PPR647 Protein Is Required for Chloroplast RNA Editing, Splicing and Chloroplast Development in Maize

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Abstract: Chloroplasts play an essential role in plant growth and development. Any factors affecting chloroplast development will lead to abnormal plant growth. Here, we characterized a new maize mutant, albino seedling mutant as-81647 (as-81647), which exhibits an entirely albino phenotype in leaves and eventually died before the three-leaf stage. Transmission electron microscopy (TEM) demonstrated that the chloroplast thylakoid membrane was impaired and the granum lamellae significantly decreased in as-81647. Map-based cloning and transgenic analysis confirmed that PPR647 encodes a new chloroplast protein consisting of 11 pentatricopeptide repeat domains. Quantitative real-time PCR (qRT-PCR) assays and transcriptome analysis (RNA-seq) showed that the PPR647 mutation significantly disrupted the expression of PEP-dependent plastid genes. In addition, RNA splicing and RNA editing of multiple chloroplast genes showed severe defects in as-81647. These results indicated that PPR647 is crucial for RNA editing, RNA splicing of chloroplast genes, and plays an essential role in chloroplast development.

Keywords: pentatricopeptide repeat protein; maize; albino-lethal; RNA editing; RNA splicing; chloroplast

1. Introduction

As the exclusive organelles for photosynthesis and energy transduction in plants, the chloroplast plays a vital role in plant growth and development. However, the process of chloroplast biogenesis and development is highly complex, and its molecular mechanisms have not been fully elucidated. Chloroplasts are semiautonomous organelles, which can encode 60–200 proteins, and their gene expression still undergoes regulation by the nuclear genome [1]. Nucleus-encoded polymerases (NEP) and plastid-encoded polymerases (PEPvc) are necessary enzymes responsible for plastid genes transcription and have significant impacts on chloroplast biogenesis [2–4]. According to the required transcriptase, chloroplast genes can be divided into three categories: Genes transcribed only dependent on PEP (mainly photosynthetic related genes, such as psbA and psbD); genes transcribed only dependent on NEP (mainly housekeeping genes such as accD, rpoa, and rpob); and genes transcribed by NEP and PEP (such as atpE and 16S rRNA [2–4]). In addition, the expression of chloroplast genes is regulated by a series of post-transcriptional processes, including C-to-U RNA editing, intron splicing, and 5′- or 3′terminal maturation [5]. Mutants of those genes often resulted in leaf color variation [6–8].

Pentatricopeptide repeat (PPR) proteins are widely found in plants and play various functions in organellar metabolism. PPR proteins are characterized by tandem repeats of a highly degenerate 35 amino acid motif [9]. According to their tandem motifs, PPR proteins can be divided into two subfamilies: P-type and PLS-type proteins. Based on their
different C termini, PLS-type PPR proteins can be further divided into E, E+, and DYW subgroups [10]. Plant PPR proteins are mainly located in chloroplasts or mitochondria [11–13]. Previous studies showed that most nucleus-encoded PPR proteins are involved in post-transcriptional gene regulation, such as RNA editing, RNA splicing, RNA stability, RNA translation, and RNA maturation [9].

Mutations in mitochondrial-localized PPR proteins are usually characterized by delayed plant growth, abnormal embryonic development, abnormal leaf shape, premature leaf senescence, and reduced seed yields [14–19]. For example, the PLS-PPR DEK36 in Arabidopsis and maize affected mitochondrial transcripts editing and seed development [15]. P-type PPR protein MTL1 affected the initiation of NAD7 gene translation and delayed plant development [14]. Mutations of ZmEMP18 and OsPPR5 caused arrested embryo and early endosperm development by controlling the editing of mitochondrial atp6, cox2, and the cis-splicing of nad4 intron 3, respectively [20]. Chloroplast-localized PPR proteins often affect chloroplasts development, resulting in plant leaf color variations. ZmPPR4 is required for intron splicing of rps12, and the absence of PPR4 can lead to an albino seedling-lethal phenotype [7]. ZmPPR5 insertion mutant had viable embryos and was deficient in chloroplast ribosomes, eventually dying at the seedling stage [8]. Null alleles of PPR protein THA8 are seedling lethal in maize and embryo lethal in Arabidopsis [21]. OSWLS4 is essential for chloroplast RNA group II intron splicing during early leaf development [22].

