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Genome editing reveals *dmrt1* as an essential male sex-determining gene in Chinese tongue sole (*Cynoglossus semilaevis*)

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Zhongkai Cui^{1,2,3,*}, Yun Liu^{4,5,*}, Wenwen Wang¹, Qian Wang¹, Ning Zhang¹, Fan Lin¹, Na Wang^{1,2}, Changwei Shao^{1,2}, Zhongdian Dong¹, Yangzhen Li^{1,2}, Yingming Yang¹, Mengzhu Hu¹, Hailong Li¹, Fengtao Gao¹, Zhanfei Wei¹, Liang Meng¹, Yang Liu^{1,2}, Min Wei^{1,2}, Ying Zhu^{1,2}, Hua Guo^{1,2}, Christopher H. K. Cheng^{4,5,†}, Manfred Scharl^{6,7,†} & Songlin Chen^{1,2,†}

Chinese tongue sole is a marine fish with ZW sex determination. Genome sequencing suggested that the Z-linked *dmrt1* is a putative male determination gene, but direct genetic evidence is still lacking. Here we show that TALEN of *dmrt1* efficiently induced mutations of this gene. The ZZ *dmrt1* mutant fish developed ovary-like testis, and the spermatogenesis was disrupted. The female-related genes *foxl2* and *cyp19a1a* were significantly increased in the gonad of the ZZ *dmrt1* mutant. Conversely, the male-related genes *Sox9a* and *Amh* were significantly decreased. The *dmrt1* deficient ZZ fish grew much faster than ZZ male control. Notably, we obtained an intersex ZW fish with a testis on one side and an ovary on the other side. This fish was chimeric for a *dmrt1* mutation in the ovary, and wild-type *dmrt1* in the testis. Our data provide the first functional evidence that *dmrt1* is a male determining gene in tongue sole.

Sex-determining (SD) genes are located on the sex chromosomes to initiate a series of signaling pathways of sex related events to induce the development of bipotential primordial gonads into testes or ovaries. So far, a number of sex-determining genes have been identified in several vertebrate species¹. Besides the *SRY* gene in mammals^{2,3}, a member of the Sox family of transcription factors, several other genes that were previously known to act in the regulatory network of sex determination and gonad development have been found to function as master sex determining gene, e.g. *dmrt1*, *Amh* or its receptor and others^{4–8}.

Most of the identified sex determining genes are from species with a XY sex determination system, while much less is known about the sex determining genes in the ZW system. The Chinese half-smooth tongue sole (*C. semilaevis*) is a very important cultured marine flatfish with a ZW sex chromosome system^{9,10}. With respect to identifying the master sex determining gene, tongue sole is an ideal model because the full genome was sequenced¹¹, female specific AFLP and SSR molecular markers were discovered^{12,13}, and high density genetic linkage groups were developed^{14,15}. However, despite having a genetic sex determination by a well differentiated sex chromosome system, ~14% of ZW genetic females undergo sex-reversal to physiological males under normal rearing conditions, and relatively high temperature (28°C) can increase the sex reversal rate of genetic females to ~73%¹⁶, which is similar to the phenomenon observed in Australian bearded dragon¹⁷.

¹Key Lab of Sustainable Development of Marine Fisheries, Ministry of Agriculture; Yellow Sea Fisheries Research Institute, Chinese Academy of Fishery Sciences, Qingdao 266071, China. ²Laboratory for Marine Fisheries Science and Food Production Processes, Qingdao National Laboratory for Marine Science and Technology, Qingdao 266273, China. ³College of Fisheries and Life, Shanghai Ocean University, Shanghai 201306, China. ⁴School of Biomedical Sciences, The Chinese University of Hong Kong, Shatin, N.T., Hong Kong, China. ⁵School of Biomedical Sciences Core Laboratory, The Chinese University of Hong Kong Shenzhen Research Institute, Shenzhen 518057, China. ⁶Physiologische Chemie, University of Würzburg, Biozentrum, and Comprehensive Cancer Center Mainfranken, University Clinic Würzburg, D-97074 Würzburg, Germany. ⁷Texas A&M Institute for Advanced Studies and Department of Biology, Texas A&M University, College Station, Texas 77843, USA. [†]These authors contributed equally to this work. [‡]These authors jointly supervised this work. Correspondence and requests for materials should be addressed to C.H.K.C. (email: chkhcheng@cuhk.edu.hk) or M.S. (email: phch1@biozentrum.uni-wuerzburg.de) or S.C. (email: chensl@ysfri.ac.cn)

Interestingly, our recent study suggested that the sex chromosome of tongue sole was derived from the same ancestral vertebrate protochromosome as the avian W and Z chromosomes¹¹. Previous studies suggested that *dmrt1* located on the Z chromosome but absent from the W acts as a dosage-sensitive sex determining gene in chicken⁴. Also in *C. semilaevis* *dmrt1* is located on the Z chromosome and thus was considered to be an excellent candidate for the gene that determines male sexual development¹¹. Only a corrupted version of *dmrt1* is present on the W and gene expression studies further supported its function as a master SD gene. However, functional evidence has not been provided so far.

Genome editing methods employing zinc-finger nucleases (ZFNs), transcription activator-like effector nucleases (TALENs) and clustered regulatory interspaced short palindromic repeats (CRISPR/Cas9) have been successively established and applied in freshwater fish such as zebrafish, medaka, rainbow trout and Nile tilapia^{18–21} making an important addition to the already well developed toolbox of producing genetically modified fish. On the contrary, a similar methodology has not been established for marine fish due to the lack of a feasible embryo microinjection method. For a single species, the Nibe croaker, stable introduction of a green fluorescence protein (GFP) transgene by embryo microinjection was achieved²². In particular genome editing techniques have not been successfully applied to marine fish.

In this study, we have for the first time developed an efficient protocol for microinjection of flatfish embryos and then constructed TALEN plasmids and developed genomic editing methods for tongue sole. With these techniques, we produced *dmrt1* mutants. Results provide functional evidence that *dmrt1* is the male sex-determining gene in Chinese tongue sole.

