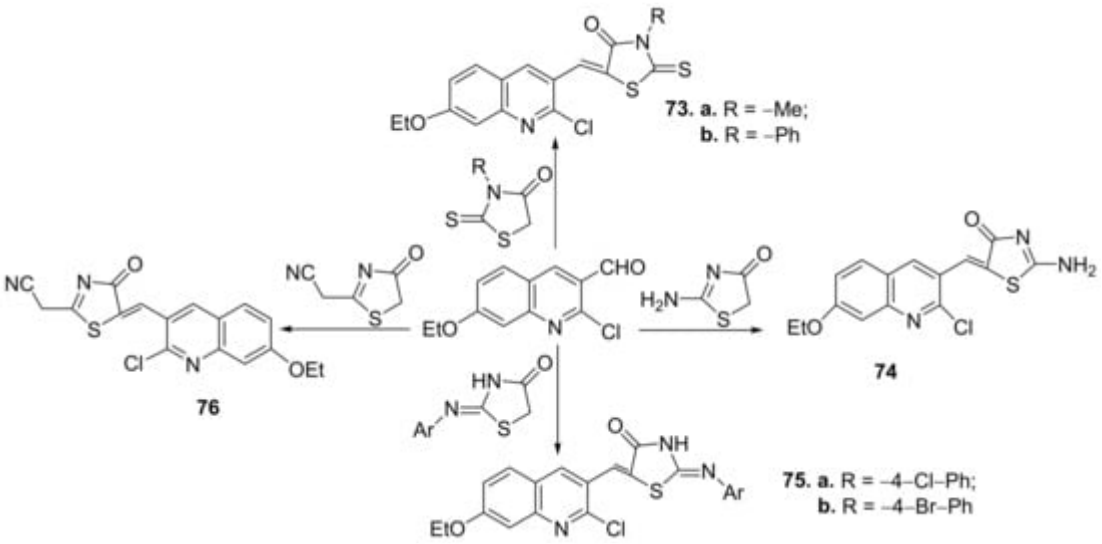
The synthesized hybrids **67a**–**u**, **68a**–**z**, **69a**–**n** and **70a**,**b** were evaluated for their in vitro antimicrobial activity against ESKAPE microbial strains (bacteria and fungus: (*Staphylococcus aureus*, *Escherichia coli*, *Acinetbacter baumannii*, *Klebsiella pneumoniae*, *Candida albicans* and *Candida neoformans*). Some of the compounds proved to have a good and broad-spectrum of antibacterial activity, against methicillin-resistant *S. aureus* (MRSA), *E. coli*, *A. baumannii*, multidrug-resistant *K. pneumoniae* and the fungus *C. albicans* and *C. neoformans* (superior to control, fluconazole). The most promising antibacterial compound was proved to be **70b** with an MIC = 75.39 µM against MRSA, *E. coli*, *A. baumannii*, and multidrug-resistant *K. pneumoniae*. The hybrid **70b** also has a very good antifungal activity against *C. albicans* and *C. neoformans* with an MIC of 37.69 and 2.36 µM, respectively, superior to control fluconazole.

Ammar et al. [26] synthesized a series of thiazole-quinoline hybrids and studied their antimicrobial properties. In order to synthesize the desired compounds, they used the condensation reaction between formil-quinoline derivatives with amino-thiazole or sulfathiazole, when the desired Schiff's base thiazole-quinoline **71** and **72**, are obtained, Scheme 18.



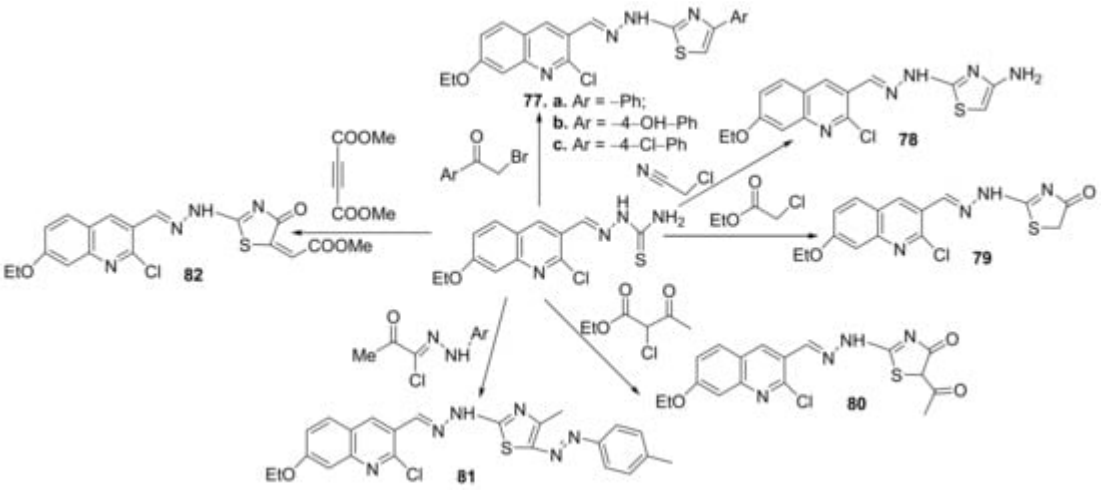
Scheme 18. Reaction pathway to obtain thiazole-quinoline hybrids **71** and **72**.

Further, the condensation reaction between formil-quinoline derivatives with different thiazolone derivatives lead to hybrid thiazolone-quinoline derivatives **73**–**76**, Scheme 19.



Scheme 19. Reaction pathway to obtain thiazolone-quinoline hybrids **73**–**76**.

Finally, the cyclization of different quinoline-thiosemicarbazone derivatives with the halogenated compounds lead to other hybrid thiazole-quinoline derivatives **77**–**82**, Scheme 20.

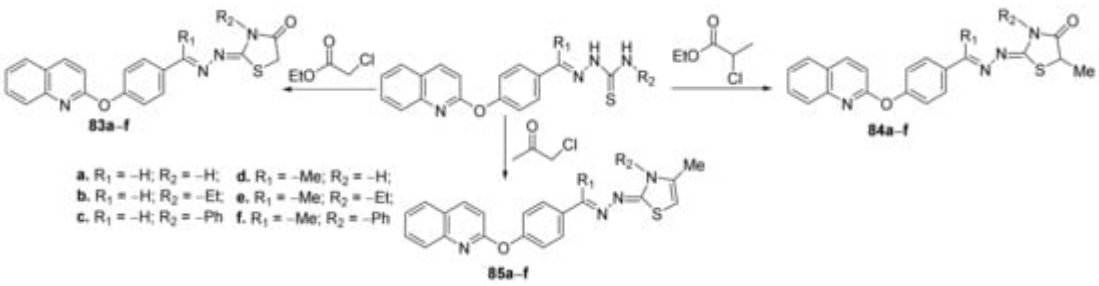


Scheme 20. Reaction pathway to obtain thiazole-quinoline hybrids **77**–**82**.

The synthesized hybrids **71**–**82**, were evaluated for theirin vitroantimicrobial activity against eight standard microbial strains, three *Gram-positive* bacteria (*Staphylococcus aureus*, *Bacillus faecalis* and *Bacillus subtilis*), three *Gram-negative* bacteria (*Escherichia coli*, *Salmonella typhi* and *Pseudomonas aeruginosa*), and two fungi (*Candida albicans* and *Fusarium oxysporum*). Some of the compounds have good antimicrobial activity, with MIC and MBC values ranging between 0.95 and 62.5 µg/mL, and 1.94 and 118.7 µg/mL, respectively. Two compounds, namely **77b** and **73a**, proved to be the most active of the series against *S. aureus* and *E. coli* having an MIC between 0.95 and 7.81 µg/mL, respectively a MBC between 3.31 and 15.62 µg/mL.

Using a similar strategy, some of the above researchers (Eissa et al. [27]) synthesized a new series of thiazole-quinoline hybrids and studied their antimicrobial properties. In order to synthesize the desired compounds, they

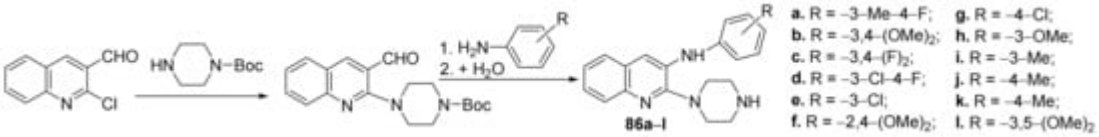
used the cyclization of quinoline-thiosemicarbazone derivatives with the halogenated compounds, when the corresponding hybrid thiazole-quinoline derivatives, **83a**–**f**, **84a**–**f** and **85a**–**f** are obtained, Scheme 21.



