An intergeneric hybrid was successfully developed between *Oryza sativa* L. (IRRI 154) and *Leersia perrieri* (A. Camus) Launert using embryo rescue technique in this study. A low crossability value (0.07%) implied that there was high incompatibility between the two species of the hybrid. The F1 hybrid showed intermediate phenotypic characteristics between the parents but the plant height was very short. The erect plant type resembled the female parent IRRI 154 but the leaves were similar to *L. perrieri*. Cytological analysis revealed highly non-homology between chromosomes of the two species as the F1 plants showed 24 univalents without any chromosome pairing. The F1 hybrid plant was further confirmed by PCR analysis using the newly designed 11 indel markers showing polymorphism between *O. sativa* and *L. perrieri*. This intergeneric hybrid will open up opportunities to transfer novel valuable traits from *L. perrieri* into cultivated rice.

**Key Words:** intergeneric hybridization, *Leersia perrieri*, molecular markers, out-group species, rice.
Development of an intergeneric hybrid between *O. sativa* and *L. perrieri*

2009). However, due to destruction of natural habitat of this plant population, this species is becoming endangered (Hutang and Gao 2017).

Several reports revealed that the species *L. perrieri* harbor valuable traits that can be exploited in rice breeding programs. Studies have been limited only to its evolutionary relationships to the genus *Oryza* (Stein et al. 2018). However, there were no extensive studies conducted to explore its morphology and traits potential to rice improvement program since it is very difficult to utilize it as a donor parent due to high incompatibility barrier with *Oryza* species. Screening for long term stagnant flooding conducted at IRRI revealed that *L. perrieri* is highly tolerant to stagnant flooding (Jena et al. unpublished).

This study is the first report on the production of an intergeneric *F*₁ hybrid between *Oryza sativa* and its outgroup species *L. perrieri*. The *F*₁ hybrid was confirmed using cytological analysis and genotyping. A set of *L. perrieri* genome specific markers confirmed the hybridity of *F*₁.

### Materials and Methods

#### Plant materials

A high-yielding rice variety developed by the International Rice Research Institute (IRRI) in the Philippines, *O. sativa* L. *spp. indica* cv. IRRI 154 was used as the female parent and *L. perrieri* (IRGC 105164) as male parent to produce an intergeneric *F*₁ hybrid. The varieties IR42 and FR13A (IRGC 122454) were used as sensitive and tolerant checks respectively for submergence tolerance screening. IRGC stands for the International Rice Genebank Collection (http://www.irgcis.irri.org/81/grc/irgcishome.html).

#### Intergeneric hybrid production

Intergeneric *F*₁ hybrid production was carried out by emasculating and pollinating spikelets of the female parent IRRI 154 with pollen of *L. perrieri*. Pollinated spikelets were immediately bagged and sprayed two times a day for three days with 0.75 ppm of 2-4D (2,4-dinitrophenylhydrazine). Fertilized spikelets after 8, 9, 10, 11 and 12 days after pollination were collected for embryo rescue. Data on number of spikelets emasculated, seed set, embryos isolated, embryos cultured and hybrid plants obtained were recorded to determine the crossability between *O. sativa* and *L. perrieri*.

#### Tissue culture

Spikelets were collected and stored in water prior to pre-treatment to prevent dehydration. Immature spikelets were dissected and filled spikelets were treated with 20% sodium hypochlorite for 20 minutes and then washed with distilled water three times. Spikelets with intergeneric hybrid embryo were isolated in an aseptic condition under stereo microscope and cultured on a quarter strength Murashige-Skoog (MS) media with 6.0 g/L agar, then incubated in dark until shoot initiation following the method of Jena et al. (1989). Germinated embryos were transferred to light incubation room until seedlings are ready for transfer to soil.

**Morphological characterization**

Phenotypes including plant height, tiller number, leaf length, leaf width, panicle length, panicle branching, and spikelet morphology were measured at the flowering stage of the parents and the *F*₁ hybrid and mean values were obtained from five plants per material. The numbers of *F*₁ plant were increased by tiller splitting propagation from the single *F*₁ plant. For spikelet-related traits, spikelets were dissected manually and photographs were taken under a stereomicroscope (Optika, model: SZ-CTV, Italy). Pollen fertility was measured by starch staining in a pollen grain using iodine solution (1% I₂KI). After staining, the pollen grains were observed under light microscope (Olympus, model: BX53, Japan) and percentage of fully stained (dark color) pollen was calculated.

