Alteration of Pseudomonas aeruginosa respiration by 4-(1-adamantyl)-phenol derivative

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Derivatives of 4-(1-adamantyl)-phenol are a promising class of antimicrobials affecting the structural integrity and functions of the bacterial cell membrane. The functioning of Pseudomonas aeruginosa respiratory chain and related system of oxidative phosphorylation was investigated before and after treatment with a derivative of 4-(1-adamantyl)-phenol (compound KVM-97). Oxygen consumption was measured polarographically with a Clark-type oxygen electrode. KVM-97 was tested at 0.5× and 1.0× MIC (minimum inhibitory concentration). Specific substrates of the respiratory chain (either 3.0 mM glutamate with 2.0 mM malonate or 3.0 mM succinate with 5.0 μM rotenone) were used. All reactions were stimulated by addition of ADP (0.2 mmol). It was found that at tested concentrations, KVM-97 inhibited the endogenous respiration and substrate oxidation in P. aeruginosa cells. The inhibiting effect was dose-dependent and more pronounced with succinate oxidation rather than glutamate oxidation. The respiratory control index value (RCI) in compound-treated cells was in average 1.5 times lower compared to the intact cells. The decrease in the RCI was related to changing the oxygen uptake rates in state 3 and state 4, which indicate the uncoupling of respiration and oxidative phosphorylation. The data obtained showed that 4-(1-adamantyl)-phenol derivative inhibits oxygen consumption and has uncoupling effects in P. aeruginosa cells.

Keywords: Pseudomonas aeruginosa, adamantane derivative, antibacterial agents, respiration

INTRODUCTION

Pseudomonas aeruginosa is an opportunistic human pathogen causing a wide range of life-threatening acute and chronic infections, including cystic fibrosis, pneumonia, bacteraemia, infections of the urinary tract, and wounds (Page, 2009; Tielen, 2013; Turner, 2014; Yayan, 2015). According to the WHO list of "priority pathogens", P. aeruginosa is one of twelve microorganisms for which new antibacterial agents are most desperately needed (Tacconelli, 2017). The main challenges in
the treatment of Pseudomonas infections are low pathogen susceptibility to the majority of antimicrobial agents, high virulence, and the ability of biofilm formation. Interestingly, the synthesis of both virulence factors and biofilm components (e.g., alginate) depends on oxygen availability and energy metabolism (Williams, 2006). Therefore, the energy-generating pathways could be considered one of the potential drug targets for the development of the novel antibacterial agents that (1) could kill microbial cells and (2) decrease the severity of disease course.

It is well-known that most antibiotics are able to alter bacterial metabolism and respiration. It was shown that bacteriostatic antibiotics decelerate cellular respiration whereas bactericidal antibiotics demonstrate an opposite action (Lobritz, 2015). Thus, chloramphenicol decreases the oxygen uptake during glucose oxidation (Duncan, 1962); quinolones (ciprofloxacin and pefloxacin) and β-lactams (meropenem) suppress the endogenous respiration in P. aeruginosa cells (Majtan, 1998).

Our previous results showed that derivatives of 4-[(1-adamantyl)]-phenol possess a strong potential to consider them antimicrobial and antibiofilm agents. Although the mode of action of the adamantane derivatives is largely unknown, our findings suggest that these compounds affect the structural integrity and functions of the bacterial cell membrane. We hypothesized that treatment of bacteria with adamantane derivatives alters oxidative metabolism and such changes may be reflected as violation of the energy generating pathways. The studies described in this paper were conducted with a view to provide the further information on the mode of antimicrobial action of 4-[(1-adamantyl)]-phenol derivatives.

MATERIALS AND METHODS

Bacteria and media
The strain Pseudomonas aeruginosa 2094 was isolated from a patient suffering from nosocomial infection. Mueller-Hinton broth (pH 7.2) (HiMedia) and Tryptic soy broth (pH 7.2) (HiMedia) were used for minimal inhibitory concentration (MIC) determination and for oxygen consumption experiments, respectively.

