Review Article

The Role of Bacterial Biofilm in Antibiotic Resistance and Food Contamination

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1. Introduction

Food contamination by foodborne pathogens is a serious public health concern that can cause foodborne diseases [1]. Foodborne diseases are continuing to be a global public health problem with an estimated 600 million people falling ill annually [2, 3]. Food contamination may occur during any step in the farm-to-fork continuum from environmental, animal, or human sources and cause foodborne disease and intoxication [4]. Biofilm formation by foodborne pathogens is an inevitable event and becomes a source of food contamination. Bacterial biofilm formation is considered to be an emergent and prevailing microbial lifestyle in natural and manmade environments and occurs on all surface types [5, 6]. Biofilm is one of the most widespread and most successful life forms on Earth [7]. In nature, microorganisms commonly exist in the shelter of highly hydrated biofilms which creates a conducive environment for cells to adhere together and onto all kinds of surfaces [8]. Because microorganisms within this community produce cement-like matrix which can act as “biological superglue” [9], to fix or trap onto different biotic or abiotic surfaces. For instance, biofilm infections on implants or indwelling devices are difficult to eradicate because of their much better protection against macrophages and antibiotics, leading to severe clinical complications often with lethal outcome. It is a critical problem in the medical sector since it is formed on medical implants, within human tissue and involved in a multitude of serious chronic infections. Generally, biofilm is a surface-attached community of microorganisms encased within a matrix of extracellular polymeric substances that can act as a barrier and recalcitrant for different hostile conditions such as sanitizers, antibiotics, and other hygienic conditions. Generally, they persist and exist in food processing environments where they become a source of cross-contamination and foodborne diseases. The other critical issue with biofilm formation is their antibiotic resistance which makes medication difficult, and they use different physical, physiological, and gene-related factors to develop their resistance mechanisms. In order to mitigate their production and develop controlling methods, it is better to understand growth requirements and mechanisms. Therefore, the aim of this review article is to provide an overview of the role of bacterial biofilms in antibiotic resistance and food contamination and emphasizes ways for controlling its production.
Pathogenic microorganisms can attach to food surfaces, grow on them, and form a biofilm that causes an increase in the food safety risk [11]. Poor sanitation of food-contact surfaces, equipment, and processing environments has been a contributing factor in foodborne disease outbreaks, especially those involving *Listeria monocytogenes* and *Salmonella* [5]. Insufficient and ineffective cleaning practices can cause food residues to remain in food processing and can facilitate bacterial attachment and biofilm formation [6]. These surfaces with adherent microbial communities are difficult to sanitize properly since cells within a biofilm are persistent or tolerant to hygienic conditions [12]. The production of biofilm and its persistence on different surfaces related to food, medical, and other sectors would be reservoirs for many pathogens that are infectious [13]. Diverse microorganisms are able to grow on food matrices and along with food industry infrastructures, and this growth may give rise to biofilms [14, 15]. Therefore, biofilms formed on these surfaces are the main cause of contamination of the final product. Once the biofilm is formed, then it will be hard to eradicate from these surfaces. This again could be a source of disease transmission and reduce shelf life and quality of foods [16, 17]. Furthermore, biofilm mode of growth induces microbial resistance to disinfection that can lead to substantial economic and health concerns [18]. For instance, a research done on *Listeria monocytogenes* indicates that its biocide resistance and ability to cooperate with other species forming heterogeneous communities allowed this bacterium to survive and struggle within the industrial areas [19].

Contaminated foods could be a serious problem for food quality, safety, public health, and economic impact [16]. For example, adherence to pathogens on the meat surface causes contamination of the meat, which leads to product collection from the market and causes huge economic loss at the industry and country level [2, 20, 21]. Food contaminations and foodborne diseases put their pressure on developing countries, especially in infants, children, and other susceptible communities and it also has burden on local and global markets [22]. Food contamination not only leads to economic crises but also food safety which is the primary criterion in our expanding market [23]. Therefore, illness and death from diseases caused by contaminated food are a continuing threat to public health and a major impediment to socioeconomic development worldwide [24]. Generally, the food sector is a sensitive issue that can provoke panic in the food industry if the food is contaminated.

