Evolution of Developmental Control Mechanisms

Boule-like genes regulate male and female gametogenesis in the flatworm Macrostomum lignano

Georg Kuales a, Katrien De Mulder a, b, Jade Glashauser a, Willi Salvenmoser a, Shigeo Takashima c, Volker Hartenstein c, Eugene Berezikov b, Walter Salzburger d, Peter Ladurner a, *

a University of Innsbruck, Institute of Zoology and CMBI, Technikerstrasse 25 A-6020 Innsbruck, Austria
b Hubrecht Institute and University medical Center Utrecht, Uppsalalaan 8, 3584 CT Utrecht, The Netherlands
c University of California Los Angeles, Department of Molecular, Cell and Developmental Biology, 621 Charles E. Young Drive, East Boyer Hall 559, CA 90095-1606 California, USA
d University of Basel, Zoological Institute, Vesalgasse 1, CH-4051 Basel, Switzerland

ARTICLE INFO

Article history:
Received for publication 21 May 2010
Revised 17 June 2011
Accepted 20 June 2011
Available online 28 June 2011

Keywords:
Boule
Planaria
Flatworms
Platyhelminthes
Germline
 Hermaphrodite

ABSTRACT

Members of the DAZ (Deleted in AZoospermia) gene family are important players in the process of gametogenesis and their dysregulation accounts for 10% of human male infertility. Boule, the ancestor of the family, is mainly involved in male meiosis in most organisms. With the exception of Drosophila and C. elegans, nothing is known on the function of boule in non-vertebrate animals. In the present study, we report on three boule orthologues in the flatworm Macrostomum lignano. We demonstrate that macbol1 and macbol2 are expressed in testes whilst macbol3 is expressed in ovaries and developing eggs. Macbol1 RNAi blocked spermatocyte differentiation whereas macbol2 showed no effect upon RNAi treatment. Macbol3 RNAi resulted in aberrant egg maturation and led to female sterility. We further demonstrated the evolutionary functional conservation of macbol1 by introducing this gene into Drosophila bol1 mutants. Macbol1 was able to rescue the progression of fly meiotic divisions. In summary, our findings provide evidence for an involvement of boule genes in male and female gamete development in one organism. Furthermore, boule gene function is shown here for the first time in a lophotrochozan. Our results point to a more diverse functional assignment of boule genes. Therefore, a better understanding of boule function in flatworms can help to elucidate the molecular mechanisms of and concomitant infertility in higher organisms including humans.

© 2011 Elsevier Inc. All rights reserved.

Introduction

The production of gametes that carry all the genetic information of their genitors by means of meiosis is a crucial step to ensure the generation of fertile offspring. Meiosis is a complex process of the genetic regulation that is not entirely understood (Handel and Schimenti, 2010; Keeney, 2009a, b; Maines and Wasserman, 1998; Orr-Weaver, 1995), despite the remarkable similarity in the generation of male gametes between e.g. Drosophila and mammals (Fuller, 1998).

The Deleted in AZoospermia (DAZ) gene family is known to be crucial for the development of the germine and the generation of male and female gametes (Brook et al., 2009; Kee et al., 2009; Kerr and Cheng, 2010; Takeda et al., 2009; VanGompel and Xu, 2010; Xu et al., 2001). Interested in the background of male infertility, Tiepolo and Zuffardi (1976) proposed the existence of an AZoospermia Factor (AZF) or a spermatogenesis gene that, when not working properly, could lead to a breakdown of sperm production. The first member of the DAZ family was isolated from deleted Y-chromosomal regions in human and was believed to be a strong candidate for AZF (Ma et al., 1993; Reijo et al., 1995). Later on, three genes belonging to the DAZ Family, DAZ, DAZL (DAZ-like) and Boule, proved to be indispensable for gametogenesis (Carani et al., 1997; Cooke et al., 1996; Eberhart et al., 1996; Houston et al., 1998; Houston and King, 2000; Johnson et al., 2001; Karashima et al., 2000; Liu et al., 2009; Maegawa et al., 1999; Ruggiu et al., 1997; Xu et al., 2003; Zhang et al., 2009). The name-giving family member DAZ (Reijo et al., 1995) emerged as a Y-linked duplicate of the autosomal DAZL about 30 million years ago in catarrhine primates (old world monkeys) (Xu et al., 2001). The second family member, DAZL, arose by duplication from boule in an ancestor of vertebrates more than 450 million years ago (Xu et al., 2009). Boule was first identified in Drosophila melanogaster (Castillon et al., 1993; Eberhart et al., 1996), and is considered the ancestor of the DAZ gene family (Haag, 2001; Xu et al., 2001).

The high evolutionary conservation of orthologues was demonstrated by rescue experiments: Spermatogenesis of sterile fruit flies carrying a bol1-mutation was partially restored by introducing human
**BOULE** (Xu et al., 2003). Interestingly, male sterility in a hybrid bovine species could be caused by reduced *boule* expression although no direct evidence has been provided (Zhang et al., 2009). *Boule* was also shown to have an additional function in the nervous system of *D. melanogaster*. Joiner and Wu (2004) reported a defect in the fly eye when Boule protein was overexpressed. Hooppfer et al. (2008) showed that the protein was a negative regulator of axon pruning. All family members are characterized by a RRM (RNA Recognition Motif) containing RNP-1 and RNP-2 sequences (RiboNucleoProtein binding sites 1 and 2) and a DAZ-motif consisting of one (*boule* and **DAZL**) or more (**DAZ** **DAZ** tandem-repeats (Reijo et al., 1995).

