Raman and Surface-Enhanced Raman Scattering for Biofilm Characterization

Seda Keleştemur, Ertug Avci and Mustafa Çulha *
Department of Genetics and Bioengineering, Yeditepe University, Istanbul 34755, Turkey; demirseda85@gmail.com (S.K.); ertugavci@gmail.com (E.A.)
* Correspondence: mculha@yeditepe.edu.tr; Tel.: +90-216-578-1587
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Abstract: Biofilms are a communal way of living for microorganisms in which microorganism cells are surrounded by extracellular polymeric substances (EPS). Most microorganisms can live in biofilm form. Since microorganisms are everywhere, understanding biofilm structure and composition is crucial for making the world a better place to live, not only for humans but also for other living creatures. Raman spectroscopy is a nondestructive technique and provides fingerprint information about an analyte of interest. Surface-enhanced Raman spectroscopy is a form of this technique and provides enhanced scattering of the analyte that is in close vicinity of a nanostructured noble metal surface such as silver or gold. In this review, the applications of both techniques and their combination with other biofilm analysis techniques for characterization of composition and structure of biofilms are discussed.

Keywords: Raman spectroscopy; surface-enhanced Raman spectroscopy; bacteria; biofilm; nanoparticles

1. Introduction

A biofilm is an assembly of microbial cells and extracellular polymeric substances (EPS) that are formed from polysaccharides, lipids, proteins, genetic materials, and humic-like substances [1]. EPS provides microorganisms irreversible attachment onto a surface and protects them from environmental stress conditions [2]. Microcolonies bury in an EPS matrix in a biofilm composition, which is surrounded by water channels. Water channels provide an exchange of nutrients and metabolites between the microorganisms and the aqueous phase [1,3]. The composition of the EPS also changes depending on the nature of the microorganism. In Gram-negative bacteria, the EPS consists of neutral and polyanionic polysaccharides, while in Gram-positive bacteria the EPS formation is cationic. On the other hand, the environmental conditions such as pH, temperature, oxygen, and nitrogen levels, and also the surface properties, affect the composition of a biofilm [3,4].

Identification of biofilm composition is very important for clarifying the interactions between the environment and microbial growth and also for understanding the molecular dynamics of the biofilm’s own chaotic system. Biofilm analysis can be performed with microscopic, spectroscopic, and molecular techniques [5]. Improvements in analytical techniques have greatly contributed towards elucidating the structure of biofilms.

Microscopy was one of the pioneering techniques used to analyze biofilm structure. Confocal laser scanning microscopy (CLSM) provides an opportunity to monitor biofilm structure and dynamics during the accumulation stage [6]. Characterization of the three-dimensional architecture of biofilm composition is possible by using specific fluorescent dyes [7]. The combination of microscopy techniques allows detailed information to be obtained, as demonstrated in a study by Strathmann et al. [8] who combined epifluorescence microscopy with CLSM and monitored carbohydrate distribution in an extracellular matrix. On the other hand, sample preparation in these techniques is extensive. Staining
brings additional cost and due to staining, simultaneous analysis of multiple components in biofilms is not an easy task. The assembly of a biofilm structure on a solid surface can be examined with scanning electron microscopy (SEM) and transmission electron microscopy (TEM) after a fixation procedure. Although the distribution of cells and the polymeric matrix can be excellently monitored, these microscopy techniques are not able to monitor living biofilm structures [9]. The other important microscopy technique is scanning transmission X-ray microscopy (STXM), which is extensively used in mapping biofilm composition without using a probe and also with reduced radiation damage on the sample. STXM is used to monitor not only the polymeric substances but also allows the metal distribution to be viewed. On the other hand, errors in reference spectra and absorption saturation limit the use of this technique [10]. Atomic force microscopy (AFM) is the most general technique used to gain information from the depth of the surface topography, and this feature allows the distribution of the extracellular polymeric substances to be displayed [11]. However, it is time consuming to use when characterizing larger areas of a heterogenic biofilm structure due to its low scanning speed.

Nuclear magnetic resonance (NMR) is another technique used to obtain structural information on the substances in biofilm matrices [12]. Although the technique gives an opportunity to acquire information in situ from a biofilm, it is not suitable for characterizing high molecular weight molecules due to the complexity of interpreting the obtained data [13]. Separating EPS from the biofilm matrix is another preferred method in characterizing the components. Extraction by centrifugation or using chemical additives, solid phase extraction with liquid chromatography (LC) and field-flow fractionation are new approaches and the main techniques used in several studies [14]. The identification of biofilm-associated genes of a microorganism can be carried out using molecular techniques, as well as analytical techniques. A combination of the widely used molecular techniques, gel electrophoresis, and polymerase chain reaction (PCR) with microscopic and spectroscopic techniques has revealed gene regulation profiles during biofilm formation [15]. Understanding polysaccharide, protein, and lipid profiles in biofilms was also achieved by separating the matrix with a chromatography column and then identifying the profiles with sensitive and high technology mass spectrometry techniques such as matrix-assisted laser desorption ionization mass spectrometry (MALDI-MS), electrospray ionization mass spectrometry (ESI-MS) or nano-ESI-MS, ESI-tandem MS, ESI-MS-MS, and LC-MS-MS [16,17]. However, chromatography and MS-based techniques are destructive, expensive, and time consuming. In addition, complex sample preparation steps are needed.

