Organelle DNA degradation contributes to the efficient use of phosphate in seed plants

Tsuneaki Takami¹, Norikazu Ohnishi¹, Yuko Kurita²⁴, Shoko Iwamura², Miwa Ohnishi², Makoto Kusaba³, Tetsuro Mimura² & Wataru Sakamoto¹*

¹ Institute of Plant Science and Resources, Okayama University, Kurashiki, Japan
² Department of Biology, Graduate School of Science, Kobe University, Kobe, Japan
³ Graduate School of Science, Hiroshima University, Higashi-Hiroshima, Japan
⁴ Present address: Faculty of Agriculture, Ryukoku University, Otsu, Japan
* e-mail: saka@okayama-u.ac.jp

Corresponding author: Wataru Sakamoto
Mailing address: Institute of Plant Science and Resources
Okayama University
2-20-1 Chuo, Kurashiki, Okayama 710-0046
Japan
Telephone and Fax: +81-86-434-1206
Email: saka@okayama-u.ac.jp

One Sentence Summary: DNA retained in the endosymbiotic organelles – chloroplasts and mitochondria – of seed plants influences growth in phosphate-limited conditions through a degradation mechanism implemented by DPD1 exonuclease.
Mitochondria and chloroplasts (plastids) both harbor extra-nuclear DNA that originates from the ancestral endosymbiotic bacteria. These organelle DNAs (orgDNAs) encode limited genetic information but are highly abundant, with multiple copies in vegetative tissues such as mature leaves. Abundant orgDNA constitutes a significant pool of organic phosphate along with RNA in chloroplasts, which could potentially contribute to phosphate recycling when it is degraded and relocated. However, whether orgDNA is degraded nucleolytically in leaves remains unclear. In this study, we revealed the prevailing mechanism, in which organelle exonuclease DPD1 degrades abundant orgDNA during leaf senescence. The DPD1 degradation system is conserved in seed plants, and more remarkably we found that it was correlated with the efficient use of phosphate when plants were exposed to nutrient-deficient conditions. The loss of DPD1 compromised both the relocation of phosphorus to upper tissues and the response to phosphate starvation, resulting in reduced plant fitness. Our findings highlighted that DNA is also an internal phosphate-rich reservoir retained in organelles since their endosymbiotic origin.
Mitochondria and chloroplasts (plastids) originate respectively from the endosymbiosis of ancestral α-proteobacterium and cyanobacterium, ca. 1.5 billion years ago. Reflecting this endosymbiotic origin is the retention of their own DNA genomes and transcription/translation machineries. During the evolution of eukaryotic cells, however, most genes from these endosymbionts have been transferred to the nucleus, and only a small proportion of the ancestral genes remain within each organelle. In the model plant Arabidopsis thaliana, for example, only 87 proteins are synthesized in chloroplasts, whereas all other constituent proteins are encoded in the nuclear genome. Present eukaryotes, therefore, require the coordinated regulation between mitochondria/chloroplasts and the nucleus to fulfill organelle functionality.

In contrast to their limited genetic capacity, organelle genomes of relatively small size are known to be highly abundant, with multiple copies in each organelle. A striking example is leaf mesophyll cells, in which chloroplast DNA (cpDNA) accounts for ca. 30% of cellular total DNA, with an estimated >1,000 copies per cell. Typically, an A. thaliana mesophyll contains ~80 chloroplasts, resulting in >10 copies per chloroplast on average. Plastid DNA (ptDNA) copy numbers appear to vary in different species and in different plastid types, and they reach up to ~10,000 in developing leaves. As a consequence of the abundant DNA and protein synthesis, plastids contain a substantial amount of nucleic acids, which constitute a major pool of total cellular phosphorus (P) in leaves. Reportedly, chloroplast ribosomal RNAs account for the largest organic P pool, making up approximately half of the total nucleic acids pool. The multiple copies of ptDNA represent a considerable P pool. Excess ptDNA can be dispensable without affecting organelle functionality or cell viability, potentially providing a source of organic P for relocation when degraded. However, whether the amount of cpDNA/ptDNA is controlled by degradation in mature leaves has long been unclear. Little is known about the enzymatic degradation mechanism and its possible impact on the efficient use of the internal P pool in endosymbiotic organelles.

In reproductive organs, several nucleases targeted to the endosymbiotic organelles have been reported to digest DNA. In animals, mitochondrial EndoG nuclease expressed during male gametogenesis has been reported, which secures maternal inheritance of mtDNA. In a green alga, uniparental disappearance of orgDNA during mating occurs, although the nuclease responsible remains unclear. In flowering plants, we reported that DPD1 exonuclease degrades orgDNA in male gametophytes. However, DNA degradation by DPD1 per se does not contribute to maternal inheritance. Therefore, we postulate that DPD1 has functions other than the control of maternal inheritance. In this study, we demonstrated that in addition to its role in pollen, DPD1 degrades orgDNA in leaves undergoing senescence.
where nutrients are recycled through various macromolecule degradation mechanisms. DPD1 presents a determinate mechanism of orgDNA degradation conserved in plants. Moreover, this orgDNA degradation was shown to affect the efficient use of phosphate (Pi) positively when exposed to starvation conditions. We discuss a novel aspect of orgDNA, likely sensing Pi availability and acting as an internal reservoir of Pi, through degradation mediated by DPD1 in seed plants.

Results
Exonuclease activity of DPD1 confined to DNA but not RNA. We have shown that DPD1 is conserved in angiosperms but it was not detected in mosses or green algae, suggesting that it emerged during the evolution of flowering plants. Our search in the PLAZA database allowed us to isolate 43 DPD1 homologues from 35 plant species (Supplementary Fig. 1). Consistently, no DPD1 homologues were present in microorganisms or bryophytes, although its presence extended to gymnosperms (coniferous plants such as Pinus, Picea, Pseudotsuga and Gnetum) (Supplementary Fig. 1), supporting this specific emergence of DPD1 in seed plants (spermatophytes).

DPD1, which exhibits exonuclease activity and is targeted to both mitochondria and plastids, is unique in that most of the cell death-associated nucleases identified previously in plants are S1-type or Staphylococcal endonucleases. Because these endonucleases digest both RNA and DNA when single-stranded, we first tested if DPD1 has substrate specificity. Our in vitro nuclease assay, conducted using a purified DPD1 C-terminally fused to histidine tag and synthesized oligonucleotides as substrates (Fig. 1a), demonstrated that DPD1 degraded only DNA and not RNA, irrespective of whether it was single-stranded or double-stranded (Fig. 1b). This activity depended on Mg²⁺ (Fig. 1c) and was inhibited when a substrate 3’-end-labeled with a fluorescent dye (6-FAM) was used (Fig. 1b–d). We inferred that DPD1 is a 3’ to 5’ Mg²⁺-dependent deoxyribo-exonuclease, whose activity can be detected in physiological conditions equivalent to chloroplast stroma (Mg²⁺ concentration of >0.02 mM, temperature 22°C and pH 7.0–8.0, Supplementary Fig. 2). Given its heterogeneous forms, DPD1 alone can degrade at least a portion of orgDNAs processively, if they have a free 3’ end.

