A Novel Allotriploid Hybrid Derived From Female Goldfish × Male Bleeker’s Yellow Tail

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Hybridization is a traditional and effective strategy to alter the genotypes and phenotypes of the offspring, and distant hybridization is a useful strategy to generate polyploids in fish. In this study, goldfish (Carassius auratus, GF, 2n = 100) and Bleeker’s yellow tail (Xenocypris davidi Bleeker, YT, 2n = 48), which belong to different subfamilies, were crossed with each other. The cross of female GF × male YT successfully obtained hybrid offspring (GFYT hybrids), while the cross of female YT × male GF was lethal, and all the fertilized eggs stopped developing before the neurula stage of embryogenesis. All GFYT hybrids possessed 124 chromosomes (3n = 124) with two sets from GF and one set from YT. The measurable and countable traits of GFYT hybrids were identified, and the genetic characteristics of 5S rDNA between GFYT hybrids and their parents were also revealed. There were, respectively, four and three different 5S rDNA types in GF (assigned as GF-Ⅰ~Ⅳ) and YT (assigned as YT-Ⅰ~Ⅲ), and GFYT hybrids specifically inherited YT-Ⅰ and YT-Ⅱ 5S rDNA types from YT and GF-Ⅲ and GF-Ⅳ from GF. In addition, there were only testis-like and fat-like gonads found in GFYT hybrids. Interestingly, there were pyknotic and heteromorphous chromatin and invaginated cell membrane observed in the spermatids of testis-like gonads, but no mature sperm were found. Furthermore, TUNEL assays indicated that, compared with control, apparent apoptotic signals, which were mainly distributed around spermatid regions, were detected in the testis-like gonads, and the expression of apoptosis pathway-related genes including p53, bcl-2, bax, and caspase9 was significantly upregulated. Moreover, the expression of meiosis-related genes including spo11, dmc1, and rad51 showed an abnormally high expression, but mns1 and meig1, two key genes involved in the maturation of spermatid, were extremely downregulated. In brief, this is the first report of allotriploid via distant hybridization between GF and YT that possessing different chromosome numbers in vertebrates. The obtainment of GFYT hybrids not only harbors potential benefits and application in aquaculture but also further extends the understanding of the influence of hybridization and polyploidization on the genomic constitution of the hybrid offspring. Furthermore, they can be used as a model to test the origin and consequences of polyploidization and served as a proper resource to study the underlying mechanisms of spermatogenesis dysfunctions.

Keywords: distant hybridization, allotriploid, inheritance, recombination, sterility
INTRODUCTION

Hybridization, including intraspecific hybridization (crossing within a species as a line or strain) and distant hybridization (interspecific or above-specific hybridization) (Bartley et al., 2001; Liu et al., 2007), is the mating of genetically differentiated individuals or groups. It is a useful strategy to alter the genotypes and phenotypes of the offspring via transferring the genome of one species to another. Hybridization is thought to facilitate adaptive radiation and speciation in animals (Mallet, 2007; Abbott et al., 2013). By hybridization, the hybrid offspring may inherit the advantageous traits from two parental species, forming a new species with novel heterosis (Liu, 2010), exhibiting advantages in growth rate, disease resistance, viability, fecundity, yield, and appearance (Birchler et al., 2006; Hu et al., 2012). Compared with intraspecific hybridization, distant hybridization is apt to generate heterosis (Chen et al., 2018). In nature, hybridization exists widely in plants and animals, and it is an impetus for speciation and biological evolution (Barton and Hewitt, 1989). Numerous species have hybrid ancestry, presumably due to hybridization between closely related species (Schumer et al., 2016).

Polyploidization, the duplication of the whole genome, is an important driving force for the emergence of evolutionary novelties (Soltis and Soltis, 2009; Zhao et al., 2018). Hybridization, especially distant hybridization, plays vital roles in triggering polyploidization (Mallet, 2007). Polyploidization is an important and frequent event in lower vertebrate evolution, especially that of fish (Venkatesh, 2003), from the sharks to the higher teleosts (Leggatt and Iwama, 2003; Liu et al., 2007; Wang et al., 2019; Wang et al., 2021). In agriculture, hybridization and polyploidization are important breeding technologies. Many artificially induced polyploid fish have been used in aquaculture to produce sterility and to improve production (Liu et al., 2001; Liu et al., 2007). Furthermore, these polyploids are useful model systems to verify the theories about the origin and consequences of polyploidization (Comber and Smith, 2004).

