Proanthocyanidin synthesis in *Theobroma cacao*: genes encoding anthocyanidin synthase, anthocyanidin reductase, and leucoanthocyanidin reductase

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Abstract

**Background:** The proanthocyanidins (PAs), a subgroup of flavonoids, accumulate to levels of approximately 10% total dry weight of cacao seeds. PAs have been associated with human health benefits and also play important roles in pest and disease defense throughout the plant.

**Results:** To dissect the genetic basis of PA biosynthetic pathway in cacao (*Theobroma cacao*), we have isolated three genes encoding key PA synthesis enzymes, anthocyanidin synthase (ANS), anthocyanidin reductase (ANR) and leucoanthocyanidin reductase (LAR). We measured the expression levels of *TcANR*, *TcANS* and *TcLAR* and PA content in cacao leaves, flowers, pod exocarp and seeds. In all tissues examined, all three genes were abundantly expressed and well correlated with PA accumulation levels, suggesting their active roles in PA synthesis. Overexpression of *TcANR* in an Arabidopsis *ban* mutant complemented the PA deficient phenotype in seeds and resulted in reduced anthocyanidin levels in hypocotyls. Overexpression of *TcANS* in tobacco resulted in increased content of both anthocyanins and PAs in flower petals. Overexpression of *TcANS* in an Arabidopsis *ldox* mutant complemented its PA deficient phenotype in seeds. Recombinant *TcLAR* protein converted leucoanthocyanidin to catechin *in vitro*. Transgenic tobacco overexpressing *TcLAR* had decreased amounts of anthocyanidins and increased PAs. Overexpressing *TcLAR* in Arabidopsis *ldox* mutant also resulted in elevated synthesis of not only catechin but also epicatechin.

**Conclusion:** Our results confirm the *in vivo* function of cacao ANS and ANR predicted based on sequence homology to previously characterized enzymes from other species. In addition, our results provide a clear functional analysis of a LAR gene *in vivo*.

Background

Flavonoids are a diverse group of plant secondary metabolites with various biological functions that play important roles during plant development. They are involved in plant defenses against insects, pathogens and microbes, in absorption of free radicals and UV light, and in attraction of beneficial symbionts and pollinators [1-4]. Proanthocyanidins (PAs, also known as condensed tannins) are components of metabolites synthesized through the general flavonoid pathway. Their main known function is to provide protection against microbial pathogens and invasions of insects and herbivores through multiple mechanisms [1]. PAs have metal chelating activity that results in severe limitation of bacterial growth; PAs can also associate with and irreversibly precipitate proteins, which is responsible for the astringent taste that repels herbivores. Furthermore, PAs can be oxidized to quinones which not only are powerful antibiotics themselves, but also can initiate cross-linking of cell walls to increase the strength of this physical barrier to pathogens [5-8]. In addition to these functional roles in plants, PAs, especially those from cacao, have recently been suggested to be beneficial to humans by improving cardiovascular health through activation of nitric oxide.

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synthesize, by providing cancer chemopreventative effects, and also through neuroprotective activities [9-11].

The biosynthetic pathways that lead to the production of flavan-3-ols ((+)-catechin and (−)-epicatechin), the building blocks of PAs, have been well studied in the model plant species maize (Zea mays) and Arabidopsis as summarized in Figure 1 [12,13]. Biosynthesis of flavan-3-ols involves three principal enzymes: leucoanthocyanidin reductase (LAR), anthocyanidin synthase (ANS; also called leucoanthocyanidin dioxygenase, LDOX), and anthocyanidin reductase (ANR; in Arabidopsis, the product of BANYULS gene). The synthesis of PAs and anthocyanins share common steps leading to flavan-3,4-diols (such as leucoanthocyanidin), which can be converted to catechin (2,3-trans-flavan-3-ol) by LAR [14] or to anthocyanidin by ANS [15,16]. Anthocyanidin then either serves as the substrate for the synthesis of epicatechin (2,3-cis-flavan-3-ol) by ANR [17], or can otherwise be converted to anthocyanin by glycosylation [18]. Both catechin and epicatechin act as the initiators for PA polymerization, with intermediates derived from leucoanthocyanidin, catechin or epicatechin added sequentially as extension units [1]. However, the details of the polymerization process are unclear and it is not known whether this is a spontaneous or an enzyme-catalyzed reaction. Recent work has identified two new enzymes downstream of flavan-3-ols that are involved in key steps of PA polymer biosynthesis, an epicatechin 3′-O-glucosyltransferase in Medicago truncatula [19] and MATE transporters from Arabidopsis and Medicago truncatula [20] that transport epicatechin 3′-O-glucoside to the vacuole in which it is likely that PA polymerization occurs [21].

ANS and ANR genes have been biochemically and genetically characterized in Arabidopsis [15,17,22,23], Medicago truncatula [17,24] and Vitis vinifera (grape) [25]. The Arabidopsis ans (ldox) mutant exhibits a deficiency in both anthocyanin accumulation in hypocotyls and PA deposition in seeds that results in a transparent testa phenotype [15]. Seeds of the Arabidopsis banyuls (anr) mutant exhibit a lack of PAs and a hyper-accumulation of anthocyanins, resulting in a dark red color reminiscent of the famous Banyuls wine produced in southern France [22]. Over-expression of ANR genes from Medicago and grape in tobacco results in a loss of

![Figure 1 Outline of the proanthocyanidin synthesis pathway (adapted from [17]). Enzymes are represented in uppercase letters. DFR, dihydroflavonol 4-reductase, EC 1.1.1.219; ANS, anthocyanidin synthase, EC 1.14.11.19; ANR, anthocyanidin reductase, EC 1.3.1.77; LAR, leucoanthocyanidin reductase, EC 1.17.1.13; OMT, O-methyltransferases, EC 2.1.1.-; UFGT, UDP-glucose: anthocyanidin/flavonol 3-O-glucosyltransferase, EC 2.4.1.115; RT, rhamnosyltransferase, EC 2.4.1.159.](image-url)
anthocyanin pigments in flower petals and elevated levels of PAs [17,25]. Antisense down-regulation of ANS in *Medicago* results in reduction of both anthocyanins in leaves and PAs in seeds [24].

*LAR* genes have been isolated from various plant species including *Desmodium uncinatum* [14], *Vitis vinifera* [25], *Lotus corniculatus* [26] and *Medicago truncatula* [24] and their corresponding protein functions have been characterized by *in vitro* recombinant enzyme assays. However, the genetic evidence for *LAR* function is rather indirect and less convincing than it is for ANR and ANS, as discussed by Pang et al. [24]. It appears that the genomic sequence of *Arabidopsis thaliana* does not contain an intact *LAR* orthologue, and correspondingly, catechin is not detected in *Arabidopsis* seed extracts [12,14,15]. *LAR* genes are expressed in other plant species that accumulate not only epicatechin but also catechin [14,25,26]. For example, grape and *Lotus* express both *LAR* and *ANR* genes and synthesize PAs consisting of both catechin and epicatechin [25-27]. In *Medicago*, although both *LAR* and *ANR* are expressed, the PAs are composed almost entirely of epicatechin [24].

