Naphthoquinones and Their Derivatives: Emerging Trends in Combating Microbial Pathogens

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Abstract: In the current era, an ever-emerging threat of multidrug-resistant (MDR) pathogens pose serious health challenges to mankind. Researchers are uninterruptedly putting their efforts to design and develop alternative, innovative strategies to tackle the antibiotic resistance displayed by varied pathogens. Among several naturally derived and chemically synthesized compounds, quinones have achieved a distinct position to defeat microbial pathogens. This review unleashes the structural diversity and promising biological activities of naphthoquinones (NQs) and their derivatives documented in the past two decades. Further, realizing their functional potentialities, researchers were encouraged to approach NQs as lead molecules. We have retrieved information that is dedicated on biological applications (antibacterial, antifungal, antiparasitic) of NQs. The multiple roles of NQs offer them a promising armory to combat microbial pathogens including MDR and the ESKAPE (Enterococcus faecium, Staphylococcus aureus, Klebsiella pneumoniae, Acinetobacter baumannii, Pseudomonas aeruginosa, and Enterobacter spp.) group. In bacteria, NQs may exhibit their function in the following ways (1) plasmid curing, (2) inhibiting efflux pumps (EPs), (3) generating reactive oxygen species (ROS), (4) the inhibition of topoisomerase activity. Sparse but meticulous literature suggests the mechanistic roles of NQs. We have highlighted the possible mechanisms of NQs and how the targeted drug synthesis can be achieved via molecular docking analysis. This bioinformatics-oriented approach will explicitly lead to the development of effective and most potent drugs against targeted pathogens. The mechanistic approaches of emerging molecules like NQs might prove a milestone to defeat the battle against microbial pathogens.

Keywords: efflux pumps; MDR; ESKAPE pathogens; naphthoquinones; plasmid curing; reactive oxygen species; topoisomerase

1. Introduction

Antibiotics represent world-class, assured molecules that have captured a gigantic share in the global market to combat ever-rising and prevalent infections. Over decades, different types of antibiotics have come into medical practice. Penicillin occupied the European and U.S. markets since its discovery in 1928 by Sir Alexander Fleming, followed by its commercial production in the 1940s [1]. Further, the world was gifted with the discovery of another antibiotic, streptomycin, by Albert Schatz, Bugie, and Waksman in
1943. This antibiotic was able to inhibit bacteria, predominantly the organisms responsible for tuberculosis [2]. After the success stories of penicillin and streptomycin, a huge number of antibiotics succeeded commercially, such that the projected rise of the global antibiotic market is up to US $67.25 billion by 2026 [3].

The challenges of resistance acquired by microbial pathogens towards existing antibiotics led to the advent of new antimicrobials. Presently, healthcare sectors are severely affected due to eternally escalating antimicrobial resistance (AMR) shown by pathogenic bacteria, parasites, viruses, and fungi. This serious threat associated with public health needs urgent attention and an immediate action plan from government policymakers, as well as private industries. It is essential to note that the challenges associated with AMR have led to a substantial cost escalation for pharmaceutical and health-care products. Patients suffering from microbial infections are ultimately the victims of long-term illness, and are therefore loaded with an additional monetary burden in the form of expensive tests and drugs [4,5]. There is also an increased morbidity and mortality rate in patients. These multidrug-resistant (MDR) pathogens are therefore referred to as “Super Bugs” [6]. MDR bacterial pathogens also comprise the ESKAPE group [7]. The abbreviation ‘ESKAPE’ has been used to designate a group having *Enterococcus faecium*, *Staphylococcus aureus*, *Klebsiella pneumoniae*, *Acinetobacter baumannii*, *Pseudomonas aeruginosa*, and *Enterobacter* spp. In the year 2008, Rice [8] had coined the terminology ‘the ESKAPE bugs’ to denote the ability of pathogens to escape from the lethal activity of antibiotics and impose severe menace to human health. These pathogens exhibit resistance to antimicrobial drugs like carbapenems, fluoroquinolones, glycopeptides, β-lactams, β-lactam–β-lactamase inhibitor combinations, lipopeptides, macrolides, tetracyclines, and polymyxins etc. [7]. In the year 2017, the World Health Organization (WHO) published a list of priority pathogens—Priority 1 (Critical), Priority 2 (High) and Priority 3 (Medium)—exhibiting resistance to antimicrobial agents [9]. These pathogens can worsen emergency situations and therefore, need urgent surveillance so that new and more effective compounds can be brought through the pipelines. Since 2015, WHO has introduced World Antibiotic Awareness Week (WAAW) to create awareness in the public community, health workforce and among policy-formulating personnel to restrict further emergence of antibiotic resistance and its spread. Since 2020, the Tripartite Executive Committee declared WAAW dates to be 18–24 November [10].

To gain AMR, pathogens acquire plasmids (R-drug resistance) or transposons and also possess multidrug efflux pumps (EPs) to drive out the drug molecules from their system [11]. Besides, other strategies are used, like (1) inactivation, alteration, or degradation of the drug by bacterial enzymes, (2) modification of drug binding sites on the bacterial cell, (3) biofilm formation (restricting the entry of the drug), and (4) reduction in intracellular drug accumulation [12]. Challenges associated with AMR encouraged researchers to explore a variety of naturally existing and chemically synthesized compounds over the decades. Since ancient times, medicinal plants have been evidenced as a great support to tackle dreaded illnesses [13]. Recent advances in the area of phytochemicals and synthesized derivatives have been looked forward to due to their multifunctional therapeutic approaches for dealing with AMR-associated challenges [14–17]. The unique structural, biological, and functional properties of naphthoquinones (NQs), along with their derivatives, have gained enormous consideration, particularly from a medicinal chemistry perspective [18]. NQs are widely distributed as natural pigments in plants, fungi, and some animals [19]. NQ derivatives bearing hydroxyl, methyl, nitrogen, sulfur, halide, phenylamino-phenylthio, or sulphide possess exceptional biological activity. Derivatives bearing hydroxyl groups are seeking more consideration due to their broad-spectrum pharmacological properties [20]. NQs possess widespread antibacterial, antiparasitic [21,22], antifungal, [23,24], antiviral [25], and antimalarial properties [26]. In the field of cancer biology, NQs are also noticeably identified for their abilities to produce reactive oxygen species (ROS) in cancer cell lines [27–29]. NQs are propitious candidates over the other chemotherapeutic drugs used currently. Until today, an ample number of NQ derivatives have been
analyzed for their functional potential against various pathogens. This encouraged us to present a comprehensive review of the structural diversity and multifunctional potentiality of NQs in medicinal chemistry. We also shed a light on the current understanding of the mechanistic roles of NQs in combating microbial infections. We also emphasize molecular docking—a powerful approach used in predicting the interactions of NQ molecules in biological systems.

2. Structural Diversity of Naphthoquinones Entities

Structurally, NQs constitute bicyclic structures with two carbonyl groups placed either in positions 1, 4 or 1, 2. The latter case is less frequent (Figure 1A). The chemistry and biological activities of quinones are primarily dependent on the position and chemical nature of the side groups attached (R). Groups like hydrogen, hydroxyl, methyl, nitrogen, sulfur, halide, etc. are attached to the NQ’s ring structure. Generally, the presence of a hydroxyl and/or methyl group in quinone structure is found in nature. These derivatives have been reported for a broad range of applications in pharmacology [19]. Recently, NQ derivatives isolated from plant sources including lawsone, juglone, plumbagin, shikonin and lapachol (Table 1) have fascinated researchers due to their (1) abundance, (2) structural diversity, and (3) broad-spectrum therapeutic potential [20]. It is imperative to state that 1,4-NQs derivatized at the 2nd and 3rd position with different chemical groups are recurrently reported for their biological properties (Table 1). NQs having oxygen entities at 1, 4 positions in the aromatic ring exhibit promising antimicrobial properties. Sparse literature is evident on 1,2-NQs. Realizing the significance of 1,4-NQ derivatives for biological applications, the major focus of this review remains on 1,4-NQ derivatives.

Table 1. Chemical structures of plant-originated naphthoquinone derivatives.

| Name and Structure of the Naphthoquinone Derivative | Source of Plant Material | Reference |
|----------------------------------------------------|--------------------------|-----------|
| (A) Juglone: 5-Hydroxy-1,4-naphthalenedione         | Caesalpinia sappan       | [30]      |
| (B) Plumbagin: 5-Hydroxy-2-methyl-naphthalene-1,4-dione | Plumbago zeylanica       | [31]      |
|                                                    | Plumbago auriculata      | [32]      |
|                                                    | Plumbago indica          | [33]      |
| (C) Lawsone: 2-Hydroxy-1,4-naphthoquinone          | Plumbago zeylanica       | [34]      |
|                                                    | Lawsonia inermis         | [35]      |
| (D) Shikonin: 5,8-Dihydroxy-2-[(1R)-1-hydroxy-4-methyl-3-penten-1-yl]-1,4-naphthoquinone | Arnebia euchroma         | [36]      |
|                                                    | Lithospermum erythrorhizon | [37]    |
| (E) Lapachol: 2-Hydroxy-3-(3-methyl-2-butenyl)-1,4-Naphthalenedione | Tabebuia ochracea        | [38,39]   |
Figure 1. Diversity of naphthoquinone molecules. (A): structures of chemically synthesized naphthoquinones. (B): production, purification, characterization, and biological applications of 1,4-naphthoquinone derivatives from plant material.
3. 1,4-Naphthoquinone Derivatives

The 1,4-NQs are redox-active compounds having resemblance to the structure of naphthalene [40]. Researchers have focused on 1,4-NQ derivatives to explore for antibacterial, antifungal, antiviral, anticancer, and antiparasitic potential. The following sections describe 1,4-NQs, both natural-origin and chemically synthesized, having immense antimicrobial potential against various pathogens, including those that are MDR or in the ESKAPE category.

