Antimicrobial Activity of Pregnenolone in vitro

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Authors’ contributions

This work was carried out in collaboration among all authors. Author TPP designed the study, performed the statistical analysis, wrote the protocol and wrote the first draft of the manuscript. Authors II and TP managed the analyses of the study. Author GD managed the literature searches. All authors read and approved the final manuscript.

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ABSTRACT

The antimicrobial effect of pregnenolone on clinical and reference strains of Escherichia coli and Staphylococcus aureus was tested. Pregnenolone was found to have inhibitory activity against all tested bacterial strains. Slightly higher sensitivity is shown by the strains of S. aureus. When applied directly, pregnenolone has a weak antimicrobial effect due to its very low water solubility, as it is in oleose state because the formulation studied in this experiment contains tocopherol (a fat-soluble vitamin) as a co-solvent. When emulsified with lecithin, as well as with methanol, its solubility in water increases and penetrates over a longer distance in the agar around the points of its application. Applied as an emulsion, it shows significantly higher inhibitory activity against E. coli and S. aureus. Even the non-emulsified version should still be useful in vivo due to the fact that intracellular environments are much more lipophilic than serum, the target of most antimicrobial substances is the intracellular space, and non-emulsified pregnenolone has been shown to have very high intracellular uptake.

Keywords: Pregnenolone; antimicrobial activity; E. coli; S. aureus.

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1. INTRODUCTION

Pregnenolone is involved in the synthesis of all steroids by cholesterol, including corticosteroids (mainly cortisol and aldosterone), androgens and estrogens. Cholesterol is converted to pregnenolone in cellular mitochondria, from which progesterone or dehydroepiandrosterone (DHEA) is then synthesized. Progesterone can be transformed into cortisol, deoxytocicosterone or aldosterone, and DHEA is a precursor to androgenic and estrogenic steroids such as testosterone, dihydrotestosterone, estradiol, estrone, etc. [1]. There is evidence of antimicrobial activity of pregnenolone [2, 3]. One of the first direct experiments with pregnenolone was performed by Lamb et al. [2], who reported successful treatment with pregnenolone acetate of Nocardia asteroides, a pathogen with very high mortality, especially in immune-compromised patients such as with AIDS disease. The authors describe a case of extremely resistant N. asteroides infection in a child, and successful treatment with pregnenolone acetate (200mg-300mg daily doses orally) after traditional treatment with sulfonamides did not give the desired effects and the patient’s condition worsened. The authors also emphasize the absolute absence of side effects in the treatment with pregnenolone (acetate salt), which is of great importance given that all known interventions for the treatment of such highly resistant bacterial infections have pronounced side effects / toxic effects. In 1966 Yotis and Stanke [3] described a pronounced antibacterial effect of pregnane-derivative steroids such as pregnenolone and progesterone against a wide range of pathogens, including Staphylococci sp. Bandy et al. [4] reported an efficient and easy synthesis of 17-chalconoyl derivatives of pregnenolone and their antimicrobial activity against various microbial strains. Lone et al. [5] also synthesized oxides of steroidal chalcones and established their activity in vitro against various bacterial and fungal strains by the agar diffusion method. The activity of the tested compounds against individual microorganisms varies due to the structural differences between them. Pregnenolone has also been used as a template for the development of new anti-cancer compounds. Mohareb and Al-Omran [6] reported cytotoxic properties of newly synthesized heterocyclic steroids against three human tumor cell lines. In 2018 Chepkirui et al. [7] isolated five previously undescribed pregnenolone-type triterpenes 1–5 from a mycelial culture of basidiomycete Fomitiporia aethiopica fermented on rice and extracted with methanol. They have shown moderate cytotoxic effects against various human cancer cell lines, but have no significant antimicrobial activity.

Steroid-antibiotic conjugates have recently been developed as potential therapeutic agents for infectious diseases. Figueroa et al. [8] reported antibacterial activity of two pregnenolone derivatives (hemisuccinate-pregnenolone and ethylenediamine-hemisuccinate-pregnenololone) on S. aureus, E. coli and Klebsiella pneumoniae, with minimal inhibitory concentrations MIC = 0.25 - 1 mg / ml. Other authors [9] synthesized a conjugate of pregnenolone with carbamazepine and found that it exhibited a significant antimicrobial effect on Proteus mirabilis. Their data indicate that the conjugate pregnenolone-carbamazepine can be used as an antibiotic for the treatment of infections caused by P. mirabilis. In 2013 Kakati et al. [10] synthesized a new class of chalconoyl pregnenolones, some of which show significant inhibitory activity against certain species of bacteria (Bacillus subtilis and E. coli) as well as against filamentous and oval fungi (Aspergillus niger and Candida albicans).

