Bacteriolyses of Bacterial Cell Walls by Zinc (II) Ions on the Basis of the Results of Halo Antibacterial Susceptibility Tests for Metallic Salt Solutions

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Abstract

Bacteriolyses of bacterial cell walls by zinc (II) ions on the basis of the results of halo antibacterial susceptibility tests were investigated for the nitrate and the sulfate solutions. From the results obtained by halo antibacterial tests of sulfate solutions against Staphylococcus epidermidis, the antibacterial order is Zn\(^{2+}\) > Cu\(^{2+}\) > Ag\(^{+}\) > Al\(^{3+}\), in which Zn\(^{2+}\) ions indicate the highest antibacterial effect. Bacteriolysis of S. aureus PGN cell wall by zinc ion is due to the inhibition of PGN elongation by the activation of PGN autolysins of amidases and side-chain endopeptidase. On the other hand, bacteriolysis of E. coli cell wall by zinc ions is attributed to the destruction of outer membrane structure due to degradative enzymes of lipoproteins at N- and C-terminals, and also is dependent on the activities of PGN hydrolases and autolysins of amidases and carboxypeptidase-transpeptidase. Zinc ions induced ROS such as \(\text{O}_2^-\), \(\text{H}_2\text{O}_2\), \(\cdot\text{OH}\), \(\text{OH}^-\) producing in bacterial cell wall occur oxidative stress.

Keywords: Halo antibacterial test; Zinc ions; PGN cell wall; Outer membrane lipoprotein; Biosynthesis and autolysin; Reactive oxygen species (ROS)

Introduction

Silver, copper, and zinc of transitional metals have highly antibacterial activities and are utilized as chemotheraphy agents. Recently, antibacterial activities of zinc and its complexes call attention to potential treatments such as prevention of serious diseases [1], exploitation during bacterial pathogenesis [2], and virus counter measure [3]. In this study, the halo antibacterial susceptibility tests of various nitrate and sulfate solutions against Staphylococcus epidermidis were carried out, wherein it was turned out that zinc ions had the highest antibacterial effects. On the basis of the high antibacterial activity for these zinc ions obtained from the halo antibacterial tests for metallic sulfate solutions, the processes of bacteriolyses and destructions by the antibacterial activities of zinc ion solutions were analyzed and considered against S. aureus peptidoglycon
and *E.coli* outer membrane cell walls. Furthermore, the bacteriolysic mechanisms by zinc (II) ions have been revealed against both Gram-positive and Gram-negative bacteria.

**Method**

**Halo Antibacterial Susceptibility Tests**

Halo antibacterial tests have been carried out for the nitrate and sulfate aqueous solutions against *Staphylococcus epidermidis* in the presence of human skin surface. The other, the antibacterial reagents were prepared metallic ions 100 mM/L aqueous solutions from metallic salt reagents. The metallic salts were used manufacturing by Wako Pure Reagents and Nissan Chemistry Ltd. The preparation method is shown in Table 1, wherein the crystalline powders of metallic salts of 0.01mol are dissolved in distilled water of 100cc, preparing metallic ion concentration of 100 mM/L as antibacterial reagents (crystalline powders of 0.005mol for silver sulfate and aluminum sulfate were used).