Results

Development of embryo microinjection technique in flatfish. To develop a microinjection technique in flatfish we first used a reporter plasmid (*psmyd1:gfp*) where GFP is expressed under control of a muscle specific promoter. Zygotes were microinjected within ~40 minutes post-fertilization beginning at 1-cell stage and up to 4-cell stage embryos.

Successful injection was confirmed by expression of GFP in G0 fry. Of approximately 2000 injected embryos, nearly 500 embryos developed normally until gastrulation stage (~17 hpf). The survival rate of un-injected eggs is about 40%. At hatching stage, 46.2% (12/26) larvae showed clear GFP expression in skeletal muscle in accordance with the tissue-specificity of the *smyd1* promoter (Fig. 1B).

TALENs effectively induce mutations in the *dmrt1* gene of *C. Semilaevis*. To functionally test the role of *dmrt1* in sexual development of *C. semilaevis* a TALEN approach was used to eliminate *dmrt1* function. TALENs directed against exon1 sequences were injected. From 60,000 injected embryos, 500 survived until free swimming larvae stage and finally 65 adults were raised. In pooled microinjected embryos and larvae, mutagenic efficiency of *dmrt1* was about 55%. The *dmrt1* mutated samples in the *T7E1* tests were divided into two parts according to the 200bp and 150bp bands, respectively, recorded after *T7E1* digestion (Fig. 2B). From sequencing of embryos showing the target site to be mutated we recorded 10 different insertion and deletion events (Fig. 2C). Thus, the *dmrt1* TALENs used in this study are indeed suitable for targeted mutation of the *dmrt1* gene of *C. semilaevis* (Fig. 2).

The mutation frequency for survived TALEN sole was 50.77% (33/65). The 33 TALEN mutated sole consisted of 16 genetic females and 17 genetic males (Figure S2). There are a number of mutants with a mutation frequency ranging from 20% to 100% (Figure S3) and even different mutations were detected from the same individual. In general, the mutation rate of gonads was similar or even higher than that of somatic tissues (fins) (Figure S4).

***Dmrt1*-deficient tongue sole have a sex-reversed phenotype and display female gene expression patterns.** In wild-type fish, testes are short and thick with a “cashew nut” shape, whereas ovaries are long, thin and have a circular shape. In *dmrt1*-deficient genotypic males, the gonads presented an ovary-like gross appearance (Fig. 3A) being significantly longer than the control testes. Histological examination of the gonads from one-year-old *dmrt1*-deficient fish revealed an abnormal structure. Compared to wild-type males, only a few spermatogonia, and few or in some cases even no spermatocytes and spermatids were observed (Fig. 3B,a and b). In addition, *dmrt1*-deficient testes showed structures typically found in normal ovaries, like ovarian lamella and a central lumen (Fig. 3B,a). Intriguingly, we detected in the mutant gonads structures that appeared to resemble oogonia (Fig. 3B,b).

Dmrt1 gene expression analyses of mutant gonads revealed far less transcript abundance than samples from control males (Fig. 4 and Figure S5B). We then examined *foxl2* and *cyp19a1a* gene expression, which are expressed in ovary and essential for the maintenance of ovary differentiation^{23,24}. And *Sox9a* and *Amh* gene expression were also examined. Compared to control males, the expression of *foxl2* and *cyp19a1a* was significantly increased in the *dmrt1*-deficient males (Fig. 4 and Figure S5A and C). Conversely, the male-related genes *Sox9a* and *Amh* were significantly decreased. Interestingly, the expression of *foxl2* in the *dmrt1*-deficient males was even significantly higher than in wild-type females (Fig. 4 and Figure S5C) ($p < 0.05$).

The *dmrt1*-deficient males showed accelerated growth. Compared to wild-type males, the body weight of the *dmrt1*-deficient males was increased 2.5 fold ($p < 0.01$) (Fig. 5 and Table S2). The body width and length of the *dmrt1* deficient males were also larger than that of wild-type male fish (Fig. 5 and Table S2). These data indicate that disruption of *dmrt1* is connected to obvious changes in the growth characteristics of male *C. semilaevis*.

Chimeric *dmrt1*-mutant fish develop as intersex with a wild-type testis and a *dmrt1* mutant ovary. Among the 33 *dmrt1* mutated tongue soles we discovered a intersex with a testis on the up-side and an ovary on the down-side of the body (Fig. 6A). Histological examination demonstrated that the upside gonad has a typical testis structure with abundant spermatozoa, while the down-side gonad has a typical ovary structure and contains many oocytes (Fig. 6B). The genetic sex of the intersex was identified to be ZW by sex linked SSR marker analysis. Usually ZW genotypes develop as phenotypic females but some can develop as phenotypic males (pseudomales or