Phylogenetic analyses support the existence of two subfamilies, one primarily responsible for early germ cell development (containing **DAZ** and **DAZL**), and the other (**Boule**) with a focus on directing meiosis (Xu et al., 2001). Whereas **DAZ** can only be found in the male function (Eberhart et al., 1996; Menke et al., 1997; Reijo et al., 1995, 2000). **DAZL** is required in male and female gonads (Cooke et al., 1996; Saxena et al., 1996). Boule was initially thought to be restricted to spermatogenesis throughout the animal kingdom, until its orthologue in *C. elegans* showed that gene family members are involved in the transition from embryonic germ cells to primordial germ cells (Cuffman et al., 2005; Kee et al., 2009). However, nothing is known on the expression and function of **boule**-like genes in more basal organisms.

Sexually reproducing flatworms are especially suitable to study germ cell differentiation. The exceptional regenerative capacity of planaria and other flatworms allows them to reconstitute germ cells from somatic stem cells during regeneration (De Mulder et al., 2009, 2010; Egger et al., 2006; Handberg-Thorsager and Salo, 2007; Newmark et al., 2008; Pfister et al., 2008; Sato et al., 2006; Wang et al., 2007; Zayas et al., 2005).

All studies so far have been performed on vertebrates or in *D. melanogaster* and *C. elegans*. In humans, it has been shown that **DAZ** gene family members are involved in the transition from embryonic stem cells to primordial germ cells (Cauffman et al., 2005; Kee et al., 2009). However, nothing is known on the expression and function of **boule**-like genes in more basal organisms.

The flatworm *M. lignano* is an emerging model system in developmental and evolutionary studies (De Mulder et al., 2009, 2010; Egger et al., 2006; Handberg-Thorsager and Salo, 2007; Newmark et al., 2008; Pfister et al., 2008; Sato et al., 2006; Wang et al., 2007; Zayas et al., 2005). However, nothing is known on the expression and function of **boule**-like genes in more basal organisms.

Sexually reproducing flatworms are especially suitable to study germ cell differentiation. The exceptional regenerative capacity of planaria and other flatworms allows them to reconstitute germ cells from somatic stem cells during regeneration (De Mulder et al., 2009, 2010; Egger et al., 2006; Handberg-Thorsager and Salo, 2007; Newmark et al., 2008; Pfister et al., 2008; Sato et al., 2006; Wang et al., 2007; Zayas et al., 2005).

The flatworm *M. lignano* is an emerging model system in developmental and evolutionary studies (De Mulder et al., 2009, 2010; Egger et al., 2006; Handberg-Thorsager and Salo, 2007; Newmark et al., 2008; Pfister et al., 2008; Sato et al., 2006; Wang et al., 2007; Zayas et al., 2005). However, nothing is known on the expression and function of **boule**-like genes in more basal organisms.

**Whole-mount in situ hybridisation (ISH)**

In *in situ* riboprobes were generated using the DIG RNA labelling KIT SP6/T7 (Roche). The PCR templates were used for probe synthesis were amplified with the primers 5′-CTCTGCGACTGAGACAGTAA-3′ and 5′-
Fig. 1. Alignment of DAZ family RRM domains and DAZ-repeats of different species, comparative protein domain organisation, and phylogenetic analyses of RRM sequences. (A) Alignment of DAZ family RRM sequences and of DAZ-repeats; the highly conserved motifs RNP-1 and RNP-2 are shaded in pink. The two amino acids specific for all boule orthologues are highlighted in yellow. Species abbreviations: Hs, Homo sapiens (DAZ: AAB02393; DAZL: NP_001342; bol: AAK58689); Mm, Mus musculus (DAZL: AAH99940; bol: AAK69026); Gg, Gallus gallus (DAZL: NP_989549; bol: XP_421917); Xl, Xenopus laevis (AAH79768); Dr, Danio rerio (CAM56544); Og, Oryzias latipes (DAZL: NP_001098269; bol: ACU31026); Dm, Drosophila melanogaster (Q24207); Ce, Caenorhabditis elegans (AAA67839); Sm, S. mansoni (XM_002575473); Macbol1, Macbol2 and Macbol3 and human Hsbol. (C) Phylogeny of DAZ family members using conserved RRM sequences as depicted in (A); the tree is based on a Bayesian inference analysis with Mr. Bayes. Numbers above the branches indicate Bayesian posterior probabilities, and numbers below the branches are maximum likelihood bootstrap values (100 replicates; WAG + f model); note that all three Macbol genes are located within the boule-clade, clustered together with the parasitic flatworm S. mansoni but are split up according to their role in male or female gametogenesis.
Preparation of in situ hybridisation sections

Macbol2 and macbol3 RNAi experiments and consecutive whole mount in situ hybridisations on treated and control animals were performed as described. After visualisation with NBT/BCIP, specimens were washed in PBS and fixed in BOUIN’s fluid or 2.5% glutaraldehyde in PBS, dehydrated in standard ethanol series and embedded in Spurr’s low viscosity resin (Spurr, 1969). Serial sections were assembled using a Diamond Histo Butler knife (Diatome, Switzerland), mounted, and examined with a Leica DM5000B microscope.