Then, choosing the right or suitable parents and screening the hybrid offspring with improved characteristics becomes one of the main directions of fish genetic breeding (Wang et al., 2015). Goldfish (Carassius auratus Var., 2n = 100, GF), within the subfamily of Cyprinidae, genus Carassius which are commonly speculated as ancient polyploids (Braasch and Postlethwait, 2012; Knytl and Fornaini, 2021), presents the advantages of a colorful body and strong disease resistance. Bleeker’s yellow tail (Xenocypris davidi Bleeker, 2n = 48, YT), within the subfamily of Xenocyprinidae, possesses the benefit of excellent meat quality. Based on some general rules regarding parental selection for the distant hybridization of fish that were revealed in our previous works (Liu, 2010; Wang et al., 2019), the reciprocal crosses between GF and YT were conducted in this study.

The SS rDNA multigene family, composed of highly conserved SS rRNA sequence and variable non-transcribed spacer (NTS) which form arrays of hundreds to thousands of tandem repeats, presents species-specific characteristics in length and the composition of nucleotide sequence (He et al., 2012; He et al., 2013). The accumulation of results from fish indicates SS rDNA as a valuable marker for identification (Martins and Galetti, 2001; Wasko et al., 2001; Pasolini et al., 2006), particularly in hybrids (Qin et al., 2010; He et al., 2012; He et al., 2013).

In addition to the embryonic development, the ploidy (DNA content and chromosome composition), and the morphological characteristics (countable and measurable traits), the heredity of SS rDNA and the gonadal development of the hybrids were also observed and assayed. Briefly, this is the first report of allotriploid via distant hybridization between GF and YT that possessing different chromosome numbers in vertebrates. The obtainment of GFYT hybrids has potential benefits and application in aquaculture. In addition, the variation analysis of SS rDNA multigene families further extended the understanding of the influence of hybridization and polyploidization on the genomic constitution of the hybrid offspring. Furthermore, they can be used as a model to test the origin and consequences of polyploidization. Moreover, this novel allotriploid, which harbors only testis-like and fat-like gonads, can also serve as a proper resource to study the underlying mechanisms of spermatogenesis dysfunctions.

MATERIALS AND METHODS

Animals

All fish, including GF, YT, and GFYT hybrids, were cultured in ponds (23–27°C, natural light cycle) at the State Key Laboratory of Developmental Biology of Freshwater Fish and artificially fed. All experiments were approved by the Animal Care Committee of Hunan Normal University and followed the guidelines of the Administration of Affairs Concerning Animal Experimentation of China.

Obtainment of GFYT Hybrids

During the breeding seasons (April to June), the crosses of GF (2n = 100, δ, n = 3) × YT (2n = 48, δ, n = 3) and YT (2n = 48, δ, n = 3) × GF (2n = 100, δ, n = 3) were conducted, respectively. Two self-crossings of GF and YT (n = 6 each, with the same sex ratio) were also performed. The zygotes were cultured in dishes at a water temperature of 19–20°C. About 2,000 embryos were chosen at random to measure the fertilization rate (number of embryos at the gastrula stage/number of eggs × 100%) and hatching rate (number of hatched fry/number of eggs × 100%). The main stages of embryogenesis were photographed using a stereoscope (Leica, MZ16FA, Wetzlar, Germany). Finally, the hatched fry was transferred to different ponds for further culture.

