Similarities and differences of X and Y chromosome homologous genes, SRY and SOX3, in regulating the renin-angiotensin system promoters.

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Abbreviated title: SOX3 and SRY regulate the renin-angiotensin system

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ABSTRACT

The renin-angiotensin system (RAS) is subject to sex-specific modulation by hormones and gene products. However, sex-differences in the balance between the vasoconstrictor/proliferative ACE/Ang II/AT1 axis, and the vasodilator/anti-proliferative ACE2/Ang-(1-7)/MAS axis are poorly known. Data in the rat have suggested the male specific Y-chromosome gene Sry to contribute to balance between these two axes, but why the testis determining gene has these functions remains unknown. A combination of in silico genetic/protein comparisons, functional luciferase assays for promoters of the human RAS, and RNAseq profiling in rat were used to address if regulation of Sry on the RAS is conserved in the homologous X-chromosome gene, Sox3. Both SRY and SOX3 up-regulated the promoter of Angiotensinogen (AGT), and down-regulated the promoters of ACE2, AT2, and MAS, likely through overlapping mechanisms. The regulation by both SRY and SOX3 on the MAS promoter indicates a cis regulation through multiple SOX binding sites. The Renin (REN) promoter is up-regulated by SRY and down-regulated by SOX3 likely through trans and cis mechanisms respectively. Sry transcripts are found in all analyzed male rat tissues including the kidney, while Sox3 transcripts are found only in the brain and testis, suggesting that the primary tissue for renin production (kidney) can only be regulated by SRY and not SOX3. These results suggest that SRY regulation of the RAS is partially shared with its X-chromosome homolog SOX3, but SRY gained a sex-specific control in the kidney for the rate limiting step of the RAS, potentially resulting in male specific blood pressure regulation.

Keywords

renin angiotensin system, hypertension, Sox3, Sry, sex differences

Abbreviations: RAS, renin-angiotensin system; AGT, Angiotensinogen; REN, Renin; ACE, carboxypeptidase angiotensin-I-converting enzyme; Ang II, angiotensin II; AT1, type-1 angiotensin II receptor; ACE2, angiotensin-I-converting enzyme 2; Ang-(1-7), angiotensin-(1-7);
HMG, homologous high mobility group; dN, rate of nonsynonymous; dS, rate of synonymous;
ECR, Evolutionary conserved region; CHO, Chinese hamster ovary; SEM, standard error of the
mean.

Introduction

The renin-angiotensin system (RAS) is one of the most studied hormonal systems in the
physiological regulation of blood pressure through renal and non-renal mechanisms (16).
Dysregulation of the RAS is considered a major factor in the development of cardiovascular
pathologies. It is generally subdivided into two major axes that act to maintain homeostasis, with
each tissue/system having a unique combination of proteins and peptides (9). There is the: 1)
vasoconstrictor/proliferative axis of the RAS consisting of the carboxypeptidase angiotensin-I-
converting enzyme (ACE), angiotensin II (Ang II), the type-1 angiotensin II receptor (AT1); and
2) vasodilator/anti-proliferative axis consisting of angiotensin-I-converting enzyme 2 (ACE2),
angiotensin-(1-7) (Ang-(1-7)), the MAS receptor. It is now generally accepted that both axes
participate in the regulation of blood pressure and metabolism, and, even more importantly, in
the pathogenesis of cardiovascular, renal, and metabolic diseases (58). Several reports have
shown differences between components such as plasma renin activity and levels of
angiotensinogen in human, mouse, and rat models between males and females as previously
reviewed in the literature (55). However, few studies have been performed to observe and
analyze the effects of sex differences in regulation and balance of the two axes of the RAS (26).

Y-chromosome consomic rats first suggested a contribution of the male specific
chromosome to elevate blood pressure (21), with crosses of the consomic rats to the testicular
feminized rats suggesting an interplay between genes on the Y-chromosome and androgen
signaling (20). Dissection of the Y-chromosome elucidated the Sry gene as a candidate to
elevate blood pressure in the rat (22, 65). Part of the blood pressure elevation in the rat was due
to changes in the renin-angiotensin system (19, 40), a regulation that is seen by human SRY on
human RAS promoters (52, 53). Electroporation of Sry into the kidney of normotensive rats elevates plasma renin activity (19), an activity that is known to be elevated in human males relative to females (28). Evidence points to a combination of sex hormones and SRY regulating promoter activity of RAS genes, suggesting an elevation of the ACE/Ang II/AT1 axis and a depression of the ACE2/Ang-(1-7)/MAS axis (19, 23, 38, 40, 52, 53). As SRY is primarily known for its role in testis determination (4), the regulation of the RAS and blood pressure was not initially expected.