Materials
A derivative of 4-[(1-adamantyl)]-phenol (compound KVM-97) was synthesized in the Institute of Organic Chemistry of the National Academy of Sciences of Ukraine (Kyiv, Ukraine) by Yu. V. Korotkiy. KVM-97 was diluted in 10% dimethyl sulfoxide (DMSO); stock solution concentration was 1 g/L. All other chemicals were obtained from commercial sources.

Susceptibility to adamantane derivative
MIC was determined by the serial broth macrodilution method according to the EUCAST recommendations. The inoculum density was 10⁶ CFU/ml culture medium, 24 h culture was used for its preparation. The lowest concentration of KVM-97 that inhibited bacterial growth was considered the MIC.

Respiration measurement
Oxygen consumption was measured polarographically with a Clark-type oxygen electrode (Hansatech, UK). Bacteria cells were grown aerobically at 37°C for 20 h in broth. Resting cell suspension was centrifuged at 5000 g for 10 min, washed twice with a buffer solution of 18 mM KH₂PO₄, 125 mM KCl, 5 mM EDTA, 1 mM MgCl₂ (pH 7.4), and resuspended in the same solution. The cell suspension (0.8 ml) was stirred in a thermostated chamber at 35°C during the measurement. The oxygen concentration in suspension at 35°C was considered as 205 μM. In the experiments with substrates, either glutamate (3.0 mM) and malonate (2.0 mM), or succinate (3.0 mM) and rotenone (5.0 μM) were added with a microsyringe to the centre of the chamber. All reactions were stimulated by addition of ADP (0.2 mM). KVM-97 was added to achieve different concentrations: 0.5× MIC and 1.0× MIC. Oxygen consumption was monitored for approximately 7 min and the linear part of the oxygen uptake curve was used to determine the respiration rates. The oxygen consumption rate was expressed as nanomoles of oxygen consumed per
minute, corresponding to the total protein of cells. Protein concentration was determined by the method of Lowry et al. with bovine serum albumin as the standard (Lowry, 1951).

The ADP-stimulated respiration rate (state 3) was registered after addition of ADP to test suspension. The slower respiration rate, after complete ADP phosphorylation to ATP, was considered as state 4. The respiratory control index (RCI), defined as the ratio of the oxygen uptake rates in state 3 compared with state 4, was used for the evaluation of the uncoupling of respiration and oxidative phosphorylation (Manole, 2006).

Statistical analysis
The data were expressed as mean ± standard deviation (SD). The statistical significance of differences between the groups was determined by the non-parametric Kruskal–Wallis test. The number of samples per group (n) is noted in the figure legend. The statistical significance threshold of p < 0.05 was applied.

RESULTS

Effect of KVM-97 on the endogenous respiration
The MIC value of KVM-97 for P. aeruginosa strain of was determined as 50 mg/L. The alterations of the endogenous respiration of compound-treated P. aeruginosa cells are demonstrated in Table. The treatment of P. aeruginosa with KVM-97 at 1.0× MIC resulted in the most pronounced effect: the complete suppression of cellular respiration was observed. The respiratory activity after addition of KVM-97 at sub-inhibitory concentration 0.5× MIC decreased from 20.0 ± 1.1 to 16.0 ± 0.97 nmol O$_2$/min/mg protein (p < 0.05). Thus, the addition of KVM-97 to endogenous respiring P. aeruginosa cells caused a deceleration in oxygen consumption. The respiration rate in compound-treated cells decreased with increasing of compound in the dose-dependent manner.

Effect of KVM-97 on substrate respiration
The effect of KVM-97 on respiration was examined in the presence of either glutamate or succinate as respiratory substrates. Malonate and rotenone were used as specific inhibitors of succinate-oxidizing and NADH-oxidizing systems, respectively.

