The aim of this work was to develop a new software tool for identifying gene mutations that determine the porin-mediated resistance to antibiotics in gram-negative bacteria and to demonstrate the functionality of this program by detecting porin-mediated resistance to carbapenems in clinical isolates of *Pseudomonas aeruginosa*.

**Materials and Methods.** The proposed algorithm is based on searching for a correspondence between the reference and the studied genes. When the sought nucleotide sequence is found in the analyzed genome, it is compared with the reference one and analyzed. The genomic analysis is then verified by comparing between the amino acid sequences encoded by the reference and studied genes. The genes of the susceptible *P. aeruginosa* ATCC 27853 strain were used as the reference nucleotide sequences encoding for porins (OprD, OpdD, and OpdP) involved in the transport of carbapenems into the bacterial cell. The complete genomes of clinical *P. aeruginosa* isolates from the PATRIC database 3.6.9 and our own collection were used to test the functionality of the proposed program. The analyzed isolates were phenotypically characterized according to the CLSI standard. The search for carbapenemase genes in the studied genomes of *P. aeruginosa* was carried out using the ResFinder 4.1.

**Results.** The developed program for detecting the genetic determinants of non-plasmid antibiotic resistance made it possible to identify mutations of various types and significance in the porin genes of *P. aeruginosa* clinical isolates. These mutations led to modifications of the peptide structure of porin proteins. Single amino acid substitutions prevailed in the OpdD and OpdP porins of carbapenem-susceptible and carbapenem-resistant isolates. In the carbapenem-resistant strains, the gene encoding for OprD porin was found heavily modified, including insertions and/or deletions, which led to premature termination of porin synthesis. In several isolates resistant to meropenem, no mutations were detected in the gene encoding for OprD, which might be associated with alternative mechanisms of resistance to carbapenems.

**Conclusion.** The proposed software product can become an effective tool for deciphering the molecular genetic mechanisms of bacterial chromosomal resistance to antibiotics. Testing the program revealed differences between the occurrences of mutations significant for carbapenem resistance in the oprD, opdD, and opdP genes.

**Key words:** antibiotic resistance; *Pseudomonas aeruginosa*; OpdP; OprD; OpdD; porin genes; carbapenems.

**How to cite:** Savinova T.A., Samchenko A.A., Bocharova Y.A., Mayansky N.A., Chebotar I.V. Computer program for detection and analyzing the porin-mediated antibiotic resistance of bacteria. Sovremennye tehnologii v medicine 2021; 13(6): 15, https://doi.org/10.17691/stm2021.13.6.02

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**Introduction**

The global occurrence of antimicrobial resistance (AMR), has become one of the most important problems of modern health care [1]. The situation with AMR is expected to get worse due to the large-scale use of antibiotics associated with the COVID-19 pandemic [2, 3]. It is for medical science to improve methods for assessing antibiotic resistance and deciphering its mechanisms. At the present stage, the commonly used phenotypic characterization of AMR is insufficient for successfully fighting the AMR; the phenotypic profile of resistant bacteria should be supplemented by an insight into the mechanisms of its formation [4].

Methods for identifying genetic determinants of resistance are among the most informative methods for describing the mechanisms of AMR formation. The main determinants of AMR are traditionally subdivided into
P. aeruginosa carbapenems in clinical isolates of P. aeruginosa. Chromosomal resistance is associated with mutations (point mutations, small insertions/deletions, extensive insertions/deletions, including insertions of mobile genetic elements) in chromosomal genes specific for this type of bacteria [8–10].

To search for plasmid genes involved in AMR, online software tools/programs that require a locally installed (standalone) version have been created and are successfully used. These include ResFinder, CARD, ARDB, ARG-ANNOT, and other resources [11–14]. They are primarily aimed at finding genes that are transmitted as mobile genetic elements, for example, genes for beta-lactamases (resistance to beta-lactam antibiotics), aminoglycoside transferases (resistance to aminoglycosides and fluoroquinolones), chloramphenicol acetyltransferases (resistance to chloramphenicol), glutathione-S-transferases (resistance to fosfomycin), genes for ribosomal protection proteins (resistance to tetracycline) and other determinants of antibiotic resistance. Some of the existing tools are capable of searching for chromosomal determinants of drug resistance as well (PointFinder, CARD).