Scheme 21. Reaction pathway to obtain thiazole-quinoline hybrids **83**–**85a**–**f**.

The synthesized hybrids **83a**–**f**, **84a**–**f** and **85a**–**f**, were evaluated for their in vitro antimicrobial activity against *Gram-positive* (five strains: *Staphylococcus aureus*, *Staphylococcus epidermidis*, *Streptococcus pyogenes*, *Bacillus subtilis* and *Enterococcus faecalis*) and *Gram-negative* bacteria (five strains: *Neisseria gonorrhoeae*, *Proteus vulgaris*, *Klebsiella pneumonia*, *Shigella flexneri* and *Pseudomonas aeruginosa*), as well as fungus (five strains: *Aspergillus fumigatus*, *Aspergillus clavatus*, *Candida albicans*, *Geotrichum candidum*, and *Penicillium marneffei*). Some of the compounds displayed good antimicrobial activity, superior to the used control. The most active compound was found to be **85e**, having a two-fold potency compared with gentamycin for inhibition of *N. gonorrhoeae*, four-fold potency compared with amphotericin B for the inhibition of *A. fumigatus*, equipotent activity compared with the reference drugs for inhibition of *S. flexneri*, *S. pyogenes*, *P. vulgaris*, *A. clavatus*, *G. candidum* and *P. marneffei*.

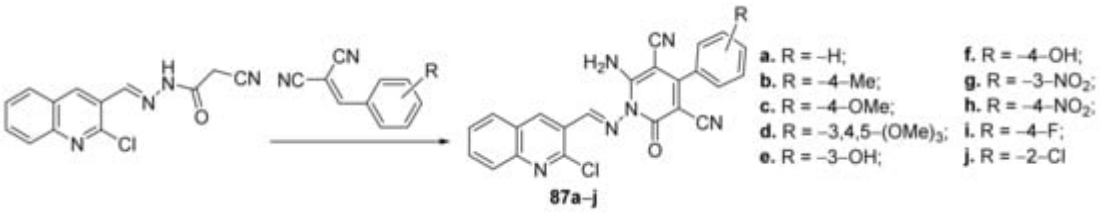
Lagdhir et al. [28] synthesized a library of piperazin-quinoline hybrids and studied their antimicrobial properties. The reaction pathway involves two steps (an alkylation and a condensation reaction), leading to the piperazin-quinoline hybrids **86a**–**l**, Scheme 22.



Scheme 22. Reaction pathway to obtain piperazin-quinoline hybrids **86a**–**l**.

The synthesized hybrids **86a**–**l** were evaluated for their in vitro antibacterial (*Staphylococcus aureus*, *Streptococcus pyogenes*, *Escherichia coli* and *Pseudomonas aeruginosa*) and antifungal (*Aspergillus clavatus*, *Aspergillus niger* and *Candida albicans*) activity, antimalarial (*Plasmodium falciparum*) and antituberculosis (*Mycobacterium tuberculosis*) activity. Some of the compounds have good antibacterial and antifungal activity against *S. aureus* and *C. albicans*. The hybrids **86a**, **86b**, **86d**, **86j** and **86k**, are the most active as an antimicrobial against *S. aureus*, having an MIC = 100 µg/mL, equal to the control drug ampicillin. The hybrid **86k** has excellent antifungal activity against *C. albicans*, having an MIC = 250 µg/mL, two folds higher compared with the control drug griseofulvin. The antimalarial and antitubercular activity proved to be moderate for the majority of compounds.

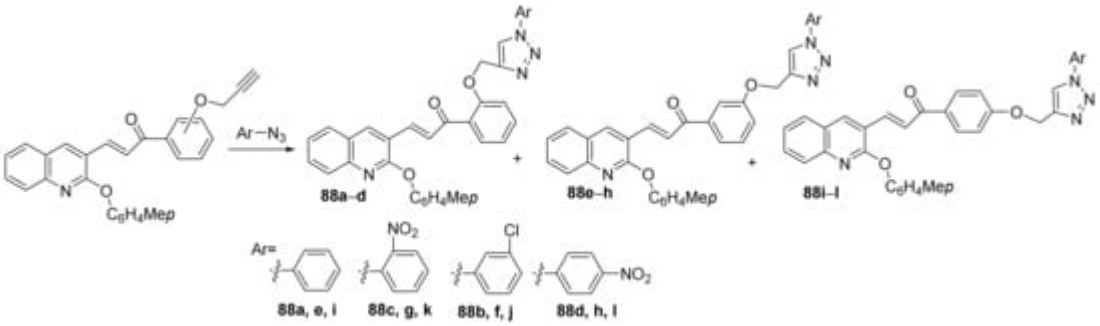
Desai et al. [29] synthesized a series of pyridine-quinoline hybrids and evaluated it for their antimicrobial properties. The reaction pathway involves a cyclocondensation reaction of quinoline derivative with benzylidene-malononitril, when the corresponding pyridine-quinoline hybrids **87a**–**j** were obtained, Scheme 23.



Scheme 23. Reaction pathway to obtain pyridine-quinoline hybrids **87a**–**j**.

The synthesized hybrids **87a**–**j** were evaluated for their in vitro antimicrobial activity against *Gram-positive* (two strains: *Staphylococcus aureus* and *Staphylococcus pyogenes*) and *Gram-negative* (two strains: *Escherichia coli* and *Pseudomonas aeruginosa*) bacteria, as well as to fungus (three strains: *Aspergillus clavatus*, *Aspergillus niger* and *Candida albicans*). Some of the compounds displayed promising antimicrobial activity. The hybrid **87i** has the best antibacterial activity against *E. coli*, *P. aeruginosa* and *S. aureus* strains, with an MIC = 12.5 µg/mL, two folds higher compared with the control drug ciprofloxacin (MIC = 25 µg/mL). The most active compound against *C. albicans* was found to be **87e**, having an MIC=25 µg/mL, much better compared with the control drug griseofulvin (MIC = 500 µg/mL).

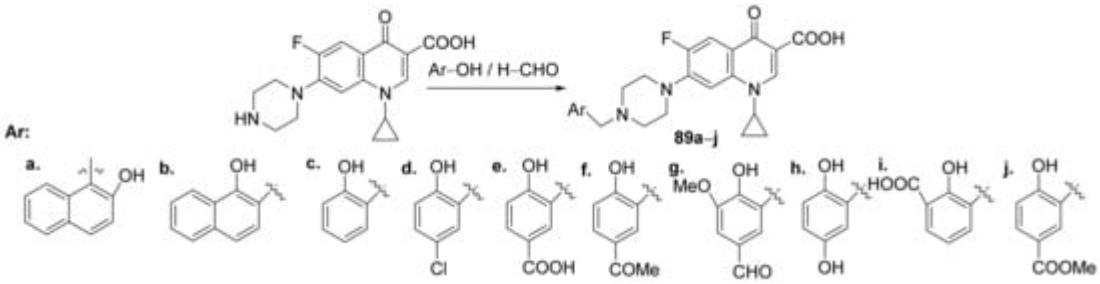
Vishnuvardhan et al. [30] synthesized a library of triazole-quinoline hybrids and studied their antimicrobial properties. The reaction pathway involves a typical click cyclocondensation reaction of quinoline with a triple bond with aryl-azide derivatives, when the corresponding triazole-quinoline hybrids **88a**–**l**, Scheme 24.



Scheme 24. Reaction pathway to obtain triazole-quinoline hybrids **88a**–**l**.

The synthesized hybrids **88a**–**l** were evaluated for theirin vitroantimicrobial activity against *Gram-positive* (*Staphylococcus aureus* and *Enterococcus faecalis*) and *Gram-negative* (*Escherichia coli* and *Pseudomonas aeruginosa*) bacteria, as well as to fungus (*Aspergillus niger* and *Candida albicans*). Most of the hybrid compounds have good antimicrobial activity. The best antibacterial activity reveals the hybrids **88d**, **88h** and **88i**, having a DIZ in the range of 16–21 mm, superior to control ampicillin (DIZ = 15 mm). The best antifungal activity reveals the hybrids **88d**, **88h** and **88k**, having a DIZ in the range of 18–27 mm, superior to control griseofulvin (DIZ = 17 mm).