SRY is a member of the SOX transcription factor family that play important roles in regulating cell specialization and differentiation (29). The SOX family is composed of 20 human transcription factors, all sharing a homologous high mobility group (HMG)-box domain first discovered in the SRY protein (6). The HMG box of SOX transcription factors binds to DNA with sequence specificity resulting in both conserved transcription factor recruitment (50) and architectural modification by inducing a bend to DNA (49). Sry evolved from the Sox3 gene (12, 59), which in human is found on the X-chromosome. Thus Sry could have either maintained the regulation of the RAS genes from Sox3 or gained this function de novo. With recent elucidation of critical genes maintained on the Y-chromosome (3), understanding the impact these genes have on various disease phenotypes is of critical importance, especially with focus on similarities and differences between the Y-chromosome genes and their homologous counterpart on the X-chromosome. We hypothesize that there is preserved promoter regulation between SRY and SOX3 on the RAS genes, with some possibility of sex differences in the regulation of RAS components and ultimately blood pressure regulation. Therefore, the aim of this study was to further evaluate mechanisms of action, especially in the promoter regions of the RAS genes that were previously shown to be modulated by SRY.

**Materials and Methods**

*Sry and Sox3 in silico comparisons*
Nucleotide sequences for Sry (321 total mammalian sequences) and Sox3 (34 total vertebrate sequences from human to zebrafish) were aligned with ClustalW codon in MEGA5 (62). Phylogenetic analysis was performed with all Sry and Sox3 sequences to confirm none of the Sry sequences clustered within the SOX3 sequences. The test statistic dN-dS was used for detecting codons that have undergone selection, where dS is the rate of synonymous substitutions and dN is the rate of nonsynonymous substitutions. A positive value for the test statistic indicates an overabundance of nonsynonymous substitutions. The normalized dN-dS values were calculated for each codon, using the joint Maximum Likelihood reconstructions of ancestral states under a Muse-Gaut model (45) of codon substitution and Tamura-Nei model (61) of nucleotide substitution. Based on sequence alignments, those amino acids conserved in all SRY and SOX3 sequences were colored red and those conserved in only SOX3 but not SRY in cyan onto the structure of SRY-DNA (pdb 1j46). Amino acids were swapped in the HMG box of SRY (using pdb 1j46) for those of SOX3. Models for both SRY and SOX3 interaction with DNA were then energy minimized with the AMBER03 force field (15) with a pKa of 7.4, water density of 0.997 g/mL, and 0.9% NaCl mass fraction in YASARA (32). Molecular dynamic simulations were performed for 3.25 nanoseconds.

Promoter, Evolutionary Conserved Region (ECR), and transcription factor binding site analysis

SOX binding sites in each promoter were detected using Genomatix MatInspector (8). All promoters cloned except the ACE promoter contain predicted SOX binding sites. ECR analysis was performed in the ECR browser (48) using the parameters listed in the figure legends for each analysis. Conserved regions were then analyzed for conserved transcription factor binding sites between human and rat using RVISTA (37) for all vertebrate transcription factor binding sites.

Cloning - The pEF1 SOX effector plasmids cloning strategy was previously published (52). Details for the various human pGL3 luciferase reporter constructs were also previously
published (53). Starting with the human REN(-1444/+8) pGL3 reporter construct, the three SOX binding sites were sequentially removed resulting in the REN(-1444/+8) SOX mut reporter vector; ie the first site was removed followed by sequence confirmation, then the second site, and finally the third site. This was performed using phosphorylated primers flanking each SOX binding site (Table 1), followed by PCR with Phusion Hot Start II polymerase (Fermentas), and vector recircularized with T4 ligase. SOX binding sites were assessed in the MAS promoter using two strategies, 1) varying the size of the promoter insert or 2) mutating a specific SOX binding site. Both were performed using primer directed mutagenesis (Table 1). All constructs were sequence confirmed using BigDye Sanger sequencing on an ABI 3130xl genetic analyzer (Applied Biosystems). The SRY-MAS electrophoresis mobility shift assay (EMSA) was performed as previously published (52).

