Comparative Analysis of Complete Chloroplast Genomes of Anemoclema, Anemone, Pulsatilla, and Hepatica Revealing Structural Variations Among Genera in Tribe Anemoneae (Ranunculaceae)

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Structural rearrangements of Anemone species' chloroplast genome has been reported based on genetic mapping of restriction sites but has never been confirmed by genomic studies. We used a next-generation sequencing method to characterize the complete chloroplast genomes of five species in the tribe Anemoneae. Plastid genomes were assembled using \textit{de novo} assembling methods combined with conventional Sanger sequencing to fill the gaps. The gene order of the chloroplast genomes of tribe Anemoneae was compared with that of other Ranunculaceae species. Multiple inversions and transpositions were detected in tribe Anemoneae. Anemoclema, Anemone, Hepatica, and Pulsatilla shared the same gene order, which contained three inversions in the large single copy region (LSC) compared to other Ranunculaceae genera. Archiclematis, Clematis, and Naravelia shared the same gene order containing two inversions and one transposition in LSC. A roughly 4.4 kb expansion region in inverted repeat (IR) regions was detected in tribe Anemoneae, suggesting that this expansion event may be a synapomorphy for this group. Plastome phylogenomic analyses using parsimony and a Bayesian method with implementation of partitioned models generated a well resolved phylogeny of Ranunculaceae. These results suggest that evaluation of chloroplast genomes may result in improved resolution of family phylogenies. Samples of Anemone, Hepatica, and Pulsatilla were tested to form paraphyletic grades within tribe Anemoneae. Anemoclema was a sister clade to Clematis. Structural variation of the plastid genome within tribe Anemoneae provided strong phylogenetic information for Ranunculaceae.

Keywords: chloroplast genome, inversion, IR expansion, phylogenomics, Ranunculaceae, transposition, Tribe Anemoneae
INTRODUCTION

Comparative analysis of chloroplast genomes can provide valuable information for phylogeny reconstruction and resolution of complex evolutionary relationships (Shaw et al., 2007; Mardanov et al., 2008; Moore et al., 2010; Park et al., 2017; Sun et al., 2017). In angiosperms, plastid genome gene number and order is conserved (Wolf et al., 1987). This is because chloroplast sequences evolve at approximately half the speed of nuclear regions (Jansen et al., 2005; Walker et al., 2014). However, sequence rearrangements in chloroplast genomes have been reported from various kinds of plants (Doyle et al., 1996; Tanghatsornruang et al., 2011; Walker et al., 2015; Sun et al., 2017). These rearrangements include large inversions in large single copy region (LSC), and inverted repeats region (IR) expansions or contractions into single copy regions with inversions (Palmer et al., 1987; Tanghatsornruang et al., 2009). Intramolecular recombination may be the reason for large inversions in plastid genomes (Ogihara et al., 1988; Hiratsuka et al., 1989). These inversion events were probably triggered by tRNA activity (Hiratsuka et al., 1989) or intragenomic recombination in regions with variable G + C content (Fuller et al., 2001; Smith et al., 2002; Walker et al., 2014). Gene rearrangements and inversions in plastid genomes are believed to have important value in phylogenetic analyses because they are rare, homology estimates are easy, and determination of inversion event polarity is easy (Johansson, 1999; Lee et al., 2007; Jansen et al., 2008; Walker et al., 2014; Yan et al., 2017). Comparison of whole plastid genomes provides the opportunity to explore sequence variation. These comparisons also permit examination of molecular evolutionary patterns associated with structural rearrangement and elucidation of the molecular mechanisms underlying those events.

Ranunculaceae, one of the earliest families that diverged from the eudicots (APG IV, 2016), is composed of more than 2000 mostly herbaceous species with a global distribution (Tamura, 1993, 1995; Ro et al., 1997). In recent years, molecular phylogenetics has provided deep insights and reassessment of Ranunculaceae taxonomy. Some genera have been reduced and a new genus (Gymnacomnium) has been proposed based on molecular phylogenetic analysis results (Compton and Hedderson, 1997; Compton et al., 1998; Ro et al., 1997; Miikeda et al., 2006; Hoot et al., 2012; Wang et al., 2013; Falck and Lehtonen, 2014; Jiang et al., 2017a). All molecular studies to date were mainly based on tandemly repeated nrDNA and several commonly used plastid regions (Compton and Hedderson, 1997; Compton et al., 1998; Wang et al., 2005, 2010, 2013; Miikeda et al., 2006; Hörandl et al., 2009; Emadzade et al., 2010, 2011; Jabbour and Renner, 2012; Falck and Lehtonen, 2014; Cossard et al., 2016; Jiang et al., 2017a). There are only a few complete chloroplast genomes published and accessible from GenBank (http://www.ncbi.nlm.nih.gov). Phylogenetic inferences for Ranunculaceae taxa based on genomic data have yet to be conducted.

Chloroplast genome structural rearrangements and inversions in Anemone and other related genera have been previously reported based on genetic mapping by restriction enzyme site methods (Hoot and Palmer, 1994; Johansson, 1999). In recent years, several complete chloroplast genomes of Ranunculaceae have been published (Chen et al., 2015; Park et al., 2015, 2017; Li et al., 2016; Park and Park, 2016; Jiang et al., 2017b; Lim et al., 2017; Szczecinska et al., 2017; Liu et al., 2018). Recently, Jiang et al. (2017b) and Liu et al. (2018) published plastome sequences of Anemoclema and Clematis s.l. (including Archilematris, Clematis, and Naravelia; Liu et al., 2018), and they also discovered striking structural rearrangements in plastome sequence of the reported genera comparing to that of other Ranunculaceae genera. However, structural variation of tribe Anemoneae plastomes were not discussed in detail by previous studies. The phylogenetic significance of the plastid genome structural variation in tribe Anemoneae and Ranunculaceae still needs to be assessed.