In *Arabidopsis* and *Medicago*, PA accumulation and gene expression is quantitatively and spatially limited to seed coats, making it remarkably difficult for biochemical analysis [12,24]. In contrast, *Theobroma cacao* (*Ts*) produces significant amounts of PAs in various tissues including leaves and beans; up to 12% of dry weight in leaves [28] and approximately 10% in mature beans [29]. Furthermore, large amounts of catechin and epicatechin monomers as well as their related polymers of different lengths have been detected in cocoa powder [28,30]. Considering the wide range of health benefits suggested for PAs and its significance for plant resistance, we targeted this pathway for molecular-genetic analysis.

This manuscript describes the isolation and expression of the *TcANR*, *TcANS*, and *TcLAR* genes encoding the key enzymes in proanthocyanidin biosynthesis. We measured PA content in different cacao tissues and performed functional characterization of the *TcANR*, *TcANS*, and *TcLAR* gene products through *in vivo* tests. The results presented here provide background and genetic tools that will be useful in the development of new cacao varieties with altered PA profiles.

**Results**

**Molecular cloning and sequence analysis of *TcANR*, *TcANS* and *TcLAR* genes**

The general pathway for the synthesis of proanthocyanidin and anthocyanins indicates the key enzymes ANR, ANS and LAR that carry out the biochemical steps at a critical metabolic branch point (Figure 1). To explore the genetic control of this important pathway in cacao, putative *TcANR*, *TcANS* and *TcLAR* cDNA sequences were identified in a collection of *Theobroma cacao* expressed sequence tags (ESTs) [31] by querying the cacao ESTtik database (http://esttik.cirad.fr/) with protein sequences of *Arabidopsis BANYULS* (NP_176365), *Arabidopsis LDOX* (Q96323) and *Desmodium LAR* (CAD79341). ESTs similar to each gene were assembled into contigs to determine consensus full-length open reading frames (ORF) by alignment with cDNAs of homologous genes from other species and predictions from the ORF Finder program (www.ncbi.nlm.nih.gov/projects/gorf/). Full-length cDNAs of each gene were amplified by reverse transcription-PCR (RT-PCR) using RNA isolated from young leaves of cacao (Scavina 6). The *TcANR* cDNA (GU324348) contained a 1,008-bp open reading frame (ORF) encoding a protein of 336 amino acids that showed a 63% identity with the *Arabidopsis BANYULS* gene at the amino acid level. The *TcANS* cDNA (GU324350) contained an ORF of 1,062-bp, which encodes a protein of 354 amino acids with 82% amino acid identity with the *Arabidopsis LDOX* gene. The *TcLAR* cDNA (GU324352) contained a 1,083-bp ORF encoding a protein of 361 amino acids with 61% amino acid identity with the *Desmodium LAR* protein.

To determine the genomic sequences of the putative cacao ANS, ANR and LAR genes, the cDNAs were used to screen a cacao BAC library by hybridization. A portion of each hybridizing BAC clone was sequenced using primers designed from the corresponding cDNAs. The genomic structure of each gene was established by alignment with its cDNA sequence (Figure 2). We also retrieved gene models of the ANS and ANR gene from *Arabidopsis* (AT4G22880 and AT1G61720) and *LAR* from *Medicago* and grape (BN000703 and NC_012007 c2622652-2619277) and compared them with the corresponding cacao genes.

The coding region of *TcANR* (GU324347) consisted of 6 exons and 5 introns distributed over 2,005-bp. The coding region of *TcANS* (GU324348) was shorter, having only 1,418-bp and consisting of 2 exons and 1 intron. The genomic organizations of these two cacao genes were nearly identical to the corresponding *Arabidopsis* genes, which had the same exon and intron numbers. Moreover, the lengths of the exons were very similar between *Arabidopsis* and cacao: for ANR, the numbers of nucleotides were precisely the same for the exons 2, 3, 4, and 5, although the lengths of introns were more variable.

Since *LAR* does not have an orthologue in *Arabidopsis*, we compared the genomic organization of *TcLAR* with *MtLAR* and *VvLAR* (Figure 2). The genomic organization of the coding region of *TcLAR* (GU324351) was similar to both *MtLAR* and *VvLAR*, consisting of 5 exons and 4 introns. As observed with the ANR gene, the middle three exons (exon 2, 3 and 4) of *LAR* from
all three species had identical lengths. *TcLAR* exhibited an extremely long third intron of 2,338 bp; similarly, *VvLAR* also featured a long third intron of 1,661 bp, while *MtLAR* contained two long introns (intron 1 and intron 2) that are 812-bp and 1,178-bp respectively.

The analysis of the complete cacao genome sequence of the criollo variety was consistent with these results and allowed us to localize the position of each gene relative to the molecular-genetic map [32]. The *ANR* gene was located on chromosome 6 (*Tc06_g018030*, GenBank: GU324347.1) and was 99.75 identical to the cDNA described above. Similarly, a single *ANS* gene was located on chromosome 3 (*Tc03_g026420*, GenBank: GU324349.1) and shared 99.3% identity to the cDNA described above. The *LAR* gene was also located on chromosome 3 (*Tc03_g002450*, GenBank: GU324351.1) and shares 98.5% identity to its corresponding cDNA. However, two more distantly related genes were identified in the cacao genome assembly that were not identified in the cacao EST collection. One gene (*Tc05_p002410*) encodes a protein that shares only 37% identity and 57% similarity with the *TcANS* protein and a second (*Tc02_g034610*) shares 63% identity and 76% similarity with the *TcLAR* protein. For this study, we focused our further work on the three genes of highest similarity to the well-characterized corresponding Arabidopsis and *Desmodium* genes.

LAR and ANR proteins belong to the reductase-epimerase-dehydrogenase (RED) superfamily, although their relationships are rather distant. ANS belongs to a different protein superfamily, the 2-oxoglutarate-dependent dioxygenase (2-ODD) superfamily, although it shares the same substrate with LAR (Figure 1). A phylogenetic tree was constructed using the neighbor-joining method with the sequences of functionally-tested proteins of LAR, ANR, ANS and DFR from various plant species. The tree construction also included all the IFR-like proteins from Arabidopsis that are most closely related to LAR proteins (Figure 3, see protein alignment in Additional file 1: Figure S1). The tree was noticeably bifurcated into two clades: all RED proteins (including ANR, LAR, DFR and IFR) constituting one clade, and all ANS proteins constituting the other. Within the RED superfamily clade, the IFR, LAR, ANR and DFR proteins are clearly divided into four distinct groups with IFR and LAR forming a subgroup that is distantly related to the subgroup formed by DFR and ANR groups. The cacao ANR, ANS and LAR proteins used in the current research all clustered within their own groups.