The emergence of antimicrobial resistance is a rapidly increasing challenge in public health worldwide [25]. Biofilm-forming bacteria are embedded in a matrix and acquire properties that render them highly tolerant to antibiotics, UV light, chemical biocides, host immune response, and other external stresses [26–30]. Biofilm can protect microorganisms from harsh environmental conditions such as extreme temperature and pH, high salinity and pressure, poor nutrients, antibiotics, etc., by acting as a barrier [31]. Structural barriers, along with persistent cells within biofilm, play a decisive role in antibiotic resistance [32]. As reports indicate, biofilm-related infections are difficult for medication and will not be cured easily [33]. Consequently, the prescription of antibiotics will not solve or remove biofilm-related infection due to their antibiotic tolerance and genetic mutation [34]. Biofilm is now considered to be a primary cause of chronic infection, and antibiotic-resistant bacteria are prevalent in biofilm form [35]. Currently, it is believed that over 80% of chronic infectious diseases are caused by biofilm, and it is known that conventional antibiotic medications are inadequate at eradicating these biofilm-mediated infections [30]. As Brackman and Coenye [36] reported, antimicrobial therapy often fails to eradicate biofilm from the site of infection. Generally, antibiotic resistance has emerged at an alarming rate and becomes an escalating public health problem. This problem is amplified by biofilm formation which creates additional bacterial tolerance to antimicrobial agents [35].

The spreads of biofilm-related infections are an intractable problem in modern medicine. Biofilm formation is the main virulence factor for a wide range of microorganisms that cause chronic infections [37]. Bacterial biofilm represents a major health concern due to the high demand for implantable medical devices and the rising numbers of bacterial resistance [38]. Pathogenic microorganisms can produce biofilm on implanted devices [39]. Many bloodstream infections and urinary tract infections are associated with indwelling medical devices and arise from a bacterial biofilm that consists of bacteria embedded within an extracellular polysaccharide matrix on the catheter surface [40, 41]. For example, *Staphylococcus aureus* and *Staphylococcus epidermidis* are considered two of the most important pathogens, and their biofilm frequently causes device-associated infections [42]. According to Otto, the biofilm phenotype that these bacteria adapt during device-associated infection facilitates increased resistance to antibiotics and host immune defenses [43]. Biofilm formation by microbial pathogens enables them to survive in hosts and causes chronic infections that result in persistent inflammation and tissue damage [30]. Therefore, biofilm formation on medical instruments, human tissues, and organs has an impact on human health and the economy.

**2. Stages of Biofilm Development**

Biofilm is an association of microorganisms that are firmly attached to the biotic or abiotic surface, encased within an extracellular polymeric substance (EPS) matrix, and that can show new character with respect to gene expression, protein synthesis, growth rate, and metabolic activities [44, 45]. Biofilm production can be influenced by a number of factors such as surface conditions, chemical and physical growth factors, cellular structures, and any other challenges. The interaction between these and other factors determines its fate [46]. As shown in Figures 1 and 2, structural and physiological change takes place after cells have been attached to conditioned surfaces. Structural polymeric substances produced are acting as a barrier [31] and prevent the entrance of antibiotics and sanitizer agents. Bacterial cell growth within biofilm is very slow and produces persistent
cells that can survive hostile conditions such as exposure to antibiotics and other biocides [6, 33] (Figure 2).

Microbial cells within a biofilm are very close to each other so that they can communicate through chemicals that enable them to coordinate and respond to any ecological, environmental, and host related cues [49]. According to Oliveira et al., biofilm formation is commonly viewed as a cooperative enterprise, where strains and species work together for a common goal [50]. For this cooperative activity, there must be cell-to-cell communication. This cell-to-cell communication mechanism within the microbial community is known as quorum sensing in which microorganisms use signaling such as acyl homoserine lactone (AHL) in Gram-negative bacteria, the autoinducing peptide (AIP) in

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**Figure 1**: Biofilm formation and structure, adapted from [46, 47] with major modification.

