

Genome Informatics: Assignment 3

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Abstract

***SRY* is a gene on the *Y* chromosome in humans which is significant for sex differentiation primarily in early development. Even though gene expression differences between the sexes are mostly attributed to hormonal differences, we here show that both *SRY* and sex chromosome complement carries weight, and importantly, that *SRY* only appears to affect gene expression when a *Y* chromosome is present. By analysing the human *SRY* sequence and transcript to their respective orthologues, and by identifying structurally important parts of the gene, we gain insights into the functionality of *SRY* and its effect on expression levels for autosomal genes.**

Y chromosome contains the so-called Sex-determining Region Y (*SRY*; also known as the Testis-determining Factor [1]) – a gene encoding for the likewise named gene-regulatory transcription factor. Also in other mammals, *SRY* plays a similar role, and is typically the one signifying gene for differentiation of bipotent organs to the male variants [2, 3].

All orthologs of *SRY* are signified by the existence of a specific High-Mobility Group (*HMG*)-box, which all tend to be involved in DNA-regulatory processes, such as looping and bending of the structure [4]. The group as a whole is very prevalent in nature, but the various kinds can be extremely diverse, with conservation of amino acids as low as 50 % between species [5].

The *SRY HMG*-box bases the common reference point for several developmentally important genes entitled *SOX* (*SRY*-like *HMG*-box) genes, which are named as such precisely because they contain the common *HMG* box, but share no other greater similarities because of this. Together, *SRY*, which is sometimes referred to as *SoxA*, and the other *SOX* genes form the so-called *SOX* gene family. *SRY* has been proven to be sufficient for testis development, but in the absence of the gene, male sexual fate commitment can still happen in some cases through complementary action of the *SOX-9* gene [6, 7].

We here investigate the structural features of *SRY* in humans and its orthologs and attempt to assess differences and similarities. We also, in imitation of Wijchers et al. [8], show that *SRY* is not the sole driving factor of differential expression in autosomal genes, but that also the sex chromosome complement is important. More precisely, our results indicate that *SRY* acts collaboratively with the *Y* chromosome in order to induce differential expression in a set of autosomal genes.

Preface

This is an assignment report in connection to the *Genome Informatics* module in the Computational Biology course at the University of Cambridge, Michaelmas term 2016. All related code is as of December 27, 2016 available per request by contacting hpa22@cam.ac.uk. Data used is acquired from NCBI's Gene Expression Omnibus, GEO Series Accession number [GSE21822](#). Ortholog and SNP data is supplied by the Ensembl database under the *SRY* gene information page. For genome related analyses, the GRCh38.p7 version of the human genome was used.

Introduction

In placental mammals and marsupials, sex differentiation is generally determined by the complementary *Y* chromosome, which is typically linked to testis development and overall commitment to male sexual fate. In humans, this happens primarily because the

Methods

Using the given approach in [8], we analyse our raw cDNA from a hybridization to Affymetrix mouse

genome 430 2.0. As samples, we have four different genotypes, consisting of sex chromosome setups XX , $X/Y\text{-}sry$, $X/Y\text{-}$ and $X/Y\text{-}sry$. The two prior represent individuals with male phenotypical characteristics, and the two latter the female counterparts. The chromosomal setups with the *sry* suffix are named as such because they have had an *SRY* transgene inserted into an autosome. Similarly, $/Y\text{-}$ represents Y chromosome variants that have been modified to not contain the *SRY* gene [9]. For every chromosomal setup, there are three biological replicates, where all individual data is gathered from earpunches of the specimen involved.

According to the authors in our study of interest, the Bioconductor RMA package in R is used for preprocessing of the raw .CEL files. However, no such package appears to exist [10]. Because of this discrepancy, we in this investigation use the Bioconductor affy package for preprocessing, using the default *rma* function with quantile normalisation and RMA background correction. Like the authors, we define differential expression as being greater or equal than a log-2 fold change of 1.2, as well as fulfilling a Student's T-test with $p < 0.05$). This we do using the *simpleaffy* package, along with quality-control by *affyQCreport*. In addition, we also filter probes using MAS5.0 absent/present calls.

Orthologues to *SRY* are taken from Chimpanzee, Cow, Macaque, Mouse, Pig (two genes), Rat and Vervet-AGM. For alignment of the these genes and their transcripts we use Clustal Omega 1.2.3 and MUSCLE 3.8 respectively. Motif identification and analysis is done with Meme-suite 4.11.2 modules MEME, Tomtom and MAST.

Results

Gene expression hints at collaborative action between *SRY* and the Y chromosome

In the quality control of our samples, no clear discrepancies can be determined to be present. We therefore choose to keep all samples for our analyses.

