The rapid alkalinization factor (RALF) gene family is essential for the plant growth and development. However, there is little known about these genes among Rosaceae species. Here, we identify 124 RALF-like genes from seven Rosaceae species, and 39 genes from Arabidopsis, totally 163 genes, divided into four clades according to the phylogenetic analysis, which includes 45 mature RALF genes from Rosaceae species. The YISY motif and RRXL cleavage site are typical features of true RALF genes, but some variants were detected in our study, such as YISP, YIST, NISY, YINY, YIGY, YVGY, FIGY, YIAY, and RRVM. Motif1 is widely distributed among all the clades. According to screening of cis-regulatory elements, GO annotation, expression sequence tags (EST), RNA-seq, and RT-qPCR, we reported that 24 RALF genes coding mature proteins related to tissue development, fungal infection, and hormone response. Purifying selection may play an important role in the evolutionary process of RALF-like genes among Rosaceae species according to the result from \( ka/ks \). The tandem duplication event just occurs in four gene pairs (Fv-RALF9 and Fv-RALF10, Md-RALF7 and Md-RALF8, Pm-RALF2 and Pm-RALF8, and Pp-RALF11 and Pp-RALF14) from four Rosaceae species. Our research provides a wide overview of RALF-like genes in seven Rosaceae species involved in identification, classification, structure, expression, and evolution analysis.

Keywords: RALF; Rosaceae species; identify; classify; evolution; expression
physiochemical properties. As such, they hold that the peptides of clade IV are not the true RALF genes, and are described as the RALF-like peptides. Collectively, the release of mature RALF peptides is important to functioning.

RALF was initially discovered in the medium of tobacco suspension cells during the peptide screen for systemin proteins (defense-related, alkalinization-inducing peptides) [2]. Homologs of RALF have already been found in the same plant kingdom; for example, five homologs were found in poplar [5], 39 homologs of RALF were found in Arabidopsis thaliana [6], and 12 of these genes from the expression sequence tag (EST) database. RALFs have been identified in a variety of species including monocots, eudicots, and early diverging lineages [2,5–11], and Campbell and Turner [4] identified 765 RALF proteins from 51 plant species.

According to previous studies, RALFs have wide expression profiles, which vary based on tissue growth stage, biotic, or abiotic treatments. In Arabidopsis thaliana, RALF-like genes showed diverse expression profiles in different organs [12], as well as tissue- and stage-specific expression [5]. RALF-like genes also respond to nematode, drought [13], nitrosative, and oxidative stresses [11]. In particular, Mecchia et al. [13] reported that RALF4/19 peptides interact with LRX proteins to control pollen tube growth in Arabidopsis. Ge et al. [14] also reported that RALF4/19 peptide ligands enhance the strength of LLG2/3B–UPS/ANX interaction to influence the integrity of pollen tube. Moussu et al. [15] reported the crystal structure of the LRX-RALF cell wall complex and demonstrated that RALF peptides were activated as folded proteins, and RALFs can also instruct LRX cell wall modules and CrRKL1L receptors through structurally different binding modes to coordinate pollen tube integrity. ScRALF23 was reported as able to affect the pollen mitosis by functioning among the process from sporophyte to gametophyte signaling [16]. Leucine-rich repeat extension proteins can combine with RALF to modify cell wall expansion and bind to the transmembrane receptor FERONIA and regulate the cell growth [17]. Haruta et al. [18] reported that PtdRALF1 and PtdRALF2 were continuously and stably expressed in several organs, whereas PtdRALF2 showed low expression in the leaves of mature poplar trees, and PtdRALF1 was hardly affected by phytohormones, nutrient concentration pathogen elicitors, and decreasing pH caused by HCl, but the expression of PtdRALF2 could be inhibited strongly by MeJA after 5 h.

In the present study, we performed a genome-wide analysis to identify RALF-like genes of seven Rosaceae species (peach, apple, wild strawberry, cultivated strawberry, European pear, sweet cherry, and Japanese apricot). The analysis revealed that the seven Rosaceae species contain 124 RALF-like genes that were classified into four clades using phylogenetic analysis. Here, we selected 45 mature RALF genes from Rosaceae species, which include the RRXL cleavage site, the YISY motif, and four highly conserved cysteines in the C-terminus. Further, we analyzed the protein structure, duplication events, and evolutionary selections. Importantly, we combined cis-regulatory elements analysis, gene ontology (GO) annotation, EST data, RNA-seq analysis, and RT-qPCR results to have a clear knowledge of these genes in some development stages, tissues, and under diverse stresses. Specifically, we analyzed the coding capacity of these genes, and picked out 24 coding mature RALF genes, which can be further investigated. The research provides us with a comprehensive knowledge of RALF-like genes in seven Rosaceae species.
2. Materials and Methods

2.1. Data Sources

The peach (Prunus persica) genome (v2.0.a1) [22], apple (Malus × domestica) genome (HFTH1 Whole Genome v1.0) [23], wild strawberry (Fragaria vesca) genome (v4.0.a1) [24], cultivated strawberry (Fragaria × ananassa) genome (v1.0.a1) [25], European pear (Pyrus communis) genome (v1.0) [26], sweet cherry (Prunus avium) genome (v1.0.a1) [27], and related annotation files were downloaded from GDR database [28].

The genome and related annotation files of Japanese apricot (Prunus mume) (v1.0) [29] were retrieved from NCBI (https://www.ncbi.nlm.nih.gov/). The Arabidopsis RALF-like proteins were downloaded from the Phytozome database [30] using “RALF” as the keyword, and the RALF genes reported in another research were added [4]. The RNA-seq data for investigating the expression profiles of RALF-like genes were retrieved from European Nucleotide Archive (https://www.ebi.ac.uk/ena).

2.2. Identification and Sequence Analysis of RALF-Like Proteins

To identify the candidate members of the RALF-like proteins in seven Rosaceae species, the RALF domain (PF05498) was downloaded from the Pfam database [31], and HMMER v3.0 [32] was used to search the conserved domain sequences. Finally, these candidate sequences were submitted to Pfam [31] to confirm whether they had the RALF domain.

The predicted subcellular localization of candidate proteins was retrieved from Plant-mPLoc [33–35] (http://www.csbio.sjtu.edu.cn/bioinf/plant-multi/). Verified RALF-like protein sequences were submitted to the ExPASy proteomics service [36,37] to screen the essential information of proteins, and the signal peptide information was obtained from SignalP-5.0 [38].