Immunocytochemistry

Macbol1 antibody was raised in rabbit against the peptide CCRGDRPGLKMYMDH (aa 398–412) by GenScript (New Jersey, USA). MSP-1 monoclonal antibody was used to stain spermatids (Ladurner et al., 2005a). Specimens were incubated with both antibodies in single and double staining at 4 °C overnight in diluted 1:1800 (Macbol1) and 1:200 (MSP-1) in BSA-T (PBS with 1% albumin fraction V (Rotth) and 0.1% Triton X-100). Secondary antibodies, swine anti-rabbit FITC-labelled (Dako) and goat anti-mouse Rhodamine-labelled (Rockland), diluted 1:200 and 1:500 in BSA-T, respectively, were applied for 1 h at RT. Photos were taken using either a Leica DM5000 light microscope or a Zeiss LSM 510.

Immunohistochemistry

EnVision + System (Dako) colour development was applied in preparation for semithin sectioning. M. lignano were relaxed in 7.14% MgCl2 for 15 min and fixed in PFA-sucrose (4% paraformaldehyde + 9% sucrose) for 30 min. Specimens were washed 3 × 5 min with PBS-T (PBS with 0.1% Triton X-100) and incubated in peroxidase-blocking reagent (Dako) for 10 min. Blocking of non-specific binding sites was done with BSA-T for 30 min, and primary antibody (Macbol1) was applied and diluted 1:1800 overnight at 4 °C. Visualisation was performed using DAB+ as chromogen (Dako), according to the manufacturer’s protocol. Dehydroxylation was done with standard ethanol series. Specimens were then embedded in Spurr’s low viscosity resin (Spurr, 1969). Blocks were trimmed and sectioned serially on a Reichert Autocut 2040 microtome with a Diamond Histo Butler knife (Diatome), yielding complete series of 1.0 μm sections.

Electron microscopy

After three weeks of RNAi, two adult specimens of each treatment group (macbol1, macbol2, and macbol3) and two control worms of the same age were relaxed in 7.14% MgCl2 and fixed in glutaraldehyde in 0.1 M cacodylate buffer (pH 7.4) for 1 h at 4 °C. Postfixation was performed for 1 h at RT in 1% osmium tetroxide in 0.05 M cacodylate buffer after several washing steps with buffer. Additional washing steps with buffer were done prior to dehydration in ascending ethanol series and embedding in Spurr’s low viscosity resin (Spurr, 1969).

Specimens were cut with the prototype of a Butler diamond knife (Diatome), yielding complete series of 1.0 μm sections that were mounted on glass slides. After drying for 2 h at RT, one section series of each experimental group was stained with Richardsions’ methylene blue Azur II resin (Richardson et al., 1960), mounted with cedar wood oil and examined for control with a Leica 5000B light microscope. One wild type animal and two animals of each RNAi group were cut with a diamond knife on an Ultracut S ultramicrotome (Leica), yielding complete ultrathin section series. Subsequent double staining with uranyl acetate and lead citrate was performed. Specimens were examined on a ZEISS Libra 120 energy filter electron microscope. Photo-documentation and analysis were done using a 2 k Vario Speed SSCD camera (Droendle) and the iT EM software (TEM imaging platform, Olympus).

Fly injection, crossing and tests analysis

Injection of wild type Drosophila embryos with the macbol1 construct was performed by BestGene (USA). Backcrossing into the bol1-mutant background ([P(ry] + [T2]P] bol1] ry[506/TM3, ry[KR] Sb[1] Ser[1]]) and analyses of transformants were performed at the Department of Molecular, Cell and Developmental Biology, UCLA, USA, and at the Department of Ultrastructural Research and Evolutionary Biology at the University of Innsbruck, Austria. For analyses, fly testes were dissected and Macbol1 antibody staining was performed as previously described (Ladurner et al., 2005a). For double staining, DAPI was applied 1:1000 along with the secondary antibody, gDNA was purified with DNeasy Tissue Kit (QIAGEN) and fragments amplified with the PCR primers 5′-AGCAGATGCAGAATACACAG-3′ and 5′-CCTCAAAATACGCCGA- GAG-3′ for wild type fly boule, 5′-CGGACTCTGCTGTCCTCTC-3′ and 5′-GCCGCAAAAGTCTCAGAAG-3′ for bol1; 5′-CCTCTGGCGACTGAGCATGAA-3′ and 5′-GCCGGAGCGTAGTAAAGTATG-3′ and 5′-AACCCCGACGGTAAACG- CATC-3′ for macbol1; and 5′-GACTCTACCGAGAAGTACGG-3′. Specimens were cut with the prototype of a Butler diamond knife (Diatome, Switzerland). Blocks were trimmed and sectioned serially on a Reichert Autocut 2040 microtome with a Diamond Histo Butler knife (Diatome, Switzerland). Sections were counterstained with Richardsions’ dye (Richardson et al., 1960), mounted, and examined with a Leica DM5000B light microscope.

Postembryonic development and regeneration

To obtain exactly staged worms for analysis of postembryonic development eggs from adult M. lignano were collected and surveyed until the embryos hatched. Hatchlings were kept under regular culture conditions until they reached the desired time points. For regeneration experiments, adult M. lignano were cut between pharynx and testes to remove the gonads completely. Cut specimens were allowed to regenerate missing parts until the defined time points. Specimens were relaxed in 7.14% MgCl2, fixed and frozen in methanol at −20 °C and then processed for in situ hybridisation and Macbol1 antibody staining. During Macbol1 antibody staining, we analysed 34, 37, 39, 21, 22, 21, 29, 43, and 54 animals at days 1, 2, 3, 4, 5, 6, 7, 10 and 16, respectively, of postembryonic development. At the same time, points 24, 23, 20, 46, 45, 40, 15, and 17 regenerates, respectively, were checked.