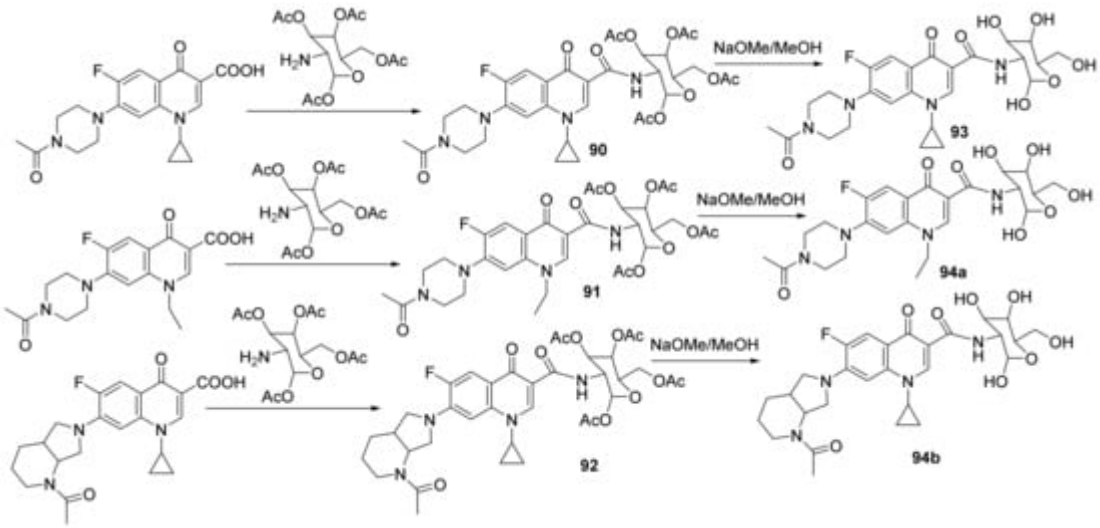
Abdel-Rahman et al. [31] synthesized a series of piperazin-quinoline hybrids derived from ciprofloxacin and studied their antimicrobial and anticancer properties. The reaction pathway involves the reaction of ciprofloxacin with the corresponding phenolic derivatives with an excess of formaldehyde, when the piperazin-quinoline hybrids **89a**–**j** are obtained, Scheme 25.



Scheme 25. Reaction pathway to obtain piperazin-quinoline hybrids **89a**–**j**.

The synthesized hybrids **89a**–**j** were evaluated for their antimicrobial and anticancer activity. The antibacterial screening was preconformed on *Gram-positive* and *Gram-negative* strains: *Staphylococcus aureus*, MRSA clinical strain, MRSA reference strain, *Escherichia coli* and *Pseudomonas aeruginosa*. The obtained results reveal that the hybrid **89d** has the best antibacterial activity against *S. aureus*, MRSA (reference strain) and MRSA (clinical strain) with an MIC of 0.57, 0.52, and 0.082 µg/mL, respectively, (compared with the reference standard drug ciprofloxacin which has an MIC of 1.63 µg/mL against *S. aureus*, an MIC of 1.45 µg/mL against MRSA reference, and an MIC of 0.84 µg/mL against MRSA clinical). The hybrid **89j** exhibited the best antimicrobial activity against *E. coli* and *P. aeruginosa*, with an MIC of 0.036 and 0.043, respectively, (compared with the reference standard drug ciprofloxacin which has an MIC of 0.056 µg/mL against *E. coli* and an MIC of 1.27 µg/mL against *P. aeruginosa*).

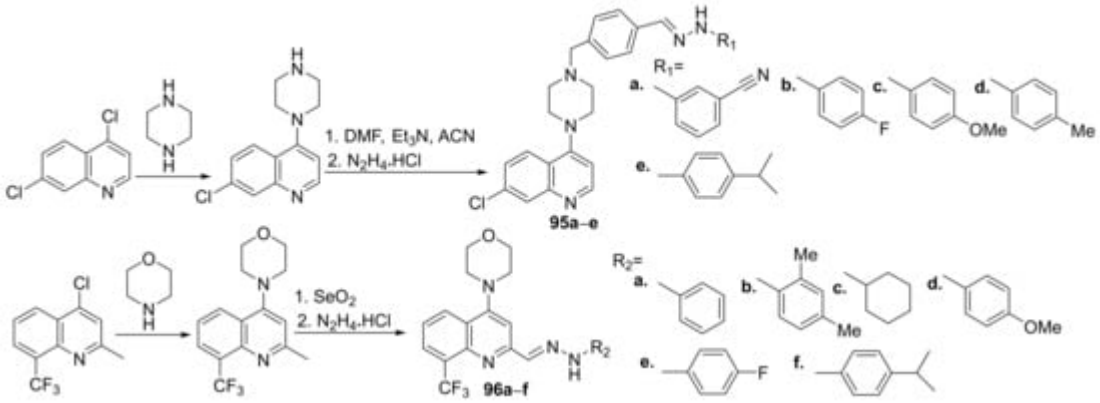
Mohammed et al. [32] synthesized a series of glycosylated-quinoline hybrids derived from fluoroquinolone and studied their antimicrobial properties. The reaction pathway involves the reaction of ciprofloxacin with the corresponding phenolic derivative with an excess of formaldehyde, when the glycosylated-quinoline hybrids **90**–**94** are obtained, Scheme 26.



Scheme 26. Reaction pathway to obtainglycosylated-quinoline hybrids **90**–**94**.

The synthesized glycosylated-quinoline hybrids **90**–**94** were evaluated for their antibacterial activity against various *Gram-positive* and *Gram-negative* bacteria: *Escherichia coli*, *Listeria monocytogenes*, *Salmonella enterica*, *Pseudomonas aeruginosa*, *Listeria monocytogenes*, *E. coli* clinical isolate (resistant to nalidixic acid, ciprofloxacin HCl and norfloxacin antibiotics), methicillin-resistant *Staphylococcus aureus* (MRSA), methicillin-sensitive *Staphylococcus aureus* (MSSA). The hybrids were also tested for their antifungal activity against fungi: *Candida albicans*, *Aspergillus flavus*, *Fusarium solani*, *Stachybotrys chartarum* and *Penicillium chrysogenum*. The hybrid compounds **90**, **91** and **94a** have excellent antimicrobial activity against a fluoroquinolone-resistant *E. coli* clinical isolate, comparable to controls ciprofloxacin and norfloxacin. The hybrid compound **91** also has good antifungal activity against *C. albicans* and *P. chrysogenum*.

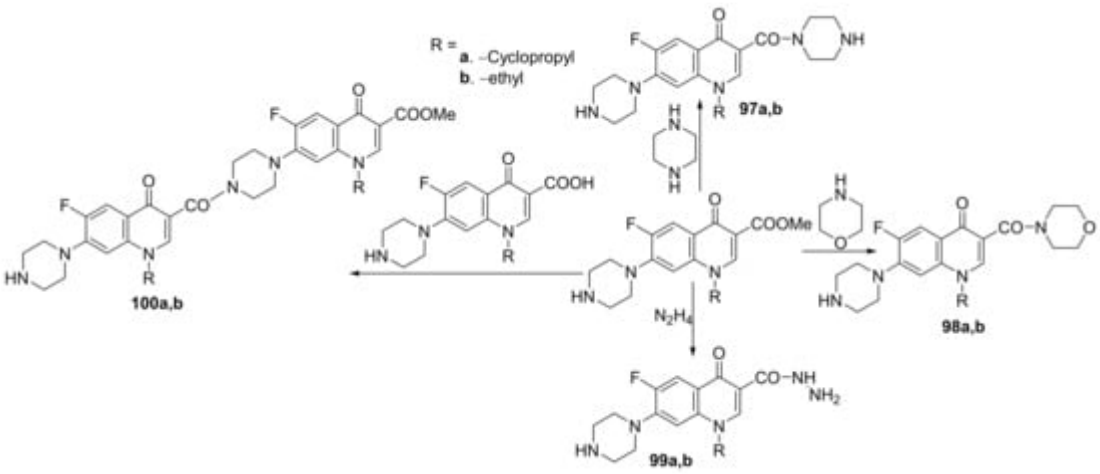
Shruthi et al. [33] synthesized a series of piperazine-quinoline hybrids **95a**–**e** and morpholine-quinoline hybrids **96a**–**f** and evaluate them for their antimicrobial properties. The reaction pathway is depicted in Scheme 27.