In addition to the techniques mentioned above, the vibrational spectroscopic techniques have been used to obtain information about the biofilm structure at a molecular level. Infrared and Raman spectroscopies are vibrational spectroscopy techniques used complementary to each other to acquire spectral information on the molecular structure of biofilms. Although infrared spectroscopy (IR) is widely used for characterizing biofilms, interference from water has limited its use in the composition of hydrated biofilms [18]. The spectral -OH band attributed to water suppresses the other characteristic bands of the components in a biofilm. Raman spectroscopy is based on Raman scattering, which occurs when photons scatter from the molecule with either lower or higher frequency than the frequency of the incident light [19]. Since the frequencies of the scattered photons depend on the types of bonds in the molecule, the Raman spectrum of each molecule is unique and can be considered as a fingerprint of that molecule. Diffraction-limited spatial resolution of Raman spectroscopy is around 1µm, which enables detailed analysis of complex structures containing various kinds of molecules and molecular groups such as proteins, carbohydrates, lipids, and nucleic acids. In addition to this remarkable feature, it is a non-destructive technique; therefore, sample preparation is easier than the other techniques. There is no need to stain samples and also no need to dry them because Raman scattering of water is very weak compared to other molecules (e.g., there is not any interference from water in Raman spectroscopy). We refer the readers to the other three reviews for detailed information about its use for characterizing biological structures [19–21]. Although Raman spectroscopy can be a useful tool for many research areas, weak Raman scattering hampers the use of this technique when a lower limit of detection is desired for a molecule of interest. In addition, acquisition of spectra from
biomolecules may take a longer time due to their poor Raman scattering nature and may require higher laser power ending up damaging the sample. In surface-enhanced Raman scattering (SERS), the Raman scattering from a molecule in the close vicinity of a nanostructured noble metal surface such as silver or gold is easily enhanced up to $10^7$ times [22,23]. The enhancement mechanism is attributed to the combined effect of both an electromagnetic (through localized surface plasmon resonance) and a chemical (through charge transfer upon contact of a molecule to nanostructured surface) enhancements. As a result, the sensitivity in SERS is significantly increased and the detection limit is lowered. For more information on the biological applications of SERS other than in biofilms, we refer the readers to review articles published recently [24,25]. Although the enhancement in scattering makes SERS suitable for many applications in several fields, there are still some challenges preventing the implementation of this technique widely including clinical applications. First, the choice of the SERS substrate is the most important factor due to the issues with the enhancement factor and spectral reproducibility. Another issue is the selective enhancement of the spectral bands observed on a spectrum. Since noble metal nanoparticles used as SERS substrates have specific affinity for certain functional groups such as thiols and amines, obtaining spectral information from low-affinity molecular components of biofilms can be limited and molecular moieties close to the noble metal surface can dominate the spectrum. To overcome this problem, SERS substrates can be functionalized with certain chemical groups to provide an affinity for the target biomolecules. Another issue is the spectral variations in the spectra of the same molecules or molecular structures from laboratory to laboratory. There is no worldwide accepted and applied protocol for spectra acquisition from cells, tissues, microorganisms, etc., which also prevents the implementation of this technique in clinics. On the other hand, we are hopeful that with fruitful collaborations at least some of the issues with the application of SERS in clinics will be solved in the near future.

In this review, Raman spectroscopy and SERS-based studies on biofilms are discussed. The first part focuses on the significant developments in the use of Raman spectroscopy for characterizing biofilm structures. The second part includes applications of the SERS technique for biofilm characterization, detection of quorum sensing molecules, and the identification and discrimination of biofilms of a variety of microorganisms.

