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Candesartan Differentially Regulates Epithelial Sodium Channel in Cortex Versus Medulla of Streptozotocin-Induced Diabetic Rats

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Abstract: Diabetes is associated with an activated renal renin-angiotensin-aldosterone system (RAAS) and it was shown that streptozotocin (STZ)-induced diabetic rats had increased whole kidney protein levels of the epithelial sodium channel subunits (α-, β- and γ-ENaC). However, the role of the RAAS on the regional, i.e., cortical versus medullary, regulation of ENaC is unclear. Male Sprague-Dawley rats were injected with STZ (intravenous, 65 mg/kg·bw, n=12/group). After 14 days, half of them received drinking water with candesartan (2 mg/kg·bw/day), an angiotensin-II type-1 receptor (AT1R) antagonist, for one week. In the medulla, i.e., inner stripe of the outer medulla (ISOM), base and/or tip of the inner medulla, immunoblotting revealed increased protein abundances of α1 Na-K-ATPase and ENaC subunits with diabetes (200-600% of controls), which were not reversed by candesartan. In fact, candesartan increased all ENaC subunits and α1 Na-K-ATPase in the ISOM and/or base in control rats. In contrast, in the cortex, diabetes did not increase these proteins. However, candesartan reduced cortical β- and γ-ENaC regardless of diabetic state. In summary, diabetes-induced increases in ENaC were seen preferentially in the medulla. These changes appeared to be due to a mechanism clearly distinct from AT1R activation, because they were not abolished by candesartan. In fact, candesartan treatment tended to increase some of these medullary proteins, perhaps in compensation for increased NaCl load. In contrast, cortical β- and γ-ENaC were reduced by candesartan regardless of diabetic state suggesting their regulation by AT1R at this site; however this did not appear to be a site of diabetes-induced ENaC up-regulation.

Keywords: Epithelial transporters, AT1R antagonist, angiotensin II, angiotensin receptor.

INTRODUCTION

Uncontrolled diabetes mellitus (DM) is associated with natriuresis and diuresis. The kidney adapts to sodium and water losses associated with DM *via* upregulation of both sodium and water transport proteins. In this regard, we have shown that streptozotocin (STZ)-induced type I diabetic rats have significantly increased abundances of renal epithelial sodium transporters and channels of the distal tubule, e.g., the thiazide-sensitive Na-Cl cotransporter (NCC) and the epithelial sodium channel (ENaC) subunits in whole kidney homogenates [1]. In addition, urea transporters, aquaporin 2 (AQP2), and the Na-K-2Cl cotransporter (NKCC2), major transport/channel proteins responsible for the production of concentrated urine, were found to be significantly increased in type 1 diabetic rats [2].

Diabetes is associated with increased activity of the renal renin-angiotensin-aldosterone system (RAAS). Activated RAAS is known to increase sodium reabsorption. Furthermore the increase in the levels of individual components of RAAS, such as angiotensin II and aldosterone, increase the protein expression of epithelial sodium transporters of the distal tubule in cell culture studies and also in non-diabetic animal models [3-6].

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Diabetic patients are commonly treated with angiotensin II type 1 receptor (AT1R) blockers to lower their blood pressure. Thus, the interaction between the diabetic state and AT1R activity as a determinant of renal epithelial sodium reabsorption is an increasingly relevant and timely area of study. Using candesartan, an AT1R blocker, we have recently demonstrated that elevated ENaC activity in type II diabetic obese Zucker rats is due to upregulated angiotensin type 1 receptor (AT1R) activity [7].

With regard to type I diabetic rats, we have recently shown that candesartan significantly increases levels of outer medullary proteins responsible for the production of concentrated urine in STZ-diabetic rats including AQP2 and NKCC2 [8]. However, a comprehensive regional analysis of epithelial sodium channel subunits and the effect of AT1R blockade in these type I diabetic rats has not been reported. Therefore the present study was designed to determine the regional (cortical versus medullary) regulation of epithelial sodium transporters of distal tubule such as ENaC and Na-K-ATPase by type 1 diabetes and the role of activated RAAS on the regional regulation of these proteins. For this, the protein abundances of ENaC subunits (-α, -β and -γ) and α1 subunit of Na-K-ATPase were determined in different regions of the kidneys i.e., cortex, outer medulla, inner-medullary base and inner-medullary tip in STZ-induced diabetic rats treated with or without candesartan. In addition, the protein abundance of NCC was also determined in the cortex of these rats.