**Cytological analysis**

Meiotic cells of parents and hybrid were observed at the right growth stage and panicles were collected at 8:00 AM and were fixed in 3:1 (v/v) ethanol-acetic acid solution with a pinch of ferric chloride as mordant. Newly fixed samples were first incubated at room temperature for 24 hours and then stored at 4°C until use. Analyses were carried out by simple anther squash technique using 1% aceto-carmine as stain. Chromosome spreads of pollen mother cells (PMC) were observed under a light microscope (Olympus BX53) with 100x magnification in oil immersion objective and pictures were taken using Image Pro 7.0 software (Media Cybernetics, USA).

**Development of DNA markers**

The bacterial artificial chromosome (BAC) end sequences of *L. perrieri* were generated through the *Oryza* Map Alignment Project and *Oryza* Genome Evolution Project (Jacquemin et al. 2013). DNA sequence alignments between *O. sativa* cv Nipponbare and *L. perrieri* were conducted using the comparative genomics tool provided by Gramene database (www.gramene.org). Based on the sequence comparison data, one indel-sequence based marker for each of the 12 chromosomes was designed (Supplemental Table 1).

**Genotype confirmation of hybrid by using PCR**

Genomic DNA was isolated from leaves of both parents and *F*₁ hybrid using a simple DNA extraction method without chloroform extraction and DNA pellet precipitation steps (Kim et al. 2016). PCR amplifications were carried out using the newly designed 12 markers and the PCR products were analyzed by electrophoresis on 3% agarose gel. Based on the predicted PCR product sizes (Supplemental Table 1) and comparison of PCR band patterns among two parents and *F*₁ plants, the intergeneric hybridity was confirmed.
Screening for submergence flooding of L. perrieri

Seeds of O. sativa (IRRI 154) and L. perrieri were pre-germinated for three days and then seeded in trays together with tolerant and sensitive checks and were allowed to grow for 21 days at normal condition. Number of seedlings germinated was recorded. Then, the trays containing normal seedling plants were placed in three independent 1.5 m-depth tanks for experimental replications. The tanks were slowly filled with water for providing submergence stress to the plants. Twenty-one days after submergence, water in the tank was slowly drained out. Another 21 days after de-submergence, number of seedlings survived were recorded.

Results

Production of intergeneric hybrid between O. sativa and L. perrieri

Spikelets of the intergeneric cross between IRRI 154 (O. sativa) and L. perrieri were collected after 8, 9, 10, 11 and 12 days after pollination (DAP) to determine the optimum number of days required for embryo rescue with viable embryos. After pollination, swollen ovary which might have embryo ranged from 51.64 to 80.92% and the highest percentage was observed 12 DAP (Table 1). However, no embryo was observed from the 140 swollen ovaries at 12 DAP. Of the 489 spikelets containing swollen ovary at 10 DAP, only five embryos were found and the rest of spikelets had only watery ovary without development of embryo and endosperm. Finally, to obtain F1 plants, in total, 933 swollen ovaries among 1,414 pollinated spikelets were dissected under stereomicroscope and seven embryos were found (Table 2). Of the seven embryos rescued on the quarter strength MS medium, two embryos germinated. However, due to mortality of germinated embryo as the slightly developed callus turned blackish, only one embryo successfully germinated and developed to viable plant. The crossability between O. sativa and L. perrieri was 0.07% (Table 2).