2. Raman Spectroscopy of Biofilms

The initial reports regarding the use of Raman spectroscopy in biofilm research focused on the investigations of diffusion and distribution processes of certain molecules in the biofilm structure. Nearly two decades earlier, Suci et al. [26] have studied the spatiotemporal distribution of an antibacterial and antifungal agent, chlorhexidine, in Candida albicans biofilms. Both attenuated total reflectance (ATR)-Fourier transform infrared (FTIR) spectroscopy and Raman spectroscopy results have shown that the diffusion coefficient for the transport of chlorhexidine through biofilms with 200 µm thickness was less than the diffusion coefficient for chlorhexidine in water. It was also shown that the chlorhexidine concentration was higher in the outer regions of the thick biofilms due to its adsorptive behavior. In another report, diffusion of polyethylene glycol and hydrogen-deuterium oxide (HOD) in Streptococcus mutans biofilms was monitored [27]. Hydrated biofilms were mapped in x, y, and z directions. HOD could diffuse everywhere in the biofilm. On the other hand, polyethylene glycol diffusion rate was diverse in different parts of the biofilm due to the heterogeneous structure of the biomass.

Raman spectroscopy has also been used for analyzing the composition and structure of bacterial colonies. Choo-Smith et al. [28] analyzed the colony structure of two strains of Staphylococcus aureus, two strains of Escherichia coli and a strain of Candida albicans by acquiring Raman spectra from various positions within the colonies after their growth for 6, 12, and 24 h. At the sixth hour, microcolonies were very homogenous, indicating that the 6-h culture is more suitable for classification purposes. On the other hand, they were heterogeneous at 12 and 24 h. Spectra obtained from different positions within single colonies revealed the presence of more than one layer in the colonies. Moreover, it was
found that RNA concentration was higher in the deeper levels and glycogen amounts were higher on the surface of the colonies.

To investigate the effects of the initial age of bacterial cells for biofilm development, *Enterococcus faecalis* was investigated as the bacterial model. It is a commensal bacterium living in the intestinal tract and can cause infections after root canal treatment [29]. Biofilm matrices were prepared using bacterial cells in the exponential, stationary, and starvation phases, and their Raman spectra were acquired. It was found that the biofilms formed with the inoculation of bacterial cells in the starvation phase included more protein but less nucleic acid than the biofilms originating from exponential and stationary phase cells.

Detailed analyses of biofilm compositions were performed by Sandt et al. [30]. They investigated the structure and composition of hydrated biofilms grown in a flow cell using, for the first time, confocal Raman microscopy. Raman mapping of *Pseudomonas aeruginosa* PAO1 biofilms were performed for 3 weeks, from the beginning of the initial attachments to the formation of EPS in the biofilm. During the initial phase of the biofilm formation, variations among the acquired spectra were minor. On the other hand, spectra became highly variable in time due to the enriched composition of the biofilm. In order to analyze the source of the peaks in the collected spectra, biofilms were washed with EDTA to remove unbound EPS, and sharp spectral features attributed to EPS were obtained in difference spectra. In the other report [31], they analyzed water content and biomass density of wild-type (WT) *Pseudomonas aeruginosa* PAO1, small colony variant (SCV) PAO1, and *Pseudoalteromonas* ssp. NCIMB 2021 biofilms. Intensity ratios of the O-H band of water at 3450 cm$^{-1}$ and C-H band at 2950 cm$^{-1}$ on the mapped biofilm surfaces were used to find water to biomass ratios of the biofilms (Figure 1). Protein (BSA) solutions were used as standards to compare relative hydration.

Figure 1. Colony in a pure small colony variant (SCV) PAO1 biofilm 21 days after inoculation, showing cavities filled with water and little biomass in the core of the colony. (a) Bright field image. Raman spectra (1084 cm$^{-1}$) were collected every 2.5 μm in a plane extending down from the line shown through the colony in a. (b) Map of CH$_3$ signal (3300–3500 cm$^{-1}$, biomass). (c) Map of OH signal (2905–2965 cm$^{-1}$, water). (d) Map of water to biomass ratios (W/BRs). The arrows indicate the presence of streamers. The intensity scale for b and c (right-hand side of the figures) goes through blue to red as the peak area/1000 increases. The intensity scale for d goes through blue to red as the W/BR increases. Reproduced with permission from Springer [31].
In this context, Samek et al. acquired Raman spectra of the biofilm-forming *Staphylococcus epidermidis* 1457 strain and its non-biofilm-forming mutant *Staphylococcus epidermidis* 1457-M10 strain after growth of each on Columbia blood agar culture plates for 24 h [32]. Although there were subtle differences between the spectra of the two strains, the principal component analysis (PCA) of the spectra clearly discriminated the two strains and showed the effects of the biofilm matrix ingredients on the spectra, especially that of polysaccharide inter-cellular adhesions (PIA). Beier et al. [33] discriminated two oral bacteria species, *Streptococcus sanguinis* and *Streptococcus mutans*, in their hydrated biofilm mixtures using a classification algorithm that was previously developed by using their dried biofilms.

