Genetic Features of the Spontaneous Self- Compatible Mutant, ‘Jin Zhui’ (Pyrus bretschneideri Rehd.)

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Abstract

‘Jin Zhui’ is a spontaneous self-compatible mutant of ‘Ya Li’ (Pyrus bretschneideri Rehd. S27S34), the latter displaying a typical S-RNase-based gametophytic self-incompatibility (GSI). The pollen-part mutation (PPM) of ‘Jin Zhui’ might be due to a natural mutation in the pollen-S gene (S24 haplotype). However, the molecular mechanisms behind these phenotypic changes are still unclear. In this study, we identified five SFL (S-Locus F-box) genes in ‘Ya Li’, while no nucleotide differences were found in the SFL genes of ‘Jin Zhui’. Further genetic analysis by S-RNase PCR-typing of selfed progeny of ‘Jin Zhui’ and ‘Ya Li’×’Jin Zhui’ progeny showed three progeny classes (S27S27, S27S34 and S24S34) as opposed to the two classes reported previously (S24S24 and S34S34), indicating that the pollen gametes of ‘Jin Zhui’, bearing either the S27- or S34-haplotype, were able to overcome self-incompatibility (SI) barriers. Moreover, no evidence of pollen-S duplication was found. These findings support the hypothesis that loss of function of S-locus unlinked PPM expressed in pollen leads to SI breakdown in ‘Jin Zhui’, rather than natural mutation in the pollen-S gene (S24 haplotype). Furthermore, abnormal meiosis was observed in a number of pollen mother cells (PMCs) in ‘Jin Zhui’, but not in ‘Ya Li’. These and other interesting findings are discussed.

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Introduction

Gametophytic self-incompatibility (GSI) is a genetic mechanism in many flowering plants that prevents self-fertilization and promotes cross-fertilization. In Solanaceae, Plantaginaceae and Rosaceae, GSI is controlled by a single multi-allelic S-locus, which is comprised of the pistil-S and pollen-S genes [1]. The pistil-S gene encodes a polymorphic ribonuclease (S-RNase) essential for recognition by multiple factor SI system’ [14,30]. Therefore, SC

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accessions reported in *Pyrus* are mostly related to *S*-allele duplications. However, mutations in *S*-locus unlinked factors (also called modifier genes) that are required for GSI had also been associated with SC in Solanaceae and some genera of Rosaceae. In apricot cultivar ‘Canino’ (*S21S23*), a mutation was found in a modifier gene unlinked to the *S*-locus which independently caused the loss of pollen-*S* function [31], [32]. Results of Wu et al. [33] indicated that the SI breakdown in PPM ‘Katy’ apricot was associated with factors unlinked to the *S*-locus. Zuriaga et al. [34] reported another SC North-American apricot cv. Zaigers ‘Katy’, in which the mutated *S*-locus unlinked PPM was located at the distal end of chr.3. Pollen modifier factors have also been identified in *Petunia, Pyrus* and *Prunus*, such as the pollen-expressed Skp1-like1 proteins. As a modifier, Skp1-like1 proteins were proposed to be involved in a SCF complex [17], [35], [36]. Nevertheless, in *Pyrus*, the loss of function of *S*-locus unlinked PPM that leads to self-compatibility has never been reported. Therefore, the identification of *S*-locus unlinked PPM in *Pyrus* will be necessary to give a more complete picture of the self-incompatible mechanism.

‘Jin Zhui’ is a spontaneous self-compatible mutant of SI pear cultivar ‘Ya Li’. Li et al. [21] used genetic analysis of a small population to show that breakdown of SI in ‘Jin Zhui’ might be caused by pollen-*S*-locus mutation [21]. However, it is not clear what form of mutation led to the breakdown of SI in ‘Jin Zhui’. In this work, genetic populations of selfed ‘Jin Zhui’ progeny and crossed progeny with ‘Jin Zhui’ were constructed. We analyzed the self-compatible ‘Jin Zhui’ using genetic and molecular approaches, with the compiled evidence suggesting that loss of function of the *S*-locus unlinked PPM was probably involved in pollen-*S*-function breakdown. Further mapping will be necessary to identify the underlying mutant *S*-locus unlinked PPM.

### Materials and Methods

**Ethics Statement**

No specific permits were required for the described observation or field studies. For locations and activities partaken, a. no specific permissions were required; b. locations were not privately-owned or protected in any way and c. no endangered or protected species were involved.

**Plant Material**

Five cultivars of Chinese pear (*Pyrus bretschneideri* Rehd.), ‘Ya Li’ (*S12S31*), ‘Jin Zhui’ (*S8*), bud mutant of ‘Ya Li’), ‘Jin Feng’ (*S1, S21S23*), ‘Han Hong’ (*S8, S22S23*), ‘Han Xiang’ (*S1, S12S23*), ‘Jin Zhui’ selfed progeny, and crossed progeny derived from ‘Ya Li’ × ‘Jin Zhui’, ‘Ya Li’ × ‘Jin Feng’, ‘Jin Zhui’ × ‘Jin Feng’, ‘Jin Zhui’ × ‘Han Hong’, ‘Ya Li’ × ‘Han Hong’, ‘Han Xiang’ × ‘Jin Zhui’, ‘Ya Li’ × ‘Jin Zhui’, ‘Jin Zhui’ × ‘Jin Feng’, ‘Ya Li’ × ‘Jin Feng’ and two self-pollinations from ‘Ya Li’ and ‘Jin Zhui’, were performed in the field. Pollination was conducted as follows: all flowers were removed before anthesis except three per inflorescence, with the anthers in these three removed and insect-proof bags used to prevent accidental contamination by foreign pollen. Fruit set was recorded about 1 month later and fruits were collected up until harvest time, with the number of seeds and weight per fruit recorded.