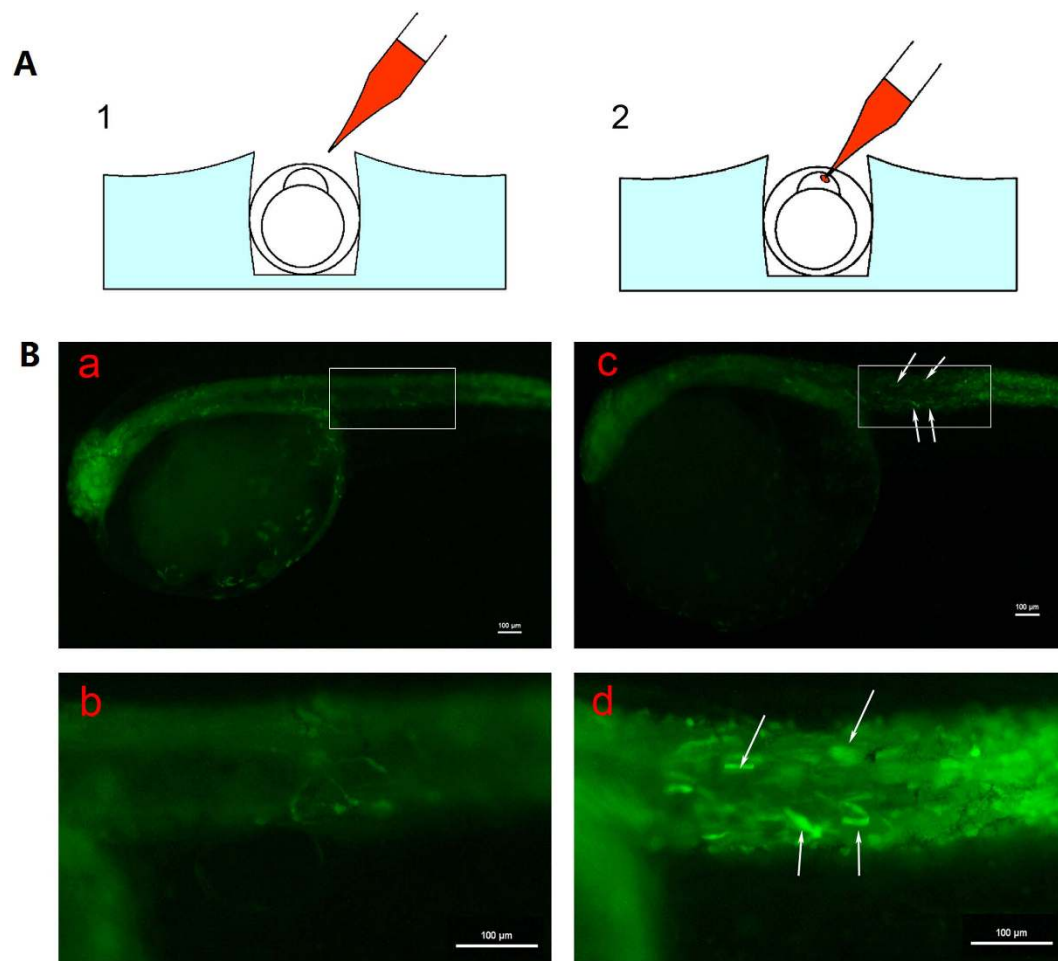


Figure 1. Microinjection method and transient expression of GFP in *C. semilaevis*. (A) Schematic drawing of microinjection. (1) A micropipette is passed through the chorion at an approximately 45-degree angle; (2) a droplet which is 1/10~1/5 of the cell volume is injected into the cell. (B) Transient expression of GFP in 2 dah *C. semilaevis*. (a) Control fish; (b) large magnification of frame area in (a); (c) transient expression of GFP in the muscles; (d) large magnification of frame area in (c) muscle fibres strongly expressing of GFP are indicated by arrows. Scale bar, 100µm.

neo-males). Notably, PCR products from fin and ovary of the intersex have the same abundance of the W and the Z derived fragment like in control females (Fig. 6C), while amplification from the testis gave a lighter W-specific band (Fig. 6C). As Z and W are of the same dosage in this fish the two sex linked SSR bands should be equal in DNA content. The weaker band for the W-linked marker might be due to the fact that sperm cells in the testis contain only Z-chromosomes because spermatocytes with a W-chromosome do not develop¹¹.

Sequencing of cloned *dmrt1* PCR products revealed *dmrt1* mutations in the ovary with a mutation rate of 56%. No *dmrt1* mutation was detected in the testis of the intersex (Fig. 6D and Figure S6). This shows that the hermaphroditic fish is a chimera for the TALEN induced *dmrt1* mutation with the wildtype part developing as a pseudomale with a normal testis while the mutant part develops towards female with an ovary as a consequence of *dmrt1* disruption. By qPCR, *dmrt1* expression was only detected in the testis, but not in the ovary (Fig. 6E and Figure S7A). And *foxl2* and *cyp19a1a* gene expression in ovary was significantly higher than in testis (Fig. 6E and Figure S7B,C), which is consistent with the expression pattern in wild-type testis and ovary (Fig. 4).

Discussion

Genome editing techniques have recently been widely applied in model fish such as zebrafish and medaka, and also several other freshwater fish, such as tilapia^{18,19,21}. In marine fish transgenic technologies have lagged behind because of the almost insurmountable problems presented by the general fragility of the embryos and high mortalities. In the present study, we developed a microinjection technique for flatfish embryos and successfully produced a gene knock out by TALEN mediated genome editing in a marine fish species.

Dmrt1 is a transcription factor that plays an important role in testis determination and differentiation in vertebrates^{5,11,25–27}. In the mouse, *dmrt1* null mutants have severely dysgenic testes in which both Sertoli cells and germ cells fail to differentiate properly after birth^{28,29}. In chicken where *dmrt1* is located on the Z chromosome³⁰, modulation of expression levels showed that this gene is necessary and sufficient for male development^{5,31}. In