Unfortunately, the functionality of these software tools does not always allow one to identify and analyze important genetic determinants of AMR associated with changes in porins responsible for transportation of antibiotics into the cell, regulatory genes for global efflux systems, and genes encoding for antibiotic targets. In particular, this applies to deciphering the molecular mechanisms of resistance to latest generation of antibiotics including carbapenems.

The aim of this work was to develop a new software tool for identifying gene mutations that determine the porin-dependent resistance to antibiotics in gram-negative bacteria and to demonstrate the functionality of this program by detecting porin-mediated resistance to carbapenems in clinical isolates of P. aeruginosa.

Materials and Methods

The overall study design included:
1) creating a database of porins’ reference genes involved in the antibiotic transport through the outer bacterial membrane into the periplasmic space;
2) obtaining the amino acid sequences of porins in the given clinical isolates (based on their full genomic data);
3) using the originally developed software to identify the genetic determinants of non-plasmid antibiotic resistance [15]; the algorithm must be able to detect the differences between the analyzed gene (or protein) sequence and the reference sequences stored in the database; testing the program using the model of porin-mediated bacterial resistance to meropenem and imipenem.

The program includes a database of reference porin genes involved in the AMR development in clinically significant species of gram-negative bacteria, including Pseudomonas aeruginosa, Klebsiella pneumoniae, Escherichia coli, Enterobacter spp., Salmonella enterica, etc. The sequences of the reference genes were obtained from the NCBI database (https://www.ncbi.nlm.nih.gov/nuccore/) by selecting the sequences of porin genes in strains that are recommended by the international standards EUCAST (European Committee on Antimicrobial Susceptibility Testing) and CLSI (Clinical and Laboratory Standards Institute, USA) as antibiotic susceptibility standards. In the present study, for the reference, we used the drug-susceptible ATCC 27853 strain of P. aeruginosa and the nucleotide sequences of the genes encoding for the respective porins OprD/OccD1, OpdD/OccK7, and OpdP/OccD3, whose participation in the mechanism of resistance to carbapenems was confirmed earlier (Table 1).

As a negative control (strains lacking species-specific P. aeruginosa porins), 10 genomes from the PATRIC 3.6.9 database (https://www.patricbrc.org/) of five different species were analyzed: Acinetobacter baumannii (SP5515, GML-KP48-AB-TR), Klebsiella pneumoniae (AR0361, AR438), Escherichia coli (AR435, AR0450), Staphylococcus aureus (AR464, AR0216), and Enterococcus faecium (VREN1530, VREN2775).

To test the capabilities of the proposed program, complete genomes (complete assemblies or contigs) of P. aeruginosa from two sources were used. The first source was the PATRIC 3.6.9 database, from which 15 genomes of clinically relevant P. aeruginosa strains were obtained together with the AMR characteristics of

### Table 1

| Pseudomonas aeruginosa porins involved in the transport of carbapenems |
|---|---|---|---|
| **Porin name** | **Alternative name** | **Substrate** | **Literature** |
| OprD | OccD1 | Meropenem, imipenem, lysine, histidine, arginine, ornithine | [16–18] |
| OpdD | OccK7 | Meropenem | [18] |
| OpdP | OccD3 | Meropenem, glycine-glutamate, arginine | [19, 20] |
these isolates. The second source of genomes (10 samples) was our own collection of DNA from clinically-relevant strains, which had their AMR spectra (sensitivity to meropenem and imipinem) previously characterized by the phenotypic CLSI standard, 2020 (https://clsi.org/standards/).

The search for adaptive genes of carbapenem resistance (genes of carbapenemases) in the studied genomes of *P. aeruginosa* was carried out using the ResFinder 4.1 resource (https://cge.cbs.dtu.dk/services/ResFinder/).

The function of the program is based on the algorithm depicted in the Figure. At the first stage, a search for a match between the reference and studied genes is performed using the BLAST program based on the best-matching choice [21]. When the desired nucleotide sequence is found in the analyzed genome, it is compared to the reference one. If they differ, the data is analyzed by the program and the result is automatically displayed in a text or tabular format to present the detected changes in the nucleotide sequence of the gene. The significance of the detected changes for the emergence of AMR is tested by analyzing the synonymy of substitutions in the amino acid sequence of porin. Synonymy analysis is performed automatically; this function is incorporated in the program code.

## Results

Genes encoding for the OprD, OpmD, and OpmP porins were found in the genomes of all *P. aeruginosa* isolates (n=25) analyzed in this study. By contrast, none of these genes were found in the genomes of non-*P. aeruginosa* isolates (n=10) used as a negative control.