**Genome DNA and RNA**

Genomic DNA from leaves and seedlings was isolated by the method described previously [37], [38], and then incubated with RNase I (Invitrogen, CA, USA) at 37°C for 2 hours to remove the RNA, after which it was quantified by spectrophotometer. RNA was extracted from leaves, styles, and pollen samples according to a modified SDS method [39] and digested with DNase I (TaKaRa; DaLian, China). RNA was detected by electrophoresis. Total RNA was used to synthesize first-strand cDNA using the SuperScript reverse transcriptase (Invitrogen, CA, USA) with Oligo-dT primer. The cDNA was then used as templates for PCR amplification.

**Genetic Linkage Analysis between Pollen-expressed SLM Alleles and *S*-RNases**

Primers were screened from the *PpSFBB* primers reported by KaKui et al. [14]. Four primer pairs (Table 1) were chosen for amplifying the pollen-expressed-SLM alleles in pollen cDNAs from ‘Ya Li’ and ‘Jin Zhui’. PCR conditions were as follows: 20 µL reaction system, 50 ng genomic DNA, 2 µL 10× Ex Taq buffer (including 2 mM MgCl2), 0.2 mM dNTPs, 10 µM of each primer and 0.2 unit Ex Taq DNA polymerase (TaKaRa). PCR products were examined by 1% agarose gel, and purified using the Gel Extraction Kit (BioDev-Tech, Beijing, China) and then cloned into the pMD18-T vector (TaKaRa). Four independent clones of each PCR product were chosen for DNA sequencing. Sequence alignment was performed with MEGA version 5.0 [40]. Phylogenetic analyses of the amino acid sequences of *SLF/SFBB* were carried out using the neighbor-joining method implemented in MEGA, with 1000 bootstrap replicates. Homologs were identified by BLASTN searches of the National Center for Biotechnology Information database (NCBI; [41]). Specific primer pairs for SF alleles were designated for genetic linkage analysis of *SLF* alleles (Table 1). A segregating population of 30 individuals from ‘Ya Li’ (*S21S34*) × ‘Jin Feng’ (*S13S14*) was used.

**Pollination Tests**

Six cross-pollinations, ‘Ya Li’ × ‘Han Hong’, ‘Jin Zhui’ × ‘Han Hong’, ‘Han Xiang’ × ‘Jin Zhui’, ‘Ya Li’ × ‘Jin Zhui’, ‘Jin Zhui’ × ‘Jin Feng’, ‘Ya Li’ × ‘Jin Feng’ and two self-pollinations from ‘Ya Li’ and ‘Jin Zhui’, were performed in the field. Pollination was conducted as follows: all flowers were removed before anthesis except three per inflorescence, with the anthers in these three removed and insect-proof bags used to prevent accidental contamination by foreign pollen. Fruit set was recorded about 1 month later and fruits were collected up until harvest time, with the number of seeds and weight per fruit recorded.

**Pollen Viability and Germination Tests**

Pollen viability was estimated as the percentage of pollen grains stained with 0.5% fluorescein diacetate (FDA). Fresh pollen grains were germinated on a solid medium (0.01% boric acid, 1% agar, 8% sucrose) in petri dish for 40 min at 28°C, and then the solid medium with germinated pollen grains was cut into small patches and placed on a glass slide. The germinated pollen grains were observed directly under a microscope (Nikon Eclipse 80i, Japan) for germination tests. For tests of pollen viability, FDA (2.5 mg/ml) was added to solid medium for 10 min, and then the pollen grains were observed under a microscope (Nikon Eclipse 80i, Japan). A total of 500 pollen grains were observed.

**Chromosome Preparation and Observation**

Upon reaching the optimal stage for meioitic studies, flower buds of ‘Jin Zhui’ and ‘Ya Li’ were fixed in canary’s fluid (3 parts of ethanol plus 1 part glacial acetic acid) for 24 hours and then stored in 70% ethanol at 4°C until used. Pollen mother cells were stained with 4’, 6-diamino-2-phenylindole (DAPI) before observation. The chromosome numbers of ‘Jin Zhui’ selfed progeny were observed.
Molecular Identification of Results

Root-tips were harvested when 1–2 cm long from germinated seeds and pretreated in 0.002 M 8-Hydroxyquinoline for 2 hours. Materials were washed twice with distilled water (Table 1). The PpSFBB4 primers were specific for the S21-haplotype of the SLF gene. The expression analysis of these SLF genes in different tissues showed they were all expressed specifically in pollen (Fig. 1 B).