Expression profiles of *TcANR*, *TcANS* and *TcLAR* genes

To assess the involvement of *TcANR*, *TcANS* and *TcLAR* in PA biosynthesis in cacao, transcript levels were investigated in various tissues including leaves, flowers and pod tissues (Figure 4A, tissue definition and collection are described in Methods). Gene transcripts levels were assessed by semi-quantitative RT-PCR. *TcActin* was chosen as a reference to normalize gene expression because both cacao microarray analyses (Z. Shi and S. Maximova, unpublished data) and data from this study suggested a relatively constant spatial and temporal expression of this gene for all cacao tissues examined. PA synthesis genes, *TcANR*, *TcANS* and *TcLAR* were expressed in all tissues examined, with relatively higher expression levels in pod exocarp and seeds and lower expression in leaves and flowers. Moreover, their expression levels were relatively similar within each tissue.
except seeds, in which the expression level of TcL AR was much lower than TcANR and TcANS.

PA levels in various cacao tissues
To determine the concentrations of PAs in different cacao tissues, samples were first extracted to obtain a soluble PA fraction. The residues left after soluble PA extraction were assayed using butanol-HCl (Yu-Guang [33]) to measure the amount of insoluble PAs represented as larger polymers. Because insoluble PA polymers will crystallize and bind to proteins and cell wall components, this interference may reduce the extraction efficiency of the insoluble PAs [24,34]. As a result, comparing the relative amount of these two fractions within the same tissue is difficult. However, the accumulation pattern of each fraction is comparable among different tissues. High levels of both soluble and insoluble PAs were detected in all tissues examined, ranging from approximately 40 mg/g DW in seeds to more than 170 mg/g DW in exocarp for soluble PAs (Figure 4B) and ranging from approximately 1.2 mg/g DW in seeds to 8 mg/g DW in exocarps for insoluble PAs (Figure 4C). The accumulation of both soluble and insoluble PAs were in good correlation with the expression patterns of
PA synthesis genes, both of which showed highest levels in fruit exocarp. However, both soluble and insoluble PA levels were much lower in seeds than in other tissues, which correlated with TcLAR but not TcANR and TcANS.

To determine the composition of monomer PAs in cacao, soluble PAs extracts prepared from all four tissues were further separated and quantified by HPLC. The data are presented for catechin and epicatechin concentrations as μg of catechin or epicatechin per g of dry tissue (Figure 4D-F). PAs consisted almost entirely of epicatechin with less than 2% catechin. In all tissues examined, the accumulation of both catechin and epicatechin were well correlated with the expression of TcLAR and TcANR. In exocarp, where both TcLAR and TcANR had the highest expression among all tissues tested, both catechin and epicatechin were also at maximal levels. In leaves and flowers, where TcANR expression was much lower, epicatechin was found at correspondingly lower levels. Likewise in seeds, which exhibited a much lower expression of TcLAR, catechin levels were also much lower.

Functional analysis of the TcANR gene
While the sequences and expression patterns of the candidate cacao PA biosynthesis genes were consistent with their candidate functions relative to enzymes from other species, we conducted in vivo functional analysis of each gene in transgenic plants to gain direct evidence for their functions. To investigate the in vivo function of TcANR, a genetic complementation experiment was performed by transferring the TcANR coding sequence under the control of an enhanced expression promoter (E12Ω, a modified CaMV35S promoter having the
omega arrangement derived from tobacco mosaic virus) into the Arabidopsis banyuls (ban) mutant, which is defective in the gene encoding ANR [22]. ban mutant seeds are pale yellow due to a lack of PAs, and there is more purple anthocyanin pigment deposited in hypocotyls compared to Arabidopsis (ecotype Columbia) control plants due to increased metabolic flux into this branch of the pathway. From eight independent transgenic TcANR-overexpressing lines tested, five lines showed white hypocotyls, and 3 lines showed reduced pigmentation in hypocotyls as compared to banyuls mutants. The lines with white hypocotyls also produced seeds exhibiting the wild-type phenotype that stained blue with DMACA reagent, suggesting the deposition of PAs in the seed coat (Figure 5A). After PA extraction and quantification, all lines showed significantly increased levels of PAs (Figure 5C). RT-PCR analysis confirmed expression of TcANR, and the expression levels positively correlated with PA accumulation (Figure 5B).

### Functional analysis of TcANS gene

To investigate the *in vivo* function of TcANS, two model plants, tobacco and Arabidopsis were utilized. We used an Arabidopsis ans (ldox) mutant to perform tests of transgenic complementation. We used tobacco as a model system for metabolic flux analysis of the equilibrium between PA and anthocyanin synthesis. Introduction of functional ANS genes into tobacco can result in flower petal color changes reflecting alterations in ANS activity.

![Figure 5](image-url)
we cannot obtain direct evidence of the TcL AR
expression of the ANR
TcL AR
confirmed high
color were chosen for further analysis. RT-PCR analysis
The two lines displaying the greatest increase in petal
increase in pink color intensity in flower petals.
The two lines displaying the greatest increase in petal
color were chosen for further analysis. RT-PCR analysis
confirmed high TcANS transcript levels in these two to-
bacco transgenic lines, which positively correlated with
the color of the petals (Figure 6A and B). Amplification
of the tobacco ribosomal RNA gene NtrRNA, which
served as an internal control, showed a relatively similar
expression level in both wild-type control and transgenic
tobacco plants. As predicted, anthocyanin levels in-
creased in the two transgenic lines (Figure 6C).
The levels of PAs in the petals of transgenic lines,
quantified by DMACA assays, were also significantly
higher as compared to untransformed Samsun plants
(Figure 6D). On average, a two-fold increase of PA was
observed in the two lines compared to wild type.
The 35S-TcANS transgene was also introduced into the Arabidopsis ans (ldox) T-DNA mutant, which pro-
duces hypocotyls that appear white to light green due to
lack of anthocyanins, and seeds that appear light yellow
due to lack of PAs. Eighteen independent hygromycin
resistant transgenic T1 seedlings were selected. From
these, 2 lines developed wild-type purple colored hypo-
cotyls (Additional file 1: Figure S2) and produced wild-
type brown-colored seeds that stained blue after reacting
with the DMACA reagent. The color suggested depos-
tion of PAs in the seed coat, however the color intensity
was lower than in wild type. RT-PCR using RNA ex-
tracted from T2 seedlings confirmed expression of
TcANS genes (data not shown).