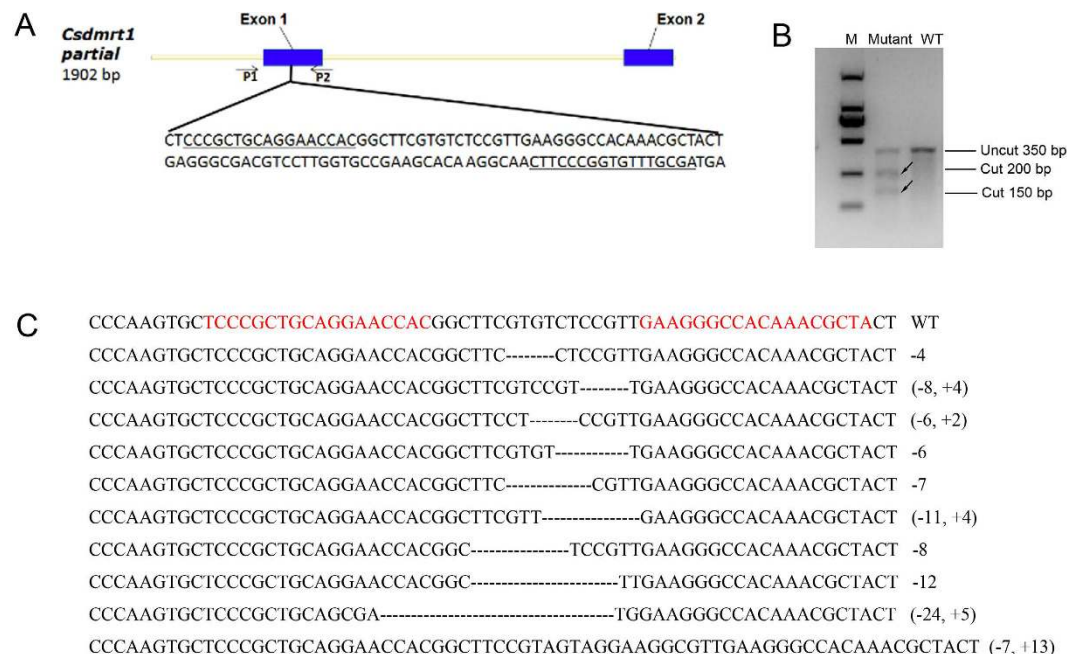


Figure 2. Genomic structure of the *C. semilaevis dmrt1* gene, design of *dmrt1*-TALENs and mutated sequences induced by TALENs. (A) *dmrt1*-TALENs designed to target exon 1 of the gene. (B) Detection of mutations in the injected embryos by *T7E1* digestion. (C) Mutant *dmrt1* sequences from TALEN injected embryos. The *T7E1* cut bands were recovered after gel electrophoresis and cloned for sequence analysis. “–” represents deletion of bp, “+” represents addition of bp. The numbers at the right side indicate the number of deleted or inserted base pairs.

several fish species, including medaka, where a duplicate of *dmrt1* is the master male sex determining gene³², over-expression of *dmrt1* resulted in masculinization, while knock-down or knock-out induced feminization^{33,34}. Here we show that *dmrt1* deficiency due to a TALEN generated gene knockout in *C. semilaevis* led to significantly compromised testis development and hypoplasia of the testes, and an overall macroscopic female-like structure of the gonad. We even detected oocyte structures in the mutant gonads indicating a high degree of feminization brought about by the lack of *dmrt1* expression. In addition, mutant *dmrt1* fish showed also feminized growth traits. As faster growth and higher body weight is a secondary sex character in tongue sole, this feature of the mutant fish is most likely a consequence of the sex reverted gonad and indicated that also the hormonal status of the mutant gonads was feminized.

In all previous experiments conducted so far in any vertebrate *dmrt1* deficiency was incompatible with normal differentiation and development of the testis. *Dmrt1* gene mutation of medaka, human and birds resulted in male-to-female sex reversal^{5,35,36}. A few spermatogonia were still observed in *dmrt1*-deficient testis of *C. semilaevis*, but few or even no spermatocytes and spermatids. The observation that in most individuals no full gonadal sex reversal was seen may be due to the fact that these fish were of the G0 generation and that they constitute a mixture of cells with homozygous and heterozygous mutant loci, and that even some wildtype cells could still be present. This possibility is supported by the detection of an intersex mosaic fish in the cohort of tongue soles that developed from the injected embryos.

So far, an intersex has not been discovered in tongue sole in nature. The intersex obtained in this study was of ZW genotype with the *dmrt1* gene being mutated in the ovary, but no *dmrt1* mutation in the testis. Under hatchery conditions genetic female soles with ZW sex chromosomes can spontaneously develop into phenotypic males, which are called pseudomales or neo-males¹¹. Thus, one explanation is that the intersex would be such a neo-male and would have developed with two testis, one on the up-side and one on the down-side. But due to the *dmrt1* mutation in the down-side part of the flatfish the gonad in this region developed into ovary.

Interestingly, we observed in the chimeric intersex the most complete sex reversal. This may be due to the genomic constitution of the pseudo-males, which are hemizygous for the Z-chromosome and thus have only a single copy of the *dmrt1* gene. In this situation the effect of the TALEN knockout of this gene should be more effective and thus apparent in the injected G0 individual.

In vertebrates, *dmrt1*, on the one side and *foxl2* and *cyp19a1a* on the other side are antagonistically expressed and cross-regulate each other during gonad development and in maintaining the identity of the adult ovary or testis^{37–39}. Consistent with numerous studies in other vertebrates^{21,40,41}, *foxl2* and *cyp19a1a* expression were significantly up-regulated and *Sox9a* and *Amh* expression were significantly down-regulated in *dmrt1*-deficient testes of *C. semilaevis* as a molecular signature of the feminization process in the mutant gonads.

In our previous studies real-time PCR analysis, methylation status across the differentially methylated region between males and females, and expression studies of *dmrt1* in normal testis and ovary of *C. semilaevis* revealed that *dmrt1* of *C. semilaevis* has convergent features that are compatible with a similar function determining male