To confirm whether the five SLF genes were linked to the S locus, a population of 30 progeny from ‘Ya Li’ × ‘Jin Feng’ was used. When 15 progeny carrying S21S34 contained a single band that was absent in the other 15 progeny which carried S21S19, the SLF gene was assigned S21-haplotype-linkage. Similarly, if a single band only appeared in the progeny carrying S21S34, the SLF gene was assigned S21-haplotype-linked. Two of the SLF genes were detected in the S21-containing progeny only, and the other three were detected in the S19-containing progeny, suggesting linkage between the PbSLFs and PbS-RNases (Fig. 1 C). These SLFs were subject to phylogenetic analysis with SLF/SFBBs from Japanese pear and Chinese pear, based on which they were classified into three types: type-1, type-3 and type-4 according to Kakui et al. [14] (Fig. 1 D). Thus, they were named as PbSLF1-S21 (GenBank accession number KC569798), PbSLF1-S34 (GenBank accession number KC569799), PbSLF2-S21 (GenBank accession number KC569800), PbSLF3-S34 (GenBank accession number KC569801) and PbSLF4-S21 (GenBank accession number KC569802).

To determine whether mutations or indels existed in the SLFs from ‘Jin Zhui’, seven SLFs including the five obtained in this study and two identified by Xu et al. [17] were cloned from pollen cDNA of ‘Jin Zhui’ and ‘Ya Li’. Comparative analysis of the seven SLFs showed no difference between ‘Ya Li’ and ‘Jin Zhui’ (data not shown).

Fruit Set and Genetic Analysis of Selfed and Crossed Progeny of ‘Jin Zhui’

Li et al. [21] found that ‘Jin Zhui’ was a pollen-part mutant of ‘Ya Li’. Genetic analysis of 29 selfed progeny of ‘Jin Zhui’ indicated the SI breakdown in ‘Jin Zhui’ might be due to the pollen-S of the S21-haplotype. To further analyze the nature of the self-compatible mutant, pollination tests were performed. ‘Ya Li’ and ‘Jin Zhui’ were self-pollinated and reciprocally pollinated. The results from pollination tests showed that ‘Jin Zhui’, which had functional style and nonfunctional pollen, was self-compatible, while ‘Ya Li’, which had functional style and pollen, was self-incompatible (Table 2). Further genetic analysis of larger populations of selfed and crossed progeny of ‘Jin Zhui’ was performed. append:dword:respectiveSegregation of S-RNases in selfed progeny of ‘Jin Zhui’ and crossed progeny between ‘Ya Li’ and ‘Jin Zhui’ was investigated by PCR using the S-haplotypespecific S-RNase primers. The results showed that both the selfed progeny of ‘Jin Zhui’ and the crossed progeny between ‘Ya Li’ and ‘Jin Zhui’ consisted of three S-genotypes: S21/S34, S21/S19 and S19/S34 (Fig. 2 A, B and Table 3). The results indicated that pollen carrying both pollen S-alleles could reach the ovule and achieve self-fertilization, which suggested that the pollen-part mutation in ‘Jin Zhui’ was unlinked to the S-locus.

To further validate these observations, additional crosses were performed and analyzed (Fig. 2 C, D and Table 3). The crossed progeny of ‘Han Xiang’ × ‘Jin Zhui’ and ‘Jin Zhui’ × ‘Jin Feng’ fell into four classes (S21S34×S21S19, S21S34×S19S34, S19S34×S21S19 and S19S34×S19S34) by S-RNase genotyping, respectively. The observed ratios for S-genotype segregations fit with the expected ratios (χ² from root-tip cells with at least 30 cells observed per sample, fresh root-tips were harvested when 1–2 cm long from germinated seeds and pretreated in 0.002 M 8-Hydroxyquinoline for 2 hours at room temperature, and then fixed with cannyol solution for 24 hours. After washing twice with distilled water, the root-tips were cut and kept in a solution of 2% cellulose and 1% pectinase at 37℃ for 2 hours. Materials were washed twice with distilled water before being fixed again with cannyol solution and then stained on a pre-cooled slide with 4’, 6-diamino-2-phenylindole pyrrolindone (DAPI). All chromosomal images were captured under the Olympus BX53 fluorescence microscope used with a microCCD camera.

Results

Molecular Identification of SLFs in ‘Ya Li’ and ‘Jin Zhui’

PCR amplification of the SLF alleles was performed using four different primer pairs as used for SFBB allele identification by Kakui et al. [14] (Table 1). Single bands were obtained when amplifying SLF alleles from the genomic DNA of ‘Ya Li’ (S21S34) using each primer pair (Fig. 1 A). Direct sequencing of these PCR products validated the five SLF candidates. Specific primer pairs for each SLF were generated on the basis of sequence alignments of the cloned SLFs (Table 1). The PpSFBB4 primers were specific for the S21-haplotype of the SLF gene. The expression analysis of these SLF genes in different tissues showed they were all expressed specifically in pollen (Fig. 1 B).