CpDNA degradation during leaf senescence. Our earlier survey of Arabidopsis transcriptome data predicted that DPD1 transcripts accumulate in senescing leaves as well as in pollen. To examine if DPD1 plays a role in vegetative tissues, detached Arabidopsis leaves were subjected to dark-induced senescence (see Methods), and orgDNA degradation
was monitored by quantitative PCR (qPCR). CpDNA levels in wild-type (ecotype Columbia [Col]) leaves declined apparently as senescence proceeded during incubation in the dark (Fig. 2a). When normalized using haploid nuclear DNA levels, we estimated the cpDNA copy number before the onset of senescence as approximately 400–600, which was similar to that reported previously (Fig. 2b) \(^{13}\). After 5 days in darkness, DNA levels declined substantially to less than 100 copies. Concomitant with cpDNA decline, our quantitative RT-PCR (qRT-PCR) analysis showed that \textit{DPD1} is upregulated (Fig. 2c), similarly or slightly earlier than senescence-related genes (Supplementary Fig. 3). Importantly, cpDNA levels did not decline in a \textit{dpd1} mutant and tended to stay constant (Fig. 2b). Retention of cpDNA in \textit{dpd1} was verified using digital PCR (Supplementary Fig. 4) and cytological observations of senescing leaves (Fig. 2d). Taken together, we concluded that DPD1 degrades cpDNA during leaf senescence.

Although the mechanism for maintaining ptDNA quantity remains unclear, a defect in DNA replication has been shown to affect ptDNA copy number adversely \(^{25}\). DNA polymerase in plant organelles is a bacterial-type pol I \(^{26}\). In \textit{Arabidopsis}, two isoforms have been reported, pol I-a and pol I-b, of which pol I-a plays the major role in ptDNA replication. Introduction of \textit{pol I-a2} into \textit{dpd1} appeared to decrease the copy number of cpDNA, whereas cpDNA stayed high during senescence (Fig. 2e). Therefore, our results revealed an epistatic effect of DPD1-mediated cpDNA degradation over DNA synthesis.

\textbf{MtDNA degradation during leaf senescence.} We next examined whether mitochondrial DNA (mtDNA) levels also declined in senescing leaves. The results showed a similar trend to cpDNA; mtDNA levels declined in Col as senescence proceeded, although they tended to stay constant in \textit{dpd1}. Therefore, we concluded that DPD1 also degrades mtDNA during leaf senescence. However, the estimated copy number was found to be very low, ranging from around a few copies per nuclear DNA even before dark incubation (Supplementary Fig. 5a). To address whether mtDNA levels decreased during leaf maturation, we examined 2-week-old seedlings grown in Murashige and Skoog (MS) plates to estimate mtDNA copy number. The result showed that approximately 20 copies of mtDNA were detected in young seedlings (Supplementary Fig. 5b). We observed a slight increase of the mtDNA copy number (approximately 25) in \textit{dpd1} compared with Col, consistent with previous reports. These results indicated that the mtDNA copy number declines before leaf maturation, which is independent of DPD1. Although the mechanism responsible for this mtDNA degradation remains unclear, the estimated copy number of mtDNA was consistent with previous reports describing that only a limited amount of mtDNA is detectable in mature leaves \(^{27,28}\). In contrast to this
shortage in mtDNA, plant mitochondria are known to undergo active fusion and fission.\(^8,29\). This dynamic behavior of mitochondria might account for the proposed sharing of genomic information between each mitochondrion. We concluded that orgDNA degradation proceeds in both organelles, but the majority occurs in chloroplasts.

**A weak stay-green phenotype in dpd1.** A careful examination of senescing leaves revealed that dpd1 displayed more greenness than Col and a dpd1 line complemented by DPD1 (G31) (Fig. 3a–b, Supplementary Fig. 6). This stay-green phenotype defines DPD1 as a factor accelerating senescence and cell death. Conversely, prolonged chloroplast functionality might be detectable in the dpd1 mutant. To address this, we first assessed the expression levels of chloroplast genes. qRT-PCR analysis revealed that the senescence-dependent decline in chloroplast-encoded transcripts was retarded in dpd1 (Fig. 3c and Supplementary Fig. 7). We inferred that the more abundant transcripts in dpd1 partly explained the stay-green phenotype, and that cpDNA degradation with DPD1 resulted in a concomitant reduction in the chloroplast RNA pool. Subsequently, we tested if the stay-green phenotype in dpd1 prolonged chloroplast functionality. Photosynthetic activity, as measured by the carbon dioxide assimilation rate in the same attached leaves grown for 2 weeks, appeared to be maintained for longer in dpd1 than Col (Fig. 3d, e). Together, these results confirmed that DPD1 accelerates senescence, although senescence still proceeds without DPD1.

**Growth defect of dpd1 in Pi starvation conditions.** The synergistic action of DPD1 on leaf senescence led us to postulate that cpDNA degradation is associated with nutrient availability. As a tradeoff between leaf longevity and nutrient deficiency, dpd1 prolongs photosynthesis, but it might impair growth in conditions with limited inorganic compounds. To test this possibility, we established hydroponic culture to grow Col and dpd1 (Supplementary Fig. 8) and observe the subsequent response to nitrogen or Pi deprivation (-N or -P, respectively) was investigated.\(^30\). First, based on our standard hydroponic conditions (1/4 MS medium), we found that Col grew better than dpd1 as estimated from the weight of aerial parts (Fig. 4a). This result was unexpected because no apparent difference was observed previously when they were grown in soil. Supplementing the hydroponic media with additional Pi rescued the defective growth of dpd1 (Fig. 4a), suggesting that dpd1 is specifically compromised in P availability. Additional Pi did not have significant effect in Col.

We next examined how Col and dpd1 respond to -N and -P in our standard hydroponic culture (see Methods). In principle, both starvation conditions attenuated plant growth by reducing the weight of aerial parts (Supplementary Fig. 9a, b). However, dpd1 appeared to
differ from Col in responding to -P. In -N conditions, both lines showed a pale color with slight anthocyanin accumulation, but no phenotypic difference was detectable. By contrast, -P conditions produced a profound growth defect in dpd1, which was characterized by reduced growth and substantial accumulation of anthocyanin (Fig. 4b). Such typical symptoms of -P conditions were not observed in Col. These results suggested that the efficient use of exogenous Pi was compromised in dpd1 by the lack of orgDNA degradation. When exposed to -N or -P conditions, qPCR demonstrated that cpDNA levels in Col leaves underwent degradation upon -N or -P similarly to dark-induced senescence, whereas degradation was inhibited in dpd1 (Fig. 4c, d).