**Cotransfection and Luciferase assays in CHO Cells**

Twenty four hours prior to transfection of CHO cells, 5x10^4 cell/well were seeded on 24 well plates and incubated overnight in HAM’s F12K media (Sigma) supplemented with 10 mM HEPES and 10% fetal bovine serum (Atlanta Biologicals) in a humidified atmosphere at 37°C and 5% CO₂. CHO cells were selected for the analysis due to previously suggested reasons (53). Each well was transiently cotransfected with 50 ng effector plasmid (pEF1 vector), 500 ng firefly luciferase reporter construct (pGL3 vector), and 500 pg of control Renilla vector. Transfections with both SRY and SOX3 together were performed by using 50 ng effector plasmid total, for example 25 ng of SRY with 25 ng of SOX3 for SRY+SOX or 25 ng of SRY with 25 ng of empty pEF1 vector for SRY alone. Luciferase assays were performed using Stop & Glo luciferase kit (Promega) according to the manufacturer. Statistical analysis was performed on the standard error of the mean (SEM) of 3-4 independent experiments followed by Student’s t-test to determine significant differences (p ≤ 0.05) relative to the experimental control.

*Sry and Sox3 expression in the rat*
Initial work in the Fisher 344 Rat BodyMap project (71) did not annotate either \textit{Sry} or \textit{Sox3} transcripts. Therefore, we used the \textit{Sry} (EU984075.1) and \textit{Sox3} (XM_006227591.2) sequences from the rat to BLAST reads from the various Sequence Read Archive (SRA) datasets of the Rat BodyMap project (35). BLAST (2) was performed using only reads that were 98\% homologous with an E-value better than 1e-20. The number of positive reads was converted into the RPKM, reads per kilobase per million \([\text{# of mapped reads}] /[\text{length of transcript in kilobases}] / [\text{million mapped reads}]\). For the global expression profile, all SRA files for an individual tissue were pooled together for BLAST analysis (16 total) only separating on the sex of the animal (male vs. female). Three tissues were picked (testis, brain, and kidney) for profiling the error between animals \((n=4)\) and at each of four ages. The error was calculated using the standard error of the mean \((\text{SEM})\). Primate \textit{SRY} \((\text{NM}_0003140.2, \text{KP141769.1, NM}_0001032836.1, \text{DQ977342.1, EU294189.1})\) and \textit{SOX3} \((\text{NM}_005634.2, \text{XM}_004065361.1, \text{NM}_001193752.1, \text{XM}_001143467.3, \text{XM}_005227560.1)\) sequences were analyzed for Sra datasets of the kidney \((\text{SRX081994, SRX081951, SRX081930, SRX081966, SRX196347})\) and testis \((\text{SRX081999, SRX081954, SRX081933, SRX081969, SRX358238})\) for RPKM.

\textbf{RESULTS}

\textit{SRY} has rapidly evolved from \textit{SOX3}, resulting in differences in all domains outside of the DNA binding HMG-box. Among 34 species sequences for the \textit{SOX3} HMG-box (representing a wide range of vertebrates) only one of the 76 amino acids was not conserved (Fig. 1A). However, in the 321 sequences of \textit{SRY} (found only in mammals) 26 amino acids are not conserved. The larger number of \textit{SRY} sequences, compared to \textit{SOX3}, represents the previous resources invested in Sanger sequencing \textit{Sry} throughout mammals primarily for Y-chromosome phylogenetic work, while the few \textit{Sox3} sequences are mostly from whole genomes sequences of species. In order to address whether \textit{SRY} and \textit{SOX3} DNA binding sites remain similar, codons of \textit{Sox3} that are under selective pressure (more negative dN-dS, Fig. 1B) were
identified. Amino acids 11, 30, 45, 60, and 66 have dN-dS around -10 and are conserved in all
or most of the SRY sequences. When conservation of amino acids for SRY and SOX3
sequences are mapped onto the protein structure, a highly conserved DNA contact is revealed
(Fig. 1C), suggesting both proteins bind to DNA with similar sequence specificity. Modeling the
SOX3 protein reveals similar molecular dynamics for all amino acids between SRY and SOX3
(Fig. 1D).