Morphological characterization

As a common modern rice cultivar, a plant of IRRI 154 variety exhibits erect stem and semi-tall plant type. In contrast, L. perrieri has a creeper type of shoot structure (Fig. 1). Overall plant phenotype of the F1 hybrid was more close to cultivated rice, O. sativa although the plant height was much smaller compared to IRRI 154 (Fig. 1). Leaf width and length of the hybrid plant was similar to those of L. perrieri. Panicle length of the hybrid was a little bit longer than L. perrieri. Panicle branching of the F1 hybrid plant was poor like L. perrieri (Fig. 2, Table 3). Spikelets of IRRI 154 had sterile lemmas but there was no awn, while L. perrieri had a long awn at the tip but absence of sterile lemmas (Fig. 3). The spikelets of the hybrid plant possessed both sterile lemmas and awn. This observation suggested that the sterile lemmas and awn on the spikelets of the hybrid plant might be inherited from IRRI 154 and L. perrieri, respectively. Pollen fertility of the F1 hybrid was observed although it was low (27.35%) compared to parents O. sativa (91.50%) and L. perrieri (86.24%) (Table 3).

Chromosome analysis of parents and F1 hybrid

Chromosome analysis of both O. sativa and L. perrieri revealed 12 pairs of bivalents in all the pollen mother cells (PMCs) observed during meiosis stage. The F1 hybrid exhibited 24 univalents without any evidence of chromosome pairing (Fig. 4).
Development of an intergeneric hybrid between *O. sativa* and *L. perrieri*

**Genotype confirmation of F₁ hybrid**

Of the 12 putative polymorphic DNA markers between *O. sativa* and *L. perrieri*, 11 markers showed the predicted PCR band sizes and polymorphism between *O. sativa* and *L. perrieri*. In the F₁ hybrid plant, all the markers of each chromosome except for chromosome 4 showed two PCR bands which were originated from the each parent allele respectively (Fig. 5). This result confirmed hybridity for the F₁ plants. Monomorphic band pattern of the chromosome 4 marker (Lp04-00646) between IRRI 154 and *L. perrieri* might be caused by insignificant sequence difference between the parents although a sufficient indel (33 bp gap) was predicted between Nipponbare and *L. perrieri* (Supplemental Table 1).

**Screening for submergence tolerance**

Screening for long period (21 days) stagnant flooding was conducted to determine the response of *L. perrieri* and IRRI 154. During the period of submergence, *L. perrieri* had rapidly elongated internodes, resulting that the upper-most shoot emerged above the 1.5 m of water level and survived. In contrast, IRRI 154 plants collapsed during submergence and the plants completely died during recovery period after draining out water (Supplemental Fig. 1). All the three replications showed same results.

---

**Fig. 2.** Panicle phenotype of IRRI 154 (A), *L. perrieri* (B) and F₁ hybrid (C). Panicle of *L. perrieri* (B) is normally closed (left) and it was artificially opened for the phenotype observation (right).

**Table 3.** Comparison of morphological characteristics between two parents and F₁ hybrid

<table>
<thead>
<tr>
<th>Traits</th>
<th><em>O. sativa</em> (IRRI 154)</th>
<th><em>L. perrieri</em> (IRGC 105164)</th>
<th>F₁ hybrid</th>
</tr>
</thead>
<tbody>
<tr>
<td>Plant height (cm)</td>
<td>103.0 ± 7.5</td>
<td>41.0 ± 14.1</td>
<td>17.5 ± 2.4</td>
</tr>
<tr>
<td>Tiller number</td>
<td>15.2 ± 2.3</td>
<td>-</td>
<td>5.0 ± 1.4</td>
</tr>
<tr>
<td>Leaf length (cm)</td>
<td>31.6 ± 7.1</td>
<td>2.3 ± 0.7</td>
<td>11.0 ± 2.5</td>
</tr>
<tr>
<td>Leaf width (cm)</td>
<td>1.23 ± 0.2</td>
<td>0.4 ± 0.1</td>
<td>0.5 ± 0.1</td>
</tr>
<tr>
<td>Panicle length (cm)</td>
<td>26.3 ± 6.8</td>
<td>2.9 ± 0.8</td>
<td>5.5 ± 2.4</td>
</tr>
<tr>
<td>Panicle branching (no.)</td>
<td>12.0 ± 3.4</td>
<td>0–3</td>
<td>3.1 ± 1.1</td>
</tr>
<tr>
<td>Stigma color</td>
<td>white</td>
<td>light purple</td>
<td>light purple</td>
</tr>
<tr>
<td>Plant type</td>
<td>erect</td>
<td>creeper</td>
<td>erect</td>
</tr>
<tr>
<td>Pollen fertility (%)</td>
<td>91.5 ± 4.2</td>
<td>86.2 ± 7.5</td>
<td>27.4 ± 11.5</td>
</tr>
<tr>
<td>Awn</td>
<td>-</td>
<td>+</td>
<td>+</td>
</tr>
</tbody>
</table>

**Fig. 3.** Spikelet morphology of IRRI 154, *L. perrieri* and F₁ hybrid. Phenotypes were observed under stereomicroscope (Scale bar = 0.5 mm).