RNA interference (RNAi)

RNAi was performed during postembryonic development and regeneration for macbol1 and during homeostasis of M. lignano for macbol1, macbol2, and macbol3 by soaking the animals in a solution of dsRNA (3.0 ng/μl) in 1/2 culture medium. dsRNA was generated with an in vitro transcription system (RiboMax™ Large Scale RNA Production System T7, Promega), using the following primer sets: 5′-GGATCCTTAATACGACTCACTATAGG-3′ and 5′-GCCGGAGCGTAGTAAAGTATG-3′.
3′ + 5′-GGATCCCTAATACGACTCACTATAGGGAGATCTGATAC-3′ (macbol1), 5′-GGATCTATTAGGATCTATAGCTAGTGGAGTGGAGGCT-3′.

During RNAi, animals were incubated in 250 µl dsRNA in groups of 25, and supplied with food in 24-well plates. RNAi solution was changed once a day. In previous studies, it was shown that luciferase RNAi treated control animals did not show any mock effect (De Mulder et al., 2008; Sekii et al., 2009).

In the present study, control animals were only kept with 250 µl f/2 culture medium. All solutions were treated with ampicillin (1:1000) and kanamycin (1:1000) prior to application. During analyses of macbol1 RNAi, each batch of 25 worms/well was split into ten, 10 and 5 specimens for in situ hybridisation (ISH), antibody/spermatozoon staining and interference contrast (IC) microscopy, respectively. After three weeks, two adult worms each, treated and control, were prepared for transmission electron microscopy and only three analysed by interference contrast imaging. Macbol2- and macbol3-treated batches were only split in 20, 3, and 2 specimens for ISH, IC, and TEM, respectively.

Results

Identification and characterization of macbol genes

Full-length M. lignano boule paralogues macbol1, macbol2 and macbol3 were obtained from the M. lignano EST, genome and transcriptome databases. Macbol1, Macbol2 and Macbol3 have an ORF of 417, 503, and 282 amino acids, respectively, and comprise the two characteristic domains of DAZ family members, an RNA recognition motif (RRM) and the DAZ-repeat (Fig. 1A). The three boule genes show high sequence similarity within the conserved RNP-domains. In our alignment, two amino acids appeared characteristic for all boule proteins (Fig. 1A). In addition, we have performed phylogenetic analyses (Fig. 1C) that revealed two well-supported clades, a DAZL and a boule clade. The three boule sequences retrieved from M. lignano clearly clustered within the boule clade in all analyses (Fig. 1C). Further, there is only one DAZ-repeat present in Boule- and DAZL orthologues but not in DAZ proteins, which contain seven repeats in humans (Reijo et al., 1995). Accordingly, Macbol1, Macbol2, and Macbol3 possess one DAZ-repeat only, notably with only few conserved amino acids (Fig. 1A). However, a modest homology was also observed for the DAZ-repeat when human family members were compared with the fly homologue (Xu et al., 2001). When the cnidarian Nemastella vectensis, the parasitic flatworm Schistosoma mansoni, or the nematode C. elegans were included into the alignment, the DAZ-repeat showed only weak sequence homology (Fig. 1A). However, the overall protein domain organisation of Boule-like proteins is conserved and is consistent with the M. lignano boule domain structures (Fig. 1B). In summary, the conserved amino acids within the RRM, the structural assembly of its conserved domains, and the overall high amino acid identity of the RRM to other Boule proteins, specify M. lignano boule-like genes as true bouleorthologues.

Macbol1 is located in primary spermatocytes

M. lignano (Figs. 2A, B) possesses paired testes and ovaries at the lateral mid-body region. The testes are elongated oval structures (Fig. 2C), surrounded by thin somatic tunica cells and extra cellular matrix. The testis periphery consists of a small number of spermatogonia and numerous spermatocytes I+II (Fig. 2C). This peripheral region is most prominent at the anterior tip (the proliferative centre) that ends in a pointed anterior extension.

The outer region of the testis centre where maturing spermatids are located (Fig. 2C) and the testis periphery appear rather compact, although towards the lumen vivid mature spermatoozoa are present. They discharge into the vas deferens at the posterior end of the testes. Sperm is stored in the seminal vesicles within the tail plate.

To study macbol1 expression during homeostasis, we applied whole-mount in situ hybridisation (ISH) (Figs. 2D, D′) and antibody staining (Figs. 2E–I) in adult animals. Macbol1 ISH staining revealed a strong labelling within distinct cells at the testis periphery (Fig. 2D, D′). Specificity of ISH staining was confirmed by using macbol1 sense probe, which showed no signal.

Macbol1 protein was located in the cytoplasm of cells restricted to the periphery of the testes, as revealed by fluorescence labelling (Figs. 2E, E′) or horseradish peroxidase staining (Figs. 2F, F′). Histological semi-thin sections of the latter showed the presence of Macbol1 in spermatoocytes 1 in the testis periphery (Figs. 2G–I). Spermatoocytes 1 can be recognised by their nuclear morphology showing typical meiotic chromatin organisation representing leptotene or pachytene stages (Fig. 2G′). No signal was detected in spermatoegonia or in later stages towards the testis centre.