The data presented in Table shows that the endogenous respiration rate was slightly stimulated by addition of both substrates. In the presence of glutamate, the oxygen consumption rate was 25.2 ± 1.07 nmol O$_2$/min/mg protein, whereas succinate slightly increased the respiration rate to 27.6 ± 1.1 nmol O$_2$/min/mg protein (Fig. 1A, B).

The rate of glutamate oxidation was decreased to 18.0 ± 1.02 and 15.5 ± 0.96 nmol O$_2$/min/mg protein (p < 0.05) under action of KVM-97 at concentrations 0.5× and 1.0× MIC, respectively (Fig. 1A). The oxidation of succinate in the respiratory chain of P. aeruginosa was inhibited by KVM-97 in dose-dependent manner, the most pronounced decrease of oxygen consumption was observed in the presence of KVM-97 at concentration 1.0× MIC (Fig. 1B).

### Table. Oxygen consumption of the P. aeruginosa in the presence and absence of different substrates (n = 5)

| Experimental conditions              | Activity, nmol O$_2$ min$^{-1}$ mg$^{-1}$ |
|-------------------------------------|------------------------------------------|
|                                      | Control | KVM-97 |
|                                      |   | 0.5× MIC | 1.0× MIC |
| Without substrates (endogenous respiration) | 20.0 ± 1.1 | 16.0 ± 0.97 # | ND |
| Glutamate                           | 25.2 ± 1.07 | 18.0 ± 1.02 # | 15.5 ± 0.96 # |
| Succinate                           | 27.6 ± 1.1 | 25.0 ± 0.93 | ND |

Note: # p < 0.05 comparatively to control intact cells in each group, ND – respiration activity not determined, complete depression of respiration.
It was found that, at the highest concentration, KVM-97 did not cause complete depression of substrate respiration in the presence of glutamate, unlike both succinate oxidation and non-substrate endogenous respiration.

Effect of KVM-97 on the oxidative phosphorylation

Oxidative phosphorylation was assessed by state 3, state 4 respiration rates, and the RCI in compound-treated and intact cells.

It was observed (Fig. 1 A) that in the presence of glutamate, addition of the test-compound in both concentrations led to reduction of the respiration rates of state 3 and state 4 by 36.5–49.8% and 16.8–25.0%, respectively. Uncoupling of the respiratory chain was manifested in RCI decrease from 2.37 ± 0.19 to 1.45 ± 0.15 (p < 0.05) (Fig. 2 A).

In contrast, in the presence of succinate and KVM-97 (0.5× MIC), state 3 respiration rate decreased by 25% whereas state 4 respiration...
rate increased slightly (up to 11%) (Fig. 1B). The RCI value decreased from 2.65 ± 0.18 to 1.80 ± 0.30 (p < 0.05) (Fig. 2B). The higher KVM-97 concentration (1.0× MIC) completely inhibited *P. aeruginosa* respiration.

Therefore, KVM-97 depressed ADP-stimulated substrate respiration of *P. aeruginosa* in a dose-dependent manner. The inhibiting effect was more pronounced with succinate oxidation rather than glutamate oxidation. The RCI value in compound-treated cells was in average 1.5 times lower compared to the intact cells, indicating the uncoupling of the respiration and oxidative phosphorylation.

**DISCUSSION**

The decrease of the efficacy of antibiotic therapy and the spread of resistant pathogens constantly demand discovery and development of new antibiotics with novel mechanism of action. One of the potential targets for pharmacotherapy is energy generation process in microbial cells. *P. aeruginosa* aerobic respiratory chain is assembled in the cytoplasmic membrane. It has at least 17 primary dehydrogenases, ubiquinone and b- and c-type cytochromes, as well as five terminal oxidases (Matsushita, 1980; Williams, 2006). *Pseudomonas* cells can synthesize virulence factors (e.g., cyanide) in the process of energy generation in the respiratory chain. Apparently, the same metabolic pathways may be involved in the biofilm development whereas alginate biosynthesis, one of the main components of *P. aeruginosa* biofilm matrix, is an energy-dependent process. Moreover, numerous data suggest the major role of oxygen availability for the biofilm growth and metabolic activity.