2.3. Maximum Likelihood Phylogeny of RALF-Like Proteins

The protein sequences of the seven Rosaceae species and Arabidopsis thaliana were aligned using Muscle [39]. The model was calculated by using ModelFinder (VT+R6) [40]. Then, a phylogenetetic tree was inferred under maximum likelihood with IQ-TREE [40,41]. The resulting tree file was visualized with FigTree version 1.4.4 (http://tree.bio.ed.ac.uk/software/figtree/) and then submitted to iTOL [42] (https://itol.embl.de/) for modification.

For future understanding of the function among 51 mature RALF genes, we selected the RALF genes with four conserved cysteine residues, YISY motif, and RRXL cleavage site to align, and build the phylogenetic tree (Best-fit model: JTT+I+G4) following the methods mentioned above.

2.4. The Conserved Domain and Motif Analysis

RALF-like proteins of peach, apple, wild strawberry, cultivated strawberry, European pear, sweet cherry, Japanese apricot, and Arabidopsis were aligned using Jalview version 2 (muscle with defaults) [43] to analyze the conserved regions and verify the RALF-like proteins twice.

In this research, six previously identified motifs [9] were submitted to the MEME suite (http://meme-suite.org/) [44], and the FIMO (Find Individual Motif Occurrences) service was used to screen among RALF-like proteins.

2.5. The Coding Capacity of RALF-Like Genes in the Seven Rosaceae Species

All the RALF-like coding sequences (CDs) from the seven Rosaceae species were submitted to coding potential calculator (CPC) (Center for Bioinformatics, Peking University, China) (http://cpc.cbi.pku.edu.cn/programs/run_cpc.jsp) [45] to analyze their coding abilities.
2.6. Duplication Event Analysis and ka and ks Values among RALF-Like Genes

Potential homologous gene pairs were identified in each of the genomes of the seven Rosaceae species using BLASTP [46] (E < 1·10^{-5}, top 5 matches). The results of BLASTP and the arranged CFF files were then used as the input files of Mscanx [47] to analyze the duplication events. The RALF-like CDs sequences of *Arabidopsis* and seven Rosaceae species were translated to proteins in MEGA7 [48] and aligned using ClustalW, then we submitted the aligned CDs sequences to DnaSPv6.12.03 [49] to calculate the ka and ks values.

2.7. The Expression Analysis Using Expression Sequence Tag Data

The EST databases of strawberry, apple, and peach (only these three species’ EST data can be found) were downloaded from PlantGDB (http://www.plantgdb.org/) [50]. A local BLASTN (E-value < 1·10^{-5}, top 5) was performed against pear EST libraries to get the hits for each RALF-like gene.

2.8. Gene Ontology and cis-Regulatory Elements Analysis

All 163 RALF-like proteins were submitted to Blast2GO [51] to obtain the gene ontology annotation information, with all the options as default. The 2 kb region upstream of the transcriptional start site was used for cis-regulatory elements analysis with the Plant Care [52] online service (http://bioinformatics.psb.ugent.be/webtools/Plantcare/html/), and the result was depicted using TBtools [53].

2.9. The Expression Profiles of RALF-Like Genes in Strawberry and Apple from RNA-Sequencing

To analyze the wide expression profiles, RNA-seq data were used, and the FASTQ files were downloaded from European nucleotide archive (https://www.ebi.ac.uk/ena). Here, *Fragaria vesca, Fragaria × ananassa, and Malus × domestica* (only these three species in the present study can be found the available RNA-seq data to analyze the expression profiles about diverse tissues and stresses) were considered when investigating the RALF-like gene expression of diverse tissues, organs, developmental stages, and stresses. The study accessions from NCBI were SRP096282, SRP098567, ERP004230, SRP035308, SRP018410, ERP013896, SRP111905, SRP125281, SRP091754, and SRP139480 [54–61], which were involved in the development of flower tissues, fruit tissues, seed tissues, and fungal stresses. Ultimately, the FPKM (fragments per kilobase per million) [62] of RALF-like genes was extracted from the GTF files and submitted to TBtools [52] to draw the heatmaps with the log2 translation of FPKM.

2.10. Gene Validation Experiments by Quantitative Real-Time PCR (RT-qPCR)

*F. × ananassa* cv. Benihoppe plants cultured in MS (Murashige and Skoog) solid medium for four weeks to perform, and 0.2 mM NAA (naphthylacetic acid), 0.2 mM ABA (abscisic acid), 0.2 mM GA4 (gibberellin 4), 0.2 mM MeJA (methyl jasmonate), and 1.0 mM SA (salicylic acid) were sprayed on the tissue of cultured seedlings. NAA, ABA, and GA4 will influence the growth of plants, and MeJA and SA are related to the stresses of plants. Three plants were used for one treatment, and the leaves of strawberry seedlings were immediately flash frozen using liquid nitrogen and stored in a −80 °C freezer after treating for 12, 24, and 48 h.

Total RNA was isolated according to the instructions of the RNA extraction kit (Tiangen, Beijing, China), and RNA was reverse-transcribed into complementary DNA (cDNA) using the PrimeScript RT reagent kit (TaKaRa, Dalian, China) for quantitative real-time PCR (RT-qPCR). RT-qPCR was performed on an ABI 7300 Real-Time PCR System (Applied Biosystems, Foster City, CA, USA). The gene-specific primers were designed by the Beacon Designer 7 program (Premier Biosoft, Palo Alto CA, USA). *EF-1α* (XM_004307362) was used as an internal control to normalize the expression level of target genes. All primers are listed in Table 1. The cycling conditions were maintained as follows; 95 °C for 4 min, 40 cycles at 95 °C for 25 s, 60 °C for 20 s, and 73 °C for 43 s to create a melting curve. The reaction was performed in a 20 µL reaction mixture including a diluted cDNA sample as a template, a SYBR premix.
ExTaq (2×) (TaKaRa, Kyoto, Japan), and primers. The relative expression level of genes was calculated by the \(2^{-\Delta\Delta CT}\) method [63].