Macbol1 expression during postembryonic development and regeneration

Groups of worms were examined from one to 16 days post hatching (dph). First ISH signals appeared in five days old animals in a group of testis cells (Figs. S1E, E′). The staining intensified after seven dph (Figs. S1F, F′) and positive cells were localised to the testis periphery at 10 days (Figs. S1G, G′). The overall morphology of adult testes was accomplished at day 16 (Figs. S1H, H′). Macbol1 antibody staining revealed that no protein can be found in freshly hatched worms (n = 34) at one dph (Figs. S2A, A′). Macbol1 positive cells could be identified in four of 37 specimens at two dph (Figs. S2B, B′). We assume that the increased sensitivity of immunocytochemistry compared to ISH allowed the identification of the few Macbol1 positive cells in these early stages. This observation is in accordance with histological serial sections of two days old hatchlings that exhibit primary spermatocytes in the testis anlage (Salvenmoser, unpubl.). A cluster of two to four Macbol1 positive cells was present in all (39) three days old hatchlings (Figs. S2C, C′). The size of the cluster increased with the progression of development and remained a compact mass of cells until day five (Figs. S2C–E′). From days seven to 10 a central cavity became apparent (Figs. S2F–G′).

We next addressed the time course of the reappearance of Macbol1 positive cells during regeneration. M. lignano is capable to fully rebuild testes and ovaries from the remaining pool of stem cells after gonads were completely removed by amputation (De Mulder et al., 2009; Egger et al., 2006; Pfister et al., 2008). We analysed macbol1 mRNA expression (Fig. S1I–P′) and Macbol1 protein localization (Figs. S2I–P′) from one to 16 days post amputation (dpa). Macbol1 protein was not detected until day six after amputation. Depending on the level of amputation, the size until complete regeneration varies (Egger et al., 2006). In the present experiment Macbol1 protein and macbol1 mRNA were localised in the testes from day seven on (panels N–P′ in Figs. S1 and S2).

Macbol1 RNAi obstructs meiotic progression

We have performed macbol1 RNA interference knock-down during postembryonic development, regeneration, and in adult animals. We analysed macbol1 RNAi effects using macbol1 ISH (panels A–F′ in Figs. 3, S3, S4), Macbol1 antibody staining (panels G–I in Figs. 3, S3, S4), spermatid labelling (panels C–L′ in Figs. 3, S3, S4), and interference contrast microscopy (panels M–R in Figs. 3, S3, S4), after one week (Suppl. Fig. S3), two weeks (Suppl. Fig. S4), and three weeks (Fig. 4) of macbol1 RNAi treatment. In summary, our results demonstrate that macbol1 RNAi in every individual animal of all...
treatment groups effectively reduced macbol1 mRNA and led to a drastic deficiency in Macbol1 protein and a severe reduction in sperm production (for details see supplemental information). Macbol1 RNAi had no effect on the level of macbol3 expression (Suppl. 7I). Spermatocytes I, however, were arrested in response to macbol1 RNAi prior to the onset of macbol2 expression. Consequently, macbol2 ISH signals were not detected (Suppl. Fig. S7E).

We next analysed the effect of macbol1 RNAi on testis morphology and seminal vesicle content of adult animals. After three weeks of treatment, the overall size of the testes had not changed dramatically

Fig. 2. M. lignano morphology and macbol1-expression. (A) interference contrast (IC) image of an adult worm with corresponding scheme in (B); (C) IC image of adult testis with dotted lines bordering the different zones of spermatogenesis; (D, D’) macbol1 in situ hybridization; (E, E’) Macbol1 protein localisation using FITC-labelling; (F, F’) horseradish peroxidase staining; (G–I) histological semi-thin sections of Macbol1 antibody stained testis visualised using horseradish peroxidase (brownish); cell nuclei are counterstained with Richardson’s dye (blue); dotted red lines in the insets indicate cutting planes within the testis; solid diamonds (♦) indicate background in the glands surrounding the female opening; sc, spermatocytes; st, spermatids; sz, spermatozoa; t, testes; tc, testis centre; tp, testis periphery; scale bars 100 μm in A, B, D, E, F; and 20 μm in C, D’, E’, F’–I.
compared to control animals (Figs. 4A, D). However, squeezing preparations (n = 3) using interference contrast (IC) microscopy revealed that the testis periphery of control animals was normally sized, although it was greatly extended in RNAi animals. Furthermore, the testis centre in control animals was large, but it was small in size and devoid of mature spermatozoa after *macbol1* RNAi (Figs. 4B, E). In control animals, the false seminal vesicle (basically a widening of the vas deferens) and the true seminal vesicle (which is surrounded by...
Two adult animals were processed for transmission electron microscopy to further address the question of which cell types were present in the enlarged testis periphery in macbol1 RNAi treated animals. The testis periphery of wild type animals comprised one to two cell layers consisting of spermatogonia and spermatocytes I + II. Only the proliferative centre was expanded and led into the narrow anterior extension (Fig. 5A) containing mostly spermatogonia and few early spermatocytes I. The testis centre comprised spermatids on the outer margin and gradually maturing spermatids further inwards. Mature spermatocytes were in the testis centre (Fig. 5A). In contrast, macbol1 RNAi led to an accumulation of spermatocytes I and a drastically enlarged testis periphery (Fig. 5B). The proliferative centre appeared unaffected. The small-sized testis centre did not contain any differentiated stages of spermatogenesis but debris of disintegrated cells (Fig. 5B and see below). We therefore conclude that macbol1 RNAi interferes with the progression of spermatogenesis.