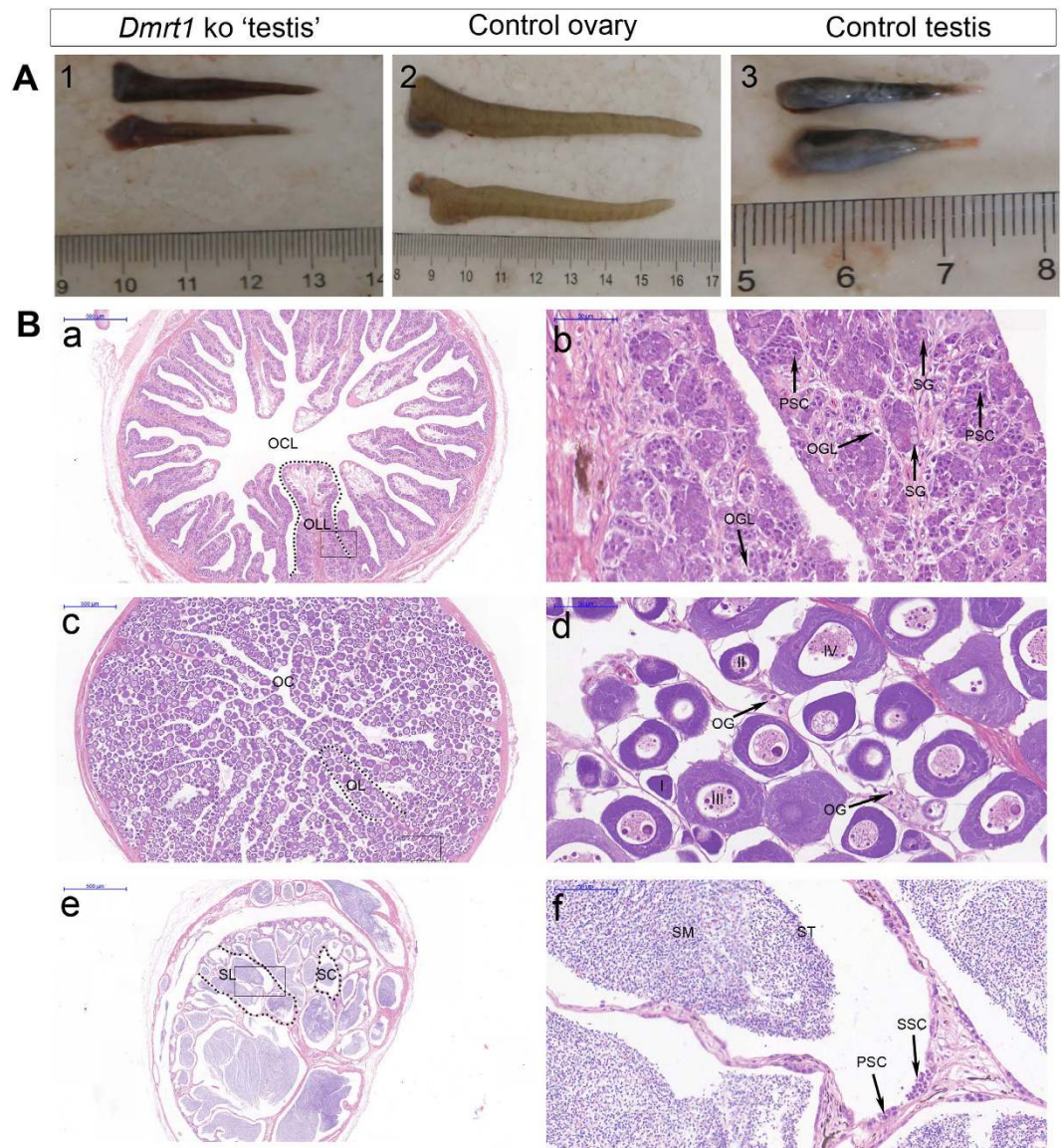


Figure 3. Effects of *dmrt1* disruption on gonad phenotype, sex differentiation. (A) Gross morphology of gonads from approximately one year old fish. (1) *dmrt1*-deficient 'testes'; (2) wild-type ovaries; (3) wild-type testes. (B) histology of gonads from approximately one year old fish. (a) *dmrt1*-deficient testis. The development of testis is ceased. The shape of the *dmrt1*-deficient testes in transverse sections is similar to control ovaries, and there are structures resembling ovarian cavity and ovarian lamella in the gonad of the mutant male fish. ovarian cavity-like (OCL), ovarian lamella-like (OLL); (b) large magnification of frame area in a. No secondary spermatocytes, spermatids and sperm are observed. oogonia-like (OGL), spermatogonia (SG) and primary spermatocytes (PSC). (c) Ovary of control female, including ovarian cavity (OC), ovarian lamella (OL); (d) large magnification of frame area in (c). Four stages of oocytes: stage I - IV and oogonia (OG). (e) Testis of control male. seminiferous lobuli (SL), seminiferous cyst (SC); (f) larger magnification of frame area in (e). Secondary spermatocytes (SSC), spermatids (ST) and sperm (SM). Scale bar is shown in the figures.

as in birds¹¹. The successful knock-out of the Z-linked *dmrt1* gene in tongue sole reported here, which resulted in female development, confirmed our previous hypothesis¹¹ that *dmrt1* is an important male-determining gene in *C. semilaevis*.

As a secondary effect, the mutant fish showed an enhanced growth as a typical female feature. As this is an important trait of high economic importance for the aquaculture industry, future studies will reveal the relationships between *dmrt1* as a primary male sex determining gene and its secondary functions on growth and body size.

In our study, we have developed methods for mRNA, DNA or anti-sense oligonucleotide microinjection into *C. semilaevis* embryos. The technique can be applied for studying many developmental processes *in vivo* and the functions of genes in flatfish. We successfully applied a genome editing technology to marine fish for the first time to study sex determination mechanism and to produce favorable economic traits, thus opening new avenues in the application of genome editing nucleases in both basic and applied research.

