Manganese superoxide dismutase activity in the rat adrenal

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Abstract

In the light of studies suggesting that transcription of the gene coding for manganese superoxide dismutase (MnSOD) is induced by ACTH in the rat adrenal gland, northern blot analysis was used to determine its mRNA distribution. It was found that mRNA coding for MnSOD is primarily present in the inner zones of the rat adrenal cortex, and not the glomerulosa. To investigate the functional relationships between MnSOD activity and expression and adrenocortical function, adrenals and blood were taken from animals pretreated with corticotrophin or betamethasone (Betnesol), or subjected to a low-sodium diet. MnSOD activity in inner zone mitochondrial fractions was enhanced by corticotrophin and by a low-sodium diet, but suppressed by betamethasone. Apparent cytosolic MnSOD activity, total cytosolic MnSOD and CuZnMnSOD, and glomerulosa mitochondrial MnSOD all were unaffected. Steroid assays showed a clear correlation between circulating corticosterone and inner zone mitochondrial MnSOD, but none between aldosterone and glomerulosa MnSOD.

Immunoblot analysis of MnSOD showed two apparent isoforms, at approximately 25 kDa and 75 kDa. There was a partial relationship between expression of the 75 kDa isoform and MnSOD activity, in that it was induced by corticotrophin. However, there was also a slight induction with betamethasone, and a low-sodium diet had no effect. The 25 kDa MnSOD isoform was unaffected by the treatments. The results suggest that MnSOD may have a specific role in the steroidogenic function of the fasciculata/reticularis of the rat adrenal, but not in that of the glomerulosa.

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Introduction

The relationship between redox status and adrenocortical function has been the object of speculation since the discovery that ascorbic acid depletion is an early response to corticotrophin (ACTH) stimulation, and this has been used as a sensitive and highly specific bioassay for corticotrophin (Sayers et al. 1948, Chayen et al. 1976). Several attempts have been made to interpret this in functional terms (Kitabchi 1967, Hornsby et al. 1985, Yanagibashi et al. 1990). In this respect, the finding that manganese superoxide dismutase (MnSOD) gene transcription may be specifically enhanced by corticotrophin stimulation lends support to the concept of a physiological role (Suwa et al. 2000).

Superoxide dismutases (SODs) are antioxidants that protect the cell from cytotoxic reactive oxygen species (ROS; McCord & Fridovich 1969). ROS are generated as byproducts of metabolic oxidation, causing extensive cellular injury. In mammals, the cytosolic CuZnSOD and the mitochondrial MnSOD (Weisiger & Fridovich 1973a, b) are part of a cascade of components within the cell’s defensive machinery that inactivate these damaging radicals. It is possible that steroid-producing organs, such as the adrenal cortex, may require additional protection, as steroidogenesis is known to produce unwanted byproducts that may indeed be toxic. These include isocaproic aldehyde, derived from the cholesterol side chain discarded by the action of CYP11A, and free radicals that may be generated by ‘leaky’ P450s. It has been suggested that enzymes such as the aldose reductase-like AKR1B7 and MnSOD may be present in steroidogenic tissues for this reason (Sahut-Barnola et al. 2000, Suwa et al. 2000).

There may be alternative interpretations. Recent studies have also recognised that ROS may act as intracellular signalling molecules, suggesting that the SODs may have roles in cell signalling, with critical roles in cellular senescence, growth and apoptosis (Gamaley & Klyubin 1999).

Finally, Suwa and co-workers (2000) have proposed that ACTH induces an imbalance between MnSOD and glutathione peroxidase, resulting in the formation of excess hydrogen peroxide, which they postulate may contribute to the downregulation of CYP11B2 and aldosterone synthesis in the glomerulosa.
The present paper describes studies on the distribution and regulation of MnSOD expression in rat adrenal tissue, aimed at further interpretation of its function.