<table>
<thead>
<tr>
<th>Metallic salt reagents</th>
<th>Nitrate and Sulfate</th>
<th>Molecular weight</th>
<th>Mass(g) of metallic salt reagents dissolved in the distilled water of 100cc. Metallic ion conc. in water</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>Nitrate</strong></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Silver nitrate; AgNO₃</td>
<td>169.87</td>
<td>1.6987g</td>
<td>10,800mg/L</td>
</tr>
<tr>
<td>Copper nitrate; Cu(NO₃)₂·3H₂O</td>
<td>241.60</td>
<td>2.4160g</td>
<td>6,350</td>
</tr>
<tr>
<td>Zinc nitrate; Zn(NO₃)₂·6H₂O</td>
<td>297.47</td>
<td>2.9747g</td>
<td>6,540</td>
</tr>
<tr>
<td>Aluminum nitrate; Al(NO₃)₃·9H₂O</td>
<td>375.13</td>
<td>3.7513g</td>
<td>2,698</td>
</tr>
<tr>
<td>Lead nitrate; Pb(NO₃)₂</td>
<td>331.21</td>
<td>3.3121g</td>
<td>20,720</td>
</tr>
<tr>
<td><strong>Sulfate</strong></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Silver sulfate; Ag₂SO₄</td>
<td>311.8</td>
<td>1.5590g</td>
<td>10,800mg/L</td>
</tr>
<tr>
<td>Copper sulfate; CuSO₄·5H₂O</td>
<td>249.68</td>
<td>2.4968g</td>
<td>6,350</td>
</tr>
<tr>
<td>Zinc sulfate; ZnSO₄·7H₂O</td>
<td>287.56</td>
<td>2.8756g</td>
<td>6,540</td>
</tr>
<tr>
<td>Aluminum sulfate; Al₂(SO₄)₃·14~18H₂O</td>
<td>324.14</td>
<td>1.6207g</td>
<td>2,698mg/L</td>
</tr>
</tbody>
</table>

Table 1: Preparing metallic ions 100 mM/L aqueous solutions from metallic salt reagents.

Firstly, *Staphylococcus epidermidis* that were collected from the inside of the arms were incubated in physiological saline aqueous water solution of salt at a thermostat of constant temperature of 35℃ through a week. Afterwards the bacteria were incubated in the standard planar agar-medium, the appearing pepite colonies are incubated in gradient medium above a week. Secondly, the incubated cells were suspended in physiological saline solution, in which they were painted and swabbed at parts of 100μL to newly prepared planar medium. Finally, the paper discs that the antimicrobial liquid was swabbed and spread were placed on the center of planar medium at 35℃ for a week. Afterwards, measurement of the inhibition halo must be done always with ruler. Inhibitory zone width W is represented as W=(X−8 mm)/2 (in mm) from measured inhibitory diameter X and paper-disc diameter of 8 mm, in which W is calculated from measured X.

**Search for Peptidoglycan (PGN) Biosyntheses and Autolysins**

The molecular structure and the action sites for PGN biosyntheses and PGN autolysins were enzymatically searched against *S.aureus* and *E.coli*.

**Results**

**Halo Antibacterial Tests**

Figure 1 indicates the bar-graphs of the relationships between various metallic ions and halo inhibitory zones width. From these values of inhibitory zone width for the nitrate solutions, it is found that the antibacterial effect has nothing for alkali metals, alkali earth metals, but, various ions of Al³⁺, Zn²⁺, Pb²⁺, Cu²⁺, Ag⁺ indicate the antibacterial effects. The order of the antibacterial effect is as following: Cu²⁺>Zn²⁺>Ag⁺>Pb²⁺>Al³⁺. On the other
hand, in sulfate solutions, Al$^{3+}$, Zn$^{2+}$, Cu$^{2+}$, Ag$^+$ have higher antibacterial activities.

Figure 1: Relationship of various metallic ions of aluminum, zinc, lead, copper and silver nitrates and sulfates, and halo inhibitory zone (in mm) by halo antibacterial susceptibility tests against *Staphylococcus epidermidis*.

Figure 2 shows the sample surface appearances of inhibitory zone after halo antibacterial tests of sulfate solutions of ZnSO$_4$, CuSO$_4$, Ag$_2$SO$_4$, Al$_2$(SO$_4$)$_3$ against *Staphylococcus epidermidis*. From these observations, the antibacterial order is Zn$^{2+}$>Cu$^{2+}$>Ag$^+$>Al$^{3+}$, in which Zn$^{2+}$ ions indicate the highest antibacterial effect. It is thought that this effect results from strong Lewis acid of Zn$^{2+}$ ion and facility to bond with S-content cysteine.

Figure 2: Sample appearance after halo antibacterial susceptibility tests in the sulfates against *Staphylococcus epidermidis*.