We next determined which cell types in the testes were specifically affected by macbol1 RNAi treatment. Consequently, we compared the ultrastructure of successive stages of spermatogenesis (Fig. 6). It became apparent that spermatogonia (Figs. 6A, B) and early spermatocytes I (Figs. 6C, D) showed an unaffected morphology in macbol1 RNAi treated animals. Late spermatocytes I, however, exhibited a modified fine structure and showed clear signs of degeneration (Figs. 6E–H). They acquired a change in nuclear morphology with lobes and foldings. Therefore, in ultrastructural sections, cytoplasmic inclusions can be seen within the nucleus (Fig. 6F), marking the onset of morphological degeneration. Cells beyond the spermatocyte I stage appeared disintegrated and carried a highly aberrant cellular content with membrane fragments (Fig. 6H) and a fragmented nucleus.

Detailed electron microscopical analyses revealed that macbol1 RNAi had no detectable effect on the morphology of cells and tissues such as nerve cells (Suppl. Figs. S5A–H), oocytes (Suppl. Figs. S5I, J), epidermal cells and muscle cells (Suppl. Figs. S5K, L), rhabdite forming cells (Suppl. Figs. S5M, N), gut cells and their associated cilia (Suppl.
Figs. S6A–D), mucus gland cells (Suppl. Figs. S6E, F), or protonephridial cells (Suppl. Figs. S6G, H). Finally, we determined reproductive success of the adult control versus RNAi-treated animals (n=25/group) by counting freshly hatched worms. After three weeks of macbol1 RNAi, controls had fathered 30 but treated animals only nine offspring.

Macbol1 shows partial functional equivalency to Drosophila boule

To address the evolutionary conservation of macbol1 function we introduced macbol1 into bol1-mutant Drosophila. We prepared a macbol1 carrying construct (Fig. 7A) for injection and generated a macbol1-transgenic bol1-mutant Drosophila strain. Testes (n=10 each) from wild type, mutant and rescued/uniFB02ies were dissected and analysed by PCR, DAPI and Macbol1 antibody staining. PCR experiments with primers specifically detecting macbol1, D. melanogaster boule, the bol1 mutation, or GAPDH were performed. These data corroborated that the rescue-phenotype could only result from the introduced flatworm gene (Fig. 7B). In wild type testes, interference contrast microscopy (Fig. 7C1) and DAPI staining (Figs. 7D1, E1) revealed numerous bundles of elongated spermatozoa and single sperm in the seminal vesicle. Macbol1 protein was not detected in wild type testes (Figs. 7F1–H1). In bol1 mutant testes no stages of spermatogenesis were observed (Figs. 7C2–E2) and Macbol1 was absent (Figs. 7F2–H2). In macbol1 rescue flies, bundles of sperm were observed in the testes (Fig. 7C3). However, the number was lower compared to wild type animals. Furthermore, in the proximal region of the testis tubule sperm nuclei were observed (Fig. 7D3). The shape of sperm nuclei was different from the wild type condition in that they were broader and not as sharply pointed (Fig. 7E3). In a distal testis area, Macbol1 positive cells were identified (Figs. 7F3 arrowhead; G3), probably accounting for spermatocytes. Tails of maturing spermatids and sperm were also labelled (Figs. 7F3 arrows; H3).

Macbol2 is expressed in spermatocytes II and spermatids

Whole mount ISH and histological sections revealed that the second M. lignano testis-specific gene, macbol2, was localised to spermatocytes II and spermatids (Figs. 8A–D). Spermatocytes II can be recognised by a large nucleus surrounded by a broad rim of cytoplasm (Fig. 8B). Spermatids are located more to the centre of the testis and show nuclear condensation and elongation of the cell. The proliferating centre and the anterior extension – regions that contain spermatogonia and spermatocytes I – lack any ISH signal (Figs. 8B–D). Likewise, the testis periphery

Fig. 5. Comparison of testis morphology of wild type and macbol1 RNAi treated adult animals by transmission electron microscopy. (A) The wild type testis centre is filled with late stages of spermatogenesis; The testis periphery is a slim layer containing spermatogonia and spermatocytes I, and only the anterior proliferating centre is expanded; (B) macbol1 RNAi leads to a drastic expansion of the testis periphery; the testis centre contains only degrading cells; ae, anterior extension; ci, cilia; ep, epidermis; g, gut; o, ovary; pc, proliferative centre; rh, rhabdites; tc, testis centre; tp, testis periphery; scale bars 50 μm.
and the testis centre did not show any macbol2 staining (Fig. 8D). After three weeks of macbol2 RNAi treatment, no macbol2 expression could be detected (Figs. 8E, S7F); however, no phenotype was observed. Accordingly, after three weeks, 20 offspring were counted for both, adult RNAi-treated and control worms. Macbol2 RNAi had no effect on the expression of macbol1 and macbol3 (Suppl. Figs. S7B, J).