Utilizing the previously designed luciferase vectors controlled by the various RAS
promoters, SRY and SOX3 were tested for promoter regulations (Fig. 1E). The promoter of AGT
was up-regulated by both SRY and SOX3, while the promoters of ACE2, AT2, and MAS were
down-regulated by both SRY and SOX3. The only promoter that was differentially regulated was
that of Renin, REN, which was up-regulated by SRY and down-regulated by SOX3.

The MAS promoter was previously shown to be regulated by multiple SOX subfamily
members and shown to be directly bound by SRY (52), therefore we elected to characterize the
shared binding sites between SRY and SOX3 on this promoter. The MAS promoter contains no
ECRs, with sequence divergence in all species for the promoter and the 5'UTR (Fig. 2A),
reducing our ability to use evolutionary genomics to decipher the molecular binding sites.
Therefore, the binding sites for SOX were tested using two different approaches; 1) a cleavage
series to generate MAS promoters of various sizes, and 2) point mutations within two of the
SOX binding sites. The full MAS promoter used [MAS(-2222/+4)] contains 10 predicted SOX
binding sites. This full length construct is regulated by all of the SOX A/B protein family
members (Fig 2B). Cleaving part of the MAS promoter such that there are 7 possible SOX
binding sites [MAS(-1123/+4)], or 1 possible binding sites [MAS(-377/+4)] had little effect on
relative luciferase production (Fig 2C), with the exception being the 5 possible binding site
MAS(-847/+4) construct that had significant up regulation of the promoter. The addition of SRY
or SOX3 to the cells treated with the various sized promoter constructs showed a complete loss
of SRY/SOX3 repression only on the shortest of the constructs (MAS (-377/+4), Fig 2D) suggesting that the cluster of SOX binding sites between -847 and -377 are important for promoter regulation.

To assess if the SOX binding sites outside of the -847 and -377 region are also critical to MAS promoter regulation, but were missed due to the nature of promoter cleavage series, we performed mutations at the distal (-1855) triple SOX binding site and the proximal SOX binding site (-86). Both mutations result in up-regulation of the MAS promoter in CHO cells, with significance seen with the mutation at -1855 and a trend seen at -86 (Fig. 2E). To further validate the in vitro confirmed SOX binding site in MAS at -86, we used the short MAS promoter construct [MAS(-377/+4)], again mutating the SOX binding element, recovering the mutations, and mutating it to a different sequence than the first mutation (Fig. 2F). Mutation at the SOX binding site for both changes (MutSOX-86 and Mut2SOX-86) resulted in around 5 fold up-regulation of the MAS promoter, while recovery of the mutation (RecSOX-86) yielded similar values as the wild type MAS promoter. DNA probes for the MAS promoter containing this SOX binding site can be bound by both 6X His tagged and GST tagged SRY protein (Fig 2G). The data on the MAS promoter indicates that SRY and SOX3 regulate the MAS promoter in cis through a combination of multiple SOX binding sites. The ENCODE data that the MAS promoter is regulated through the GATA 2/3 transcription factors, reaffirms our interests of the SOX gene regulation, as many promoters have been shown to contain competitive SOX and GATA mechanisms (34, 42, 43).

Moving from MAS to the other promoters of renin angiotensin system, for two of the promoters that are similarly regulated by SRY and SOX3, the next step was to determine if both proteins operate using the same mechanism or if they result in synergistic or additive regulation. The up-regulated AGT promoter had a concentration dependent activation by SRY and SOX3, with SRY&SOX3 simultaneous treatment resulting in a similar activation as SRY or SOX3 alone
The down-regulated promoter of ACE2 was also regulated in a concentration dependent manner with SRY&SOX3 again resulting in a similar activation as SRY or SOX3 alone (Fig. 3B). Both results suggest that SRY and SOX3 regulation have an overlapping mechanism in promoter control.

In order to address possible mechanistic insight into the different regulation of the Renin promoter by SRY and SOX3, an analysis of conserved transcription factor binding sites was performed on the Renin promoter to determine if there were any Sox consensus binding sequences (Fig. 4). The 5’ region of the Renin gene is located around 30kB from its close syntenic gene Kiss1, with very few conserved ECRs located in the intergenic space (Fig. 4A). Addressing the proximal promoter of Renin, an approximate 500 bp ECR (Fig. 4B) is found to contain 12 conserved transcription factor binding sites (Fig. 4C) none of which are SOX binding elements. The REN(-1444/+8) promoter driving our luciferase production contains a total of 3 SOX binding elements located >1kB from the transcriptional start site (TSS), outside the ECR (Fig. 4D). A TATA box is located 29 bps away from the TSS suggesting proper distance for transcriptional initiation (Fig. 4D).