3.1. 1,4-Naphthoquinone Derivatives of Natural Origin

Naphthoquinones, phenol-rich compounds, are seen abundantly in a variety of plants [30,32], animals [41], and fungi [42]. The functional potency of any antimicrobial agent is uniquely determined from its minimum inhibitory concentration (MIC). The MIC represents the lowest concentration that hampers the visible growth of microbes. Minimum bactericidal concentrations (MBCs) indicate the lowest concentration of a compound that prevents the growth of an organism by killing it. Therefore, MICs and MBCs are imperative for any antimicrobial agent to be considered for pharmacological applications. MICs of plant-derived NQs against pathogenic bacteria are represented in Table 2.

Table 2. Naphthoquinones extracted from plants along with their minimum inhibitory concentration (MIC).

| Type of Naphthoquinone | Active against Bacterial Strains | MIC (µg/mL) | Reference |
|------------------------|----------------------------------|-------------|-----------|
| Shikonin from *Arnebia euchroma* root extract | Staphylococcus aureus | 128 | [36] |
| | Streptococcus agalactiae | 128 |
| | *Escherichia coli* | 256 |
| | *Salmonella* isolates | 256 |
| | *Pseudomonas aeruginosa* | 512 |
| Plumbagin from *Plumbago zeylanica* | Staphylococcus aureus | 0.5 | [43] |
| | *Escherichia coli* | 8 |
| | *Klebsiella pneumoniae* | 2 |
| | *Pseudomonas aeruginosa* | 8 |
| Plumbagin from *Plumbago zeylanica* | *S. aureus* (MRSA) | 4–8 | [31] |
| | *S. aureus* (MSSA) | 7.8 |
| Lawsone from *Plumbago zeylanica* root extract | *Staphylococcus aureus* | 400 |
| | *Salmonella typhi* | 200 |
| | *Bacillus cereus* | 200 |
| | *Bacillus subtilis* | 200 |
| | *Pseudomonas aeruginosa* | 800 |
| | *Escherichia coli* | 800 |
| | *Shigella dysenteriae* | >1600 |
| | *Serratia marcescens* | >1600 |
| | *Proteus mirabilis* | 800 |
| | *Klebsiella pneumoniae* | 800 |
| | *Enterobacter* | 800 |
| | *Acinetobacter baumannii* | 800 |
| Naphthoquinone pigments from *Onosma visianii* | *Bacillus megaterium* | 9.54–54.28 |
| | *Monrichardia arborescens* | 6.82–38.10 |
| | *Micrococcus luteus* | 6.82–76.20 |
| | *Staphylococcus epidermidis* | 6.82–76.20 |
| | *Enterococcus faecalis* | 6.82–76.20 |
| | *Citrobacter koseri* | 6.82–76.20 |
| | *Hafnia alvei* | 6.82–38.10 |
| | *Pseudomonas proteolytica* | 4.77–6.82 |
| | *Stenotrophomonas maltophilia* | 4.77–6.82 |
| | *Yersinia intermedia* | 4.77–6.82 |
| Shikonin from *Lithospermum erythrorhizon* | *Staphylococcus aureus* (MRSA) | 7.8–31.2 | [37] |
| | *S. aureus* (MSSA) | 7.8 |
| *Plumbago auriculata* root extracts | *Proteus vulgaris* | Strong activity |
| | *Klebsiella pneumoniae* | Strong activity |
| | *Escherichia coli* | Moderate activity |
| | *Pseudomonas aeruginosa* | Less activity | [32] |
This section is devoted to the structural diversity and functional potentiality of NQs of plant origin. Various plants, namely, Plumbago zeylanica, Plumbago indica, Lawsonia inermis, Plumbago auriculata, Caesalpinia sappan, Onosma visianii, and Impatiens glandulifera, and solvents like methanol, chloroform, acetone, ethanol, petroleum ether, diethyl ether, benzene, acetonitrile (ACN), dimethyl sulfoxide (DMSO), dimethylformamide (DMF), ethyl acetate (EA), isopropyl alcohol (IPA), and water etc. have been reported to extract NQs. Among those organic solvents, ethanolic and methanolic extracts appear to result in a relatively purer form of NQs with limited interfering compounds. The overall protocol used to produce, purify, and characterize 1,4-NQs from plant material is illustrated in Figure 1B. Juglone, plumbagin, lawsone, shikonin, and lapachol are a few examples of 1,4-NQs found profusely in nature and are also recognized for their substantial antimicrobial potential [46]. Lim et al. [30] extracted 5-hydroxy-1,4-naphthoquinone (juglone) (A in Table 1) from Caesalpinia sappan heartwood and examined its activity against intestinal cultures, viz. Bifidobacterium bifidum, Bifidobacterium breve, Clostridium perfringens, Escherichia coli, and Lactobacillus casei. Authors had purified those bioactive molecules using different solvents, like methanol, EA, butanol, chloroform, hexane, and water. The powerful growth-inhibition activity of NQ was obtained from EA, butanol, and methanol extracts against B. bifidum and C. perfringens. Chloroform fractions resulted in moderate antimicrobial activity, whereas water and hexane fractions were ineffective against both pathogens. The scenario presented here is quite interesting and justifies the selectivity and specificity of bioactive compounds isolated from C. sappan heartwood using different solvents. Authors also screened commercially accessible compounds like 5-hydroxy-1,4-NQ, 5-hydroxy-2-methyl-1,4-NQ, 1,4-NQ, and 1,2-NQ at concentrations of 0.25, 0.5, 1, 2, and 5 mg/disk. Their analysis recommended the dose-dependent activity of 1,4-NQ, and 1,2-NQ against C. perfringens, rather than other isolates used by them. Additionally, the 1,4-NQ derivative showed superior activity against four other organisms (B. bifidum, B. breve, E. coli, and L. casei) as compared to other derivatives (5-hydroxy-1,4-NQ and 5-hydroxy-2-methyl-1,4-NQ). Results also confirmed that the antibacterial activity of 1,4-NQ was reduced significantly due to the presence of a \(-\text{CH}_3\) group at the 2nd and an \(-\text{OH}\) group at the 5th position.
Plumbagin (B in Table 1) is another widely explored secondary plant metabolite. Plumbagin is of immense interest due to its abilities to (1) generate ROS, (2) inhibit EPs, (3) accumulate drugs, and (4) cure plasmids from bacteria. Thus, due to such exceptional properties, it has been utilized as an effective antibacterial agent [47]. Periasamy et al. [31] evaluated the antibacterial activity of plumbagin (from Plumbago zeylanica) against 100 methicillin-resistant S. aureus (MRSA) strains, including MDR phenotypes. P. zeylanica root extract was obtained using ACN, DMSO, DMF, IPA, methanol and water. All fractions were used to analyze their antibacterial potential where active constituents of ACN and (32 to 128 µg/mL) against the other three biofilm-forming pathogens. The activity of those three antibiotics was enhanced up to 2-fold when they were used in combination with plumbagin. Surprisingly, the enhancement of the 2-fold activity of ciprofloxacin was found against P. aeruginosa, whereas the activity of amoxicillin and ampicillin was reduced to 2-fold against the same isolate. For K. pneumoniae, the activity of amoxicillin and ampicillin was found to be enhanced up to 6-fold. At the same time, the efficiency of ciprofloxacin was reduced by 2-fold for K. pneumoniae [43]. The synergistic approaches are unquestionably valuable to prove the potential of plumbagin as a favorable drug. Adusei et al. [43] further reported the considerable antibiofilm activity of plumbagin (32 to 128 µg/mL) against the above-mentioned five pathogens. The antibiofilm activity of plumbagin was noticeably higher against S. aureus as compared to ciprofloxacin. Both plumbagin and ciprofloxacin displayed antibiofilm activity (>50%) against the other three pathogens (S. aureus, E. coli, and K. pneumoniae) at 128 µg/mL concentration. The combination of plumbagin with other drugs undoubtedly impede the growth and infection of biofilm-forming pathogens.

The literature suggests that plumbago species has always been a choice NQ among the ones of natural sources. Patwardhan et al. [32] documented the effectiveness of P. auriculata root extract against 23 nosocomial pathogenic strains including P. aeruginosa, P. vulgaris, E. coli, and K. pneumoniae. Solvent extracts of plumbago roots were evaluated at various concentrations from 250 to 4000 µg/mL. Plumbago extract obtained using ethanol displayed the highest antimicrobial activity as compared with other solvents. Ethanolic based extracts impeded the growth of P. vulgaris and K. pneumoniae, followed by E. coli. It is important to mention that P. aeruginosa was found to be the least affected with the same ethanolic plumbago extract.

A report documented by Kaewbumrung and Panichayupakaranant [33] explains the stability and the yield of plumbagin from P. indica roots. For the extraction procedure, authors used ethanol, EA, DCM, diethyl ether, and isopropanol. Out of those five solvents, ethanol was found to be the most suitable to result extract (11.5% w/w) having a high amount (5.79 mg/g) of the plumbagin. Further purification and elution of pure plumbagin was achieved using a silica gel column with a mixture of hexane: EA (9.2%:0.8% v/v). This step enhanced the total plumbagin content (13.26% w/w). Impurities soluble in hexane and EA were removed to result in a purified form of a derivative. The promising bactericidal activity of plumbagin extract was observed against S. aureus, S. epidermidis, and P. acnes. Like EA, petroleum ether has been the solvent of choice to extract NQs from plant sources. Vukic et al. [44] used petroleum ether along with EA to isolate NQs
from \textit{Onosma visianii}. Thus, the selection of appropriate solvents is important to obtain biologically active components from plant materials. From the above discussed literature, we underscore the pivotal role of solvents to extract NQs from natural sources. Attention to this concept could be helpful to enhance the overall yield of NQs from the desired biological sources. In conclusion, plumbago species are one of the preferred natural sources to isolate NQs. Among various solvents, ethanol is a virtuous choice for the extraction of bioactive compounds, followed by EA and hexane to demonstrate the assured biological activity of NQs.

