

## Cloning and Characterization of the *DMRT* Gene Homologue in the Sawfly, *Athalia rosae ruficornis* (Hymenoptera)\*

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The system of sex-determination has been well studied in *Drosophila melanogaster*. The *doublesex* (*dsx*) gene functions at the downstream-most of the sex-determining cascade, and the female- and male-specific proteins (DSX<sup>F</sup>, DSX<sup>M</sup>) regulate expression of genes engaged in sexual differentiation such as yolk protein genes (Burtis and Barker, 1989; Schutt and Nothiger, 2000). The *dsx* homologues have been identified in many species other than *Drosophila*: *Bmdsx* from *Bombyx mori*, *mab-3* from *Caenorhabditis elegans*, *DMRTs* from human, and *Dmrts* from mouse and chicken (Shen and Hodgkin, 1988; Ohbayashi *et al.*, 2001; Ottolenghi *et al.*, 2002; Smith *et al.*, 2002). These *dsx* homologues were considered to regulate sexual differentiation and have been evolutionary conserved as the sex-determining genes (Raymond *et al.*, 1998). A common characteristic of these gene homologues is the presence of a DNA binding domain termed the DM domain (Zhu *et al.*, 2000).

In the present study, we cloned and sequenced two *dsx* gene homologues from the sawfly, *Athalia rosae* (*ArDMRT1* and *ArDMRT2*). Reverse transcription PCR demonstrated that the *ArDMRT1* was expressed in embryos, larvae and pupae of both sexes, while the *ArDMRT2* was expressed only in embryos of both sexes. A molecular phylogenetic analysis demonstrated that *ArDMRT1* and *ArDMRT2* were more similar to homologous genes of vertebrates, rather than *dsx* of *D. melanogaster*. Recent progress of the genome research in the honey bee *Apis mellifera* revealed that there are some *dsx* homologues similar to that of *D. melanogaster*. Investigation of these genes in *A. rosae* is under way. We plan to analyze the function of these genes using the germline transformation system that we have recently established in *A. rosae* (Sumitani *et al.*, 2003).

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