The reported chloroplast-localized PPR proteins in maize often affect the RNA editing efficiency or the intron splicing efficiency of some chloroplast genes, such as the splicing of trnG affected by PPR5 [6]; rps12 requires PPR4 [8] and EMB2645 [23], and ycf3 and trnA is affected by THA8 [21]. The editing of atpA-1148 requires ZmPPR26 [24], and rps8 is affected by ATP4 [25]. The maize genome encodes more than 600 PPR proteins. Most of their function remains unclear, and no chloroplast-localized PPR protein affecting these two functions at the same time has been reported in maize. To identify additional PPR proteins and elucidate their functions in organelles is vital for understanding plant growth and development. This study identified a novel PLS-type PPR protein, PPR467, which targeted the chloroplast. PPR467 is involved in RNA editing and splicing of plastid genes in developing leaves. Disruption of PPR467 function impaired chloroplast development and plastid gene expression and resulted in an albino-lethal phenotype.

2. Results
2.1. Phenotypic Characterization of as-81647

A lethal albino mutant was isolated from inbred line 81647 and designated as-81647. The as-81647 mutant was albino from germination (Figure 1a–c) and only survived for about ten days. Consistent with the albino phenotype, the total chlorophyll content in as-81647 was significantly lower than that in WT (Figure 1d). Detailed analysis revealed that the chlorophyll a (Chl a), chlorophyll b (Chl b), and carotenoid (Caro) contents of as-81647 were only 5.8%, 1.64%, and 1.60% in WT plants. The ratio of Chl a to Chl b in as-81647 was only 1/10 that in WT (Figure 1e).

2.2. Abnormal Chloroplast Morphology in as-81647

We examined chloroplast structure from two-leaf stage seedlings of WT and as-81647 by TEM (Figure 2a–h). Under normal conditions, chloroplasts in WT plants were crescent-shaped and contained well-developed thylakoid membranes consisting of stroma thylakoids and grana thylakoids (Figure 2a–d). In contrast, numerous vacuole cells without chloroplasts were found in as-81647 (Figure 2a–d). In contrast, numerous vacuole cells without chloroplasts were found in as-81647 (Figure 2e,h) and the remaining chloroplasts were small and severely deformed (Figure 2b,f). The granum lamellae were severely degraded with increased osmiophilic bodies and peroxides (Figure 2c,g).

2.3. Increased Reactive Oxygen Species (ROS) Levels in as-81647

To explore the reason of cell death in as-81647, young leaves were stained with trypan blue and DAB (Figure 2i–l). Deep blue staining showed that the whole blade of as-81647
was staining, while a few areas in WT (Figure 2j), indicating that a large number of cells died in the as-81647 leaves. DAB staining showed a large amount of brown precipitate in as-81647 (Figure 2k) but not in WT (Figure 2l), indicating excessive accumulation of H$_2$O$_2$ in as-81647. These results showed that the cells of as-81647 leaves had died or were dying accompanied by the abnormal accumulation of ROS and that abnormal accumulation of ROS may be the direct cause of cell death in as-81647.

2.4. PLS-Type Pentatricopeptide Repeat (PPR) Protein PPR647, Is Responsible for the as-81647 Albino Phenotype

F$_2$ populations derived from the crosses between B73 and as-81647/+ were used for genetic analysis and gene mapping. Among this monohybrid cross, all F$_1$ plants had typical green leaves and approximately half of the F$_1$ had phenotype-segregating ears. Moreover, the ratio of green to albino leaves in F$_2$ offspring of the segregating ears was 3:1 (Table 1). This result suggested that as-81647 is a recessive mutant.

Figure 1. The phenotype of as-81647. (a−c) Phenotypes of as-81647 after germination, at two days after planting (DAP) (a), 4 DAP (b), and 7 DAP (c). (d) Chl a, Chl b, Caro and total Chl content of WT and as-81647. (e) The ratio of Chl a to Chl b of WT and as-81647. Bar = 1 cm. ** p < 0.01 (Student’s t-test).
Figure 2. Electron microscopy observations of as-81647. (a,e) The number of chloroplasts per unit area in WT (a) and as-81647 (e) plants. (b,f) Chloroplast structure in mesophyll cells and vascular bundle sheath cells in WT (b) and as-81647 (f) plants. (c,g) Comparison of chloroplast structure between WT (c) and as-81647 (g) plants. (d,h) Granum structure in WT (d) and as-81647 (h) plants. M Mesophyll cell, BS Bundle sheath cells, ch chloroplasts, g granum thylakoid, og osmiophilic globule. Bar = 1 μm (b,c,f,g) and bar = 10 μm (a,e,d,h). (i,j) DAB staining of WT (i) and as-81647 (j) leaves. (k,l) Trypan blue staining of WT and as-81647 leaves. Bar = 1 cm.

