

## Article

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## Denaturation of Protein by Chlorine Dioxide: Oxidative Modification of Tryptophan and Tyrosine Residues<sup>†</sup>

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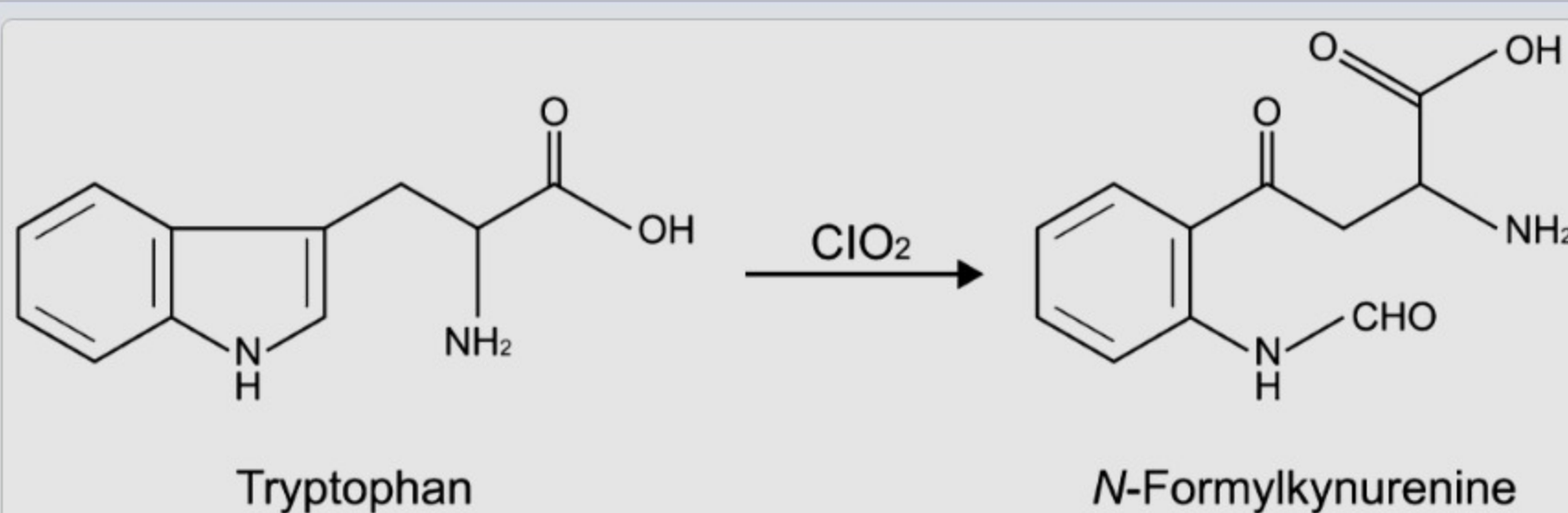
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## Abstract



Oxychlorine compounds, such as hypochlorous acid (HOCl) and chlorine dioxide ( $\text{ClO}_2$ ), have potent antimicrobial activity. Although the biochemical mechanism of the antimicrobial activity of HOCl has been extensively investigated, little is known about that of  $\text{ClO}_2$ . Using bovine serum albumin and glucose-6-phosphate dehydrogenase of *Saccharomyces cerevisiae* as model proteins, here I demonstrate that the antimicrobial activity of  $\text{ClO}_2$  is attributable primarily to its protein-denaturing activity. By solubility analysis, circular dichroism spectroscopy, differential scanning calorimetry, and measurement of enzymatic activity, I demonstrate that protein is rapidly denatured by  $\text{ClO}_2$  with a concomitant decrease in the concentration of  $\text{ClO}_2$  in the reaction mixture. Circular dichroism spectra of the  $\text{ClO}_2$ -treated proteins show a change in ellipticity at 220 nm, indicating a decrease in  $\alpha$ -helical content. Differential scanning calorimetry shows that transition temperature and endothermic transition enthalpy of heat-induced unfolding decrease in the  $\text{ClO}_2$ -treated protein. The enzymatic activity of glucose-6-phosphate dehydrogenase decreases to 10% within 15 s of treatment with 10  $\mu\text{M}$   $\text{ClO}_2$ . Elemental analyses show that oxygen, but not chlorine, atoms are incorporated in the  $\text{ClO}_2$ -treated protein, providing direct evidence that protein is oxidized by  $\text{ClO}_2$ . Furthermore, mass spectrometry and nuclear magnetic resonance spectroscopy show that tryptophan residues become *N*-formylkynurenine and tyrosine residues become 3,4-dihydroxyphenylalanine (DOPA) or 2,4,5-trihydroxyphenylalanine (TOPA) in the  $\text{ClO}_2$ -treated proteins. Taking these results together, I conclude that microbes are inactivated by  $\text{ClO}_2$  owing to denaturation of constituent proteins critical to their integrity and/or function, and that this denaturation is caused primarily by covalent oxidative modification of their tryptophan and tyrosine residues.

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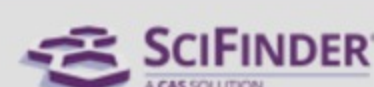


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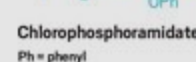
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