Like plumbagin, another NQ—namely lawsone (C in Table 1)—has stimulated researchers to explore its medicinal potential. The foremost report on the extraction of lawsone (2-hydroxy-1,4-NQ) from \textit{P. zeylanica} was accomplished by Patwardhan et al. [34]. Authors extracted lawsone via the solvents acetone, benzene, chloroform, cyclohexane, diethyl ether, ethanol, methanol, and petroleum ether. Consequently, Patwardhan and collaborators [34] focused on the biological activity, purification, and characterization of lawsone extracted using ethanol. The antibacterial potential of ethanolic root extract (ranging from 200 to 800 $\mu$g/mL) was effective against three Gram-positive cultures (\textit{S. aureus}, \textit{B. cereus}, and \textit{B. subtilis}) and six Gram-negative cultures (\textit{E. coli}, \textit{S. typhi}, \textit{Enterococcus} spp., \textit{K. pneumonia}, \textit{A. baumannii}, and \textit{S. dysenteriae}). However, the same ethanolic extract was operative against \textit{S. marcescens} and \textit{Proteus mirabilis} at higher concentrations (>1600 $\mu$g/mL). Patwardhan et al. [34] also examined the plasmid-curing efficiency of lawsone, which is discussed briefly in Section 5.1.

A liposoluble red-colored pigment, shikonin (D in Table 1), is usually extracted from the roots of various plants like \textit{Alkanna tinctorial}, \textit{Lithospermum erythrorhizon}, or \textit{Arnebia decumbens} L. Shikonin is one of the unique bioactive NQs routinely extracted from the roots of \textit{L. erythrorhizon}, and is popular for several biological activities.

Lee et al. [37] isolated shikonin from the roots of \textit{L. erythrorhizon} and examined its antibacterial potency against seven MRSA strains. This study revealed the MIC of shikonin to be 7.8 to 31.2 $\mu$g/mL. MRSA strains were found to be more susceptible to shikonin, as compared to ampicillin and oxacillin (antibiotics of the penicillin class, cell wall attacking). Shikonin was evaluated in combination with Tris and Triton X-100 (membrane-permeabilizing agents), sodium azide and N,N'-dicyclohexylcarbodiimide (ATPase inhibitors), and peptidoglycan (derived from \textit{S. aureus}). The work was supported through experiments like (1) the broth microdilution method (to analyze the susceptibility of bacteria to antibacterial agents), (2) the time–kill test (to determine the bactericidal/bacteriostatic activity of antibacterial agents over time), and (3) transmission electron microscopy (predicting the underlying mechanism of antibacterial agents affecting cell morphology). Studies have discovered the enhanced antibacterial activity of shikonin in the presence of Tris and Triton X-100 and ATPase inhibitors. Lee et al. [37] suggested that shikonin can be proposed as a natural antibiotic and even to realize the underlying mechanism responsible for antimicrobial action. Al-Mussawi [48] had also isolated shikonin from \textit{Arnebia decumbens} L. and purified it using column chromatography. Researchers proved the antibacterial activity of shikonin against \textit{P. aeruginosa}, \textit{E. coli}, \textit{S. aureus}, and \textit{K. pneumonia}. Moreover, shikonin has been widely explored for anti-inflammatory, wound-healing, antithrombotic, and anticancer properties [49]. Large-scale production of any bioactive molecule is mandatory to utilize them for pharmacological purposes. When NQs are extracted from natural plant sources, the yield and purity cannot be neglected. Recently, Huang et al. [36] developed an ultrasound-assisted extraction (UAE) technique to isolate shikonin from \textit{A. euchroma}. The response surface methodology (RSM) was used to design an appropriate extraction set-up for the production of shikonin. This experiment set-up can encourage researchers to extract shikonin at ease using an energy-saving approach. Authors magnificently reported around a 1.26% yield of shikonin under the ideal extraction protocol using ultrasound power at 93 W (in 87 min at 39 $^\circ$C) with a liquid–solid ratio of 11:1. The same studies supplemented the antimicrobial activity of shikonin with clinical isolates (three) along with standard ones (five) at MICs of 128–1024 $\mu$g/mL. Extracts obtained from the medicinal plants find
applications in the manufacturing of ointment to treat patients with burn infections. Aljanyby [50] isolated an aqueous extract having alkannin esters and shikonin from A. tinctoria and demonstrated antibacterial activity against drug-resistant bacterial strains (395) at a 300 mg/mL concentration.

Among different NQs mentioned above, lapachol (E in Table 1) has been regularly reported to have antibacterial and anticancer properties [51]. Lapachol has been derivatized to produce a pharmacologically important molecule. Zani et al. [38] had isolated lapachol from Tabebuia ochracea (Bignoniaceae family), which was further derivatized by Souza et al. [39] to produce thiosemicarbazone and semicarbazone. In an antibacterial assay, E. faecalis and S. aureus were found to be susceptible to thiosemicarbazone lapachol (0.05 µmol/mL) and semicarbazone lapachol (0.10 µmol/mL). Along with Bignoniaceae, many other families like Verbenaceae, Proteaceae, Leguminosae, Sapotaceae, Scrophulariaceae, and Malvaceae show broad scope to extract lapachol.

An interesting report by Balachandran et al. [52] documented the isolation of an antibacterially bioactive 1,4 NQ molecule from Streptomyces sp., named bluemomycin, using EA. The EA extract displayed potent antibacterial activity against both Gram-negative bacteria as well as Gram-positive bacteria.

In the view of the present scenario, it is also vital to communicate that the microorganisms used for antimicrobial assays are S. aureus, P. aeruginosa, and E. coli, which belong to the ESKAPE category. At lower MICs, most NQs of natural origin have demonstrated better activities against Gram-positive (Staphylococcus spp., Bacillus spp.) than Gram-negative bacteria. Among Gram-negative bacteria, Pseudomonas spp. and E. coli have been frequently tested to determine the antibacterial potency of NQs (Figure 2).

3.2. Chemically Synthesized 1,4-Naphthoquinone Derivatives

Just like those of a natural origin, chemically derivatized 1,4-NQs are widely investigated for antimicrobial potential. Literature suggests that most of the additions or deletions of the chemical groups have been carried out at 2nd and 3rd positions in NQ moieties. The groups preferred for additions are H, OH, CH₃, Cl, Br, N, and S. Most of the chemical elements chosen for derivatization are placed from 14 to 17 in the periodic table. These atoms are typically reactive non-metals and strong oxidizing agents.

Chemical modifications in the NQ moieties have been facilitated to explore them for a wide range of biological applications. Ravichandiran et al. [53] synthesized a series of NQs containing phenylamino-phenylthio moieties and evaluated their antibiotic activity against S. aureus, Listeria monocytogenes, E. coli, P. aeruginosa, and K. pneumonia with MICs ranging between 15.6 to 500 µg/mL. Most of the synthesized NQ derivatives were able to inhibit S. aureus and E. coli. However, NQ derivatives were found to be less effective against the other three strains. The structure–activity relationship (SAR) suggested that the introduction of the thiophenol group in one of the structures can exhibit moderate antibacterial activity as compared to its ligand. This study revealed that the introduction of a substituted amide group, acid chlorides (aliphatic and aromatic), in one of the compounds shows superior antibacterial properties as compared with its parent compound. Likewise, a compound containing a 3,5-dinitro aryl moiety displayed enhanced antibacterial activity against selected pathogens. Contrary to this, synthesized compounds having a bulky moiety or electron-withdrawing groups (NO₂ and methyl) showed low inhibitory actions against test cultures. This can be further justified by the fact that the bulky groups cannot enter or penetrate easily into the bacterial cell and thus, encounter challenges to fit at the target sites [53]. The synthesis of quinones substituted with N-, S-, O- and the evaluation of their antimicrobial and anticancer activities was performed by Kurban et al. [18]. For the first time, researchers documented the efficiency of benzo- and NQ derivatives to inhibit an enzyme, catalase (which is responsible for the decomposition of H₂O₂ into H₂O and O₂ to protect cells from oxidative damage).

From the literature, it is noteworthy that the bacterial species of the ESKAPE category are extensively tested to assess the efficacies of chemically synthesized NQ derivatives. NQs
can impose greater effects against Gram-positive bacteria. However, it is also important to highlight the exceptional cases wherein Gram-negative strains like K. pneumoniae and E. coli are also susceptible to some of the NQ derivatives [53,54]. The mechanism of action might be influenced by the constituents of the cell wall. Also, the interaction of the functional groups of NQ derivatives at the target position is another parameter which cannot be neglected. The addition or deletion of a single chemical group (for example OH, Cl, Br, S, N, or O) to or from a specific position in a NQ structure can affect the efficacies of the NQ against targeted pathogens. Table 3 presents the antibacterial potential of chemically synthesized NQ derivatives along with their MICs.

![Figure 2](image-url)

**Figure 2.** Percentage-wise distribution of bacterial species used to evaluate the antibacterial activity of (A): naturally derived and (B): chemically synthesized 1,4-naphthoquinones.
Yap et al. [55] put forth findings that demonstrated the synergistic relationship between 1,4-NQ and antibiotics like imipenem, cefuroxime, and cefotaxime against methicillin-resistant *Staphylococcus aureus* (MRSA). Authors suggest that 1,4-NQ could be developed into an adjuvant that will help in enhancing the function of antibiotics against drug-resistant bacteria as a part of combination therapy.