Testing the antimicrobial potential of steroids such as pregnenolone and its derivatives is a new and promising approach in the search for effective remedies for infections caused by multidrug-resistant bacteria, which are a major problem in medical practice today. In this regard, the aim of the present work is to test the antibacterial activity of pregnenolone with tocopherols against Gram-negative (E. coli) and Gram-positive (S. aureus) pathogenic bacteria.

2. MATERIALS AND METHODS

Antimicrobial agent. The antimicrobial effect of Pregnenolone (Sigma-Aldrich) with tocopherols (Sigma-Aldrich) was tested. Soya Lecithin 1360 MG (SXENB1611201B, Solgar, UK, Ltd., Herts, HP 23, Inc., Leonia N. J.), as well as 70% methanol were used as an emulsifiers in some of the experiments.

Control. As a positive control, the broad-spectrum antibiotic thiamphenicol (Nikovet - Sofia) was used, to which the tested microorganisms did not show resistance.

Microorganisms. Pure cultures of 10 pathogenic strains were tested - five Gram-negative (Escherichia coli) and five - Gram-positive (Staphylococcus aureus). Two of them are
reference, obtained from the Bulgarian National Bank for Industrial Microorganisms and Cell Cultures (NBIMCC): *Escherichia coli* ATCC- 8739 (NBIMCC 3397) and *Staphylococcus aureus subsp. aureus* ATCC - 6538 (NBIMCC 3359).

The others are four strains of *E. coli* and four of *S. aureus* are isolated from inflammatory skin secretions from dogs in the laboratory of microbiology at the University Clinic in the Faculty of Veterinary Medicine at the University of Forestry in Sofia.

*Nutrient media.* Mueller Hinton agar and broth (BUL BIO NCIPD - Sofia) and Columbia blood agar (Biolab Zrt. H-1141, Budapest Ov. Utra 43) were used, as well as selective media: Endo agar (Antisel - Sharlau Chemie SA, Spain) for *E. coli* and Mannitol Salt agar for *S. aureus* (Biolab Zrt. H-1141, Budapest Ov. Utra 43).

The *cultivation of microorganisms* was carried out at 35-37° C for 24-48 hours under aerobic conditions.

The studies were performed by the *classical agar diffusion method* of Bauer et al. [11] and according to NCCLS [12,13]. Suspensions of test microorganisms were inoculated in exponential growth phase at a dose of 2,10^6 cells / ml in an amount of 0.1 ml in 9 cm diameter Petri dishes on Mueller-Hinton agar with a pH of 7.2 - 7.4 and a layer thickness 4 mm. Endo agar and Mannitol Salt agar for the respective bacteria were also used in some of the experiments. Pregnenolone (with and without emulsifier) and the control antibiotic were administered by instillation of 0.1 ml in 9-mm wells in the agar at concentrations of the active substances per well of 0.5 mg for pregnenolone and 30 μg for thiamphenicol, respectively. After incubation for diffusion for 3-4 hours at room temperature, the dishes were cultured at 35-37° C for 18-24 and 72 hours. The results were read by measuring the diameters of the inhibitor zones in millimeters, including the diameter of the well to the nearest 1 mm, with a transparent ruler on the outside of the bottom of the plates, with full growth inhibition being taken as the boundary zone. According to the three-stage Bauer-Kirby system, an inhibitory effect of pregnenolone to the tested bacteria was found depending on whether it was administered in pure form or in emulsion.

The data in Table 1 show that when testing the pure product without emulsification, the growth of *S. aureus* strains was inhibited by a minimum dose of 150 mg (inhibitory zone diameters averaged 12.0 ± 0.6 mm at this dose), while of those of *E. coli* - from 200 mg (diameters of the inhibitory zones on average 13.0 ± 0.9 mm). However, the differences in the mean diameters of the zones without growth between the strains of *S. aureus* and *E. coli* were not significant - P> 0.05, t-criteria of Student.