**Table 1.** Primers used to perform quantitative real-time PCR (RT-qPCR) for RALF-like genes.

| Symbol | Gene ID | Forward Sequence (5′ to 3′) | Reverse Sequence (5′ to 3′) |
|--------|---------|-----------------------------|----------------------------|
| Fa-RALF1 | snap_masked-Fvb4-2-processed-gene-95.31-mRNA-1 | CAACCTCCTCATTCTCCTCAC | TGCTGGTGGCTAAGATGCC |
| Fa-RALF2 | augustus_masked-Fvb4-2-processed-gene-134.10-mRNA-1 | TCCCTCCCCATTCTTCAATC | CCTCTTTCTAGTCGTTAGC |
| Fa-RALF5 | snap_masked-Fvb2-1-processed-gene-164.41-mRNA-1 | AGCGGTGAGCTTTGGTGCT | GCGGCAAGATGCTGGTTGAG |
| Fa-RALF8 | snap_masked-Fvb6-1-processed-gene-108.14-mRNA-1 | CATTGGACGCAATGCTCAAG | AATTACTCGAGTACCCGAG |
| Fa-RALF10 | augustus_masked-Fvb6-1-processed-gene-322.10-mRNA-1 | AGCTGGTGAGACCTCTCATC | CGGGTATAGGCGTTGAG |
| Fa-RALF12 | augustus_masked-Fvb3-3-processed-gene-31.4-mRNA-1 | ACTCTCTCTGGTCTCCACCC | GAGACTCTTGACGAGTCC |
| Fa-RALF13 | augustus_masked-Fvb3-3-processed-gene-31.5-mRNA-1 | CATCAGCTACGGTGCCCTAAT | CTGCAAGGCTCGATAGGT |
| Fa-RALF14 | augustus_masked-Fvb1-4-processed-gene-196.6-mRNA-1 | TCTTCTACGGTGCTCTCGAG | GCAGACTCTTGAGTTAGG |
| Fa-RALF18 | augustus_masked-Fvb6-3-processed-gene-378.1-mRNA-1 | TGATACCCACCAAGGGTT | GCTTAGGGGGTTGAG |
| Fa-RALF19 | augustus_masked-Fvb1-4-processed-gene-61.5-mRNA-1 | TGACTCTTTACACTCTACCCACTA | GCCGTGGGCGAAGGAGTCAGG |
| Fa-RALF24 | augustus_masked-Fvb1-3-processed-gene-90.5-mRNA-1 | GCTTCCGCTTGATCTCCGAC | GCGCTATAGGANCGTTGAG |
| Fa-RALF25 | snap_masked-Fvb6-4-processed-gene-50.20-mRNA-1 | TTCCGCTGCTGAGATGCCCTGA | ATACGATGCTGGCTGAG |
| Fa-RALF29 | augustus_masked-Fvb2-3-processed-gene-83.11-mRNA-1 | TAATCTTCTACCTTGAGTCGAGCT | TTTGAGTGGCATGAGTCTAG |
| Fa-RALF31 | snap_masked-Fvb4-3-processed-gene-153.15-mRNA-1 | GCAATGGGTCTTCAGTTCTG | GCAATGGGTCTTCAGTTCTG |
| FXARAL32 | maker-Fvb4-3-snap-masked-Fvb4-3-processed-gene-192.46-mRNA-1 | GCAATGGGTCTTCAGTTCTG | GCAATGGGTCTTCAGTTCTG |
| Fa-RALF33 | augustus_masked-Fvb1-2-processed-gene-103.4-mRNA-1 | ATGACTTCTACCTTGAGTCGAGCT | AATTACCTCGGCTCGCTTGA |
| Fa-RALF35 | snap_masked-Fvb3-2-processed-gene-47.50-mRNA-1 | GCAATGGGTCTTCAGTTCTG | GCAATGGGTCTTCAGTTCTG |
| Fa-RALF39 | snap_masked-Fvb4-4-processed-gene-106.36-mRNA-1 | TCTTCTACCTTGAGTCGAGCT | GCAATGGGTCTTCAGTTCTG |
| Fa-RALF40 | augustus_masked-Fvb4-4-processed-gene-145.10-mRNA-1 | AAGTGCGCGGACGGCAC | TCGTGTAATGACCTCCAGC |
| EF-1α | XM_004307362 | CATGCCGACACTGTTGGTTG | GACGGCTCAGAATCTACTAG |

3. Results

3.1. Identification, Coding Capacity, Classification, and Structural Features of RALF-Like Genes in Rosaceae Species

The genome-wide search for and verification of the seven Rosaceae species reference genomes indicated that there are 124 sequences with the RALF domain, with 41 from cultivated strawberry, 13 from wild strawberry, 20 from sweet cherry, 14 from peach, and 10 from European pear, in addition to 39 Arabidopsis RALF-like protein sequences that were retrieved from Phytozome [23] and previous research [4], among these 163 sequences, there are five common protein pairs (Pp-RALF1 and Pp-RALF2, Fv-RALF5 and Fa-RALF10, Fa-RALF7 and Fv-RALF6, Fa-RALF12 and Fv-RALF10, and At-RALF7 and At-RALF37).

The RALF-like proteins’ primary structure information is listed in Table S1. The molecular weight ranges from 7 to 121 kDa, the number of amino acids ranges from 64 to 747, and the minimum theoretical isoelectric point (pI) is 4.42 and the maximum is 10.31. In the seven Rosaceae species, there were 81% (101/124) RALF-like proteins with a pI higher than 7, and others lower than 7. Therefore, most are basic amino acids. There are ~86% (107/124) of these proteins which have a signal peptide, which is essential for secreted proteins. The majority of RALF-like proteins located extracellularly are also involved in other positions, such as cell membrane, cytoplasm, endoplasmic reticulum, mitochondrion, and the nucleus, which implies these proteins mainly function outside the cell.
The results of CPC [45] showed that Fa-RALF22 and Fa-RALF23 have no coding capacity, and 32 have a weak noncoding capacity, 58 have the weak coding ability, and 32 have a clear coding capacity (Table S1).

According to the phylogenetic analysis (Figure 1), these RALF-like proteins were divided into four clades (I–IV), and clade I was divided into nine subclades (A–G, S1, and S represents the single sequence) and clade III was divided into two subclades (A and B). The majority (~57% (71/124)) of RALF-like proteins in Rosaceae species belong to clade I, 24 RALF-like proteins to clade II, 26 to clade III, and three to clade IV. Clade I–D, clade I–F, clade II, and clade III–B contain all the Rosaceae species, clade I–A contains five Rosaceae species, and clade I–C and clade I–E contain six Rosaceae species. Clade I–G and clade III–A contain four Rosaceae species. Others contained one Rosaceae species, which implied that most RALF-like proteins identified in the present study have conserved functional structures.

Figure 1. The maximum-likelihood phylogenetic tree of RALF-like proteins among Arabidopsis and seven Rosaceae species.
Combining the alignment from Jalview Version 2 (muscle with defaults) [43] (Figure S1) and the analysis of motifs retrieved from MEME [44] (Figure S2), we found some structural features among clades or subclades according to the alignment and motif results.