**Table 3.** Antibacterial potential of chemically synthesized naphthoquinones along with their minimum inhibitory concentration (MIC).

| Type of Naphthoquinone | Antibacterial Activity Against | MIC (µg/mL) | Reference |
|------------------------|-------------------------------|-------------|-----------|
| Plumbagin              | *Mycobacterium tuberculosis*  | 4           | [56]      |
|                        | *Mycobacterium smegmatis*     | 4           |           |
| Lawsone                | *Mycobacterium tuberculosis*  | >16         |           |
|                        | *Mycobacterium smegmatis*     | >32         |           |
| Lawsone methyl ether   | *Staphylococcus aureus* (MRSA)| 62.5–125    | [57]      |
| Lawsone derivatives    | *Staphylococcus aureus* (MRSA)| 32–128 & >128| [58]      |
|                        | *Staphylococcus aureus* (MSSA)| 0.6–128 & >128|           |
| Lapachol               | *Staphylococcus aureus*       | 1.25 mM     | [59]      |
|                        | *Staphylococcus aureus* (MRSA)| 30–500 & >500| [60]      |
| Imidazole derivatives  | *Staphylococcus aureus*       | 8–512       | [61]      |
| of 1,4-naphthoquinone  | *Bacillus subtilis*           | 32–512      |           |
|                        | *Pseudomonas aeruginosa*      | 8–256       |           |
|                        | *Proteus vulgaris*            | 16–256      |           |
|                        | *Escherichia coli*            | 32–256      |           |
| 1,4-naphthoquinone derivatives | *Staphylococcus aureus* | 4–256 | [62] |
|                        | *Bacillus subtilis*           | 128–512     |           |
|                        | *Pseudomonas aeruginosa*      | 32–252      |           |
|                        | *Proteus vulgaris*            | 128–512     |           |
|                        | *Escherichia coli*            | 256–512     |           |
| Phenylamino-phenylthio derivatives of 1,4-naphthoquinone (no. of derivatives synthesized) | *Staphylococcus aureus* | 31.25–500 | [53] |
|                        | *Listeria monocytogenes*      | 62.5–500    |           |
|                        | *Pseudomonas aeruginosa*      | 62.5–500    |           |
|                        | *Escherichia coli*            | 15.6–500    |           |
|                        | *Klebsiella pneumoniae*       | 62.5–500    |           |
| Sulfide derivatives of 1,4-naphthoquinone | *Staphylococcus aureus* | 7.8–250 & >250 | [24] |
|                        | *Escherichia coli*            | 31.3–250 & >250|       |
| Menadione               | *Staphylococcus aureus*       | 128         | [63]      |
|                        | *Pseudomonas aeruginosa*      | 64          |           |
|                        | *Escherichia coli*            | 128         |           |
|                        | *Klebsiella pneumoniae*       | 128         |           |
| Arylsulfanyl-methyl-[1,4]-naphthoquinone derivatives | *Staphylococcus aureus* | 32–256 | [64] |
|                        | *Staphylococcus epidermidis*  | 16–256      |           |
|                        | *Staphylococcus simulans*     | 32–256      |           |
|                        | *Escherichia coli*            | 256 (less activity) | 32–256|
| 1,4-naphthoquinone substituted at positions 2 and 3 | *Staphylococcus aureus* | 7.8–500 | [65] |
|                        | *Salmonella bongori*          | 125–500     |           |
|                        | *Pseudomonas aeruginosa*      | 125–500     |           |
|                        | *Proteus vulgaris*            | 250–500     |           |
|                        | *Escherichia coli*            | 125–500     |           |
|                        | *Klebsiella pneumoniae*       | 62.5–500    |           |
|                        | *Enterococcus faecalis*       | 125–500     |           |
|                        | *Enterobacter cloacae*        | 250–500     |           |
| Shikonin derivatives   | *Staphylococcus aureus*       | 1.1–45.4    | [66]      |
|                        | *Bacillus subtilis*           | 2–50 & >50  |           |
|                        | *Escherichia coli*            | 3.1–50 & >50|           |
|                        | *Pseudomonas aeruginosa*      | 4–50 & >50  |           |
Table 3. Cont.

| Type of Naphthoquinone                      | Antibacterial Activity Against                      | MIC (µg/mL) | Reference |
|---------------------------------------------|----------------------------------------------------|-------------|-----------|
| Naphthoquinone derivatives                  |                                                    |             | [19]      |
| Staphylococcus aureus                       |                                                    | 16–256      |           |
| Staphylococcus epidermidis                  |                                                    | 128         |           |
| Bacillus cereus                             |                                                    | 256         |           |
| Salmonella enterica                         |                                                    | 256         | [67]      |
| Listeria monocytogenes                      |                                                    |             |           |
| Pseudomonas aeruginosa                      |                                                    | 128         |           |
| Escherichia coli                            |                                                    | 128         |           |
| Enterococcus faecalis                       |                                                    | 256         |           |
| Vibrio alginolyticus                        |                                                    | 256         |           |
| 2-bromo-5-hydroxy-1,4-NQ                    | Staphylococcus aureus                             | 16          |           |
| Juglone                                     |                                                    |             |           |
| Staphylococcus aureus                       |                                                    | 32–256      |           |
| Staphylococcus epidermidis                  |                                                    | 128         |           |
| Bacillus cereus                             |                                                    | 256         |           |
| Salmonella enterica                         |                                                    | 256         | [67]      |
| Listeria monocytogenes                      |                                                    |             |           |
| Pseudomonas aeruginosa                      |                                                    | 128         |           |
| Escherichia coli                            |                                                    | 128         |           |
| Enterococcus faecalis                       |                                                    | 256         |           |
| Vibrio alginolyticus                        |                                                    | 256         |           |
| Nitrogen and sulfur derivatives of 1,4-naphthoquinone | Bacillus subtilis                              | 1.4–19.3    | [68]      |
| Plumbagin, juglone, lawsonex, menadione and their analogues | Staphylococcus aureus (MRSA) | 3.9–125 | [69] |
| Plumbagin, juglone, lawsonex, menadione and their analogues | Pseudomonas aeruginosa | No significant activity | |
| Menadione                                   |                                                    |             | [70]      |
| Staphylococcus aureus                       |                                                    | 3.1         |           |
| Bacillus anthracis                          |                                                    | 6.25        |           |
| Streptococcus pyogenes                      |                                                    | 25          |           |
| Streptococcus agalactiae                    |                                                    | 6.25        |           |
| 1,4-naphthoquinone                          |                                                    |             |           |
| Staphylococcus aureus                       |                                                    | 6.25        |           |
| Bacillus anthracis                          |                                                    | 12.5        |           |
| Streptococcus pyogenes                      |                                                    | 50          |           |
| Streptococcus agalactiae                    |                                                    | 12.5        |           |
| 2-hydroxy, 1,4 naphthoquinone (Lawsonex) and its 2-hydroxy naphthoquinone derivatives | Staphylococcus aureus | 16–512 & >512 | [71] |
| Plumbagin derivatives                       | Mycobacterium smegmatis                           | 13.3–30.4   | [72]      |
| Plumbagin derivatives                       | Mycobacterium tuberculosis                        | 15.6–77.4   |           |
| Nitrogen, sulfur groups substitution at 2,3 positions of 1,4-naphthoquinone | Staphylococcus aureus | 6.25–50 & >50 | [73] |
| 5-Hydroxy-2-methyl-1,4-NQ                   | Clostridium perfringens                           |             |           |
| 1,4-naphthoquinone                          |                                                    |             |           |
| Lactobacillus casei                         |                                                |             |           |
| Bifidobacterium bifidum                     |                                                |             |           |
| Bifidobacterium breve                       |                                                |             |           |
| Clostridium perfringens                     |                                                |             |           |
| Escherichia coli                            |                                                |             |           |
| 1,2-naphthoquinone                          |                                                |             |           |
| Clostridium perfringens                     |                                                |             |           |
| Bifidobacterium bifidum                     |                                                |             |           |
| Bifidobacterium breve                       |                                                |             |           |
| 5-Amino-8-Hydroxy-1,4-NQ                    | Staphylococcus aureus                            | 50          | [74]      |
| (L)-a-amino acid methyl ester, heteroalkyl and aryl substituted 1,4-naphthoquinone derivatives | Staphylococcus aureus | 12.5–50 & >50 | [54] |
| 1,4-naphthoquinone                          | Staphylococcus aureus                            | 10          |           |

MRSA/MSSA: methicillin-resistant/sensitive Staphylococcus aureus.
From the overall literature survey, 1,4-NQs extracted from natural sources and synthesized by chemical means propose their extraordinary candidature against pathogens. The percentage-wise distribution of bacterial species evaluated to demonstrate antibacterial potential of 1,4-NQs is depicted in Figure 2. *Staphylococcus* spp. belonging to the ESKAPE and MDR pathogen groups has been reported frequently for its susceptibility towards natural (~20%) and chemically synthesized (~27%) NQs. Followed by *Staphylococcus* spp., other Gram-positive bacteria, namely, *Bacillus* spp. (~16%), have been used to validate the antibacterial activity. Among Gram-negative bacteria, *E. coli* (~11%) and *Pseudomonas* spp. (~9%) seemed to be used for in vitro assays. In addition to the above-mentioned prominent bacteria, several other organisms have been utilized in experiments. Like the natural NQs, chemically synthesized NQ derivatives have generally been tested against *Pseudomonas* spp. and *E. coli* (both members of the ESKAPE and MDR pathogen groups). Organisms, viz. *E. faecium*, *K. pneumoniae*, *A. baumannii*, and *Enterobacter* spp. of ESKAPE, have been assessed for the antibacterial efficiency of NQs (Figure 2). The pragmatic analysis of 1,4-NQs, overtly showcases them as an emerging drug against the targeted pathogens.