Low fitness and P relocation of dpd1 in Pi starvation conditions. To ascertain the compromised response to -P, we measured seed production in Col and dpd1. First, seed numbers in plants grown in soil or hydroponic culture were counted. The hydroponic culture reduced seed production rate in Col, even in control conditions (1/4 MS), to approximately 80% of that grown in soil (Fig. 4e and Supplementary Fig. 9b). We also observed that the seed production rate was lower in dpd1 than Col. Remarkably, -P reduced seed production, even in Col, to approximately 50% of that in the control conditions, whereas dpd1 consistently showed a greater reduction in seed set than Col. Although our data indicated that dpd1 exhibited lower fitness in -P conditions than Col, it was possible that this resulted from the delayed senescence in dpd1, as evidenced by its weak stay-green phenotype. To compare this effect, we also measured seed set in -N conditions. The results showed no significant difference in seed set between Col and dpd1 (Fig. 4e). Therefore, nutrient starvation per se did not alter the sink capacity, instead dpd1 had a lowered fitness confined to -P conditions.

To examine whether the reduced fitness in dpd1 resulted from altered P remobilization, we measured the P content in leaves of Col and dpd1 in -P conditions. Assuming that degradation products of orgDNA contribute to seed set by relocating the catabolic products to reproductive organs, we expected to have lower P levels in the lower leaves of Col than in dpd1. ICP-MS measurement of the total P concentration indeed showed that Col leaves at 2 weeks after Pi deprivation relocated a significant greater amount of P from the lower leaves (leaves preexisting before P deprivation) to upper leaves (leaves that emerged after the start of -P treatment) (Fig. 4f). By contrast, no significant P relocation was detected in dpd1. The adverse effect of P redistribution between Col and dpd1 in -P conditions was, as expected, shown to correlate with our fitness results. We concluded that orgDNA degradation contributes to efficient P relocation, particularly when plants face P-limited conditions.
Global response to nutrient starvation in *dpd1*. We next investigated global changes in the transcriptome in -P conditions using RNA seq (Fig. 5, Supplementary Fig. 10 and Supplementary Fig. 11). RNA was isolated from Col and *dpd1* leaves either subjected to continuous growth in 1/4 MS or to Pi deprivation (*n* = 3, dataset is presented as Supplementary Table 1 and Supplementary Table 2). Comparison of the gene expression profiles revealed that the response to -P starvation was profoundly altered between Col and *dpd1* (Fig. 5a). The number of genes that were differentially expressed upon -P treatment was 766 in Col, of which 655 genes were upregulated. In contrast, only 114 were differentially expressed in *dpd1*; 96 genes were upregulated (Supplementary Fig. 10a). Col and *dpd1* shared 99 genes, among which 86 upregulated genes had GO terms related to phosphate starvation, photosynthesis, flavonoid biosynthesis and dephosphorylation.

To investigate these genes further, we specifically examined a set of genes that were categorized as related to Pi starvation response (PSR), mainly connected by a limited supply of inorganic Pi in the root environment (Supplementary Table 3) [3]. Of 193 genes specified as being involved in the PSR, we were able to extract 192 genes; among these, 123 and 40 genes were shown to be upregulated in Col and *dpd1*, respectively, at the significance level of false discovery rate [FDR] < 0.05 (Fig. 5c). We also specifically examined the gene set that was reported as being under the control of the PHR1 transcription factor (PHR regulon, Supplementary Table 4) [3]. Of 161 genes, 74 and 39 genes were upregulated in Col and *dpd1*, respectively (Supplementary Fig. 10b). Based on these results, we inferred that the global gene expression in response to -P conditions was compromised in *dpd1*. It was conceivable that orgDNA degradation and PSR are mutually interconnected, and that proper PSR requires orgDNA degradation.

We scrutinized the PSR genes differentially expressed between Col and *dpd1* (Supplementary Table 3 and Supplementary Fig. 10c). Upregulated genes in Col included those encoding Pi transporter (PHT1;9, PHT5, PHT3;2, PHT2, PHT1;4), purple acid phosphatase (PAP23, PAP7, PAP2, PAP22, PAP25, PAP17, PAP24, PAP14, PAP12), enzymes for lipid biosynthesis (MGD2, DGD2, MGDC, SQD1, SQD2) and RNase (RNS1), with which phosphate uptake or utilization is shown to be maximized in -P conditions. It was noteworthy that in *dpd1*, upregulation of the Pi transporter genes is limited to PHT1;9 and PHT5, although most of the purple acid phosphatase genes (*PAP22, PSP25, PAP2, PAP23, PAP12, PAP17, PAP24*) are also upregulated. In contrast to the genes preferentially upregulated in Col, we found several genes with expression levels that were higher in *dpd1* than Col (PPCK2, FHL and PLDZETA2). Overall, our RNA seq analysis revealed that the response to -P conditions was disturbed severely by the loss of orgDNA degradation.
Suppression of transporter genes in dpd1, but not the PAP gene, implied that the impact of orgDNA degradation in PSR is complex and is correlated with intra-cellular and inter-cellular Pi relocation.

To assess whether the altered transcriptome in dpd1 was rather specific to PSR, we also performed RNA seq with the plants exposed to -N conditions (n=3, dataset is presented as Supplementary Table 5 and Supplementary Table 6). Both Col and dpd1 presented many genes that were differentially regulated (1,768 for Col and 961 for dpd1, Supplementary Fig. 11a), suggesting that -N conditions generally impacted a broad range of genes (Fig. 5b). To investigate the -N response specifically, we focused on two sets of genes that have been reported previously to respond to -N conditions (Fig. 5d, Supplementary Fig. 11b, Supplementary Table 7, and Supplementary Table 8)\textsuperscript{32,34}. Comparison of these transcriptomes indicated that both Col and dpd1 displayed similar expression profiles based on the values at the median and upper/lower quartile, although dpd1 had a slightly reduced number of the genes than Col (Fig. 5d). These results were consistent with the growth defect and fitness observed in -N conditions (Supplementary Fig. 9a). Taken together, we concluded that dpd1 was compromised in PSR and that orgDNA degradation acts in the efficient use of Pi.