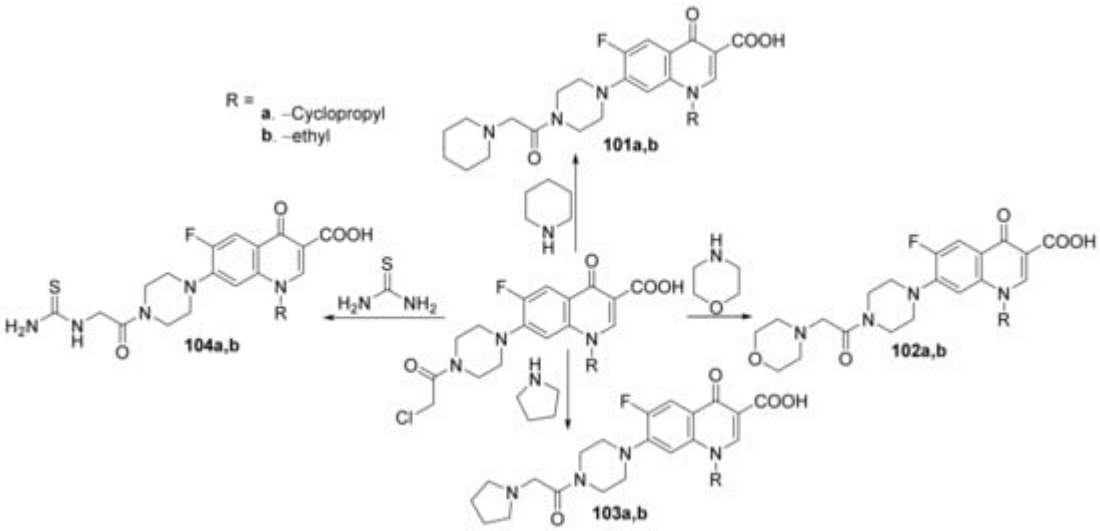
Scheme 27. Reaction pathway to obtain piperazine- and morpholine-quinoline hybrids **95a**–**e** and **96a**–**f**.

The synthesized hybrids **95a**–**e** and **96a**–**f** were evaluated for their antibacterial (*Acinetobacter baumanii*, *Enterococcus faecium*, *Klebsiella pneumonia*, *Pseudomonas aeruginosa*, *Escherichia coli* and *Staphylococcus aureus*) and antitubercular (*Mycobacterium tuberculosis*) activity. Hybrid **95b** has the best antibacterial activity against *E. coli* and *S. aureus* strains with an MIC of 4, respectively, 2 µg/mL, compared to standard drug vancomycin (MIC of 16, respectively, 0.5 µg/mL). Hybrids **95d**, **95e** and **96f** exhibited the best antibacterial activity against *A. baumaniistains* with an MIC in the range of 1–2 µg/mL, compared to standard drug vancomycin (MIC = 0.5 µg/mL). Hybrids **95b**, **95d** and **95e** also have promising antitubercular activity with an MIC of 4 µg/mL.

Kaur et al. [34] synthesized a series of 3- and 7- substituted-quinoline hybrids derived from fluoroquinolone and studied their antimicrobial properties. The reaction pathway involves the reaction of fluoroquinolone derivatives with the corresponding reagents, when the quinoline hybrids **97**–**104a**,**b** are obtained, Scheme 28 and Scheme 29.



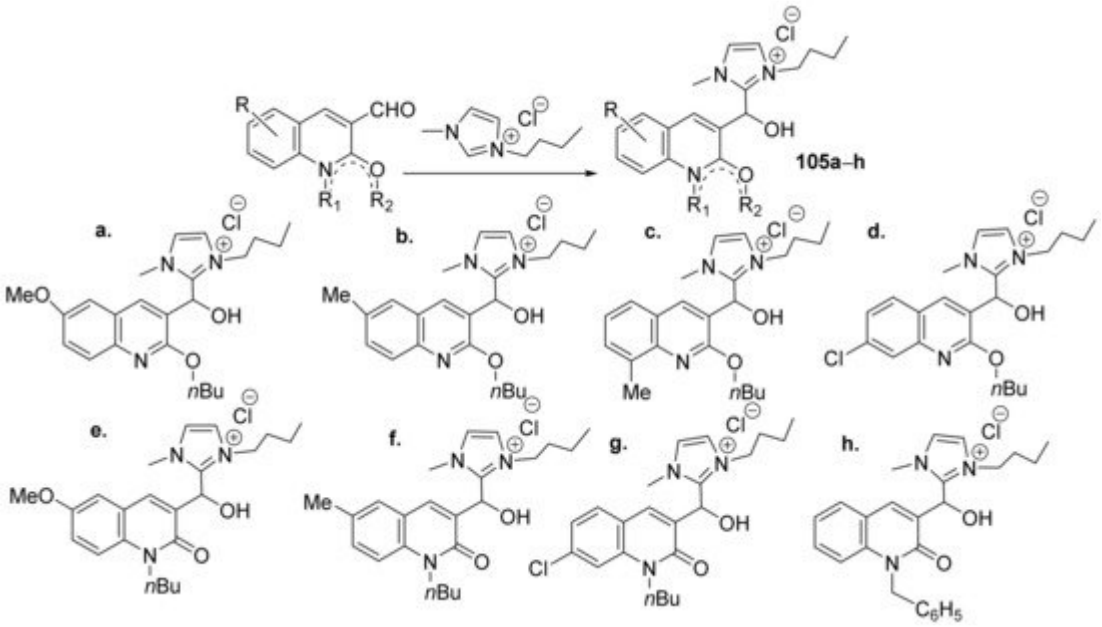
Scheme 28. Reaction pathway to obtain piperazino-quinoline hybrids **97**–**100a**,**b**.



Scheme 29. Reaction pathway to obtain 7-substituted-quinoline hybrids **101**–**104a**,**b**.

The synthesized quinoline hybrids **97**–**104a**,**b** were evaluated for their antibacterial activity against four bacterial strains: *Bacillus subtilis*, *Pseudomonas aeruginosa*, *Escherichia coli* and *Staphylococcus aureus*. All hybrids **97**–**104a**,**b** have proved to be active against all bacterial strains, with an MIC value of 25 µg/mL which is fourfold more active compared to the standard drug ciprofloxacin (MIC = 100 µg/mL).

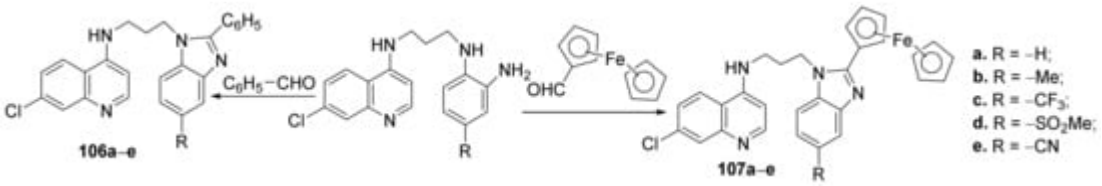
Insuasty et al. [35] synthesized a series of imidazolium-quinoline hybrids and studied their antimicrobial properties. The reaction pathway involves the reaction of 3-formyl-quinolone derivatives with the corresponding imidazolium salts, when the imidazolium-quinoline hybrids **105a**–**h** are obtained, Scheme 30.



Scheme 30. Reaction pathway to obtain imidazolium-quinoline hybrids **105a**–**h**.