Table 1. Segregation of leaf color in the F2 population.

| Number | Total | Normal Plants | Mutant Plants | Ratio of Segregation | χ²0.05 (1) |
|--------|-------|---------------|---------------|----------------------|------------|
| 1      | 410   | 309           | 101           | 3/1                  | 0.029      |
| 2      | 426   | 316           | 110           | 3/1                  | 0.153      |
| 3      | 392   | 296           | 96            | 3/1                  | 0.05       |
| 4      | 370   | 283           | 87            | 3/1                  | 0.436      |
| 5      | 398   | 295           | 103           | 3/1                  | 0.164      |

An F2 mapping population from a cross between B73 and as-81647/+ was used to map the candidate gene responsible for the as-81647 phenotype. By assaying 496 F2 albino mutants, we found bnlg1536 and umc1641 on the long arm of chromosome 3 were linked to as-81647, with genetic distances being 14.6 cM and 2.7 cM, respectively. Two extended populations with 840 and 1800 albino plants from different F2 populations were used for fine mapping, and the as-81647 gene was ultimately narrowed down to a 55.08 kb region between markers as-239 and as-254 (Figure 3a). Two candidate genes (Zm00001d044496 and Zm00001d044497) were identified in this region (Figure 3a). Sequencing analysis revealed four SNPs in these two adjacent genes. Further study found that the base substitution at +1034th base of Zm00001d044496 resulted in an amino acid replacement (Asp to Lys). The base substitution at the second splice site of Zm00001d044497 caused a Gly to Ser substitution.
To determine which gene was responsible for the as-81647 phenotype, EMS mutant stock ems4-05741c of Zm00001d044496 and ems4-057444 of Zm00001d044497 were obtained from the Maize EMS Stock Center (http://www.elabcaas.cn/memd; last accessed 8 September 2019). Phenotype analysis revealed that ems4-05741c exhibited albino leaves, and the offspring of (as-81647/+)/(ems4-05741c/+)) showed a 3:1 ratio with green and albino leaves (Figure 3d), demonstrating that they were allelic. However, ems4-057444 plants and the offspring of (as-81647/+)/(ems4-057444/+)) all had green leaves. These results indicated that ems4-05741c could not complement the as-81647 phenotype. Zm00001d044496 is the target gene for as-81647.

Another mutant allele of Zm00001d044496 designated as-cas9-1 was produced by CRISPR-Cas9. The gRNA spacer sequence in the first exon was selected as the target site for Cas9 cleavage (Figure 3e). as-cas9-1 had a 58 bp deletion at the guide RNA (gRNA), creating a premature stop codon in the mature transcript (Figure 3e). The phenotype of as-cas9-1 homozygous produced similar albino seedling to as-81647 (Figure 3f). These results further confirmed that Zm00001d044496 is the target gene.
Zm00001d044496 was predicted to encode a PLS-type PPR protein with 11 PPR motifs using Prosite, named PPR647 (Figure 3c). Among the 11 PPR motifs of PPR647, 10 are in tandem, and another one is interrupted by stretches of several amino acids. To examine the subcellular localization of PPR647, a vector containing 35S: PPR647-YFP was transiently transformed into maize protoplasts. As shown in Figure 4a, the YFP signal was co-localized with the chloroplast, indicating that PPR647 is a chloroplast-localized protein. Phylogenetic tree analysis showed that PPR647 was highly conserved in monocotyledons (Supplementary Figures S1 and S4b). The tree demonstrated that PPR647 is the orthologue of PDM2 of Arabidopsis [26]. Like as-81647 albino seedlings, the seedlings of pdm2 showed a pigment-defective phenotype which nicely supports the phylogenetic link [26]. To assess the role of PPR647 in maize development, we examined its transcript expression pattern by real-time quantitative PCR (qRT-PCR) analysis, the result showed that PPR647 was widely expressed in the root, stem, seed of DAP 20, ear, seven-day leaves (Figure 4c), consistent with the expression pattern shown in the publicly available Maize Gene Expression database (qTell) (https://qteller.maizegdb.org/; last accessed 6 October 2021) (Supplementary Figure S2). At the same time, we found that the expression of PPR647 was significantly down-regulated in as-81647 (Figure 4d).

Figure 4. Gene expression pattern of PPR647. (a) Subcellular localization of PPR647 protein. bar = 5 μm. YFP, Yellow fluorescent protein, BF, Bright field. (b) Phylogenetic tree analysis of PPR647 protein. The red asterisk and arrows indicate PPR647 protein (in B73 background) and PDM2 protein, respectively. PWZ22396 is the PPR647 in Mo17 background. (c) The relative expression level of PPR647 in various tissues was measured by qRT-PCR. RNA was isolated from the DAP20 seeds (S), roots (R), stem (ST), ears (E), seven-day leaves (L). (d) The relative expression level of PPR647 in as-81647 and WT at the three-leaf stage was measured by qRT-PCR. Values and bars represent the mean and standard obtained from three biological replicates, respectively. RNA level was normalized to that of the maize ZmActin gene (Zm00001d010159). Significant differences are indicated. ** p < 0.01 (Student’s t-test).
2.5. PPR647 Mutation Affects Chloroplast-Associated Gene Expression during Leaves Development