Materials and Methods

Animals and tissue preparation
Adult male Wistar rats 12–14 weeks old and weighing 180–220 g were obtained from commercial suppliers, and maintained briefly at Queen Mary, University of London under standard conditions of light and temperature, in accordance with appropriate guidelines for animal care. Rats were maintained on a diet of wholemeal flour (Sainsbury’s Ltd, UK) supplemented with 1% CaCO₃ and 1% NaCl, with access to distilled water for 3 weeks. The low-sodium diet omitted the 1% NaCl. ACTH-treated animals were injected subcutaneously with 100 µg Depot Synacthen (Ciba-Geigy) on 5 days, and rats treated with betamethasone (Betnesol, Glaxo-Wellcome) were given 25 ml distilled water supplemented with 0·05 g betamethasone daily for 7 days. For confirmation of the actions of the treatments, blood was collected for steroid assay. Rats were pretreated with heparin (1000 units/animal), then killed by stunning and cervical dislocation. Blood was collected from major vessels, centrifuged at 3000 g for 10 min, and plasma was frozen for storage. Adrenals were snap-frozen and stored in liquid nitrogen. When required, zona glomerulosa (with capsule) and zona fasciculata/reticularis (with medulla) fractions were separated under pressure between two glass plates.

Steroid assays
Plasma corticosterone and aldosterone concentrations were assayed using kits provided by Diagnostic Systems Laboratories Inc. (Webster, TX, USA).

Northern blot analysis
Total RNA was extracted from the zona glomerulosa and inner zones (with medulla) from rat adrenals as described before (Halder et al. 1998). RNAs (10 µg) were separated by electrophoresis and transferred on to a nylon membrane, Hybond-N+ (Amersham). Membranes were hybridised with [32P]-labelled MnSOD cDNA screened before (Halder et al. 1998). RNAs (10 µg) were separated by electrophoresis and transferred on to a nylon membrane, Hybond-N+ (Amersham). Membranes were hybridised with [32P]-labelled MnSOD cDNA screened and isolated from a rat adrenal cDNA library, the housekeeping gene glyceraldehyde-3-phosphate dehydrogenase (G3 PDH), and the zona glomerulosa and fasciculata/reticularis-specific markers, preadipocyte factor-1 (Pref-1) and inner zone antigen (IZ-Ag) (Halder et al. 1998, Raza et al. 2001), in 5 × SSPE containing 50% formamide, 0·5% (w/v) SDS, 0·1% (w/v) polyvinylpyrrolidone, 0·1% (w/v) Ficoll and 0·1% (w/v) BSA at 42 °C for 16 h. After hybridisation, the membranes were washed sequentially in 2 × SSPE/0·1% (w/v) SDS, 1 × SSPE/0·1% (w/v) SDS, and 0·2 × SSPE/0·1% (w/v) SDS at 60 °C, and then exposed to X-ray film (Amersham) at −80 °C with an intensifying screen.

Subcellular fractionation
For subcellular fractionation, adrenals were placed in 0·05 M potassium phosphate buffer (PBS) pH 7·8. Capsule/glomerulosa and inner zone fractions were separated and homogenised in PBS on ice. Samples were subjected to subcellular centrifugation, the nuclear pellet was removed at 800 g (15 min at 4 °C), the supernatant centrifuged at 12 000 g (15 min at 4 °C) and the mitochondrial pellet and the membrane/cytosolic supernatant fractions were retained. The mitochondrial pellet was resuspended in 100 µl 0·05 M PBS, pH 7·8. Protein content for both the mitochondrial and cytosolic/membrane fractions was determined by the method of Lowry et al. (1951). Both samples were divided into two aliquots, for total SOD activity and for MnSOD activity after inhibition of CuZnSOD.

SOD activity studies
The xanthine–xanthine oxidase nitroblue tetrazolium (NBT) assay (Spitz & Oberley 1989) provides an indirect measurement of SOD activity. This measurement of total SOD (CuZnSOD and MnSOD) and MnSOD activity alone is based on a modification of the method described by White et al. (1993).

Inhibition of CuZnSOD activity by diethyldithiocarbamate
The mitochondrial and membrane/cytosol aliquots were treated with 50 mM diethyldithiocarbamate (DDC) for 1 h at 30 °C, inactivating CuZnSOD activity. The samples were dialysed using three changes of 400 volumes of 50 mM PBS buffer pH 7·8 with 0·1 mM EDTA for 12–16 h at 4 °C. Dialysed samples were then assayed for MnSOD activity.