The tribe Anemoneae (as defined by Tamura, 1995) traditionally includes three subtribes (Kingdoninae, Anemoninae, and Clematidinae), based on previous molecular phylogenetic studies that did not include the subtribe Kingdoninae (Hoot et al., 2012; Jiang et al., 2017a). For subtribe Clematidinae, almost all satellite genera of Clematis (such as Naravelia and Archilematris) were nested within Clematis by previous studies (Miikeda et al., 2006; Xie et al., 2011; Lehtonen et al., 2016; Jiang et al., 2017a; Liu et al., 2018). In subtribe Anemoneinae, Hoot et al. (2012) reduced Hepatica, Pulsatilla, Oreithales, Knowltonia, and Barneoudia to the genus Anemone using molecular phylogenetic results inferred from nrITS and atpB-rbcL regions. This result was consistent with subsequent findings by Zhang et al. (2015). However, using six plastid DNA regions, Jiang et al. (2017a) disputed that Anemone s.l. (sensu Hoot et al., 2012) was confirmed to be a paraphyletic group and argues that Hepatica should not be included in Anemone s.l. These two competing phylogenetic hypotheses need to be reconciled using phylogenomic data.

In this study, we report on complete chloroplast genomes from five tribe Anemoneae species (Anemoclema, Anemone, Pulsatilla, and Hepatica). The plastomes of Anemone and Hepatica are reported for the first time. Together with the plastome sequence analyses of Clematis s.l., the aims of this study are to present whole chloroplast genome data for these species; to compare the plastid genomic structure and sequence variation within the tribe Anemoneae; to test two alternative hypotheses (tRNA activity or G + C content variation) that may cause gene rearrangement events; to test competing phylogenetic hypotheses within tribe Anemoneae using plastid phylogenomic data; and to clarify the phylogenetic significance of plastome structural variation of tribe Anemoneae. We also identified repeat sequences and SSRs inside these five plastomes. The data presented in this study should be useful for future phylogenetic studies of tribe Anemoneae and possibly the rest of the buttercup family.

MATERIALS AND METHODS

Plant Sampling

We chose to sample five species, Anemone tomentosa, A. trullifolia, Hepatica henryi, Pulsatilla chinensis, and
Anemoclema glaucifolium, for this study. Fresh young plant leaves were collected from the field for DNA extraction and were dried with silica-gel. Vouchers were deposited at the Beijing Forestry University (BJFC) Herbarium. In our previous study (Liu et al., 2018), complete chloroplast genome sequences of Clematis, Archiclematis, and Naravelia were reported. Thus, in this study, we included samples representing all of the major clades (Hepatica clade, Anemone s.l. clade, Anemoclema, and Clematis s.l. clade) of tribe Anemoneae (Hoot et al., 2012; Zhang et al., 2015; Jiang et al., 2017a) to check plastome structural variation. We also obtained whole chloroplast genomes of other Ranunculaceae species and outgroup species of Berberis available from Genbank for structural comparison and phylogenomic analysis (Table 1).

Sequencing, Plastome Assembly, Annotation, and Visualization

Approximately 50 mg dried leaves were ground for each species, and total DNAs were extracted using cetyl-trimethylammonium bromide (CTAB; Doyle and Doyle, 1987) with the quality of DNAs assessed by agarose gel electrophoresis. Total DNAs were subsequently sent to Novogene (http://www.novogene.com, China) for short insert (350 bp) library construction and next-generation sequencing. Pair end reads of 2 x 150 bp for all tested species were generated on an Illumina Hiseq 4000 genome analyzer platform. Original reads were filtered using the FASTX-Toolkit (http://hannonlab.cshl.edu/fastx_toolkit) to acquire high-quality data by deleting adaptors and low quality reads.

We used BLAT analysis with a Python script (Weitemier et al., 2014) to exclude nuclear and mitochondrial reads using published plastomes from Ranunculaceae as references. Putative chloroplast reads were used for de novo assembly to reconstruct the samples’ complete plastid genomes using Geneious R11 (Kearse et al., 2012) with a low sensitivity setting. Contigs from de novo assembly were annotated in Geneious R11 and then were concatenated into larger contigs based on Pulsatilla vernalis (KR297062) plastomes. All gaps were bridged using conventional Sanger sequencing. The IR region was determined using the Repeat Finder function in Geneious R11. The IR region was subsequently manually inverted and copied to construct the complete plastome sequence. The IR and SC boundaries for all three species were checked using Sanger sequencing.