OrgDNA degradation mediated by DPD1 in natural conditions. To verify the role of orgDNA degradation, we questioned if it occurs in the natural environment. Seasonal remobilization of nutrients such as N and P from senescing leaves has been documented as being important in deciduous trees\textsuperscript{35}. In Populus alba, we have shown previously that about 60% of P in leaves was remobilized before the autumn leaf fall\textsuperscript{36,37}. CpDNA degradation has also been reported in a tree\textsuperscript{38}. Therefore, we considered that P. alba is suitable to test if DPD1-mediated orgDNA degradation coincides with P remobilization. We conducted leaf sampling from a field-grown P. alba tree (Supplementary Fig. 12), every month from the stage of bud break (April) up to complete leaf fall (November) (Fig. 6a). Estimation of cpDNA copy number by qPCR revealed that, in general, cpDNA was more abundant in spring and decreased gradually in autumn (Fig. 6b). Similarly to the case in Arabidopsis, mtDNA levels were much lower throughout the season (Fig. 6c). A small spike in orgDNA levels detected in the summer was likely due to leaf regeneration, which was consistent with P measurements in our previous study\textsuperscript{36}. Cytological observation of cpDNA was consistent with qPCR, showing holistic disappearance of cpDNA in autumn leaf samples (Fig. 6d).

qRT-PCR analysis of these samples, designed based on the available reference sequence from P. trichocarpa, demonstrated that a poplar DPD1 homologue was highly
upregulated toward leaf fall, with the highest level observed in November (Fig. 6e). We obtained these data from two consecutive seasons (2015 and 2016), which all showed upregulation of DPD1 that accompanied concomitant upregulation of senescence-related genes (Supplementary Fig. 14). We also confirmed DPD1 upregulation in laboratory conditions, which mimicked natural seasonal changes and leaf fall with three defined growing conditions (Fig. 6f); DPD1 was specifically upregulated at Stage 3, corresponding to autumn/winter (Fig. 6g and Supplementary Fig. 15). Therefore, all of these experiments confirmed the contribution of the DPD1 system during natural leaf fall, during which Pi is redistributed.

Discussion

CpDNA degradation in leaf tissues has been documented for more than two decades. However, whether DNA is degraded nucleolytically has been controversial, partly because of technical limitations of qPCR, variation within species and tissues, and a lack of mechanistic insights. Our studies with DPD1 uncovered the prevailing degradation mechanism among seed plants. We focused initially on male gametophytes (pollen), because the disappearance of orgDNA in male germ cells is often related to maternal inheritance, as evidenced in animal EndoG. Although we identified DPD1 exonuclease through forward-genetic mutant screening, orgDNA was shown to be degraded mainly in pollen vegetative cells, which deliver sperm cells to ovules but do not contribute to fertilization. In fact, we did not observe a contribution of DPD1 to the maternal inheritance mode of mtDNA, which led us to reconsider the physiological role of orgDNA degradation mediated by DPD1. Here, we demonstrated that both in an annual plant (Arabidopsis) and a deciduous tree (P. alba), the DPD1 system operates on orgDNA degradation in vegetative tissues, toward the final stage of leaf lifespan for Pi availability. These results revealed that the primary role of the DPD1 system is associated with the efficient use of Pi, rather than orgDNA inheritance.

During leaf senescence, relocating internal macronutrients to the upper and reproductive tissues plays a critical role in maximum fitness. Catabolism of macromolecules stored in chloroplasts is well described, including Rubisco, photosynthetic antenna protein, lipids and pigments, which act mainly in relocating N. Our finding adds orgDNA to the macromolecules undergoing degradation. It is noteworthy that a substantial portion of orgDNA resides in chloroplasts of fully expanded leaves, whereas only a limited number of mtDNAs exist. Although DPD1 is dual targeted to both organelles, its dominant role seems to be in chloroplasts/plastids. Conceivably, orgDNA degradation is beneficial in pollen vegetative cells because male gametophytes, once formed, are isolated
from other part of tissues, which hampers their ability to receive external P efficiently.

Conservation of the DPD1 system even in evergreen coniferous species (Supplementary Fig. 1) implicates that DPD1 has emerged during the evolution of microsporophytes, rather than the evolution of leaf senescence.

Several lines of evidence were presented to demonstrate the correlation of orgDNA with Pi starvation. First, dpd1 showed defective growth in our standard hydroponic culture, which was then rescued by supplementing with additional Pi (Fig. 4a). Second, dpd1 showed reduced fitness as well as typical symptoms in -P conditions (Fig. 4e). Third, these deficiencies in dpd1 were not detected in -N conditions but rather specific to -P conditions. Finally, RNA seq analysis in -P conditions indicated that dpd1 had compromised accumulation of PSR genes (Fig. 5). All these results suggested that orgDNA degradation participates in the efficient use of Pi. The most likely model to explain these results is that orgDNA itself acts as a P pool and is subjected to degradation for the redistribution of Pi (Supplementary Fig. 15, model 1). Consistent with this, upregulation of DPD1 during autumn leaf fall in P. alba coincided with the relocation of P from leaves (Fig. 6), which accounts for 60% of total P 36. Although whether orgDNA serves as a significant pool of internal P awaits further investigation, we inferred that the lower fitness in dpd1 could be explained by this reservoir model. The alternative model is that some orgDNA degradation product(s), such as nucleotides or their catabolic components, act as a positive sensor of PSR (Supplementary Fig. 15, model 2). Lack of these products may prevent plants in -P conditions from responding to P deficiency properly, which leads to lower fitness. Although we cannot exclude these two possibilities mutually, our data revealed an interconnection between orgDNA degradation and the efficient Pi use.

In leaves, nucleic acids constitute the most abundant Pi esters along with phospholipids 15,50. Breakdown of nucleic acids and/or enzymes for the biosynthesis of galacto- and sulpho-lipids to remodel phospholipids are induced as a part of PSR, along with purple acid phosphatases that hydrolyze Pi monoesters 51,52. We confirmed these PSR in our RNA seq data 33. Based on the results presented in this study, we considered that DNA degradation in endosymbiotic organelles also contributes to PSR. The nucleic acid P pool, representing 40–60% of the total internal organic P, consists of RNA and DNA, with ribosomal RNA as the largest pool 15,50. To degrade these large P pools, endonucleases are upregulated. RNS1 and RNS2 are the major ribonuclease that supposedly degrade cytosolic or extracellular RNAs 53. BFN1 has been reported to be upregulated during leaf senescence and to degrade single-stranded DNA/RNA 54,55. These findings imply RNA as a major P pool for relocation, whereas DNA has been considered as a minor P pool because of its indispensability. By
contrast, DPD1 is unique in that it is confined to plastids and mitochondria and degrades 'dispensable' orgDNA. Given that total DNA represents 20–30% of total nucleic acids in leaves, we estimated that orgDNA comprises 6–9% of total nucleic acid pool. Although minor, the dispensable orgDNA pool may serve as a safe guard of the Pi reservoir, consequently giving an advantage in -P conditions. In principle, extra internal Pi is considered to be stored in vacuoles. While transporting free Pi out of vacuoles plays a critical role in -P management, the contribution of chloroplasts remained elusive. During leaf senescence, dismantling of chloroplast compounds through enzymatic degradation and/or autophagic processes are recognized to be crucial. Our finding reinforces the importance of chloroplasts for relocating macronutrients, particularly for P.