The synthesized imidazolium-quinoline hybrids **105a**–**h** were evaluated for their antibacterial (*Klebsiella pneumoniae*, *Escherichia coli* and *Staphylococcus aureus*), antifungal (*Cryptococcus neoformans*) and antitubercular (*Mycobacterium tuberculosis H37Rv* and *Mycobacterium bovis BCG*) activities. Hybrid derivatives **105c,d** demonstrated a remarkable antifungal activity against *C. neoformans* (MIC in the range of 15 µg/mL) while for the other fungal strains the activity is weak. The hybrids have modest antibacterial activity (both against *Gram-positive* and *Gram-negative* bacteria) as well as antitubercular activity.

Baartzes et al. [36] synthesized a series of benzimidazole-quinoline and ferrocenyl-quinoline hybrids and studied their antimicrobial properties. The reaction pathway involves the reaction of amino-quinoline derivatives with the corresponding formyl derivatives, when the benzimidazole-quinoline hybrids **106a**–**e** and ferrocenyl-quinoline hybrids **107a**–**e** are obtained, Scheme 31.

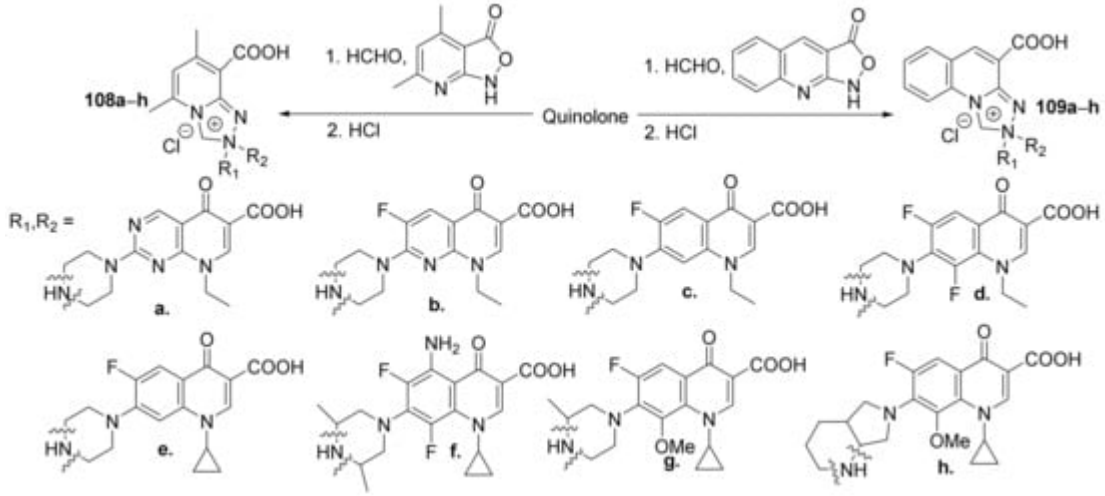


Scheme 31. Reaction pathway to obtain benzimidazole-quinoline and ferrocenyl-quinoline hybrids **106a**–**e** and **107a**–**e**.

The synthesized quinoline hybrids **106a**–**e** and **107a**–**e** were evaluated for their antimalarial (*Plasmodium falciparum* and *Plasmodium berghei*) and antitubercular (*Mycobacterium tuberculosis*) activity. All hybrid derivatives are active against tested malaria strains and have modest activity against them. The most active hybrids against

malarial strains have proved to be **106c** and **107b**, with an IC_{50} of 0.43, respectively, 0.32 μM, compared with the standard drug chlorquine (IC_{50} = 0.01 μM).

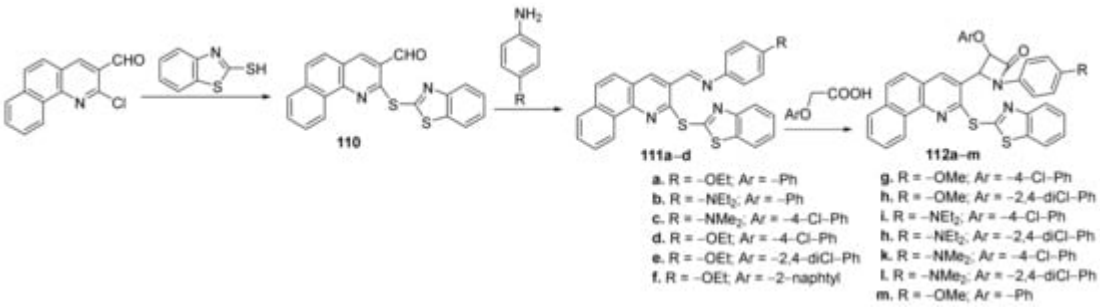
Fedorowicz et al. [37] synthesized a series of zwiterionic hybrids pyridine-fluoroquinolone **108a**–**h** and quinoline-fluoroquinolone **109a**–**h** and studied their antimicrobial properties. The reaction pathway involves a tandem Mannich-electrophilic amination reaction of isoxazolones derivatives and fluoroquinolone bearing a secondary amino group at position 7 of the quinoline ring, Scheme 32.



Scheme 32. Reaction pathway to obtain zwiterionic pyridine-fluoroquinolone and quinoline-fluoroquinolone hybrids **108a**–**h** and **109a**–**h**.

The synthesized quinoline hybrids **108a**–**h** and **109a**–**h** were evaluated for their antibacterial activity against *Gram-positive* and *Gram-negative* bacterial strains (laboratory and clinical: *Staphylococcus aureus ATCC 6538*, *Staphylococcus aureus MRSA N315*, *Staphylococcus epidermidis ATCC 14990*, *Bacillus subtilis ATCC 6633*, *Escherichia coli ATCC 8739*, *Pseudomonas aeruginosa ATCC 9027*, *Proteus vulgaris NCTC 4635*, *Staphylococcus aureus MRSA 6347*, *Staphylococcus epidermidis MRSE 13199* and *Serratia marcescens 12795*) as well as for antibiofilm activity. The hybrid derivatives proved to have bactericidal and antibiofilm activity. The most active hybrids were found to be **109d** and **109e**, exhibiting good inhibition against all strains, with the IC_{50} values in the low micromolar range.

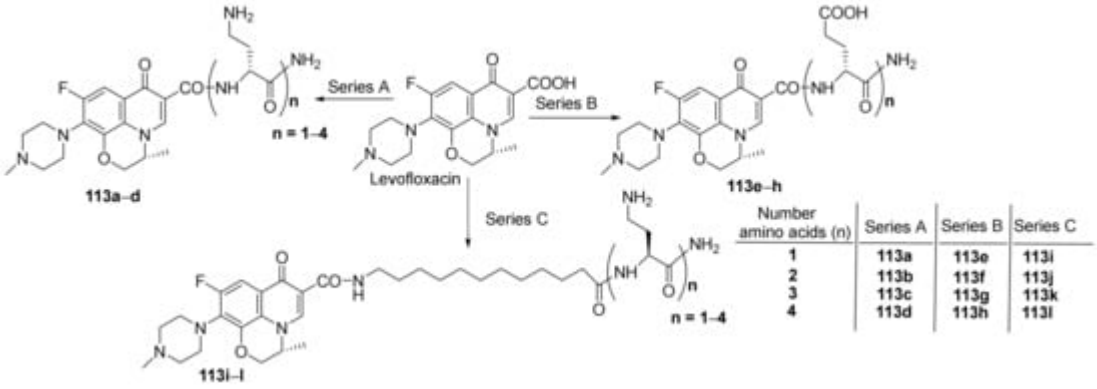
Borazjani et al. [38] synthesized a library of quinoline hybrids (benzothiazole-benzo-quinoline **110**, imino-benzothiazole-benzo-quinoline **111a**–**d**, β-lactam-benzo-thiazole-benzo-quinoline **112a**–**m**) and studied their antimicrobial properties. The reaction pathway involves a [2+2]-cycloaddition reaction of imines **111a**–**d** and ketenes derived from substituted acetic acids, Scheme 33.



Scheme 33. Reaction pathway to obtain benzothiazole-benzo-quinoline hybrids **110**, **111a**–**d** and **112a**–**m**.

The synthesized quinoline hybrids **110**–**112** were evaluated for their antimicrobial activity against *Gram-positive* and *Gram-negative* bacterial strains: *Staphylococcus aureus*, *Bacillus subtilis*, *Enterococcus faecalis*, *Salmonella typhi*, *Escherichia coli* and *Pseudomonas aeruginosa*. From the β-lactam class, the assay indicates that the most active hybrids against *E. coli* and *P. aeruginosa*, are **112k** and **112m**, with an MIC of 42, respectively, 20 µg/mL, compared to standard drug gentamycin (MIC of 90, respectively, 5 µg/mL). From the imino-benzothiazole-benzo-quinoline class, the most active hybrids against *P. aeruginosa* and *S. aureus*, are **111a**–**c**, with an MIC of 42 µg/mL, compared to standard drug gentamycin (MIC of 5, respectively, 90 µg/mL).