Complete plastid genomes were manually edited to remove ambiguous sites. The three plastomes were then annotated using the Unix program Plann (Huang and Cronk, 2015) and the annotations were verified using the online program DOGMA (http://dogma.cbb.utexas.edu/; Wyman et al., 2004). If ambiguous annotations were present between Plann and DOGMA, we determined gene boundaries using the online program Blast (Gish and States, 1993). Illustrations of circular plastomes were generated using the Organellar Genome DRAW tool (Lohse et al., 2013). Final plastid genomes were deposited in GenBank (Table 1).

| Species | Sample locality | Voucher (Herbarium) | Genbank accession | Reference |
|---------|-----------------|---------------------|-------------------|-----------|
| Aconitum chinense | Incheon, Korea | VP0000494117 (NIBR) | KT820665 | Lim et al., 2017 |
| Anemone tomentosa | Barkam, Sichuan, China | H. J. Liu 1-1080 (BJFC) | M0001339 | This study |
| Anemone trifoliola | Dinggye, Xizang, China | PE2013 Tibet 2588 (PE) | MH205608 | This study |
| Anemoclema glaucifolium | Shangrila, Yunnan, China | B.XU-M417-090 (SWFC) | MH205609 | This study |
| Archiclematis alternata | Nyalam, Xizang, China | PE2010 Tibet 963 (PE) | MG675221 | Liu et al., 2018 |
| Berberis amurensis | NA | NA | KM057374 | Unpublished |
| Clematis brevicaudata | Jiuteng, Beijing, China | L. Xie 20140706 (BJFC) | MG675223 | Liu et al., 2018 |
| Clematis fusca var. coreana | NA | NA | KM652489 | Park and Park, 2016 |
| Clematis repens | Emei, Sichuan, China | L. Xie 2015EM24 (BJFC) | MG675222 | Liu et al., 2018 |
| Clematis terniflora | Huzhou, Zhejiang Prov. China | Unknown number (HZU) | KJ066785 | Li et al., 2016 |
| Coptis chinensis | NA | NA | KY120323 | Unpublished |
| Gymnacocharmis gymmandrum | NA | NA | KT964697 | Unpublished |
| Hepatica henryi* | Emei, Sichuan, China | L. Xie 2015EM309 (BJFC) | MG001340 | This study |
| Hydrastis canadensis | NA | NA | KY085918 | Unpublished |
| Megaleranthis saniculifolia | Mt. Sobaek, Korea | Unknown number (Korea University Herbarium) | FJ597983 | Kim et al., 2009 |
| Naravelia pilulifera | Yangchun, Guangdong, China | L. Xie & S. Liao 2014022 (BJFC) | KY120887 | Liu et al., 2018 |
| Naravelia zeylanica | Machanbaw, Kachin State, Myanmar | PT-ET 1281 (PE) | MG675224 | Liu et al., 2018 |
| Pulsatilla chinensis* | Yangqing, Beijing, China | L.Xie 2015YQ002 (BJFC) | MG001341 | This study |
| Pulsatilla vernalis | NA | NA | KR297062 | Unpublished |
| Ranunculus macranthus | NA | NA | DQ59689 | Raubeson et al., 2007 |
| Thalictrum coreanum | Gangwon-do, Korea | NA | KM205688 | Park et al., 2015 |
| Trollius chinensis | NA | NA | KX752098 | Unpublished |

Species with asterisks were collected by this study, whereas others were obtained from Genbank. NA, not applicable.
Genome Comparisons
We obtained other complete chloroplast genomes of Ranunculaceae and an outgroup species *Berberis* from Genbank (Table 1) for comparative analyses. The IR/SC boundary regions of species from tribe Anemoneae were illustrated and compared to other Ranunculaceae species and *Berberis*. MAFFT was used to compare the similarity of plastid genome sequences (Katoh et al., 2005) and mVISTA was used to export visual results to evaluate similarity (Frazer et al., 2004). Visual results from mVISTA were further analyzed using two alignment programs: LAGAN, which produces true multiple alignments regardless of whether they contain inversions or not, and Shuffle-LAGAN, which can detect rearrangements and inversions (Brudno et al., 2003a,b). Detailed gene inversions and transpositions were identified by comparing the gene order of Ranunculaceae samples to *Berberis* with a whole plastome alignment method that used Mauve v2.3.1. (Darling et al., 2010).

Analysis of G + C Content at Inversion/Transposition Borders and Sliding Window Analysis
The G + C content was calculated for the spacer regions boundary each of the major inversions and for transpositions of the tribe Anemoneae' plastomes. Flanking regions were defined as the noncoding sequence between the nearest coding genes on either side of the inversion/transposition boundary (Walker et al., 2014). We also conducted a sliding window analysis to identify the nucleotide variability (Pi) in the inversion/transposition regions using DnaSP version 5 (Librado and Rozas, 2009).

Characterization of Repeat Sequences and SSRs
We used REPuter (Kurtz et al., 2001) to identify and locate the repeat sequences for the newly sampled species, including direct, reverse and palindromic repeats, within the plastid genome. For repeat identification, the following parameters were used: (1) 30 bp minimum repeat size and (2) 90% or greater sequence identity (Hamming distance = 3). In order to avoid redundancy, repeat sequence analysis was carried out with a single IR region. SSRs were determined using MISA (Thiel et al., 2003) and parameters were set to 10 repeat units ≥ 10 for mononucleotide SSRs, six repeat units ≥ 6 for dinucleotide, five repeat units ≥ 5 for trinucleotide, four repeat units ≥ 4 for tetranucleotide, and three repeat units ≥ 3 for pentanucleotide and hexanucleotide SSRs.

Phylogenomic Analysis
We used the plastome sequences of all the other genera of Ranunculaceae available from GenBank for phylogenomic analysis. The sampling covered 15 of the family’s genera, seven of which belong to tribe Anemoneae. There is an accession of *Actaea* plastome (NC034704, unpublished) sequence available in GenBank. However, after extracting genes from this plastome and doing Blast in GenBank, we found that this sample belongs...
to Apiaceae and was mis-identified as *Actaea*. Thus, we did not include this sequence in the present analysis. *Berberis amurensis* plastome was chosen as the outgroup (**Table 1**).

