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**Key Words**

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## Superoxide Dismutase Gene Expression in the Heart of Deoxycorticosterone Acetate-salt-induced Hypertensive Rats

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**ABSTRACT**

**Background.** Reactive oxygen species (ROS), such as superoxide anion radicals ( $\bullet\text{O}_2^-$ ), hydrogen peroxide ( $\text{H}_2\text{O}_2$ ), hydroxyl radicals ( $\bullet\text{OH}$ ), and others, may play critical roles in the pathogenesis of hypertension and other pathological processes. A previous study showed increased superoxide dismutase (SOD) gene expression in rat hearts of spontaneously hypertensive rats (i.e., congenital hypertension). SOD is a naturally existing antioxidant which plays an important role in scavenging oxygen free radicals.

**Aim.** To determine if changes in the SOD gene occur in the rat heart during a hypertensive state, the present study compared this gene in deoxycorticosterone acetate-salt (DOCA-salt)-induced hypertensive rats with normotensive Wistar-Kyoto rats (WKY).

**Methods.** We measured the mRNA and protein levels of 2 main subtypes of SOD (Mn-SOD and Cu, Zn-SOD) in hearts of 6, 9, and 12-week-old DOCA-salt-induced hypertensive rats and compared the values with those of age-matched normotensive WKY rats.

**Results.** By using Northern blot analysis of SOD-mRNA, DOCA-salt-induced hypertensive rats and normotensive WKY rats showed no statistical differences. The protein levels also showed similar changes.

**Conclusions.** This study shows that SOD gene expression is similar in acquired hypertensive (DOCA-salt) rats as well as in normotensive rats, which indicates that SOD gene expression is possibly not well correlated to changes in blood pressure in these rats with acquired hypertension.

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## INTRODUCTION

Reactive oxygen species (ROS), such as superoxide anion radicals ( $\bullet\text{O}_2^-$ ), hydrogen peroxide ( $\text{H}_2\text{O}_2$ ), hydroxyl radicals ( $\bullet\text{OH}$ ), and others, may play critical

roles in the pathogenesis of hypertension and other pathological processes.<sup>1</sup> It is well known that ROS can result in cell damage by reacting with various cellular constituents, including membrane lipids, proteins, and DNA. There is also evidence that ROS can influence

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