Raman spectroscopy has also been used to investigate the formation of different EPS types during biofilm growth. EPS is divided into two groups: soluble EPS and bound EPS. The latter again includes two groups: tightly bound and loosely bound EPS. Chen et al. [34] investigated the characteristics of the functional groups of these four types of EPS obtained from biofilms grown on two different biomass carriers in reactors and found that protein and polysaccharide amounts of tightly bound EPS in one kind of carrier were higher than the other.

Ivelva et al. [35] made a systematic study by acquiring Raman spectra of various biofilm ingredients such as polysaccharides, alginates, proteins, and spectra from free *Escherichia coli* and *Raoultella planticola* cells and their encapsulated forms, and used these data to gain in-depth knowledge about the origins of Raman spectra of multispecies biofilms.

The comparison of Raman spectra of planktonic cells and their biofilm forms gives invaluable information about biofilm formation and development. Huang et al. [36] compared Raman spectra of planktonic cells of *Pseudomonas fluorescens* SBW25 and cells obtained from its biofilm form were grown in King’s B medium for 3 days. The principal component-discriminant function analysis was used to discriminate between the spectra of the two groups and showed that metabolic status and environmental conditions had significant effects in that discrimination. Sharma et al. [37] characterized planktonic and biofilm cells of *Cronobacter sakazakii*, which cause infant meningitis. Comparison of the spectra of the two different forms of this opportunistic bacteria showed that the intensity of the bands assigned to carbohydrates, lipids, carotenoids, DNA, and tyrosine were more prominent in biofilm spectra.

Kusić et al. [38] studied planktonic cells and biofilms of six *Legionella* strains, as well as *Pseudomonas aeruginosa*, *Klebsiella pneumoniae*, and *Escherichia coli* species. Comparison of the Raman spectra of *L. pneumophila* and *Pseudomonas aeruginosa* grown in tap water and artificial media revealed an increased lipid and decreased DNA content in bacterial cells as an adaptation to nutrient-limited environmental conditions. Biofilms of *Legionella* strains had higher lipid content whereas biofilms of *Pseudomonas aeruginosa*, *Klebsiella pneumoniae*, and *Escherichia coli* had higher polysaccharide content relative to their planktonic cellular states. Subsequent support vector machine (SVM) analyses on Raman spectra of biofilms and planktonic cells resulted in more than 90% accuracies in their classifications. In another related report [39] by the same group, bacterial cells of *Legionella* species were isolated from their biofilms using immunomagnetic separation and identified on a single cell level using multivariate analysis. Implemented techniques were not involved in any cultivation step. An overall accuracy of 98.6% was achieved for the discrimination of isolated *Legionella* cells from *Pseudomonas aeruginosa*, *Klebsiella pneumoniae*, and *Escherichia coli* cells. In addition, an accuracy of 86.3% was obtained for the classification of five different *Legionella* strains.

Using different instruments for an experiment always brings new dimensions and gives comprehensive information about the matter of investigation. In this context, when the power of Raman spectroscopy is combined with that of CLSM, a better understanding of biofilm composition and its structure could be obtained. A good example of this was demonstrated by Wagner et al. [40]. For 92-day-old biofilms, CLSM analysis could not detect any composition change of the EPS. On the other hand, Raman spectroscopy showed a change in the composition of polysaccharides to glycoproteins.

A detailed analysis reported by Andrews et al. [41] showed the power of combining CLSM and Raman spectroscopy for a deeper understanding of cell attachment and biofilm growth. A total of
11 bacterial strains including species of *Rhodococcus*, *Pseudomonas putida*, *Sphingomonas*, *Acidovorax*, and *Duganella* were used for the study. It was found that attachment behavior of bacterial cells on different surfaces and subsequent biofilm growth was influenced by several extracellular macromolecules such as lipids, proteins, and nucleic acids, and the effects of these molecules were genus specific.

Feng et al. [42] developed a microfluidic lab-on-a-chip device to study *Pseudomonas aeruginosa* biofilms using both Raman spectroscopy and CLSM. They measured biofilm thickness using CLSM at different time periods and characterized biofilm composition with Raman spectroscopy. Comparison of the acquired spectra at different growth stages revealed that the major constituents of the biofilm at a mid-stage of biofilm development were similar to an early stage. After 48 h, the biofilm was in the late stage and production of carbohydrates and proteins was much more than nucleic acids. Further validation was achieved by discrimination of the Raman spectra of each stage using PCA.

CLSM was not the only technique used in combination with Raman spectroscopy to elucidate the composition and the structure of biofilms. Recently MALDI-MS was used with Raman spectroscopy and it was found that although nucleic acid bands were prominent for planktonic cells, glycolipids, rhamnolipids, and carbohydrate bands were more dominant in the case of biofilms [43]. MALDI-MS was used to obtain chemical information about rhamnolipid species; nine different species were found. In another study, the same group showed the presence of quinolone signaling molecules in *Pseudomonas aeruginosa* biofilms using Raman spectroscopy for the first time and analyzed the distribution of quinolone species with secondary ion mass spectrometry (SIMS) [44].

