Analysis of the SDS-PAGE patterns of outer membrane proteins from Escherichia coli strains that have lost the ability to form K1 antigen and varied in the susceptibility to normal human serum

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Abstract We used SDS-polyacrylamide gel electrophoresis to investigate the outer membrane proteins (OMPs) band composition of 19 Escherichia coli K1 strains that have spontaneously lost the ability to form K1 polysaccharide capsule (E. coli K1−) and demonstrated different degrees of susceptibility to the bactericidal action of normal human serum. Presented results showed that there were differences between E. coli K1− strains in OMPs expressing capacity. The analysis performed on OMPs has not revealed a direct association between the different OMPs band composition and the susceptibility of these strains to the serum.

Abbreviations

C Complement
NHS Normal human serum
OMP Outer membrane protein
UPEC Uropathogenic Escherichia coli strains
UTIs Urinary tract infections

Uropathogenic Escherichia coli strains are the major cause of urinary tract infections (UTIs) and are responsible for approximately 70–95% of community acquired and 50% of nosocomial UTIs (Kucheria et al. 2005). Mostly, these infections arise in the bladder (cystitis); however, in cases where the kidneys (pyelonephritis) are involved, the course of the disease is usually more severe. Frequent causative agent of childhood pyelonephritis is K1 capsular polysaccharide expressing E. coli rod (Johnson 1991). Moreover, E. coli K1 strains are a particularly important cause of neonatal meningitis (more than 80% of the cases) and are commonly associated with neonatal septicaemia. Typically, these pathogens are capable of invading the brain of the newborns as a complication of the bloodstream dissemination following the colonization of the gastrointestinal tract (Nassif et al. 2002; Wooster et al. 2006; Parthasarathy et al. 2007). In order to invade the meninges, the bacteria must multiply in the blood and reach a high degree of bacteraemia (>10^6 colony forming units (CFU)/mL of blood) and, after this, cross the blood–brain barrier. Pathogens presenting brain tropism must avoid the host’s innate defence mechanisms, such as the complement (C) system and opsonophagocytosis, to survive in the bloodstream. This suggests that these microorganisms possess very specific virulence factors which are essential for meningeal invasion.

One of the significant factors in the pathogenicity of E. coli K1 strains is their resistance to the action of the C. The mechanism of the resistance to the bactericidal effect of the serum is not completely explained but it is known that this phenomenon has a multifactor basis. The organisation of the bacterial outer membrane and the structures such as bacterial capsule, outer membrane proteins (OMPs), and O-specific side chains of lipopolysaccharides (LPS) are the factors determining the susceptibility of the bacteria to the bactericidal action of the C (Taylor 1992, 1995).

The capsular polysaccharide produced by E. coli K1 strains is a homopolymer of α-2,8-linked sialic acid (N-acetylneuramininic acid, Neu5Ac, NeuAc) residues and is structurally and immunologically identical to the capsular polymer of Neisseria
meningitidis group B, Pasteurella haemolytica A2, and Moraxella nonliquefaciens (Ferrero and Aparicio 2010). This polysaccharide can mimic the mammalian polysialic acid structure that can be found predominantly on the neural cell adhesion molecule (NCAM), which is responsible for mediating several neuronal functions by controlling intercellular adhesion, neurite outgrowth, cell migration, proliferation, and survival. Due to biochemical similarity with polysialylated form of humans NCAM, bacterial α-2,8-linked sialic acid is poorly immunogenic in humans and, hence, its immunological tolerance is observed.

The presence of NeuAc on the surface of the bacteria impedes the action of the C by enhancing the binding of factor H to C3b on the outside of the cell and, consequently, by reducing the amplification of C3b production and downregulation of the activity of the alternative pathway (Rautemaa and Meri 1999; Kugelberg et al. 2008). The protection against serum killing is also the result of the interaction between C4b-binding protein (C4bp), the predominant serum inhibitor of C3b activation via the classical pathway, and the outer membrane protein A (OmpA) of E. coli K1 rods (Wooster et al. 2006; Prasadara et al. 2002). Moreover, the component of the bacterial capsule, α-2,8-linked Neu5Ac, can provide resistance against the C attack by inhibiting the insertion of membrane attack complex (MAC) into the bacterial membrane. Therefore, an increased synthesis of K1 capsule has a significant impact on the activation of the C, opsonization, and virulence of E. coli K1 rods. On the other hand, E. coli K1 strains demonstrate different degrees of sensitivity to the bactericidal action of the serum, which indicates that other outer membrane structures may also play a crucial role in the determination of bacterial resistance to the lytic action of the serum. Subsequently, we have obtained K1– mutant forms (E. coli K1–) that have spontaneously lost the ability to produce K1 capsular antigen. In the present study, we have used sodium dodecyl sulphate-polyacrylamide gel electrophoresis (SDS-PAGE) in order to investigate outer OMPs band composition of E. coli K1– strains and to discern the relationship between OMPs pattern and susceptibility to NHS.