Berry et al. [39] synthesized a series of peptide-fluoroquinolone hybrids and studied their antimicrobial properties. In order to synthesize the desired hybrids, the researchers used solid-phase peptide synthesis, from levofloxacin fluoroquinolone with the corresponding peptide (oligopeptide), when the desired peptide-fluoroquinolone hybrids **113a**–**l** are obtained, Scheme 34.

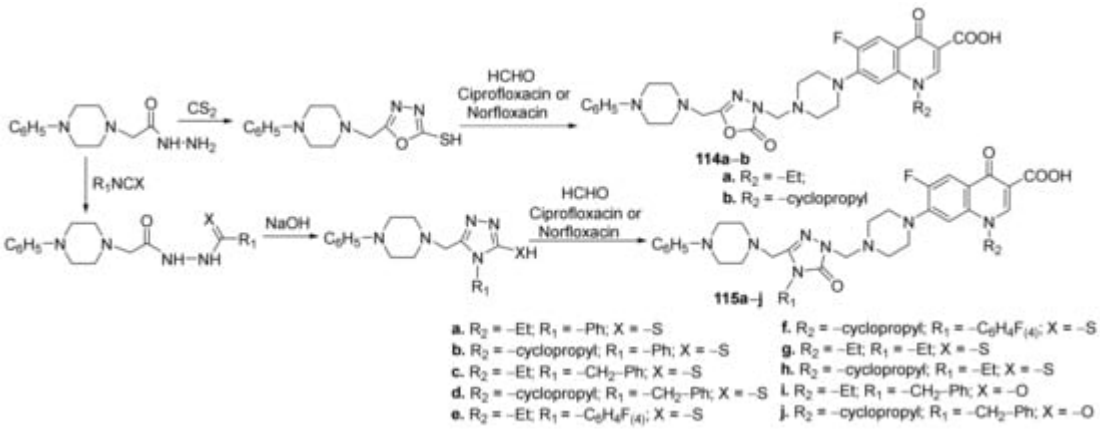


Scheme 34. Reaction pathway to obtain peptide-quinolone hybrids **113a**–**l**.

The synthesized peptide-fluoroquinolone hybrids **113a**–**l** were evaluated for their antimicrobial activity against MDR bacterial strains, *Gram-negative* and *Gram-positive*: *Pseudomonas aeruginosa*, *Escherichia coli*, *Klebsiella pneumoniae*, *Acinetobacter baumannii*, methicillin-resistant *Staphylococcus aureus* (MRSA), methicillin-sensitive *Staphylococcus aureus* (MSSA), methicillin-resistant *Staphylococcus epidermis* (MRSE), *Enterococcus faecalis*, *Enterobacter cloacae*, *Stenotrophomonas maltophilia*. The assay indicates that all the peptide-hybrids have weak antibacterial activity. If the hybrids are mixed with fluoroquinolone (ciprofloxacin, levofloxacin and moxifloxacin)

drugs, the resulting conjugates possess antimicrobial activity against MDR *Gram-negative* bacteria (clinical isolates, *P. aeruginosa*, *E. coli*, *K. pneumoniae*, *A. baumannii*), superior to reference levofloxacin.

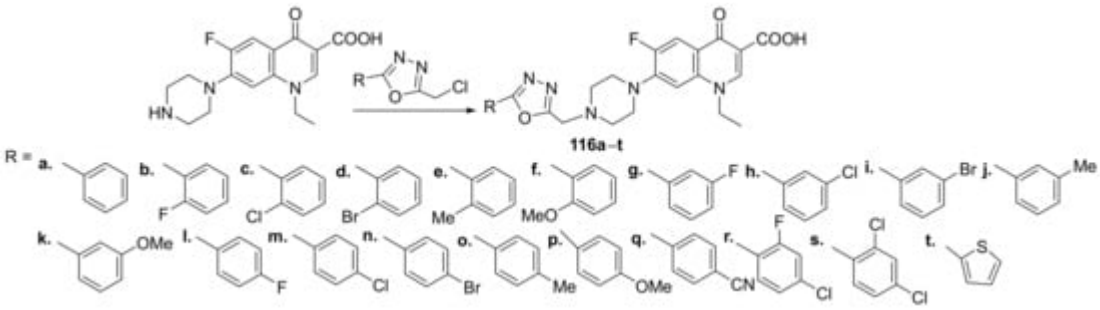
Mermer et al. [40] synthesized a library of triazole- and oxadiazole-fluoroquinolone hybrids and studied their antimicrobial properties. The reaction pathway took placeviaseveral steps of sequential reactions, starting from phenyl piperazine. Finally, the corresponding triazole-fluoroquinolone **114a**,**b** and oxadiazole-fluoroquinolone **115a**–**j** hybrids were obtained *via* a *one-pot* three-component Mannich reaction, Scheme 35. The reactions were performed both under conventional thermal heating and microwave, the last pathway being more advantageous.



Scheme 35. Reaction pathway to obtain oxadiazole- and triazole-fluoroquinolone hybrids **114a**,**b** and **115a**–**j**.

The synthesized hybrids **114a**,**b** and **115a**–**j** were tested for their antimicrobial activity (against *Gram-positive* and *Gram-negative* strains: *Staphylococcus aureus*, *Enterococcus faecalis*, *Escherichia coli*, *Pseudomonas aeruginosa*, *Klebsiella pneumoniae*, *Acinetobacter haemolyticus*), DNA gyrase and Topoisomerase IV inhibition potentials. The hybrids have good antimicrobial activity and displayed excellent DNA gyrase inhibition. The hybrids **114b**, **115b** and **115h** exhibited the best antimicrobial activity against the tested strains. Thus, the hybrids have excellent activity against *K. pneumoniae* with an MIC of 0.25 µg/mL, compared with the standard drug gentamycin (MIC = 0.25 µg/mL). The hybrids also have excellent activity against *A. haemolyticus* and *P. aeruginosa* with an MIC in the range of 0.5–2 µg/mL, compared with the standard drug gentamycin (MIC = 0.78 µg/mL, respectively, MIC = 1.56 µg/mL). Against *Gram-positive* strain *E. faecalis* the hybrids have excellent activity with an MIC in the range of 0.5–8 µg/mL, compared with the standard drug ampicillin (MIC = 12.5 µg/mL).

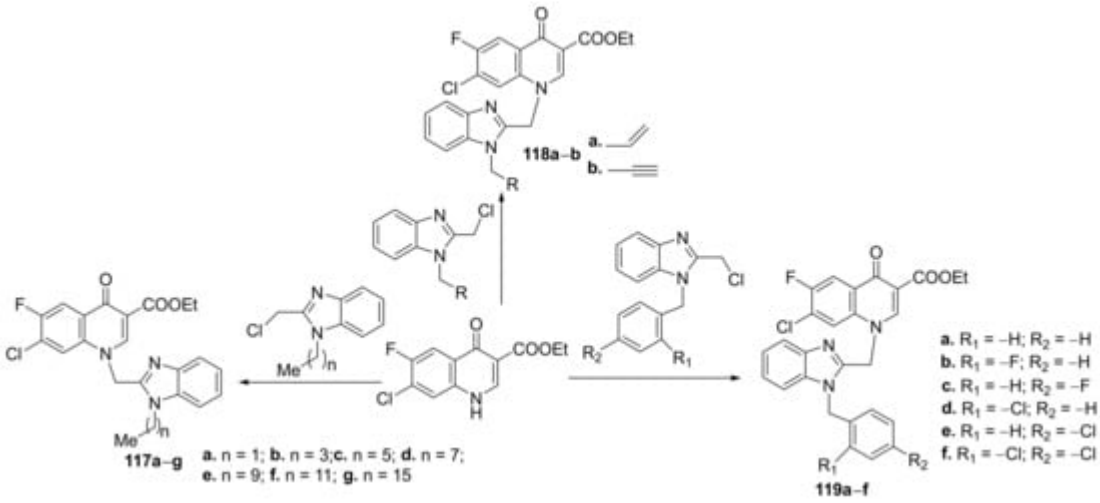
Guo et al. [41] synthesized a library of oxadiazole-quinoline hybrids and studied their antibacterial properties. The reaction pathway is straight, involving an alkylation reaction of fluoroquinolone with the corresponding oxadiazole, when the desired oxadiazole-fluoroquinolone hybrids **116a**–**t** were obtained, Scheme 36.