Spectrophotometric analysis of SOD activity
Assays for total and MnSOD activity in the different cell fractions from the treated animals were carried out in 1 ml aliquots with different amounts of protein, from 0 to 100 µg. The assay mixture contained the following reactants: 1 mM DTPA, 1 unit catalase, 5·6 × 10⁻⁵ M NBT, 10⁻⁴ M xanthine, 0·13 mg/ml BSA, 50 µM bathocuproinedisulphonic acid, 0·025% Triton X-100 and xanthine oxidase (an amount sufficient to achieve an absorbance rate change of 0·025 absorbance units/min at 560 nm). The assay depends on the inhibition of the xanthine oxidase by SOD activity in added tissue samples. Homogenates were assayed for total SOD activity
(CuZnSOD + MnSOD) and MnSOD activity alone (after inactivation of CuZnSOD with DDC).

Data were plotted as percentage xanthine oxidase inhibition against added tissue protein concentration, and 1 unit of activity was defined as the amount of protein required to decrease the reference rate by 50% of maximum inhibition. Results were expressed as units of activity per mg protein.

**SDS-PAGE gel electrophoresis**

Protein fractions were subjected to SDS-PAGE. A vertical slab gel electrophoresis unit (Biorad) incorporating a discontinuous (multiphasic) buffer system was used. The stacking gel (pH 6-8, Tris 125 mM, 4% (w/v) acrylamide/bisacrylamide, 0-1% (w/v) SDS) was polymerised on top of the resolving gel (pH 8-8, Tris 375 mM, 12% (w/v) acrylamide/bisacrylamide, 0-1% (w/v) SDS). Protein samples were loaded onto the stacking gel. Gel electrophoresis was run in a reservoir buffer (pH 8-3, Tris 25 mM, glycine 192 mM, SDS (0-1% (w/v)) at 35 V overnight and then increased to 200 V for 3-4 h at 4 °C. Prestained SDS-PAGE molecular weight standards (14-200 kDa; Amersham International plc) were run on each gel.

After electrophoresis, the gel was equilibrated in transfer buffer (pH 8-3, Tris 29 mM, glycine 150 mM and 20% (v/v) methanol) for 30 min. The separated proteins were electrotransferred to Hybond-enhanced chemiluminescence (ECL) nitrocellulose membrane (Amersham) in transfer buffer at 4 °C using a Transphor unit at 35 V overnight or 100 V for 2-3 h at 4 °C. Transfer to polyvinylidene difluoride membranes required that the membranes be rinsed in methanol for 1 min, then washed for 10 min in transfer buffer before electrotransfer as described above. Membranes were removed and rinsed in Tween buffer (150 mM NaCl, 10 mM Tris–HCl pH 7-6 and 0-05% Tween 20) twice for 15 min, removing excess glycine and methanol. Membranes were then subjected to immunoblotting.

**Immunoblotting**

Membranes with the transferred protein were blocked for 2 h in Tween buffer with 5% casein and 0-1% sodium azide at room temperature. Membranes were incubated with a rabbit anti-rat MnSOD polyclonal antibody (1:500; generous gift from Professor Naoyuki Taniguchi at Osaka University Medical School) in blocking buffer with 0-3% BSA overnight at 4 °C. Membranes were washed in Tween buffer three times for a total of 30 min before the addition of a secondary swine anti-rabbit horseradish-peroxidase-linked antibody (Amersham) diluted 1:1000 in Tween buffer with 1% casein for 1-5 h at room temperature. The secondary antibody was removed and the membrane washed in Tween buffer for 45 min; the solution was changed at 10-min intervals at room temperature. Luminescent bands were developed using the ECL (Amersham) reagents, and visualised using Hyperfilm-ECL (Amersham).

**Statistical analysis**

Student’s t-test with Bonferroni correction and Pearson’s method for calculation of correlation were used as appropriate.

**Results**

**Northern blot analysis of MnSOD mRNA**

RNA extracted from the zona glomerulosa and inner zones (with medulla) were hybridised with 32P-labelled cDNA probes for IZ-Ag, MnSOD, Pref-1 and the housekeeping gene, G3 PDH (Fig. 1). Using Pref-1 as a zona-glomerulosa-specific control (expression in the inner zones derives from the medulla, in which Pref-1 is also present) and IZ-Ag as inner zone marker, it was shown that MnSOD was localised to the inner zones (Fig. 1).