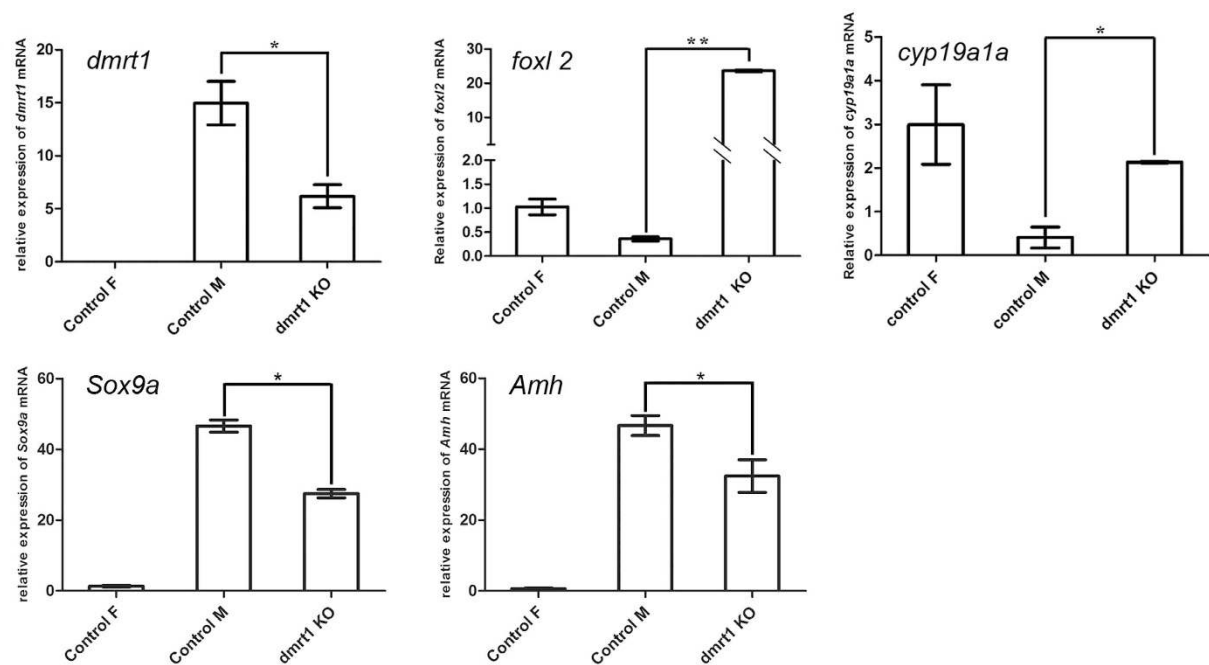


Figure 4. Gene expressions of sex differentiation markers on *dmrt1*-deficient gonad. *Dmrt1* and *foxl2* are key transcription factors in testicular and ovarian differentiation, respectively. *Cyp19a1a* encodes for aromatase which is responsible for estrogen production. *Sox9a* and *Amh* are male related genes. Relative mRNA expression of *dmrt1*, *foxl2*, *cyp19a1a*, *Sox9a* and *Amh* in *dmrt1*-deficient gonads at one year of age from one *dmrt1*-deficient and three wild-type gonads. β -actin was used for calibration.

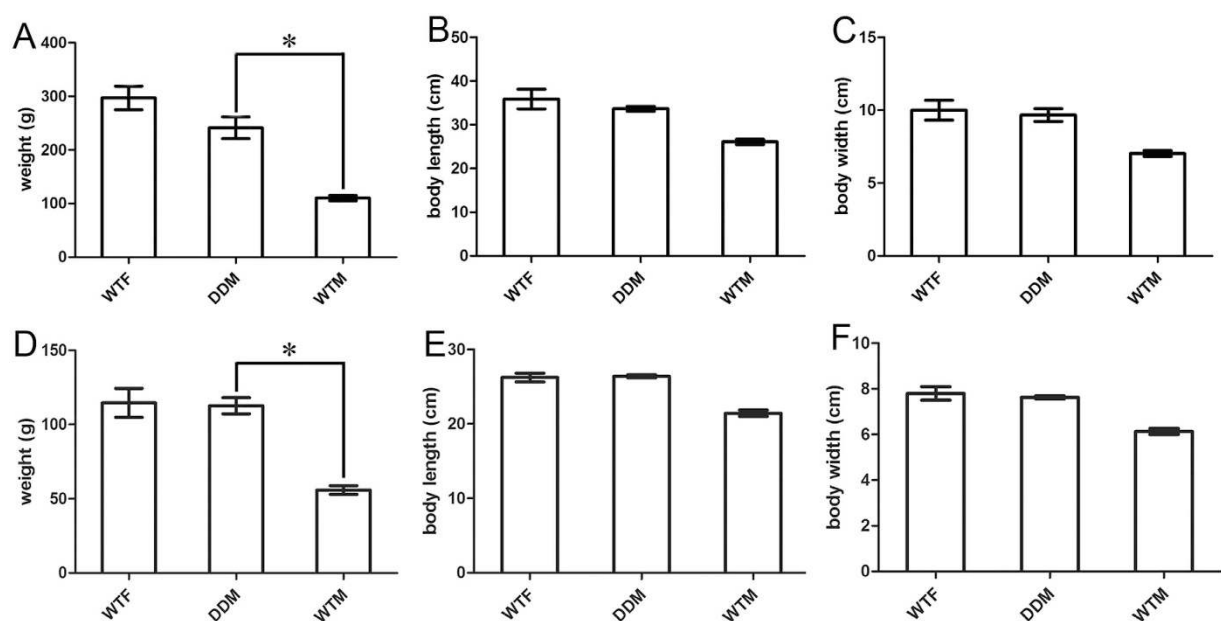


Figure 5. Growth parameters of *dmrt1*-deficient *C. semilaevis*. (A) Weight of *dmrt1*-deficient male *C. semilaevis* (DDM) in 2014 (1 year old) was significantly higher than of wild-type males (WTM) and similar to that of wild-type females (WTF). (B) Body length and (C) width of *dmrt1*-deficient *C. semilaevis* were larger than of wild-type male and close to the values of females. (D) Weight of *dmrt1*-deficient male *C. semilaevis* (DDM) in 2015 (8 months old) was also higher than of wild-type males (WTM) and similar to that of wild-type females (WTF). (E) Body length and (F) width of *dmrt1*-deficient *C. semilaevis* were larger than of wild-type male and close to the values of females.