Functional analysis of the TcLAR gene
Because the LAR gene is not present in Arabidopsis [15],
we cannot obtain direct evidence of the in vivo function of
TcLAR through genetic complementation analysis in Arabidopsis. Therefore, to functionally characterize
TcLAR, we used tobacco as a model system. As over-
expression of the ANR gene can divert the metabolic
flow from anthocyanin synthesis to PA synthesis [17],
we predicted that over-expression of the cacao LAR gene
in tobacco would result in a decrease in anthocyanin
pigment and an increase in PA accumulation in flower
petals. Transgenic tobacco plants were generated that
constitutively expressed the ORF of TcLAR under the control of the E12Ω promoter. Twenty-two independent
transgenic lines that are resistant to hygromycin were
generated. Nine of these exhibited a decrease in intensity
of the visible pink color of petals. Two lines (Lines
LAR7C1 and LAR9A2) exhibited virtually white petals
(Figure 6E). RT-PCR showed that both lines expressed
high levels of TcLAR transcripts (Figure 6F). Quantifica-
tion of soluble PA and anthocyanin levels indicated that
anthocyanin levels in these two lines were about half
those of WT controls, and that total PAs accumulated to
an approximately 5-fold higher level than in controls
(Figure 6G and H).
Moreover, the levels of PAs in transgenic tobacco
petals were inversely proportional to the concentrations
of anthocyanin, indicating diversion of metabolic flow
from anthocyanin to PA synthesis. Insoluble PA levels
were also estimated, but the levels were insignificant
(data not shown). To confirm the increased accumula-
tion of catechin, as it is the predicted product of TcLAR,
we separated and quantified the PA extracts using
HPLC. The HPLC analysis showed that there is indeed
an approximately 2-fold increase of catechin levels in
both transgenic lines. Surprisingly, there is also a signifi-
cant increase of epicatechin levels (Figure 6I).

We also took advantage of the Arabidopsis ans (ldox)
T-DNA mutant to examine LAR function. Because Arabidopsis lacks an LAR gene [12,14,15] and the ldox
mutant is deficient in cyanidin (the substrate for ANR),
the ldox mutant exhibits a significant decrease of
epicatechin and PA synthesis. We reasoned that since
the ldox mutant accumulates leucoanthocyanidin which
can provide the substrate for a heterologous LAR pro-
tein, the ldox mutant was potentially a good in vivo
model to analyze LAR function. We predicted that over-
expression of TcLAR should result in the synthesis of
catechin in developing siliques of the ldox mutant Arabi-
dopsis, even if PAs were not produced due to lack of
epicatechin synthesis. HPLC separation and quantifica-
tion of PA extracts from Arabidopsis transgenic lines
over-expressing TcLAR gene in the ldox mutant back-
ground revealed not only a significant increase of
catechin, but also surprisingly, a modest increase of epi-
catechin (Figure 7). Quantification of total PAs extracted
from mature seeds also revealed significant PAs in-
creases in all transgenic lines compared to the ldox
mutant (Additional file 1: Figure S4).

These unexpected results prompted us to check the
possibility that TcLAR may carry a dual enzymatic
activity, capable of converting leucoanthocyanidin to both
catechin and epicatechin. Thus, recombinant TcLAR
Figure 6 (See legend on next page.)
protein was expressed in *E. coli*, purified and assayed using $^3$H-labeled leucocyanidin as substrate in the presence of NADPH, followed by analysis of products by HPLC-UV and radioactivity detection. The negative control reaction (boiled protein) showed no product formation (Figure 8A and C), whereas a peak with the same retention time and UV-spectrum as that of the pure $^3$H-leucocyanidin (Figure 8B and D). However, no epicatechin product was detected (the retention time of 26–27 min), suggesting that TcLAR does not exhibit duel functionality in this *in vitro* assay.

**Discussion**

The distribution of cacao PA accumulation correlates with expression of the PA biosynthesis genes

The localization of PAs in plant tissues has been well studied in the model plants Arabidopsis, the legume *Medicago truncatula*, and in the fruit species grape (*Vitis vinifera*). In Arabidopsis, the expression of the *AtANR* (*BANYULS*) gene and PA accumulation is limited to the seed coat [22,35]. Similarly in *Medicago* the major localization of PAs is also in the seed coat with very small amounts present in flowers, leaves, roots and stems [24]. Although *Theobroma cacao* and Arabidopsis are phylogenetically closely related, the PA accumulation profiles are quite different. Our results demonstrated that in cacao leaves, flowers and fruits, PAs accumulate at high levels and this is correlated with expression of genes encoding the key enzymes involved in PA synthesis. This is similar to *Vitis*, in which both PA accumulation and genes involved in PA synthesis are found in leaves, fruit skins and seeds [25].

In cacao, the expression of *ANS*, *ANR*, and *LAR* were co-regulated and their expression correlated well with PA accumulation in most of the tissues, suggesting significant roles in PA synthesis for both ANR and LAR. The only exception is in seeds, in which the expression level of *TeANS* is higher than *TeANR* and *TeLAR* is much lower. The high expression level of *TeANS* may contribute to anthocyanin synthesis, because at the time when seed tissues were harvested (18 WAP), anthocyanins are actively synthesized [36-38]. The PA level in seeds is also much lower than in other tissues, well correlated with the *TeLAR* but not *TeANR*. The LAR enzyme seems to be a rate-limiting factor of the PA synthesis. To further explore the individual contributions of these proteins to catechin and epicatechin synthesis, flavan-3-ol monomer composition was examined by HPLC analysis. Cacao flavan-3-ol monomers were composed almost entirely of epicatechin units with catechin units comprising less than 2% in all tissues examined. This result is consistent with previous study of PA synthesis in cacao beans [30], in which PA extension units were reported to be composed exclusively of epicatechin, and terminal units were mostly epicatechin with about 1% catechin. Similarly, *Medicago* PAs consist almost entirely of epicatechin units [24]. In grape, both ANR (encoded by *VvANR*) and LARs (encoded by *VvLAR1* and *VvLAR2*) contribute to PA synthesis in the fruits, in which PAs consist of both catechin and epicatechin. However, in grape leaves, although catechin is still present, LAR genes are expressed at very low levels.