**Figure 2**: The mechanisms of antibiotic resistance in biofilms, source: [48] (with own modification).
Gram-positive bacteria, and the autoinducer-2 (AI-2) in both Gram-negative and -positive bacteria for a different purpose [36, 51]. Quorum sensing (QS) system is a mechanism by which bacteria regulate the gene expression profile according to the size of the microbial population, causing the formation of different forms of biofilm [7]. As a general quorum sensing is a process by which bacteria produce and detect signal molecules and thereby coordinate their behavior in a cell-density-dependent manner [36]. In addition to communication, these close contacts microbial communities enable them to exchange genetic material, and even the frequency of gene transfer is high when compared to their free form [52]. Therefore, horizontal microbial gene transfer and biofilm formation are interrelated [53]. For biofilm formation, microorganisms should transit from their free form into a sessile form which requires stepwise physiological and structural changes [47, 54]. Thus, these stepwise and dynamical process comprises (a) initial or reversible attachment on the conditioned surface, (b) irreversible attachment (c), microcolony or early development of biofilm structure, (d) maturation of biofilm which forms mushroom or tower-like structure, and (e) dispersion or detachment in which cells slough off from the matrix and return to their original free form [47, 55] (Figure 1). Therefore, the aim of this review article is to provide an overview of the role of bacterial biofilm in antibiotic resistance and food contamination.

2.1. Initial or Reversible Attachment. Bacterial surface attachment represents a turning point from planktonic life to the biofilm mode [56]. Reversible attachment involves an interaction of planktonic microorganisms with a conditioned surface [57–59]. But the interaction is very weak which involves van der Waals, electrostatic forces and hydrophobic interactions. It has been reported that the attachment will be best on surfaces that are rough, hydrophobic, and coated with different organic substances [44]. Bacterial structures such as the fimbiae, pili and flagella give strength to the interaction between bacteria and the surface of attachment [60]. Generally, cell appendages involved in the reversible attachment and bacteria at this stage commit to the biofilm lifestyle or leave the surface and return to the planktonic lifestyle [56].

2.2. Irreversible Attachment. At this stage, loosely bound organisms consolidate the attachment process by producing extracellular polymeric substances that complex with surface materials and/or receptor-specific ligands located on pili, fimbiae, and fimbriate or both [57–59]. After microorganisms are attached on preconditioned and permissive surfaces, then the cell starts an irreversible adhesion and accumulates as multilayered cell clusters [61]. As recent studies revealed biofilm formation is commenced with a layer of polymeric substances (EPS) in which microbial cells are swarming on the surface with subsequent growth of the biofilm [62]. During this step, a number of physiological and structural changes have occurred, such as nonmotility of the attached cells [58].

2.3. Microcolony Formation. Microbial cells embedded within the extracellular matrix undergo coordinated community growth that leads to the formation of microcolonies. According to Dunne, microcolony formation results from simultaneous aggregation and growth of microorganisms and is accompanied by the production of EPS [57]. Microcolonies which are basic units of biofilm are compartmentalized by channels with different distinct microenvironments [29] (Figure 1). After cells are firmly attached to conductive surfaces, then numerous microorganisms will come up and secrete polymeric substances that can act as a “glue” to fix microorganisms on different surfaces. After these sequential events, microcolonies are produced.

2.4. Biofilm Maturation. If conditions are suitable for sufficient growth and differentiation, a biofilm may develop into spatially well-arranged, three-dimensional mature biofilm structures [61] such as mushroom or tower-like structures interspersed with fluid filled channels in which nutrients, oxygen, and essential substances can be diffused and circulate in each microenvironment [51] (Figure 1 and 2). The development of biofilm is a cooperative group behavior mediated by density-dependent chemical signals released by bacterial populations embedded in a self-produced extracellular matrix [63]. This signaling mechanism is known as quorum sensing which is used to communicate and orchestrate group behaviors, including virulence factor secretion and biofilm formation [64, 65]. Quorum sensing activates the maturation and disassembly of the biofilm in a coordinate manner [63]. Generally, cell-to-cell signaling plays a tremendous role in cell attachment and detachment from biofilm [66].