In this study, we first separated the complete plastome sequences into coding regions (protein-coding genes, as well as tRNA genes and rRNA genes), intergenic spacer regions, and introns. Gene orders for all samples were tested using mVISTA, and then were shuffled in the same order with a *Berberis amurensis* plastome sequence whenever gene inversions/transpositions were found. All data sets were then aligned using MAFFT v6.833 ([Katoh et al., 2005](#)) and manually adjusted by MEGA 7.0 ([Kumar et al., 2016](#)). The ambiguous alignments were removed from the data sets for phylogeny reconstruction. Substitution models and data partitions for Bayesian analysis was determined by PartitionFinder v2.1.1 ([Lanfear et al., 2016](#)) and the best scheme selected by Akaike information criterion (AIC; [Posada and Buckley, 2004](#)).

Phylogeny reconstruction occurred using data sets of LSC, SSC, IR, and complete plastomes with Parsimony (MP) and Bayesian methods. Each data set was further separated into CDs, intron, intergenic spacer regions ([Ma et al., 2014](#)).

Parsimony (MP) analysis was conducted for all the separated data sets and the complete plastome data set using PAUP* 4.0b10 ([Swofford, 2003](#)). All characters were treated as unordered and equally weighted, and gaps were set as missing data. We used Branch-and-Bound or heuristic search (1000 replicates), simple addition, and tree bisection-reconnection branch swapping with MUL-trees to search the MP tree(s). Branch support values were assessed by performing 1000 bootstrap replicates using 1000 random taxon addition replicates with 10 trees held at each step and TBR swapping.

Bayesian inference (BI) was conducted with MrBayes v3.2.3 ([Ronquist and Huelsenbeck, 2003](#)) using partitioned substitution models tested by PartitionFinder. Two independent Markov chain Monte Carlo (MCMC) chains were run, each with three heated and one cold chain for 2,000,000 generations and sampling trees every 100 generations. The MCMC convergence in Bayesian inference was checked by AWTY ([https://github.com/danlwarriner/RWTY, Warren et al., 2017](#)). The first 20% of trees were discarded as burn-in with the remaining trees being used for generating the consensus tree.

**RESULTS**

**Strategies for Assembling Plastomes**

We obtained 2.7 Gb of average NGS clean data for each species, with minimums and maximums of 2.2 Gb (*Pulsatilla chinensis, 6,309,984 reads*), and 3.3 Gb (*Anemone tomentosa, 9,512,414 reads*), respectively. Blat analysis selected out 542,906 putative plastid reads for *Anemone tomentosa*, 451,685 reads for *A. trullifolia*, 100,695 reads for *Pulsatilla*, 128,187 reads for *Hepatica*, and 335,675 reads for *Anemoclome*. *Anemone tomentosa* reads obtained three large contigs from *de novo* assembly (83,809 bp, 27,444 bp, and 20,496 bp). Three gaps were bridged using Sanger sequencing. For *A. trullifolia* data, only one large contig (126,074 bp) was derived that included complete LSC, IR, and SSC regions and regions without gaps. For *Pulsatilla chinensis* plastid reads, eight large contigs ranging from 3,287 bp to 38,885 bp were obtained. We filled seven gaps for this sample. For *Hepatica henryi*, two large contigs (45,926 bp and 65,671 bp) were obtained and one large gap (ca. 1.5 kb) was filled by designing two pairs of primers. For *Anemoclome* data, three large contigs (92,619 bp, 33,225 bp, and 4,489 bp.) were concatenated. Two gaps were bridged using Sanger sequencing. Information for all of the gaps and primers for Sanger sequencing are presented in the Supplementary Materials.

**Size and Structure of Plastomes of the Samples**

The genome size of the five newly sequenced samples ranged from 157,096 bp (*Anemone trullifolia*) to 163,669 bp (*Pulsatilla chinensis*; **Figure 1**), and the overall GC content varied from 37.2% (*Pulsatilla chinensis*) to 37.9% (*Anemoclome glaucifolium* and *Hepatica henryi*; **Table 2**). All five plastid genomes consisted of a pair of IRs (31,022–31,490 bp) separated by the LSC (78,795–82,339 bp) and SSC (16,257–19,100 bp) regions, respectively (**Table 2**). The plastomes of the five samples encoded an identical

**Table 2 | Base compositions of five newly sequenced Anemoneae plastomes.**

| Features             | Anemone tomentosa | Anemone trullifolia | Anemoclome glaucifolium | Pulsatilla chinensis | Hepatica henryi |
|----------------------|-------------------|---------------------|-------------------------|---------------------|-----------------|
| Genome size          | 162,213           | 157,096             | 160,939                 | 163,669             | 160,283         |
| Length of LSC        | 82,237            | 78,796              | 80,250                  | 82,339              | 80,558          |
| Length of SSC        | 16,906            | 16,257              | 17,637                  | 19,100              | 17,647          |
| Length of IR         | 31,490            | 31,022              | 31,256                  | 31,115              | 31,039          |
| Total G + C content (%) | 37.6%             | 37.6%               | 37.9%                   | 37.2%               | 37.9%           |
| Total number of genes| 112               | 112                 | 112                     | 112                 | 112             |
| Protein encoding     | 79                | 79                  | 79                      | 79                  | 79              |
| tRNA                 | 29                | 29                  | 29                      | 29                  | 29              |
| rRNA                 | 4                 | 4                   | 4                       | 4                   | 4               |
| Genes with introns   | 18                | 17                  | 18                      | 18                  | 17              |
| Duplicated in IRs    | 25                | 25                  | 25                      | 25                  | 25              |

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**Legend:**

- **Actaea:** Species that was mis-identified in the study.
- **Table 1:** Features of the five newly sequenced Anemoneae plastomes.
- **Table 2:** Base compositions of five newly sequenced Anemoneae plastomes.
set of 112 genes, including 78 protein-coding genes, 29 transfer RNAs, four ribosomal RNAs, and 25 genes are duplicated in IRs. There are 18 genes with introns in the plastomes of *Anemoclema*, *Anemone tomentosa*, and *Pulsatilla chinensis*, whereas, 17 genes with introns are present in *Anemone trullifolia* and *Hepatica henryi* plastomes (no introns in *rps16*).