Scheme 36. Reaction pathway to obtain oxadiazole-fluoroquinolone hybrids **116a**–**t**.

The synthesized oxadiazole-fluoroquinolone hybrids **116a**–**t** were tested for their antibacterial activity against methicillin-resistant *Staphylococcus aureus* (MRSA) and laboratory *Staphylococcus aureus*. The hybrids displayed good antibacterial activity, one of the compounds **116k** exhibited excellent antibacterial activity against both methicillin-resistant *S. aureus* and laboratory *S. aureus*, with an MIC in the range of 0.25–2 µg/mL, superior to control drug vancomycin (MIC = 2 µg/mL).

Wang et al. [42] synthesized a series of benzimidazole–quinoline hybrids and studied their antibacterial and antifungal properties. The reaction pathway involves an *N*-alkylation reaction of fluoroquinolone with the corresponding benzimidazole, when the desired benzimidazole-fluoroquinolone hybrids **117a**–**g**, **118a**,**b** and **119a**–**f**, were obtained, Scheme 37.

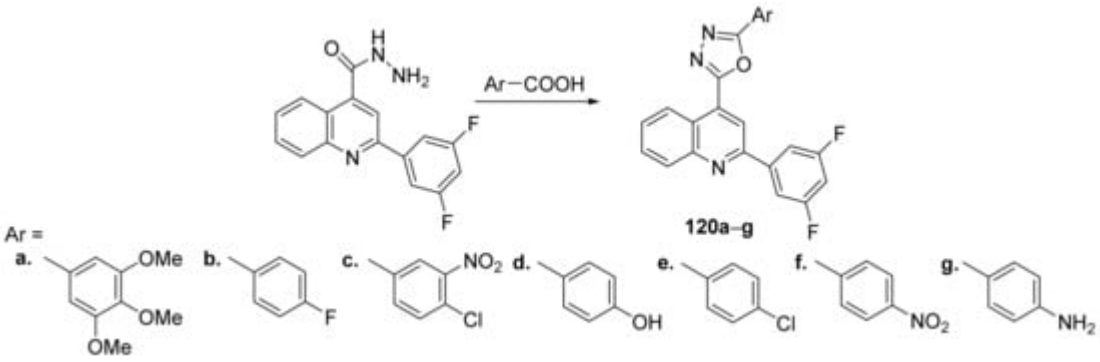


Scheme 37. Reaction pathway to obtain benzimidazole-quinoline hybrids **117a**–**g**, **118a**,**b** and **119a**–**f**.

The synthesized benzimidazole-fluoroquinolone hybrids **117a**–**g**, **118a**,**b** and **119a**–**f** were screened against *Gram-positive* and *Gram-negative* bacteria, respectively, fungus (methicillin-resistant *Staphylococcus aureus* (MRSA), *Enterococcus faecalis*, *Staphylococcus aureus*, *Staphylococcus aureus ATCC25923*, *Staphylococcus aureus ATCC29213*, *Klebsiella pneumonia*, *Escherichia coli*, *Pseudomonas aeruginosa*, *Acinetobacter baumanii*, *Pseudomonas aeruginosa ATCC27853*, *Escherichia coli ATCC25922*, *Candida albicans*, *Candida tropicalis*, *Aspergillus fumigatus*, *Candida albicans ATCC90023*, *Candida parapsilosis ATCC22019*). The results of the assay

were promising, with some hybrids having excellent antibacterial activity. The most active hybrids against *K. pneumonia* are **117a** and **117c**, with an MIC of 8 µg/mL, compared to the standard drug norfloxacin (MIC > 512 µg/mL).The most active hybrids against *S. aureus* are **119a** and **119f**, with an MIC of 4 µg/mL, compared to the standard drug norfloxacin (MIC = 64 µg/mL).

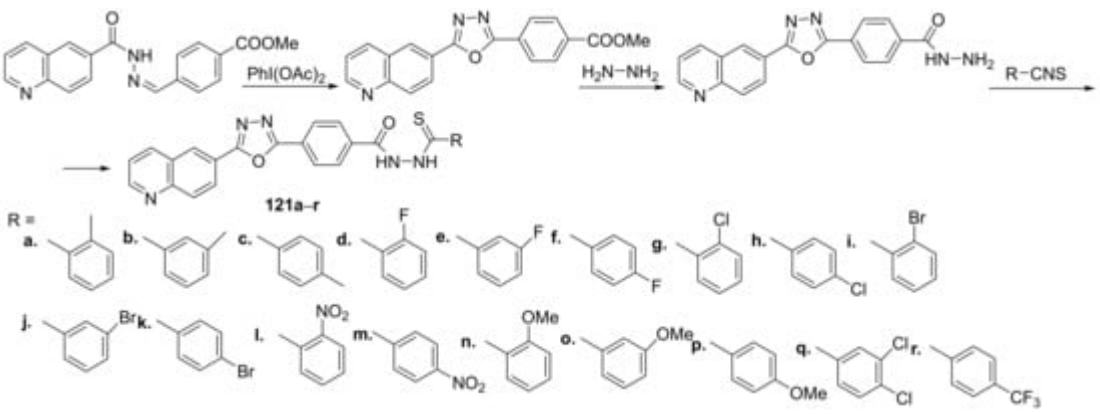
Bharadwaj et al. [43] synthesized a series of oxadiazole–quinoline hybrids and studied their antibacterial and antifungal properties. The reaction pathway involves a cyclocondensation reaction of hydrazinyl-quinoline derivative with the corresponding aromatic acids, when the desired oxadiazole–quinoline hybrids **120a**–**g** were obtained, Scheme 38.



Scheme 38. Reaction pathway to obtain oxadiazole-quinoline hybrids **120a**–**g**.

The synthesized oxadiazole–quinoline hybrids **120a**–**g** were tested against clinical isolates *Gram-positive* and *Gram-negative* bacteria (*Staphylococcus aureus*, *Bacillus cereus*, *Escherichia coli*, *Serratia marcescens*), respectively, fungus (*Aspergillus niger*, *Trichophyton mentagrophytes*, *Candida albicans*, *Candida parapsilosis*). The antimicrobial activity of oxadiazole–quinoline derivatives was good, the hybrids **120a** and **120f** having the best antimicrobial activity against *B. cereus* with an MIC of 17, respectively, 24 µg/mL, compared to standard drug ampicilin (MIC = 16 µg/mL).

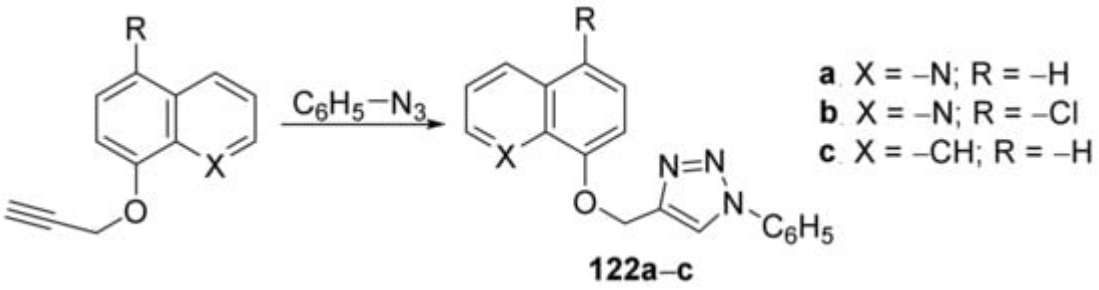
Tahaab et al. [44] synthesized a series of oxadiazole–quinoline hybrids and studied their leishmanicidal potential. The reaction pathway to obtain the oxadiazole–quinoline hybrids **121a**–**r** is depicted in Scheme 39.



Scheme 39. Reaction pathway to obtain oxadiazole-quinoline hybrids **121a**–**r**.