To explore the function of PPR647, qRT-PCR and RNA-seq analysis were conducted with the leaves at the two-leaf stage. Transcript levels of chloroplast-associated genes investigated by qRT-PCR results showed that the expression levels of all tested PEP-dependent photosynthesis genes (PEPs) were significantly downregulated (Figure 5a), while the expression of four tested NEP-dependent genes (NEPs) were slightly increased (Figure 5b). Expression levels of other photosynthesis-associated genes encoded by the nucleus were significantly reduced compared with WT (Figure 5c).

From the RNA-seq analysis, 3473 DEGs were identified between WT and as-81647 leaves, including 1766 up and 1671 down-regulated DEGs (Excel S1). Verification of the expression patterns of eleven DEGs via qRT-PCR revealed highly positive correlations between the RNA-seq data and qRT-PCR results (Table S2). According to the RNA-seq result, we found that most chloroplast genome genes were down-regulated, including PEP mediated genes and NEP, PEP co-mediated genes, while only NEP dependent genes were up-regulated (rpoB, rpoA) (Figure 6c). Combining the results of RNA-seq and the qRT-PCR
results led to the conclusion that the mutation of PPR647 affected chloroplast-associated gene transcription, especially PEP mediated chloroplast genes.

KEGG analysis showed that those DEGs were involved in 123 metabolic pathways, 28 of which were significantly enriched (corrected p-value < 0.05) (Figure 6b, Excel S2). Photosynthesis, carbon fixation, and carbon metabolism are the most significant enrichment pathways (Figure 6b). This result showed that PPR647 mainly affected the chloroplast development and photosynthesis of maize, consistent with the qRT-RCR result and albino phenotype.

Furthermore, expression levels of genes encoding ascorbate peroxidase and glutaredoxin (two major scavenger enzymes in the ROS degradation pathway) were mostly up-regulated (Figure 6d). This corresponds to the results of histochemical staining (Figure 6e).

Based on GO analysis, these DEGs were classified into different biological processes or molecular functions. In the cell components, DEGs were mainly concentrated in membrane parts, such as thylakoid, thylakoid membrane, photosynthetic membrane, chloroplast, chloroplast thylakoid, etc. In the biological process, DEGs were mainly enriched into photosynthesis, light reaction, generation of precursor metabolites and energy, and the oxidation-reduction process. At the molecular function level, the main functional items were oxidoreductase activity, tetrapyrrole binding, and chloroplast binding (Figure 6a). KEGG analysis showed that those DEGs were involved in 123 metabolic pathways, 28 of which were significantly enriched (corrected p-value < 0.05) (Figure 6b, Excel S2). Photosynthesis,
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Furthermore, expression levels of genes encoding ascorbate peroxidase and glutaredoxin (two major scavenger enzymes in the ROS degradation pathway) were mostly up-regulated (Figure 6d). This corresponds to the results of histochemical staining (Figure 2i–l). In as-81647, the increase of ROS production in cells activates the response of ROS scavenging genes. However, this response cannot balance the abnormal increase of ROS which eventually leads to cell death.

2.6. PPR647 Is Required for C-to-U RNA Editing of Multiple Chloroplast Transcripts

Accumulating evidence shows that PLS-PPR proteins are required for RNA editing [19,20,27]. To explore the function of PPR647, we investigated the editing efficiency of 27 identified RNA editing sites in maize chloroplast transcripts. The results showed that the editing efficiency of most sites was changed in as-81647 and ems4-05741c (Figure 7, Supplementary Figure S3, Supplementary Table S2). Among them, the editing efficiency of rpoc2-926, ycf3-63, rps8-62, ndhB-197, ndhB-205, ndhB-278, ndhB-495 sites, which were 100% edited in WT and significantly reduced in as-81647 and ems4-05741c (Figure 7). In addition, all sites of rpoB were completely edited in WT and as-81647 plants, but their editing efficiency was zero in ems4-05741c (Figure 7). For this phenomenon, we speculate that the different mutation sites of as-81647 and ems4-05741c bring about inconsistent effects on rpoB editing. These data suggested that PPR647 is required for C-to-U RNA editing of multiple chloroplast transcripts.