**Fig. 4.** Chromosome configuration at metaphase I of IRRI 154 (A), *L. perrieri* (B) and F₁ hybrid (C) showing their chromosome numbers.

**Fig. 5.** Genotype confirmation of F₁ hybrid using the newly designed markers for discriminating *O. sativa* and *L. perrieri* alleles.
Discussion

Wild *Oryza* species as well as the nearest relatives of the genus *Oryza* has untapped valuable traits that can be used in rice improvement programs. Although extensive studies have been made to discover the value-added traits of wild *Oryza* species and were successfully utilized in rice breeding programs, the traits from distantly related genera have not been explored yet (Jena and Nissila 2017, Sanchez et al. 2014). One limitation on production of intergeneric hybrid for crop improvement includes high incompatibility between the genomes of two different genera being crossed. Reproductive barriers that restrict intergeneric hybridization to become successful may happen before or after fertilization. Pre-fertilization barriers may involve failure of pollen germination as well as pollen tube growth (Morgan et al. 2010). On the other hand, post-fertilization barriers include failure of zygote development after fertilization, degeneration of hybrid embryo, non-development of endosperm and female sterility in the hybrid plants, or even lethality of the hybrid (Kaneko and Bang 2014). This eventually results to abortion of embryos. In this study, we successfully obtained intergeneric hybrid by embryo rescue. Excision of embryo in an aseptic condition has been executed and embryo was placed on a special MS media with 1/4 strength of nutrients. It was established that the genotype, developmental stage of the embryo at excision as well as embryo culture media composition are the three main factors for the success and efficiency of embryo rescue (Jena and Khush 1989, Shen et al. 2011). In attempting embryo rescue of highly incompatible crosses, it is critical that the tissue culture process is initiated prior to the period where embryo abortion starts (Reed 2005). In this study, the optimum number of days before embryo abortion in the intergeneric cross between *O. sativa* and *L. perrieri* was 10 days after pollination took place.

Incorporating economically important genes into cultivated rice is a way of increasing genetic diversity as well as improvement of modern rice varieties to overcome the adverse effect of climate change. Currently, only interspecific hybridization has been successfully carried out between cultivated rice and wild *Oryza* species. Unlike other crops, rice has never been used for intergeneric hybridization. This is the first report on a successful development of a hybrid between *O. sativa* and *L. perrieri*.

The genus *Oryza* and *Leersia* diverged 14.2–15 million years ago (Guo and Ge 2005, Stein et al. 2018). In the phylogenetic tree of *Oryzaeae*, the morphological evidence that separates *Leersia* to *Oryza* is the loss of sterile lemmas in the genus *Leersia* (Kellogg 2009). Very low crossability between *O. sativa* and *L. perrieri* clearly indicates that there is high incompatibility between the two species. This is expected since two parents are very distantly related and from different genera. The low percentage of seed set and resulting embryo abortion might be due to introduction of excessive exotic genetic material from *L. perrieri* as well as the presence of genetic imbalance leading to somatic incompatibility (Liu et al. 2005).

Although it has been established that *L. perrieri* has 24 chromosomes, its genome has not yet been classified clearly in relation to the other wild *Oryza* species. A study by Katayama (1995) observed that there were no genomic relationship between *L. perrieri* and *O. punctata* (BB genome) and *O. latifolia* (CCDD genome). It can be deduced that *L. perrieri* has no chromosome homology with the BB and CCDD genomes of *Oryza*. Our findings also show that *L. perrieri* is not even related with the AA genome since there was no chromosome association observed to the recurrent parent IRRI 154. Non-homologous chromosome pairing was observed in the F1 hybrid obtained in this study clearly (Fig. 4) suggested that the two parents are highly incompatible genetically with no chromosome associations between the species. Meiotic data observed in our study might suggest that there are no genomic relationships between *L. perrieri* and *O. sativa*, and several wild relatives in the secondary/tertiary gene pool of *Oryza*. Nonetheless, the likelihood of similar sets of chromosome number between *O. sativa* and *L. perrieri* indicates the potential for backcrossing to take place and fertility could be improved than that of hybrids with parents with different chromosome numbers. However, the lack of chromosome pairing usually results in little or no homologous recombination, which might be caused by large DNA blocks inherited in the progenies. These DNA blocks introgressed in the progenies sometimes contain unfavorable alleles that might be a hindrance to produce genotypes that can be employed in future breeding programs.