**FIGURE 2** | Comparison of the LSC, IRs and SSC boundary regions of tribe Anemoneae, published Ranunculaceae genera, and *Berberis* plastomes.
Plastomes of Tribe Anemoneae

Chloroplast Genome Comparison

We compared the IR/SC boundary regions of tribe Anemoneae to other Ranunculaceae species as well as an outgroup from Berberis. The junction positions are similar and conserved in tribe Anemoneae, yet differ from other Ranunculaceae species and Berberis (Figure 2). For example, rps19-infA (seven genes) is in the LSC region of Berberis, Hydrastis, Coptis, Thalictrum, Megaleranthis, Ranunculus, Aconitum, and Trollius plastomes, but is located in IR in Clematis (including Archiclematis and Naravelia), Anemoclema, Anomone, Pulsatilla, and other species (Figure 2).

![Sequence alignment of tribe Anemoneae, published Ranunculaceae genera, and Berberis plastome using the mVISTA program. A cut-off of 70% similarity was used for the plot and the Y-scale represents the percent similarity ranging from 50-100%. Blue represents coding regions, and pink represents non-coding regions. (A) LAGAN method (as described in Materials and Methods); (B) Shuffle LAGAN method (as described in Materials and Methods).]
and Hepatica). This difference makes the IR region of tribe Anemoneae roughly 4.4 kb longer than in other genera of the family. The gene orders located within the IR-SSC and IR-LSC boundaries are similar among tribe Anemoneae samples but different from those in other Ranunculaceae genera (Figure 1).

To investigate levels of genome divergence, multiple alignments of plastid genomes were performed (Figures 3A,B and Supplementary Materials). For mVISTA analysis, LAGAN, and Shuffle-LAGAN program results differed because of gene inversion/transposition occurring in tribe Anemoneae (Figures 3A,B). When using the LAGAN method (Figure 3A), plastomes of Anemoclema, Anemone, Pulsatilla, and Hepatica showed consistency in gene order, but carried large unalignable regions in LSC comparing to Clematis and other Ranunculaceae species. Plastomes of Clematis s.l. also showed some unalignable regions compared to other Ranunculaceae species. Conversely, all of the sequences aligned well when Shuffle-LAGAN methods were used (Figure 3B). This approach also revealed high sequence similarity across the coding region accompanied by more variability in non-coding regions.

tribe Anemoneae samples exhibited inversion and transposition regions that were detected by MAUVE (Supplementary Materials), compared to other Ranunculaceae plastomes. In Anemoclema, Anemone, Hepatica, and Pulsatilla, three inversions are present in the LSC region (Figure 1). The first inversion (INV 1) is located between trnS-GGA and trnS-GCU (ca. 40 kb in length). The second inversion (INV 2) was between rps16 and trnH-GUG (ca. 5k in length). The third inversion region (INV3) was located in rps4 (ca. 600 bp in length). Two inversions and one transposition were detected in Clematis s.l. plastomes, with INV2, and INV3 being similar to those in Anemone s. l. plastomes. The transposition region 1 (TP1) was located between trnL UAA and ndhC (ca. 3k in length; Figure 1). No inversions or transpositions were found in published plastomes of other Ranunculaceae genera or in comparison to other plastomes of angiosperm species like Amborella trichopoda, Berberis amurensis, and Nicotiana tabacum.

### G + C Content at Inversion and Transposition Borders and Sliding Window Analysis

The G + C content of the sequence in boundary regions was detected to be lower than the average G + C content of whole plastome and all the noncoding regions (Table 3). The sliding window analysis revealed that higher nucleotide variability (Pi) was exhibited at SC regions in comparison to IR regions (Figure 4). Genetic variation was particularly high at the boundary of INV1, which existed in Anemoclema, Anemone, Hepatica, and Pulsatilla plastomes. The INV2 and TP1, found in Clematis s.l. plastomes, has a high nucleotide variability at the borders as well. However, we did not find high genetic variation in INV3, which is shared among all the Anemoneae species.