All microorganisms studied showed high sensitivity to thiamphenicol used as a positive control. The differences in the diameters of the inhibitory zones of all strains between the antibiotic and pregnenolone administered directly without emulsification were statistically significant (P <0.01, t-criteria of Student).

Significantly higher sensitivity to pregnenolone was shown by the studied Gram-negative and
Gram-positive strains when it is applied in an emulsion. The summarized results are presented in Tables 2 and 3.

The data show that in an emulsion in physiological saline with methanol (Table 2) pregnenolone exhibited a significantly higher inhibitory effect on the studied bacterial strains than the oily product.

The differences in the diameters of the inhibitory zones with pregnenolone tested without emulsification were significant (P > 0.05, t-criteria of Student). When administered at a dose of 200 mg, the inhibitory zones of the Gram-positive and Gram-negative strains tested were comparable to those of the control broad-spectrum antibiotic (P > 0.05, t-criteria of Student).

The results of the test of the inhibitory activity of pregnenolone were highest when it was emulsified in saline with lecithin and methanol (Table 3).

The differences with the results obtained in the other test variants were statistically significant (P < 0.001, t-criteria of Student). As can be seen from the aggregated data, when pregnenolone in this emulsion was administered at a dose of 112 mg, the inhibitory zones of the studied Gram-positive and Gram-negative strains were comparable to those of the control broad-spectrum antibiotic (P > 0.05, t-criteria of Student).

In all experimental variants used, S. aureus strains showed higher susceptibility to pregnenolone compared to those of E. coli, although the differences were not significant (P > 0.05, t-criteria of Student).

4. DISCUSSION

The results of the present studies show that pregnenolone exhibits inhibitory activity against both Gram-positive and Gram-negative bacteria. Of great practical importance is the high susceptibility of the tested E. coli strains - both clinical and control reference strains. Slightly higher susceptibility is shown by all studied strains of S. aureus, which is also of great importance for clinical practice. However, our results show that pregnenolone administered directly has a weak antimicrobial effect. This is due to its very low solubility in water, as it is in an oily state due to the content of tocopherol (fat-soluble vitamin). However, upon emulsification, its solubility in water increases and penetrates over a longer distance into the agar around its application points. The present results show that methanol is effective as an agent in this regard, but by the use of lecithin a better emulsion and higher water solubility of pregnenolone is achieved. However, when the conditions of the intracellular environment are considered (e.g. in vivo effects), the lipophilic nature of non-esterified pregnenolone is actually an advantage for its use as an anti-microbial intervention. The intracellular environment is much more lipophilic than serum, and most anti-microbial substances (e.g. antibiotics) exert their beneficial effects inside the cell. McManus et al. [14] have demonstrated that non-esterified steroids have very good intracellular uptake. In fact, in their experiment non-esterified pregnenolone had the highest cellular uptake than any other of the nine common steroids they tested, and it reached intracellular concentrations ~100 times higher than the concentrations in the culture media. Therefore, non-esterified formulations of pregnenolone such as the tested tocopherol formulation are likely to display strong anti-microbial effect in vivo comparable to the effects in vitro demonstrated in the present study.

Our results are in accordance with those of other authors. Figueroa et al. [8] found that the effect caused by pregnenolone derivatives may be due to the interaction with certain bacterial membrane factors that are specific for bacterial resistance. In this sense, the antibacterial activity of pregnenolone derivatives may depend on the nature of the functional groups involved in their chemical structure.

<table>
<thead>
<tr>
<th>Microorganisms</th>
<th>Inhibitory zones in mm</th>
<th>Thiamphenicol</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>50 mg</td>
<td>100 mg</td>
</tr>
<tr>
<td>Escherichia coli</td>
<td>10.0 ± 0.0</td>
<td>11.2 ± 0.8</td>
</tr>
<tr>
<td>Staphylococcus aureus</td>
<td>10.6 ± 0.5</td>
<td>11.4 ± 0.5</td>
</tr>
<tr>
<td>Total</td>
<td>10.3 ± 0.5</td>
<td>11.3 ± 0.6</td>
</tr>
</tbody>
</table>
affects its antibacterial activity. In order for it to

danazol

indicate that the degree of lipophilicity may affect

growth of the test microorganisms has been

inhibited in a dose

conjugate on

substance to read the antimicrobial ef

require higher water solubility of the examined

experiments, because the

microorganisms would be facilitated and

pregnenolone in direct contact with

hydrophobic region of the steroid to interact with

vitamin B1 requires the

lipophilicity of pregnenolone may not be an issue

and may even be an advantage considering its extremely high intracellular uptake by both active

and passive transport.