| primer | Sequence(from 5’ to 3’) | note |
|-------|------------------------|------|
| S12-F | AGTTGGAATTTTTGACGAGC | This work |
| S12-R | TCAACACCTTCGATCATCTGTC | This work |
| S19-F | GACCAAAATATGCCAAGGCG | [17] |
| S19-R | TTGGTTCACTTGAAAGGAGAC | [17] |
| S21-F | ATATTGGAAGGAGAATCAG | [17] |
| S21-R | ATATTGGAAGGAGAATCAG | [17] |
| S31-F | AAGACCACAGAGTTGCGAAGAACA | This work |
| S31-R | TTTCCAACTGGGTTCTCAGATTGTC | This work |

| primer | Sequence(from 5’ to 3’) | note |
|-------|------------------------|------|
| PbSLF3-S34-F | F: TTATCTTCTACATTTATGACATGAG | [17] |
| PbSLF3-S34-R | R: ATACAGAAAAATGGTGTCG | [17] |
| PbSLF6-S21-F | CATACAATTTAATTAAGAAGAGATG | [17] |
| PbSLF6-S21-R | AACTCCTCATTGATGCTTAGC | [17] |
| PbSLF1-S21-F | ATGTCCTCCGTGATGAgAT | This work |
| PbSLF1-S21-R | ATGTTATTCCTCCGATCGGG | This work |
| PbSLF1-S34-F | ATGTCCTCCGTGATGAgAT | This work |
| PbSLF1-S34-R | CAATCCATTGCAATGCCG | This work |
| PbSLF2-S34-F | AAATTTGGTCAGGTTGCGC | This work |
| PbSLF2-S34-R | GCATCCAAATCAAACACCTCTGC | This work |
| PbSLF3-S34-F | TCTTTGATGACATCTCAG | This work |
| PbSLF3-S34-R | CAATAACAAATACCCCTG | This work |

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values of 1.31, $P>0.05$ and 0.29, $P>0.05$). These results supported self-compatibility (SC) in ‘Jin Zhui’ was associated with S-locus unlinked PPM rather than S-allele duplications.

The homozygotes from selfed progeny of ‘Jin Zhui’ and crossed progeny of ‘Ya Li’ x ‘Jin Zhui’ were much fewer in number than heterozygotes (Table 3). The average number of seeds per fruit produced from selfed progeny of ‘Jin Zhui’ and crossed progeny of ‘Ya Li’ x ‘Jin Zhui’ was 3.91 and 4.13, respectively. This was much lower than the two cross-pollinations ‘Jin Zhui’ x ‘Han Hong’ (7.09) and ‘Ya Li’ x ‘Han Hong’ (7.75) (Fig. 2 E).

Figure 1. Molecular identification of SLFs in ‘Ya Li’. A. PCR products from genomic DNA of ‘Ya Li’ for SLF allele identification. Lanes 1–5 represent PCR products using primers PpSFBB1, PpSFBB2, PpSFBB3, and PpSFBB4, respectively. M, DNA marker. B. Expression of PbSLFs in leaf, style and pollen in ‘Ya Li’. RT-PCR was performed with SLF-specific primers and synthesized cDNA was used as templates. Pear actin gene was used as internal control. C. Linkage analysis of PbSLFs and PbS-RNases. Genomic DNAs from 30 crossed progeny of ‘Ya Li’ ($S_{J2}S_{K2}$) and ‘Jin Feng’ ($S_{J2}S_{K2}$) were used as templates for linkage analysis. PbS$_{21}$-RNase, PbS$_{26}$-RNase and PbS$_{34}$-RNase (top panel) and PbSLFs (lower panel) were detected by PCR using gene specific primers. Lanes 1–30 represent a progeny population of 30 individuals. D. Phylogenetic tree of deduced amino acid sequences of PbSLFs and other SLFs/SFBBs. A neighbor-joining tree was constructed from 49 SLF/SFBB proteins from Japanese pear and 11 PbSLF proteins from Pyrus bretschneideri. PhSLF-S3 and PhSLF-S3L from Petunia hybrida were used as an outgroup. Numbers on the branches showed bootstrap values above 50% from 1000 bootstrap replicates. Eight types of the SLF/SFBB proteins were classified by Kakui et al. [14]. PbSLFs obtained in this study were marked in red.

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Figure 2. Segregation of $S$-RNases in selfed and crossed progeny of 'Jin Zhui'.

A. Segregation analysis of $S_{21}$- and $S_{34}$-RNases in selfed progeny of 'Jin Zhui'. Lanes 1–94 represent a progeny population of 94 individuals.

B. Segregation analysis of $S_{21}$- and $S_{34}$-RNases in crossed progeny between 'Ya Li' and 'Jin Zhui'. Lanes 1–59 represents a progeny population of 59 individuals.

C. Segregation analysis of $S$-alleles in crossed progeny between 'Han Xiang' and 'Jin Zhui'. Lanes 1–45 represent a progeny population of 45 individuals.

D. Segregation analysis of $S$-alleles in crossed progeny between 'Jin Zhui' and 'Jin Feng'. Lanes 1–31 represent a progeny population of 31 individuals.

E. Average numbers of seeds per fruit from selfed and crossed progeny of 'Jin Zhui'. JZ, Jin Zhui; YL, Ya Li; HH, Han Hong.