Lack of orgDNA degradation in dpd1 caused a weak stay-green phenotype, but leaf senescence proceeded almost normally. Therefore, we considered that orgDNA degradation is not a decisive factor controlling the onset of leaf senescence. As a consequence of more cpDNA being retained (Fig. 3c), chloroplasts showed prolonged functionality because of the retarded decline in chloroplast transcripts. We inferred that orgDNA indirectly determines leaf lifespan, by balancing a tradeoff between prolonged photosynthesis and nutrient demand (Supplementary Fig. 15). In general, leaf senescence is associated with nutrient starvation, and an overlap between PSR and senescence-induced genes has been reported. Given the fact that orgDNA declines in response to both -N and -P but PSR is predominantly compromised in dpd1 (Fig. 5), we considered that the primary role of orgDNA degradation is likely to be to maximize P availability in leaves. One possibility of orgDNA contributing to leaf senescence could be a 'point of no return', which is proposed to define the stage that the senescence process cannot be reversed. It is known that senescence is reversible up to a certain point by providing additional nutrients. Conceivably, senescence is no longer reversible when cpDNA is completely lost.

DPD1 is homologous to DnaQ, an epsilon proofreading subunit of E. coli DNA polymerase III. Given that orgDNA replication adopts Pol I, it remains unclear how DPD1 emerged during evolution. In principle, DPD1 alone can degrade orgDNA given their heterogeneity, as advocated by Bendich: many orgDNAs are nicked, linearized and have a free 3' end. Whether algae or mosses have other types of exonucleases remains elusive, although the salvage function of orgDNAs was postulated earlier for Chlamydomonas. TREX1 is a mammal DPD1 homologue, which has been shown to be associated with inflammatory disease. Unlike TREX1, which degrades foreign pathogenic DNA, DPD1 has evolved to degrade endogenous DNA for salvage. In agriculture, the use of excess N and P fertilizers has drawn considerable attention, owing to the fact that over-fertilization of crop
fields disturbs the environment, and there is concern over whether the mining of P fertilizers will compromise their availability in the future. Our findings highlight orgDNA as a potential source of P storage, and future engineering for the efficient use of P in crop production.

Methods

Arabidopsis growth conditions and sampling

*Arabidopsis thaliana* ecotype Columbia (Col) was used as the control throughout this study. *dpd1* mutants (*dpd1-1* and *dpd1-5*) and a G31 transgenic line (*dpd1-1* complemented with the *DPD1* genomic sequence) were described previously. For growing plants, surface-sterilized seeds were placed on 0.8% (w/v) agar plates supplemented with MS medium (Sigma) and 1% (w/v) sucrose for 3 days at 4°C, followed by further growth in MS plates for 18 days at 23°C, at a photoperiod of 10 h light and 14 h darkness. Seedlings were then transplanted to soil and were grown for a further 4–5 weeks. Dark-induced leaf senescence was induced in these mature plants, from which we excised all leaves. We placed the leaves in darkness in a sealed chamber containing wet paper to maintain humidity.

Hydroponic culture

Hydroponic culture of *Arabidopsis* plants was performed as described by Conn et al. with a slight modification: the device used in the culture is shown in Supplementary Fig. 8 along with a detailed description in the legend. We used 1/4 MS medium as the hydroponic medium. Sterilized and cold-treated seeds were germinated on top of 1.5-mL microtubes immersed with 1/4 MS liquid medium (rack culture). After continuous growth for 1 month, whole plants with the microtubes were transferred and plugged into a new 15-mL tube filled with 1/4 MS media (tube culture). Growth was conducted in P- or N-depleted conditions by replacing the medium with medium lacking the corresponding elements 10 days after initiating tube culture. We prepared medium lacking potassium dihydrogen phosphate for -P, and lacking ammonium nitrate and potassium nitrate for -N. Hydroponic culture medium was exchanged with fresh medium every week. Phenotypes and responses to -P or -N deprivation were examined two weeks after plants were subjected to nutrient deprivation.

Poplar sampling

Periodic sampling of leaves from a white poplar tree (located at Uji Campus, Kyoto University, 34°91'N, 135°80'E, altitude 24 m above sea level, see Supplementary Fig. 12) was conducted during April–November in 2015 and 2016. Sampling was done every month between 13:30
and 14:30. Leaves in the area between 1.0 and 2.5 m from the ground were collected randomly. For each sampling, three sets were prepared, consisting of five leaves, which were subjected to RNA isolation followed by qRT-PCR. Meteorological data were acquired from the database of the Japan Meteorological Agency (http://www.jma.go.jp/jma/menu/menu/report.html).

For sampling of leaves from a shortened seasonal cycle system in growth chambers, poplar plants were cultivated initially from potted cuttings (shoots with five leaves at a height of 10 cm) with subsequent incubation at Stage 1 (1 month at 25°C in a 14 h/10 h light/dark cycle), Stage 2 (1 month at 15°C in an 8 h/16 h light/dark cycle) and Stage 3 (2–3 months at 5°C in an 8 h/16 h light/dark cycle). Stages respectively mimic spring/summer, autumn and winter in natural field conditions. Other growth conditions were similar to those reported previously (26). For sampling, fifth to seventh leaves from apical meristems on the respective plants were collected. Three sets were prepared and subjected to qRT-PCR.

**PCR analysis**

For studies of *Arabidopsis*, total DNA was isolated as described previously. For qPCR of organelle genes, primers were designed as listed in Supplementary Table 9. The reactions were performed using a kit (Thunderbird SYBR qPCR Mix; Toyobo Co. Ltd.) and Light Cycler 2.0 software (Roche Diagnostics Corp.) with 40 cycles of denaturation (95°C for 5 s) and extension (60°C for 30 s). Quantitative data were obtained from at least three biological replicates and were analyzed using LightCycler version 4.0 software (Roche Diagnostics Corp.). To normalize qPCR data from orgDNA over nuclear DNA, *DPD1* was used as a control of single-copy nuclear DNA, except that *18S rRNA* was used in Fig. 2a to follow Zoschke et al.13 (see corresponding figure legends). For studies on *Populus*, we conducted the same experiment as described above, but the primers were designed specifically based on the whole genome sequence of *Populus trichocarpa* (taxid: 3694, Phytozome 12, ver. 3.1) for nuclear genes, chloroplast genome sequence of *Populus alba* (taxid: 43335, Accession: AP008956) for chloroplast genes, and mitochondrial genome sequence of *Populus tremula* (taxid: 113636, Accession: KT337313) for mitochondrial genes. To minimize the amplification of mtDNA or ptDNA sequences included in the nuclear genome, we selected *rpoC1* for ptDNA, and *matR* and *cox3* for mtDNA as a reference. As a control nuclear gene, popular CAD gene (Potri.009G095800.1) was selected as a single-copy gene.