Delivery of SRY only at 50 ng, and not 25 ng, significantly up-regulated the renin promoter, while both concentrations of SOX3 result in down-regulation of the promoter (Fig. 5A). Surprisingly, delivery of SRY&SOX3 results in a repression similar to SOX3 alone (Fig. 5A), suggesting that regulation mechanisms differ between SRY and SOX3 on the Renin promoter. To further understand this difference, the REN(-1444/+8) promoter was tested with members of each SOX subfamily (Fig. 5B). The promoter was significantly regulated by SRY (up-regulated), SOX2 (down-regulated), and SOX3 (down-regulated). It should be noted that SOX2 and SOX3 are both members of the SOXB1 subfamily, with highly conserved sequence and function (30). Removal of all three SOX binding elements in this promoter construct resulted in an increase in promoter activity for all SOX constructs (except SOX30) with complete loss of promoter
repression seen in the SOX2 and SOX3 treated cells (Fig. 5C). This suggests that Renin
promoter repression seen by SOX3 and SOX2 is a cis-acting function while the activation by
SRY is likely a trans-acting function.

In order to understand the context for SRY and SOX3 regulation of the RAS, an
analysis of tissue expression was performed in the rat, an organism with extensive work done to
colorize the protein products of the RAS (51). Using unannotated (for Sry and Sox3) RNAseq datasets from the rat BodyMap project (71), Sry and Sox3 expression profiles were
determined using 320 RNAseq experiments for 10 tissues of male and female Fisher 344 rats
(Fig. 6). Global expression showed identifiable transcripts for Sry in all male tissues; however,
Sox3 transcripts were only detected in male and female brain and the male testis (Fig. 6A). The
testis (Fig. 6B) and brain (Fig. 6C) both had expression of Sry and Sox3 in the four age groups
of male rats, while in the kidney only Sry transcripts were detected (Fig. 6D). The kidney is the
primary source for Renin found in the blood, and thus the top tissue of interest for Renin
regulation control in blood pressure. The rat expression profile for Sox3 closely matches that of
the human protein atlas for human SOX3 (http://www.proteinatlas.org/ENSG00000134595-
SOX3/tissue). In some of the human protein atlas RNA and protein expression samples, SRY
was detectable in the kidney (http://www.proteinatlas.org/ENSG00000184895-
SRY/tissue/kidney#img). We therefore tested primate expression of SRY and SOX3 using
publically available Sra datasets for RNAseq of kidney and testis. While SRY and SOX3 were
both detected in the testis of primates, only SRY could be detected in the kidney (Fig. 6E).

DISCUSSION

There are sex differences in the prevalence of many conditions including cardiovascular
diseases, asthma, autoimmune disorders, birth defects, neurological disorders, and cancers
(47). Of these diseases, the RAS has been associated with the cardiovascular, respiratory,
neurological, reproductive and immune systems along with cancer progression (24, 27, 41, 54,
Sry transcripts/proteins have been identified in all of these tissue types in the rat and in the human (64, 67). Part of the sex disparities for blood pressure is due to hormonal contributions (57), but additional evidence points to sex chromosome contributions (10, 11, 60). Previous research from our group has established animal models for Y-chromosome specific elevation of blood pressure (21) and behavior alteration (14, 63) in the rat, narrowing down the entire Y chromosome to the Sry gene (19). This has opened up the question of whether SRY function in blood pressure control might be conserved from the divergence of the mammalian X-chromosome gene SOX3, or whether the regulation is novel to SRY.

Genes on the Y-chromosome evolve much faster than their X-chromosome homologs or other autosomal genes due to a loss of crossover, the founders effect and increased mutagenesis in the testis (25). In our data we confirm this, with SOX3 protein sequences from a broad range of vertebrates highly conserved for the HMG-box; however, the HMG-box of SRY found in a small subset of vertebrates (mammals) has a higher rate of divergence (Fig. 1). Still, structural and functional assessment of these sequences confirms that DNA binding through the HMG box is similar between SRY and SOX3, maintaining the critical amino acids for proper DNA contact, with sequence divergence between SRY and SOX3 found at amino acids that do not contribute to the folding of the domain or DNA contact.