A majority of the RALF-like proteins have four conserved cysteines, except 22 sequences from the Rosaceae species (sequences marked with green background in Table S1). There are 51 sequences which have the YISY motif, the RRXL cleavage site, and four conserved cysteine residues the typical features of mature RALF genes and 45 from the Rosaceae species. All the sequences in clade I-F and most sequences of clade I–D, clade I–E, and clade II (seven from Rosaceae species in Clade I–D, eight from Rosaceae species in Clade I–E, 18 from Rosaceae species in Clade II) have typical features. These mature RALF-like genes may be more meaningful for future research, whereas other sequences have none or one of these features. Searching these sequences, some YISY motifs may have some mutations, such as YISP, YIST, NISY, YINY, YIGY, YVGY, FIGY, YIAY, and RRXL, which may mutate to RRVM.

These protein sequences were searched for the presence of six motifs (Figure 2), and motif1 was found to exist in 141 sequences and 110 sequences from Rosaceae species. The second and third most common motifs are motif3 (84 sequences with 71 from Rosaceae species) and motif2 (75 sequences with 63 from Rosaceae species). Motif1 distributed widely among all the clades.

As depicted in the phylogenetic tree of 51 mature RALF genes (Figure 3), we selected 10 homologous genes pairs (Md-RALF4 and Pc-RALF2; Pp-RALF5, Pm-RALF5, and Pa-RALF10; Fa-RALF5, Fa-RALF29, and Fa-RALF35; Pm-RALF7 and Pp-RALF6; Fa-RALF2, Fv-RALF2, and Fa-RALF40; Md-RALF14 and Md-RALF5; Pp-RALF12 and Pa-RALF3; Pm-RALF6, Pp-RALF8, and Pa-RALF2; Pc-RALF9 and Md-RALF15; Fa-RALF19, Fv-RALF1, Fa-RALF33, Fa-RALF24, and Fa-RALF14) to analyze.
Figure 3. The maximum-likelihood phylogenetic tree of 51 mature RALF genes examined in the present study. RALF genes from *Arabidopsis* marked with the same color and background, and the same gene pair with the same mark.

### 3.2. Chromosome Location, Duplication Events, and Divergence Rates

As depicted (Figure S3), there is an expanding distribution among Rosaceae species according to the chromosomal location results. In wild strawberry, there are six chromosomes, except Fvb7, in which 13 RALF-like genes were located. In cultivated strawberry, there are 21 chromosomes which have RALF-like gene sites. The following numbers of RALF-like genes were identified in specific chromosomal locations: 10 in apple, 8 in sweet cherry, 6 in peach, and 9 in European pear. In Japanese apricot, which has eight chromosomes, Pltd and Unplaced Scaffold regions, RALF-like gene sites were found on six chromosomes, whereas *Pm-RALF9* is located in Un region. As shown, most of the RALF-like genes are distributed around the telomere, and others around the centromere.

Genes can be duplicated by numerous mechanisms, including through whole genome duplication or segmental (WGDs), dispersed gene duplication (DD), tandem duplication (TD), proximal duplication (PD), and singleton (single-copy, SC) events. Among the 124 sequences, 45.2% (56/124) involved WGD events, 33.1% (41/124) dispersed gene duplication events, 9.7% (12/124) tandem events, 5.6% (8/124) proximal duplication events, and 6.4% (7/124) singleton events (Table S1). Only four couples of tandem duplication genes, *Fv-RALF9* and *Fv-RALF10*, *Md-RALF7* and *Md-RALF8*, and *Pm-RALF2*, and *Pm-RALF8*, *Pp-RALF11*, and *Pp-RALF14* were found (Figure S3).

The ka and ks were calculated by DnaSP [49] using all the RALF-like CDs sequences identified. As is calculated, the ka range is 0 to 4.8, ks range is 0 to 4.1, and the ka/ks is 0 to 22.1. The data distribution map was displayed through box plots which were constructed via Origin (OriginLab...
Corporation, Northampton, MA, USA) and the outlines were eliminated automatically (Figure 4). The ka and ks values were stably distributed (Figure 4a,b), which implies there were stable evolutionary counts and mutation frequencies between and among these species. The ka/ks orients the evolution direction. As depicted (Figure 4c), all the average mean values of genes pairs were lower than one, which implies that purifying the selection may promote evolution. The values of ka, ks are listed in Table S2.

![Figure 4](image-url)

**Figure 4.** The ka, ks, and ka/ks value distributions. (a) ka values; (b) ks values; (c) ka/ks values.

### 3.3. Gene Ontology and Cis-Regulatory Elements Analysis

Among 124 sequences (RALF-like genes in *Arabidopsis* not included), 17 GO terms were found in 51 RALF-like genes. Approximately 85% (138/163) of RALF-like genes in the present study (conclude the sequences from *Arabidopsis thaliana*) were enriched in the GO category of cellular component (CC). The gene number of each GO term and functional class were counted and depicted in Figure 5. The GO categories of the biological process (BP) and molecular function (MF) accounted for 74% (120/163) and 33% (54/163), respectively. The calcium-mediated signaling (GO: 0019722), plasmodesma (GO: 0009506), protein phosphorylation (GO: 0006468), hormone activity (GO: 0005179), cell–cell signaling (GO: 0007267), and the integral component of the membrane (GO: 0016021) are major GO categories.

There are 21 cis-regulatory elements in total, such as abscisic acid (ABA), anaerobic (Ana), auxin (Auxin), cell cycle (Cyc), circadian control (Cir), defense and stress (DS), differentiation of the palisade mesophyll cells (DPMC), drought (Drought), endosperm specific expression (Endosperm), flavonoid biosynthetic (Fla), gibberellin (Gibberellin), light (Light), low temperature-related (LT), methyl jasmonate (MeJA), meristem-specific expression (Meristem), phytochrome downregulation-related (Phy), (Root), salicylic acid (SA), seed-specific expression (Seed), wound-related (Wound), and zein metabolism-related (Zein), which are involved in stress and tissue-specific expression (Figure S4 and Table S1). The major elements contain light-related elements, with 998 counts, followed by anaerobic...
(256), abscisic acid (195), drought (112), and gibberellin (95). The quantity of every RALF-like gene ranges from 0 to 12 (Table S1).

Figure 5. The gene ontology (GO) term distribution and functional clustering among seven Rosaceae species.

