Antimicrobial Activity of Minocycline against Bacteria and Fungi

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Abstract: Minocycline is a semi-synthetic tetracycline derivative and appears to be as effective as other tetracyclines and analogues. The present research work is on anti-microbial assay to demonstrate the minocycline is more effective on selected gram-positive than the gram-negative bacterial strains. It showed the highest zone of inhibition on Bacillus subtilis than Staphylococcus aureus when cultured in Mueller-Hinton agar medium.

Keywords: Antibacterial, antifungal and minocycline

1. Introduction

Minocycline has the spectrum of antibacterial activity in vitro¹. It has proven that the minocycline shows greater effect than the tetracycline and its analogues². Minocycline is indicated for the treatment of several diseases including acne vulgaris, urinary tract infections, and central nervous system³. In regard with the antibacterial activity of minocycline, Staphylococcus aureus has isolated from the aural discharge of purulent otitis was subjected against Benzylpenicillin, Streptomycin, Kanamycin, Erythromycin, Lincomycin, Oleandomycin, Spiramycin, Leucomycin and Chloramphenicol than all these antibiotics, minocycline was more active. As like any other tetracycline, the minocycline is found to be bacteriostatic against the susceptible organisms. The antibacterial activity in vitro is influenced by the inoculum’s size, pH of the culture medium and the presence of the serum⁴. Mueller-Hinton agar medium creates the optimum environment for the minocycline to be more active⁵. Therefore, in our present study we used Mueller-Hinton agar medium to test against the bacterial strains and a fungi. The minocycline was believed to have the highest activity against staphylococcus aureus but in our observation Bacillus subtilis had highest zone of inhibition than staphylococcus aureus.

2. Materials and Method

Test-pathogenic microorganisms
Two Gram-negative Klebsiella pneumonia, Pseudomonas aeruginosa, and two Gram-positive Bacillus subtilis, Staphylococcus aureus bacterial pathogens and one fungal pathogen Candida albicans were used in vitro antimicrobial activity. These selected pathogenic strains were obtained from Microbial Type Culture Collection (MTCC), Chandigarh, Punjab, India.

In vitro antimicrobial activity
The antibacterial activity was determined by well diffusion method⁶. About 25 ml of molten Mueller Hinton agar was poured into a sterile Petri plate (Himedia, Mumbai, India). The plates were allowed to solidify, after which 18 h grown (OD adjusted to 0.6) 100 µl of above said pathogenic bacteria cultures were transferred onto plate and made culture lawn by using sterile cotton swab. After five minutes setting of the pathogenic bacteria, a sterile cork borer was used to make 5 mm well on the agar. The test samples were dissolved in DMSO and loaded in to wells with various concentrations such as 25 µg/well, 50 µg/well, 75 µg/well and 100 µg/well. The Streptomycin added well served as positive control for bacteria and clotrimazole served as control for fungi. The solvent alone served as negative control. The plates were incubated at 37°C in a 40 W florescent light source (~ 400 nm) for 24 h. The antibacterial activity was determined by measuring the diameter of the zone of inhibition around the well using antibiotic zone scale (Himedia, Mumbai, India).

3. Result

Minocycline effectively killed all the test pathogens at all tested concentrations. The lowest concentration of 25 µg/well showed zone of inhibition against all the tested bacterial pathogens ranged between 12mm and 28mm whilst, the highest concentration of 100 µg/well showed zone of inhibition ranged between 19mm and 33 mm using Minocycline. Bacillus subtilis had the highest zone of inhibition with 33mm at 100µg/well than Staphylococcus aureus, which showed 31mm of zone of inhibition at 100µg/well. In case of Candida albicans 6mm zone of inhibition observed for the highest concentration of 100 µg/well.

Antimicrobial activity of Minocycline

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<thead>
<tr>
<th>Bacteria</th>
<th>Concentration per disc</th>
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<tbody>
<tr>
<td></td>
<td>25µg</td>
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<tr>
<td>Staphylococcus aureus</td>
<td>25</td>
</tr>
<tr>
<td>Bacillus subtilis</td>
<td>28</td>
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<tr>
<td>Klebsiella pneumonia</td>
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<tr>
<td>Pseudomonas aeruginosa</td>
<td>12</td>
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<tr>
<td>fungi</td>
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<tr>
<td>Candida albicans</td>
<td>10</td>
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4. Discussion

The laboratory studies have experimented the potentiality of Minocycline over the Tetracycline and its other analogues in inhibiting the growth of clinically isolated strains of Tetracycline resistant Staphylococci in many countries.
around the world. Minocycline showed little more effective in killing the β-haemolytic streptococci against Streptomyces faecalis (enterococci), the activity of Minocycline inclined to have more than that of the other Tetracycline. The greater activity was also observed against Streptomyces Viridians, were the Minocycline was very effective even at the low concentration. Tetracycline, Oxytetracycline and Demethylchlorotetracycline was less effective against D. pneumonia when compared with Minocycline, Doxycycline, Methacycline and Chlortetracycline. In our present study Minocycline was very effective on Bacillus subtilis, having the zone of inhibition of 33mm at the concentration of 100 μg/well and 28mm of zone of inhibition at 25 μg/well whereas Minocycline against, Staphillococcus aureus showed only 25mm and 31mm zone of inhibition at 25 μg/well and 100 μg/well respectively. On the other hand, Minocycline showed less activity on gram-negative when compared with gram-positive bacterial strains. Against, Escherchia. Coli and Haemophilus influenzae, Minocycline was less active than that of Methycline, chlorotetracycline or doxycycline. In our assessment, as per the observation on Pseudonas aeruginosa against Minocycline showed the zone inhibition with 12mm at lowest concentration of 25 μg/well and at highest concentration 100 μg/well showed 19mm zone of inhibition. Klebsiella pneumonia showed only 18mm zone of inhibition at 25 μg/well and 22mm zone of inhibition at 100 μg/well. The activity of Minocycline in vitro against the gram-negative species is essentially similar to that of other tetracycline analogues.

5. Conclusion

Minocycline, has long been established as the safe drug for treating human diseases. It is used as antibiotics and has the strong potential to treat multiple microbial infections. In vitro antimicrobial assay on minocycline reveals that it is effective over gram-positive than gram-negative strains.

References