Since many studies have reported on the application of Raman spectroscopy for biofilm structure and composition, it was very unlikely that the effects of antimicrobial agents on biofilms using the same technique had not been investigated. For example, Lu et al. [45] showed antimicrobial effects of diallyl sulfide on the biofilms of *Campylobacter jejuni*, a bacterium that can cause gastrointestinal diseases. The chemical compositions of planktonic cells and biofilms were compared and EPS recalcitrance modes against antimicrobial agents (ciprofloxacin, erythromycin, and diallyl sulfide) were demonstrated. It was found that diallyl sulfide was effective on biofilms at a concentration 100-fold less than the other two antimicrobial agents.

The effects of antibiotics on *Pseudomonas aeruginosa* biofilms have also been shown [46]. Biofilms grown in a microfluidic device were monitored with CLSM and their composition was characterized by Raman spectroscopy in the presence of three antibiotics (ceftazidime, patulin, and epigallocatechin gallate; EGCG). Upon exposure to antibiotics, an intensity decrease in the protein and nucleic acid bands were observed. Moreover, in ceftazidime and patulin-treated biofilms, carotenoid accumulation was detected. In another study, the antimicrobial activity of papain functionalized polycaprolactam as a food packing material was tested against *Escherichia coli* biofilms [47]. Intensity changes of many Raman bands attributed to functional groups showed inhibition of biofilm formation.

Since Raman scattering of many biomolecules is very weak, implementation of resonance Raman spectroscopy (RRS) provided another dimension for in-depth analysis [48]. Resonance Raman scattering occurs when the excitation wavelength of the laser couples with the wavelength at which the targeted molecule absorbs the light at maximum level, and is at least 100 times more intense than Raman scattering. Therefore, use of a suitable laser wavelength makes it a lot easier to find certain molecules in biofilms. One of these molecules is cytochrome c. Pätzold et al. [49] used resonance enhancement of cytochrome c to find the distribution of nitrifiers and Annamox bacteria in their natural environment. In addition, bacteria could be grouped at the strain level using the heterogeneity of spectrum of cytochrome c in different exposure times and culture conditions. In this context, Kniggendorf et al. [50] developed coexisting microcolonies of *Nitrosomonas* on granules from two sequencing batch reactors and mapped with resonance Raman spectroscopy. The spectra obtained were compared with reference spectra of bacteria. As a result, *Nitrosomonas communis* Nm02 was identified with 94% certainty and *Nitrosomonas europaea* with 80% certainty.

Thus far, the application of Raman spectroscopy and its special form resonance Raman spectroscopy were reviewed. Both techniques provide fingerprint molecular information on the
analyte of interest and hence are extremely useful for bacterial biofilm studies. On the other hand, weak Raman scattering for biomolecules and the limited number of molecules that can be used for resonance studies limit the use of both techniques. Surface-enhanced Raman scattering (SERS), on the other hand, provides more enhancement and lowers the detection limits of the molecule of interest. In the next section, applications of SERS for biofilm analyses are provided.

3. Surface-Enhanced Raman Scattering of Biofilms

3.1. Identification of Spectral Signatures in a Biofilm

The application of SERS in biofilm characterization has become promising with the studies published for the last 10 years. Clarification of the biochemical origins of the SERS signals in a biofilm and determining the spectral signatures and biomarkers specific to the microorganisms could help widen the use of SERS in different clinical settings. The first study showing the applicability of SERS in biofilm characterization was published by Ivleva et al. [51] in 2008. In the study of Ivleva et al., protozoa cells and their biofilm structure were identified with SERS. The biofilm was grown in a reactor and was cultivated on a glass slide. Hydroxylamine hydrochloride-reduced silver nanoparticles (AgNPs) were used as SERS substrates and the biofilm-coated glass slides were immersed into the silver colloidal suspension and the SERS measurements were performed. The peaks at 1382 cm\(^{-1}\) and 1585 cm\(^{-1}\) were identified in the SERS spectrum of protozoa cells belonging to COO\(^-\) symmetric and asymmetric stretching vibrations of amino acids and polysaccharides. Apart from the cell spectra, the bands at 734 cm\(^{-1}\) and 1355 cm\(^{-1}\) attributed to polysaccharides were observed in the SERS spectra of protozoa biofilms. It was defined that these two dominant peaks could be used as a polysaccharide marker in a biofilm structure. Characterization of a heterotrophic biofilm structure was also demonstrated by Ivleva et al. [52] under the same conditions as the previous study mentioned above. Dominant peaks on the biofilm spectra were consistent with their previous study [51]. The dominant peaks were identified at 1555, 1380, and 1280 cm\(^{-1}\) with maximum intensities. The peak at 1380 cm\(^{-1}\) belonging to polyanionic polysaccharides distributed in the biofilm matrix and this specific peak was assigned as a marker for polyanionic polysaccharide in the biofilm matrix. Also, the peak at 1280 cm\(^{-1}\) was attributed to COH, HCO, and HCC deformation, COO\(^-\) stretching symmetric, and C-O stretching. In addition, the phenylalanine ring breathing peak observed at 1000 cm\(^{-1}\) could be used as a protein signature in a biofilm.