2.7. PPR647 Affects the Splicing of rpl2 Transcripts in Chloroplasts

To determine the function of PPR647 in RNA splicing of chloroplast genes, we performed RT-PCR analysis using primers located in exons flanking intron and then compared the lengths of the amplified products between WT and mutant plants. Nearly no mature rpl2 transcripts were detected in as-81647 and ems4-05741c, and compared with WT, the splicing efficiency of atpF, ndhB, ndhA, and ycf3-2 significantly decreased in as-81647 and ems4-05741c (Figure 8). This result indicated that PPR647 is essential for RNA splicing of chloroplast genes (especially the rpl2 gene).

2.8. PPR647 Might Affect RNA Editing by Interacting with ZmMORF2

Recent studies have shown that MORF2 and MORF9 are located in plastid and are required for chloroplast RNA editing [28]. To further investigate the role of PPR647 in RNA editing, we examined the physical interaction of PPR647 with ZmMORF2 (encoded by Zm00001d026243), ZmMORF9 (encoded by Zm00001d024674) using the yeast two-hybrid assay. From the result, we found PPR647 was determined to interact with ZmMORF2 but not ZmMORF9 (Figure 9), which suggested that PPR647 might affect RNA editing by interacting with MORF2.
Figure 7. RNA editing analysis of chloroplast transcripts. The editing efficiency of multiple chloroplast transcripts was severely affected in as-81647 and ems4-05741c plants. The box marks the editing site. T (A) stands for edited, C (G) stands for not edited, and T/C stands part edited in WT.
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Figure 8. Splicing analysis of chloroplast transcripts in WT and mutants. From left to right, the template is DNA of WT, cDNA of WT, cDNA of as-81647, and cDNA of ems4-05741c. S, spliced; U, unspliced. The genes marked in red are genes affected.
Figure 8. Splicing analysis of chloroplast transcripts in WT and mutants. From left to right, the template is DNA of WT, cDNA of WT, cDNA of as-81647, and cDNA of ems4-05741c. S, spliced; U, unspliced. The genes marked in red are genes affected.

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Figure 9. The interaction between PPR647 and MORF2 protein. Yeast two−hybrid assays indicated the interactions between PPR647 and MORF2. Proteins were obtained on the selective medium SD−Trp−Leu and SD−Trp−Leu−His−Ade. AD, activating domain; BD, binding domain.

3. Discussion

There are hundreds of PPR proteins in maize, but only a few have been identified. Mutants of PPR proteins exhibit different phenotypes, such as leaf hypoplasia, growth restriction, photosynthetic pigment deficiency, and embryo or seed development defects [16,28–31]. In this study, a lethal albino seedling mutant (as-81647) was isolated and studied. In contrast to other reported albino mutants (al1 and al2) [32,33], whose Chl deletion phenotype were transient, as-81647 exhibits an albino-lethal phenotype throughout plant development. Corresponding to its phototype, we found that the chlorophyll content of as-81647 is almost zero and most cells in as-81647 had no chloroplast formation (Figure 2). Staining results showed that excessive reactive oxygen species accumulated in as-81647 leaf and caused cell death (Figure 2i–l). Map-based cloning and transgenic analysis confirmed PPR647, a novel PLS-type PPR protein, is responsible for the albino phenotype of as-81647. Subcellular localization demonstrated it is function in chloroplasts (Figure 3). Studies have shown that the biogenesis of Chl is tightly with chloroplast development and
its function [34]. So, those results indicated that PPR647 may play an important role in maize chloroplast development.

PEP is a multisubunit polymerase that drives the transcription of photosynthetic genes in chloroplasts, such as psbA, psbD, rbcL [2]. NEP is a single subunit polymerase that responsible for the accurate transcription of plastid housekeeping genes, such as rpoa, rpoB, rpoC1, and rpoC2 [2]. In our study, qRT-PCR and RNA-seq confirmed the transcription level of chloroplast gene mediated by PEP (PEPs) decreased significantly (Figure 5a) and genes mediated by NEP (NEPs) increased slightly (Figure 5b), suggested that PPR647 is involved in regulating the gene expression of PEPs, and the reduced PEPs transcripts may be the reason of arrested chloroplast development in as-81647. Similar results have been established in previous studies, such as other PPR proteins AtACM1 [35], AtPDM3 [36] and OsWSL [37] in which the transcription levels of NEPs elevated and transcription levels of PEPs were decreased. In acm1, the accumulation of chloroplast rRNAs and ribosome subunit RPS14 was disrupted, leading to the deletion of plastid ribosomes [35]. In pdm3, there is an increased steady-state level of rpoB transcripts but a reduced level of RpoB leading to a disrupted PEP complex [36]. In wsl, the low splicing efficiency of chloroplast transcript rpl2, leads to an aberrant transcript accumulation and the reduction of Rpl2 proteins, resulting in the lack of plastid ribosomes [37]. Chloroplast rpl2 encodes the L2 subunit of 50S ribosomal protein, an essential component of the chloroplast ribosome, and the absence of this protein is a very sensitive marker for the absence of ribosomal function [38]. Splicing efficiency experiment showed that PPR647 dysfunction resulted in ineffect splicing of rpl2 intron and reduced the splicing efficiency of atpF, ndhA, ndhB, and ycf3.2 (Figure 8). RNA-seq data showed that the expression of these genes decreased in varying degrees, indicating that their transcription was affected (Supplementary Figure S4). Therefore, we speculate that the lack of plastid ribosomes is due to the ineffective splicing of rpl2 and is one possible reason for the decrease of PEP-dependent transcript accumulation in as-81647. Multiple chloroplast genes (especially rpl2) require PPR647 to undergo splicing in maize.