**Search and Analysis**

**Action Sites of PGN Biosyntheses of Transglycosylase TG and Transpeptidase TP, and PGN Autolysins**

*S.aureus* surface cell envelop consists of teichoic acids, lipoteichoic acids, and thick PGN cell wall [4], whereas *E.coli* cell wall is comprised of lipid A, lipopolysaccharide (LPS), porin proteins, outer membrane of lipoprotein, and thinner 2-7 nm PGN layer in 30-70 nm periplasmic space [4]. Figure 3 shows the molecular structure of *S.aureus* PGN cell wall, including that indicates the action sites of PGN biosynthesis enzymes of TG/TP, and PGN forth autolysins and Lysostaphin enzyme.

Figure 3: The molecular structure of *S.aureus* PGN cell wall and the action sites of PGN biosynthesis TG/TP, PGN forth autolysins, and Lysostaphin enzyme.
Discussions

Bacteriolysis of S.aureus PGN Cell Wall by Zinc Ions

For the sake of growth of S.aureus PGN cell wall, there is necessarily required for the adequate balance between PGN syntheses and PGN autolysins. When the balance is broken, the bacteriolysis and the destruction of bacterial cell wall should become to occur. Wall teichoic acids are spatial regulators of PGN cross-linking biosynthesis TP [5], however, it is not explicit whether zinc ions could inhibit both TG and TP enzymes of PGN, whereas is due to uncertain relation between wall teichoic acids biosynthesis and PGN biosynthesis.

Zn\(^{2+}\) binding Rv3717 showed inactivity on polymerized PGN and however, it is induced to a potential role of N-Acetylmuramyl-L-alanine Amidase [6] in PGN fragment recycling. The novel zinc-induced cell wall hydrolases are effective for PGN murein hydrolase activity and generalized autolysis; PGN hydrolase Amidase MurA [7], Lytic Amidase LytA [8], enzymatically active domain of autolysin LytM [9], zinc-dependent metalloenzyme of active amidase domain AmiE [10] as prevention of the pathogen growth, Lysostaphin-like PGN hydrolase, and glycylglycine endopeptidase LytM [11]. It is thought that the activations of these PGN autolysins could be enhanced.
in the inhibition of PGN elongation, in inducing simultaneously bacteriolysis and destruction of *S. aureus* PGN cell wall. Accordingly, it has become apparent that Zn\(^{2+}\) ions inhibit the PGN elongation due to the activations of the PGN autolysins.

**Production of Reactive Oxygen Species (ROS) against *S. aureus***

For the penetration of zinc ions to PGN cell wall, the ROS production such as superoxide anion radical O\(_2^-\), hydroxyl radical ·OH, hydrogen peroxide H\(_2\)O\(_2\) occurred from superoxide radical O\(_2^-\) molecular [12]. O\(_2^-\) and H\(_2\)O\(_2\) permeate into membrane and cytoplasm, which DNA molecular is damaged by oxidative stress [13].

\[
\begin{align*}
O_2+H^+ + e^- & \rightarrow \cdot HO_2 \\
\cdot HO_2 & \rightarrow H^+ + O_2 \\
O_2^- + 2H^+ + e^- & \rightarrow H_2O_2 \\
H_2O_2 + e^- & \rightarrow HO^- + \cdot OH \\
\cdot OH + e^- + H^+ & \rightarrow H_2O
\end{align*}
\]

Thus, from above-mentioned discussion, a process of bacteriolysis of *S. aureus* PGN cell wall by Zn\(^{2+}\) ions is shown in Table 2.

<table>
<thead>
<tr>
<th>Zn(^{2+}) ions solution</th>
<th><em>S. aureus</em> Cell Wall</th>
</tr>
</thead>
<tbody>
<tr>
<td>Zn(^{2+})</td>
<td>Teichoic acid, Lipoteichoic acid, Peptidoglycan layer, Proteins</td>
</tr>
<tr>
<td>Zn(^{2+})</td>
<td>O(_2^-), H(_2)O(_2), ·OH, ·NO, ONOO(^-)</td>
</tr>
<tr>
<td>Zn(^{2+})</td>
<td>Teichoic acids are spatial regulators of biosynthesis of PGN cross-linking TP enzyme</td>
</tr>
<tr>
<td>Zn(^{2+})</td>
<td>It is unknown whether zinc ions inhibit the PGN biosynthesis TG/TP enzyme</td>
</tr>
<tr>
<td>Zn(^{2+})</td>
<td>Activations of PGN autolysins</td>
</tr>
<tr>
<td>Zn(^{2+})</td>
<td>Bacteriolysis and destruction of PGN cell wall due to inhibition of PGN elongations</td>
</tr>
<tr>
<td>Zn(^{2+})</td>
<td>ROS productions and the oxidative stress</td>
</tr>
</tbody>
</table>