In the submergence experiment, *L. perrieri* showed dramatic internode elongation to avoid submergence stress while other *O. sativa* materials collapsed during submergence. In the *O. sativa* species, deep water rice varieties are able to escape submergence stress by using elongation mechanism because of the presence of *SNORKEL1* and *SNORKEL2* genes encoding single ethylene response factor (ERF) domain proteins (Hattori et al. 2009). However, the genetic factors of *L. perrieri* for elongation mechanism need to be studied in future which might have either superior alleles to the known genes such as *SNORKEL1* and *SNORKEL2* or new genes which are absent in *O. sativa* species. In addition to elongation mechanism, three *ERF* genes related to *SUB1* locus conferring ‘quiescence strategy’ to submergence stress were also detected in chromosome 9 of *L. perrieri* (Dos Santos et al. 2017). However, our phenotype observations under submergence stress suggest that *L. perrieri* uses an elongation mechanism as primary submergence tolerance mechanism rather than quiescence mechanism. This particular trait can be transferred to cultivated rice once the F1 can be further backcrossed to *O. sativa* parent for the production of monosomic alien addition lines and disomic introgression lines. The development of these valuable materials will be able to provide an avenue of improving rice varieties in rice ecosystems that
Development of an intergeneric hybrid between *O. sativa* and *L. perrieri*

are prone to flooding.

Intergeneric hybridization is an effective method to broaden the genetic base of a cultivated crop species. It has been reported that intergeneric hybrids have been developed between yellow mustard and rapeseed (Brown et al. 1997), turnip and radish (Lou et al. 2017) and many other crops (Bang et al. 2007, D’Hont et al. 1995, Hu et al. 2002). Tissue culture is necessary to successfully obtain intergeneric hybrids. Production of intergeneric hybrids and their progenies can also be a tool to conserve valuable traits of species that are threatened to be extinct by incorporating genes into the existing gene pool of cultivated species. Rice has been used together with barley to produce an intergeneric hybrid using protoplast fusion (Kisaka 1998). Intergeneric hybridization was conducted between *O. sativa* and *Luziola peruviana* but the focus of this study was to observe chromosomal aberrations and cytological alterations that exist in the intergeneric hybrid (Moreno et al. 2014).

In this study, as a first step, intergeneric hybridization between *O. sativa* and *L. perrieri* was successfully carried out. To transfer DNA segments harboring valuable traits from this grass species to the cultivated rice, chromosome introgression of *L. perrieri* into *O. sativa* is an essential procedure. In the wide hybridization between different genomes, obtaining BC₁ or F₂ plants from the initial F₁ hybrid is one of the most difficult steps. To overcome this, we will amplify the number of F₁ plants using tiller splitting propagation and make the F₁ plants very healthy through growth condition optimization. Furthermore, constant pollination and embryo rescue will be required to increase the possibility like obtaining F₁ hybrid plants. Fortunately, our F₁ hybrid showed some pollen fertility (Table 3) and this might be helpful for the production of the next generation plants. Moreover, production of these lines can be a good material to unravel the genetic mechanism of *L. perrieri* for its survival in stress conditions like flooding.

**Acknowledgments**

We would like to thank Mr. Patricio Carandang for the excellent technical assistance provided during the conduct of this study. We are grateful to CGIAR global rice science partnership program (GRiSP: Grant No. DRPC 2011-134) of IRRI for supporting this study. We thank the IRRI editorial team for editing this manuscript.

**Literature Cited**


**Ballesfin, Vinarao, Sapin, Kim and Jena**