4. Structure–Activity Relationship and Bioinformatics Approach: Determining the Possible Mechanistic Role of 1,4-Naphthoquinones at the Molecular Level in a Cell

Currently, computational tools like artificial intelligence and machine learning have been suggested to enhance the simulation and modeling processes for nanotherapeutics [75–77]. SAR studies aid in understanding the probable mechanistic role of NQs at the molecular level in a biological system. It can be figured out that the chemical modifications of NQs, through the addition or removal of a specific chemical group like the halides, thiolated at particular positions (C-2 and C-3) in a parent molecule, can drastically affect its activity. Research conducted by Wellington et al. [24] highlights the SAR of the NQ derivatives with an antibacterial effect on *E. coli*. The activity of these compounds was dependent on the fluoro group present on the phenyl ring. The fluoro group at metaposition exhibits better activity (MIC 31.3 µg/mL) than that at para-position (93.3 µg/mL). The addition of two fluoro groups, one at meta- and other at para-position, drastically reduced the antibacterial activity of chemically synthesized NQ derivatives (MIC of 187.5 µg/mL). Also, the addition of 3-sulfanylp propaneic acid to the original compound (1,4-NQs) displayed the weakest activity (250 µg/mL) against *E. coli*. Another study conducted by Sánchez-Calvo et al. [19] presented exciting data on the presence of halogen derivatives (Cl, Br) at the C-2, and C-3 positions, while the carbonyl group at the C-1 and C-4 positions influence the antimicrobial activity of NQs. Furthermore, side chains with >10 carbon atoms can diminish the antimicrobial potential of NQs due to the upsurge in lipophilic character. The hydroxyl group at C-5 and/or C-8 also enhances the activity of NQ derivatives up to a noticeable limit. If it is further replaced by other functional groups, then antimicrobial activity is reduced considerably. This proves the importance of core rings of NQ with respect to antibacterial activity. Ravichandiran et al. [68] evaluated nitrogen- and sulfur-bearing quinone derivatives for antibacterial activity. Heterocyclic quinone derivatives bearing a nitro group (with strong electron-withdrawing power) exhibit momentous antibacterial and anticancer potential.

The molecular docking approach is another hopeful area to reveal better understanding about the interactions of chemical entities at the molecular level. Ravichandiran et al. [53] successfully studied the possible molecular interactions of the selected NQ derivative (5a) against the crystal structure of the cytoplasmic chaperone, DmsD protein, (which is required for the synthesis of dimethyl sulphoxide-DMSO reductase) belonging to *E. coli*. Structurally, the 5a derivative has a hydrogen atom at a significant position. The docking analysis confirmed the binding of the ligand in the cavity of the DmsD protein molecule. Authors documented the formation of a hydrogen bond with ARG A15 (arginine A15 residue), with a corresponding active site, along with π–π stacking. Authors also dragged readers’ attention towards electrochemical studies on the reduction potential of the derivatives. The compound 5a has the highest reduction potential, which correlates to
the lower MIC value. This justifies the hypothesis of the significance of redox behavior in governing the antibacterial activity of NQs.

Computational modeling is also a trustworthy approach to predict the functioning of a complex system supported through computer simulation. Figueredo et al. [60] investigated the pharmacological potential of 2-(2-hydroxyethylamine)-3-(3-methyl-2-butenyl)-1,4-dihydro-1,4-naphthalenedione, 2-(2-hydroxy-ethylamine)-3-(2-methyl-propenyl)-(1,4) naphthoquinone, and 2(3-hydroxy-propylamine)-3-(3-methyl-2-butenyl)-(1,4) naphthoquinone alkaloid analogues of lapachol and nor lapachol. The antibacterial-potential study was also supplemented with computational prediction modeling data. In silico studies (through ChEMBL database), it has been revealed that 2-(2-hydroxy ethylamine) -3-(2-methyl-propenyl)-(1,4) naphthoquinone could be a possible drug to target replicative DNA helicase and RecA. Another report published by Figueredo et al. [78] explained the possible interaction of alkaloid analogues derived from lapachol and nor lapachol with the NorA efflux pump (in \textit{S. aureus}). This association study, conducted with molecular docking, indicated noticeable reduction in MICs (Table 4) and also the inhibition of the activity of the NorA efflux pump with high affinity.

A few years back, Qiu et al. [66] used a docking simulation study to approach the shikonin compound as a tyrosyl-tRNA synthetase (TyrRS) inhibitor. Scaffold modification of shikonin was carried out through in vitro primary screening. This study continued the demonstration of the inhibitory effect of modified shikonin against TyrRS and selected bacterial strains. One of the derivatives, namely, PMM-154, was identified as the most potent compound against \textit{S. aureus} ATCC 25923 (MIC of 1.1 mg/mL) with the apoptosis effect. The outcome of this study confirmed the modification of the phenyl ring of phenylpiperazine at the third position by trifluoromethyl to propose PMM-154 as a promising TyrRS inhibitor to deal with infections caused by microbial pathogens.

5. Possible Mechanistic Roles of Napthoquinones

The multifunctionalities of NQs, like antibacterial, antifungal, antiparasitic, etc. functions, make them a propitious armory to combat MDR and ESKAPE pathogens. Sparse but meticulous literature is available on the mechanistic roles of NQs against pathogens. NQs are expansively studied for their anticancer activities through the production of ROS and subsequent apoptosis of cancer cells [79,80]. NQs are also recognized to inhibit topoisomerases in cancer cells [81]. However, surprisingly for their antimicrobial potential, they have been reasonably less discovered. Therefore, a few questions related to the mechanistic roles of NQs in biological systems remain unanswered. Here, we list out some of the possible mechanisms responsible for antibacterial potential of NQs in biological systems. Literature suggests that among all NQ derivatives known, juglone has been most studied in depth for mechanistic perspectives, like the (1) ability to cure plasmids from bacteria, (2) interference with the activity of EPs, (3) generation of ROS, and (4) activity of the topoisomerase enzyme. Bioactive molecules from \textit{Plumbago} species have fascinated scientific fraternities due to their abilities to cure plasmids from bacteria. It is also remarkable to state that the molecular weights (ranging from 170 to 300 gm/mol) of natural sourced NQs like lawsone, plumbagin, juglone, shikonin, and menadione are less than the molecular weights of conventional antibiotics like penicillin, streptomycin, ciprofloxacin, and tetracycline (ranging from 330 to 582 gm/mol). This is a striking indication that these NQs are not bulky molecules and consequently can enter easily in the cell provided there is a favorable charge on the compound with a compatible chemical nature of side groups attached. NQs can perform their function through the above-mentioned mechanisms, which is explained in the following sections.

5.1. Efficiency to Cure Plasmids from Bacteria

Plasmids are well accepted as a “vehicle” to communicate genetic information among bacteria [82]. Plasmids are generally circular, extrachromosomal DNA elements where characteristic copy numbers are maintained in the host cell. Based on their functional prop-
erties, plasmids are classified as either (1) fertility (F-plasmids), (2) resistance (R-plasmids), or (3) virulence, (4) degradative, or (5) col plasmids. These plasmids encode properties like resistance to various antibiotics, virulence, the ability to degrade heavy metals or aromatic compounds, etc. Some plasmids bear the capacity to degrade hydrocarbons, synthesizing bacteriocins [83,84]. Resistance to antibiotics is spread through vertical (from parents to the next generation/offspring) and horizontal transfer (across genera or species). The spread of infection or antibiotic resistance can be restricted by inhibiting the conjugal transfer of plasmids. If a bacterial population loses plasmid/s, they may lose plasmid-borne characteristics. This process is known as curing. This process can be demonstrated in the laboratory with the help of curing agents like ethidium bromide (EtBr), acridine orange (AO), acriflavine (AF), and sodium dodecyl sulphate (SDS). None of the plasmid-curing agents can obviate all plasmids present in diverse hosts. Therefore, it becomes mandatory to investigate innovative molecules that can work as effective curing agents. Plasmid-curing compounds impose activity through (1) disrupting the replication of plasmids by integrating into the DNA, (2) producing breaks in the DNA, and (3) preventing conjugation or influencing plasmid supercoiling. These consequences lead to a reduction in the occurrence of plasmids within the population over time [85]. Replication of plasmids can be prohibited at various stages, and this concept has been well established through the “rolling circle” model [86]. Plasmid curing—the process of removal or obviating plasmids from bacteria—is depicted in Figure 3.

The literature suggests that NQs have abilities to cure plasmid/s from bacterial populations (Figure 3). The curing of plasmids from bacterial communities could be one of the conceivable mechanisms of action of NQs. Over the last 25 years, researchers have gained interest in NQs and their derivatives for plasmid-curing activities. Lakshmi and Thomas [87] reported plumbagin to cure F-like plasmid TP18 as well as attacking roles on DNA gyrase. Authors also revealed that the plasmid-curing abilities of plumbagin might be due to interference with the replication of miniTP18. Subsequently, there is a reduction in the copy number of plasmids along with an enhanced rate of segregation. The efficiency of plasmid curing is calculated from the number of colonies exhibiting the reversal of plasmid-borne character (for example, resistance to antibiotics) per 100 colonies tested (Figure 3). The efficiency of plasmid curing certainly varies and is dependent not only on the plasmid but also its typical host. Beg and Ahmad [88] reported 14% plasmid (pUK651 from E. coli x+)-curing efficiency in P. zeylanica (root) ethanolic lawsone extracts. Patwardhan et al. [34] showed lawsone could cure plasmids pBR322 and pRK2013 with curing efficiency of 11 and 20% respectively. Plasmid R136 seen in S. typhi was cured at 4.2% curing efficiency without any effect on plasmid RP4, found in E. coli. The plasmids having a molecular weight ranging from 4.3 to 56.4 Kb were cured fruitfully. Similarly, Patwardhan et al. [32] have also documented the effect of P. auriculata root extracts on nosocomial pathogens. Authors also proved the reversal of plasmid resistance to antibiotics in pathogens. P. auriculata root extract contains a variety of NQs, including plumbagin. Plasmids from P. aeruginosa (with 13%), E. coli (with 15%), P. vulgaris (with 32%), and K. pneumoniae (with 30%) were cured successfully. The curing efficiencies of root extracts stated here were significantly higher than AO (a reference curing agent). Therefore, the plasmid-curing abilities of NQs are noteworthy strategies if used in combination with antibiotics. The plasmid-curing abilities of a chemical will not kill pathogens directly; nevertheless, it will increase the susceptibility of pathogens towards antibiotics due to the loss of plasmid-borne resistance.