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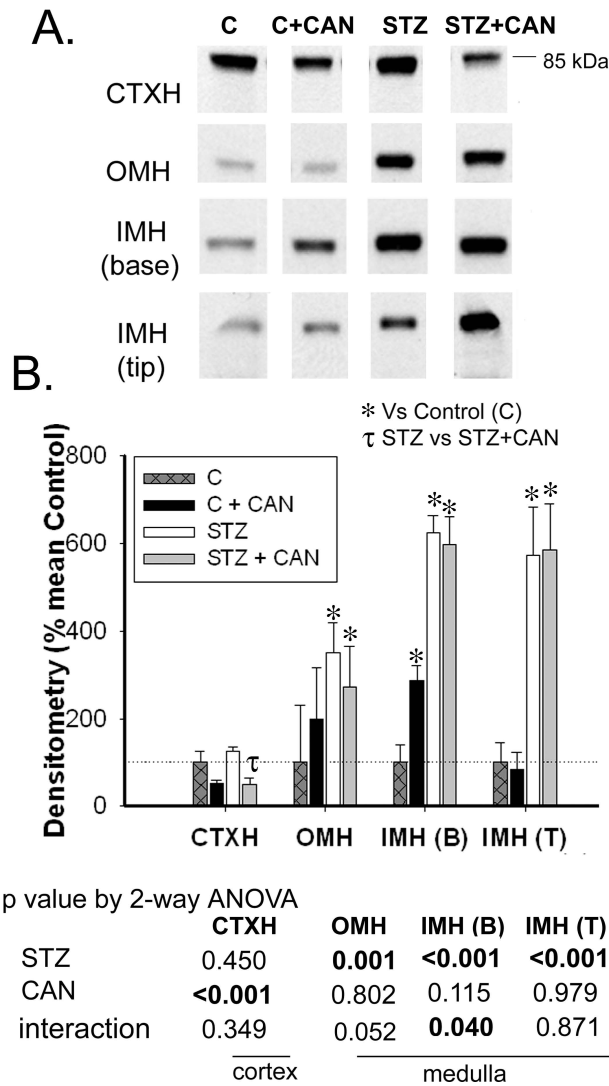


p value by 2-way ANOVA

	CTXH	OMH	IMH (B)	IMH (T)
STZ	0.450	**0.001**	**<0.001**	**<0.001**
CAN	**<0.001**	0.802	0.115	0.979
interaction	0.349	0.052	**0.040**	0.871
	cortex	medulla		

Fig. (3). Effect of diabetes and candesartan on 85kDa band of γ-ENaC protein abundance in different kidney regions. **A**) Representative lanes from immunoblots for the γ-ENaC subunit in the cortex homogenate (CTXH), inner stripe of outer medulla homogenate (OMH), inner-medullary base homogenate (IMH (B)) and inner medullary tip homogenate (IMH (T)) from non-diabetic (C) and diabetic (STZ) rats treated with or without candesartan (CAN). **B**) Densitometric summary of the blots (n = 6/group). For immunoblotting, each lane was loaded with an equal amount of total protein from a different rat sample (confirmed *a priori* by equal Coomassie-staining of representative protein bands). * indicates mean is significantly different (p<0.05) from that of the untreated non-diabetic (C) and τ from untreated diabetic rats (STZ) by unpaired t-test. Results from two-way ANOVA (STZ X CAN) are given below the graph. Significant (p<0.05) factors are in **BOLD**.

nificant change was observed in the CTXH or in the IMH (T) (Fig. **5**).