Functional characterization of cacao PA biosynthetic genes ANS, ANR and LAR

*In vivo* genetic analysis of *TeANS* and *TeANR* verified their roles in PA biosynthesis. Over-expression of *TeANS* in tobacco resulted in elevated levels of both anthocyanins and PAs in flower petals. It also restored anthocyanin synthesis in hypocotyls as well as PA accumulation in seeds of the Arabidopsis *ans* (*ldox*) mutant. Similarly, over-expression of *TeANR* in the Arabidopsis *banyuls* (*anr*) mutant restored PA synthesis in seeds. Moreover, ectopic over-expression of *TeANR* in hypocotyls diverted the anthocyanin synthesis branch and resulted in decreased anthocyanin pigments related to gene expression levels. The catalytic activities of LAR proteins from grapevine, *Medicago* and *Desmodium* have been verified.
by *in vitro* assays of recombinant proteins [14,24,25]. *Lotus* LAR showed weak activity in a recombinant DFR protein coupled assay [26]. However, none of the enzyme assays was able to recover sufficient flavan-3-ol products for stereo chemical analysis to allow specific determination of catechin and epicatechin. The *in vivo* activities of *Medicago* and *Desmodium* LAR proteins was also tested in tobacco and white clover [14,24]. In the former study [14], the authors found that transgenic plants expressing the DuLAR protein showed slightly elevated LAR enzymatic activity compared to extracts from untransformed plants, but they did not demonstrate changes in metabolite synthesis. In a more recent study of *MtLAR* overexpressing transgenic tobacco plants, there were no detectable changes in anthocyanin and PA levels [24]. The results presented in this work shows that a cacao *TcLAR* gene is co-regulated with *TcANR* in all tissues examined, providing a simple model to investigate LAR enzyme functions. In our results, *TcLAR* showed clear *in vivo* activities in both transgenic tobacco and Arabidopsis plants by converting metabolic flow from anthocyanin synthesis to PA synthesis. This
data provides a direct genetic evidence for a clear role of LAR in PA biosynthesis. The metabolic flow divergence resulting from TcLAR over-expression could be due to perturbations of a hypothesized metabolic channeling mechanism that suggests that multiple enzymes in each subsequent step of the synthesis pathway interact as a means to increase the efficiency and throughput of the pathway [39].

To our surprise, ectopic expression of TcLAR resulted in elevated levels of both catechin and epicatechin units in tobacco flowers (Figure 6H) in contrast to the prediction that only catechin should be formed. Based on the sequence similarity between LAR and ANR, it is possible that LAR could function redundantly to ANR and convert cyanidin to epicatechin. Alternatively, LAR may perform as a dual functional enzyme and convert leucoanthocyanidin to both catechin and epicatechin. To test the possibility that LAR performs the same activity as ANR and uses cyanidin as a substrate to form epicatechin, we took advantage of the Arabidopsis idox (ans) mutant. Since Arabidopsis does not have an LAR gene and synthesizes only epicatechin through ANS and ANR pathway, the idox mutants have a significantly reduced PA (epicatechin) level due to lack of a supply of

Figure 8 Enzymatic assay of recombinant TcLAR. A, HPLC analysis of products from incubation of leucocyanidin with boiled protein as control, monitored by UV spectroscopy. B, as above, for incubation of purified recombinant TcLAR with 3H-leucocyanidin. C, As in (A), but monitoring radioactivity of fractions; D, As in (B), but monitoring radioactivity of fractions; (E), UV spectrum of catechin product from (B). F, HPLC analysis of an authentic standard of (−)-catechin. mAU, milliabsorbance units; CPM, counts per minute.
cyanidin, the epicatechin precursor. Thus, the ldox mutant provides a system that has leucoanthocyanidin but not cyanidin. Over-expression of TclAR in the ldox mutant resulted in synthesis of catechin confirming its predicted enzymatic function. However, HPLC quantification showed that there was only a slight elevation of epicatechin levels compared to the significant elevation of catechin, suggesting that LAR was less likely to perform dual function and convert leucoanthocyanidin to epicatechin.

When recombinant TclAR was assayed in vitro, while we observed the production of catechin from 2,3-trans-3,4-cis-leucoanthocyanidin, we did not detect even trace amounts of epicatechin. The failure to detect epicatechin in assays with the cacao LAR enzyme suggests that it functions solely in converting leucoanthocyanidin to catechin. However, it is possible that LAR enzyme can synthetize epicatechin from 2,3-cis-3,4-trans-leuco cyanidin as suggested by [14]; but we could not test this possibility in the recombinant TclAR enzyme assay because 2,3-cis-3,4-trans-leuco cyanidin is not commercially available. The increase in epicatechin in TclAR over-expressing Arabidopsis and tobacco plants could be due to the possible existence of a gene(s) encoding a catechin-epicatechin epimerase. One possible candidate gene that could be involved is ANR. Vitis vinifera ANR protein expressed and purified from E.coli showed epimerase activities in vitro and converted cyanidin to a mixture of catechin and epicatechin [40]. Similarly, two recombinant ANR proteins cloned from Camellia sinensis also showed epimerase activities in vitro [41]. However, we regard this as a remote possibility because Arabidopsis itself does not synthesize catechin and thus it would be surprising if a catechin-epicatechin-specific epimerase were expressed in the absence of its substrate. Nevertheless, we cannot exclude the possible existence of such an epimerase that may also serve other functions important in plant development. A second alternative hypothesis to explain epicatechin formation would be racemization of catechins by polymerization to proanthocyanidins followed by nonstereo-specific depolymerization. An earlier report lends some support to this idea through the analysis of transgenic apple lines in which the

Conclusions

We successfully isolated three genes encoding key PA synthesis enzymes in Theobroma cacao, TclANS, TclAR, and TclANR. In vivo genetic analysis of TclANS and TclANR in tobacco and complementation in Arabidopsis verified their roles in PA biosynthesis. In vitro enzyme assays of TclAR recombinant protein verified its predicted function. Moreover, in vivo overexpression of TclAR in tobacco and Arabidopsis ldox mutants successfully diverted metabolic flow to PA synthesis, which provide direct evidence for a clear role of the TclAR gene in PA biosynthesis. Our results provide new knowledge and genetic tools for development of cacao varieties with novel PA profiles through conventional breeding or genetic approaches.

Methods

Plant material

Two Theobroma cacao varieties: Scavina 6 and Amelonado were used for this study. Cacao plants were grown in greenhouse as previously described [44]. Leaf and flower tissues were collected from Scavina 6 plants. For leave tissues, young red stage leaves were collected. The definition of leaves stages were previously described [45]. Cacao pods were obtained by hand pollinating Amelonado (a self-compatible variety), in order to reduce the effect of genetic variation on seed traits. Pods harvested 18 weeks after pollination were dissected into exocarp (outer fruit tissue) and seeds for separate analysis. Exocarp samples represent the outer 1–3 mm layer of the fruit obtained using a fruit peeler. All samples were frozen in liquid nitrogen upon collection and stored at −80°C until extraction.

Transgenic and wild-type tobacco plants (Nicotiana tabacum var. Samsun) provided by Wayne Curtis, Department of Chemical Engineering, The Pennsylvania
State University) were grown in a greenhouse under the same condition as cacao plants. Arabidopsis plants (Arabidopsis thaliana) were grown in soil at 22°C, 50% humidity and a 16 h/8 h light/dark photoperiod in a growth chamber (Conviron, Pembina, ND, USA). Plants grown aseptically were plated on MS medium [46] with 2% (w/v) sucrose solidified with 0.6% (w/v) agar. Arabidopsis ecotype Columbia (Col –0) plants were used as the wild type. T-DNA insertion mutants ban (SALK_040250) and ldox (SALK_028793) were obtained from The Arabidopsis Biological Resource Center (Columbus, OH, USA).