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seggregation ratios of the three classes (S$_{21}$S$_{21}$; S$_{21}$S$_{34}$; S$_{34}$S$_{34}$) from selfed progeny of ‘Jin Zhui’ and crossed progeny between ‘Ya Li’ and ‘Jin Zhui’ were 9:74:1 and 6:45:9, respectively. These segregation ratios did not fit the expected ratios of 1:2:1, with χ² values of 31.11, P<0.01 and 14.56, P<0.01 (Table 3). The segregation distortion showed that there were significantly fewer segregation ratios of the three classes (S$_{21}$S$_{21}$; S$_{21}$S$_{34}$; S$_{34}$S$_{34}$) from selfed progeny from ‘Jin Zhui’ and crossed progeny between ‘Ya Li’ and ‘Jin Zhui’ selfed 150 119 79.3 50

| Crosses          | Pollinated flowers | Fruits set | Fruit-set per. (%) | Pollinated inflorescence No. |
|------------------|--------------------|------------|--------------------|------------------------------|
| ‘Ya Li’ selfed   | 117                | 3          | 2.6                | 39                           |
| ‘Jin Zhui’ selfed| 150                | 119        | 79.3               | 50                           |
| ‘Jin Zhui’ × ‘Ya Li’ | 106             | 5          | 4.7                | 36                           |
| ‘Ya Li’ × ‘Jin Zhui’ | 150            | 124        | 82.7               | 53                           |
| ‘Ya Li’ × ‘Han Hong’ | 133           | 103        | 77.4               | 50                           |
| ‘Jin Zhui’ × ‘Han Hong’ | 120          | 104        | 86.7               | 42                           |

Significant Difference in Abortive Pollen between ‘Jin Zhui’ and ‘Ya Li’

To investigate the pollen-part mutation of ‘Jin Zhui’, we observed the pollen germination ability of ‘Jin Zhui’ and ‘Ya Li’ (Fig. 3 A), where a significant difference was found. The number of abortive pollen grains was found to be much increased in ‘Jin Zhui’, with the abortive pollen grains incapable of both hydration and germination. The abortive pollen grains of ‘Jin Zhui’ and ‘Ya Li’ could not be stained by FDA, indicating they were inviable (Fig. 3 B). The percentage of abortive pollen grains in ‘Jin Zhui’ (about 20%) was much higher than in ‘Ya Li’ (about 1.6%) (Fig. 3 C), which was consistent with the results observed during pollen germinations. Further DAPI staining showed that the abortive pollen grains had no apparent nucleus (Fig. 3 D), and that they had already degenerated into shells. The increase in these types of abortive pollen grains in ‘Jin Zhui’ implied partial disruption of normal meiosis during the development of pollen.

Abnormal Meiosis of Pollen Mother Cells (PMCs) in ‘Jin Zhui’

In order to investigate the abortive pollen grains in ‘Jin Zhui’, we analyzed the meiotic behavior of pollen mother cells (PMCs). As a self-incompatible cultivar, ‘Ya Li’ had normal meiotic behavior (Fig. 4 A-J). ‘Ya Li’ possesses 34 chromosomes, which during meiotic prophase I, undergo leptotene (not shown), zygote (not shown), pachytene (Fig. 4 A), diplotene (not shown) and finally, the 17 bivalents condense through diakinesis (Fig. 4 B). The 17 bivalents then distribute on the equatorial plate at metaphase I (Fig. 4 C). During anaphase I, each group of homologous chromosomes separated from each other (Fig. 4 D), with each group reaching one pole of the cell (Fig. 4 E). Through meiotic prophase II (Fig. 4 F) and metaphase II (Fig. 4 G), the sister chromatids in each group were then separated in anaphase II (Fig. 4 H), giving four pools of 17 chromosomes (Fig. 4 I), which gave rise to four tetrads of microspores (Fig. 4 J). As a self-compatible pollen-part mutant of ‘Ya Li’, ‘Jin Zhui’ had abnormal meiosis in a number of PMCs (Fig. 4 K–T). Univalents appeared at metaphase I (Fig. 4 K and L), and laggards prevailed during anaphase I (Fig. 4 M, N, and O). During metaphase II, univalents also appeared (Fig. 4 P and Q), and then laggards and unbalanced separation were observed in anaphase II (Fig. 4 R, S and T). Unbalanced separation produced many abortive pollen grains which had no nucleus nor vitality. The number of meiotic abnormalities in ‘Jin Zhui’ PMCs was recorded (Table 4), and showed that abnormal meiosis existed in a number of PMCs of ‘Jin Zhui’. To test whether the abnormal meiosis affected the selfed progeny, we observed chromosome numbers of selfed progeny from ‘Jin Zhui’. Both the homozygotes and heterozygotes in the progeny had the usual (34) number of chromosomes (Fig. 5). Though abnormal meiosis was found in ‘Jin Zhui’ PMCs, the selfed progeny from ‘Jin Zhui’ had normal chromosome numbers.