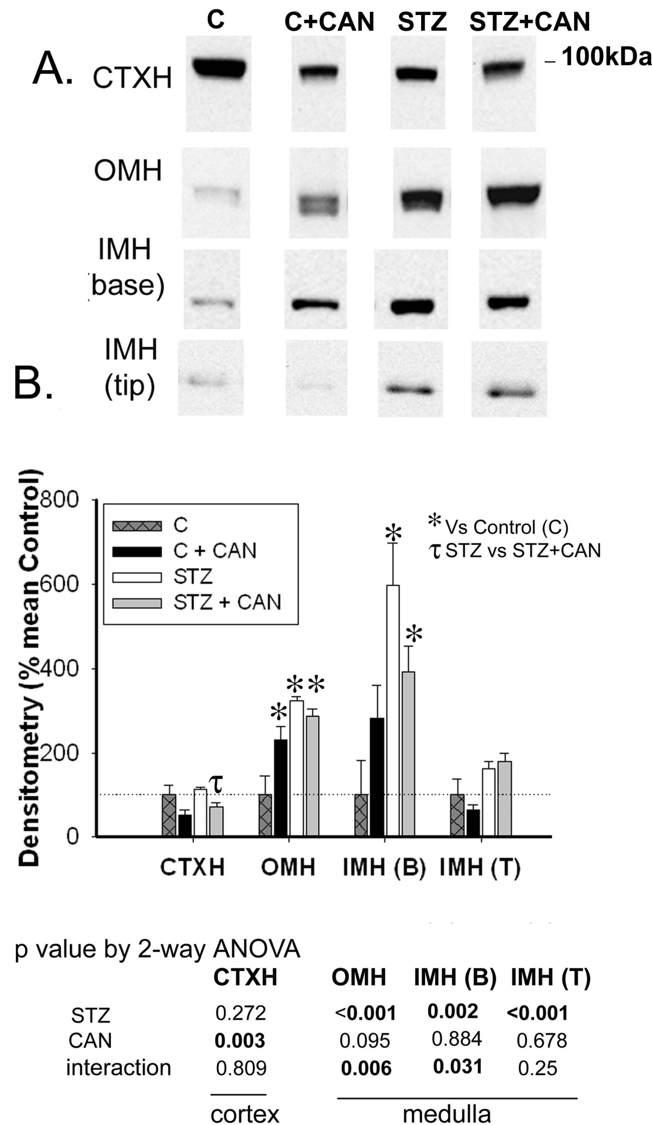
Regulation by Candesartan

Treatment with candesartan significantly decreased the protein abundance of α1 subunit of Na-K-ATPase in the CTXH of diabetic rats. Similarly, for non-diabetic rats it trended in same direction (p=0.07). Candesartan led to a significant increase in OMH α1 Na-K-ATPase with a trend for an increase in the IMH (B), p = 0.015, in non-diabetic rats (Fig. **5**).

Fig. (4). Effect of diabetes and candesartan on NCC protein abundance in different kidney regions. **A**) Lane from representative immunoblots for the NCC in the cortex homogenate (CTXH), inner stripe of outer medulla homogenate (OMH), inner medullary base homogenate (IMH (B)) and inner-medullary tip homogenate (IMH (T)) from non-diabetic (C) and diabetic (STZ) rats treated with or without candesartan (CAN). **B**) Densitometric summary of the blots (n = 6/group). For immunoblotting, each lane was loaded with an equal amount of total protein from a different rat sample (confirmed *a priori* by equal Coomassie-staining of representative protein bands). * indicates mean is significantly different (p<0.05) from that of the untreated non-diabetic (C) and τ from untreated diabetic rats (STZ) by unpaired t-test. Results from two-way ANOVA (STZ X CAN) are given below the graph. Significant (p<0.05) factors are in **BOLD**.

DISCUSSION

In this study, we evaluated the regulation of major renal epithelial sodium transport-related proteins including ENaC subunits, NCC, and Na-K-ATPase in the type I diabetic rat. We further determined whether blockade of the AT1R with candesartan would abolish changes that resulted from STZ-induced diabetes. We demonstrated that there are striking differences in the regulation of ENaC subunit protein abundances in cortex versus medullary regions. Diabetes increased ENaC subunit expression in the medulla, but not in the cortex. Candesartan did not reduce this medullary upregulation; and in contrast it resulted in an increase in β- and γ-ENaC in the non-diabetic control rats. The thiazide-sensitive Na-Cl cotransporter, NCC, expressed in the cortex,



p value by 2-way ANOVA

	CTXH	OMH	IMH (B)	IMH (T)
STZ	0.272	**<0.001**	**0.002**	**<0.001**
CAN	**0.003**	0.095	0.884	0.678
interaction	0.809	**0.006**	**0.031**	0.25
	cortex	medulla		