To determine whether sex-chromosome sensitivity (SCS) affects autosomal gene expression over the whole genome, we produce the comparison seen in fig. 1A and B where differences between the genotypes of the males (A) and females (B) are shown. Using our definition of significant expression values, we are able to round up 126 SCS genes, as opposed to Wijchers et al. who find 369 differentially expressed genes under the same conditions. Nevertheless, we are able

to produce the same trends in the expression, namely that our SCS genes are biased towards an XX configuration at its core in males, and the contrary in females. Similarly, we establish a set of sex-sensitive and dimorphic genes (SSD) comparing XX and $X/Y\text{-}sry$ individuals, and are able to find a selection of 46 (175) such genes. We also identify 287 (401) dit in either sex. A list of the genes contained in each group can be provided upon request.

We can furthermore see hints of the effects of *SRY* in fig. 3D and E. Notably, in the XY comparison, *SRY* introduces a trend towards the samples without the transgene, whereas in our XX samples the opposite happens. However, the difference in subfigure E is significantly smaller, suggesting that the *SRY* acts collaboratively with the Y complement, i.e. we have an effect from *SRY* primarily when we also have the Y chromosome. It is also evident that some genes have clear preferences towards the Y chromosome (cf. the outliers in fig. 3A, B, C, F), giving fairly constant differential expressions between the comparisons, and subsequently not being affected categorically by the presence of *SRY*. Because of these results we converge on our previously mentioned set of sex sensitive dimorphic genes (see fig. 1B) in determining the effects caused solely by our transgene.

When instead limiting our analysis to our new set of genes, we see from the blue triangles in fig. 3D and E that the effect from *SRY* has vanished. We nonetheless retain that a part our genes are being affected mostly by the sex chromosome, most notable seen in fig. 3C and F. Again, we also see that the trend shift for most of the SSD genes only occurs when *SRY* is put in combination with the Y chromosome.

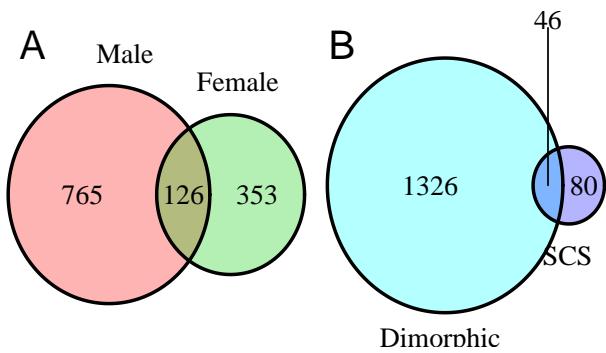


Figure 1: Venn representations of genes differentially expressed in various groups of interest, and their corresponding overlaps.

***SRY* is poorly conserved apart from the HMG-box structure**

In doing sequence alignments of *SRY* and its orthologues, we find that the sequences differ significantly, with almost no conservation of the overall structure. The one conserved structure is the region coding for the transcript HMG-box; when mapping the human *SRY* HMG-box to the 9 orthologues the result is a ca. 90 % identity in the (non-fragmented) HMG-box region. In contrast, when only the gene sequences are set to match against each other, they do so with ca. 70 % conservation in the same region. The transcripts perform slightly better with ca. 85 % complete or near global match. Alignments can be provided upon request.

When piped through motif detection and analysis software, the conserved and matched motifs typically either correspond to other *SOX* genes or indeed HMG-boxes, further establishing the notion of the HMG-box being the structurally most significant part of the gene and transcript. It also suggests that the gene is first and foremost modulatory, i.e. having the role of binding to parts of the DNA and structurally affecting it in various ways.

By analysing SNP data, we find that point mutations which give rise to disease or a phenotypic change are focused to the HMG-box region in humans. This can be seen in figure fig. 2. Other SNPs (not shown here) are still concentrated towards the region, although a far greater amount of mutations are distributed outside of this area.

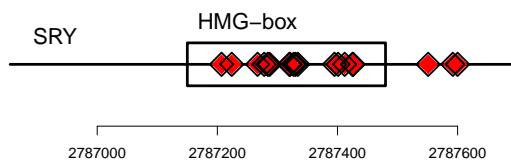


Figure 2: Known SNPs related to disease or changes in phenotype. Axis shows chromosome coordinates.

Discussion

Analysing the sequence and its structural traits hints at *SRY* indeed being mostly modular, and functionally revolving around its ability to bind to and affect DNA structure. Also the SNP investigations reinforce this, as it appears as the transcript to a higher degree tends to be lost or malfunctioning when the modulatory region is altered.

Our expression analysis shows that, in addition to being regulated by *SRY*, a fair amount of genes are

dependent primarily on the sex chromosome complement. In addition, we see that the genes being affected in their expression mostly by *SRY* in fact only appear to be so when there is also a Y chromosome prevalent, and interestingly enough almost exclusively in a repressive manner. Further investigations are clearly required, but even from our analysis it appears plausible that some genes are driven mostly by chromosomal complement, whereas some instead are directly or indirectly repressed by *SRY*. Importantly, it appears as if the modulatory significance of the *SRY* transcript is mainly in relation to the parts of the Y chromosome, and seemingly not a genome-wide phenomenon. Because of the repressive effect, it could be the case that *SRY* hampers transcription of genes directly through its structurally affecting role, but this phenomenon could also be indirect; further analysis is needed.