### Table 2. Fruit setting rates in self- or cross-pollination of three pear cultivars.

| Crosses          | Pollinated flowers | Fruits set | Fruit-set per. (%) | Pollinated inflorescence No. |
|------------------|--------------------|------------|--------------------|------------------------------|
| ‘Ya Li’ selfed   | 117                | 3          | 2.6                | 39                           |
| ‘Jin Zhui’ selfed| 150                | 119        | 79.3               | 50                           |
| ‘Jin Zhui’ × ‘Ya Li’ | 106             | 5          | 4.7                | 36                           |
| ‘Ya Li’ × ‘Jin Zhui’ | 150            | 124        | 82.7               | 53                           |
| ‘Ya Li’ × ‘Han Hong’ | 133           | 103        | 77.4               | 50                           |
| ‘Jin Zhui’ × ‘Han Hong’ | 120          | 104        | 86.7               | 42                           |

### Table 3. Segregation of S-genotypes in progeny of different self- or cross-pollinations.

| Crosses          | Total | S-genotypes observed in progeny | Expected segregation ratio* | χ² | P-value |
|------------------|-------|--------------------------------|-----------------------------|----|---------|
| ‘Jin Zhui’ selfed| 94    | S$_{21}$S$_{21}$, S$_{21}$S$_{34}$, S$_{34}$S$_{34}$, S$_{21}$S$_{21}$, S$_{21}$S$_{34}$, S$_{34}$S$_{34}$, S$_{21}$S$_{31}$, S$_{31}$S$_{34}$ | 1:2:1                        | 31.11 | P<0.01 |
| ‘Ya Li’ × ‘Jin Zhui’ | 59    | 9                              | 1:2:1                        | 14.56 | P<0.01 |
| ‘Jin Zhui’ × ‘Jin Feng’ | 31    | 17                             | 1:1                          | 0.29 | P>0.05 |
| ‘Han Xiang’ × ‘Jin Zhui’ | 45    | 10                             | 1:1:1                        | 1.31 | P>0.05 |

*Expected ratios for a single mutation unlinked to the S-locus.

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Discussion

Identification and Comparative Analysis of SLF Genes between ‘Ya Li’ and ‘Jin Zhui’

In gametophytic self-incompatible (GSI) species of Solanaceae, Rosaceae and Plantaginaceae, multiple F-box genes within the S-locus region have been reported [6–8], [12], [42]. Through mutant analysis and transgenic approaches, the pollen S-genes had
Figure 3. Significant difference in abortive pollen grains between ‘Jin Zhui’ and ‘Ya Li’. A. Pollen grains germination of ‘Jin Zhui’ (left) and ‘Ya Li’ (right). Abortive pollen grains without hydration and germination are indicated with arrows. Scale bar: 100 μm. B. FDA staining of germinated pollen grains from ‘Jin Zhui’ (left) and ‘Ya Li’ (right). Abortive pollen grains not stained by FDA are indicated by arrows. Scale bar: 100 μm. C. Percentages of abortive pollen grains in FDA staining between ‘Jin Zhui’ and ‘Ya Li’. A total of 500 fresh pollen grains were examined in each case. D. DAPI staining of pollen grains from ‘Jin Zhui’ (left) and ‘Ya Li’ (right). Abortive pollen grains without apparent nucleus are indicated by arrows. Scale bar: 50 μm.

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Figure 4. Meiosis of ‘Ya Li’ and ‘Jin Zhui’. (A–J). DAPI staining of ‘Ya Li’ PMCs during meiosis. (A)–(J) show pachytene, diakinesis, metaphase I, anaphase I, telophase I, prophase II, metaphase II, anaphase II, telophase II and tetrad, respectively. (K–T) DAPI staining of ‘Jin Zhui’ PMCs during meiosis. (K) and (L) show univalents in metaphase I, (M), (N) and (O) laggard chromosomes at anaphase I, (P) and (Q) univalents in metaphase II, (R), (S) and (T) laggard chromosome and unbalanced separation in anaphase II, respectively. Scale bar: 5 μm.

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been determined experimentally [9–11], [24], [43]. In Petunia and Pyrus, multiple SLFs/SFBs as pollen S factors were proven to collaborate in recognizing non-self S-RNases and mediate their degradation [14,30]. In previous studies, mutations or indels in pollen-S genes were always considered a reasonable explanation for self-compatible PPM, where Li et al. [21] reported that ‘Jin Zhui’ was a pollen-part mutant. The breakdown of self-incompatibility (SI) in ‘Jin Zhui’ might be caused by mutant pollen-S34 genes. Our study on pollen-part mutation of ‘Jin Zhui’ should help us to understand the function of the pollen-S gene. We cloned five F-box genes from pollen cDNA of ‘Ya Li’, with linkage analysis showing them to be S-Locus F-box (SLF) genes. However, comparative analysis of seven SLFs, including the two SLFs identified by Xu et al. [17], showed no differences between ‘Ya Li’ and ‘Jin Zhui’. Therefore, these seven SLFs were probably not directly responsible for the SI breakdown of ‘Jin Zhui’. Subsequent genetic analysis further supported this hypothesis.