Fig. (5). Effect of diabetes and candesartan on protein abundance for α1 subunit of Na-K-ATPase in different kidney regions. **A**) Representative lanes from immunoblots for the Na-K-ATPase in the cortex homogenate (CTXH), inner stripe of outer medulla homogenate (OMH), inner-medullary base homogenate (IMH (B)) and inner medullary tip homogenate (IMH (T)) from non-diabetic (C) and diabetic (STZ) rats treated with or without candesartan (CAN). **B**) Densitometric summary of the blots (n = 6/group). For immunoblotting, each lane was loaded with an equal amount of total protein from a different rat sample (confirmed *a priori* by equal Coomassie-staining of representative protein bands). * indicates a mean is significantly different (p<0.05) from that of the untreated non-diabetic (C) and τ from untreated diabetic rats (STZ) by unpaired t-test. Results from two-way ANOVA (STZ X CAN) are given below the graph. Significant (p<0.05) factors are in **BOLD**.

was not statistically increased by diabetes likely due to high variability; however the mean was increased by 78%. Candesartan decreased NCC in both diabetic and non-diabetic rats. Finally, the α1 subunit of Na-K-ATPase was increased by diabetes most strongly in regions expressing thick ascending limb i.e., OMH or the early portion of the IMCD (base). These regions would be expected to have high energy demands and might be expected to be the most severely affected by diabetes-induced oxidative stress [17, 18]. Candesartan did not reverse this upregulation.

ENaC

The increase in ENaC subunits in diabetic rats is in agreement with results in whole kidney homogenate from our previous studies in type I and type II diabetes [1, 7]. In the present study, in addition to cortex and outer medulla, a comprehensive study of the regulation of ENaC was also done in the inner medullary tip (predominantly terminal IMCDs) and base (predominantly initial IMCDs), which was not examined previously. We found that all three subunits of ENaC were significantly increased by diabetes in both base and tip of the inner medulla but that candesartan regulated these subunits differentially in these two segments.

Our finding of differential regional regulation of sodium transporters by type I diabetes is in agreement with our previous finding from a long term study (12 week) of ovariectomized rats (supplemented with estrogen) with STZ-induced diabetes [19]. Regulation of β- and γ-ENaC found in that study was similar to the results demonstrated here (however, inner medulla was not subdivided into base and tip in the previous study). In contrast to our present finding, α–ENaC was found to be decreased in cortex and was not affected in the medulla in our previous long term study [19]. Furthermore, a recent study of long term (8 weeks) diabetic rats by Vidotti *et al.* [20], also showed reduced expression of α-ENaC relative to non-diabetic control rats. The reason for this disparity is not clear; however, the changes in protein abundances found after long term diabetes [20, 28], could be a compensatory response rather than an effect of diabetes found early on in diabetes. Reduced α-ENaC protein levels found in these earlier studies [19, 20] could also be due to the nephropathy or renal hypertrophy associated with long term diabetes [21].

We found that candesartan treatment did not attenuate the elevated levels ENaC subunits in the medulla of diabetic rats. Thus upregulated AT1R does not appear to have a role in the increased ENaC abundance that is observed in diabetes. We speculate that increased ENaC subunit protein levels in diabetes could be due to increased presence of vasopressin (AVP), since diabetes is associated with elevated AVP levels [22, 23]. In this regard, AVP has been shown to increase ENaC subunits protein abundances [24]. However, why diabetes increased ENaC subunits only in medulla and not in cortex is not clear. We speculate that the lower number of vasopressin receptors present in the cortex versus medulla could be one of the reasons [25, 26].

Like diabetes, candesartan also appeared to have a differential effect in medulla vs cortex of both diabetic and non diabetic rats. Candesartan did not affect ENaC subunits in the medulla whereas in cortex, candesartan reduced the abundance of ENaC subunits in both diabetic as well as non-diabetic rats. One plausible explanation for the blunted or absent effects of candesartan on medullary ENaC subunits in diabetic rats could be a compensatory response, as these proteins may be required to reduce the loss of electrolytes in the diabetic condition. Moreover, the levels of ENaC subunits were markedly higher in the medulla of diabetic rats to begin

with. The downregulatory effect of candesartan observed in cortex is in agreement with our recent study in lean Zucker rats, where we demonstrated that candesartan was able to significantly reduce whole kidney abundances of β and γ-ENaC in non-diabetic lean rats [7]. In this regard, angiotensin II has been shown to directly increase the activity as well as abundance of ENaC subunits [3, 4]. However, Beutler *et al*. [3] found that in response to candesartan infusion, the expression levels of β and γ-ENaC were significantly increased in the cortex region of the rats with NaCl restriction [3]. One possible explanation for this discrepancy could be the use of NaCl restricted rats, a model of high plasma renin activity, in their study. In addition, these rats were exposed to candesartan for a relatively shorter time compared to our study, i.e., two days versus 7 days.