**Regulation of MnSOD in the rat adrenal**

The effects of ACTH pretreatment, low dietary sodium and betamethasone were studied, as these treatments have well recorded effects on both structure and function of the zona glomerulosa and inner zones (Vinson et al. 1992). Adrenal weights after treatment were (mean ± s.e.m.): controls 52.2 ± 5.5 mg, ACTH-treated 72.8 ± 3.6 mg (P<0.01 compared with controls, t-test with Bonferroni correction), low-sodium diet 65.2 ± 2.5 mg, betamethasone 43.5 ± 3.9 mg. The effectiveness of the treatments is best illustrated by the changes in circulating steroid concentration: ACTH significantly stimulated corticosterone, and aldosterone was increased in the animals fed a low-sodium diet. ACTH also marginally increased aldosterone concentrations, whereas both steroids were reduced by betamethasone (Fig. 2).

**MnSOD activity**

Values for mitochondrial MnSOD activity, compared with total SOD (not shown), confirmed that MnSOD in fact accounts for virtually all the SOD activity assayed in these preparations. In inner zone mitochondria, ACTH pretreatment produced a significant increase in MnSOD activity, and a smaller but still significant increase in inner zone mitochondrial MnSOD was also seen in the animals fed a low-sodium diet. Betamethasone had no significant effect (Fig. 3). Total MnSOD (MnSOD with CuZnSOD) in the cytosolic fraction was unaffected by
any of the treatments (Fig. 4). Data for glomerulosa fraction mitochondrial MnSOD showed that activity was low at all times, and not significantly affected by the treatments (Fig. 3).

**Immunoblotting analysis**

To relate activity with protein expression, immunoblotting analysis was performed on SDS gels of inner zone mitochondrial protein. In both glomerulosa and inner zone fractions, two bands were revealed, of approximately 25 kDa and 75 kDa (Fig. 6). The 25 kDa band was unaffected by any of the treatments, but the 75 kDa band changed in a manner that partly correlated with the activity studies, in that it was increased by ACTH pretreatment, but there was no change in the animals fed a low-sodium diet. Betamethasone also produced a modest increase in this component.

There was a clear correlation between circulating plasma corticosterone concentrations and inner zone MnSOD activity, but none between plasma aldosterone and glomerulosa fraction MnSOD (Fig. 5).
Discussion

The sensitivity and specificity of the adrenocortical response to corticotrophin and, in the present context, the exquisite tissue specificity of the redox response, has been reported over 50 years, and the present results extend this series of studies (Sayers et al. 1948, Kitabchi 1967, Chayen et al. 1976, Hornsby et al. 1985, Yanagibashi et al. 1990).

In demonstrating the corticotrophin-dependence of the transcription, expression and activity of MnSOD specifically in the zonae fasciculata/reticularis of the rat adrenal, the results presented here extend those of Suwa et al. (2000), who showed induction of MnSOD transcription by corticotrophin; similar findings in bovine adrenocortical cells were reported by Chinn et al. (2002).

There are several different possible interpretations of the significance of MnSOD distribution and activity in the rat adrenal. One is that the zonal specificity of MnSOD expression (Fig. 1) reflects the very much greater capacity for steroidogenesis, and P-450 mediated oxidation, in the fasciculata/reticularis. Accordingly, it could be that the induction of MnSOD is a purely protective device. With increased steroidogenesis, leakage of electrons from the adrenal mitochondrial electron transport chain may be enhanced, generating oxygen radicals (Hanukoglu & Rapoport 1995). Hence adrenocortical MnSODs may be critical in catalysing the dismutation of the ROS to hydrogen peroxide, which in turn would be detoxified to water by glutathione peroxidase or catalase. Others (Azhar et al. 1995) also have considered MnSOD to be related to steroidogenesis, although they assumed that its role was in cholesterol transport.