**Self-compatibility in ‘Jin Zhui’ is Associated with S-locus Unlinked PPM**

‘Ya Li’ (S21S34) is a traditional Chinese pear (Pyrus bretschneideri) cultivar exhibiting self-incompatibility, and ‘Jin Zhui’ (S21S34) is a pollen-part mutant of ‘Ya Li’. Genetic analysis of 29 selfed progeny in ‘Jin Zhui’ showed that the pollen S34 allele might be mutated in ‘Jin Zhui’ [21].

To investigate the genetics of SC in ‘Jin Zhui’, genetic populations were constructed. ‘Jin Zhui’ (S21S34) was self-pollinated and reciprocally crossed with ‘Ya Li’ (S21S34). It is noteworthy that ‘Jin Zhui’ pollen tubes bearing either the S21R haplotype were able to grow through ‘Jin Zhui’ and ‘Ya Li’ pistils and complete fertilization, producing three S-genotype classes (S21S21, S21S34 and S34S34) instead of the two classes observed previously (S21S21 and S21S34). However, no progeny were obtained in the reciprocal cross when using ‘Jin Zhui’ as the female parent. These results should support the PPM in ‘Jin Zhui’ was unlinked to the S-locus. The crosses between ‘Han Xiang’ (S21S34) and ‘Jin Zhui’ (S21S34), ‘Jin Zhui’ (S21S34) and ‘Jin Feng’ (S21S21) reinforced this conclusion.

Interestingly, in the ‘Jin Zhui’ × ‘Jin Zhui’ and ‘Jin Zhui’ × ‘Ya Li’ populations the number of seedlings homozygous for both the S21R and the S34R haplotype is significantly lower than that for the heterozygote S21R S34R (see Table 3). Some explanations for this phenomenon may include linkage in coupling between the mutated allele of the S-locus unlinked PPM and the S-allele, or postzygotic selection against homozygous embryos. In the case of the former, the segregation ratios observed in different populations do not support linkage between the mutated factor and the S-locus. Whereas postzygotic selection would explain the significantly reduced numbers of S21S21 and S34S34 genotypes, and could also be a reasonable explanation that no S21S34 homozygotes were detected previously by Li et al. [21] in a small population from selfed progeny of ‘Jin Zhui’.

In Solanaceae, self-compatible pollen-part mutants may arise from S-allele duplications located in a centric fragment, in a non-S chromosome, or linked to the S-locus leading to the formation of S-heteroallelic pollen [27]. According to the segregations obtained in the crosses (including ‘Jin Zhui’ selfed, ‘Ya Li’ × ‘Jin Zhui’, ‘Jin Zhui’ × ‘Jin Feng’, ‘Han Xiang’ × ‘Jin Zhui’) performed here, S-allele duplications did not seem likely in ‘Jin Zhui’ (all descendants should have had the S21S34 genotype) (Fig. 2, Table 3). S-allele duplications may also result from polyploidy, but ‘Jin Zhui’ was confirmed as diploid by chromosome observation in selfed progeny (Fig. 5). These results rule out competitive interaction resulting from S-heteroallelic pollen as the cause of SC in ‘Jin Zhui’. Taken together, these results supported the hypothesis that an S-locus unlinked PPM was required in GSI. Loss of function in this S-locus unlinked PPM was responsible for the SI breakdown of ‘Jin Zhui’.

**Possible Role of the S-locus Unlinked PPM in Meiosis**

Abnormal meiotic behaviors in PPM ‘Jin Zhui’ were observed in many PMCs (Fig. 4 and Table 4). Univalents, laggards and unbalanced separation were detected during the process of meiosis (Fig. 4). In anaphase II, unbalanced separation produced a number of abnormal microspores, which were degraded into shells (Fig. 3). Meanwhile, to test whether the abnormal meiosis affected the selfed progeny, we observed chromosome numbers of selfed progeny from ‘Jin Zhui’, and found that both the homozygotes and heterozygotes in the progeny had the usual 34 chromosomes (Fig. 5). Though abnormal meiosis was found in ‘Jin Zhui’ PMCs, the selfed progeny from ‘Jin Zhui’ had normal chromosome numbers and were not influenced by abnormal meiosis. Abnormal meiosis in PMCs was consistently caused by related mutant genes. In an Arabidopsis male sterile mutant, abnormal meiosis caused by a Ds insertion in the SKP1-LIKE1 gene was reported [44]. While the SKP1-LIKE1 homolog SSK (SLF-interacting Skp1-like1) was

### Table 4. Meiotic abnormalities in ‘Jin Zhui’.

| Phases      | Total number of cells | Number of abnormal cells | Abnormalities                  | Percentage of abnormal cells (%) |
|-------------|-----------------------|--------------------------|--------------------------------|---------------------------------|
| Metaphase I | 625                   | 283                      | univalents                     | 45.28                           |
| Anaphase I  | 731                   | 172                      | Laggards                       | 23.53                           |
| Metaphase II| 563                   | 169                      | univalents                     | 30.01                           |
| Anaphase II | 612                   | 244                      | laggers and unbalanced separation | 39.87                          |