5.2. Inhibition of Efflux Pumps (EPs)

As a strategy of drug resistance, Gram-negative bacteria possess EPs that facilitate the movement of drug molecules outside the cell (generally in a nonspecific manner that confers resistance to bacteria against multiple antibiotics) [89,90]. EPs have the capability to extrude not only antibiotics but also other several organic pollutants and heavy metals.
The entire mechanism of action of EPs under the active and inhibited state (due to the action of NQs) is portrayed in Figure 4.

Figure 3. Plasmid curing—a possible mechanism of action for naphthoquinones. (A): Replication and multiplication of plasmid/s is affected at each stage of cell duplication. (B): Exposure of a bacterial culture to varied concentrations of curing agents, followed by the incubation, serial dilution, and plating of the culture on media with and without antibiotic. (C): Scoring of cured derivatives—by their failure to grow in presence of antibiotics. Calculation of plasmid curing efficiency—number of colonies displaying reversal of resistance to antibiotic/s per 100 colonies tested. Performance of agarose gel electrophoresis—to confirm the removal/obviation of plasmid DNA from the strain treated with a curing agent.
Figure 4. Interference with the activity of efflux pumps through the action of naphthoquinones. (A): efflux pumps under an active state conferring resistance against antibiotic molecules by pumping them outside the bacterial cell. (B): inhibition of efflux pumps by naphthoquinone molecule, resisting it from the efflux of antibiotic molecules and therefore, making the cell sensitive to antibiotics.
Plant-derived compounds, quorum-sensing signal molecules, and bacterial metabolites are reported to interfere with the activity of EPs. The functional status of EPs plays a decisive role in bacterial response to the surroundings. EP-mediated multidrug resistance is advantageous for bacterial communities in intrinsic, acquired, and phenotypic ways. Bacterial EPs are classified in the (1) resistance-nodulation-division (RND) family, (2) small multidrug resistance (SMR) family, (3) major facilitator superfamily (MFS), (4) multidrug and toxic compound extrusion (MATE) family, and (5) the adenosine triphosphate (ATP) binding cassette (ABC) superfamily. A single EP can pump out a wide range of antibiotics [91]. The inhibition of EPs guarantees the enhanced susceptibility of bacteria towards antimicrobial agents and also make them suitable targets.

Compounds like piperine [92], 5'-methoxy-hydnocarpin [93], quinoline derivatives [94] etc. have been reported to interfere with or affect EPs in several bacteria. An interesting fact about these chemicals is that they contain a ring structure that is a crucial part of NQs. Among those in the ESKEAPE and MDR category, *S. aureus* shows the presence of EPs. Zmantar et al. [67] reported the ability of juglone (at 182–256 µg/mL) to inhibit (50%) the EPs of *S. aureus* and the accumulation of chemicals in the cells. It is worth noting that juglone at the same concentration (182–256 µg/mL) inhibits the efflux of EtBr. Like juglone, plumbagin and shikonin also interfere with the functionalities of EPs. Ohene-Agyei et al. [95] provided evidence that plumbagin and shikonin can inhibit the EPs of *E. coli*. EPs are mainly utilized for the removal of the substrate from bacteria. It has been observed that AcrB-mediated substrate EPs become affected and inhibited in the presence of NQs. Antibiotics or other antimicrobial-agent-based therapy, in combination with NQ, result in the inhibition of EPs in bacteria. Table 4 represents the MICs of NQ derivatives when used individually and in combination with antimicrobial compounds inhibiting the activity of EPs. The MIC of antimicrobial compounds may be reduced several-fold when they are used in combination or in a synergistic way with other compounds like NQs while performing biological assays.

### Table 4. Naphthoquinone derivatives inhibiting the activity of efflux pumps in bacteria.

| Test Organism | MIC of Antimicrobial Compound | MIC of NQs/Substances (S)/Derivatives (Individual) | MIC of Antimicrobial Compound with NQs | Reference |
|---------------|-------------------------------|--------------------------------------------------|-------------------------------------|-----------|
|               | Name of the Antimicrobial Compound | MIC of AC (Individual) | Norfloxacin (NF) | MIC in µg/mL | With 1/8th MIC of S3, S4 and S5 |
| *Staphylococcus aureus* SA-1199 (wildtype -WT) | Ethidium bromide (EtBr) | ≥1024 (S3) | 32 (EtBr + S3) | S. aureus (WT) | 16 (EtBr + S3) |
| *Staphylococcus aureus* SA-1199B strain (expressing norA gene expressing NorA efflux protein) | EtBr | ≥1024 (S4) | 32 (EtBr + S4) | S. aureus (NorA) | 32 (EtBr + S3) |
|               | | ≥1024 (S5) | 64 (EtBr + S5) | | 128 (EtBr + S4) |
|               | | | 64 (EtBr + S5) | | 64 (EtBr + S5) |
| *Staphylococcus aureus* SA-1199B strain (expressing norA gene expressing NorA efflux protein) | Norfloxacin (NF) | ≥1024 (S3) | S. aureus (WT) | 1 (NF + S3) |
|               | | ≥1024 (S4) | 2.51 (NF + S4) | | 4 (NF + S5) |
|               | | ≥1024 (S5) | 80 (NF + S3) | S. aureus (NorA) | 50.8 (NF + S4) |
|               | | | 101.6 (NF + S5) | | |
| Test Organism | MIC of Antimicrobial Compound | MIC of NQs/Substances (S)/Derivatives (Individual) | MIC of Antimicrobial Compound with NQs | Reference |
|---------------|-------------------------------|-----------------------------------------------|----------------------------------------|-----------|
| -             | Erythromycin                  | Juglone                                      | With \( \frac{1}{2} \) MIC of Juglone  |           |
| Bacillus cereus | 128                          | 256                                          | 32                                     | [67]      |
| Escherichia coli | 64                           | 128                                          | 16                                     |           |
| Salmonella enterica | 4                          | 256                                          | 2                                      |           |
| Staphylococcus aureus | 128                      | 128                                          | 16                                     |           |
| Listeria monocytogenes | 64                        | 256                                          | 16                                     |           |
| Enterococcus faecalis | 64                         | 256                                          | 32                                     |           |
| Staphylococcus epidermidis | 32                     | 128                                          | 4                                      |           |
| Staphylococcus aureus | 16–256                      | 32–256                                       | 4–32                                   |           |
| (Oral strains = 8) | -                           | -                                            | (2 to 8-fold decrease in MIC)         |           |
| Tetracycline | Juglone | With \( \frac{1}{2} \) MIC of Juglone          |                                        |           |
| Bacillus cereus | 2                            | 256                                          | 1                                      |           |
| Vibrio alginolyticus | 256                        | 256                                          | 64                                     | [67]      |
| Salmonella enterica | 128                        | 256                                          | 64                                     |           |
| Staphylococcus aureus | 4                           | 128                                          | 2                                      |           |
| Enterococcus faecalis | 128                        | 256                                          | 64                                     |           |
| Staphylococcus epidermidis | 8                          | 128                                          | 2                                      |           |
| Pseudomonas aeruginosa | 64                         | 128                                          | 32                                     |           |
| Staphylococcus aureus | 2–32                        | 32–256                                       | 1–16                                   |           |
| (Oral strains = 8) | -                           | -                                            | (2 to 8-fold decrease in the MIC)     |           |
| Benzalkonium chloride | 16                          | 256                                          | 4                                      |           |
| Bacillus cereus | 16                            | 256                                          | 8                                      |           |
| Escherichia coli | 16                           | 256                                          | 4                                      |           |
| Salmonella enterica | 16                          | 256                                          | 4                                      |           |
| Staphylococcus aureus | 16                         | 128                                          | 8                                      |           |
| Listeria monocytogenes | 2                           | 256                                          | 1                                      |           |
| Enterococcus faecalis | 8                           | 256                                          | 2                                      |           |
| Staphylococcus epidermidis | 4                         | 128                                          | 2                                      |           |
| Pseudomonas aeruginosa | 32                         | 128                                          | 16                                     |           |
| Staphylococcus aureus | 4–16                        | 32–256                                       | 1–4                                   |           |
| (Oral strains = 8) | -                           | -                                            | (2 to 8-fold decrease in the MIC)     |           |
| MIC in mg/L     |                                |                                |                                        |           |
| Escherichia coli | Erythromycin                  | Plumbagin Plumbagin (64) + AC               |                                        | [95]      |
| Chloramphenicol | 256                          | 128 (Individual MIC)                         |                                        |           |
| Tetraphenylphosphonium | 4                         | 10 (Individual MIC)                          |                                        |           |
| Escherichia coli \(\Delta\)AcB (Drug sensitive) | Erythromycin | Plumbagin Plumbagin (10) + AC               |                                        |           |
| Chloramphenicol | 16                           | 10 (Individual MIC)                          |                                        |           |
| Tetraphenylphosphonium | 16                         | 16 (Individual MIC)                         |                                        |           |
| Escherichia coli | Tetraphenyphenolphosphonium   | >256                                         | 64                                     |           |
| Escherichia coli \(\Delta\)AcB (Drug sensitive) | Tetraphenyphenolphosphonium | >256                                         | 16                                     |           |
| Escherichia coli | Tetraphenyphenolphosphonium   | >256                                         | 64                                     |           |
| Escherichia coli | Tetraphenyphenolphosphonium   | >256                                         | 16                                     |           |
Table 4. Cont.

| Test Organism                  | Name of the Antimicrobial Compound | MIC of AC (Individual) | MIC of NQs/Substances (S)/Derivatives (Individual) | MIC of Antimicrobial Compound with NQs | Reference |
|-------------------------------|-----------------------------------|------------------------|----------------------------------------------------|---------------------------------------|-----------|
|                                | Plumbagin                        | 64                     | NDGA (256)                                         | Plumbagin NDGA (256) + AC             | [95]      |
| **Escherichia coli**           | Erythromycin                     | 256                    | 512                                                | 64                                    | [95]      |
|                               | Chloramphenicol                   | 4                      | 1                                                  |                                       |           |
|                               | Novobiocin                        | 512                    | 64                                                 |                                       |           |
|                               | Tetraphenyl-Phosphonium           | 1024                   | [96]                                               |                                       |           |
|                               | Tetracycline                      | 1                      | 0.25                                               |                                       |           |

Antimicrobial compound: AC; ethidium bromide: EtBr; norfloxacin: NF; plumbagin nordihydroguaretic acid: NDGA; substances/derivatives: S3, S4, S5.