2.5. Biofilm Dispersal. Biofilm formation is a cyclical process in which bacterial cells are detached from the mature biofilm and enter into their previous mode of life, i.e., planktonic state. As shown in Figure 1, detached bacterial cells will seek new surfaces to attach and start up a new round of biofilm formation. In this step, microbial cells will decide based on the environmental cues whether they live together or “fall apart” [46]. From a food contamination point of view, this step is important to disseminate microorganisms into food products. Biofilm cells can be detached from actively growing cells or from the deprived environment, communication, or removal of aggregates. It has been reported that nutrient limitation forces microorganisms to seek new environments [29, 46].

3. Biofilm and Its Impact on Antibiotic Resistance

The emergence and spread of antimicrobial resistance among bacteria are the most important health problems worldwide [67–69]. Antibiotic resistance is one of the consequences of the bacterial biofilm communities which contribute to chronic infections [67]. Biofilm-forming Klebsiella pneumoniae is an important multidrug-resistant (MDR) pathogen affecting humans and a major source for hospital infections associated with high morbidity and
mortality due to limited treatment options [70]. It has been reported that biofilm formation is a means for a bacterium to resist hostile environmental influences such as antibiotics and antimicrobial agents [70–73]. As Verderosa et al. reported, biofilm is recalcitrant to antibiotic therapy and a major cause of persistent and recurrent infections by clinically important pathogens worldwide [74]. This is because the formation of biofilms and subsequent encasement of bacterial cells in a complex matrix can enhance resistance to antimicrobials and sterilizing agents making these organisms difficult to eradicate and control [75–77]. The extracellular polymeric substances (EPS) matrix protects bacteria from antibiotics, avoiding drug penetration at bactericidal concentrations [38] (Figures 1 and 2). Bacteria within a biofilm are several orders of magnitude more resistant to antibiotics, compared with planktonic bacteria [78]. For instance, biofilms can tolerate antimicrobial agents at concentrations of 10–1000 times that needed to inactivate genetically equivalent planktonic bacteria [79]. As shown in Figures 1 and 2, the nature of biofilm structure and other physiological changes such as slow growth rate assists them to be resistant to antimicrobial agents [66, 80] (Figures 1 and 2). As reported microorganisms in a biofilm are resistant due to the following suggested factors: (a) polymeric matrix that can restrict diffusion of antibiotics (b) interaction of antibiotics with a polymeric matrix which lowers their activity, (c) enzyme-mediated resistance such as β-lactamase [73], (d) changes in metabolic activity inside the biofilm (Figure 2), (e) genetic changes on target cells or hiding the target sites, (f) extrusion of antibiotics using efflux pumps [73], and (g) the presence of outer membrane structure, such as in Gram-negative bacteria [81]. These mechanisms are critical for antibiotic resistance and survival of biofilm bacteria [73, 82]. The antibiotic resistance used by bacteria in biofilm is distinct and different from natural or innate resistance mechanisms [48] (Figure 2). As similar findings revealed bacteria within biofilm develop different molecular strategies to protect their cells from hostile conditions such as the interaction of biofilm matrix with antibiotics that can retard or lower their activities, slow growth rates in which antibiotic will not be effective, genetic related resistance, and producing persistent cells which are tolerant to different antibiotics [38] (Figure 2). In biofilm-forming bacteria, there is a high rate of mutation that enables them to develop resistant mechanisms, and this, in turn, gives an opportunity for their genes to produce enzymes that inactivate the antibiotics or expel the antibiotics using efflux pumps [34, 83]. Bacteria within biofilm produce persisters cells that are metabolically inert and it is one of their mechanisms to escape from antibiotics and even they have the ability to survive in high concentration of antibiotics [84] (Figure 2). Biofilm plays a critical role in the spread of antibiotic resistance. Within the high dense bacterial population, efficient horizontal transfer of resistance and virulence genes takes place [85]. The number of microorganisms within the matrix is too dense so that there is close contact between different microorganisms which enable them to exchange resistant genes and finally, the whole community may acquire that resistant gene [68] (Figures 1 and 2). Therefore, genetic diversification of microorganisms in biofilm is largely responsible for shaping antibiotic resistance [7]. As studies have suggested that biofilm is important for the transfer of conjugative plasmids due to the high proximity of cells within this multicellular structure [86]. The resistance of biofilm to antibiotics depends on different factors such as physical, physiological, and gene-related factors [34]. Thus, this multifactorial nature of biofilm development and drug tolerance imposes great challenges for the use of conventional antimicrobials [37]. To sum up, bacterial biofilm is a key player in the development of antimicrobial resistance [38].