5. CONCLUSION

Pregnenolone exhibits inhibitory activity against

both Gram-positive (Staphylococcus aureus) and

Gram-negative bacteria (Escherichia coli).

The inhibitory activity of pregnenolone in the

oleic state (in combination with tocopherol) is

lower due to lack of water solubility. However, the

antimicrobial activity of pregnenolone is significantly increased in emulsion in saline and methanol. Even higher water solubility of

pregnenolone is achieved in an emulsion in saline with lecithin and methanol, which results in a significant increase in its antimicrobial properties in vitro in the agar-gel diffusion method. Pending future in-vivo experiments, we suspect that in living organisms even non-

esterified pregnenolone would display strong antibacterial effects due to its high intracellular uptake.

CONSENT

It is not applicable.

ETHICAL APPROVAL

It is not applicable.

Table 2. Antimicrobial effect of pregnenolone emulsified with methanol against Gram-positive and Gram-negative bacteria in the agar-gel diffusion method

<table>
<thead>
<tr>
<th>Microorganisms</th>
<th>Inhibitory zones in mm</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>50 mg</td>
</tr>
<tr>
<td>Escherichia coli</td>
<td>15.4 + 1.9</td>
</tr>
<tr>
<td>Staphylococcus aureus</td>
<td>14.8 + 1.7</td>
</tr>
<tr>
<td>Total</td>
<td>15.1 + 1.8</td>
</tr>
</tbody>
</table>

Table 3. Antimicrobial effect of pregnenolone emulsified with lecithin and methanol against Gram-positive and Gram-negative bacteria in the agar-gel diffusion method

<table>
<thead>
<tr>
<th>Microorganisms</th>
<th>Inhibitory zones in mm</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>14 mg</td>
</tr>
<tr>
<td>Escherichia coli</td>
<td>13.8 + 1.8</td>
</tr>
<tr>
<td>Staphylococcus aureus</td>
<td>14.4 + 1.0</td>
</tr>
<tr>
<td>Total</td>
<td>14.1 + 1.4</td>
</tr>
</tbody>
</table>

Kakati et al. [10] found that the presence of α, β-
unsaturated carbonyl moiety in the chalconoyl
pregnenolones synthesized by them is essential
for their antimicrobial activity. Prabpayak et al. [15] also reported antibacterial action of
pregnenolone compounds. Valverde et al. [16]
synthesized pregnenolone derivatives and found
antibacterial activity in vitro on S. aureus, K.
pneumoniae and E. coli. Growth of test
microorganisms has been inhibited in a dose-
dependent manner. These results suggest that
the quaternary amine group of hemisuccinate-
pregnenolone-vitamin B1 requires the
hydrophobic region of the steroid to interact with
certain components of the bacterial cell, thereby
violating bacterial growth and causing cell death.
These data indicate that in oily form the effect of
pregnenolone in direct contact with
microorganisms would be facilitated and
correspondingly higher than found in our
experiments, because the in vitro test methods
require higher water solubility of the examined
substance to read the antimicrobial effect.

Figueroa-Valverde et al. [9] also reported
antibacterial activity of the synthesized by them
pregnenolone-danazol-ethylenediamine
conjugate on S. aureus and Vibrio cholerae. The
growth of the test microorganisms has been
inhibited in a dose-dependent manner. Their data
indicate that the degree of lipophilicity may affect
the inhibitory activity of the pregnenolone-
danazol-ethylenediamine conjugate. Our results
also show that the lipophilicity of pregnenolone
affects its antibacterial activity. In order for it to
manifest in vitro, it is necessary to obtain a
water-soluble emulsion. However, in vivo, the
lipophilicity of pregnenolone may not be an issue
and may even be an advantage considering its extremely high intracellular uptake by both active
and passive transport.
COMPETING INTERESTS

Authors have declared that no competing interests exist.

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