

Interrelationship between oxygen consumption, superoxide anion and hydrogen peroxide formation in phagocytosing guinea pig polymorphonuclear leucocytes.

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The paper presents an experimental procedure for a simultaneous assay of oxygen consumption, O₂⁻ release and H₂O₂ accumulation at a very early stage of the respiratory burst that is induced by phagocytosis in guinea pig polymorphonuclear leucocytes. The main findings are as follows: (a) The oxygen consumption that is measurable does not correspond to all oxygen that is reduced. The relationship between the actual oxygen consumed and the amount that is reduced depends on the fate of the intermediate products O₂⁻ and H₂O₂. (b) O₂⁻ is measurable extracellularly by the reduction of cytochrome c. When cytochrome c oxidizes the extracellular O₂⁻, molecular oxygen is formed. This fact is shown by a decrease of oxygen consumption. The molar ratio between the O₂⁻ detected and the oxygen given back is 1. (c) The amount of O₂⁻ released from the cells accounts for only a small part of oxygen actually reduced. (d) H₂O₂ is detectable only in the presence of NaN₃. In this condition almost all oxygen consumed is recovered in the form of H₂O₂. The molar ratio O₂/H₂O₂ is near unity. The amount of H₂O₂ derived from dismutation of O₂⁻ released is only an aliquot of the total H₂O₂ accumulated. Thus, most of H₂O₂ is derived from intracellular sources. (e) In the absence of inhibitors of H₂O₂ degrading reactions, no detectable accumulation of peroxide occurs. Under these conditions, the main part of H₂O₂ formed is degraded in almost equal amount by catalase and myeloperoxidase, while only a small aliquot is degraded by NaN₃ insensitive reactions.

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