Our previous results showed that derivatives of 4-(1-adamantyl)-phenol possess a strong potential to consider them as antimicrobial and antibiofilm agents. Adamantane derivative KVM-97 is lipophilic compound and rapid integration into the cellular membranes is likely to occur.

In this paper we presented the evidence that KVM-97 significantly depressed the endogenous respiration and succinate oxidation in *P. aeruginosa* cells. Compound treatment resulted in alteration of state 3 and state 4 respiration, and RCI as well, indicating the uncoupling of the respiration and oxidative phosphorylation.

Interestingly, the addition of KVM-97 at 1.0× MIC to *P. aeruginosa* cells did not cause complete inhibition of glutamate-dependent respiration, unlike the non-substrate endogenous respiration and succinate oxidation. We hypothesized that KVM-97 affect glutamate-dependent metabolic processes as glutamate excess prevent compound-induced inhibition of respiration.

It is well known that glutamate plays the central role in a wide range of metabolic processes in bacterial cells. This nonessential amino acid is involved in protein synthesis and other fundamental processes such as glycolysis, gluconeogenesis and the citric acid cycle. It is also a key metabolite, because it serves as a link between nitrogen and carbon metabolism (Berezov, 2004). Glutamate is important for the ammonia assimilation as well. Proteins or amino acids are implicated as the source of ammonia, the end product of endogenous respiration of *P. aeruginosa* (Warren, 1960).

Therefore, the antimicrobial and antibiofilm mechanism of action of the adamantane derivatives may be associated with a violation of the bacterial respiratory chain. Used in clinical practice, antibiotics may also alter bacterial cellular metabolism and respiration. Thus, chloramphenicol decreased the oxygen uptake in *P. aeruginosa* during glucose oxidation and appeared to inhibit the oxidation of α-ketoglutaric acid to glutamate (Duncan, 1962).

**CONCLUSIONS**

In the present study we demonstrated the alteration of oxygen consumption in *P. aeruginosa* cells caused by the 4-(1-adamantyl)-phenol derivative. The results suggest that KVM-97 inhibits the endogenous respiration and substrate oxidation and has an uncoupling effect in *P. aeruginosa* cells in dose-dependent manner. The mode of action of KVM-97 is related to the effect on glutamate-dependent metabolic processes, because glutamate excess prevented compound-induced inhibition of respiration in *P. aeruginosa*.
cells. Further studies are obviously required, which might provide valuable information about the role of adamantane derivatives in the disturbance of energy-dependent pathways.

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slopinimo koncentracijoje). Naudoti specifiniai kvėpavimo grandinės substratai – 3,00 mM glutamatas su 2,00 mM malonatu arba 3,00 mM sukcinatas su 5,0μM rotenonu. Visos reakcijos buvo sužadintos ADP (0,2 mmol). Tyrimo metu nustatyta, kad KVM-97 matuotomis koncentracijomis slopino endogeninį kvėpavimą ir substrato oksidaciją P. aeruginosa ląstele. Slopinantis poveikis priklausė nuo dozės dydžio ir buvo ryškesnis sukcinato nei glutamato oksidacijoje. Kvėpavimo kontrolės indeksas (RCI) junginiu paveiktose ląstelėse buvo 1,5 karto mažesnis nei nepaveiktose. RCI sumažėjimas yra susijęs su deguonies įsisavinimo pokyčiu 3 ir 4 būsenoje ir liudija kvėpavimo ir oksidacinio fosforilinimo blokavimą. Gauti duomenys rodo, kad 4-(1-adamantil)-fenolio junginiai slopina P. aeruginosa ląstelių deguonies su naudojimą ir pasižymi blokuojančiu poveikiu.

**Raktažodžiai:** Pseudomonas aeruginosa, adamantano junginys, antibakteriniai preparatai, kvėpavimas