Measurement of Measurable and Countable Traits

For measurable traits, one-year-old GF, YT, and GFYT hybrids (230–350 g, n = 30 each, including parents) were randomly selected to check total length (TL), body length (BL), body height (BH), head length (HL), head height (HH), caudal peduncle length (CPL), and caudal peduncle height (CPH) (with 0.1cm accuracy). Then, the ratios of BH/BL, BL/TL, HL/BL, HH/HL, CPH/CPL, and HH/BH were further calculated. In addition, the number of lateral line scales (LS), scale rows above the lateral line (ALS), scale rows below the lateral line (BLS), dorsal fin rays (DFR), anal fin rays (AFR), and pelvic fin rays (PFR) were also counted.
Flow Cytometry

The peripheral blood, taken from the caudal veins of 5 GF, 5 YT, and 30 GFYT hybrids which were older than 6 months, were filtered and stained with 4', 6-Diamidino-2-phenylindole DNA-staining solution, cystain DNA 1 step (Partec, Görlitz, Germany). The mean DNA content of GF, YT, and GFYT hybrids was measured with flow cytometry (Ploidy Analyzer PA, Partec, Münster, Germany). Finally, a comparison of mean DNA content between GFYT hybrids and their parents was tested by the $\chi^2$ test with SPSS (version 17.0, the Chi-square test with Yates correction was used for testing deviation from the expected ratio values).

Chromosome Spreads

The preparation of metaphase chromosome spreads was performed with the kidney tissues and the protocols were carried out according to Wang et al. (2014).

Fluorescence In Situ Hybridization

A specific probe of 263 bp centromere repeat sequence, amplified from the genomic DNA of GF, and labeled with Dig-11-dUTP via using a PCR DIG probe synthesis kit (Roche, Penzberg, Germany), was hybridized with the metaphase chromosome spreads of GFYT hybrids and their parents (n = 3 each) according to the previous method described by He et al. (2012). The primer pair (5'-AAG CTTTTTCTCTAAGAGAAGGC-3'; 5'-TTGAGCAGATTT GGGCTTGATT-3', sequence number: JQ086761) was used to amplify this specific probe and at least 30 metaphases from each sample were analyzed.

Histological Analysis of Gonadal Tissues

Compared with one-year-old female GF and male YT, 20 GFYT hybrids at the same age were randomly selected to check the histological structure of gonads. First, the gonadal tissues were fixed in Bouin’s solution overnight. The paraffin sections (6 μM) were sequentially cut, deparaffined, and further stained with hematoxylin and eosin. The microstructure of gonadal tissues was photographed and performed under a light microscope (Olympus, CKX41-32PH, Tokyo, Japan). Meantime, some other gonadal tissues were sequentially fixed in 3% glutaraldehyde solution, washed with phosphate buffer, transferred into osmic acid solution, dehydrated in a graded acetone series, and finally embedded in Epon 812. Ultrathin sections (60 nM) were cut and stained with uranyl acetate and lead citrate. The ultrastructure of gonadal tissues was observed and photographed with an electron microscope (HITACHI, HT7800, Tokyo, Japan).

Terminal Deoxynucleotidyl Transferase dUTP Nick End-Labeling Assay

Gonadal tissues from one-year-old testis-like GFYT hybrids and male GF (n = 3 each) were embedded in the optimal cutting temperature compound (SAKURA, Torrance, CA, United States). The apoptotic analysis was performed on sections (6 mm) by using the in situ cell death detection kit (Roche, Penzberg, Germany) according to the manufacturer’s

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**TABLE 1 | List of primers used in this study.**

| Gene   | Sequence (5’→3’) | Annealing temperature (°C) | Primer purpose |
|--------|------------------|----------------------------|----------------|
| 5s rDNA | F: GCTATGCCCCGATCTGCTGTA | 60 | PCR |
|        | R: CAAGTTTCTGATGGCCCTAGTAAAGC | | |
| dmc1   | F: GAAGATGAGGAATCTCTCTTAA | 56 | qPCR |
|        | R: GCCTCAAGACGGCTCCTTTA | | |
| spo11  | F: GAAGATGAGGAATCTCTCTTAA | 55 | qPCR |
|        | R: GCCTCAAGACGGCTCCTTTA | | |
| rad51  | F: GATCCATGGAAGAGGAGAGAAGA | 56 | qPCR |
|        | R: TCTACTGAGGAGGGGGTTGGGGTC | | |
| mns1   | F: CCAACCCTGCTGAGAAGAAATAT | 55 | qPCR |
|        | R: GATCCATGGAAGAGGAGAGAAGA | | |
| meig1  | F: GATCGAGAGAATCTGACATCC | 60 | qPCR |
|        | R: ACAAGATGAGGCTGAGACGACGA | | |
| p53    | F: CCATCGCTAATGATCAGATCCT | 56 | qPCR |
|        | R: TCTCTGATGAGGGTTCGGTTGTC | | |
| bcl-2  | F: CGATATAGGAGTGAAGTACACCC | 55 | qPCR |
|        | R: TTCGCCAGATGAGTCAGG | | |
| bax    | F: ATTTACTGCTCTCCTCTGTCATCC | 55 | qPCR |
|        | R: AGAAGACGAGGGCTGAGTTG | | |
| Caspase9 | F: GATGAGAGGGCTTCTCTTGCTG | 60 | qPCR |
|        | R: GAGAGCGGGAGGCTGAAAGCT | | |
| β-actin | F: GCCCTGCCCCAGGCGATCCT | 62 | qPCR |
|        | R: AGTGGGATCGCATCGCTGCA | | |