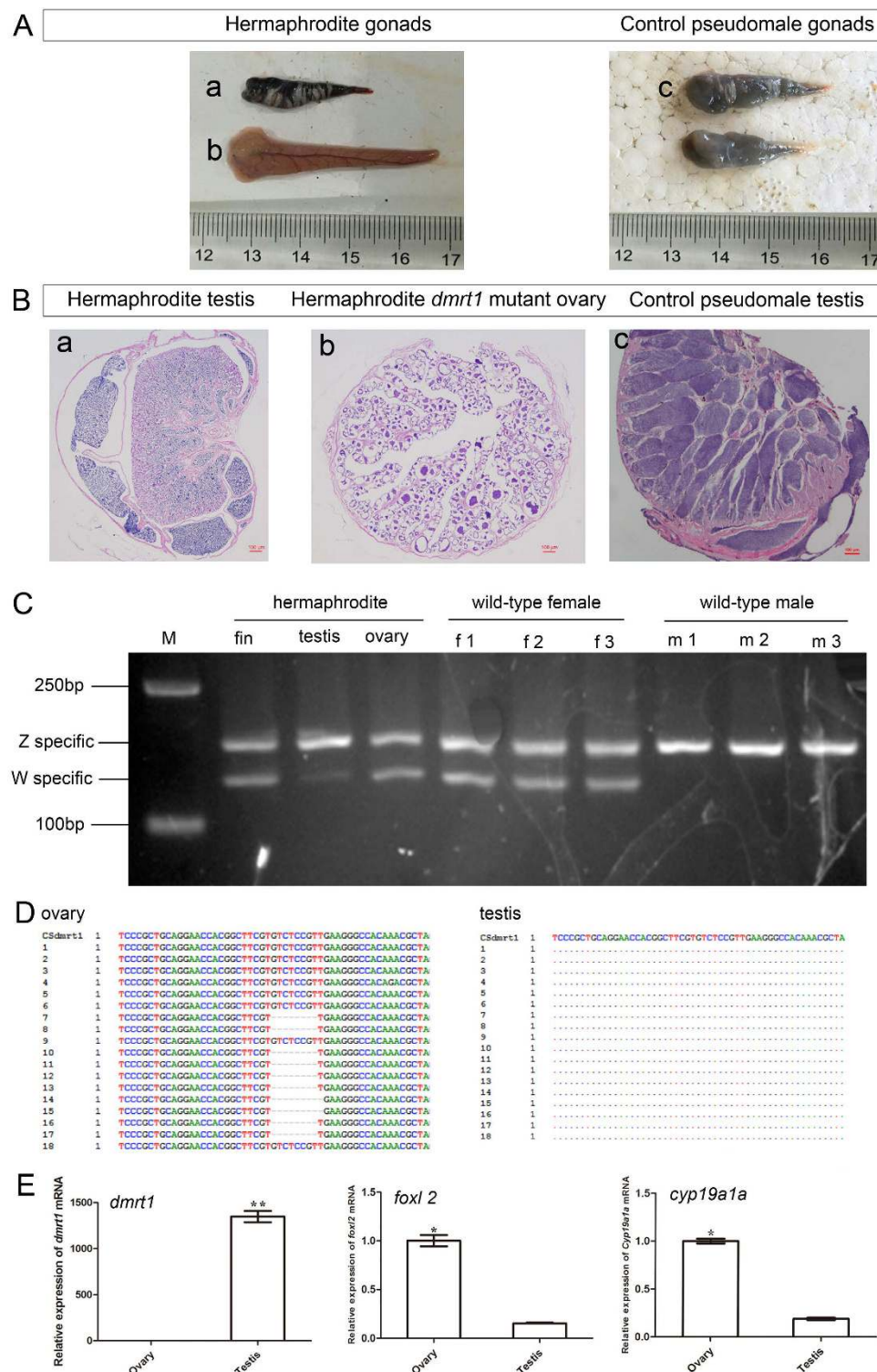


Figure 6. Effect of *dmrt1* disruption on intersex gonad phenotype, sex differentiation and gene expression.

(A) Phenotype of gonads in the intersex. (a) Testis shaped up-side gonad of the intersex; (b) ovary shaped down-side mutant gonad of the intersex; (c) testes of a pseudomale. (B) Histology of the gonads. (a) Testis of the intersex showing normal male structures; (b) *dmrt1* mutant gonad (ovary) of the intersex; (c) testis of a pseudomale. Scale bar, 100 μ m. (C) Determination of the genetic sex of the intersex by SSR PCR yielding different sized products for the Z (169 bp) and W (134 bp) chromosomes. (D) Sequences of wild-type and mutated *dmrt1* retrieved from ovary, partial; and only wild-type *dmrt1* in testis, partial. (E) Relative mRNA expression of *dmrt1*, *foxl2* and *cyp19a1a* in the *dmrt1*-deficient gonad at one year of age. β -actin was used as internal standard.

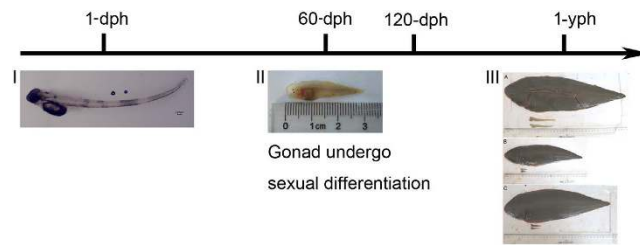


Figure 7. The Schedule of sampling. Sexual differentiation happens at approximately 60-dph. The tissues for gonadal histology and gene expression studies were sampled at about approximately 1-yph. (I) 1-dph *C. semilaepis*. (II) 60-dph *C. semilaepis*. (III) 1-yph *C. semilaepis*. (A) Wild-type female *C. semilaepis*; (B) wild-type male *C. semilaepis*; (C) *dmrt1*-deficient male *C. semilaepis*.

In conclusion, we have confirmed that *dmrt1* is an essential gene in determining male sexual development of *C. semilaepis*. To further show that it is the single master male sex-determining gene in tongue sole evidence could be provided that this gene is also sufficient to induce male development, e.g. by transgenic expression in WZ embryos. A situation in which *dmrt1* is the male sex determining gene in this fish analogous to the situation in birds will support the hypothesis (known as the “some chromosomes and some gene are better at sex” hypothesis) that certain genes and ancestral chromosomes are repeatedly evolutionary “selected” to serve as master regulators of genetic sex determination⁴².

Materials and Methods

Ethics statement. The collection and handling of the animals in the study was approved by the Chinese Academy of Fishery Sciences’ animal care and use committee, and all experimental animal protocols were carried out in accordance with the guidelines for the care and use of laboratory animals at the Chinese Academy of Fishery Sciences.

Experimental fish production and sampling. *C. semilaepis* were kept in the Haiyang High-Tech Experimental Base (Haiyang, China). The parental fish were strengthening reared and spawned by induction in seawater at 21–23 °C. Zygotes were obtained by artificial insemination and cultivated in sterilized seawater at 22–23 °C.

In Chinese tongue sole, sexual differentiation happens at ~60 days after hatching (60-dph). The tissues for gonadal histology and gene expression studies were sampled at about 1 year after hatching. Owing to the limited number of the injected fish obtained and the 60-dph gonads being sampled difficultly, we have to skip the 60-dph sample and only sampled at 1-yph (Fig. 7).

Development of microinjection techniques for tongue sole embryos. Because so far no methods are available for embryo microinjection in flatfish, the corresponding techniques had to be developed for tongue sole.

