Scanning probe microscopy investigation of the bacteriophage effect on bacterial biofilms
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Background
Biofilms are surface-associated bacterial communities immersed in a hydrated matrix of extracellular polymeric substance (EPS), comprised of polysaccharides, proteins, nucleic acids, and lipids (Hall-Stoodley et al., 2004). Biofilms could grow on natural, industrial and hospital surfaces (Bridier et al., 2011), and are associated with up to 60% of all human infections (Spoering & Lewis, 2001). At the same time polysaccharide capsule of bacterial biofilms increases their resistance to antibiotics (Stewart, 1996). Due to their ability to digest the EPS capsule of the host bacterial cell and reduce biofilms (Chai et al., 2014), depolymerase containing bacteriophages are considered as promising biofilm disruptive agents (Yan et al., 2014). Though many aspects of depolymerase associated bacteriophage induced biofilm degradation have been investigated, the physical mechanisms of this process remain not fully understood. The purpose of this work was direct investigation of bacteriophage effects on biofilms at a nanoscale and microscale using high-resolution scanning probe microscopy methods such as atomic force microscopy (AFM) and scanning ion conductance microscopy (SICM). The medically relevant \textit{A.baumannii} cells and their specific bacteriophages were used in this study.

Experimental
Bacteriophage vB\textsubscript{AbaP}\textsubscript{APK2} (APK2) (A.V. Popova et al., 2020) and its bacterial host, \textit{A. baumannii} strain belonging to the capsular type K2 (Senchenkova et al., 2014) were used in this study. Biofilms were prepared by deposition of 50-1000 μl \textit{A. baumannii} cells suspension in stationary or exponential growth phase on a substrate surface (bare or modified glass, mica, highly oriented pyrolytic graphite, polycarbonate) for 10 minutes – 72 hours at 37°C or 21°C. Then the samples were rinsed with distilled water followed by drying (for imaging in air) or addition of the imaging buffer (for imaging in liquid). For AFM experiments we have used a multimode atomic force microscope with a Nanoscope V controller (Veeco, USA) operating in a tapping-mode using commercial cantilevers. AFM images were processed in Femtoscan software (Advanced Technologies Center, Russia). Biofilm topography measurements were performed in a Petri dish (35 mm) on scanning ion-conductive microscope (ICAPPIC Ltd, UK). Borosilicate nanocapillaries made of borosilicate blanks (Sutter Instruments, USA) with an outer diameter of 1.2 mm, an inner diameter of 0.69 mm and a length of 7.5 cm were used as a probe.

Results and Discussion
Optical images of \textit{A. baumannii} cells adsorbed on a glass surface demonstrate formation of biofilms already within 10 minutes after deposition (Figure 1b; the image is obtained in air). The surface area of biofilms ranges from 5 to 100 μm\textsuperscript{2}. Similar biofilms have been formed on mica, highly oriented pyrolytic graphite (HOPG) and polycarbonate surfaces. AFM images of \textit{A. baumannii} biofilms obtained in air reveal tightly packed individual cells with round and slightly elongated shape (Figure 1b), which is typical for \textit{A. baumannii} cells. The lateral size of individual cells varied from 0.5 to 1.2 μm depending on the growth conditions, whereas the height was considerably lower, in the range of 200 – 700 nm, depending on
preparation, indicative of cell shrinkage caused by sample drying. Typical SICM images of *A. baumannii* biofilm obtained in Hanks solution (Figure 1c) demonstrate the biofilm height of ~1 μm and multiple values, which correspond to one and several bacterial layers, respectively. Exposition of bacterial biofilms to bacteriophage containing solution leads to changes in the biofilm morphology and appearance of adsorbed bacteriophage particle on bacterial surface (Figure 2a). The apparent packaging of individual cells in a biofilm becomes sparser. Also under certain conditions biofilms become rich in small holes ~40–100 nm in diameter. These observations may be associated with digestion of EPS capsule by bacteriophages. Incubation of *A. baumannii* biofilms with bacteriophages for 25 minutes at 37°C leads to a complete destruction of some cells (e.g., top left cells in Figure 2b) that is connected with a cell lysis caused by bacteriophage (the lytic cycle of this bacteriophages is 20 minutes). However, the major fraction of the cells remains undistorted due to small multiplicity of infection and peculiarities of lytic cycle in the adsorbed bacterial cells (Abraham et al., 2020).

**Conclusions**

*A. baumannii* biofilms formed on different substrates have been characterized using AFM and SICM. Specific bacteriophages induce apparent changes of *A. baumannii* biofilms such as EPS capsule digestion, bacteriophage adsorption and cell lysis.

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**References**