Biofilm formation consists of adhesion, colony formation, and maturation phases [53]. Monitoring the structural changes in biofilms at different time points of their growth with SERS can provide in-depth understanding of biofilm formation processes. In this context, Chao et al. [54] studied the formation Escherichia coli, Pseudomonas putida, and Bacillus subtilis biofilms between 0 and 72 h of growth with SERS using hydroxylamine hydrochloride-reduced AgNPs. Biofilms were grown on quartz windows and then 200 µL AgNPs was added on the substrates, which were incubated until dry before SERS measurements. The determined main peaks on the SERS spectra were 479, 730, 980, 1320, and 1145 cm\(^{-1}\) belonging to carbohydrates, nucleic acids, proteins, and lipids. Indeed, the biofilms formed by Gram-negative bacteria (Escherichia coli and Pseudomonas putida) and Gram-positive bacteria (Bacillus subtilis) could be discriminated based on the peaks observed at 659, 1112, and 1366 cm\(^{-1}\), which were attributed to tyrosine, lipids, and proteins. The main goal of the study was to monitor changes in the SERS spectral pattern during biofilm formation. The deterministic peaks of Escherichia coli were observed at 659, 730, 1320, and 1445 cm\(^{-1}\) with pattern differences from the previous studies [55–59]. The differences were explained in that study by changes in the bacterial strain, growth conditions, instrumentation, and SERS substrates. In the study of Chao et al., the regulation of biofilm processes was also evaluated based on the expression profiles of carbohydrates, proteins, DNA content, and lipids. The key peaks of the polysaccharides were defined as the peaks observed at 409, 479, and 565 cm\(^{-1}\) with an increase in the intensity profile during maturation. Protein enrichment was observed after 72 h incubation with an intensity increase in the peaks of tyrosine, phenylalanine, amide I, amide II, and amide III at 659, 1005, 1616, 1571, and 1242 cm\(^{-1}\). Also, extracellular DNA
was found that Gram-negative bacteria expressed high amounts of lipid during maturation. Polysaccharide-rich matrix, protein is the major component in the bacterial biofilms [60].

2018 distinctive spectral feature for monitoring bacterial death. RNA components being released after disintegration of the bacterial cell wall, which can be used as a distinctive spectral feature for monitoring bacterial death. In particular, the peak observed at 678 cm$^{-1}$ could be attributed to DNA/RNA components because these peaks were significantly observed on the spectrum of pristine oligonucleotides. Indeed, it was demonstrated that the general spectral patterns in a two-microorganism model were significantly different during biofilm formation. Additionally, microorganism death was observed with SERS in another study published by our group [61]. *Escherichia coli* was selected as the model microorganism and was exposed to ampicillin for 2, 4, and 6 h. Changes in molecular activity of the bacteria were clearly observed with increasing exposure time to the antibiotic, as seen in Figure 2. In particular, the peak observed at 678 cm$^{-1}$, which shows a significant change in the spectrum after 6 h, is thought to be attributed to DNA and RNA components being released after disintegration of the bacterial cell wall, which can be used as a distinctive spectral feature for monitoring bacterial death.

**Figure 2.** Surface-enhanced Raman scattering (SERS) spectra obtained during the death of *E. coli* with ampicillin exposure. NA: Nutrient agar [61].
3.2. Detection of Quorum Sensing Molecules

Quorum sensing is a chemical communication system for microorganisms in a biofilm structure. The microorganisms produce chemical signals called autoinducers that diffuse across the cell membrane. When these signal molecules reach a threshold concentration, they bind to receptors on bacteria, which leads to gene regulation [62]. Quorum sensing is essential for the production of virulence factors and for growth of the biofilm. These molecules could be used as a biofilm reagent in SERS measurements. In this context, Pearman et al. [62] acquired SERS spectra of N-acyl homoserine lactones (AHLs), which are expressed by Gram-negative bacteria and possess acyl chain lengths from 4 to 12 carbons, using colloidal borohydride and citrate-reduced AgNPs. The peaks of amide I, amide II, amide III, and amide IV at 1648, 1570, 1308 and 771 cm$^{-1}$ were observed clearly in all of the AHLs' spectra; the most significant peak was determined as amide I. The observed spectral differences at around 1200 cm$^{-1}$ and at 1533 cm$^{-1}$ among the spectra of AHLs with different acyl chain lengths were found as useful for their identification and discrimination. The obtained spectra of autoinducers in this study can be useful for their detection in biofilm matrices in future work.