**TABLE 2 | Fertilization rates and hatching rates of different crosses.**

| Crossing group | Fertilization rates | Hatching rates |
|----------------|--------------------|----------------|
| GF(♀) × GF(♂) | 94.3% | 86.7% |
| YT(♀) × YT(♂) | 90.2% | 83.4% |
| GF(♀) × YT(♂) | 75.3% | 64.6% |
| YT(♀) × GF(♂) | 5.0% | 0 |

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instructions. Apoptotic signals were observed using a microscope (Leica, DM6000B, Wetzlar, Germany).

5S rDNA Assay
Total genomic DNA was, respectively, isolated from the blood cells of GF, YT, and GFYT hybrids with a DNA extraction kit by following the manufacturer’s instructions (Sangon, Shanghai, China). The reaction mixture (25 μl) consisted of 20 ng genomic DNA, 1.5 mM MgCl₂, 0.2 mM of each dNTP, 0.4 μM of each primer, 1 × PCR buffer, and 1.25 U Taq polymerase (Takara, Dalian, China). The amplification process started with an initial denaturation at 94°C for 4 min, then followed by 30 cycles of 94°C for 30 s, 60°C for 30 s, and 72°C for 1 min. A final extension step was performed at 72°C for 10 min. The PCR products were sequentially separated on a 1.2% agarose gel, purified using a gel extraction kit (Sangon, Shanghai, China), ligated into a pMD18-T vector, and transferred into E. coli DH5α. Finally, the positive clones were then sequenced using an automated DNA sequencer (ABI PRISM 3730; Applied Biosystems, Carlsbad, CA, United States). Ten clones of each PCR band were sequenced. The primers used for 5S rDNA amplification were listed in Table 1.

Real-Time Quantitative PCR
Total RNA was extracted from the gonads of male GF and testis-like GFYT hybrids (n = 3 each) with TRIzol™ Reagent (Invitrogen, Carlsbad, CA, United States). The concentration and purity of RNA were determined using spectrophotometry and agarose gel electrophoresis. After treatment with RNase-free DNase (Promega, Madison, WI, United States), the total RNA was reversed transcribed to complementary DNA (cDNA) with Rever Tra Ace M-MLV (TOYOBO, Osaka, Japan). RT-qPCR was carried out by using 23SYBR Green mix (TOYOBO, Osaka, Japan) and ABI 7900HT RT-PCR System (Applied Biosystems, Carlsbad, CA, United States). β-actin was used as the internal reference and the expression levels were calculated with the 2^-ΔΔCt method. The primers used here were listed in Table 1.

Statistical Analysis
All statistical analyses were conducted by using SPSS software (version 17.0). Differences between multiple groups were analyzed by one-way analysis of variance (ANOVA) (IBM Corp. New York, United States). Sigma Plot 10.0 (San Jose, CA, United States) was used to present the basic graph. All data were shown as means ± SD. A p-value less than 0.05 was considered to be statistically significant.