The synthesized oxadiazole-quinoline hybrids **121a**–**r** were tested for their leishmanicidal activity against *Leishmania major* promastigote. Most of the synthesized hybrids have a good leishmanicidal activity, compound **121r** was found to be the most active (IC_{50} = 0.10 μM) from the series, being 70 times more active than the standard drug (pentamidine, IC_{50} = 7 μM).

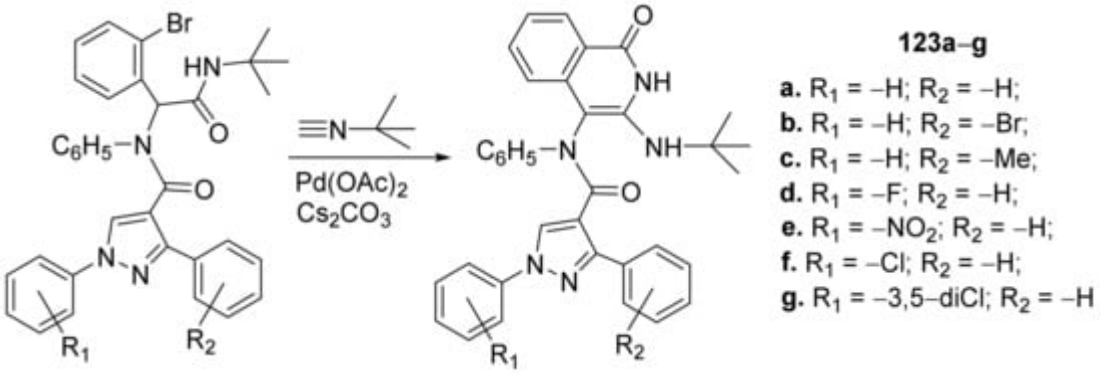
Irfan et al. [45] synthesized a series of triazole–quinoline hybrids and studied their antifungal properties. The reaction pathway involves a typical click cyclocondensation reaction of azide with a compound with a triple bond, when the desired triazole–quinoline hybrids **122a**–**c** were obtained, Scheme 40.



Scheme 40. Reaction pathway to obtain triazole-quinoline hybrids **122a**–**c**.

The synthesized triazole–quinoline hybrids **122a**–**c** were tested against fungus *Candida albicans*, both clinical isolates and laboratory strains [three FLC susceptible strains (*C. albicans D27*, *C. albicans D31* and *C. albicans D39*) and one FLC resistant strain (*C. albicans D15.9*)]. The best antifungal activity was found for the hybrids **122a** and **122b**, having an MIC of 25 μg/mL for **122a** and an MIC of 250 μg/mL for **122b**, compared to control FLC (MIC >1 μg/mL)

Pandya et al. [46] synthesized a library of pyrazole–isoquinoline hybrids and studied their antimicrobial properties. The reaction pathway involves a palladium-catalyzed reaction of pyrazole derivatives with *t*-butyl-isocyanide, when the corresponding pyrazole–isoquinoline hybrids **123a**–**g**, were obtained, Scheme 41.

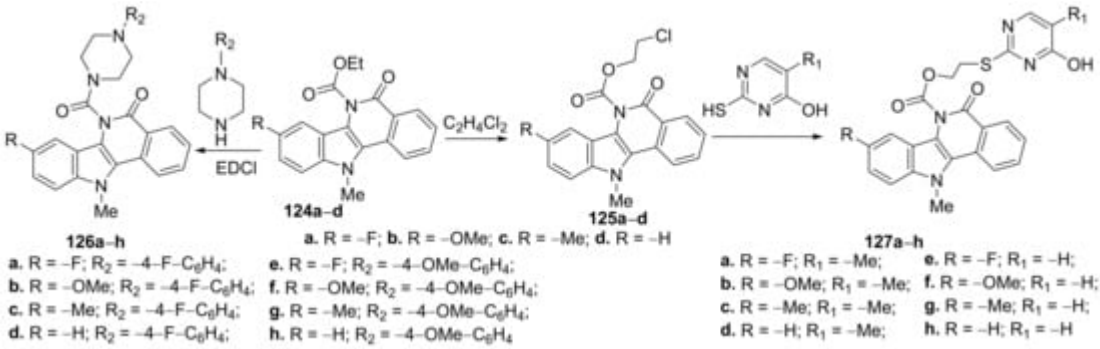


Scheme 41. Reaction pathway to obtain pyrazole-isoquinoline hybrids **123a**–**g**.

The synthesized pyrazole–isoquinoline hybrids **123a**–**g** were evaluated for their antimicrobial activity against different pathogenic strains: bacterial strains (*Staphylococcus aureus*, *Escherichia coli*, *Enterococcus faecalis*,

Streptococcus pyogens and *Vibrio cholera*), fungal strains (*Candida albicans*, *Candida glabrata*, *Candida krusei*, *Candida tropicalis* and *Candida parapsilosis*) and tubercular strain (*Mycobacterium tuberculosis*). The antimicrobial activity of hybrids was very good, the hybrids **123e** and **123g** having the best antimicrobial activity, compared to standard drugs kanamycin and amphotericin B. Thus, the most active hybrids against *S. aureus* are **123e** and **123g**, having an MIC of 20 µM, respectively, 37 µM, compared to standard drug kanamycin (MIC of 31 µM). The most active hybrids against *V. cholera* are **123e** and **123g**, having an MIC of 41 µM, respectively, 90 µM, compared to the standard drug kanamycin (MIC of 62 µM). The hybrids **123e** and **123g** have the best antitubercular activity against *M. tuberculosis* with an MIC of 30 µg/mL, respectively, 32 µg/mL, compared to standard drugs rifampicin and isoniazide (MIC of 90 µg/mL).

Verma et al. [47] obtained a series of piperazine- and pyrimidine- isoquinoline hybrids and studied their antimicrobial properties. The piperazine-isoquinoline hybrids **126a**–**h** were synthesized by condensation of the carboxylic acid intermediates **124a**–**d** with appropriate aryl-piperazines, Scheme 38. The pyrimidine-isoquinoline hybrids **127a**–**h** were synthesized in two steps: an *O*-alkylation of the carboxylic acid intermediates **124a**–**d** (with ethylene dichloride), followed by an *S*-alkylation of the obtained compounds **125a**–**d** (with thio-pyrimidine), Scheme 42.



Scheme 42. Reaction pathway to obtain piperazine- and pyrimidine-isoquinoline hybrids **126a**–**h** and **127a**–**h**.

The synthesized piperazine- and pyrimidine-isoquinoline hybrids **124a**–**h** and **125a**–**h** were evaluated for their antibacterial and antifungal (*Escherichia coli*, *Klebsiella pneumoniae*, *Staphylococcus aureus*, *Bacillus subtilis*, *Aspergillus niger*, *Aspergillus oryzae*, *Candida albicans* and *Pencillium chrysogenum*), antioxidant, anticancer and antituberculosis (*Mycobacterium tuberculosis*) activities. The antibacterial assay indicates that three hybrids, namely **124a**, **125a** and **126e** have the best activity against *E. coli* (with an MIC in the range of 1–3 µg/mL) and *K. pneumoniae* (with an MIC in the range of 1.5–3 µg/mL), compared with the standard drug ciprofloxacin (MIC = 1.5 µg/mL). The hybrids **125a**, **126a** and **127a** also have excellentactivity against *S. aureus* (with an MIC in the range of 1–3 µg/mL) and *B. subtilis* (with an MIC in the range of 1.5–3 µg/mL), compared with the standard drug ciprofloxacin (MIC = 1.5 µg/mL, respectively, MIC = 3 µg/mL). The hybrids **125a**, **126a** and **127a** have excellent activity against fungus *A. niger*, *C. albicans*, *A. oryzae*, and *P. chrysogenum* (with an MIC of 1.5 µg/mL), compared with the standard drug fluconazole (MIC = 1.5 µg/mL for *A. niger* and *C. albicans*, respectively, MIC = 3 µg/mL for *A. oryzae*, and *P. chrysogenum*). The hybrids **127b** and **127e** have the best activity against *M. tuberculosis* (MIC 1.0 mg/mL), compared with the standard drug rifampicin (MIC = 0.1mg/mL. The antioxidant and anticancer activity proved to be modest.

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