Characterisation of the M-locus and functional analysis of the male-determining gene in the housefly
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Document Version
Publisher's PDF, also known as Version of record

Publication date:
2018

Link to publication in University of Groningen/UMCG research database

Citation for published version (APA):

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English summary
The housefly, *Musca domestica* is particularly suited to investigate the evolution of sex determination and sex chromosomes because it has a polymorphic sex determination system. The male-determining *M*-locus, typically located on the Y-chromosome, can also be present on any of the five autosomes or even the X-chromosome. Recently, based upon differential expression analysis, a male-determining gene was identified and termed *Mdmd* for *Musca domestica* male determiner. *Mdmd* appears to have arisen as a duplication of the splicing regulatory gene *CWC22*, called *nucampholin* (*ncm*) in insects. To further characterise the *M*-loci in terms of genomic organisation and function, I addressed several questions about *Mdmd* structure and function: What is the genomic organisation of *M*-loci on different chromosomes? What is the coding sequence of *Mdmd*? To what extent are the different *M*-loci conserved? What is the evolutionary relationship between *Mdmd* and its paralog *CWC22/ncampholin*? When and where is *Mdmd* expressed? Is *Mdmd* sufficient for male function?

Although *Mdmd* was identified as the male-determining gene in the housefly, the complete sequence of *Mdmd* and its embedding in the *M*-locus remained unknown. In Chapter 2, I investigated the complex nature of *M*-loci in two autosomal *M* strains, *M iii*(M-locus on autosome III) and *M v*(M-locus on autosome V). I found that the *M*-loci contain multiple copies of different sequences of *Mdmd* sequences, with various levels of homology to each other. Interestingly, the *M iii*-locus and the *M v*-locus share common sequences. On the basis of these common sequences, I identified an open reading frame (ORF) that is part of the *Mdmd* gene (Chapter 3). Sequences with high similarity to the *Mdmd* ORF were also detected in *M ii* (M-locus on autosome II) and *M y* (M-locus on Y-chromosome) strains, but not in the *M i*(M-locus on autosome I) strain, which probably has a different male-determining gene (s). This ORF is assumed to be the coding sequence of *Mdmd*, the functional male-determining gene.

The liability and turnover of sex chromosomes is a remarkable aspect of sex determination evolution. Sex chromosomes are believed to evolve from ordinary autosomes that lost recombination after having acquired a sex-determining role. What drives the evolution of new sex chromosomes is not yet well understood. My results in *M. domestica* provide support for the birth-decay-rebirth model of sex chromosome evolution. The high sequence similarity of *Mdmd ii*, *Mdmd iii*, *Mdmd v* and *Mdmd v* suggests that all *Mdmd* genes originated from a common ancestral sequence. A comparison of *Mdmd* protein sequences and its paralog *CWC22/ncampholin* in Chapter 3 suggests a scenario of *M*-locus evolution, whereby the male-determining gene *Mdmd* evolved after a single duplication event of *Md-ncm* generating a proto-Y chromosome. Whether this happened on the ancestral Y or on an autosomal pair that was not yet involved in sex determination cannot be
answered at this moment.

The next stage of Y-chromosome evolution would be the reduction of recombination in the surrounding *Mdmd* region, followed by accumulation of repetitive sequences and transposons due to the lack of recombination on the proto-sex chromosome. Consistent with this model, I found that *M*-loci in the *MIII* and *MV* strain contain transposable elements and repetitive sequences (Chapter 2). Subsequent amplification of *Mdmd* appears to have led to the complex structure of the *M*-locus, as multiple tandemly repeated copies of *Mdmd* are found in *MIII* and *MV* males in Chapter 2. After amplification, the *M*-locus may have translocated multiple times as a cluster from the Y to an autosome and/or subsequently between autosomes, generating novel Y-chromosomes. In addition, the data presented in Chapter 2 revealed that to some extent different sequences exist in different autosomes, indicating that after translocation, the *M*-locus underwent further independent genomic changes on each autosome. The existence of multiple different autosomal *M* variants in the housefly provides a unique opportunity for further study of early stages of sex chromosome evolution.

As *Mdmd* is a crucial gene for male development, localising *Mdmd* mRNA in different embryonic developmental stages is needed to understand its regulation in the sex determination pathway. In Chapter 4, I demonstrate the ubiquitous expression of *Mdmd* mRNA throughout embryonic development. This suggests that *Mdmd* acts at a very early embryonic stage and that it needs to be continuously active in embryos to sustain male development. Sharma et al. (2017) showed that targeted disruption of *Mdmd* turns genotypic males into females. Although this indicated that *Mdmd* plays a crucial role in male development, it did not proof that *Mdmd* is sufficient for male determination. To test whether *Mdmd* is solely sufficient to perform the male-determining function, in Chapter 4, I introduced *MdmdV* mRNA into early blastoderm stage embryos from the *MIII* strain and tested for sex-reversal. Transient expression of *MdmdV* mRNA in female embryos did not yield any masculinised flies, although an insignificant bias towards more males was observed in injected offspring. These results either indicate that expression of *MdmdV* alone is not sufficient to turn genotypic females into males, or alternatively, it is caused by an experimental shortcoming, i.e. insufficient translation of *MdmdV* mRNA. An alternative approach to determine whether expression of *Mdmd* is sufficient to turn genotypic females into males, would be to use *piggyBac* germline transformation to repeatedly express *MdmdV* during development. In Box 4.1, I describe how I constructed a pBac[3×P3-EGFP, hsp70-*MdmdV*] transgene. This transgene will be used in future experiments to assess the masculinising activity of *MdmdV*.
My work has shed light on the complex structure of the $M$-loci in the housefly and on the evolution of sex chromosomes in the housefly and in insects in general.