3.4. The Expression Information from Expression Sequence Tag Data

As listed in Table S3, according to the results from ESTs, we reported that twelve plant tissues (flower, bud, fruit, fruit mesocarp, fruit core, fruit cortex, phloem, xylem, leaf, shoot internodes, young root, and young shoot), six abiotic stresses (cold, water, drought, heat, and salt), and two biotic stresses (*Venturia inaequalis* and *Choristoneura rosaceana*) are involved (Table 2).

| Plant Tissues and Stresses | Genes Retrieved from Expression Sequence Tag (EST) Data |
|---------------------------|---------------------------------------------------------|
| Sault                     | Fo-RALF10, Fo-RALF11, Fo-RALF12                        |
| Drought                   | Fo-RALF10                                              |
| Heat                      | Fo-RALF11, Fo-RALF12                                    |
| Red fruit/Fruit           | Fa-RALF5, Fa-RALF9, Fa-RALF35, Mf-RALF2, Mf-RALF3, Mf-RALF5, Mf-RALF6, Mf-RALF7, Mf-RALF11, Mf-RALF12, Mf-RALF13, Mf-RALF14, Mf-RALF15, Mf-RALF19, Mf-RALF20, Pp-RALF12, Pp-RALF13 |
| Flower                    | Md-RALF11, Md-RALF12, Md-RALF13, Md-RALF14, Md-RALF15, Md-RALF19, Md-RALF20 |
| Bud                       | Md-RALF2, Md-RALF3, Md-RALF7, Md-RALF11, Md-RALF13, Md-RALF15, Md-RALF19, Md-RALF20 |
| Young root                | Md-RALF3, Md-RALF7, Md-RALF11                          |
| Xylem                     | Md-RALF6, Md-RALF7, Md-RALF13, Md-RALF19, Md-RALF20    |
| Young shoot               | Md-RALF2, Md-RALF7, Md-RALF13, Md-RALF15, Md-RALF19, Md-RALF20 |
| Leaf                      | Md-RALF12, Md-RALF13, Md-RALF15, Md-RALF19, Md-RALF20, Pp-RALF5 |
| Shoot internodes          | Md-RALF13, Md-RALF19                                   |
| Fruit cortex              | Md-RALF13, Md-RALF19                                   |
| Fruit core                | Md-RALF13, Md-RALF19                                   |
| Phloem                    | Md-RALF20                                             |
| Mesocarp                  | Pp-RALF6, Pp-RALF8, Pp-RALF13                          |
| *Venturia inaequalis*     | Md-RALF5, Md-RALF14                                    |
| *Choristoneura rosaceana* | Md-RALF20                                             |
3.5. The RNA- Sequencing Expression of RALF-Like Genes in Strawberry and Apple

The downloaded RNA-seq data were used to analyze the expression profiles of RALF-like genes and the basic information of these data listed in Table S4. In wild strawberry, Fv-RALF1, Fv-RALF2, Fv-RALF3, Fv-RALF5, and Fv-RALF12 were more highly expressed than other genes (Figure S5a). Fv-RALF1 shows a lower expression level in the ghost development stages and wall 4, wall 5 stages, which means it may affect the development of ghost and wall. Fv-RALF2 and Fv-RALF3 all have different expression profiles in the cortex compared with other tissues, which implies the potential functions on cortex development. Fv-RALF5 and Fv-RALF12 both have lower expression in embryo than other tissues and may influence the development of the embryo. Fv-RALF4, Fv-RALF6, Fv-RALF7, Fv-RALF8, Fv-RALF9, Fv-RALF10, and Fv-RALF11 all have higher expression levels in specific tissues (Figure S5d). Fv-RALF4, Fv-RALF9, and Fa-RALF10 are more highly expressed pollen-specific genes. These genes may play an important role in the development of pollen. The receptacle fruit at green stage (15 days post-anthesis) and white (turning) stage (22 days post-anthesis) with achenes removed were used to analyze the RALF-like gene expression profile (Figure S5b). As depicted, Fv-RALF1, Fv-RALF5, Fv-RALF12, and Fv-RALF13 all show higher expression levels in some ways. To retrieve the expression profile of RALF-like genes in achene and receptacle fruit, the RNA-seq data of one red-fruited and two natural white-fruited strawberry varieties in two tissues and three ripening stages were used (Figure S5e). Fv-RALF1, Fv-RALF2, Fv-RALF5, Fv-RALF10, and Fv-RALF12 were more highly expressed in some cultivars and tissues, and showed variable expression profiles, which means they may have an effect on the development of achene and receptacle fruit. The powdery mildew-infected leaf sample from Hawaii was used to analyze the expression level under fungal stress (Figure S5c). Fv-RALF12 always showed a high expression levels, though this does not exclude it from having an essential function in resistance from infection by powdery mildew. However, Fv-RALF1 is more highly expressed one day after infection, which may represent a response to infection.

In cultivated strawberry, the RNA-seq data of the development achene and receptacle fruit were also analyzed (Figure S6a). Fa-RALF1, Fa-RALF18, and Fa-RALF39 are more highly expressed at the white receptacle stage, whereas, Fa-RALF14, Fa-RALF24, and Fa-RALF33 exhibit a higher level of expression in leaf than other tissues, and Fa-RALF29 and Fa-RALF35 may be highly expressed root-specific genes. In diverse development stages of cultivated strawberry (Figure S6b), we reported that Fa-RALF1, Fa-RALF10, and Fa-RALF33 were all more highly expressed at the turning stage compared to other stages. Fa-RALF6, Fa-RALF17, Fa-RALF26, and Fa-RALF32 were more highly expressed at the green fruit stage.

According to the RNA-seq data of apple, we reported that Md-RALF7 and Md-RALF8 may have an effect on the development of flower (Figure S7a). Md-RALF3, Md-RALF4, Md-RALF13, and Md-RALF16 are more highly expressed in root tip (Figure S7b). Md-RALF3, Md-RALF4, and Md-RALF13 were more highly expressed than other isotypes in during infection by *Alternaria alternata* (Figure S7c). Md-RALF3 responds to infection after 12 h. Md-RALF4 responds to the infection after 18 h. The expression level of Md-RALF13 remained high until 72 h, in which it began to decline after.

3.6. RT-qPCR Analysis of Nineteen RALF-Like Genes in Cultivated Strawberry Related to Hormones

Nineteen RALF-like genes of cultivated strawberry under different hormone stresses were analyzed by RT-qPCR (Figure 6). Herein, we reported that Fa-RALF1 (GA), Fa-RALF2 (ABA), Fa-RALF5 (NAA), Fa-RALF8 (NAA), Fa-RALF10 (SA), Fa-RALF12 (GA), Fa-RALF13 (ABA), Fa-RALF14 (ABA and SA), Fa-RALF18 (MeJA), Fa-RALF31 (ABA and SA), Fa-RALF32 (ABA), Fa-RALF33 (MeJA), Fa-RALF35 (MeJA), and Fa-RALF39 (ABA) have an obvious response to one or two hormones, whereas Fa-RALF19, Fa-RALF24, Fa-RALF25, Fa-RALF29, and Fa-RALF40 evidently respond to all the stresses, which implies a role for RALF genes in hormone response. Here, fourteen genes (Fa-RALF1, Fa-RALF2, Fa-RALF5, Fa-RALF10, Fa-RALF12, Fa-RALF13, Fa-RALF14, Fa-RALF19, Fa-RALF24, Fa-RALF29, Fa-RALF32, Fa-RALF35, Fa-RALF39, and Fa-RALF40) are mature RALF genes and seven genes (Fa-RALF5, Fa-RALF10, Fa-RALF14, Fa-RALF19, Fa-RALF24, Fa-RALF29, and Fa-RALF35) have coding capacity.
and seven genes (Fa-RALF5, Fa-RALF10, Fa-RALF14, Fa-RALF19, Fa-RALF24, Fa-RALF29, and Fa-RALF35) have coding capacity.