For qRT-PCR, total RNA was isolated using an RNeasy Plant Mini Kit (Qiagen), followed by reverse transcriptase and PCR reactions with a ReverTra Ace qPCR RT Kit (Toyobo Co. Ltd.) in accordance with the manufacturer's instructions. For *Arabidopsis*, we used Histone
variant gene H3.3 as an internal control, as described previously (16). For P. alba, we used ACTIN2 (Potri.001G309500.1) as an internal control, to measure the expression levels of DPD1 (Potri.005G020600.1), SAG12 (Potri.004G055900.1) and SGR1 (Potri.003G119600.1).

For digital PCR, a QuantStudio 3D Digital PCR System (Thermo Fisher Scientific Inc.) was used. DNAs were labeled using the Taqman probe method with primers designed accordingly (Supplementary Table 9). We used PsbA and DPD1 to measure the respective levels of cpDNA (FAM labeled) and nuclear DNA (VIC labeled), and adopted 40 cycles of denaturation (98°C for 30 s) and extension (60°C for 2 min) for the PCR reaction. Post-reaction chips were subjected to a QuantStudio 3D digital PCR system. DNA levels were quantified using AnalysisSuite Cloud Software.

All primers used for PCR analyses in this study are listed in Supplementary Table 9, with the accession numbers of the corresponding genes. All quantitative data included at least three biological replicates and are presented with SD in the graphs (statistical analysis is indicated in the corresponding figure legends).

Nuclease assay

The recombinant DPD1-His protein was purified as described previously 22 with a slight modification. Overexpression of proteins was conducted at 28°C. Ni²⁺-affinity purification was performed with Ni-NTA agarose (GE Healthcare). After purification, the imidazole-containing buffer was exchanged for 2× DPD1 storage buffer (100 mM Tris-HCl [pH 7.5], 200 mM NaCl) using a gel filtration column midiTrap G-25 (GE Healthcare). The obtained fractions containing the desired protein were subjected to centrifugation with AmiconUltra-4 (10K) (Millipore Corp.) to concentrate the recombinant protein to >2.0 µg µL⁻¹. The protein solution was diluted to adjust the concentration to 2.0 µg µL⁻¹, and was subsequently mixed with an equal amount of glycerol to make a 1.0 µg µL⁻¹ stock solution. Stock solution aliquots were stored at -30°C until use. Either aliquots of soluble proteins extracted from E. coli cells or purified recombinant proteins were solubilized by incubation at 75°C for 5 min in the presence of 2% SDS and 0.1 M DTT. The protein samples were centrifuged for 1 min at >20,000 × g and were then subjected to SDS-PAGE with 12.5% (w/v) polyacrylamide gels. The proteins in the gel were subsequently visualized by staining (CBB Stain ONE; Nacalai Tesque Inc.).

For the in vitro nuclease assay, we used 6-FAM-labeled oligonucleotides purchased from Hokkaido System Science as substrates. Oligonucleotides of all types (dsDNA, ssDNA and ssRNA) were designed based on the sequence (5’-CGAACACATACTTCACAAGC-3’) derived from one primer used earlier for amplifying a ptDNA fragment (ndhI gene). The nuclease assay was performed in a 12.5 µL reaction mixture that consisted of 40 mM Tris-HCl
(pH 7.5), 2 mM MgCl₂, 1.6 μM oligonucleotides and 17.5–175 ng of purified DPD1-His protein. Each reaction was terminated by the immediate addition of stopping buffer (1% [w/v] SDS, 50% [v/v] glycerol, 0.05% [w/v] bromophenol blue). After each reaction, the digestion products were separated electrophoretically on 20% (w/v, acrylamide: bis = 29:1) polyacrylamide gels. For double-stranded DNA, reaction mixtures with no treatment were loaded on a polyacrylamide gel. Reaction mixtures containing single-stranded DNA or RNA were supplemented with an equal amount of denaturing buffer (TBE buffer containing 10 M urea, 20% [v/v] glycerol and 0.1% [w/v] bromophenol blue) and were then heated at 65°C for 5 min. Subsequently, the samples were subjected to denaturing polyacrylamide gel electrophoresis in the presence of 7 M urea. The separated fragments were detected using an image analyzer (LAS4000; Fuji).

Cytological observation

For observing DNA with 4,6-diamido-2-phenylindole (DAPI), leaves were simultaneously fixed and stained with 1 μg/mL DAPI (3% [w/v] glutaraldehyde). The leaves were observed directly using a microscope (BX61; Olympus Optical Co. Ltd.) equipped with a disc scan unit. When necessary, sections were prepared using a vibratome VT 1200S (Leica Biosystems) with samples embedded in either 4% (w/v) gelatin or 1.5% (w/v) agarose, setting blade speed 0.4 mm/s, blade vibration 1.5 mm, thickness 70–100 μm and blade angle of 12–15°.

Phylogenetic analysis

Protein sequences homologous to DPD1 were obtained from the PLAZA database (https://bioinformatics.psb.ugent.be/plaza/). Multiple alignment of the extracted homologues was performed using MUSCLE software with the MEGA7 database. An unrooted tree was constructed using the Maximum Likelihood method based on the JTT matrix-based model with the default settings in MEGA7.