In the genomes of the 25 studied isolates of *P. aeruginosa*, mutations of various types and significance were identified. In 15 isolates, mutations of the oprD gene were detected, in 22 isolates — mutations of the opmD gene, and in 13 isolates — mutations of the opmP gene (Table 2). In total, 4 groups of mutations were identified.

### Table 2

**Peptide structure modifications in porins of Pseudomonas aeruginosa genomes (n=25)**

| Modifictions                                      | The number of genomes in which mutations corresponding to porin changes were found |
|---------------------------------------------------|----------------------------------------------------------------------------------|
|                                                   | OprD | OpmD | OpmP |
| No changes in the product (genes are identical to the reference, or synonymous mutations in the gene) | 10   | 3    | 12   |
| AAS                                               | 3    | 18   | 12   |
| Stop/AAS + stop                                   | 1    | —    | —    |
| Frameshift + stop/AAS + frameshift + stop         | 5    | 4    | 1    |
| AAS + INS                                         | 6/25 | —    | —    |

*Here: AAS — amino acid substitution; stop — stop codon leading to premature stopping of porin synthesis; frameshift — shift of the reading frame; INS — amino acid insertion without frameshift.*
found: those led to modifications of the porin peptide structure.

Amino acid substitutions in the OpdD and OpdP porins were detected most often. Porins OprD and OpdD were characterized by a greater variety of modifications, including insertions and/or deletions accompanied by the appearance of a stop codon.

We then analyzed whether the phenotypic manifestation of AMR correlated with the presence of mutations in the porin genes or the presence of carbapenemases genes (genotype/phenotype ratio). The AMR was assessed by the data on participation of porins in the antibiotics transport through the outer membrane (see Table 1; Table 3). In 7 out of 8 isolates sensitive to meropenem, either no mutations were observed in the genes for the OprD and wild-type OpdP porins, or 1 to 4 mutations were detected in these genes. One meropenem susceptible isolate had an OprD mutant porin (16 amino acid substitutions). The most frequent oprD mutations in isolates resistant or with intermediate resistance to carbapenems were mutations that led to amino acid substitutions and insertion of two amino acids (6/22), as well as a frameshift mutation leading to premature termination of porin synthesis (5/22). In 3 isolates insensitive to meropenem, no mutations in oprD were detected.

Isolated substitutions (1–8 AAS) predominated in porins OpdD and OpdP, and this pattern of AAS was observed in both carbapenem-sensitive and carbapenem-insensitive isolates. Deletion of one nucleotide in the opdD gene, leading to premature termination of porin synthesis, was detected in 4 carbapenem-insensitive isolates (in 3 — resistant to meropenem and imipenem, and in 1 — with an intermediate level of resistance to meropenem). In 1 isolate resistant to meropenem and imipenem, the loss of functional porin OpdP resulted from the appearance of a stop codon in the nucleotide sequence encoding for position 81 of the protein.

Discussion

In most cases, P. aeruginosa clinical isolates exhibit resistance to beta-lactam antibiotics, including carbapenems [22, 23]. Methods for detecting beta-lactamase-mediated resistance are quite simple and well implemented in the routine laboratory practice [24, 25]. At the same time, the analysis of chromosomal determinants of resistance is associated with difficulties with data processing. The proposed computer program is aimed at identifying the genetic determinants of non-plasmid antibiotic resistance and mutations in the oprD, opdD, and opdP genes of porins that transport carbapenems into the bacterial cell. This program also allows us to estimate changes in the amino acid structure of the respective proteins associated with resistance to carbapenem.

The analysis of 35 genomes (25 genomes of P. aeruginosa and 10 genomes of five other bacterial species) provided evidence for the usefulness of the proposed program in the search for oprD, opdD, and opdP genes and their mutations. The presence of these genes in all studied isolates of P. aeruginosa and their absence in the genomes of A. baumannii, K. pneumoniae, E. coli, S. aureus, and E. faecium indicates the specificity of the developed program in a search for target nucleotide sequences. Among P. aeruginosa isolates, the most diverse changes were found in the oprD gene, which corroborated with reports on the frequent occurrence of this gene in the carbapenem-resistant phenotype in P. aeruginosa [26].

The proposed program allowed us to identify the entire spectrum of theoretically possible types of mutations, including mutations leading to nonsynonymous amino acid substitutions, nonsense mutations (leading to the appearance of stop codons), insertions, deletions, and frameshifts.