RESULTS
The Formation of GFYT Hybrids and Their Embryonic Development
The crosses of GF (♀) × YT (♂) in which the number of maternal chromosomes is more than that of paternal chromosomes generated four types of hybrid offspring, including gray or red single-tail individuals (ST GFYT hybrids) and gray or red twin-tail individuals (TT GFYT hybrids) (Figure 1A). While the crosses of YT (♀) × GF (♂) in which the number of maternal chromosomes is less than that of paternal chromosomes were lethal (Figure 1B). In addition, embryogenesis was also observed. Most of the embryos from the cross of GF (♀) × YT (♂) showed normal development, but all the embryos from the cross of YT (♀) × GF (♂) died before the end of the neurula stage (Figure 1C). Meantime, the fertilization rates and hatching rates were further analyzed (Table 2). There were high fertilization rates and hatching rates in both the self-crossing of GF (94.3 and 86.7%, respectively) and YT (90.2 and 83.4%, respectively). Also, the fertilization rates and hatching rates of GF (♀) × YT (♂) were 75.3 and 64.6%, respectively. However, the fertilization rates and hatching rates of YT (♀) × GF (♂) were severely reduced to 5.0% and 0.

Measurable and Countable Traits of GFYT Hybrids
The comparisons of measurable and countable traits among GFYT hybrids and their parents were also conducted (Figures 1D, E). There was no significant difference between single-tail and twin-tail individuals. Compared with parents GF and YT, the rates of the main characteristics, including body height/body length and head length/body length of GFYT hybrids presented obvious intermediate properties. But the rate of body length/total length of GFYT hybrids showed an apparent bias to that of YT, and the rates of head height/head length and caudal peduncle height/total length of GFYT hybrids showed an apparent bias to that of GF. The countable traits of GFYT hybrids were also more similar to that of GF, excepting BLS.

DNA Content and Chromosomal Constitution of GFYT Hybrids
To reveal the construction of genetic materials (nucleus), the mean DNA contents of GFYT hybrids and their parents were checked. There was no significant difference between the value of GFYT/(GF + 0.5YT) and the expected value (Figures 2 A–C, Table 3). Then, we further confirmed the chromosomal constitution of GFYT hybrids and their parents. There were 100 and 48 chromosomes in GF and YT, respectively. All GFYT hybrids possessed 124 chromosomes which was equal to the number of GF chromosomes and one-half of YT chromosomes (Figures 2D–F). To further confirm the chromosomal heredity of GFYT hybrids, a specific probe from GF was applied to hybridize the metaphase chromosome spreads of GFYT hybrids and their parents. There were 100 signals in GF and none in YT. As expected, only 100 signals were found in the GFYT hybrids. Then, a novel allotriploid population with 124 chromosomes, with two sets from GF and one set from YT, was successfully obtained from the cross of GF (♀) × YT (♂) × YT (♀) (Figure 1C).
ST GFYT hybrids and four (188, 206, 340, and 482 bp) in TT GFYT hybrids (Table 4), respectively. Based on BLASTn analysis (https://blast.ncbi.nlm.nih.gov/Blast), all fragments were confirmed to be 5S rDNA repeat units. According to the length, the similarities of 5S rDNA repeat units between GF and YT were first checked. There were extremely low similarities between GF-168 and YT-188 (72.8%), GF-203 and YT-206 (76.5%), and GF-340 and YT-376 (70.8%). So, the 5S rDNA composition of GF and YT were, respectively, assigned as GF-I~IV and YT-I~III (Table 4). Then, the sequence alignment between GFYT hybrids and their parents was conducted. The 188 bp and 206 bp 5S rDNA of GFYT hybrids (including ST and TT), respectively, showed a high similarity to that of YT-I and YT-II but obviously low to that of GF-I and GF-II (values in the brackets). Inversely, the 340 bp 5S rDNA of GFYT hybrids presented a high similarity to that of GF-III, but apparently low to that of YT-III (value in the brackets). The ~480 bp 5S rDNA of GFYT hybrids exhibited high similarity to that of GF-IV (Table 5). In addition, the 5S rDNA sequences with the same or similar length between ST and TT GFYT hybrids showed high similarity (Table 5). Given the high conservation of coding region and variability of NTS in 5S rDNA sequences, the alignment of NTS sequences with the same length between GFYT hybrids and parents was further conducted to confirm their composition, which presented high constance except several nucleotide mutations (Figure 3B). These results indicated that GFYT hybrids, including single-tail and double-tail individuals, both specifically inherited YT-I and YT-II 5S rDNA types from YT, GF-III, and GF-IV from GF (Figure 3C).
Reproduction Dysfunction of GFYT Hybrids