Material and methods

Bacterial strains Eighteen E. coli K1 strains isolated from urine of children with urinary tract infections and a standard E. coli B2095 K1 strain were used to obtain unencapsulated E. coli K1−. K1− mutant forms were derived by selection for resistance to the K1-specific bacteriophage.

Detection of capsular antigen K1 in E. coli The K1–specific bacteriophage (K1 A) was used for the detection of capsular antigen K1 in accordance with the procedure used by Devine et al. (1990). The E. coli strains were grown overnight on nutrient agar, and then bacterial cells were inoculated into 4 mL of nutrient broth and incubated at 37 °C for 4 h. Afterwards, a single streak of each culture was applied to nutrient agar and allowed to dry. Once the streak was dry, 10 μL of bacteriophage suspension (8 × 10^10 PFU/mL) was placed in the centre of the streak. After incubation at 37 °C for 18 h, the presence of K1 was indicated by a clear zone of bacterial lysis at the site of the placement of bacteriophage suspension. E. coli K1− forms were obtained from colonies growing in an otherwise clear zone of bacterial lysis where the K1 A specific bacteriophage was used. Once isolated, these colonies were tested three times to check whether the loss of the ability to produce K1 capsule is a permanent feature. The E. coli K1− colonies were inoculated into nutrient broth and incubated at 37 °C for 4 h; following this, a single streak of each culture was applied to nutrient agar and
allowed to dry. Next, the phage K1 A suspension was placed in the centre of the streak. After incubation at 37 °C for 18 h, no zone of bacterial lysis was observed.

Phage K1 A and standard strain *E. coli* B2095 (O2:K1) used for the proliferation of the phage were obtained from the collection of Dr. G. Schmidt (Forschungsinstitut Borstel, Institut für Experimentelle Biologie und Medizin, Germany). The bacteriophage suspension was stored in 2-mL portions at 4 °C.

Serum NHS was obtained from healthy adult volunteers untreated with any antimicrobial drugs. The samples of NHS were collected, pooled, and stored in 0.5 mL portions at −70 °C. A suitable volume of the serum was thawed immediately before experiments. Each portion was used only once.

Bactericidal activity of serum The bactericidal activity of NHS was determined as described previously (Cisowska et al. 2004). Briefly, the strains were grown overnight and then bacterial cells of early exponential growth phase were transferred to fresh nutrient broth and incubated at 37 °C for 0.5 h. After incubation, the bacterial cells were centrifuged (4,000 rpm for 20 min). Next, 0.5 mL of bacterial suspension was added to the dividing NHS (serum was diluted with 0.1 mol/L NaCl) giving the final serum concentration of 50%; the samples were immediately collected and diluted and then cultured on nutrient agar plates for 18 h at 37 °C (0 min). Bacteria with NHS were incubated in a water bath at 37 °C; the samples were collected after 30, 60, and 180 min and then they were cultured and handled in the same manner as described above. The number of CFU at 0 min was assumed as 100%. Strains with a survival ratio of not less than 50% after 180 min of incubation in the sera were regarded as resistant.