Nucleic acid purification and cDNA synthesis
Total RNA from leaves of Theobroma cacao (Scavina 6) was isolated using a modified cetyl trimethyl ammonium bromide (CTAB) extraction method as previously described [47] with the following modifications. RNA isolated from the CTAB extraction LiCl precipitation was further purified and concentrated using RNeasy columns (Qiagen, Valencia, CA, USA), but the phenol/chloroform extraction and sodium acetate/ethanol precipitation step was omitted. The quality of RNA was verified by observing absorbance ratios of A260/A280 (1.8 to 2.0) and A260/A230 (1.8 to 2.2) and by separating 200 ng RNA samples on 0.8% agarose gels to examine intact ribosomal bands. First strand cDNA was synthesized using the SMART RACE cDNA amplification kit (Clontech, Mountain View, CA, USA).

Isolation of cDNA and genomic clones from Theobroma cacao
The putative expressed sequence tag (EST) sequences of cacao anthocyanidin reductase (TcANR), anthocyanidin synthase (TcANS) and leucoanthocyanidin reductase (TcLAR) genes were obtained by searching the EST database (http://esttik.cirad.fr/) [31] using the tBLASTn program [48]. The query sequences were used as the protein sequences of BANYULS and LDOX from Arabidopsis thaliana and DuLAR from Desmodium uncinatum respectively (Accession numbers: NP_176365, Q96323 and CAD79341). Based on the sequences of the ESTs from the EST database (EST Treatment and Investigation Kit; http://estтик.cirad.fr), PCR primers were designed to amplify the entire coding sequences of each gene: ANR_F (5’-AGCCATGGGACG- CAGACCGTAGG-3’) and ANR_R (5’-GCGGCGCTCA CTTGAGCAGCCCCCTTAGC-3’), ANS_F (5’-CCATGTT GACTTCAATGGCCCCAG-3’) and ANS_R (5’-GCG GCGCGCTCAATTAGACAGGCCCATC-3’) and LAR_F (CCATGATATGAAAATCAACAAACATGAATGGTTC) and LAR_R (GCGGCGCGCTATGTCATATCCAGTG). NcoI sites were added to the 5’ end of each start codon and NotI sites were added to the 3’ end of each stop codon to facilitate the subsequent cloning into binary T-DNA vectors. The coding sequences were amplified from cacao cDNA prepared from young leaves (genotype Scavina 6) with the Advantage cDNA PCR Kit (Clontech, Mountain View, CA, USA) using these primers. PCR reactions were carried out in a total volume of 20 μL at 94°C for 5 min; 5 cycles of 94°C for 30 sec, 55°C for 30 sec, and 72°C for 1 min; then another 23 cycles of 94°C for 30 sec, 60°C for 30 sec, and 72°C for 1 min; followed by a final extension at 72°C for 5 min. PCR products were gel purified and cloned into the pGEM-T easy vector (Promega, Madison, WI, USA). The correct open reading frames (ORFs) of each of the resulting constructs (pGEMT-TcANR, pGEMT-TcANS and pGEMT-TcLAR) were confirmed by DNA sequencing. The DNA sequences of the TcANR, TcANS and TcLAR genes were obtained by isolation and sequencing of genomic clones. Briefly, 2 high-density filters were arrayed with 18,432 colonies of Theobroma cacao (genotype LCT-EEN 37) bacterial artificial chromosome (BAC) clones on each (library and filters constructed by The Clemson University Genomics Institute (CUGI, https://www.genome.clemson.edu/)). The filters were hybridized to full-length cDNA of each gene labeled with P32 using the MEGA Labeling Kit (GE Healthcare, Piscataway NJ). DNA hybridizations were carried out at 65°C in 1 mM ethylenediaminetetraacetic acid (EDTA), 7% sodium dodecyl sulfate (SDS), 0.5 M sodium phosphate (pH7.2) for 16–18 h. Filters were washed twice at 65°C in 1 mM EDTA, 1% SDS, 40 mM Na2HPO4 for 20 min, twice at 65°C in 1.5× sodium chloride/sodium citrate (SSC), 0.1% SDS for 20 min and twice at 65°C in 0.5× SSC, 0.1% SDS for 20 min. Two or more BAC clones were identified for each gene and confirmed by PCR using plasmid DNA from individual colonies and gene specific primers. High purity plasmid DNA from individual BAC clones was then isolated using a NucleoBond BAC 100 kit (Macherey-Nagel Inc., Bethlehem, PA, USA) and both strands of DNA were sequenced using primers designed from the cDNAs and genomic DNA. DNA sequencing results were analyzed and assembled using Vector NTI software (Invitrogen, San Diego, CA), application Contig Assembly. The DNA sequence of each gene was then compared to the corresponding coding sequence by using the BLAST2 online tool (www.ncbi.nlm.nih.gov/blast/bl2seq/wblast2.cgi) to obtain exon and intron locations for the gene organization analyses.

Phylogenetic analysis
Deduced protein sequences of all Arabidopsis IFR-like genes were retrieved from The Arabidopsis Information Resource (TAIR) database (http://www.arabidopsis.org/) by querying the TAIR protein database with the Desmodium LAR protein sequence (CAD79341) using the WU-BLAST2 (BLASTP) program. Protein sequences
from other species were retrieved from the GenBank (http://www.ncbi.nlm.nih.gov.Genbank/). Accession numbers are indicated in the figure legend.

Multiple sequence alignment of proteins was performed by ClustalX algorithm [49] with default parameter settings (gap opening penalty: 10, gap extension penalty: 0.2, delay divergent cutoff: 30%, protein weight matrix: Gonnet series) and this alignment was used to construct the phylogenetic tree using the neighbor-joining method in the MEGA package [50]. Five hundred bootstrapped datasets were used to estimate the confidence of each tree clade.

**Determination of proanthocyanidins (PAs) and anthocyanins**

For quantitative analysis of anthocyanin levels in transgenic tobacco flowers, fresh petals (0.3-0.5 g fresh weight) from three flowers were immersed in 5 mL ethanol: 6 M HCl (1:1) and incubated at 4°C overnight. The extract solution was transferred to a new tube and the petals were extracted for the second time using the same method. Absorbance of the pooled extract solution was then measured at 526 nm and the total anthocyanin levels were calculated using a standard molar absorbance curve prepared using cyanidin-3-glucoside (Sigma-Aldrich, MO, USA).