Photosynthetic activity measurement

Photosynthetic activity of Col and dpd1 leaves of plants grown in soil was measured as the transpiration rate (LI-6400XT; Li-Cor Inc.). The same leaves were subjected to measurement to estimate the decline in photosynthetic activity at 1 and 2 weeks after the initial measurements. CO₂-dependent photosynthesis curves were obtained at a light intensity of 1,000 μmol m⁻² s⁻¹. For each measurement, the relative moisture of the chamber was adjusted to 60–70%.
**Arabidopsis RNA seq analysis**

For RNA seq in Arabidopsis, total RNA was isolated from leaves either in P depletion or control conditions as described above. RNA sequencing was conducted using a HiSeq 2500 or 4000 Illumina sequencing platform and outsourced (Macrogen Corp. Japan), including DNA library preparation using a TruSeq RNA sample Prep Kit v2 and sequencing reaction with a TruSeq rapid SBS kit, Truseq SBS Kit v4, or TruSeq 3000 4000 SBS Kit v3. Sequences were obtained as pair-end reads. At least four billion reads were obtained for each sample (n=3). Mapping of the obtained sequences was performed using the Quas/R package. The gene expression levels were detected by edge/R after normalization with the TCC package. Volcano plots were constructed using the ggplot2/R package with the dataset of all differentially expressed genes (Supplementary Table 1 and Supplementary Table 2 for -P, and Supplementary Table 5 and Supplementary Table 6 for -N). Box plots were constructed using boxplot and beeswarm/R packages with the dataset of the selected genes (Supplementary Table 3 and Supplementary Table 4 for -P, and Supplementary Table 7 and Supplementary Table 8 for -N), which was reported earlier as P-responding or as N-responding, respectively.

**Measurement of total phosphorus contents**

Plants grown in hydroponic culture, with 1/4 MS or in -P conditions, were subjected to P measurement. Before P deprivation, all leaves were marked as lower leaves, whereas newly emerged leaves in -P condition (2 weeks) were designated as upper leaves. Samples (n=6) were dried in an oven at 60°C for at least 1 day. Dried samples were then digested with 60% (w/v) nitric acid at temperatures as high as 180°C. The concentration of P in the digested solution was ascertained using ICP-mass spectrometry (7500CX; Agilent Technologies Inc.).

**Reporting summary**

Further information on experimental design is available in the Nature Research Reporting Summary linked to this article.

**Data availability**

Accession numbers of the genes used in this study are listed in Supplementary Table 9. Precise p values calculated by statistical tests in this study are listed in Supplementary Table 10. The raw data used to construct graphs in this study are presented as Supplementary Dataset. The raw transcriptomic data are deposited in the DDBJ with the accession number.
DRA007138, under the BioProject with the accession number PRJDB7233. All transcriptomic data used in Fig. 5, Supplementary Figs. 10 and 11 are available in Supplementary tables 1-8.
References


Corriveau, J. L. & Coleman, A. W. Rapid screening method to detect potential


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AUTHOR INFORMATION

Affiliations

Institute of Plant Science and Resources, Okayama University, Kurashiki, Japan
Tsuneaki Takami, Norikazu Ohnishi & Wataru Sakamoto

Department of Biology, Graduate School of Science, Kobe University, Kobe, Japan
Yuko Kurita, Shoko Iwamura, Miwa Ohnishi & Tetsuro Mimura

Graduate School of Science, Hiroshima University, Higashi-Hiroshima, Japan
Makoto Kusaba

Contributions

W.S. designed the project. T.T. performed all qPCR and qRT-PCR measurements for various environments, in addition to photosynthetic activity measurements and RNA seq analysis. N.O. performed nuclease assays. Y.K., S.I., M.O. and T.M. prepared poplar samples and conducted primary work related to poplar. T.T., M.K. and W.S. analyzed the data. W.S. wrote the manuscript with consultation among all coauthors.

Competing interests

The authors declare no competing interests.

SUPPLEMENTARY INFORMATION

Supplementary Figures 1–15
Supplementary Tables 1–8
Supplementary Table 9
Supplementary Table 10
Supplementary Dataset
Reporting Summary
Figure legends

**Fig. 1 | Exonuclease activity of DPD1.** a, Schematic representation of the DPD1 construct used for this study (top), and Coomassie-stained SDS-PAGE gel showing the induction of DPD1 fusion proteins (left) and fusion proteins purified using an Ni-NTA agarose column (right). The control sample without induction (C) and IPTG-induced samples (I) are indicated. M, molecular weight markers. b, In vitro nuclease assay of DPD1-His using 20-mer double-stranded DNA (dsDNA), single-stranded DNA (ssDNA) or ssRNA as substrate. Arrowheads indicate the positions of the substrates. c, Non-denaturing 20% polyacrylamide gel electrophoresis demonstrating Mg$^{2+}$ dependence of the nuclease activity. 20-mer ssDNA was used as the substrate (indicated by red arrows). d, Denaturing 20% polyacrylamide gel electrophoresis demonstrating 3′-to-5′ polarity of the nuclease activity. Either 5′- or 3′-end-labeled 20-mer ssDNA was used as a substrate. All nuclease assays were repeated twice with three independent sample preparations.

**Fig. 2 | DPD1 is induced by leaf senescence and degrades orgDNA in vivo.** a, Decline in cpDNA levels during dark-induced leaf senescence in Col, as estimated by qPCR (psbA was used for cpDNA and 18S rRNA for nuclear DNA). Representative images of senescing leaves on days 0–5 are presented at the top. b, Retention of cpDNA in dpd1 estimated by qPCR (open and closed circles represent dpd1 and Col, respectively). Chloroplast genes used for qPCR are presented in each graph. The copy number of cpDNA was estimated by normalization with DPD1. c, Upregulation of DPD1 transcripts in Col senescing leaves, as estimated by qRT-PCR. d, Cytological observation of chloroplasts (chlorophyll autofluorescence, Chl) and cpDNAs (stained with DAPI) in senescing leaves of Col and dpd1 (after 5 days in darkness). e, Estimation of cpDNA copy number in senescing leaves (after 5 days in darkness) of Col, dpd1, polla2 and dpd1/polla2 by qPCR (psbA/DPD1, n=3, Student’s t-test, two-sided, *P<0.05, **P<0.01, p values shown in Supplementary Table 10). All quantitative data in Fig. 2 were from three biological replicates (Supplementary Dataset) and are shown as mean values with SD error bars.

**Fig. 3 | Stay-green phenotype and prolonged leaf longevity in dpd1.** a, Detached leaves subjected to dark-induced senescence from Col, dpd1-1 and G31 (transgenic dpd1-1 complemented by DPD1). All detached leaves from the respective plants are aligned from left (younger) to right (older) before dark induction (Day 0) and after 5 days in darkness (Day 5). Representative images from three independent experiments are shown. A similar stay-green
phenotype was observed in *dpd1-5* (Supplementary Fig. S6). b, Chlorophyll contents of senescing leaves (ninth-oldest leaves among all leaves subjected to dark induction) from Col (closed) and *dpd1-1* (open) (mean value ± SD, n=4, Dunnett’s test, against day 0, two-sided, ***P<0.001, p values shown in Supplementary Table 10). c, Retarded decline in transcripts encoded in chloroplasts (*psbA, clpP*) and those encoding chloroplast-targeted proteins (*psbO, Sig2*), estimated by qRT-PCR (mean value with SD, n=3, Dunnett's test, two-sided, *P<0.05, **P<0.01, ***P<0.01, p values shown in Supplementary Table 10). Other transcripts are also indicated in Supplementary Fig. S7. d, CO₂-dependent photosynthetic activity of mature leaves from Col and *dpd1-1*, grown in normal conditions with a light intensity of 1,000 μmol m⁻² s⁻¹ (mean value ± SD, n=6, Student's-t test, two-sided). e, Photosynthetic activity as in d, but at a fixed CO₂ concentration of 1,100 ppm (mean value ± SD, n=6, Dunnett’s test, two-sided, against initial measurements). Initial measurements were conducted in mature leaves from Col and *dpd1-1* (black, left bar) with subsequent measurements taken of the same leaves after 1 week (gray, middle bar) and 2 weeks (light gray, right bar). Raw data for all quantitative analyses are shown in Supplementary Dataset.