Isolation of the OMPs The OMPs were isolated using procedure described by Murphy and Bartos (1989) with our own modifications. For the OMP isolation, we used bacteria grown in 100 mL brain heart infusion broth (Difco) at 37 °C for 18 h. After incubation, the bacterial cells were centrifuged (4,000 rpm at 4 °C for 20 min) and the resulting pellet was suspended in 2.5 mL of buffer B (1 mol/L sodium acetate, 0.001 mol/L β-mercaptoethanol; pH 4.0). Afterwards, 22.5 mL of the solution containing 5% Zwittergent Z 3-14® (Calbiochem–Behring) and 0.5 mol/L CaCl₂ was added. This solution was stirred at room temperature for 1 h. To precipitate nucleic acids, a volume of 6.25 mL of 96% cold ethanol was added very slowly. Subsequently, the mixture was centrifuged at 12,300 rpm at 4 °C for 10 min. The remaining proteins were precipitated by the addition of 93.5 mL of 96% cold ethanol and by centrifugation at 12,300 rpm at 4 °C for 20 min. The pellet was suspended in 5 mL of buffer Z (0.05% Zwittergent, 0.05 mol/L Tris and 0.01 mol/L EDTA; pH 8.0) and stirred for 1 h at room temperature. The solution was kept at 4 °C overnight and centrifuged at 8,700 rpm at 4 °C for 10 min to obtain OMPs in the supernatant. Following the isolation of the OMPs, we checked the enzymatic activity of succinic dehydrogenase, a marker of cytoplasmic protein membranes, in the soluble fraction of buffer Z using the method of Rockwood et al. (1987). The Zwittergent-extracted OMPs contained no detectable succinic dehydrogenase activity, indicating that these preparations were not contaminated by cytoplasmic membrane.

Protein quantification was performed with the bicinchoninic acid (BCA) Protein Assay Kit (PIERCE®) with bovine serum albumine (Sigma) as the standard. The Pierce BCA Protein Assay is a detergent-compatible formulation based on BCA for the colorimetric detection and quantification of total protein concentration (PIERCE Instruction). The purple-coloured reaction product (reduction of Cu²⁺ to Cu⁺ by protein in an alkaline medium) exhibits a strong absorbance at 562 nm with increasing protein concentration over a broad range of 20–20,000-μg/mL. OMPs after preparation in the sample buffer were loaded into each of the wells of the SDS-PAGE gels in comparable concentrations of 9–10 μg per well.

Polyacrylamide gel electrophoresis SDS-PAGE was carried out on slabs with 12.5% acrylamide as described by Laemmli (1970). Samples were applied to slabs after heating at 100 °C for 4 min. The Wide Molecular Weight Range Sigma Marker M4038 protein standard (6-205 kDa) was used for molar mass calibration. Electrophoresis was performed at 100 V for 25 min and then at 200 V for 45 min. After the electrophoresis, gel was stained for 1 h with a solution containing 25% methanol, 10% acetic acid, and 0.05% Coomassie Brilliant blue and decolorized with 10% acetic acid for about 3 h (we used identical procedure for *E. coli* K1+; Cisowska et al. 2005). For the molecular analysis of OMPs BIO-CAPT v. 99 computer program as well as the BIO-1D++ v. 99 program (Vilber Lourmat, France; BioRad) were used.

Results

The results concerning the susceptibility to 50% NHS of *E. coli* strains without K1 polysaccharide capsule (K1− forms) are enclosed in Table 1. The detailed results of susceptibility of *E. coli* K1+ strains to 50% NHS were previously published (Cisowska et al. 2004); therefore, in Table 1 we presented only the survival percent after 3 h of incubation in NHS for K1+ strains in order to make the comparison between K1+ and K1− forms easier. The tested *E. coli* K1+ rods showed diverse susceptibility to the bactericidal action of NHS. Among 19 *E. coli* K1+ strains, eight (*E. coli* 988, 601, 529, 544, 1000, 315, 105, and 380) were sensitive to NHS. The remaining strains were resistant to NHS and the survival ratio of these strains after 180 min of incubation in serum ranged from 82.8 to
4,250.0 % of the initial cell number. Most of *E. coli* K1− isolates were sensitive to the bactericidal action of NHS (*E. coli* 380, 763, 848, 500, 5915, 309, 988, 601, 529, 158, 481, 544, 1000, 328, 315, 34, and 105). Moreover, in the case of some of them, NHS decreased the percent of survival to <0.0001 % even after 30–60 min of incubation. Two *E. coli* K1− strains were resistant to the serum and proliferated in NHS very intensively (*E. coli* 111 and B2095).

To investigate OMPs band patterns of *E. coli* K1− strains and evaluate whether there is a relationship between OMPs composition and susceptibility of these unencapsulated forms to bactericidal action of NHS, the SDS-PAGE was used. The electrophoretic band patterns for the OMPs of these rods and the molar mass of these proteins are shown in Figs. 1, 2, and 3.