To extract soluble PAs from cacao and tobacco tissues, 0.3 to 0.5 g of frozen tissues were ground into a fine powder in liquid nitrogen and then extracted with 5 mL of extraction solution (70% acetone: 29.5% water: 0.5% acetic acid) by vortexing for 5 seconds followed by water bath sonication for 15 min using a bench top ultrasonic cleaner (Model 2510, Bransonic, Danbury, CT, USA). To extract soluble PAs from Arabidopsis seeds and siliques, the same extraction solution and method were applied, except that 100 to 500 mg dry seeds and 10 green siliques were used as grinding samples, and 500 μL extraction solution were used. After sonication, samples were vortexed again and centrifuged at 2500 g for 10 min. The supernatant was transferred to a new tube and the pellet was re-extracted twice as above. Pooled supernatants were extracted twice with hexane to remove fat and chlorophyll and then filtered through a 0.45 μm polytetrafluoroethylene (PTFE) syringe filter (Millipore, Billerica, MA, USA). Depending on availability of plant samples, different numbers of biological replicates were performed for cacao, tobacco and Arabidopsis samples. For cacao, there are at least five biological replicates, for tobacco, there are seven or more biological replicates, and for Arabidopsis, there are three biological replicates.

To quantify soluble PA levels, extracts were then quantified by reaction with p-dimethylamino-cinnamaldehyde (DMACA), which specifically interacts with PA monomers and polymers to form blue pigments [51]. Briefly, 50 μL aliquots of samples were mixed with 200 μL of dimethylaminocinnamaldehyde (DMACA; Sigma-Aldrich, MO, USA) reagent (0.1% DMACA, 90% reagent-grade ethanol, 10% HCl) in 96-well microtiter plates. Absorption was measured at 640 nm at one-minute intervals for 20 min to get the highest readings. Triple technical replicates were performed to obtain mean values. The total PA levels were calculated using the standard molar absorbance curve prepared using procyanidin B2 (Indofine, NJ, USA).

For quantitative analysis of insoluble PAs from cacao tissues, the residues from soluble PA extractions were air dried in an exhaust hood for two days, weighed, and 5 mL butanol-HCl reagent (95% butan-1-ol: 5% concentrated HCl) was added and the mixture was sonicated for one hour followed by centrifugation at 2500 g for 10 min. An aliquot of clear supernatant was diluted 40-fold in butanol-HCl reagent and absorbance was measured at 550 nm to determine the amount of background absorption. The samples were then boiled for 1 hour with vortexing every 20 min, cooled to room temperature and centrifuged again at 2500 g for 10 min. The supernatant from boiled sample was diluted 40-fold in butanol-HCl reagent and absorbance was measured at 550 nm. The values were normalized by subtraction of the background absorbance and the PA levels were calculated as cyanidin equivalents using cyanidin-3-glucoside (Sigma-Aldrich, MO, USA) as standards.

To visualize the presence of PAs in Arabidopsis seeds, dry seeds were immersed for 2 days in the 0.1% DMACA reagent described above and then washed 3 times with 70% ethanol as described previously [35]. Catechin and epicatechin content was determined by reverse-phase HPLC using an Alliance separations module (Model 2695; Waters, Milford, MA, USA) equipped with a multi λ fluorescence detector (Model 2475; Waters, Milford, MA, USA). Samples of soluble PA extracts (10 μL) were separated on a 250 mm × 4.6 mm Luna 5-μm Phenyl Hexyl column (Phenomenex, Torrance, CA, USA) and then assayed by fluorescence emission at 315 nm following excitation at 280 nm. The HPLC separation utilized a binary mobile phase gradient mixture of A+B where mobile phase A was 0.5% trifluoroacetic acid (TFA) (v/v with water) and mobile phase B was 0.5% TFA (v/v with methanol). The gradient conditions were: 0 min, 16% mobile phase B; 4 min, 16% mobile phase B; 14 min, 50% mobile phase B; 18 min, 50% mobile phase B; 22 min, 100% mobile phase B; 26 min, 100% mobile phase B; 30 min, 16% mobile phase B. The column was maintained at 30°C and the flow rate was 1 mL/min. Catechin and epicatechin standards were purchased from Sigma-Aldrich (St. Louis, MO, USA). This work was performed at the Hershey Technical Center (Hershey, PA, USA).
Transformation of tobacco and arabisopsis

The coding sequences of TcANS, TcANR and TcLAR were excised from the cloning vector (pGEM-T easy) (Promega, Madison, WI, USA) with Neol and NotI restriction enzymes and cloned into the pET2113-EGFP [44] intermediate vector to replace the original EGFP coding sequence. As a result, the cacao coding sequences are located immediately downstream of the E12-Ω, an enhanced expression promoter modified from CaMV35S [52], and upstream of the CaMV35S-terminator. The over-expression cassettes of TcANR and TcANS was excised out from pE2113-TcANR and pE2113-TcANS constructs respectively with HaelII restriction enzyme, blunt ended with T4 polymerase and then introduced into the pCAMBIA-1300 binary vector (CAMBIA, Canberra, Australia) linearized with Smal restriction enzyme; the over-expression cassettes of TcLAR was excised out from pE2113-TcLAR construct with HindIII and PvuII restriction enzyme and ligated into pCAMBIA-1300 binary vector linearized with HindIII and Smal restriction enzyme. All binary transformation constructs were introduced into Agrobacterium tumefaciens strain AGL1 [53] by electroporation as described previously [54].

Tobacco leaf disc transformation was performed as previously described [55] and transgenic shoots were regenerated on MSs (MS shooting) media supplemented with 25 mg/L hygromycin. Only one shoot was selected from each explant to ensure independent transformants. After rooting for 2 weeks in MSr (MS rooting) media supplemented with 25 mg/L hygromycin, hygromycin-resistant plantlets were transferred to soil and grown in a greenhouse as described above.

Arabidopsis transformation was carried out using the floral dip method [56], and T1 transgenic plants were selected on MS media supplemented with 2% sucrose, 0.65% agar and 25 mg/L hygromycin. Higromycin-resistant T1 seedlings were transferred to soil 7 days after germination and grown in a growth chamber as described in above.

Expression analysis of TcANS, TcANR and TcLAR

Total RNA from leaves, flowers, whole pods, pod exocarp and ovules of Theobroma cacao (Scavina 6 and Amelonado) was isolated as described above. Total RNA from young leaves of transgenic and wild-type tobacco plants as well as Arabidopsis plants was isolated using the RNeasy Plant mini kit (Qiagen, Valencia, CA, USA). cDNA was synthesized from 1 μg of total RNA in a total volume of 20 μL using M-MuLV Reverse Transcriptase (NEB, Ipswich, MA, USA) according to the supplier's protocols, and 2 μL were used in the subsequent reverse transcription-PCR (RT-PCR) reactions. The primers for RT-PCR were designed to amplify across at least one intron giving products of approximately 500 bp from cDNA and 700 bp to 1500 bp from genomic DNA. These primer sets were used to check all cDNAs for genomic DNA contamination. The primers used for TcANS were TcANSRT_F (5′-ACCTTGTAAACCTCTGAGATCTCGG-3′) and TcANSRT_R (5′-GACGGTTGTCACCAAATGTGATGAT-3′); the primers used for TcANR were TcANR_F (5′-TGCTTGAAGAGGCTACGGTGTGTTTA-3′) and TcANR_R (5′-GAAAGATGGTCGAAGGCCAATGCTG-3′); the primers used for TcLAR were TcLAR_F (5′-AATTCCATTGCAGCTTGTCGCGT-3′) and TcLAR_R (5′-GGCTTGCTACTGTTTGCGATTAT-3′). TeActin was used as an internal standard for cacao gene expression using primer set Tc6RT_F (5′-AGGCTGAGATTCCTCGTCTGCAGA-3′) and Tc6RT_R (5′-CTCAGATCACTGTTGTGAT-3′) [57], and NtrRNA_F (5′-AGGAATGGTCGAAGGCCAATGCTG-3′) and NtrRNA_R (5′-GTGCCGCCAGAACCAGTCATAAGG-3′) [58].