In non-diabetic rats, candesartan increased ENaC protein levels in the medulla (unlike cortex). A comprehensive analysis of medullary regions revealed that this regulation was mostly in outer medulla and initial IMCDs (inner medullary base). However no such change was observed in terminal IMCDs (inner medullary tip). We speculate that this could be a compensatory effect rather than a direct regulation by candesartan. It may be possible that a candesartan-induced reduction of ENaC in the cortex could increase the sodium load to the medulla, thereby prompting an increased ENaC abundance in the medullary regions, at least in vehicle-treated non-diabetic rats. However, medullas of the diabetic rats had higher abundance of these proteins to begin with thus limiting the additional candesartan effect and resembling a blunted effect of candesartan. In addition, in early diabetes (19-20 days from the onset of DM) AT1R mRNA has been shown to be increased only in cortex, which could also be the cause for a blunted candesartan effect in the medulla of diabetic rats [27]. Candesartan did not affect blood pressure in these rats as observed by tail cuff method [8] thus change in blood pressure can be ruled out as a factor for this regulation.

NCC

In our previous study we showed that induction of type I diabetes significantly increases the abundance of whole kidney NCC [1]. This was confirmed in the present study that further examined the effects of candesartan and showed that candesartan reversed the elevation of NCC suggesting a role for upregulated AT1R in the elevation of NCC in type 1 diabetes. In addition, the significant decrease in NCC levels in AT1R knockout mice demonstrated by Brooks *et al.* [28] further suggests a direct role of angiotensin II in the upregulation of NCC. Although our recent study in non-diabetic lean Zucker rats also demonstrated that candesartan was able to significantly reduce NCC [7], long term STZ-induced diabetes did not seem to affect NCC abundance [20]. The reasons for this disparity could be similar to what has been discussed above for ENaC protein.

Na-K-ATPase

In diabetic rats, the α1 subunit of Na-K-ATPase was increased in the medullary region (OM and IM) while no significant change was found in the cortex. Khadauri *et al.* [29] has shown that STZ-induced diabetes increased the activity of Na-K-ATPase activity in the proximal convoluted tubule and medullary thick ascending limb, and cortical and outer medullary collecting tubule, but not in the proximal straight tubule, cortical thick ascending limb or distal convoluted tubule. Moreover, medullary regions would be expected to have high energy demands and are the most severely affected by diabetes-induced oxidative stress [17]. Similar to ENaC protein, we have demonstrated that candesartan differentially regulate Na-K-ATPase protein also in cortex vs medulla in these rats.

We have recently published that the STZ-diabetic rats have elevated levels of urea transporters (UT-A1, UT-A3), AQP2, and NKCC2/BSC1 proteins, which were further increased (UT-A3 and NKCC2/BSC1) or remained unaltered (UT-A1 and AQP2) by candesartan treatment [8]. Thus we suggest that early type I diabetes is associated with a compensatory elevation in the levels of medullary proteins to avoid urinary loss of sodium and water.

In summary, we found that medullary ENaC and Na-K-ATPase proteins were significantly increased in type I diabetes, which is not due to upregulated AT1R activity. We suggest that elevated AVP associated with diabetes may contribute to this upregulation. Unlike ENaC and Na-K-ATPase, increased NCC in type 1 diabetic rats appears to be associated with upregulated AT1R activity. We suggest that the regulation of ENaC and Na-K-ATPase proteins may be a compensatory effect rather than a direct effect of candesartan, at least in the medulla. Thus, while candesartan reduces ENaC and Na-K-ATPase proteins in the cortex, the medulla escapes these effects in the diabetic state, perhaps to avoid excessive sodium loss.

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