The analysis performed on OMPs has not revealed a direct correlation between the various OMPs band patterns and the susceptibility of these strains to the serum. *E. coli* K1− strains sensitive to the lytic activity of NHS possessed between 12 and 22 different OMPs bands and the molar mass of these proteins ranged from 13.9 to 106.2 kDa (Figs. 1, 2, and 3). The greatest diversity in the quantity of possessed OMPs was observed for *E. coli* 763 K1− strain, which had 22 bands corresponding to proteins with molar mass ranging from 14.0 to 93.1 kDa (Fig. 3, lane 3). The lowest diversity was found for *E. coli* 1000 K1− strain. In this case, only 12 bands, corresponding to proteins with molar mass between 17.5 and 59.9 kDa, were obtained (Fig. 2, lane 3). Addition-ally, NHS-resistant K1− strains *E. coli* B2095 and 111 did not differ significantly from NHS-sensitive strains. It was noted also that *E. coli* K1− strains resistant to serum contained 13 (*E. coli* B2095; Fig. 2, lane 9) or 15 (*E. coli* 111; Fig. 1, lane 6) different OMPs with the molar mass ranging from 16.9 to 60.7 and 17.5 to 66.4 kDa, respectively. However, proteins with higher molar mass were detected only in certain K1− strains sensitive to the serum. These proteins were observed for example in the case of strains: *E. coli* 988 K1− (Fig. 1, lane 2; OMP: 102 and 75 kDa), *E. coli* 601 K1− (Fig. 1, lane 3; OMP: 102, 93, and 79 kDa), *E. coli* 529 K1− (Fig. 1, lane 4; OMP: 89 and 79 kDa), *E. coli* 315 K1− (Fig. 2, lane 5; OMP: 83 kDa), *E. coli* 34 K1− (Fig. 2, lane 6; OMP: 106, 97, 84, and 74 kDa), *E. coli* 763 K1− (Fig. 3, lane 3; OMP: 93, 86, and 77 kDa), *E. coli* 848 K1− (Fig. 3, lane 4; OMP: 105 and 78 kDa), or *E. coli* 309 K1− (Fig. 3, lane 7; 95 kDa).

### Table 1  Bactericidal activity of 50 % NHS against *Escherichia coli* K1+ and K1− strains

| Strain no. | Survival % after 180 mina | Time of incubation (min) | Survival % after 180 min |
|------------|---------------------------|--------------------------|--------------------------|
|            |                           | 0 CFU | 30 CFU | 60 CFU | 180 CFU |                           |
| 988        | 39.4                      | 2.4×10⁷ | 5.4×10⁶ | 4.6×10⁵ | 4.0×10⁴ | 0.2                      |
| 601        | 1.6                       | 9.8×10⁶ | 1.8×10⁵ | <1.0×10⁴ | <0.5×10⁴ | <0.00005                 |
| 529        | 0.0004                    | 1.4×10⁷ | 2.2×10⁶ | 2.7×10⁵ | <0.7×10⁵ | <0.00005                 |
| 158        | 480.5                     | 6.5×10⁶ | 4.7×10⁵ | 3.2×10⁴ | <0.5×10⁴ | <0.00008                 |
| 111        | 4,250.0                   | 1.7×10⁷ | 3.1×10⁷ | 4.8×10⁷ | 1.7×10⁸ | 1,000.0                  |
| 481        | 109.1                     | 1.3×10⁷ | 9.8×10⁶ | 1.1×10⁶ | 1.3×10⁶ | 10.0                     |
| 544        | 0.02                      | 1.3×10⁷ | 2.3×10⁶ | <1.0×10⁷ | <0.5×10⁷ | <0.00004                 |
| 1000       | 0.1                       | 6.9×10⁶ | <0.5×10⁴ | <0.5×10⁴ | <0.5×10⁴ | <0.00007                 |
| 328        | 82.8                      | 1.2×10⁷ | 2.7×10⁶ | <0.7×10⁷ | <0.5×10⁷ | <0.00004                 |
| 315        | 0.4                       | 2.1×10⁷ | <0.5×10⁴ | <0.5×10⁴ | <0.5×10⁴ | <0.00002                 |
| 34         | 353.8                     | 7.0×10⁶ | <1.0×10⁴ | <0.5×10⁴ | <0.5×10⁴ | <0.00007                 |
| 105        | 9.2                       | 5.7×10⁶ | <0.5×10⁴ | <0.5×10⁴ | <0.5×10⁴ | <0.00009                 |
| B2095      | 3,000.0                   | 1.8×10⁷ | 2.6×10⁷ | 1.5×10⁸ | 2.4×10⁸ | 12,973.0                 |
| 380        | 13.1                      | 2.0×10⁷ | <0.5×10⁴ | <0.5×10⁴ | <0.5×10⁴ | <0.00002                 |
| 763        | 683.3                     | 9.8×10⁶ | 1.3×10⁶ | 4.9×10⁴ | 2.3×10² | 0.02                     |
| 848        | 275.0                     | 1.1×10⁷ | <0.5×10⁴ | <0.5×10⁴ | <0.5×10⁴ | <0.00004                 |
| 500        | 418.2                     | 8.3×10⁶ | 6.8×10⁵ | 6.4×10⁴ | 7.9×10³ | 0.09                     |
| 5915       | 194.1                     | 1.9×10⁷ | 0.7×10⁴ | <0.5×10⁴ | <0.5×10⁴ | <0.00003                 |
| 309        | 769.2                     | 8.0×10⁶ | <0.5×10⁴ | <0.5×10⁴ | <0.5×10⁴ | <0.00006                 |