With a solid number of studies pointing to sex-specific differences in RAS modulation (26), our previous data showing SRY regulation of the RAS genes both in vitro (40, 53) and in vivo (19) in combination with the conserved DNA binding between SRY and SOX3, we hypothesized that the promoter regions of the RAS genes may be regulated by both SRY and SOX3 transcription factors, which may potentially contribute to sex differences in blood pressure regulation. The promoter of AGT was up-regulated by both SRY and SOX3, while the promoters of ACE2, AT2, and MAS were down-regulated by both SRY and SOX3. The only promoter that was differentially regulated was that of Renin, REN, which was up-regulated by SRY and down-
regulated by SOX3. This suggests that SRY has gained a novel protein function from its homolog, SOX3, in trans-regulating Renin (or loss of cis-regulation seen by SOX3) while maintaining the regulation of genes such as AGT, ACE2, AT2, and MAS. As many of the experiments shown in this paper are based on luciferase promoter constructs, the size of these constructs may influence the outcomes of the SRY/SOX3 regulation with potential regulation left out of the construct design. However, the constructs as designed show promoter regulation that is supported by the in vivo Sry electroporation studies in the rat (19) and thus likely contain the minimal promoter for Sry regulation in vivo. It is also possible that regions outside of the HMG-box of SRY and SOX3 control the differences seen in regulation of Renin. Renin, primarily produced in the kidney, is the rate-limiting step of the RAS. Thus the activation by SRY, which is expressed in the kidney, might contribute to elevation of blood pressure through increased Ang II peptide; however SOX3 does not appear to regulate this step. A decrease in the Ang-(1-7) arm of the RAS maintained in SRY and SOX3 treated cells, further potentiates blood pressure elevation control by SRY. The androgen receptor is known to directly interact with SRY and regulate gene promoters (70), suggesting potential for synergistic effects between Sry and androgen signaling to contribution to sex differences in blood pressure.

The rat is a well-established model organism for understanding the contributions of the Y-chromosome on phenotype. Almost 25 years ago, crosses of the Y-chromosome from the Spontaneously hypertensive rat (SHR) (21) and the SHR Stroke Prone (SHRSP) (13) with the normotensive WKY rat resulted in a blood pressure elevation, while the reciprocal crosses reduces blood pressure in the SHR rat. Crosses of the Brown Norway (BN) Y-chromosome onto the SHR were also able to reduce blood pressure (31). Recently, crosses for the Y-chromosome from the Salt Sensitive (SS) rat, with salt-dependent blood pressure increases, to the normotensive BN altered kidney function (39). These rat consomic animals establish needed model organisms to translate the literature of human Y-chromosome contributions to blood
pressure (11, 18, 60). Details in this paper provide the first molecular mechanisms for differences in homologous transcription factors found on the X- or Y-chromosome to contribute to sex differences in blood pressure regulation.

In the rat and human, Sry is expressed in numerous tissues including testis, brain, heart, and kidney (64); however, Sox3 expression appears to be limited to the brain and testis. In addition to testis development (69), Sox3 has been proposed to be critical for brain development, particularly for the hypothalamic-pituitary-adrenal axis, HPA (46, 56). In the brain, proteins of the RAS have been shown to contribute to the HPA (36), an axis with known sex differences (33, 66). In the classical RAS, Ang II acts on the pituitary to release vasopressin and ACTH. These pituitary cell lineages are developmentally regulated by SOX3 (1), such that they are likely activated by the ACE/Ang II/AT1 axis of the RAS. In concert, the ACE 2/Ang-(1-7)/MAS axis of the RAS is repressed in the pituitary (16). It is of interest to note that in the rat models of Y-chromosome crosses, behavioral differences have also been seen in males (14). With the brain RAS shown to be involved in neurological development (5, 44) and RAS inhibitors given during pregnancy associated with mental delays in the offspring (7), our data also suggests a potential impact on sex differences in brain development through the RAS regulation.

In summary, the fact that SOX3 functions in the brain and with the documented literature above on RAS in brain development, we suggest that regulation of multiple RAS proteins in the brain was inherited and selected on following the X-Y split in mammals, with SRY gaining additional expression in tissues such as the kidney and trans regulation of genes such as Renin resulting in a male specific modulation of blood pressure.