*Pseudomonas aeruginosa* is a pathogenic bacterium known to cause cystic fibrosis [63] and is generally used as a model microorganism in studies identifying virulence factors and quorum sensing molecules in biofilms. Pyocyanin is a blue-green toxic molecule that acts as a virulence factor. It is a member of the quorum sensing system of *Pseudomonas aeruginosa* and is used as a biomarker for this bacteria species. The first study about the detection of pyocyanin in biofluid samples with SERS was performed by Wu et al. [64] in 2014. Pyocyanin was detected in clinical sputum samples with 95.6% sensitivity and 93.3% specificity using a silver nanorod array as the SERS substrate. The achieved limit of detection for pyocyanin was 5 ppb, which shows the potential use of this technique in the clinic for the early diagnosis of *Pseudomonas aeruginosa* infections. The other outcome of this study was that the characteristic SERS spectral peaks of pyocyanin at 1088, 1125, 1352, and 1613 cm$^{-1}$ could be extremely useful for the identification of *Pseudomonas aeruginosa* biofilms using SERS. In this context, novel plasmonic substrates were fabricated for label-free detection of pyocyanin with surface-enhanced resonance Raman scattering (SERRS) in a recent study (Figure 3) [65]. The study showed the efficiency of three differently designed plasmonic substrates as sensors for pyocyanin detection. Macroporous poly-N-isopropylacrylamide hydrogels loaded with Au nanorods and a mesostructured substrate with a mesoporous TiO$_2$ thin film over a submonolayer of Au nanospheres were produced and their efficient use for the characterization of a biofilm at early and mature phases was demonstrated. Mesoporous silica-coated micropatterned supercrystal arrays of Au nanorods were fabricated as a third sensor, which lowered the detection limit of pyocyanin due to its better SERS enhancing property. SERS detection of pyocyanin was performed for both planktonic cultures and biofilms grown on these plasmonic substrates. In addition, the designed sensors were implanted in mice to demonstrate their potential use for detecting small biomolecules in chaotic biofilm structures and for the screening of antimicrobial drugs. In a study by Polisetti et al. [66], pyocyanin was secreted from both clinic and laboratory strains under different nutritional support and was revealed—using SERS imaging and principal component analysis—as a key molecule in the *Pseudomonas aeruginosa* biofilm. Sodium borohydride-reduced AgNPs were used as SERS substrates and biofilm samples were grown in sterile tubes. Biofilm samples with AgNPs dropped on silicon wafers were incubated until dry before characterization with SERS. In this study, C-C stretching, C-N stretching, and C-H in-plane bend modes were assigned as a marker peak of pyocyanin observed at 1354 cm$^{-1}$ from the spectral data and SERS was proposed as a high-potential technique for chemometric analysis.
were observed during biofilm maturation. The SERS spectra of the *Pseudomonas aeruginosa* biofilms collected at these time periods. The SERS peaks that were useful for the discrimination of the biofilms composed of both *Pseudomonas aeruginosa* and *Staphylococcus aureus* were adenine peaks at 733 and 1340 cm$^{-1}$, amide I band at 1704 cm$^{-1}$, C-H bending of proteins at 1028 cm$^{-1}$, a lipid peak at 1460 cm$^{-1}$, and the peak at 1330 cm$^{-1}$ attributed to DNA bases. PCA of the spectra obtained from planktonic and biofilm forms of the microorganisms were used to find dominant species in dual-species biofilms at different time periods.