5.3. Generation of Reactive Oxygen Species (ROS) in Bacteria

Reactive oxygen species like \( \text{H}_2\text{O}_2 \), \( \text{OH}^- \), and \( \text{O}_2^- \) are the accountable candidates for the emergence of lethal response in bacteria. The accumulation of \( \text{OH}^- \) radicals in an uncontrolled fashion may lead to bacterial cell death (due to the peroxidation of essential lipids in the cell). ROS have the abilities to target multiple biomolecules like proteins, DNA, and lipids (building blocks required for the survival of the cell) and exhibit severe impact on cell viability, leading to cell death [96]. This entire mechanism is conceptualized in Figure 5. Successive single-electron reductions are responsible for the production of ROS, though it is a subject of debate whether some researchers have proved the participation of ROS in the ‘antibiotic-mediated killing of bacteria’ [97].

The literature discusses the fact that compounds similar to benzoquinones are responsible for ROS generation in *Mycobacterium tuberculosis* [98]. Authors demonstrated that MIC values are highly affected due to the generation of extracellular \( \text{H}_2\text{O}_2 \) and therefore are recommended in designing an effective drug strategy. Thus, the generation of high levels of ROS in microbial cells could be a gifted strategy to treat bacterial infections [99]. Ravichandiran et al. [53] proved the action of NQ derivatives against *E. coli* through the generation of intracellular ROS. Authors also stated that redox behavior is a substantial factor and a possible mechanism of antibacterial activity of NQs. Additionally, the binding of NQs to bacterial membranes is a crucial one to demonstrate its action. Authors supported this fact through (1) the estimation of intracellular ROS, (2) the apoptosis-induced effects (3) bactericidal time–killing, (4) molecular modeling, and (5) an electrochemical study.

Researchers have utilized innovative technologies like isobaric tags for relative and absolute quantitation (iTRAQ) to evaluate the quantitative expression of proteins in cells treated with NQs. Isobaric tags (for assistance with relative and absolute quantitation) are popular as iTRAQ technology. This isobaric labeling technique uses quantitative proteomics through tandem mass spectrometry (MS) to quantify proteins present in varied sources in a single experiment set up. The technique essentially operates through the labeling (covalent interactions) of N-terminus with peptides (side-chain amines) after the digestion of protein with tags (4-plex, 8-plex) possessing speckled mass. Evaluation of data is performed at the peptide level where signals (of the reporter ions) for each MS spectrum are used to calculate the ratio of the relative abundance of the respective peptides identified by the spectrum [100,101]. More than one signal appearing in MS data might be due to the abundance of reporter ions. The collective ratios of protein/peptide positively represent the
relative quantification of a particular protein. Wang et al. [46] applied iTRAQ technology to the analysis of the expression of 53 proteins in *S. aureus* after treatment with juglone. Oxidative damage was the prime mechanism found to kill *S. aureus*. Juglone displayed the capability to upregulate the expression of oxidoreductase, resulting in an increase in the redox processes, thereby generating a peroxidative surrounding in the cell. Juglone treatment reduces cell-wall formation significantly and is also accompanied by an increase in cell permeability. Authors also suggested that juglone may impose different mechanisms of action against other bacteria. Another study documented by Linzner et al. [59] falls on parallel lines. Their study proposed the generation of ROS activity by NQs in bacterial cells. Authors also investigated the mode of action of lapachol in *S. aureus*, indicating the oxidative and quinone stress response. The generation of ROS by NQs is a noticeable antibacterial strategy through various targets like lipids, proteins, and DNA to enhance the susceptibility of bacteria towards ROS-generating compounds (Figure 5).

The literature discusses the fact that compounds similar to benzoquinones are responsible for ROS generation in *Mycobacterium tuberculosis* [98]. Authors demonstrated that MIC values are highly affected due to the generation of extracellular H2O2 and therefore are recommended in designing an effective drug strategy. Thus, the generation of high levels of ROS in microbial cells could be a gifted strategy to treat bacterial infections [99]. Ravichandiran et al. [53] proved the action of NQ derivatives against *E. coli* through the generation of intracellular ROS. Authors also stated that redox behavior is a substantial factor and a possible mechanism of antibacterial activity of NQs. Additionally, the binding of NQs to bacterial membranes is a crucial one to demonstrate its action. Authors

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**Figure 5.** Generation of reactive oxygen species (ROS)—a possible mechanism of action of naphthoquinones. Effect on (A): membrane lipids (B): cell DNA and (C): cell proteins.
5.4. Inhibition of Topoisomerase Enzyme

DNA topoisomerase enzymes are known to regulate and alter the topology of DNA in the cells. Based on their ability to make temporary single- or double-stranded breaks in DNA, topoisomerases are categorized as type I and II respectively [102]. The energy required for enzymatic-induced cleavage is achieved through the hydrolysis of ATP. DNA gyrase represents the bacterial type II topoisomerases, which are recognized for their "highly conserved" functional properties. The entire mechanism promotes DNA replication and transcription processes that are vital for cell survival. These enzymes regulate the topology of bacterial DNA through its cleavage as exemplified in Figure 6A. Malfunctioning of these events eventually results in cell death [103]. Bacterial DNA topoisomerase II (DNA gyrase) is an unquestionable target for antibacterial agents. NQ compounds inhibit the supercoiling activity of DNA. Karkare et al. [104] showed the inhibitory activity of gyrase enzyme in *S. aureus, E. coli*, and *M. tuberculosis* using commercially available diospyrin and 7-methyljuglone (Table 5). NQs bind to the N-terminal of GyrB, a novel location close to the ATPase site. These facts provide strong indication to depict a possible mode of action of NQs in organisms (Figure 6A), which can be explored further to develop an innovative antibacterial agent.

![Diagram](image)

**Figure 6.** Inhibition of topoisomerase enzyme—a possible mechanism of action of naphthoquinones. (A): interaction of topoisomerase enzyme in the bacterial system. (B): functional state of topoisomerase enzyme: a temporary single stranded break in DNA. (C): interference with the activity of the topoisomerase enzyme in the presence of naphthoquinone molecules.
Table 5. Naphthoquinone inhibiting supercoiling activity of enzyme—DNA gyrase (bacterial topoisomerase II).

| Test Organism       | IC50 (µM) of Naphthoquinone Derivative Inhibiting Supercoiling Activity of DNA Gyrase (Bacterial Topoisomerase II) Enzyme | Reference |
|---------------------|-----------------------------------------------------------------------------------------------------------------|-----------|
|                     | Diospyrin 7-Methyljuglone Menadione Shinanolone                                                                 |           |
| Staphylococcus aureus | 8 60 - -                                                                                                         | [104]     |
| Escherichia coli    | 4 30 - -                                                                                                          |           |
| Mycobacterium tuberculosis | 15 30 >200 >200                                                  |           |

The ability of NQs to constrain the biological activity of topoisomerase enzyme plays a decisive role at the stage of cell duplication in bacteria (Figure 6). This approach would result in reducing the load of bacteria and their infections drastically. Rather than a direct killing (bactericidal) effect, the inhibition of topoisomerase enzyme activity can result in bacteriostatic action to prevent the growth of bacteria.

6. Antifungal Potential of Naphthoquinones

Currently, though numerous effective antifungal agents are accessible in the market, their toxicity, development of resistance etc. have been emerging issues in confronting fungal infections. The need for a groundbreaking antifungal agent is always a priority [42]. In addition to antibacterial action, NQs also possess antifungal potential [23,24,42]. A study conducted by Carriço et al. [42] emphasized the potency of synthetic NQs against opportunistic and dermatophyte fungi. These fungi can grow on skin, nails, hair, feathers etc. where some of a body part surfaces and give rise to various diseases. Carriço et al. [42] assessed the antifungal activity of NQs against 89 fungal cultures. Among which, NQs, namely IVS320 (3a,10b-dihydro-1H-cyclopenta (b) naphtho (2,3-d) furan-5,10-dione)-dione), IVS322 (7,9a-dihydro-6bH-cyclopenta (b) naphtho (2,1-d) furan-5,6-dione), nor-α-lapachone (2,2-dimethyl-2,3-dihydrofuran-2,3-dihydrophtphtho (b) furan-4,9-dione) and nor-β-lapachone (2,2-dimethyl-2,3-dihydrofuran-2,3-dihydronaphtho (1,2-b) furan-4,5-dione), one NQ IVS320 (3a,10b-dihydro-1H-cyclopenta [b] naphtho (2,3-d) furan-5,10-dione)-dione) demonstrated the lowest MIC against dermatophytes (5–28 µg/mL) and Cryptococcus spp. (3–5 µg/mL). A preliminary study smoothed to understand the mechanistic action of IVS320 in altering cell-membrane permeability and not the fungal cell wall. Likewise, thiolated NQ derivatives can also exhibit antifungal activity at a lower MIC. Recently, Wellington et al. [24] analyzed thiolated NQ derivatives against C. albicans and revealed the utmost activity with MIC of 23.4 µg/mL. The authors further predicted that the removal of the meta-fluoro group from the same derivative can diminish its antifungal activity. Sánchez-Calvo et al. [19] demonstrated the antibacterial as well as antifungal activity of 1,4-NQs, 2-chloro-5,8-dihydroxy-1,4-NQ at the lowest MIC (2 µg/mL) against C. krusei. The antifungal properties of NQs along with antibacterial potential make them hopeful candidates for drug development.