### Table 3

| Species                     | Whole plastome | Noncoding regions | Inversion 1 | Noncoding regions | Inversion 2 | Noncoding regions | Inversion 3 | Noncoding regions | Transposition borders |
|-----------------------------|----------------|-------------------|-------------|-------------------|-------------|-------------------|-------------|-------------------|------------------------|
| Archiclematis alternata     | 38.00%         | 34.60%            | 23.70%      | 23.70%            | 23.70%      | 23.70%            | 23.70%      | 23.70%            | 23.70%                 |
| Clematis brevicaudata       | 38.00%         | 34.60%            | 23.70%      | 23.70%            | 23.70%      | 23.70%            | 23.70%      | 23.70%            | 23.70%                 |
| Clematis fusca var. cornnea | 38.00%         | 34.60%            | 23.70%      | 23.70%            | 23.70%      | 23.70%            | 23.70%      | 23.70%            | 23.70%                 |
| Clematis rubens             | 38.00%         | 34.60%            | 23.70%      | 23.70%            | 23.70%      | 23.70%            | 23.70%      | 23.70%            | 23.70%                 |
| Naravelia pilulifera        | 37.90%         | 34.40%            | 23.70%      | 23.70%            | 23.70%      | 23.70%            | 23.70%      | 23.70%            | 23.70%                 |
| Anemoclema glaucifolium     | 38.00%         | 34.60%            | 23.70%      | 23.70%            | 23.70%      | 23.70%            | 23.70%      | 23.70%            | 23.70%                 |
| Pulsatilla chilensis        | 37.90%         | 34.40%            | 23.70%      | 23.70%            | 23.70%      | 23.70%            | 23.70%      | 23.70%            | 23.70%                 |
| Hepatica henryi             | 37.90%         | 34.40%            | 23.70%      | 23.70%            | 23.70%      | 23.70%            | 23.70%      | 23.70%            | 23.70%                 |
| Mean                        | 37.83%         | 34.29%            | 23.36%      | 22.80%            | 23.70%      | 23.70%            | 23.70%      | 23.70%            | 23.70%                 |

NA, not applicable.
Repetitive Sequences
We detected a total of 173 repeats including direct, reverse, palindromic and complement repeats in the five newly sequenced plastomes (Figure 5). The most common repeat types are direct repeats, which account for 45% of the total repeats, followed by palindromic repeats (35%) and reverse repeats (16%). The only two complement repeats were found in Pulsatilla plastomes. Most of the repeats were short, ranging from 30–59 bp. However, a few much longer direct and reverse repeats (up to more than 100 bp) were found in Hepatica and Anemone plastomes. The majority of repeats were located in noncoding regions (84%), among which only 3% were found in introns. There were 16% repeats detected in CDs.

We also investigated repeats that are shared among plastomes of the five samples by using strict criteria, i.e., repeats which are identical in length and located in homologous regions were defined as shared repeats. Under this criteria, there were four repeats shared by all five species. All tested plastomes possessed their own repeats, with the number varying from 13 (Hepatica henryi) to 32 (Pulsatilla chinensis; Figure 5). These repeats may serve as potential population genetic markers for further studies.

SSR Polymorphisms
We identified 57, 57, 43, 51, and 61 SSRs in Anemone tomentosa, A. trullifolia, Anemoclema glaucifolium, H. henryi, and P. chinensis, respectively (Table 4). A mononucleotide repeat unit (A/T) was found to be the most abundant, accounting for 65–95% among the five species. This was followed by a dinucleotide repeat unit (AT/AT) with particular numbers of four, six, one, three and two SSRs, respectively. The trinucleotide repeat unit (AAT/ATT) was detected in Anemone trullifolia, Hepatica henryi, and Pulsatilla chinensis, and only one tetranucleotide repeat (AAGT) was found in Anemone tomentosa. Multiple pentanucleotide repeats were detected in Anemone tomentosa, Hepatica henryi, and Pulsatilla chinensis. One hexanucleotide repeat was present in Pulsatilla chinensis plastome. The mononucleotide repeat unit C/G was also identified in all five species. Within the five plastomes, SSR loci were mainly located in IGS, followed by locations in CDS and introns. As expected, most SSRs

FIGURE 4 | A sliding window analyses of the whole plastid genomes of Clematis s.l. (upper) and Anemoclema, Anemone, Hepatica, and Pulsatilla samples (lower). Circles identify the regions bordering inversion sites, and lines parallel to the x-axis identify the positions of the LSC, SSC and IR regions.
were located in the LSC region, followed by SSC and IR regions.

**Phylogenomic Analysis**

Phylogenies, reconstructed with each data set and by both methods were consistent with each other and only differed for some nodes’ supporting values. Because complete plastome sequence data provided the most robust phylogeny, we used this result (Figure 6). Excluding ambiguous alignments, the complete chloroplast genome data set alignment was 123,519 bp in length (including 16,916 informative characters). Only one parsimony tree with 58,536 steps was searched, along with a consistency index (CI) of 0.75, and a retention index (RI) of 0.75. The Bayesian analysis used partitioned substitution models checked by PartitionFinder. The length of each coding region, intron, intergenic spacer, and models of each partition subset tested by PartitionFinder are provided in the Supplementary Material.

Parsimony analysis of the major Ranunculaceae clades did not resolve the sister relationship of *Aconitum* clade and *Ranunculus* + tribe Anemoneae clade. The *Thalictrum*, *Trollius*, and *Megaliranthis* clade was also not fully supported by MP analysis (Figure 6). The Bayesian analysis using partitioned models resolved all of the family’s major clades. Except for the *Clematis* s.l. clade, all clades were fully supported (PP = 1) by the Bayesian method.

Tribe Anemoneae was supported by our phylogenomic analyses and was closely related to *Ranunculus*. There was strong support for grouping *Archiclematis*, *Clematis*, and *Naravelia* as

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**FIGURE 5** | Analyses of repeated sequences in five newly sequenced plastomes. (A) Number of four repeat types; (B) Frequency of direct repeats by length; (C) Frequency of reverse repeats by length; (D) Frequency of palindromic repeats by length; (E) Location of repeats; (F) Summary of shared repeats among the five plastomes.
TABLE 4 | Simple sequence repeats (SSRs) for five newly sequenced plastome samples.