Figure 6. Relative expression profiles of 19 RALF-like genes from Fragaria × ananassa under different hormone treatments (a–s). Error bars represent standard deviation (SD). Columns with stars marked the significantly different. The numbers of stars stand for the difference extent. The more stars, the higher the significant difference; three stars represents an extreme difference. Here, all the treatments are compared with “control”, and * represents $p \leq 0.05$, ** represents $p \leq 0.01$, *** represents $p \leq 0.001$. MeJA-12,-24,-48: the leaves treated by MeJA after 12, 24, and 48 h; ABA-12,-24,-48: the leaves treated by ABA after 12, 24, and 48 h; NAA-12,-24,-48: the leaves treated by NAA after 12, 24, and 48 h; GA-12,-24,-48: the leaves treated by GA after 12, 24, and 48 h; SA-12,-24,-48: the leaves treated by SA after 12, 24, and 48 h.
4. Discussion

4.1. Expression Profile of RALF-Like Genes in Seven Rosaceae Species

As reported, RALF genes are not only involved in several biotic and abiotic stresses but also influence the growth and development of plants. Here, we found that the expression profiles are mostly similar with the researchers reported, such as the drought response [13], root growth [64], hormone response [18], fungi response [18–20], cell wall development [15,17], and the flower development (especially the development of pollen tube) [14,16]. In the present study, we combined the results of RNA-seq, EST data, cis-regulatory elements analysis, and RT-qPCR to identify several genes which have identical expression profiles in two or more analyses, which may give some instructions of the research of RALF-like genes among Rosaceae species.

_Fv-RALF10_ exhibits drought responses according to the analysis of EST data and cis-regulatory elements. RNA-seq consistent with cis-regulatory element analysis involved in _Fv-RALF1_, _Fv-RALF2_, _Fv-RALF12_, _Fv-RALF15_ (seed, embryo, and endosperm), which have no research reported, may be the special findings in our study among Rosaceae species. The RALF genes most reported were the role they play in root growth and pollen tube development [14,16]. Here, the comparison between RNA-seq and EST data reveal that _Md-RALF3_ may influence on the growth of young root, and _Md-RALF7_ and _Md-RALF8_ may affect the development of flowers. As reported that MeJA may influence the expression of RALF genes [18]; here, we tried more hormones, which may influence the tissue development and stresses response, the results from RT-qPCR and cis-regulatory elements showed that _Fa-RALF1_, _Fa-RALF2_, _Fa-RALF12_, _Fa-RALF13_, _Fa-RALF14_, _Fa-RALF18_, _Fa-RALF19_, _Fa-RALF24_, _Fa-RALF25_, _Fa-RALF29_, _Fa-RALF31_, _Fa-RALF33_, and _Fa-RALF40_ have identical expression profiles, in other words, they may have influence on the growth, development and stresses response of Rosaceae species.

In strawberry, the _FaRALF-33-like_ gene, which is homologous to _FvRALF-33-like_ (GenBank accession No.: XM_011460413.1) and _AtRALF33_ (named _At-RALF1_ here), responded to infection by three different fungi [20]. Local BLASTP [46] was used to search the similar genes among our 163 sequences, using _FvRALF-33-like_ and _AtRALF33_ as the query sequences. We revealed that _Fa-RALF1_ (GenBank accession No.: XP_011458715) and _Fv-RALF5_ are the best-match sequences. As shown (Figure 7), except for _Fa-RALF1_, other sequences all possess the typical features and the integral YISY, YYNC, RCR motifs.

![Figure 7](Image)

_Fv-RALF5_ belongs to clade II, and among the clade, there are seventeen other mature RALF genes ( _Pc-RAL2_, _Md-RALF4_, _Pc-RALF6_, _Md-RALF11_, _Pa-RALF10_, _Pm-RALF5_, _Pp-RALF5_, _Fa-RALF35_, _Fa-RALF29_, _Fa-RALF5_, _Pp-RALF6_, _Pm-RALF7_, _Pa-RALF1_, _Pc-RALF7_, _Md-RALF7_, _Md-RALF12_, and _Fa-RALF10_ ) from Rosaceae species, and no the RCR motif was found in _Pc-RALF6_. Therefore, the 16 _RALF-like_ genes, except for _Pc-RALF6_, may have similar functions as _Fv-RALF5_, _At-RALF1_, and _FaRALF-33-like_ genes. As depicted in the phylogenetic tree, we reported that mature RALF genes in clade II, except _Pc-RALF6_, _At-RALF4_, and _At-RALF1_, were similarly divided within the same clade, which is consistent with previous reports. Stegmann et al. [65] already investigated that _RALF33_ ( _AT4G15800_, named _At-RALF1_ here), _RALF23_ ( _AT3G16570_, named _At-RALF4_ here), and _RALF34_ ( _AT5G67070_, named _At-RALF14_ here) negatively regulate immunity, whereas _RALF32_ ( _AT4G14010_, named _At-RALF5_ here) did not. _Fv-RALF5_ clustered with _Fa-RALF10_, which implies that they have a similar function. They may also respond to the fungal infection. Furthermore, according to the RNA-seq results and cis-regulatory element analysis, we reveal that they may have
influence on the development of receptacle and be involved in some hormone stresses (ABA, MeJA, etc.) and low temperature stress. Compared with the expression profiles of Arabidopsis in Campbell and Turner [4], we found that the clade IV (clade I-A and clade III-A concluded here) are not consistent with our research mostly. The expression profiles are involved in hormone, fruit development, flower development, leaf growth, salt stress, powdery mildew response, and achene development.