The differences in genes sensitive to our different variables ought to be mainly because of the seemingly different approach in preprocessing of the data. Still, this gives us relatively different results in considering the effects of *SRY*, as we see an effect related to the transgene in addition to just the sex-sensitivity, which is the result of the authors. Other results are difficult to compare due to the representation of the data points found in the reference paper, as the authors have chosen to visualize the set on top of the subset and thus effectively preventing an accurate comparison.

Our results here have given a further insight into the impact of *SRY* with respect to autosomal gene expression, and suggested that *SRY* might work collaboratively with the Y chromosome in order to affect this. Ultimately, more rigorous studies on *SRY* transcript targets and functionality is naturally what can help unravel the true nature of these results.

Acknowledgements

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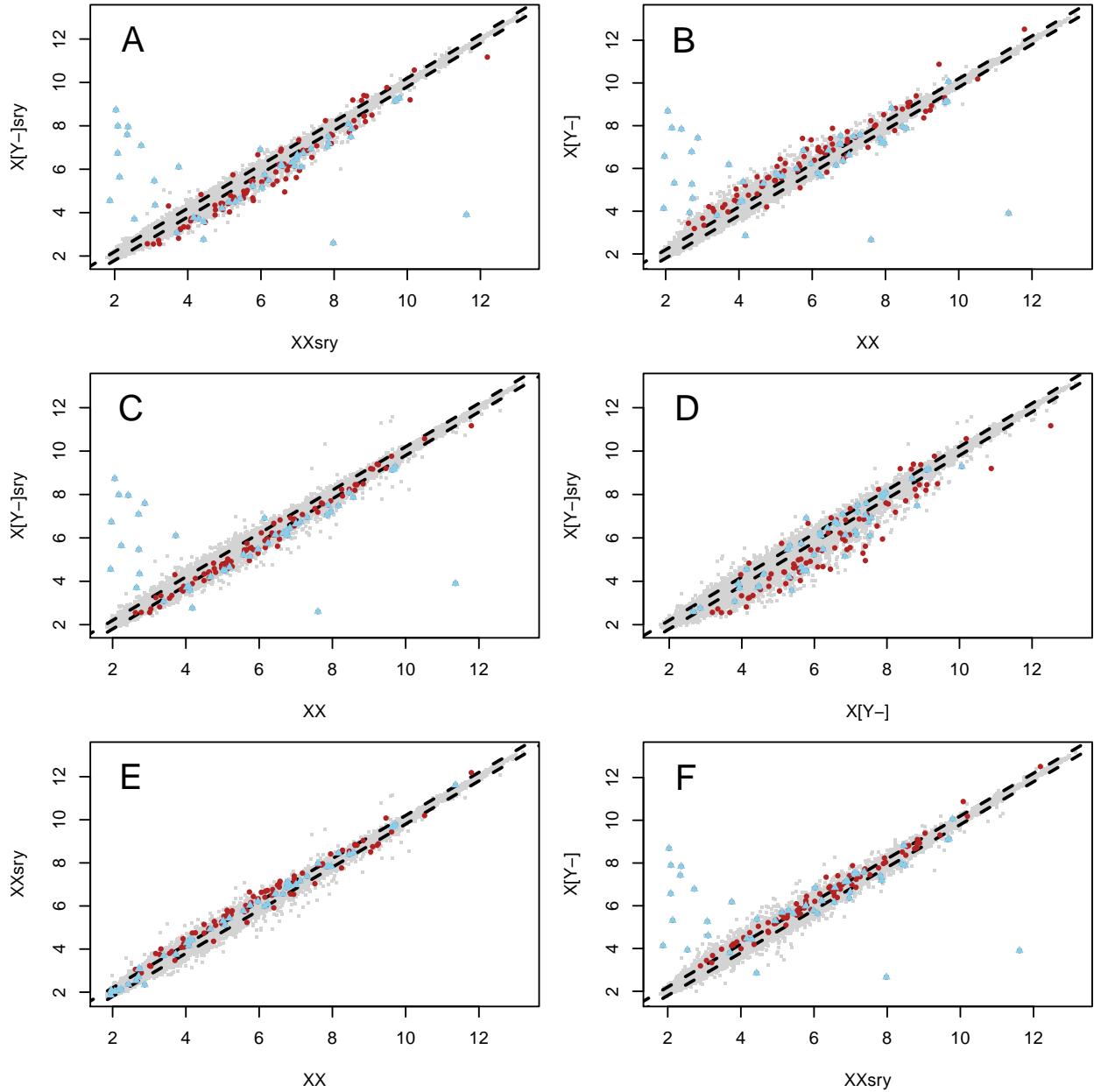


Figure 3: Pairwise comparisons between log-2 gene expression values of the four core chromosomal pairs. The dashed lines represent a log-2 fold change of 1.2. Red dots represent the SCS genes, whereas the blue triangles correspond to the sex-chromosome sensitive XY dimorphic genes. Note how subfigures **A**, **B** and **C** are involved in determining the sets of SCS and SSD genes (see fig. 1).