**Macbol3 regulates oocyte maturation**

We found the third *M. lignano boule* ortholog to be expressed in oocytes and developing eggs (Figs. 8F–I). *Macbol3* expression was lacking in oogonia and early oocytes, although it was present in later oocytes that commenced vitellogenesis (Figs. 8G–I). *Macbol3* RNAi
resulted in macbol3 knock-down (Fig. 8J, S7K), but did not show any cross effects on macbol1 and macbol2 (Suppl. Figs. S7C, G). Macbol3 RNAi treatment resulted in aberrant egg maturation and caused female sterility with complete penetrance within the treatment group (Figs. 9A, E). After three weeks of macbol3 RNAi, treated animals had not produced any offspring compared to 27 in the control group. Early stages of oogenesis like oogonia and early oocytes I seemed not to be affected by macbol3 RNAi treatment compared to the wild type control. The oocytes of macbol3 treated animals showed typical meiotic figures including synaptonemal complexes. Later stages such as oocytes I in vitellogenesis, however, showed distinct changes in their ultrastructure. As a striking difference, these oocytes lacked the prominent rER compartments (Figs. 9B–H). In addition, the cells appeared smaller and less developed in their cytoplasmic maturation (such as yolk and egg-shell granule production). Finally, egg maturation could not be completed and eggs exhibited an abnormal phenotype. Animals ceased egg laying and they became female sterile hermaphrodites.

Discussion

Evolution of DAZ family proteins

Up to now, there is a complete lack of knowledge on the expression and function of boule-like genes in Lophotrochozoa. In M. lignano, we were able to identify three distinct boule paralogues. Previously, two potential boule paralogues have also been reported from the sea anemone N. vectensis. One of them perfectly matched a Boule protein.
consensus sequence (Shah et al., 2010), and the other showed both similarities and differences in critical positions. Nevertheless, there is the possibility that multiple boule paralogues – just as in our cricket model – do exist already in Cnidaria. However, there are no indications that multiple boule genes are present in Deuterostomia. In M. lignano all three reported boule paralogues match the consensus sequence of Shah et al. (2010). Our findings demonstrate that the paralogues corroborate the well-known function in male meiosis but also act during female gametogenesis. Nevertheless, the single and differently employed boule orthologues that we can identify today in various taxa like vertebrates, insects and nematodes do not necessarily represent the same ancestral gene. They could be the result of preservation of particular ancient paralogues and loss of the others. To follow up this hypothesis, more invertebrate species of different phyla, especially deuterostome invertebrates but also other cnidarian and flatworm species, should be investigated and potential boule sequences should be analysed and compared in detail.

Functional conservation of boule-like genes

Human BOULE’s importance in female embryos and the notion that it may have a role in primordial germ cell (PGC) migration (Kee et al., 2009), together with expression of fish boule in PGCs (Xu et al., 2009), suggest an early role of boule in female germ cell development of vertebrates. As in D. melanogaster and C. elegans, we could neither detect PGC-specific boule expression for any of the paralogues of M. lignano, nor an additional paralogue in its genome. Taken together, these findings could either hint at a loss of DAZ family mediated regulation of PGCs in...
Protopistomia or its evolution in Deuterostomia. In the latter scenario, however, the question of how an upstream regulatory function has been acquired remains elusive. However, although in fish and mice boule shows a certain degree of functional redundancy with Dazl (Li et al., 2011; VanGompel and Xu, 2010; Xu et al., 2009), in invertebrates, boule genes are indispensable for the first meiotic divisions (Eberhart et al., 1996; Karashima et al., 2000). Molecular evolution may be rapid in reproductive genes (Wyczółkowski et al., 2000), which also seems to be the case in DAZ family members that have undergone two consecutive duplication events (Bielawski and Yang, 2001; Haag, 2001; Tung et al., 2006). The cause may be positive Darwinian selection or a relaxation of functional constraints (Bielawski and Yang, 2001). Overall, DAZ family proteins act during gametogenesis in all organisms studied so far. They promote translation initiation of target mRNAs with only a short poly-A tail upon binding to their 3′-UTR. It has been shown that twine mRNA, the cd25 homologue in Drosophila is a potential substrate for boule (Maines and Wasserman, 1999). Twine or stringshows differential expression in male and female gonads of D. melanogaster (Alphey et al., 1992; Courtot et al., 1992; Sgrissi et al., 1995). In the M. lignano transcriptome (build 1000918) we have identified four twine or string cd25 phosphatases (RNA328_6352, RNA328_80531, RNA328_24244, RNA328_2512) that could be potential Macbol interaction partners. Furthermore, proteins such as PUM2 and PABP (Poly-Adenylation Binding Protein) in humans have also been shown to interact with Boule (Brook et al., 2009; Collier et al., 2005; Moore et al., 2003; Urano et al., 2005). Brook et al. (2009) gives a complete list of DAZ family protein interaction partners. According to a proposed model, interaction is mediated between the DAZ family protein’s RRM and DAZ repeat and the response elements in the 3′-UTR of targeted mRNAs (Collier et al., 2005; Meagawa et al., 2002). Like poly-A tails, DAZ-family proteins are able to recruit PABPs that would not be attracted sufficiently by short poly-A tailed mRNAs alone. As a consequence, the 5′- and the 3′-UTR can be bridged by the numerous proteins of the initiation complex, forming a “closed loop”, and can be joined by the ribosomal subunits. In general, the role of boule orthologues, i.e. to participate in the proper propagation of first meiotic divisions, seems to be conserved throughout the animal kingdom (Luetjens et al., 2004). This is underlined by the fact that both human BOULE (Xu et al., 2003) and macbol1 are able to substitute to the same extent for the defect allele in Drosophila. These cross-species rescues could indeed reflect conservation of boule function. Alternatively, biochemical interactions that are not based on evolutionary relation of boule genes could be responsible for the partial rescue. However, to fundamentally understand such rescue experiments, a detailed knowledge on the molecular function of boule proteins in these species would be required. When introduced into the invertebrate background, vertebrate Xdazl can also act during meiosis (Houston et al., 1998). It is thought that meiosis represents an evolutionary bottleneck during male gamete production (Xu et al., 2003), and boule and DAZL seem to be under similar functional constraints (Tung et al., 2006). Despite the number of identified interacting molecules, virtually nothing is known about factors that specifically direct DAZ family mediated stimulation into male and/or female gametogenesis in M. lignano and other organisms.