*CFU* colony forming units

a Detailed results of the susceptibility of *E. coli* strains K1+ to NHS published in Cisowska et al. (2004)
The comparison of the outer membrane proteins patterns of the two serum-resistant K1− strains, E. coli B2095 (Fig. 2, lane 9) and 111 (Fig. 1, lane 6), shows a close correspondence. These rods mostly produced the same OMPs (60, 47, 26, 23, and 16 kDa) and differed only in respect to a few of them (E. coli 111 – 66, 40, and 30 kDa proteins; E. coli B2095, 26 kDa protein). Since the bands corresponding to proteins with such molar masses were also detected in the group of NHS-sensitive K1− strains, we could not relate the occurrence of a specific OMP to serum susceptibility.

Several differences can be seen when SDS-PAGE patterns of the outer membrane proteins of both NHS-resistant K1+ and K1− E. coli B2095 forms are examined (Fig. 2, lanes 8 and 9). The number of OMPs generated by K1− mutant was comparable to that of the K1+ strain, but differed qualitatively. OMPs with the molar masses: 113, 93, 71, 50, 46, and 38 kDa were not produced by E. coli B2095 K1− form, and, additionally, some proteins (52, 47, 26, 23, and 16 kDa) were not formed by K1+ rods. Moreover, E. coli K1− mutant over-expressed some of the proteins (22, 33, and 35 kDa) as compared to K1+ form.

**Discussion**

During the immune responses, the ability to avoid killing by the serum C is an important evasion strategy of bacterial pathogens. The mechanism of the bacterial resistance to the bactericidal effect of the serum is not fully understood; however, it is known that the OMPs are one of the factors determining the resistance or sensitivity of bacteria to the bactericidal action of the C (Taylor 1992, 1995).

In the bloodstream, N. meningitidis is exposed to the antibody/complement-mediated lysis, as well as to opsonophagocytosis, and it seems that the quantity of the polysaccharide capsule that is produced influences the resistance of this pathogen to killing by the host (Uria et al. 2008). Only the
Nop protein, another OMP commonly expressed by resistance of these pathogens (Madico et al. 2006). Also, the regulator of the alternative C pathway, which promotes serum by all meningococcal strains, can recruit C factor H, a key bacteria from MAC-mediated lysis. Furthermore, the major bacterial surface inhibits MAC formation and protects the survival (Hill et al. 2010). Factor H-binding protein, expressed cascade (factor H, C4bp), potentially leading to their increased a real but too small to interact with some negative regulators of the C cascade (factor H, C4bp), potentially leading to their increased survival (Hill et al. 2010). Factor H-binding protein, expressed by all meningococcal strains, can recruit C factor H, a key regulator of the alternative C pathway, which promotes serum resistance of these pathogens (Madico et al. 2006). Also, the Omp protein, another OMP commonly expressed by N. meningitidis strains, is able to bind vitronectin (Vn), which inhibits C5b-7 complex formation and C9 polymerization (Singh et al. 2010). As a consequence, Vn deposited on the bacterial surface inhibits MAC formation and protects the bacteria from MAC-mediated lysis. Furthermore, the major OMP of meningococci Por A can bind to the C regulator C4-binding protein and influence serum resistance; although, it may be inhibited by the capsule (Jarva et al. 2005). It must be mentioned, however, that Drogari-Apiranthitou et al. (2002) did not find a significantly reduced C3b deposition in two PorA(−) mutant strains and showed similar amounts of MAC deposition in PorA(+) and PorA(−) variants, regardless of the variance in their serum sensitivity.