The PEP complex consists of core subunits encoded by plastid gene (rpoa, rpoB, rpoC1, and rpoC2) and accessory protein encoded by nuclear gene (PAPs) [2]. Until now, many PAP mutants have been identified, such as wls3, wlp2 in rice, pap1, pap2, pap3, pap5, pap6, and pap10 in Arabidopsis, in which the transcription levels of NEP-dependent genes were also elevated and transcription levels of PEP-dependent genes were decreased [39–43]. At the same time, some proteins also caused this phenomenon due to their interaction with some proteins which directly interact with PEP complexes. For example, DG1 protein of Arabidopsis is involved in regulating PEPs transcription by interacting with SIG6 protein, which is necessary for PEP plastid gene transcription [44]. Therefore, another possibility is PPR647 may directly or indirectly bind to some components of PEP, that participate in the regulation of PEP transcription mechanism. Multiple chloroplast genes (especially rpl2) require PPR647 to undergo splicing in maize.

RNA editing, a post-transcriptional process, alters the RNA sequence by converting specific target cytidines in plastid transcripts to uridine. Our present study demonstrated that PPR647 affected the C to U editing efficiency of multiple plastid editing sites and interact with MORF2 (Figure 7, Supplementary Figure S3). Interestingly, we found that most of the affected sites were more influenced in ems4-05741, especially the rpoB (Figure 7, Supplementary Table S3). AS ems4-05741 was a strong allele compared with as-81647, we speculate that the site affecting rpoB splicing in PPR647 may not be affected in as-81647. Sequence comparisons and phylogenetic analyses identified PPR647 as the orthologue of PDM2 from Arabidopsis [26], which shows 46.3% sequence identity orthologue and 61.1% sequence similarity with all tested PEP-dependent transcripts, and higher expression of all NEP-dependent transcripts [26]. PDM2 is responsible for multiple RNA editing sites in plastid by interacting with MORF2 and MORF9. However, different from as-81647, embryogenesis
was also affected in pdm2, and the splicing of plastid genes in PDM2 was not affected, indicated that there may be some functional differences between PDM2 and PPR647.

In summary, the maize chloroplast localized PPR protein PPR647 affected the transcription of PEP-dependent plastid genes, and simultaneously plays roles in both the RNA editing and intron splicing in plastid genes. Although the functions of many PPR proteins have been widely investigated, few of them simultaneously play roles in both RNA editing and intron splicing (except for rice WLS4 protein) [22]. Studies have shown that there is a feedback mechanism between intron splicing and RNA editing in chloroplast genome, altered editing is most likely an indirect effect of defective splicing of chloroplast introns in as-81647. At present, there is no PPR protein in maize that simultaneously affects the splicing and editing efficiency of chloroplast genes have been reported. As a PLS-family PPR protein, studies on PPR647 will help further explore the function of the PPR protein family and lay a foundation for revealing the relationship between RNA editing and splicing in plastid.

4. Materials and Methods

4.1. Plant Materials and Growth Conditions

The albino (as-81647) mutant was isolated from maize inbred line 81647. ems4-05741c and ems4-057444 mutant lines were obtained from Maize EMS-induced Mutant Database (MEMD, http://www.elabcaas.cn/memd/; last accessed 8 September 2019) [33]. The preservation of as-81647 depends on heterozygous plants (as-81647/+), and as-81647/+ used as the male parent, was crossed with B73 and YQ165 to generate F1 and F2. Albino plants in the F2 population were used for genetic mapping. All materials used for mapping were grown at the Experimental Station of Shandong Agricultural University in China. For other purposes, the plants were grown in growth chambers with a photoperiod comprising 16 h light at 28 °C (day), 8 h dark at 22 °C (night), and 70%~80% relative humidity.