Ten homologous genes pairs were selected from the phylogenetic trees of mature peptides (Md-RALF4 and Pc-RALF2; Pp-RALF5, Pm-RALF5, and Pa-RALF10; Fa-RALF5, Fa-RALF29, and Fa-RALF35; Pm-RALF7 and Pp-RALF6; Fa-RALF2, Fv-RALF2, and RALF40; Md-RALF14 and Md-RALF5; Pp-RALF12 and Pa-RALF3; Pm-RALF6, Pp-RALF8, and Pa-RALF2; Pc-RALF9 and Md-RALF15; Fa-RALF19, Fv-RALF1, Fa-RALF33, Fa-RALF24, and Fa-RALF14), and except for Fv-RALF5 and Fa-RALF10, which were analyzed above, we examined whether these gene pairs had similar expression profiles using RNA-seq, EST, and RT-qPCR combined with cis-regulatory element analysis. The integrative analysis of RT-qPCR and cis-regulatory element implied that the Fa-RALF5, Fa-RALF29, and Fa-RALF35 may respond to auxin stress, Fa-RALF2, Fv-RALF2, and Fa-RALF40 to ABA, and Fa-RALF19, Fv-RALF1, Fa-RALF33, Fa-RALF24, and Fa-RALF14 to auxin and ABA. Based on the RNA-seq data, Fa-RALF2, Fv-RALF2, and Fa-RALF40 may affect the development of cortex, achene, and receptacle fruit; Fa-RALF19, Fv-RALF1, Fa-RALF33, Fa-RALF24, and Fa-RALF14 were mainly expressed in leaf and may also be expressed in achene, receptacle fruit, ghost, wall and may all be involved in response to fungal infection. As for the EST data, we reported that Pp-RALF5, Pm-RALF5, and Pa-RALF10 are expressed in leaf; Fa-RALF5, Fa-RALF29, and Fa-RALF35 in red fruit; Pm-RALF7 and Pp-RALF6 in mesocarp; Pp-RALF12 and Pa-RALF3 in fruit; Pm-RALF6, Pp-RALF8, and Pa-RALF2 in mesocarp; Md-RALF14 and Md-RALF5 in fruit and flower; and Pc-RALF9 and Md-RALF15 in fruit, flower, young shoots, and leaf. Notably, Md-RALF14 and Md-RALF5 expression is also altered upon fungal infection.

The genes mentioned above are essential to the growth, development, and stress response. In particular, the mature RALF genes were already found to exhibit obvious expression differences. In our research, there are 24 mature RALF genes with coding capacity (Fv-RALF5, Fa-RALF1, Fa-RALF19, Fa-RALF33, Fa-RALF14, Fa-RALF24, Md-RALF2, Md-RALF4, Md-RALF15, Md-RALF11, Md-RALF12, Fa-RALF35, Pa-RALF10, Pc-RALF1, Pa-RALF2, Pc-RALF6, Pc-RALF9, Pp-RALF5, Pp-RALF6, Pm-RALF7, Pp-RALF8, Pm-RALF6, and Pm-RALF5). As mentioned above, these 24 coding mature RALF genes are widely expressed, and 18 genes (Fv-RALF1, Fa-RALF14, Fa-RALF19, Fa-RALF24, Fa-RALF33, Fa-RALF35, Md-RALF2, Md-RALF11, Md-RALF12, Md-RALF15, Pa-RALF10, Pp-RALF9, Pm-RALF5, Pm-RALF6, Pm-RALF7, Pp-RALF5, Pp-RALF6, and Pp-RALF8) may be involved in the development of tissues and organs, which include 12 tissues or organs. In particular, seven genes have a tissue- or organ-specific expression profile. For example, Pa-RALF10, Pp-RALF5, and Pm-RALF5 may be leaf-specific genes, whereas Pm-RALF6, Pm-RALF7, Pp-RALF6, and Pp-RALF8 showed mesocarp-specific expression. Meanwhile, five genes (Fa-RALF35, Fa-RALF19, Fa-RALF33, Fa-RALF14, and Fa-RALF24) may be involved in hormone response. Notably, there are three genes (Fv-RALF5, Fa-RALF33, and Md-RALF14) related to fungal infection.

The GO terms give a clear function cluster of RALF-like genes, which involved in the pollen tube development, root development, negative regulation of growth, cell wall growth, hormone activaty, brassinosteroid-mediated signaling pathway, and so on. According to the classification, we found they are very consistent with previous researches about the functions of RALF genes [1,14–18,64]. Fungal infection response and pollen tube development are important for Rosaceae species, and the RALF-like gene expression in seed and fruit among Rosaceae species specially.

4.2. Classification, Duplication Events and Evolution of RALF-Like Proteins in Seven Rosaceae Species

In several researches, the classification and evolution of RALF-like proteins were analyzed. Cao et al. [9] characterized RALF proteins in Arabidopsis, rice, poplar, and maize, which were divided into ten groups, and elucidated that tandem duplication played a dominant role in the expansion of
RALF gene family and RALF-like proteins mainly through purifying selection, which is consistent with the conclusions of our research. However, we found that the dominant duplication event was WGD or segmental, not tandem. Sharma et al. [11] divided RALF-like proteins of Arabidopsis, rice, maize, and soybean into seven groups, and the RALF-like genes in 51 plant species also diverged into four major clades [4]. In our work, four clades were also divided among RALF-like genes between Arabidopsis and seven Rosaceae species.

The number of RALF-like genes was not consistent with the previous. Cao et al. [9] identified 33 RALF-like genes in Arabidopsis; Sharma et al. [11] identified 39; Campbell and Turner [4] identified 37; and, in our research, we retrieved 39 from analysis. Due to the reference genomes we selected not being identical with Campbell and Turner [4], we identified 13, and not 9, RALF-like genes in Fragaria vesca, 20, and not 33, in Malus × domestica, and 14, and not 13, in Prunus persica. Meanwhile, Fragaria × ananassa and not Arabidopsis had the most RALF-like genes (41). The classification in our study among RALF-like genes in Arabidopsis largely confirmed previous reports.

The alignment of RALF-like genes was implemented using Jalview Version 2 (muscle with defaults) [43], and the residue conserved within the mature peptide region of the four major clades was demonstrated using WebLogo 3 [66]. As depicted in Figure S8 compared with the analysis of Campbell and Turner [4], clade I–C, clade I–G, clade II, and clade IV were similar to clade I and II (A), clade I–D, clade I–E, clade I–F to clade III, clade I–A to clade IV, clade III–B to clade II (B), and clade III–A to clade IV, whereas clade I–B did not share any similarity to other clades and has none of the features of RALF genes. Campbell and Turner [4] reported that the genes in clade IV were not mature RALF genes. They also reported the RALF-like genes in clade I–A and clade III–A to be RNA-like genes, which is consistent with our research.

5. Conclusions

This study provides extensive information on the RALF-like genes of seven Rosaceae species, and involved, including their identification, an analysis of their coding capacity, evolutionary relationships, duplication events, motifs, domains, GO annotation, cis-regulatory elements, and EST data, as well as use of RNA-seq and RT-qPCR to characterize expression profiles. These RALF-like genes are divided into four clades, and purifying selection plays a main role in the evolutionary process. In particular, the 24 RALF genes coding for mature proteins among the Rosaceae species deserve to be further researched in regard to hormone response, fungal infection, growth, and development.