On the basis of the assumption that ROS are inherently toxic, Suwa et al. (2000) postulated that the known downregulation of aldosterone synthase brought about by extended corticotrophin treatment might be attributed to increased ROS generation in the glomerulosa. This is less plausible, however, when it is recalled that MnSOD is only one fraction of available SOD activity, and CuZnSOD and soluble SOD are expressed throughout the gland and appear to be unaffected by the treatments (Fig. 4). Moreover, although ACTH, and to a lesser extent betamethasone, stimulated expression of the 75 kDa MnSOD isoform in glomerulosa preparations, this was not associated with significant changes in MnSOD activity in the glomerulosa fraction (Fig. 3). Indeed, the evidence for MnSOD in this fraction could well be accounted for by the inevitable fasciculata contamination of this fraction. Finally, there was no correlation, either positive or negative, between plasma aldosterone and glomerulosa mitochondrial MnSOD activity (Fig. 5).

(a) **MnSOD activity/inner zones**

(b) **MnSOD activity /capsule, glomerulosa**
Yet another interpretation arises from the work of Okada et al. (1995), who have postulated a key role for peroxidase–ascorbate reaction in the generation of progesterone from pregnenolone. The hydrogen peroxide generated by MnSOD in the mitochondrial matrix is believed to pass through the mitochondrial membrane and may be used to oxidise ascorbic acid to ascorbyl radicals. These radicals may in turn oxidise NADH to NAD\(^+\), which in turn is used as a hydrogen acceptor, enabling the conversion of pregnenolone to progesterone by 3β-hydroxysteroid dehydrogenase/isomerase. It is possible that such a mechanism may account for the relationship between plasma corticosterone and MnSOD, but it does not suggest why aldosterone output should be so independent of glomerulosa fraction MnSOD, which in any case is never high (Figs 3–5). It is intriguing that Okada et al. (1995) have associated MnSOD with the tubulovesicular conformation of mitochondria characteristic of the zona fasciculata/reticularis in the rat, and we have argued that this morphology is associated, not only with the production of corticosterone, but also with the generally high level of steroidogenesis in these cells (Vinson 2004). All this suggests that MnSOD has a highly specific function in the regulation of fasciculata function and glucocorticoid secretion by corticotrophin. A role for MnSOD must be postulated that explains such specificity.

There remain two features of these findings that suggest more complex regulation. One is that a low-sodium diet, normally without effect on the adrenocortical inner zones, stimulates inner zone MnSOD activity (Fig. 3). Apart from anything else, this strongly suggests an integration between the functions of the glomerulosa, normally considered the part of the gland responsive to the sodium status, and the fasciculata, normally not regarded as sodium sensitive, although we have argued that it must be (Vinson et al. 1995). The second feature is that betamethasone treatment has a complex action in that, like ACTH, it enhances expression of the 75 kDa isoform (Fig. 6), although clearly suppressing plasma corticosterone, and, presumably, endogenous ACTH (Fig. 2). One interpretation here is that part of the action of ACTH may be indirect, as glucocorticoids are known to induce MnSOD in some tissues such as glomeruli (Kawamura et al. 1991) although they are inhibitory in many others, including hepatocytes, macrophages and intestinal epithelial cells (Valentine & Nick 1994). Hence betamethasone, although suppressing corticosterone, does not suppress MnSOD activity to the same extent (Fig. 3).

Taken together, the data strongly suggest a highly specific role for MnSOD activity in the inner zones of the rat adrenal. On balance, we think this is more likely to be related to aspects of steroidogenesis that are specific to the inner adrenocortical zones, fasciculata and reticularis. These links require further investigation.

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References


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Figure 6 (a) Immunoblotting analysis of immunoreactive MnSOD in mitochondria from rat adrenal inner zones. Example blots: two isoforms were seen, at approximately 25 kDa and 75 kDa. Densitometry: expression of the 75 kDa isoform was induced by ACTH and by betamethasone (Betameth.) (comparison with controls: *P<0·05, **P<0·01, t-test with Bonferroni correction, n=4), whereas the 25 kDa isoform was unaffected by the treatments. (b) Immunoblotting analysis of immunoreactive MnSOD in mitochondria from rat glomerulosa fraction. Example blots: as in the inner zone fractions, two isoforms were seen, at approximately 25 kDa and 75 kDa. Densitometry: expression of the 75 kDa isoform was induced by ACTH and by betamethasone (comparison with controls: *P<0·05, t-test with Bonferroni correction, n=4), whereas the 25 kDa isoform was unaffected by the treatments. Sod. Dep., low-sodium diet.


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