The detection of carbapenem-resistant isolates containing the wild-type OprD can be due to the existence of alternative resistance mechanisms and, first of all, carbapenemases of the OXA groups (see Table 3). In addition, overexpression of the efflux system genes and mutations in genes of penicillin-binding proteins cannot be ruled out [23]. Further studies on molecular mechanisms of antibiotic resistance in bacteria are needed to provide a more comprehensive insight into the genetic basis of the resistant phenotypes.

Interestingly, different types of mutations, as well as their different numbers, were found in the oprD, opdD, and opdP genes. At the present stage, we have no tangible explanation for this intriguing statistical result. A mechanistic explanation of this phenomenon requires the knowledge of DNA repair in the genomes of P. aeruginosa, which is not available today.

Several strains, although resistant to imipenem, were found susceptible to meropenem. To explain this observation, we refer to the ability of meropenem to enter the cell via several alternative routes mediated by porins OprD, OpdD, or OpdP. For instance, if the oprD gene is mutated, meropenem can be transported via the non-mutated OpdD and OpdP porins. By contrast, imipenem has only one known “gateway” — porin OprD, which explains the discrepancy between imipenem and meropenem regarding the drug-susceptibility in the studied isolates of P. aeruginosa.

The proposed software product may have limitations associated with errors in sequencing and/or genome assembly, in which case the program produces the result “Gene not found”.

Overall, the developed algorithm allows for simultaneous search for resistance determinants in whole-genome sequences, the number of which is limited only by the abilities of the computer used.
### Table 3

Mutations in *Pseudomonas aeruginosa* porin genes and phenotypic manifestations of antimicrobial resistance (n=25)