In the breeding season, compared with the parents, none of the ovum or sperm could be squeezed out from GFYT hybrids. Then, further histological observations were employed to confirm the gonadal development (Figure 4). Both GF (Figures 4A,B) and YT (Figures 4C,D), respectively, present normal oogenesis and spermatogenesis. However, only testis-like (Figures 4E,F) and fat-like gonads (Figures 4G,H) were found in GFYT hybrids. In the testis-like gonads, there were spermatogonia, spermatocytes, and spermatids, while no mature sperm was found (Figures 4E,F). Interestingly, the electron microscope did not observe abnormality in the structure of spermatogonia and spermatocytes (Figures 4I,J) but found that spermatids presented an abnormal organization with pyknotic and heteromorphous chromatin and invaginated cell membrane (Figures 4K,L), indicating the undergoing degeneration.

Apoptosis of Testis-Like Gonads in GFYT Hybrids

The newly formed allotriploid were proved to be sterile and the spermatids presented obvious degeneration (Figure 4). These results suggested that the infertility of GFYT hybrids may result from the developmental arrestment of spermatogenesis. Apoptosis, a programmed cell death, which is crucial for the normal...
development and organismal homeostasis maintenance (Singh et al., 2019), holds the highest possibility. To uncover the underlying mechanisms, the TUNEL assays were conducted. Compared with male GF whose testis showed nearly no signal, apparent apoptotic signals were mainly distributed around spermatid regions in the testis-like gonads of GFYT hybrids (Figure 5A). Then, the mRNA expression of apoptosis pathway-related genes, including p53, bcl-2, bax, and caspase9 that play vital roles in the normal development of germ cells and the reproductive system (Singh et al., 2019) were further identified (Figure 5B). Compared with control, though there was only an increasing trend in the mRNA expression of bcl-2, the expression of p53, bax, and caspase9 of testis-like gonads in GFYT hybrids showed an observable upregulation (Figure 5B). In addition, to explore the possible factors causing the prevention of spermatogenesis, the mRNA expression of several key meiosis or spermatid maturation related genes was checked. The expression of meiosis-related genes, including spo11, dmc1, and rad51 showed an abnormality of high expression in the testis-like gonad of GFYT hybrids (Figure 5C). However, the expression of mns1 and meig1, two key genes involving the maturation of spermatid, were extremely downregulated (Figure 5C).

**DISCUSSION**

A novel and sterile allotriploid was successfully obtained via crossing female GF and male YT, which belonged to different subfamilies in this study. The composition and similarity assays of 5S rDNA were also conducted to reveal the genetic characteristics of GFYT hybrids. In addition, the gonadal dysfunctions and spermatogenesis disorders were further analyzed to explore the underlying mechanisms.

**Novel Alloploid Produced by Hybridization**

Distant hybridization is a valuable approach for producing hybrid offspring with improved characteristics in fish breeding. It may break through reproductive barriers, integrate advanced characteristics from different species, expand the genetic variation, or even produce new variants or species (Coyne and Orr, 2004; Mallet 2007). However, most hybridization resulted in hybrid incompatibility (Coyne and Orr, 2004; Mallet, 2005; Maheshwari and Barbash, 2011; Abbott et al., 2013). Many genetic and environmental factors are the cause of hybrid incompatibility. Among them, the genetic factors included conflict in the interaction of different genes, allele variants or epigenome, the conflict between the maternally inherited cytoplasm and the hybrid genome, and conflict coming from the biased expression of either parental alleles in the hybrid offspring (Vaid and Laitinen, 2019). In this study, GF and YT, respectively, possess 100 and 48 chromosomes, only the cross of GF (♀) x YT (♂) in which the number of maternal chromosomes is more than that of paternal chromosomes generated surviving.
hybrid offspring, but the cross of YT (♀) × GF (♂) in which the number of maternal chromosomes is less than that of paternal chromosomes was lethal. These results were consistent with the previous studies which stated that it is more likely to obtain hybrid offspring when the number of maternal chromosomes is larger than that of paternal chromosomes, but the hybrids are unlikely to survive if the number of maternal chromosomes is fewer than that of paternal chromosomes (Liu, 2010; Song et al., 2012; Wang et al., 2019).