**Comparison of expression and function of macbol genes**

Concerning macbol1 and macbol2 expression, our data line up M. lignano with most other animals investigated so far, since boule genes are thought to be mainly involved in male meiosis (Cheng et al., 1998; Eberhart et al., 1996). Macbol1 mRNA and Macbol1 protein were exclusively found in primary spermatocytes whilstmacbol2 was expressed in secondary spermatocytes and early spermatids (Fig. 10). Likewise, boule-like genes have been shown to be vital players in the process of male gamete production in fly (Eberhart et al., 1996), bovineae (Zhang et al., 2009), mouse (VanGompel and Xu, 2010) and human (Luetjens et al., 2004; Xu et al., 2001). Moreover, BOULE isoforms were detected in the testes of four bat species (Yuan et al., 2009). However, in mice, boule is essential in the wake of meiosis for spermatid differentiation (VanGompel and Xu, 2010).

We now found macbol3 to be expressed in the ovaries and developing eggs of M. lignano. From the three boule genes that so far have been identified during oogenesis – the orthologues of medaka, rainbow trout, and C. elegans (Fig. 10) – only the nematode’s gene was demonstrated to be functionally refined to egg production. However, there was also faint staining in male gonads (Karashima et al., 2000; Otori et al., 2006). Thus, macbol3 represents the first boule homologue that, in expression and function, is specific to oogenesis only. It is therefore evident that different boule genes regulate male and female meiosis in M. lignano.

**Germline specificity of macbol paralogues**

The experimental accessibility of germ cell differentiation in flatworms (De Mulder et al., 2009, 2010; Egger et al., 2006; Handberg-Thorsager and Salo, 2007; Pfister et al., 2008; Sato et al., 2006; Seki et al., 2009; Wang et al., 2007; Zayas et al., 2005) during regeneration and postembryonic development renders these animals as model systems to study the function of DAZ family genes during the stem cell – germ line transition and germ cell differentiation. Due to M. lignano’s translucent body, its germline is amenable to experimentation and perfectly visible even in living adult specimens. Hence, the animal is used as a model for sex allocation studies – the differential investment into male and/or female function (Brauer et al., 2007; Janicke and Schärer, 2009b; Schärer, 2009; Vízoso and Schärer, 2007). It has previously been shown in M. lignano that an RNAI knock-down of mel2, a gene involved in spermatid differentiation, can be used to alter gonadal development (Seki et al., 2009). In contrast, genes such as wasa and piwi, although well characterized in germline development in many organisms, cannot serve as gonad specific markers in flatworms because they have also been shown to be present in somatic stem cells (De Mulder et al., 2009; Palakodeti et al., 2008; Pfister et al., 2007, 2008; Reddien et al., 2005; Rossi et al., 2006; Shibata et al., 1999). In M. lignano, our findings demonstrate the tight interdependency of macbol1 and meiotic divisions during spermatogenesis in adults during postembryonic development and...
regeneration. Our detailed ultrastructural analyses of germ cells and various other cells and structures did not reveal any role of macbol1 other than male meiotic control. Macbol1 RNAi led to male sterility whilst macbol3 RNAi resulted in female sterile animals. Therefore, macbol1 and macbol3 would provide versatile tools to artificially alter the amount of energy dedicated to gametogenic cascades and to address reproductive biology in *M. lignano*. As a corollary, this will also change our understanding of boule’s significance for general animal reproduction.

Conclusions

The recent success in sequencing the genomes of diverse invertebrate species will be the next step in answering important evolutionary questions. As we have demonstrated, the DAZ family homologues of *M. lignano* are true boule genes by primary sequence, domain structure and function. Further, we report for the first time different boule homologues that direct oogenesis and spermatogenesis in the same animal. Because *M. lignano* is the only invertebrate species where boule genes are essential for both gametogenic pathways, the animal is amenable for genetic manipulation, and there is an ongoing genome and transcriptome project that will provide access to potential Boule interaction partners, and the *M. lignano* boule paralogues are especially suited to study DAZ gene family evolution and the role of boule during male and female reproduction and (in)fertility.

Supplementary materials related to this article can be found online at doi:10.1016/j.ydbio.2011.06.030

Acknowledgments

The authors especially want to thank L. Mandal for help with the fly rescue experiment. Further, we want to thank F. Marx for support in gene isolation, A. Lusser for help with fly biology. This work was supported by an FWF grant 18095 to P.L. (Austria), an FWO grant to K.D.M. (Belgium), and SPA02-81-81/Plattwimmer to G.K.

References