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FIGURE LEGENDS

Figure 1. Functional conservation in SRY and SOX3. A) Sequence alignment of the HMG box of human SOX3 and SRY with conservation shown below each in multiple species. Those sites marked with * are 100% conserved in all species and those with : are functionally conserved. Amino acids in red are conserved in all SRY and SOX3 while those in cyan are conserved in SOX3 only. Amino acids highlighted in yellow are those in which mutations in SRY are associated with sex reversal (50). B) Normalized values for each amino acids’ (x-axis) non-synonymous rate (dN) minus the synonymous rate (dS) for the 34 sequences of Sox3. Values more negative suggest codon selection. C) Known structure of SRY bound to DNA (gray, pdb file 1j46) with the amino acids conserved among all SRY and SOX3 sequences shown in red and those conserved in SOX3 but not SRY shown in cyan. DNA is shown in gray. D) Root-mean squared fluctuation (RMSF) for each amino acid over a 3.25 nanosecond molecular dynamic simulation for the known structure of SRY (red) or the modeled SOX3 (cyan) indicate similarities of the 2 proteins. E) The pEF1 expression vectors for either human SRY (light gray) or SOX3 (darker gray) were transfected into CHO cells along with a pGL3 vector containing the human promoter (shown on the bottom) of various RAS genes driving the production of luciferase. Each sample was normalized to an empty vector control of pEF1 (black). Error bars are shown as the SEM. Each sample is compared to the control for each promoter. Significant differences ($p \leq 0.05$) are marked with *.
Figure 2. Functional SOX binding sites in the MAS promoter. A) ECR browser analysis of the MAS pGL3 promoter construct for the rat (rn4), mouse (mm10), dog (canFam2), and monkey (rh2Mac2). Regions in red and green share conservation with human based on 20 bp stretches sharing >50% homology. Below the ECR is the location of various landmarks of the promoter vector such as sites used to create cleavage series (-2222, -1123, and -377), predicted SOX binding sites (blue), in vitro confirmed SOX binding site (red), the location of the TATA box (-290), proposed transcriptional start site (TSS, -266), translational start site (ATG, +1), and ENCODE transcription factor binding with relative score and cell line of observation. B) The human MAS promoter treated with various SOX A/B proteins, all of which significantly repress the promoter. C) Luciferase production of four MAS promoter constructs with differing lengths, all made relative to the full length MAS (-2222/+4) construct regulation level. D) Promoter analysis of four MAS promoter constructs transfected with control (black), SRY (light gray) or SOX3 (dark gray) vectors. Total number of SOX binding sites in each construct is shown below the graph. E) Mutations to the MAS(-2222/4) promoter construct at two of the ten SOX binding sites (-1855 or -86) result in elevation of promoter activity. Data is shown as the raw luciferase:Renilla ratio. F) As the MAS(-2222/+4)MutSOX-86 showed a trend in altering regulation and was previously shown to be an in vitro binding site for SRY the site was investigated with the smaller MAS minimal promoter, MAS(-377/+4). The SOX binding site was mutated to two separate sequences, MAS(-377/+4)MutSOX-86 and MAS(-377/+4)Mut2SOX-86, resulting in elevation of promoter activity. One of the mutations was repaired to recover the SOX binding site, MAS(-377/+4)RecSOX-86, confirming only this site was altered in the promoter mutagenesis. G) The sequence for the SOX binding site at -86 was synthesized with a biotin tag and EMSA assays performed using both His and GST tagged Sry proteins confirming that SOX proteins can bind to this element. Error bars for all graphs are shown as the SEM. Each sample is compared to the control for each promoter. Significant differences ($p \leq 0.05$) relative to the control are marked with a *.
Figure 3. Similar mechanisms of regulation by SRY and SOX3 on promoters of AGT and ACE2. A) Activation of the AGT promoter with two concentrations of transfected SRY (light gray) or SOX3 (dark gray). In addition, a double transfection of SRY and SOX3 (checkered) shows non-additive activation suggesting shared mechanisms of promoter control. B) Repression of the ACE2 promoter with two concentrations of transfected SRY (light gray) or SOX3 (dark gray). In addition a double transfection of SRY and SOX3 (checkered) shows non-additive repression suggesting shared mechanisms of promoter control. Error bars are shown as the SEM. Each sample is compared to the control for each promoter. Significant differences ($p \leq 0.05$) are marked with *.