**Table 4. Meiotic abnormalities in ‘Jin Zhui’**

![Figure 5. Normal chromosome numbers in selfed progeny of ‘Jin Zhui’. A. 34 chromosomes of an S21S21 homoyzogote representing a total of 12 progeny examined. B. 34 chromosomes of an S21S34 heterozygote representing a total of 21 progeny examined. C. 34 chromosomes of an S34S34 homoyzogote representing a total of 13 progeny examined. Scale bar: 5 μm. doi:10.1371/journal.pone.0076509.g005](http://www.plosone.org/figure/5-normal-chromosome-numbers-in-selfed-progeny-of-jinzhui-a-34-chromosomes-of-an-s21s21-homoyzogote-representing-a-total-of-12-progeny-examined-b-34-chromosomes-of-an-s21s34-heterozygote-representing-a-total-of-21-progeny-examined-c-34-chromosomes-of-an-s34s34-homoyzogote-representing-a-total-of-13-progeny-examined-scale-bar-5-μm-doi10.1371journal.pone.0076509g005)
identified as a putative canonical SCF(SL) complex, and proposed to degrade non-self S-RNase in the degradation model of Pyrus, prunus and Antirrhinum [17, 33, 36, 45]. In this work, we suspect the mutant S-box linked PPM might influence the pollen development of ‘Jin Zhui’. Two S-boxes, which had been identified in ‘Ya Li’ [17], were cloned from pollen cDNAs of ‘Jin Zhui’ and ‘Ya Li’, but no nucleotide difference was found (data not shown). Further studies focusing on the identification of S-box linked PPM should help us to gain deeper insights into self-incompatibility in Pyrus.

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Author Contributions

Conceived and designed the experiments: JKWF MFL. TZL. Performed the experiments: JKWF MFL. Analyzed the data: JKWF MFL. Contributed reagents/materials/analysis tools: JKWF MFL. Wrote the paper: JKWF MFL. TZL.

References

1. De Nettancourt D (1997) Incompatibility in angiosperms. Sex Plant Reprod 10: 115–199. doi:10.1007/BF02982745.
2. Anderson MA, Cornish EC, Mau SL, Williams EG, Hoggart R, et al. (1986) Cloning of a cytoplasmic B-style self-incompatibility gene from Nicotiana alata. Nature 321: 38–44. doi:10.1038/321038a0.
3. McClure BA, Haring V, Ebert PR, Anderson MA, Simpson RJ, et al. (1989) Self-incompatibility gene products of Nicotiana alata are ribonucleases. Nature 342: 953–957. doi:10.1038/3429535a0.
4. Sassa H, Hirano I, Ikemashi H (1992) Self-incompatibility-related RNases in styles of Japanese pear (Pyrus serotina Rehd.). Plant Cell Physiol 33: 811–814.
5. Xu C, Zhang B, Zhao Z, Wang X, Skirpan AL, Wang Y, Dowd PE, et al. (2004) Identification of a canonical SCFSLF complex, and proposed localization of an S haplotype-specific F-box protein gene, SFB, is defective in self-compatible mutants, Yan Zhuang and Jin Zhui. Plant Biol (Stuttgart) 11: 774–783. doi:10.1111/j.1438-2379.2008.00180.x.
6. Sijacic P, Wang X, Skirpan AL, Wang Y, Dowd PE, et al. (2004) Identification of a novel protein with an F-box motif that is very tightly linked to a gene for S-RNase in two species of cherry, Prunus cerasus and P. avium. Plant Cell Physiol 45: 764–769. doi:10.1093/pcp/pci053.
7. Okada K, Nakata N, Moriya Y, Norioka N, Sawamura Y, et al. (2008) Deletion of a 236 kb region around S-RNase in a self-pollinator mutant S4-RNase of Japanese pear. Plant Mol Biol 66: 789–800. doi:10.1007/s11103-007-9277-1.
41. Altschul SF, Gish W, Miller W, Myers EW, Lipman DJ (1990) Basic local alignment search tool. J Mol Biol 215: 403–410. doi:http://dx.doi.org/10.1016/S0022-2836(05)80360-2.
42. Wheeler D, Newbigin E (2007) Expression of 10 S-class SLF-like genes in Nicotiana alata pollen and its implications for understanding the pollen factor of the S locus. Genetics 177: 2171–2180. doi:10.1534/genetics.107.076885.
43. Tsukamoto T, Hauck NR, Tao R, Jiang N, Iezzoni AF (2006) Molecular characterization of three non-functional S-haplotypes in sour cherry (Prunus cerasus). Plant Mol Biol 62: 371–383. doi:10.1007/s11103-006-9026-x.
44. Yang M, Hu Y, Lodhi M, McCombie WR, Ma H (1999) The Arabidopsis SKP1-LIKE gene is essential for male meiosis and may control homologue separation. Proc Natl Acad Sci 96: 11416–11421. doi:10.1073/pnas.96.20.11416.
45. Chen G, Zhang B, Liu L, Li Q, Zhang Y, et al. (2012) Identification of a ubiquitin-binding structure in the S-Locus F-Box protein Controlling SRNase-based self-incompatibility. J Genet and Genom 39: 93–102. doi:http://dx.doi.org/10.1016/j.jgg.2012.01.001.