Table 2: Bacteriolysis and destruction of *S. aureus* PGN cell wall by permeability and antibacterial activities of Zn\(^{2+}\) ions.

**Bacteriolysis and Destruction of *E. coli* Cell Wall by Zinc Ions Permeability of Zinc Ions into *E. coli* Cell Wall**

*E. coli* cell wall is constituted of the lipopolysaccharide (LPS), the lipoproteins (LP), and the thinner layer of PGN within the periplasmic space. The first permeability barrier of zinc ions in the *E. coli* cell wall is highly anionic LPS with hydrophobic lipid A, core polysaccharide, O-polysaccharide, in which zinc ions may be possible for the inhibition of LPS biosynthesis, owing to that promotes formation of metal-rich precipitates in a cell surface [14]. In zinc ion uptake across the outer membrane, the lipoproteins of OmpA, OmpC, OmpF porins have a role for at least some of these proteins in Zn\(^{2+}\) uptake, in which the lipoproteins have metallic cation selective and hydrophilic membrane crossing pore, to be effective for zinc transfer [15]. Zinc (II) ions react with -SH base, generating simultaneously H\(^+\). Zinc bivalent is unchangeable, forming as −SZn − S − bond 4-coordinated.

\[
Zn^{2+} + 2(-SH) \rightarrow (-)Zn (II) \rightarrow S + 2H^+
\]

** Destruction of Outer Membrane Structure in *E. coli* Cell Wall by Hydrolases of Lipoproteins at C-, N-Terminals**

ZnPT (zinc pyrithione) and Tol (Tol proteins)-Pal (Protein-associated lipo-protein) complex are widely used as antimicrobial agents, however, it has recently been demonstrated to be essential for bacterial survival and pathogenesis that the outer membrane structure may be destroyed [16,17].

**Inhibition of PGN Elongation due to the Damage of PGN Synthesis of Zinc-Protein Amidase in Periplasmic Space, and the Activations of PGN Autolysins**

The zinc-induced decrease of protein biosynthesis led to a partial disappearance of connexin-43 of protein synthesis in neurons [18], but it is poorly understood whether PGN biosynthesis is inhibited. Further, it is also unclear whether the both TG/TP should be inhibited by the zinc ions [19-21]. On the other hand, zinc ions were accumulated in *E. coli* periplasmic space, whereas the zinc
ions may be spent to the activation of bacteriolysis of the cell wall. Zinc depending PGN autolysins, amidase Peptidoglycan Recognition Proteins (PGRPs) [22], zinc metalloenzymes AmiD [23], amidase zinc-containing amidase; AmpD [24], zinc-present PGLYRPs [25] serve as a substitute for the PGN autolysins. It is particularly worth noting that the enhancement of the activities of autolysins is characterized on PGN carboxypeptidase-transpeptidase I1I2 [26] requiring divalent cations. Accordingly, the inhibition of PGN elongation has occurred by zinc ion induced activities of PGN hydrolases and autolysins.

**ROS Production and Oxidative Stress against E.coli**

In E.coli, free radicals (O2−, H+, OH−, ·OH) and H2O2 are formed as follows[27]:

\[
\begin{align*}
O_2 + e^- & \rightarrow O_2^- \\
2O_2^- + 2H^+ & \rightarrow H_2O_2 + O_2 \\
O_2^- + H_2O_2 & \rightarrow OH^- + \cdot OH + O_2^-
\end{align*}
\]

In the cell wall, reacting with polyunsaturated fatty acids:

\[
\begin{align*}
& \text{LH} + \text{OH} \cdot \rightarrow \text{L} \cdot + \text{HOH} \\
& \text{L} \cdot + \text{O}_2 \rightarrow \text{LOO} \cdot \\
& \text{LH} + \text{LOO} \cdot \rightarrow \text{L} \cdot + \text{LOOOH}
\end{align*}
\]

Zinc-containing PGRPs induce ROS production of H2O2, ·O2−, HO−, in which the ROS occur the oxidative stress and killing by stress damage [28].