The *smgd1:gfp* plasmid (50 ng/μl) driving GFP expression by a 5.3-kb muscle-specific *smgd1* promoter fragment²² was microinjected into tongue sole embryo at the 1–4 cell stage using a microinjector system (PV820, WPI, USA). For the injection procedure, tongue sole embryos were held in troughs (width: 0.95 mm ± 0.05 mm, depth: 1 mm ± 0.05 mm) made with a plexiglass mould in 1% agarose and aligned using forceps. The chorion of tongue sole embryos is very tough, thus thick pointed injection needles were prepared using a P-97 Flaming/Brown micropipette puller (Sutter Co, USA) (Fig. S1). The microinjection method is shown in Fig. 1A and suppl 1. In order to avoid the occurrence of reflux during microinjection, the holding pressure of the injector was adjusted to 0.1–3 psi (pounds per square inch). To avoid injury of the zygotes by excessive compression, special care was taken that they were not in the troughs for more than 3 minutes. The injected zygotes were transferred from the troughs to aseptic seawater (2 μl 1% methyleneblue in 100 ml aseptic seawater). ~1000 embryos in a 500 ml beaker with aseptic seawater were placed in a 22 °C–23 °C incubator. The injected embryos were transferred from the incubator to 30 L glass fiber-reinforce plastic tank and hatched in 22 °C–23 °C filtered seawater. The inflating volume was 0.9 L/min–3 L/min. GFP fluorescence was observed and documented using a Nikon Eclipse 80i microscope (Nikon, Japan).

Design and construction of *dmrt1*-TALENs. The target sites were selected following routine rules and the TALENs for *Csedmrt1* were constructed using the Golden Gate method as described previously⁴³. Briefly, the modular plasmids recognizing each nucleotide were digested and ligated into the backbones of two middle array plasmids. Then the middle array plasmids and the last repeat plasmid were cloned into the backbones of the two optimized TALEN expression plasmids (the pCS2-TALEN-ELD and pCS2-TALEN-KKR) developed by the team of H. Cheng⁴⁴.

The TALEN target sites of *Csedmrt1* are located in exon 1 (Fig. 2A) with the following recognition sequences: left TALEN 5'-TCCCGCTGCAGGAACCAC-3' and right TALEN 5'-GAAGGGCCACAAACGCTA-3', leaving between the two binding sites a 17 bp spacer sequence for cutting by the *Fok I* nuclease (Fig. 2A).

TALEN mRNA preparation and microinjection. Plasmids were prepared using a plasmid midi kit (Omega, USA). pCS2-TALEN-ELD and pCS2-TALEN-KKR were linearized with *Not I* and recovered as transcription templates with a gel extraction kit (ZYMO RESEARCH, USA). *In-vitro* transcription was performed with the Sp6 mMESSAGE mMACHINE Kit (Ambion, USA). The mRNA was purified with the MEGAclear Kit

(Ambion, USA) and RNA concentration was determined with NanoVue plus (Thermo Scientific, USA). The TALEN mRNAs were mixed yielding a final concentration of each arm of 100 ng/μl. TALEN mRNAs were stored at −80 °C. Microinjection of 100–300 pg of TALEN mRNAs into one to four cell stage embryos was conducted as described above.

Determination of TALEN activity, mutation analysis, *dmrt1*-knockout fish screening and genetic sex identification. For an initial test of TALEN activity, injected embryos were collected at hatching stage. DNA was extracted with the TIANamp Marine Animals DNA Kit (TIANGEN, China) and quantified using NanoVue plus. A *dmrt1* fragment containing the TALEN target sites was amplified with primers *dmrt1*-TALEN-F and *dmrt1*-TALEN-R (Table S1). PCR, *T7E* I analyses, cloning and sequencing followed standard protocols (see Suppl 2).

Small pieces of tail fin from 6 months old injected *C. semilaevis* were used for mutant founder screening with the *T7E* I assay and sequencing as described in Suppl 2. According to the sequencing results, injected fish with high mutation rate were selected. Gonads were dissected from three *dmrt1*-deficient fish. The mutation rate of each gonad tissue was calculated as described in Suppl 2. Genetic sex of tongue sole was identified by PCR with the primers described previously⁴⁵.

Growth trait analyses of *dmrt1*-deficient *C. Semilaevis*. Body length and width of *dmrt1*-deficient *C. semilaevis* and control female and male fish were recorded. The weight was determined with an electronic scale.

Histological and RT-PCR analyses of *dmrt1* knockout individuals. Histological sections of *dmrt1*-deficient fish (1 year old) and control fish were performed. Gonads were dissected and fixed in Bouin's solution for 12 h~16 h at 4 °C, and stored in 70% ethanol. For sectioning, tissues were dehydrated and embedded in paraffin. Samples were serially sectioned at 6~8 μm thickness and stained using hematoxylin-eosin (HE).

Total RNA was extracted from gonads of *dmrt1*-deficient and control fish using RNAfast200 Kit (FASTAGEN, China). DNase I treatment and cDNA preparation for RT-PCR and real-time PCR were carried out according to the suppliers instructions (Takara) using 7500 Real-Time PCR System (Applied Biosystems, USA). *Dmrt1*, *foxl2*, *cyp19a1a*, *Sox9a* and *Amh* mRNA expression were determined by RT-PCR and real-time PCR in three replicates. *β-actin* was used as the internal control⁴⁶. Relative abundance of transcripts was calculated as $R = 2^{-\Delta\Delta C_t}$. Primer sequences used for RT-PCR and real-time PCR are listed in Table S1. In addition, a forward primer was designed from the target sequence of *dmrt1* TALEN for identifying mutant fish by lower abundance of the expected product.

Data analysis. Data were tested using one way ANOVA followed by Duncan multiple comparison tests with GraphPad Prism 5.0 (GraphPad, USA), significance was set at $p < 0.05$. Histograms were generated by GraphPad Prism 5.0 (GraphPad, USA).

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Author Contributions

S.C. and M.S. conceived the research; C.H.K.C. and Yun. L. designed and developed TALEN vectors. Z.C., W.W., Q.W., N.Z., F.L., Z.D., Yang. L., Y.Y., M.H., Z.W., Y.Z.L. and M.W. performed the experiments; N.W., C.S., H.L., F.G., L.M., Y.Z. and H.G. shared reagents; Z.C. and S.C. analysed the data; Z.C., M.S., S.C., C.H.K.C., Yun. L. and Q.W. wrote the paper.

Additional Information

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