**Fig. 4** | Hydroponic culture of *dpd1* exhibited attenuated P response and reduced fitness in phosphate-deprivation conditions. a, Growth rate of Col (red bars) and *dpd1-1* (blue bars) estimated by the weight of aerial part in standard 1/4 MS conditions, either with standard (1/4) or additional (1/2 and 1) P concentration for 2 weeks (4 biological replicates). Weights are presented as the mean values ±SD (Col 1/4P: 423 ± 78 (n=12); *dpd1-1* 1/4P: 259 ± 107 (n=15); Col 1/2P: 401 ± 89 (n=12); *dpd1-1* 1/2P: 350 ± 103 (n=18); Col 1P: 344 ± 126 (n=13); *dpd1-1* 1P: 388 ± 72 (n=19), FW: fresh weight. **P<0.01 calculated using Dunnett’s test, two-sided, p value shown in Supplementary Table 10). b, Leaves from Col, *dpd1* and G31 exposed to -P conditions for 2 weeks (upper panels). Representative images from three independent experiments are shown. Typical symptoms showing purple pigmentation in *dpd1* are indicated by arrowheads. Lower panels depict representative images of a *dpd1* leaf showing anthocyanin accumulation and a young silique showing aborted seed development (arrowheads). c and d, Estimation of cpDNA copy number by qPCR, in leaves from Col and *dpd1-1* subjected to -P (c) or -N (d) for 2 weeks. Leaf number (1, 2 and 3) denotes the younger (upper) leaf as Leaf 1, in a plant grown hydroponically for 2 months (mean value ±SD, 3 biological replicates, n=3, Dunnett’s test, two-sided, against leaf 1, *P<0.05, p value shown in Supplementary Table 10). Examples of leaves used for cpDNA measurement are shown in Supplementary Fig. S8d. e Fitness of Col and *dpd1* plants estimated by seed set. Plants grown in normal conditions in soil, in standard hydroponic
culture (1/4 MS), -P in hydroponic culture (-P) and -N in hydroponic culture (-N) are compared by the number of seeds set per silique (n=50, Games-Howell’s test, two-sided, *P<0.05, ***P<0.01, p values shown in Supplementary Table 10). Lower whisker, bottom of box, center line of box, top of box and upper whisker shows minimum, lower quartile, median, upper quartile and maximum, respectively. f, Remobilization of P from lower to upper leaves (schematically illustrated at the top), grown in either control (1/4 MS, bottom) or -P (top) hydroponic conditions for 2 weeks (see Methods). P remobilization was estimated by the ratio of P concentration in upper leaves over that in lower leaves (maen value ±SD, 3 biological replicates, n=6, Student’s-t test, two-sided, **P<0.01, p value shown in Supplementary Table 10). Raw data for all quantitative analyses are shown in Supplementary Dataset.

**Fig. 5** | RNA seq analysis showing compromised response of dpd1 to -P. a, Volcano plots showing the genes significantly upregulated in -P conditions (2 weeks) in Col (top) and dpd1-1 (bottom). Data are obtained from three independent samples. Each dot in the graphs represents a single gene, and those significantly upregulated (FDR <0.05, calculated by Benjamini-Hochberg procedure included in edgeR package) are highlighted in red. b, Volcano plots as in a, except that the data are from in -N conditions. c, Box plot of PSR genes extracted from RNA seq data. 192 PSR genes were detected in our RNA sequence data. Differential expression of these genes (Log2 fold change) after -P treatment is shown. Each dot represents a single gene. Those showing significant alteration (FDR <0.05) are highlighted in red. Lower whisker, bottom of box, center line of box, top of box and upper whisker shows minimum, lower quartile, median, upper quartile and maximum, respectively. d, Box plot as in c, except that the data are from genes extracted as responding to -N, in accordance with Krapp et al.32.

**Fig. 6** | CpDNA decline and upregulation of DPD1 during leaf fall in a deciduous tree *Populus alba*. a, Example of leaf samples from a *P. alba* tree used in this study. Sampling dates are indicated above each panel (in 2014). b, Decline in cpDNA copy number estimated by qPCR (*rpoC1* as a reference gene for cpDNA and *CAD* for nuclear DNA; see Methods). Mean values with error bars as SE (n=3 for Oct, n=5 for other samples) are shown. c, Decline in mtDNA copy number in leaves estimated by qPCR, as in b. As a reference gene, *matR* (closed circle) and *cox3* (open circle) are used. d, Cytological observation of chloroplasts (chlorophyll autofluorescence, Chl) and cpDNA (DAPI stained) in *P. alba* leaf samples collected in April (top) and October (bottom). Merged images are shown on the left. Results...
presented in b, c and d are from samples prepared in 2015. Representative images from three independent leaf samples are shown. e, Expression of poplar DPD1 analyzed by qRT-PCR in 2015 (left) and 2016 (right). Mean values with error bars as SE (n=3 for Oct 16th, n=5 for other samples) are shown. Expression of other genes and meteorological data for each year (irradiance and average temperature) are shown in Supplementary Fig. S14. f, Outline of P. alba plants grown in a shortened annual-cycle cultivation system using controlled-condition growth chambers. Photographs of representative plants (n=3 independent samples) used in our study are shown. g, Expression of DPD1 and SGR, as in e. Mean values with error bars as SE (n=3 independent samples) are shown. Raw data for all quantitative analyses are shown in Supplementary Dataset.
**a**

DNA sequence comparison showing T7 promoter and 6x His tag.

**b**

Time-course experiment with varying concentrations of DPD1 (ng): 175, 70, 35, 17.5 and +EDTA. Results show dsDNA degradation over time.

**c**

Effect of MgCl₂ concentration on DPD1 activity. +MgCl₂ (mM): 2, 0.2, 0.02, 0.002. Time (min): 0, 15, 30.

**d**

DPD1 activity on 5' and 3' labeled ssDNA. Time (min): 0, 15, 30.