4. Biofilm and Its Impact on Food Contamination

Food contamination by pathogenic microorganisms has been a critical public health problem and a cause of huge economic losses worldwide [4]. Microbial biofilm contains both food spoiler and disease-causing bacteria and results in postprocessing contamination which lowers the quality and shelf life of products and could be a means for disease transmission [87–89]. For example, Escherichia coli O157: H7 attached to beef-contact surfaces found in beef fabrication facilities may serve as a source of cross-contamination [90]. Among many pathogens, Staphylococcus aureus and Pseudomonas aeruginosa are capable of constructing the biofilm on materials and equipment [91]. Friedlander et al. reported that biofilm-forming bacteria, which colonize the surfaces of equipment in the dairy industry, may adversely affect the safety and quality of the milk and its products [92]. Biofilm production by bacteria such as Listeria monocytogenes is supposed to be one of the ways that confer its increased resistance and persistence in the food chain [93]. The formation of biofilms on biotic and abiotic surfaces is a potential hazard, contributing to the constant circulation of pathogens in the conditions of food production and contamination of foods [94]. Pathogenic bacteria penetrate food production areas and may remain there in the form of a biofilm covering the surfaces of machines and equipment [95]. Therefore, biofilm formation by pathogenic bacteria leads to severe contamination problems in food, food processing, and other areas that directly affect human health and life [10, 96]. In a hygienic point of view, the attachment of pathogenic microorganisms to food-contact surfaces can lead to potential sanitation problems since it is persistent for long periods in hostile conditions and reservoir for contamination [16, 23, 96, 97]. In a research conducted on Cronobacter sakazakii, it has been reported that this bacterium is able to adhere to different surfaces such as silicon, latex, polycarbonate, stainless steel, glass, and polyvinyl chloride (PVC). Biofilm formation on stainless steel surfaces of food processing plants, leading to foodborne illness outbreaks, is enabled by the attachment and confinement of pathogens within microscale cavities of surface roughness (grooves, scratches) [98]. The attachment of microorganisms on the food preparation surface could enable microorganisms to form biofilm and become a source of contamination [87]. Generally, the growth of pathogenic bacteria such as Escherichia coli O157: H7 and Salmonella enterica can result
Biofilm-forming microorganisms present a serious problem in the medical sector. Biofilm-forming bacteria are encased in a matrix that enables them to exclude antibiotics and host immune response. In addition to having structural barriers, biofilm-forming bacteria can undergo physiological changes such as slow growth rate and producing persistent cells. In these occasions, antibiotics cannot inhibit, kill, or eradicate these slow-growing and persistent cells which are found inside the biofilm matrix. Therefore, chronic infections caused by biofilms are often difficult to treat effectively in part due to the recalcitrance of biofilms to antimicrobial therapy. In general, antimicrobial resistance along with
biofilm formation becomes an escalating and intractable problem in the health sector and food safety.

Data Availability

The data or information used to write this review article is available from the corresponding author upon request.

Conflicts of Interest

The author declares that there are no conflicts of interest in regard to the publication of this paper.

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