In the present work, the OMPs of 19 E. coli K1− mutant strains were isolated and subjected to SDS-PAGE in order to characterise OMPs. In our opinion, the enhancement of the resistance to C-mediated bactericidal activity of the serum may be associated with the presence of the specific OMPs in bacterial outer membrane. It has been demonstrated that OmpA increases the resistance to bactericidal effect of the serum by the activation of the C classical pathway and is the source of K1 virulence (Weiser and Gotschlich 1991). Relationship between the resistance to the C, virulence of E. coli strains, and OMPs banding pattern was also noted by Chaffer et al. (1999). These authors observed that highly resistant to bactericidal effect of the C and highly virulent E. coli rods had a weak peptide band located at 35 kDa, while the occurrence of a strong 35 kDa OMP band was associated with a lower virulence and C susceptibility. In our previous work, we noticed the presence of the peptide band located at 35 kDa both in E. coli strains that were resistant and sensitive to the normal bovine serum and normal cord serum (Cisowska et al. 2005). Also, in the present study both of the NHS-resistant K1− mutant strains, E. coli B2095 and 111, and NHS-sensitive K1− strains (158, 309, 481, 848, 5915, 105, 315, 328, 380, 544, 601, 988, and 1000) possessed peptide band located at ~35 kDa. Only four K1− strains (34, 500, 763, and 529) did not have ~35 kDa OMP, while their K1+ forms generated a protein band with such molar mass (Cisowska et al. 2005).

Other investigators associated the resistance of E. coli rods to the C-mediated serum killing with 25 kDa OMP (Moll et al. 1980). In the case of the tested K1− mutant strains, a peptide band located at ~25 kDa was detected only in a few NHS-sensitive E. coli (158, 380, 544, 988, 1000, 34, 5915, and 848). However, both of the serum-resistant K1− strains, E. coli B2095 and 111, did not produce this protein. Moreover, in our previous work, we noticed the presence of the peptide band located at ~25 kDa in E. coli K1+ strains both resistant (111, 481, 763, 34, and 5915) and sensitive (328, 529, 380, 544, 988, and 1000) to bactericidal effect of the serum (Cisowska et al. 2005).

The comparison of the OMPs patterns of E. coli K1+ (Cisowska et al. 2005) and their K1− mutant forms, both the NHS-resistant and NHS-sensitive strains, showed that they differed in the amount and the type of produced OMPs. Regardless of the susceptibility of E. coli K1+ rods to bactericidal activity of NHS, the majority of K1− mutants produced a smaller amount of OMPs. In particular, this difference can be seen in the case of E. coli 1000, 544, 328, and 158 K1− rods that produced even 23–26

![Fig. 3 SDS-PAGE patterns of OMPs from E. coli K1− strains. Molar mass marker, 2 E. coli 380, 3 E. coli 763, 4 E. coli 848, 5 E. coli 500, 6 E. coli 5915, 7 E. coli 309](image-url)
OMPs less than the corresponding K1+ variants. In contrast, in only three unencapsulated K1− forms the number of distinct electrophoretic components increased. Serum-resistant E. coli 34 K1+ and 763 K1+ as well as NHS-sensitive E. coli 601 K1+ generated 16, 17, and 15 OMPs, respectively, while their K1− forms produced 21, 22, and 19 OMPs, correspondingly.

In conclusion, in our study, we have not found any similarities between OMPs’ band patterns of analysed group of E. coli K1− rods. Our study did not find a OMPs band pattern characteristic for the analysed group of E. coli K1− rods. The diversity of OMPs may indicate a significant role that these proteins play in bacterial resistance to the bacteriolytic action of the C. On the other hand, presumably, the proteins that sensitise the bacterial cell to the bactericidal action of the C can be located among the OMPs that were analysed in this study.

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