In our recent publication, we focused on monitoring biofilm formation of clinically relevant microorganisms [69]. Citrate-reduced AgNPs were used as SERS substrates and the biofilm formations of *Pseudomonas aeruginosa*, *Staphylococcus epidermidis*, and *Candida albicans* were evaluated on agar plates at different time points between 4 and 120 h of biofilm growth. Citrate-reduced AgNPs were used in this study. A colloidal suspension (10 μL) was dropped on the biofilm samples and then incubated for 10 min for interaction of the AgNPs with the biofilm components. The objectives of the study were to define the biofilm formation steps of the model microorganisms at a molecular level and to show the efficiency of the SERS technique for classifying the different biofilms based on the unique metabolic processes of microorganisms using PCA. Significant changes in the SERS spectra of biofilms were observed during biofilm maturation. The SERS spectra of the *Pseudomonas aeruginosa* biofilm at different time periods are given as an example in Figure 4. For the biofilm at different time points between 4 and 120 h of biofilm growth. Citrate-reduced AgNPs were used as SERS substrates and the biofilm formations of *Pseudomonas aeruginosa*, *Staphylococcus epidermidis*, and *Candida albicans* were evaluated on agar plates at different time points between 4 and 120 h of biofilm growth. Citrate-reduced AgNPs were used in this study. A colloidal suspension (10 μL) was dropped on the biofilm samples and then incubated for 10 min for interaction of the AgNPs with the biofilm components. The objectives of the study were to define the biofilm formation steps of the model microorganisms at a molecular level and to show the efficiency of the SERS technique for classifying the different biofilms based on the unique metabolic processes of microorganisms using PCA. Significant changes in the SERS spectra of biofilms were observed during biofilm maturation. The SERS spectra of the *Pseudomonas aeruginosa* biofilm at different time periods are given as an example in Figure 4.
different time points are given as an example in Figure 4. For the Candida albicans biofilm, a significant protein, lipid, and carbohydrate enrichment was observed at the incubation times of 24 and 48 h. These molecular composition changes were effective for discriminating yeast biofilm spectra from bacterial biofilm spectra. The findings of this study can be helpful for the potential use of SERS for identifying discriminatory biofilms of pathogenic microorganisms in clinical settings. A summary list of the SERS applications in biofilm detection is provided in Table 1.

![Figure 4. SERS spectra of biofilm formation of Pseudomonas aeruginosa at cultivation times between 4 and 120 h [69].](image)

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<thead>
<tr>
<th>Type of Microorganism</th>
<th>Output</th>
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<tr>
<td>Protozoa cells</td>
<td>Characteristic peaks of polysaccharides (1382, 1585 cm⁻¹) [51]</td>
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<tr>
<td>Heterotrophic biofilm</td>
<td>Polyanioc polysaccharides (1380 cm⁻¹), phenylalanine (1000 cm⁻¹) [52]</td>
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<td>Escherichia coli</td>
<td>Microorganisms were discriminated based on the peaks of tyrosine (659 cm⁻¹),</td>
</tr>
<tr>
<td>Pseudomonas putida</td>
<td>lipids (1112 cm⁻¹), proteins (1366 cm⁻¹),</td>
</tr>
<tr>
<td>Bacillus subtilis</td>
<td>Identification of the changes in the spectral pattern during biofilm formation [54]</td>
</tr>
<tr>
<td>Algae and</td>
<td>Algae biofilms contain a polysaccharide-rich matrix</td>
</tr>
<tr>
<td>Pseudomonas aeruginosa</td>
<td>Bacteria biofilms contain a protein-rich matrix [60]</td>
</tr>
<tr>
<td>Escherichia coli</td>
<td>DNA/RNA components (678, 802, and 1570 cm⁻¹) [22]</td>
</tr>
<tr>
<td>Staphylococcus aureus</td>
<td>Distinctive spectral features for monitoring bacterial death (DNA/RNA components,</td>
</tr>
<tr>
<td></td>
<td>678 cm⁻¹) [61]</td>
</tr>
<tr>
<td>Pseudomonas aeruginosa</td>
<td>Detection of pyocyanin (1088, 1125, 1352 and 1613 cm⁻¹) [64]</td>
</tr>
<tr>
<td>Pseudomonas aeruginosa</td>
<td>Plasmonic substrates were fabricated and used in label-free detection of pyocyanin [65]</td>
</tr>
<tr>
<td>Pseudomonas aeruginosa</td>
<td>Discrimination peaks for the two-species model were assigned as a marker</td>
</tr>
<tr>
<td>Brevundimonas diminuta</td>
<td>peak of pyocyanin [66]</td>
</tr>
<tr>
<td>Staphylococcus aureus</td>
<td>Identification of biofilm formation steps of the model microorganisms</td>
</tr>
<tr>
<td>Candida albicans</td>
<td>Detection of pyocyanin (1355 cm⁻¹)</td>
</tr>
<tr>
<td></td>
<td>Discrimination of biofilms with using PCA [69]</td>
</tr>
</tbody>
</table>
4. Conclusions

Raman spectroscopy and SERS are promising techniques for identifying the chaotic composition of a biofilm at the molecular level. The discovery of spectral features specific to biofilm components and microorganisms could widen the use of the techniques in different areas such as in clinics, food and marine industries. Both techniques are also convenient to detect and characterize biofilms with their short analysis time and easy applicability relative to the other conventional techniques used in these fields. As the reports and the collaborations among the laboratories throughout the world increase, the creation of a comprehensive database for band assignments will provide new opportunities to obtain more reliable results. The combination of these techniques with other microscopic, spectroscopic, and molecular biofilm characterization techniques will also lead to a better understanding of the complex nature of biofilms.

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References


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