Supplementary Materials: The following are available online at http://www.mdpi.com/2073-4425/11/2/174/s1, Table S1: The basic information of RALF-like genes among Arabidopsis thaliana and seven Rosaceae species; Table S2: The ka, ks and ka/ks values among Arabidopsis and seven Rosaceae species; Table S3: The EST data analysis results; Table S4: The RNA-seq datasets’ information; Figure S1: The conserved protein structures analyzed in the research. The conserved alignment results from Jalview. The blue dotted line marks the YISY motif, the single arrow down represents the cysteine, the green line marks the RRXL cleavage site, and the shear shows the cutting site; Figure S2. The motif location and arrange from MEME; Figure S3: The chromosome location and tandem events among Rosaceae species. (a) Fragaria vesca; (b) Fragaria × ananassa; (c) Malus × domestica; (d) Prunus avium; (e) Prunus mume; (f) Prunus persica; (g) Pyrus communis; Figure S4: Distribution of identified cis-regulatory elements among the seven Rosaceae species. Cis-regulatory are indicated by respective colors; Figure S5: The heatmap of Fragaria vesca from diverse RNA-seq data. (a) Tissue- and stage-specific transcriptome during reproductive development of strawberry. The study accession ID is SRP018410; (b) the RNA-seq data from wild strawberry receptacle fruit (F. vesca) at green stage (15 days post-anthesis) and white (turning) stage (22 days post-anthesis) with removed achenes. The study accession ID is SRP096282. Yellow Wonder 5AF7 and Rügen (F7–4) were used. YW-15d: the fruit of Yellow Wonder 5AF7 at green stage (15 days post-anthesis); YW-22d: the fruit of Yellow Wonder 5AF7 at white (turning) stage (22 days post-anthesis); RG-15d: the fruit of Rügen (F7–4) at green stage (15 days post-anthesis); RG-22d: the fruit of Rügen (F7–4) at white (turning) stage (22 days post-anthesis); (f) Fragaria vesca ssp. vesca accession Hawaii 4 at 1 and 8 d after powdery mildew infection. 1DAF: 1 d after powdery mildew infection; 8DAF: 8 d after powdery mildew infection. The study accession ID is ERP004230. (d) Comprehensive transcriptomic dataset for flower development in woodland strawberry Fragaria vesca. The study accession ID is SRP05308; (e) comparative transcriptome analyses of one red-fruited and two natural white-fruited strawberry varieties in two tissues and three ripening stages. Hawaii_10GA: the green achenes approximately 10 d post-anthesis from Hawaii; Hawaii_10GR: the green receptacles approximately 10 d post-anthesis from Hawaii; Hawaii_25IA: the achenes at intermediate stage approximately 25 d post-anthesis from Hawaii; Hawaii_25IR: the receptacles at intermediate...
stage approximately 25 d post-anthesis from Hawaii; **Hawaii_35RA**: the achenes at ripe stage approximately 35 d post-anthesis from Hawaii; **Hawaii_35RR**: the receptacle at ripe stage approximately 35 d post-anthesis from Hawaii; **YW_10GA**: the green achenes approximately 10 d post-anthesis from Yellow Wonder; **YW_10GR**: the green receptacles approximately 10 d post-anthesis from Yellow Wonder; **YW_25IA**: the achenes at intermediate stage approximately 25 d post-anthesis from Yellow Wonder; **YW_25IR**: the receptacles at intermediate stage approximately 25 d post-anthesis from Yellow Wonder; **YW_35RA**: the achenes at ripe stage approximately 35 d post-anthesis from Reine des Valleeas; **YW_35RR**: the green achenes approximately 10 d post-anthesis from Reine des Valleeas; **RV_10GA**: the green achenes approximately 10 d post-anthesis from Reine des Valleeas; **RV_25IA**: the achenes at intermediate stage approximately 25 d post-anthesis from Reine des Valleeas; **RV_25IR**: the receptacles at intermediate stage approximately 25 d post-anthesis from Reine des Valleeas; **RV_35RA**: the achenes at ripe stage approximately 35 d post-anthesis from Reine des Valleeas; **RV_35RR**: the receptacle at ripe stage approximately 35 d post-anthesis from Reine des Valleeas. The accession ID SRP098567; Figure S6: The heatmap of Fragaria × ananassa from RNA-seq databases. (a) The heatmap of different development of strawberry achenes and receptacle with the leaf and root as controls. Four stages, the green fruit stage, white fruit stage, turning stage, and red fruit stage included, and every stage contained two tissues, receptacle, and achene. Leaf and root were used to as the control tissues. The study accession ID is ERP013896; (b) the heatmap of strawberry fruit removed achenes from diverse development stages. **GF**: green fruit; **WF**: white fruit; **TS**: turning stage; **RF**: red fruit. The study accession ID is SRP111905; Figure S7: The Heatmap of Malus × domestica from different RNA-seq data. (a) Different tissues expression profile in different tissues of Malus × domestica Golden Delicious. The study accession ID is SRP125281; (b) the heatmap of diverse tissues. **SP**: stem phloem; **ST**: shoot tip; **RT**: root tip; **YF**: young fruit. The study accession ID is SRP139480; (c) the heatmap of Malus × domestica Starking Delicious at different time infected by Alternaria alternata. 12h: infected by Alternaria alternata after 12 h; 18h: infected by Alternaria alternata after 18 h; 36h: infected by Alternaria alternata after 36 h; 72h: infected by Alternaria alternata after 72 h. The study accession ID is SRP091754; Figure S8: Divergence of RALF proteins according to the phylogenetic subclades. WebLogo 3 plots to demonstrate residue conservation within the mature peptide region of these subclade. The left of the plot marks the classification in the present study, and right marks the clustering in Campbell and Turner’s (2017) report.

**Author Contributions:** Conceptualization, H.Z. and Y.Q.; methodology, H.Z. and Y.Q.; formal analysis, H.Z., X.J., Y.C., Y.X., and Z.L.; investigation, H.Z., X.J., Y.C., Y.X., and Z.L.; writing—original draft preparation, H.Z.; writing—review and editing, Y.Q. All authors have read and agreed to the published version of the manuscript.

**Funding:** This research was funded by National Natural Science Foundation of China, grant number 31872056.

**Conflicts of Interest:** The authors declare no conflicts of interests. The funders had no role in the design of the study; in the collection, analyses, or interpretation of data; in the writing of the manuscript, or in the decision to publish the results.

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