To conquer hybrid incompatibility, the hybrid offspring may rearrange the genome (Osabe et al., 2012). The 5SrDNA multigene family presents species-specific characteristics in length and the composition of the nucleotide sequence. The accumulation of results from fish indicates 5SrDNA could be used to identify species and analyze the evolution and genetic relationship (Sajdak et al., 1998; Campo et al., 2009), especially hybrid progeny (Pendas et al., 1995; Qin et al., 2010; He et al., 2012). Herein, GFYT hybrids including single-tail and twin-tail individuals specifically inherited the paternal-specific YT-I and YT-II 5SrDNA types, maternal-specific GF-III and GF-IV. However, the paternal-specific YT-III, maternal-specific GF-I, and GF-II 5SrDNA types were lost in the hybrid offspring. In addition, several variations of nucleotide sequence were also found in NTS sequences. These variations of NTS sequences suggested that 5SrDNA can be used as suitable genetic markers for the identification of GFYT hybrids and their parents.

### TABLE 5 | Similarities of 5SrDNA sequences between hybrids and their parents.

| Length (bp) | ST GFYT/GF | ST GFYT/YT | TT GFYT/GF | TT GFYT/YT | ST GFYT/TT GFYT |
|------------|-------------|-------------|------------|------------|-----------------|
| 188 (168) | (72.8%) | 96.8% | (75.5%) | 97.3% | 99.4% |
| 206 (203) | (77.4%) | 98.1% | (78.4%) | 98.1% | 100% |
| 340 (376) | 96.1% | (66.3%) | 96.7% | (70.6%) | 97.1% |
| ~480bp | 99.5% | ---- | 99.3% | ---- | 98.9% |

*The approximate length of 5SrDNA. The numbers or values of percentage in the brackets, respectively, show the base length of 5SrDNA, similar to GF, or YT, and similarities between them and GFYT hybrids.*

![FIGURE 4](image-url)  
**FIGURE 4** | Histological structure of gonads in GFYT hybrids and their parents. (A–H) H&E-stained gonads and (I–L) ultrastructure of testis-like gonads of GFYT hybrids. (A) Ovary of GF. (B) Testis of GF. (C) Ovary of YT. (D) Testis of YT. (E,F) Testis-like gonad of GFYT hybrids. (G,H) Fat-like gonad of GFYT hybrids. (I,J) Spermatogonia and spermatocytes in the testis-like gonad of GFYT hybrids. (K,L) Spermatids with heteromorphous nuclei in the testis-like gonad of GFYT hybrids. F, H, J, and L, respectively present the amplification of the dashed box in E, G, I, and K.
Polyploidization, an important approach to hybrid speciation (Soltis and Soltis, 2009), resulted in a drastic change in the genome mainly through the expulsion inhibition of the second pole, the impeding of first cleavage, or double fertilization (Ramsey and Schemske, 1998). There may be theoretically four types of offspring in the cross of GF ($2n$) × YT ($\delta$), the allodiploid ($2n = 74$) with one set of GF and YT, respectively; allotriploid ($3n = 98$) derived from the double fertilization between one egg from GF and two sperms from YT; allotriploid ($3n = 124$) from the inhibition of the expulsion of the second pole (Figure 6). Given the fact that GFYT hybrids were all triploids with two sets of maternal chromosomes and one set of paternal chromosomes, we discreetly concluded that the GFYT hybrids resulted from the inhibition of the expulsion of the second pole. The remaining three types of chromosomal constitution may also exist, but these genetic architectures are lethal during early embryogenesis. So, this is why there were
low fertility rates (75.3%) and survival rates (64.6%) in GFYT hybrids.

In brief, this is the first report of allotriploid via distant hybridization between GF and YT which possess different chromosome numbers in vertebrates. The obtainment of GFYT hybrids has potential benefits and applications in aquaculture. In addition, the variation analysis of 5S rDNA multigene families further extended the understanding of the influence of hybridization and polyploidization on the genomic constitution of the hybrid offspring. Furthermore, they can be used as a model to test the origin and consequences of polyploidization.