The Boraginaceae family has been identified as one of the potential sources to isolate bioactive compounds having immense biological properties. Microbes associated with medicinal plants could also prove to be a potential resource to obtain bioactive molecules like NQs. This interesting concept has been presented by Mollaei et al. [105], where they used endophytic fungi associated with a medicinal plant (L. officinale of the Boraginaceae family) as a new source to extract shikonin. Such a study could prove an exceptional example to motivate the production of shikonin from endophytic fungi for industrial applications.

7. Antiparasitic Potential of Naphthoquinones

Along with antibacterial and antifungal activities, NQs are also recognized for antiparasitic activity [106,107]. Currently, parasitic diseases are a gigantic world-wide challenge for the medical sector [15]. Parasitic infections are receiving medical attention due to their severity. Salas et al. [21] documented that every year ~15,000 deaths occur due to the
infection caused by the parasite *Trypanosoma cruzi*. This clinical condition is known as Chagas disease. Naturally occurring NQs, viz. lapachol, β-lapachone, and its α-isomer have prevalent trypansomidal potential. For antiparasitic drug development, NQs seem to be a suitable class of chemicals [108]. These researchers have reported not only the synthesis but also the antiparasitic activity of semisynthetic NQs (similar to isolapachol). These semisynthetic NQs have comparatively better antimalarial potential than lapachol when they were tested against *Plasmodium falciparum*. Salas et al. [21] also described the molecular mechanism, commenting that the compound represses the formation of hemozoin crystals in parasites when they were treated with NQs. Thus, NQs inhibit the β-hematin polymerization process in *P. falciparum*. Bis-NQ derivatives also exhibit antiparasitic activities against *Toxoplasma gondii* and *T. brucei* [22]. The antiparasitic activities of NQs is another fascinating approach to explore NQs in the field of medicinal chemistry.

8. Future Prospects

The versatile nature of NQs (biological and chemical) has been explored as an emerging warrior in the battle against microbial pathogens. Even though several limitations and challenges are associated in the current area of research, we need to uplift these dynamic molecules as superlative opportunities to achieve our breakthrough. There are many more unexplored natural sources (plants, animals, microorganisms, etc.) that can be utilized to extract diverse types of NQs. The Boraginaceae family representing ~2000 sp. (trees, shrubs, herbs) are present ubiquitously and act as a potential source of bioactive compounds for medical applications. Microorganisms associated with medicinal plants could also prove to be a potential resource to extract NQs. NQ entities possess unique spatial configurations that enable substitution, addition, and modification/alteration. Even though various organic solvents are used to extract NQs from plant material, it is important to note that the selection of an appropriate solvent is crucial in order to enhance quality. Therefore, conscious efforts would facilitate the improvement of the overall yield of NQs from the chosen natural sources.

In the scientific community, 1,4-NQs are esteemed in comparison to 1,2-NQs and are therefore available as another entity to explore. We admire the tremendous work reported on the synthesis of several NQ derivatives along with their functional properties through MIC and IC₅₀ based assays. However, only sparse information discusses the mechanistic role of NQs in the biological system. Looking at the literature, we strongly feel that there are some weaknesses associated with the research being carried out on the analysis of functional activities of NQ derivatives against several pathogens. The main hindrance to perform those techniques is the requirement of highly sophisticated microscopic facilities. Subsequent improvement in this area would accelerate focused applications of the available therapeutic compounds. Information regarding the identification of the exact molecular target of NQs is deficient. From the ESKAPE and MDR groups, *Staphylococcus* spp. (Gram-positive) has been successfully inhibited in the presence of both natural and chemically modified NQs. Thorough knowledge of the mechanistic approaches of NQs against Gram-negative organisms belonging to the ESKAPE and MDR categories would be feasible. The multifunctional potentiality of NQs can be utilized solely, synergistically, or in combination. The combination of NQs with antibiotics and nanoparticles are hopeful strategies to combat pathogens. Extensive in vitro and in vivo analysis along with successful clinical trials are necessary. Like antimicrobial strategies, cancer therapy is one of the leading and challenging fields of medicinal biology. Several NQs display distinctive cytotoxic properties, as well as anticancer potential, as observed in different cell lines like adeno, breast, colon, lung carcinoma etc. The development of novel anticancer agents is the topmost priority for the healthcare system. Most cancers are not controllable using radiotherapy and surgical means. Therefore, chemotherapeutic agents involving pioneering small molecules are in the spotlight of cancer biology research. Interestingly, NQs are widely explored and the foremost focus for therapeutic purposes. Commercially, several quinone-based anticancer agents like bleomycins, dactinomycin, daunorubicin, doxorubicin, idarubicin,
mitoxantrone, and mitomycin-C are in use. Similarly, NQs impose effects on multiple singling pathways in tumor cells, through interfering in the cell-division cycle, DNA topoisomerases, and kinase signaling. Thus, there is a need for new-fangled molecules like NQs to treat cancer. The targeted drug synthesis can be achieved by considering the specific site of action of NQs in a biological system. This approach can be supplemented through valuable inputs using molecular docking analysis at the preliminary stage. The molecular docking perspective assuredly opens broad avenues to understanding the interactions of chemical entities at the molecular level. Computational modeling also offers a supportive role to predict the functionality of compounds in a complex system. These approaches would precisely lead us to develop effective or perhaps the most potent drugs for the medical sector.

9. Conclusions

Naphthoquinones, especially 1,4-NQs, stand as multifunctional molecules due to their extraordinary potential to inhibit bacteria, fungi, parasites, and cancer. It is also remarkable to note that the mechanism of action for each NQ molecule might differ. This could be possible chiefly because of the SAR and variation in the target sites. The addition or removal of a specific group from NQs largely affects its activity against pathogens. Curing of plasmids, production of ROS, inhibition of EPs, and topoisomerase enzymes remain the leading causes of NQs as antibacterial agents. Gram-positive pathogens are susceptible targets for NQs. Staphylococcus spp.—representative bacterium of the ESKAPE and MDR pathogen groups—has been frequently reported for naturally occurring (~20%) and chemically synthesized 1,4-NQs (~27%). Followed by Staphylococcus spp., Bacillus spp. (~16%) has been seen to be another susceptible target for naturally occurring 1,4-NQs. From Gram-negative bacterial genera, E. coli (~11%) and Pseudomonas spp. (~9%) have been identified as a vulnerable target. Additionally, researchers have used several other microbial pathogens to conduct laboratory experiments. The ability of NQs to cure plasmids and cause the inhibition of EPs could be a wise approach for combinatorial or synergistic therapies. Overall, the multifunctional potential of NQs and their derivatives present them as a future molecule of medicinal chemistry. Powerful evidence about the mechanistic approaches of emerging trends like NQs could be a hopeful milestone to defeat the battle against microbial pathogens.

Note: Chemical structures were designed using ChemDraw software (12, PerkinElmer, Waltham, MA, USA). Based on information available in the literature, all figures have been constructed using Paint 3D (6.2009.30067.0, Microsoft Lift London, Redmond, WA, USA) and BioRender (Professional Science Figure creator, BioRender, Toronto, ON, Canada) applications.

Author Contributions: S.K.S. and N.N.N., designed the concept. N.S.M., D.S., R.H.P. and S.A.B. collected essential literature and designed all figures. M.C. contributed towards chemical structure design using software. P.Z. guided the therapeutic approach of naphthoquinone derivatives. All authors have contributed towards executing those ideas and writing the manuscript. Finally, All authors have read and agreed to the published version of the manuscript.

Funding: S.K.S. express a deep sense of gratitude towards the Department of Science and Technology (DST), Govt. of India for financial support. (Ref: EMR/2016/007912). S.K.S. also expresses special thanks to Rashtriya Uchchatar Shiksha Abhiyan (Ref: RUSA-CBS-TH-3.2) for financial support. NNN acknowledges the financial support from Dnyandeo Yashwantrao Patil Vidyapeeth, Pune (Ref: DPU/755(8)/2017).

Institutional Review Board Statement: Not applicable.

Informed Consent Statement: Not applicable.

Data Availability Statement: This is a review article, and the entire data is presented within the article.

Acknowledgments: Not applicable.

Conflicts of Interest: The authors declare no conflict of interest.
Abbreviations
NQ, naphthoquinones; ROS, reactive oxygen species; WHO, world health organization; AMR, Antimicrobial resistance; MDR, multidrug-resistance; MIC, minimum inhibitory concentration; MBC, minimum bactericidal concentration; SAR, structure–activity relationship; EPs, efflux pumps; MRSA, methicillin-resistant Staphylococcus aureus; MSSA, methicillin-sensitive Staphylococcus aureus; DMSO, dimethyl sulfoxide; IPA, isopropyl alcohol; ACN, acetonitrile; DME, dimethyl formamide; DmsD: TaT proofreading chaperon D, iTRAQ, Isobaric tags for relative and absolute quantitation. tyrosyl-RNA synthetase (TyrRS) inhibitor; EtBr, Ethidium bromide; AO, acridine orange; AF, acriflavine; SDS, sodium dodecyl sulphate; RND, resistance- nodulation-division; SMR, small multidrug resistance; MFS, major facilitator superfamily; MATE, multidrug and toxic compound extrusion; ABC, adenosine triphosphate (ATP) binding cassette.

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