As mentioned above, the bacteriolysis and the destruction of E.coli cell wall by the process of permeability and antibacterial activities of Zn2+ ions are summarized as in Table 3.

<table>
<thead>
<tr>
<th>Zn2+ ion solution</th>
<th>E.coli cell wall</th>
</tr>
</thead>
<tbody>
<tr>
<td>Lipopolysaccharide (LPS) Lipid A, Core polysaccharide</td>
<td>Outer Membrane Lipoprotein, Porins Omp F,A,C Periplasmic Space PGN layer</td>
</tr>
<tr>
<td>Zn2+, H+</td>
<td>Zn2+, O2−, H2O2</td>
</tr>
<tr>
<td>· Negative charge</td>
<td>· Porin proteins of hydrophilic channels</td>
</tr>
<tr>
<td>· Hydrophobic Lipid A</td>
<td>· Zn binding proteins</td>
</tr>
<tr>
<td>· Inhibition of LPS biosynthesis</td>
<td>· Destruction of outer membrane structure due to degradative hydrolases of lipoprotein at C- and N-terminals</td>
</tr>
<tr>
<td>· Zn2++Z(-SH)→-SZn−-S−+2H+</td>
<td>· LOO−, L·(Fatty acid)</td>
</tr>
</tbody>
</table>

**Table 3: Bacteriolysis and destruction of the E.coli cell wall by permeability and antibacterial activities of Zn2+ ions**

**Conclusion**

Halo-antibacterial susceptibility tests are carried out on the nitrate and the sulfate solutions of metallic ion concentration of 100 mM/L against Staphylococcus epidermidis. In the nitrate solutions, the order of bacterial effect is as follows: Cu²⁺ > Zn²⁺ > Ag⁺ > Pb²⁺ > Al³⁺. The other, in the sulfate solutions, the order is Zn²⁺ > Cu²⁺ > Ag⁺ > Al³⁺. The appearance of the highest antibacterial activity is found to be the zinc sulfate solution. Bacteriolysis of S.aureus PGN cell wall by Zn²⁺ ions, whereupon the wall teichoic acids control PGN biosynthesis cross-linking TP, is due to the inhibition of PGN elongation by the activation of PGN autolysins; amidase AmiA and AmiE, PGN hydrolase Lysostaphin-like endopeptidase (Glycine cutting). On the other hand, bacteriolysis and destruction of E.coli cell wall by Zn²⁺ ions are due to destruction of outer membrane structure by degradation of lipoprotein at C-,N-terminals and by inhibition of PGN formation due to the activations of PGN autolysins of amidases and carboxypeptidase-transpeptidase.

In zinc ion induced S.aureus cell wall, the productions of O2−, H+, H2O2, ONOO− occurs. The other, in E.coli cell wall, the productions of O2−, H+ in outer membrane and H2O2, OH− in periplasmic space occur. These ROS and Zn²⁺-H2O2 damage cell membrane proteins and DNA molecules by oxidase stress. Finally, the biomedical roles of this zinc ions having the highest antibacterial effect should be included as potential treatments for neurological diseases, brain-delivery, Parkinson’s and Alzheimer’s disease as well as schizophrenia, Wilson’s and Pick’s disease, amyotrophic lateral sclerosis, Down’s
syndrome, and multiple sclerosis [1]. It may become particularly important that the role of Zn²⁺ chelators in the brain is raised increasingly.

**Acknowledgements**

Halo antibacterial susceptibility tests were carried out on the Subject Research Group of Saitama Prefectural Kumagaya Girl’s High School, under the leader of Teacher Reiko KOBAYASHI. The author acknowledges Ms. KOBAYASHI and Subject Research Group.

**References**