| Isolate name   | Resistance phenotype (CLSI) | OprD (meropenem, imipenem) | OprD (meropenem) | OpdD (imipenem) | Correspondence of phenotype and genotype | Notes |
|----------------|-----------------------------|----------------------------|------------------|-----------------|----------------------------------------|-------|
| PATRIC_MRSN8915 | R R                         | N43D E57S R59S V127L E185Q P186G V189T S267A T276A G310E G312R A315G L347M L359V K387S, frameshift and L432stop | WT              | P31S            | Yes                                    | Gene blaOXA-395 found |
| PATRIC_MRSN994  | R R                         | N107R, frameshift and D113stop | WT              | WT              | Yes                                    | Gene blaOXA-395 found |
| PATRIC_MRSN8139 | I R                         | N43D V127L E185Q, frameshift and D231stop | A10V S132T T247A D401N | A29E A424T P445T | Yes                                    |       |
| PATRIC_MRSN8912 | R R                         | N43D E57S R59S T103S K115T Q142stop | T247A A3O4G     | A29E H159L      | Yes                                    |       |
| PATRIC_MRSN4841 | I S                         | N43D E57S R59S T103S K115T F170L E185Q P186G V189T Q202E A210I K230E T240S T262N S267A G281A Q296K E301Q G310E A315G L359V V372M insS S374N S375N S376V S377G A379K G380N L381Y insG G423A | F19S L72S, frameshift and L117stop | A29E T30S | Not obvious | Gene blaOXA-483 found |
| PATRIC_MRSN443463 | S S                      | WT                                 | A183T P245Q T247A | WT              | Yes                                    |       |
| PATRIC_MRSN9718  | R S                         | WT                                 | T247A R268H      | A29E            | Yes                                    | Gene blaOXA-396 found |
| PATRIC_MRSN435288 | S I                       | WT                                 | V77I            | WT              | Yes                                    | Gene blaOXA-50 found |
| PATRIC_MRSN321   | R R                         | S278P                              | S132T T247A A301V | WT              | Yes                                    | Gene blaOXA-396 found |
| PATRIC_MRSN25678 | I S                         | N43D E57S R59S V127L                | S132T D163G N182H V240M L246F T247A D249G S287T | L15M N215T       | Yes                                    |       |
| PATRIC_MRSN373401 | I R                       | WT                                 | V77I            | WT              | Yes                                    | Gene blaOXA-486 found |
| PATRIC_MRSN29192 | R I                         | N43D E57S R59S Q202E A210I K230E T240S T262N S267A G281A Q296K E301Q G310S L359V V372M insS S374N S375N S376V S377G A379K G380N L381Y insG | V77I T247A      | WT              | Yes                                    | Gene blaOXA-485, blaOXA-486 found |
| PATRIC_MRSN409337 | R S                       | N43D E57S R59S Q202E A210I K230E T240S T262N S267A G281A Q296K E301Q G310R L359V V372M insS S374N S375N S376V S377G A379K G380N L381Y insG | WT              | WT              | Yes                                    | Gene blaOXA-50 found |
| PATRIC_MRSN358800 | R R                       | N43D E57S R59S T103S K115T M141A, frameshift and D218stop | D231G T247A A29E, frameshift and M81stop | Yes |       |       |
| Isolate name | Resistance phenotype (CLSI) | Molecular structure of porin | Correspondence of phenotype and genotype | Notes |
|--------------|-----------------------------|-----------------------------|------------------------------------------|-------|
| PATRIC_MRSN401528 | S S WT | V77I H110Y WT | Yes |  |
| Pa_135-3 | S R WT | V77I | Yes | Gene blaOXA-486 found |
| Pa_212-2 | S R | N43D E57S R59S V127L E185Q P186G V189T S267A T276A G310E G312R A315G L347M L359V S401A Q422E | V77G WT | Yes |
| Pa_224-1 | I R WT | A183T P245Q T247A | WT | Yes | Gene blaOXA-50 |
| Pa_3-2 | S S WT | V77I P202S T247A A304G | WT | Yes |  |
| Pa_316-2 | I R | N43D E57S R59S T103S K115T F170L E185Q P186G V189T Q202E A210I K230E T240S T262N S267A G281A Q310E K310E A315G L359V V372M insS S374N S375N S376V S377GA379K G380NL381YinsG G423A | F19S L72S, frameshift and L117stop | A29E T30S | Yes | Gene blaVIM-2 |
| Pa_124-1 | S R WT | V77I S114N R173S V377M | WT | Yes | Gene blaOXA-484 |
| Pa_124-2 | R R Frameshift S153P and G193stop | V77I S114N R173S V377M | N103K | Yes | Gene blaOXA-38 |
| Pa_41728-1 | S R WT | T247A T344I N345D | WT | Yes | Gene blaOXA-48 |
| Pa_41748-3 | R R | N43D E57S R59S T103S K115T F170L E185Q P186G V189T Q202E A210I K230E T240S T262N S267A G281A Q310E K310E A315G L359V V372M insS S374N S375N S376V S377GA379K G380NL381YinsG G423A | F19S L72S, frameshift and L117stop | A29E T30S | Yes | Gene blaVIM-2 |
| Pa_98-3 | R R | N43D E57S R59S T103S K115T F170L E185Q P186G V189T Q202E A210I K230E T240S T262N S267A G281A Q310E K310E A315G L359V V372M insS S374N S375N S376V S377GA379K G380NL381YinsG G423A | F19S L72S, frameshift and L117stop | A29E T30S | Yes | Gene blaVIM-2 |

Notes: isolates from the PATRIC database (n=15) are designated by the PATRIC prefix, the clinical isolates from our own collection (n=10) — by the Pa prefix; R — resistant isolates; I — isolates with an intermediate level of resistance; S — susceptible isolates (according to CLSI criteria); frameshift — shift of the reading frame; stop — replacement of the coding codon with a stop codon, leading to the termination of protein synthesis; insS — serine insertion; insG — glycine insertion; WT — wild-type porin, in which there are no amino acid substitutions compared to the reference porin; carbapenemase genes are highlighted in bold (according to ResFinder).
Conclusion

The proposed novel computer program is an effective tool for deciphering the molecular genetic mechanisms of bacterial (chromosomal) resistance to antibiotics. Its use made it possible to detect differences between the occurrences of mutations significant for resistance to carbapenems in the oprD, opdD, and opdP genes. The oprD genes were found to be more prone to mutations than the opdP genes. Due to the shunting of the porin-mediated transport of meropenem, conclusions about the resistance of a given strain to this antibiotic should be based not only on the detection of porin OprD abnormalities but also on the presence of significant mutations in the opdP and opdD genes. The present data suggest that meropenem is a more promising antibiotic than imipenem in the fight against P. aeruginosa strains with carbapenem resistance associated with impaired porin function.

Acknowledgments. We thank the Center for Precision Genome Editing and Genetic Technologies for Biomedicine (Moscow, Russia) for the genetic research methods.

Research funding. The study was supported by the Russian Science Foundation (Project ID 20-15-00235).

Conflicts of interest. The authors declare no conflicts of interest.

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