The number of PCR cycles was optimized between 20 and 32 to select a cycle number such that amplification was in the linear range; 28 cycles were chosen for all the RT-PCR reactions. The PCR reaction was carried out in a total volume of 20 μL at 94°C for 5 min; 28 cycles of 94°C for 30 sec, 55°C for 30 sec, and 72°C for 45 sec; followed by a final extension at 72°C for 5 min. The PCR products were visualized on 1% agarose gels stained with ethidium bromide (EtBr) and documented using Molecular Imager Gel Doc XR + System equipped with a 16-bit CCD camera (Bio-Rad Laboratories, Hercules, CA) and bands were quantified using Quantity One 1-D Analysis Software (Bio-Rad Laboratories, Hercules, CA).

Assay of LAR activities

The open reading frame (ORF) of the TcLAR gene was PCR amplified from pGEMT-TcLAR using Advantage 2 polymerase mix (Clontech, Mountain View, CA) and the following primers: TcLARCDF1 (5′-CCCACATCAACCAGACTTTGAGTTC-3′) [57], and TcLARCDR2 (5′-AGGCCCTCAACTGTTGTGAT-3′) [57], and NtrRNA_F (5′-AGGAATGGTCGAAGGCCAATGCTG-3′) and NtrRNA_R (5′-GTGCCGCCAGAACCAGTCATAAGG-3′) [58].

The open reading frame (ORF) of the TcLAR gene was PCR amplified from pGEMT-TcLAR using Advantage 2 polymerase mix (Clontech, Mountain View, CA) and the following primers: TcLARCDF1 (5′-CCCACATCAACCAGACTTTGAGTTC-3′) [57], and TcLARCDR2 (5′-AGGCCCTCAACTGTTGTGAT-3′) [57], and NtrRNA_F (5′-AGGAATGGTCGAAGGCCAATGCTG-3′) and NtrRNA_R (5′-GTGCCGCCAGAACCAGTCATAAGG-3′) [58].
expression, a single bacterial colony was inoculated into Luria-Bertani medium (10 g/L tryptone, 5 g/L yeast extract, 10 g/L NaCl) containing 100 μg/mL ampicillin and grown at 37°C overnight. An overnight culture was then diluted into terrific broth (TB) medium (12 g/L Tryptone, 24 g/L Yeast Extract, 0.4% glucose, 2.31 g/L KH2PO4, 12.54 μg/mL/L K2HPO4) containing 100 ampicillin and grown at 37°C until the OD600 reached 0.6-0.8, at which time IPTG (isopropyl β-D-1-thiogalactopyranoside) was added to a final concentration of 1 mM to induce protein expression. Recombinant TcLAR protein with a 6-His tag at the C terminus was purified using a MagneHis kit (Promega, Madison, WI, USA) and the protein concentration measured by the Bradford method [59]. This work was performed at the Samuel Roberts Noble Foundation (Ardmore, Oklahoma, USA).

3H-3,4-cis-leucocyanidin was synthesized as described by [60]. Assay of recombinant TcLAR protein with 3H-3, 4-cis-leucocyanidin was carried out in a final volume of 100 μL containing 10% (w/v) glycerol, 100 mM potassium phosphate (pH 7.0), 4 mM dithiothreitol (DTT), 0.5 mM NADPH, 0.4 mM 3H-leucocyanidinoglucomoi of recombinant 30 TcLAR protein. The reaction was initiated by the addition of enzyme and incubated at 30°C for 1 h. The assay was terminated by the addition of 20 μL of methanol followed by centrifugation. Products were analyzed by HPLC, with absorbance monitoring at 280 nm. Products eluting at retention times between 13 to 31 min were collected (1 min/tube) and the fractions containing labeled products were identified by liquid scintillation counting. Boiled pure protein was used as a control.

Reverse-phase HPLC analysis of enzymatic products was performed using an Agilent HP1100 HPLC (Agilent Technologies, Inc., Santa Clara, CA, USA) with the following gradient using solvents A (1% phosphoric acid) and B (acetonitrile) at a 1 mL/min flow rate: 0 to 5 min, 6% B; 5 to 10 min, 6% to 10% B; 10 to 20 min, 10% to 11% B; 20 to 25 min, 11% to 12.5% B; 25 to 45 min, 12.5% to 37% B; 45 to 48 min, 37% to 100% B; 48 to 58 min, 100%, 58 to 60 min 100% to 6% B. Absorbance data were collected at 280 nm. Identifications were based on comparison of chromatographic behavior and UV spectra with authentic standards. This work was performed at Samuel Roberts Nobel Foundation (Ardmore, Oklahoma, USA).

Sequence data from this article can be found in the GenBank/EMBL data libraries under accession numbers GU324347 (TcANR), GU324349 (TcANS) and GU324351 (TcLAR).

Additional file 1: Figure S1. Multiple sequence alignment of the LAR, ANS and ANR proteins as well as related IFR and DFR proteins of the RED superfamily. Figure S2. Complementation of the PA and anthocyanin deficient ans (ldox) mutant phenotype by constitutively expressing TcANS. Figure S3. HPLC analysis of extracts from transgenic Arabidopsis and tobacco key samples with and without standard spikes to validate the peak of catechin and epicatechin. Figure S4. Complementation of the PA deficient Arabidopsis idox mutant phenotype by constitutive expression of TcLAR.

Competing interests
The authors declare that they have no competing interests.

Authors’ contributions
YL performed most of the experiments, i.e., sequence analysis, gene cloning, gene expression studies, transgenic tobacco and Arabidopsis generation, phenotypic analysis of transgenic lines, and drafted the manuscript. ZS participated in gene expression analysis, transgenic Arabidopsis lines generation and analysis, SNM participated in the design of the study, directed the vector construction and tobacco transgenic lines generation, and participated in drafting of the manuscript. MJP developed the catechin and epicatechin HPLC quantification assays and directed PA quantification analysis. MJG conceived the study, drafted the manuscript and gave advice on experimental design, data analysis and execution. All authors read and approved the final manuscript.

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