**Hybrid Sterility**

The complex processes involving the pairing, synapsis, recombination, and the separation of homologous chromosomes would sequentially occur in meiotic prophase I to form secondary meiocytes (Pawlowski and Cande, 2005). In this study, there were two sets of chromosomes from GF and one set of chromosomes from YT in GFYT hybrids, which resulted in the failed synaptonemal complex (SC) and irregular division in meiotic prophase I. Then, the sister chromosomes separated and formed aneuploidy spermatids. So, spermatids with pyknotic and heteromorphous chromatin and invaginated cell membrane were found in the testis-like gonads of GFYT hybrids. In addition, apoptotic signals that are mainly distributed around spermatid regions further confirmed this speculation. Furthermore, the BCL-2 family of proteins, whose homeostasis of anti-apoptotic and pro-apoptotic compositions determines the final fate of cells (Deng et al., 2007; Chonghaile et al., 2011; Sarosiek et al., 2013; Sarosiek and Letai, 2016; Singh et al., 2019), are the key regulators of the intrinsic apoptosis (Fuchs and Steller, 2011; Galluzzi et al., 2018). Pro-apoptotic protein Bax activates the caspase signaling pathway to initiate apoptosis, and the anti-apoptotic protein BCL-2 suppresses Bax to prevent apoptosis (Agca et al., 2012; Liman et al., 2013; Lucinda et al., 2013). In this study, the expression of p53, bax, and caspase9 significantly increased; however, there was only an increasing trend of bcl-2 without significance. Interestingly, the meiosis-related genes (spo11, dmc1, and rad51) presented an obvious abnormality of high expression, while mns1 and meig1, two key genes involving the maturation of spermatid, showed extreme downregulation. Why there was a high expression of these meiosis-related genes? One potential explanation is compared with male GF, who finished meiosis at the reproduction season, delayed spermatocytes in the testis-like gonads of GFYT hybrids were still undergoing meiosis, even though this was abnormal. So, these meiosis-related genes were transcribed persistently. These results suggested that heterology and triploidy were more likely the reasons for the sterility of GFYT hybrids.

In addition, the conflicts between the growing demand for aquatic food consumption and the global decline of wild fishery stocks make it imperative to develop highly efficient aquaculture practices to increase fishery production. Then, farming infertile fish is a desirable and most effective genetic-containment strategy for aquatic food availability and the development of environmentally responsible aquaculture (Wong and Zohar, 2015). First, there are close linkages between growth and reproduction of fish (Hill, 2013). Sterilization not only increases the growth of muscle by reducing energy input to gonad development but also inhibits sexual maturation, which would deteriorate flesh quality and enhance susceptibility to disease and stress (Zohar, 1989). Second, the escape of non-native selectively bred or genetically modified (transgenic) fish from aquatic farms would propagate or interbreed with wild stock, thus finally greatly threatening the ecosystem and environment. The sterile trait could significantly alleviate these risks (Muir and Howard, 1999). Third, there are different processes, such as commonly used chromosome manipulation, interspecies hybridization, or newly invented bath-immersion technology (Wong and Zohar, 2015), to induce sterile fish. However, these manipulations would be time-consuming and/or could not be produced on a big scale or the hybrid offspring is fertile. GFYT hybrids, obtained via direct distant hybridization, are all allotriploid and completely sterile, well conquering these defects.

**DATA AVAILABILITY STATEMENT**

The original contributions presented in the study are included in the article/Supplementary Material, further inquiries can be directed to the corresponding author.

**ETHICS STATEMENT**

The animal study was reviewed and approved by the Animal Care Committee of Hunan Normal University.

**AUTHOR CONTRIBUTIONS**

JW: conceptualization, writing—original draft, project administration, and funding acquisition. WH: conceptualization, writing—original draft, and funding acquisition. WW: investigation and resources. ZL: investigation and data curation. LH: investigation and data curation. MC: conceptualization, writing—review and editing, and formal analysis. SL: conceptualization, writing—review and editing. TL: writing—review and editing. RZ: writing—review and editing. KL: writing—review and editing. RZ: writing—review and editing. SL: conceptualization, writing—review and editing, supervision, and funding acquisition.

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