4.2. Photosynthetic Pigment Content Analysis

Approximately 100 mg fresh leaves of wild-type (WT) and as-81647 plants were cut into pieces and placed into 95% ethanol for 48 h in darkness at 4 °C. After centrifugation, the supernatant was measured at 663 nm, 645 nm, and 470 nm with a UV-2450 instrument (Hitachi, Tokyo, Japan). Three biological repeats were measured for each sample. The pigment contents were calculated according to the following equations described by Arnon [45].

\[
\text{Chl a (mg/g)} = \left[\frac{(12.7 \times \text{OD663} - 2.69 \times \text{OD645}) \times V}{W \times 1000}\right] \\
\text{Chl b (mg/g)} = \left[\frac{(22.9 \times \text{OD645} - 4.68 \times \text{OD663}) \times V}{W \times 1000}\right] \\
\text{Caro (mg/g)} = \left[\frac{\text{OD470} \times (V/W) - 3.27 \times \text{Chl a-104} \times \text{Chl b}}{198}\right]
\]

4.3. Transmission Electron Microscopy (TEM)

Leaves from as-81647 and WT plants were fixed with 2.5% glutaraldehyde in phosphate buffer (pH = 7.4) followed by osmium tetroxide and then dehydrated in an ethanol series before being infiltrated with Spurr’s resin. Polymerization was performed at 70 °C for 8 h. The specimens were sliced to yield ultrathin sections, stained with uranyl acetate and alkaline lead citrate before being examined with a JEM-1400Plus (JEOL, Tokyo, Japan) transmission electron microscope.

4.4. Histochemical Analysis

Trypan blue staining was used to detect cell death in leaves according to the methods of the previous study [46]. Fresh leaves were immersed in trypan blue solution (2.5 mg/mL trypan blue, 25% [w/v] lactic acid, 23% water-saturated phenol, 25% glycerol) at 70 °C for 10 min, then heated in boiling water for 2 min and incubated overnight at room temperature. Next, the sample was decolonized in a chloral hydrate solution (25 g in 10 mL of H2O)
for three days. They were stored in 70% glycerol and analyzed with a stereomicroscope (Olympus sxz12, Tokyo, Japan). For DAB staining, leaf segments (approximately 5 cm in length) were immersed in DAB solution (1 mg/mL, pH = 3.8) for 8 h to react with the H₂O₂. They were placed in 75% ethanol, heated for 15 min and transferred to 10% glycerol for microscopic examination (Olympus sxz12, Japan).

4.5. Map-Based Cloning and Allelism Test

Bulked segregant analysis (BSA) and simple sequence repeat (SSR) molecular markers were used for genetic mapping of the as-81647 locus. For preliminary mapping, 308 publicly SSR markers distributed over the whole genome from MaizeGDB were used for polymorphism screening. For fine mapping, new markers designed by SSRHunter and Primer 6.0 were used. The PCR procedure was as follows: 95 °C for 5 min, followed by 35 cycles of 94 °C for 30 s, annealing for 30 s, 72 °C for 30 s, and a final elongation step at 72 °C for 10 min. The primer sequences are listed in Supplementary Table S1.

For the allelism test, as-81647/+ was crossed with ems4-05741c/+ and ems4-057444/+.

4.6. Generation of CRISPR-Cas9-Edited Mutant Alleles

The CRISPR-Cas9 vector of Zm00001d004446 was constructed following a simplex editing strategy. The 20 bp target sequence (AAGCGGGAGGCAGCGAGCAT) for editing is located in the first exon of Zm00001d004446. Immature zygotic embryos (1.5–2.0 mm) of B104 were used for Agrobacterium-mediated maize transformation. T0 lines were hybridized to B104. For molecular identification of 15 F₁ transgenic plants, a marker was designed to identify the editing effect of Zm00001d004446. The BAR gene was used to identify the stability of editing (the primers used are shown in Supplementary Table S1). The PCR products of the target were then cloned into a pMD18-T vector for sequencing analysis.

4.7. RNA Sequencing (RNA-seq) and Data Analysis

Total RNAs were extracted from the two-leaf stage leaves with three biological replicates (10 individuals per pool). RNA-seq libraries construction and sequencing were both performed on the Illumina HiSeq4000 platform at Novogene Bioinformatics Technology Co. Ltd. (Beijing, China). RNA-seq data were deposited in the National Center for Biotechnology Information (NCBI) Sequence Read Archive (SRA) under accession number SRP334160 (BioProject ID: PRJNA757925). After sequencing, low-quality reads were filtered to obtain clean reads, and these clean reads were then mapped to the reference genome (http://www.maizegdb.org; last accessed 6 October 2021, Zea mays.AGPv4). Differential expression analysis of six samples was performed using the DESeq R package (version 4.1.1, U.S.A), and p-values were adjusted to control the false discovery rate. Unigenes with a revised p (q) value < 0.05 identified by DESeq R were considered to be differentially expressed. GO annotation and GO enrichment analysis (corrected p-value < 0.05) of DEGs were performed to investigate their functions further. GO enrichment analysis of DEGs was conducted using GOseq R packages (version 2.1.2, USA) based on Wallenius non-central hypergeometric distribution [47]. GO terms with corrected p (q) value < 0.05 were considered to be significantly enriched among the DEGs. Pathway-based analysis was conducted using KEGG [48].