Figure 4. Promoter analysis of Renin. A) ECR browser analysis of the intergenic space (roughly 30kB) between Renin (REN) and the syntenic KiSS-1 metastasis-suppressor gene (KISS1) for the rat (rm4), mouse (mm10), dog (canFam2), and monkey (rh2Mac2). Regions in blue are protein coding exons, yellow are UTRs, and those in red and green share conservation with human based on 20 bp stretches sharing $\geq 90\%$ homology. B) ECR browser analysis for the proximal promoter of Renin with regions in red and green showing conservation of 50 bp stretches with 50% homology. C) The large ECR in the proximal promoter of Renin was analyzed for shared TF binding sites between human and rat using the RVISTA software selecting for all possible TFs. D) The sequence of REN(-1444/+8) pGL3 promoter construct identifying the TATA box (red, highlighted yellow) located within the proper distance of the 5'UTR (highlighted yellow). Additionally the conserved TF sites are colored relative to that in panel C. The three potential SOX binding sites (highlighted in red) fall outside of the ECR.

Figure 5. Regulation of the Renin promoter by various SOX proteins. A) Activation of the REN promoter with two concentrations of transfected SRY (light gray) or repression by two concentrations of SOX3 (dark gray). In addition a double transfection of SRY and SOX3 (checkered) showed down-regulation of the promoter similar to that of SOX3 transfected cells.
B) Transfection of CHO cells with either the control pEF1 vector (black) or one of the SOX pEF1 constructs (gray) along with the pGL3 REN(-1444/+8) luciferase vector. Error bars are shown as the SEM. Each sample is compared to the control for each promoter. Significant differences ($p \leq 0.05$) are marked with *. C) The REN(-1444/+8) promoter construct was mutated to remove each of the three SOX binding sites; this construct was named REN(-1444/+8) SOX mut. The mutant pGL3 construct (gray) was transfected with each of the SOX pEF1 expression vectors resulting in an increase in promoter activity relative to the wt pGL3 promoter construct (black) for all but the SOX30 vector ($n = 1$).

**Figure 6. Expression profile for Sry and Sox3 in the rat using data from the Rat BodyMap project.** Expression profile for Sox3 (gray) and Sry (black) in female (F) and male (M) RNAseq data for 10 tissues in the Fisher 344 rat (A). Each tissue is a pool of 16 individual RNAseq experiments (four animals at four ages). Data is shown as RPKM- reads per kilobase per million. While Sry is ubiquitously expressed in male only, Sox3 transcripts are found at all four ages (2, 6, 21, and 104 week old rats) in the male Testis (B) and Brain (C). The Kidney showed expression of Sry at all four ages with no detectable Sox3 reads (D). Error bars for the individual tissues (B-D) represent the SEM of four independent RNAseq datasets at each age. E) Analysis of primate (*Homo sapiens, Gorilla gorilla, Macaca mulatta, Pan troglodytes*) kidney and testis Sra datasets for expression of SOX3 (gray) or SRY (black). Expression of both SRY and SOX3 are seen in testis while only SRY expression is found in kidney.
A

**AGT (-1576/+34)**

- Control
- 25ng SRY
- 50ng SRY
- 25ng SOX3
- 50ng SOX3
- 25ng SRY & SOX3

![Graph showing relative luciferase units for AGT](chart)

B

**ACE 2 (-1899/+34)**

- Control
- 25ng SRY
- 50ng SRY
- 25ng SOX3
- 50ng SOX3
- 25ng SRY & SOX3

![Graph showing relative luciferase units for ACE 2](chart)
Table 1. Primers used for mutagenesis of luciferase promoters. To make each of the mutated promoter constructs the primers in this table were used. The first column shows the construct name as used in the paper, the second column the individual primer names, and the third column the primer sequences. For mutations, the SOX binding site is underlined and bolded with the DNA bases to be mutated shown in red.

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<tr>
<th>Construct name</th>
<th>Primer name</th>
<th>Primer sequence</th>
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<tr>
<td><strong>Renin SOXdel1</strong></td>
<td>R1-RenSoxdel</td>
<td>5’-CAT CAG GTT TGA CTG ACA AA-3’</td>
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<td>5’-CTT GAA TGT AAT CAG ACA GAG-3’</td>
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