| Genomes            | Repeat units | Number | Location | Region |
|--------------------|--------------|--------|----------|--------|
|                    |              |        | Intron   | IGS    | CDS    | LSC  | SSC  | IR  |
| Anemone tomentosa  | A/T          | 48     | 4        | 39     | 5      | 38   | 6    | 4   |
|                    | C/G          | 1      | 1        |        |        |      |      |     |
|                    | AT/AT        | 4      | 4        | 4      |        |      |      |     |
|                    | AAGT/ACTT    | 1      | 1        |        |        |      |      |     |
|                    | AAAAG/CTTT   | 2      | 2        |        |        |      |      |     |
|                    | AATAT/ATATT  | 1      | 1        |        |        |      |      |     |
| Anemone trullifolia| A/T          | 47     | 8        | 31     | 8      | 38   | 5    | 4   |
|                    | C/G          | 2      | 1        | 1      | 2      |      |      |     |
|                    | AT/AT        | 6      | 1        | 5      | 4      | 2    |      |     |
|                    | AAT/ATT      | 2      | 2        |        |        |      |      |     |
| Hepatica henryi    | A/T          | 33     | 5        | 23     | 5      | 28   | 5    |     |
|                    | C/G          | 1      | 1        |        |        |      |      |     |
|                    | AT/AT        | 3      | 1        | 2      | 2      | 1    |      |     |
|                    | AAT/ATT      | 1      | 1        |        |        |      |      |     |
|                    | AAATT/AATT   | 1      | 1        |        |        |      |      |     |
|                    | AAGAT/ATCTT  | 1      | 1        |        |        |      |      |     |
|                    | AATAT/ATATT  | 1      | 1        |        |        |      |      |     |
| Pulsatilla chinensis| A/T         | 46     | 7        | 32     | 7      | 36   | 8    | 2   |
|                    | C/G          | 4      | 2        | 2      | 2      |      |      |     |
|                    | AT/AT        | 2      | 1        |        |        |      |      |     |
|                    | AAT/ATT      | 3      | 3        | 1      | 2      |      |      |     |
|                    | AAATT/AATT   | 1      | 1        |        |        |      |      |     |
|                    | AAGAT/ATCTT  | 1      | 1        |        |        |      |      |     |
|                    | AATAT/ATATT  | 1      | 1        |        |        |      |      |     |
| Anemoclema glaucifolium| A/T      | 41     | 5        | 28     | 8      | 27   | 9    | 4   |
|                    | C/G          | 1      | 1        |        |        |      |      |     |
|                    | AT/AT        | 1      | 1        |        |        |      |      |     |

a sister clade to Anemoclema. Samples of Anemone, Pulsatilla, and Hepatica did not group as a clade but as a paraphyletic grade in tribe Anemoneae. The Hepatica henryi + Anemone trullifolia (sect. Omalocarpus) clade was a sister clade to Clematis + Anemoclema. Anemone tomentosa (sect. Rivularidium) + Pulsatilla clade was found to be the first diverged clade within the tribe.

**DISCUSSION**

**Structural Rearrangements of Chloroplast Genome Detected in Tribe Anemoneae**

We found two derived types of chloroplast genomes in tribe Anemoneae compared to other genera within Ranunculaceae, with one type (with two inversions and one transposition regions) in Clematis s.l. and the other (with three inversions) in the rest of the tribe's genera (Figures 1, 6). These gene rearrangements clearly bear important phylogenetic information. Two inversions (INV2 and INV3) in the Clematis s.l. plastome were also present in Anemoclema, Anemone, Hepatica, and Pulsatilla plastomes. Thus, the presence of INV2 and INV3 could be considered as a synapomorphy of tribe Anemoneae. The largest inversion (INV 1) is present in Anemoclema, Anemone, Hepatica, and Pulsatilla plastomes, and these samples were paraphyletic to Clematis s.l. clade (Figure 6). This suggests that the presence of INV 1 may be a pleisiomorphy within the tribe. In contrast, the only transposition region (TP 1) present in the Clematis s.l. clade may represent a synapomorphy for Clematis, Archiclematis, and Naravelia. The phylogenomic results suggest that the ancestor of tribe Anemoneae may have a plastome sequence similar to that of Anemone, Anemoclema, Hepatica, and Pulsatilla which carried three inversions in its LSC regions. Two steps of gene rearrangements subsequently occurred in the plastome sequence of the Clematis s.l. ancestor. One of these was the loss of INV 1, and the other was the addition of TP1.

Inversion and transposition events in the chloroplast genome may be triggered by tRNA activity (Hiratsuka et al., 1989; Walker et al., 2014) or intragenomic recombination at regions with variable G + C content (Fullerton et al., 2001; Smith et al., 2002). In this study, we evaluated the G + C content at the boundaries
of each inversion and relocation (Table 3). The regions flanking inversions and relocations had lower G + C contents than non-coding regions of the whole plastome. In contrast, the flanks of all inversion/transposition regions had tRNA genes, as well as higher genetic variation (Figure 4). The tRNA activity, higher genetic variation, and lower G + C content present in flank regions could be the key factors promoting gene rearrangements in chloroplast genomes.

**IR Expansion, Gene Duplications, and Other Genomic Features in Tribe Anemoneae**

Although genome size and overall genomic structure are highly conserved in land plants, IR expansion/contraction is common in plastid genomes and is the main outcome of plastid genome length variation in angiosperms (Kim and Lee, 2004). Gene duplications in plastid genomes are mainly caused by the expansion of the IR region to single copy regions (Goulding et al., 1996). Expansion events that result in the duplication of single genes, parts of genes, or several genes, have been documented in several plant taxa. This includes a 12-kb expansion in Nicotiana acuminate (Goulding et al., 1996), 4-kb expansion in Jasminum nudiflorum (Lee et al., 2007), and a remarkable 50-kb expansion in Pelargonium (Chumley et al., 2006). In Ranunculaceae, the termini of two genes, rps19 and trnH-GUG, were previously reported to have migrated into adjacent IRs (Park et al., 2015).