4.8. RT-PCR and qRT-PCR Analysis

Total RNA was isolated from WT root, stem, leave, ear, developing seed at 20 DAP, and as-81647 leave using an EasySpin plus Plant RNA Kit (Aidlab, Beijing, China). A HiFiScript gDNA Removal cDNA Synthesis Kit (CwBiotech, Beijing, China) was used to generate high-quality first-strand cDNA. RT-PCR was used to examine RNA splicing efficiency and RNA editing using specific primers as described previously. qRT-PCR was performed with cDNA dilutions to detect the expression of the target gene and verify the accuracy of sequencing using SYBR Premix ExTaqTM Kit (TaKaRa, Shiga, Japan) on an prism
7500 Real-Time PCR (ABI, Vernon, CA, USA) in a 20 µL reaction volume with 40 cycles. The primers used for RT-PCR and qRT-PCR are listed in Supplementary Table S1. The maize ACTIN (Zm00001d010159) gene was used as an internal control in the experiment.

4.9. Analysis of RNA Editing and Splicing of Chloroplast Genes

Specific primers were used to generate RT-PCR products covering each editing site, and the products were sequenced directly. RNA editing efficiency was estimated by the relative height of nucleotide peaks in the analyzed sequence. Specific primers used for chloroplast editing sites were quoted from the Hammani study [23]. RT-PCR was performed using particular primers situated in exons flanking the intron of each gene. Primers used for chloroplast editing and splicing are listed in Supplementary Table S1.

4.10. Phylogenetic Analysis

Homologous sequences were identified in NCBI (http://www.ncbi.nlm.nih.gov/; last accessed 6 October 2021) by performing a BLASTP search with PPR647 protein sequences. Amino acid sequences were aligned with MUSCLE in the MEGA7.0 software package using the default parameters. Evolutionary distances were calculated using Poisson correction analysis. The bootstrap method with 1000 replicates for phylogeny testing was used.

4.11. Y2H Assays

Full-length cDNA sequences of PPR647, ZmMORF2 (Zm00001d026243) and ZmMORF9 (Zm00001d024674) were cloned into the pGBKKT7 and pGADT7 vectors (Promega, Madison, WI, USA). Constructs were subsequently co-transformed in pairs into yeast (strain AH109) following previously described methods [49].

4.12. Subcellular Localization of PPR647

The full-length CDS sequence of PPR647 without the stop codon was amplified by PCR from B73 and cloned into the transient expression vector pSAT6-EYFP-N1 to generate the fusion genes PPR647-YFP driven by the CaMv 35S promoter. The PPR647-YFP fusion product was introduced into Agrobacterium tumefaciens strain Gv3101 and introduced into maize protoplast as described. Fluorescence signals were detected using a Leica TCS SP5 II (Leica, Germany) laser scanning confocal microscope.

5. Conclusions

In this study, we identified a novel maize PLS-type PPR protein, PPR467, which is located within the chloroplast. Down-regulation of PPR467 impaired chloroplast development and PEP-dependent plastid genes expression and results in an albino-lethal phenotype. To summarize, PPR467 is functioning in chloroplast RNA editing, splicing, and is crucial for chloroplast development in maize.

Supplementary Materials: The following are available online at https://www.mdpi.com/article/10.3390/ijms222011162/s1. Figure S1: Alignment of amino acid sequences with the highest identity with the PPR647 protein. The Arabidopsis homolog is annotated as PDM2; Figure S2: The expression pattern of PPR647 from the publicly available Maize Gene Expression database (qTell). Different colors represent different processing conditions; Figure S3: RNA editing analysis of various target sites. The box is the editing site, T (A) stands for edited, C (G) stands for unedited, T/C stands part edited in WT; Figure S4: Relative expression of chloroplast genes which the intron splicing was affected in as-81647 according to the RNA-seq dates; Table S1: Markers used in this research; Table S2: The expression value of 11 selected genes in RNA-seq and real-time PCR; Table S3: Editing efficiency of 27 sites in WT, as-81647, and ems4-05741c; Excel S1: The DEGs data of as-81647 leaves vs WT leaves; Excel S2: KEGG enrichment of DEGs.

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