We observed an approximately 4.4-kb expansion of the IR toward the LSC region in tribe Anemoneae. This expansion
caused duplicate copies of six ribosomal protein genes (rps8, rpl14, rpl16, rps3, rpl22, rps19). These are single genes in other genera of the family, such as Ranunculus, Thalictrum, Megetaleranthus, Trollius, and Aconitum (Hoot and Palmer, 1994; Chen et al., 2015; Park et al., 2015), as well as most other angiosperms. For this reason, this IR expansion could also be considered as a synapomorphy of tribe Anemoneae.

Simple sequence repeats (SSRs), also known as microsatellites, are often used as genetic markers for population genetics studies. This is because they provide rich information for population genetics and evolutionary studies (Powell et al., 1995). However, plastid SSRs were rarely used in tribe Anemoneae. We identified 43 to 61 chloroplast SSRs in the five samples we evaluated (Table 4). Our results showed that Anemone tomentosa, Hepatica henryi, and Pulsatilla chinensis have pentanucleotide repeats in their plastomes, and that the Pulsatilla chinensis plastome has hexanucleotide repeats. The rich diversity of chloroplast SSR loci provides opportunities to survey the population genetic structure of those species.

Phylogenomic Inference
All previous phylogenetic studies of Ranunculaceae were based on small numbers of DNA regions (Ro et al., 1997; Wang et al., 2009, 2016; Cossard et al., 2016), and there is a need for improved resolution of Ranunculaceae phylogeny needs to be further improved. In this study, phylogeny inferred from complete chloroplast genomic data was better resolved and more rigorous than previous studies (Figure 6), thereby demonstrating that plastome sequences may provide the ideal data sets for resolving family phylogenies. tribe Anemoneae was supported and tested to be sister to the genus Ranunculus, which is in agreement with morphological classifications (Tamura, 1995) and previous molecular phylogenetic studies (Wang et al., 2009, 2016; Cossard et al., 2016). Although the gene order of Anemoclema plastome was found to be identical with Anemone (Figure 1), this genus has a sister relationship to Clematis clade (Figure 6) as previously reported by molecular phylogenetic analyses (Zhang et al., 2015; Jiang et al., 2017a). Unlike results by Hoot et al. (2012), samples of Anemone, Hepatica, and Pulsatilla did not form a monophyletic group. Thus, our results did not support classification by Hoot et al. (2012), which included Hepatica into Anemone s.l.

In this study, Hetapica henryi, and Anemone trullifolia (sect. Omalocarpus) are grouped together, whereas Pulsatilla chinensis and Anemone tomentosa (sect. Rivularidum) are grouped and these two clades were paraphyletic to the Anemoclema + Clematis clade (Figure 6). These results are somewhat similar to the phylogenetic analyses by Jiang et al. (2017a), but still different with their phylogenetic topology. In Jiang et al. (2017a), Pulsatilla + sect. Rivularidum clade was sister to Anemoclema + Clematis clade, and Hepatica + sect. Omalocarpus clade was outside. In the present study, Hepatica + sect. Omalocarpus clade was sister to Anemoclema + Clematis clade, whereas Pulsatilla + sect. Rivularidum clade was outside. Statistical support of sister relationship of Pulsatilla + sect. Rivularidum clade and Anemoclema + Clematis clade by Jiang et al. (2017a) was not very strong. However, the sister relationship of Hepatica + sect. Omalocarpus and Anemoclema + Clematis clade was fully supported by both MP and Bayesian analyses by plastome phylogenomic analyses in this study.

Within tribe Anemoneae, generic relationship within subtribe Clematidinae (sensu Tamura, 1995) was clear. All the small genera like Archiclematis and Naravelia should be included into Clematis s.l. (Miikeda et al., 2006; Xie et al., 2011; Zhang et al., 2015; Liu et al., 2018; Jiang et al., 2017a), and its sister relationship was supported by all the analyses (Zhang et al., 2015; Jiang et al., 2017a; and this study). However, subtribe Anemoniinae (sensu Tamura, 1995) showed very complicated evolutionary patterns. Simply grouped together and treated them as Anemone s.l. should be reconsidered. According to this study, subtribe Anemoniinae should be separated at least three genera (Anemoclema, Anemone s.l. including Pulsatilla and Pulsatilloides, and Hepatica including sect. Omalocarpus, sect. Anemonidium, and sect. Keiskea) as suggested by Jiang et al. (2017a), and Anemoclema is better to be treated as a member of subtribe Clematidinae.

AUTHOR CONTRIBUTIONS
HL and JH contributed equally. LX and JC initiated the project. LX, JH, HL, CD, and LP conceived and designed the experiments. HL, JH, and RL performed the experiments. JH and HL analyzed the data. HL, JH, and LX wrote the manuscript.

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SUPPLEMENTARY MATERIAL
The Supplementary Material for this article can be found online at: https://www.frontiersin.org/articles/10.3389/fpls.2018.01097/full#supplementary-material

Figure S1 | MALVE alignment of Ranunculaceae plastomes. Homologous regions are shown in the same color.
Table S1 | Detailed informations of gaps between contigs in de novo assembly, and primers for Sanger sequencing that bridges the gaps.
Table S2 | Results of substitution model and data partition by PartitionFinder.
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**Conflict of Interest Statement:** The authors declare that the research was conducted in the absence of any commercial or financial relationships that could be construed as a potential conflict of interest.

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