

## **Plastid Primers for Angiosperm Phylogenetics and Phylogeography**

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Source: Applications in Plant Sciences, 3(6)

Published By: Botanical Society of America

URL: <https://doi.org/10.3732/apps.1400085>

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## PLASTID PRIMERS FOR ANGIOSPERM PHYLOGENETICS AND PHYLOGEOGRAPHY<sup>1</sup>

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- **Premise of the study:** PCR primers are available for virtually every region of the plastid genome. Selection of which primer pairs to use is second only to selection of the genic region. This is particularly true for research at the species/population interface.
- **Methods:** Primer pairs for 130 regions of the chloroplast genome were evaluated in 12 species distributed across the angiosperms. Likelihood of amplification success was inferred based upon number and location of mismatches to target sequence. Intraspecific sequence variability was evaluated under three different criteria in four species.
- **Results:** Many published primer pairs should work across all taxa sampled, with the exception of failure due to genomic reorganization events. Universal barcoding primers were the least likely to work (65% success). The list of most variable regions for use within species has little in common with the lists identified in prior studies.
- **Discussion:** Published primer sequences should amplify a diversity of flowering plant DNAs, even those designed for specific taxonomic groups. “Universal” primers may have extremely limited utility. There was little consistency in likelihood of amplification success for any given publication across lineages or within lineage across publications.

**Key words:** comparative sequencing; complete chloroplast genome; cpDNA.

Whole genome sequencing is more available and less expensive than ever before, yet most scientists continue to rely on targeted, comparative sequencing for phylogenetics and phylogeography. Identifying the most appropriate markers to employ has been challenging. Information for model organisms abounds (e.g., grasses; Saski et al., 2007; Bortiri et al., 2008; Leseberg and Duvall, 2009), and a few studies have specifically screened the same set of markers across a diversity of plant groups, ranking the utility of these markers either explicitly or implicitly (Shaw et al., 2005, 2007, 2014). These studies are exceedingly valuable, demonstrating there is no one-size-fits-all answer to the question “which markers?”. The second critical question to “which markers” is “which primers?”. Hundreds of primer sequences have been published, many designed for specific taxonomic groups. The work presented here was inspired by “The Tortoise and the Hare II” (Shaw et al., 2005), which was the first study to pull together information on a large number of regions commonly in use (at that time) for plant phylogenetics. Our laboratory was also compiling such information, as were many others.

<sup>1</sup>Manuscript received 8 September 2014; revision accepted 24 March 2015.

This work was inspired by phylogenetic combs obtained for *Guzmania* (Bromeliaceae) and *Monardella* (Lamiaceae), and by the work of Joey Shaw. Some plant material or DNA samples were provided by The Desert Botanic Garden, S. Eliason, E. A. Friar, Y.-L. Qiu, S. Vanderplank, and G. Wallace. Cris Martinez and Bill Waggoner assisted with primer screening. Harith Alappat assisted with whole chloroplast genome and primer alignments. Rancho Santa Ana Botanic Garden provided financial support.

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doi:10.3732/apps.1400085

The Tortoise and the Hare II paper was revolutionary in assessing sequence variability for all regions studied across a broad diversity of flowering plants, and providing a ranking of that variability. In the mid-2000s, a small number of complete chloroplast genome sequences were available for land plants and some of those were not annotated (e.g., *Medicago truncatula* Gaertn. [GenBank NC\_003119]; Saski et al., 2005). Grivet et al. (2001) were visionary when they moved beyond analyzing regions commonly being used to design primers for lesser-known and potentially faster-evolving regions of the chloroplast genome. They were the first to take advantage of the new genomic data boom, providing a set of 20 universal chloroplast primers designed around the complete chloroplast data from seven flowering plant species. Around the same time, I developed nondegenerate primers for 36 noncoding regions in the large and small single-copy regions of the chloroplast genome (published here). These near-universal primers were designed based on the complete chloroplast genome sequences of 16 flowering plant species (see Appendix 1).

Grivet et al. (2001) and I designed primers, but Shaw et al. (2007) took an even more applied approach when they examined sequences for three different taxon pairs (*Atropa/Nicotiana*, *Lotus/Medicago*, and *Saccharum/Oryza*), specifically searching for faster-evolving regions. Shaw et al. (2014) go one step further, comparing complete chloroplast genome sequences for 25 (primarily congeneric) sister species pairs. They examined sequence diversity for 107 single-copy noncoding regions, providing the most comprehensive analysis to date.

There are now at least 150 primer pairs available to amplify almost every intergenic, intron, and exon region of the chloroplast genome, including portions of the inverted repeats, thanks to the efforts of Shaw et al. (2005, 2007, 2014) and others (Ebert and Peakall, 2009; Scarcelli et al., 2011; Dong et al., 2012,

2013). Not surprisingly, although all worked independently, many of the same regions were explored (Appendix 2) and, in some cases, identical or nearly identical primers were designed. The push to identify faster-evolving regions was, in part, spurred by groups of organisms with exceptionally slowly evolving chloroplast genomes such as Bromeliaceae (Gaut et al., 1992) and Arecaceae (Asmussen and Chase, 2001). Heinze provided access to a comprehensive database of chloroplast primers in 2007 (Heinze, 2007). The database is periodically updated (last update 18 March 2014) and is available at <http://bfw.ac.at/200/2043.html>.

In the absence of taxon-specific complete chloroplast genome data, it is possible to mine the wealth of genomic data available in international databases such as GenBank (National Center for Biotechnology Information), EMBL-Bank (European Molecular Biology Laboratory), and DDBJ (DNA Data Bank of Japan). Primer pairs for 130 regions of the chloroplast genome were evaluated relative to representatives of 12 genera, spanning the diversity of flowering plants. Exon regions were avoided because they generally evolve more slowly than intron and intergenic spacer regions. The primers of Shaw et al. (2005, 2007), Scarcelli et al. (2011), and Dong et al. (2012), as well as the primers provided here, were evaluated. Many of the Shaw et al. (2005, 2007) and Scarcelli et al. (2011) primers are degenerate, improving the breadth of taxa they can be used on, but reducing their efficiency during the amplification process. The Dong et al. (2012) primers are primarily used for barcoding, thus amplify a diversity of taxa, but may not target the most quickly evolving regions of the genome. The likelihood of amplification success was estimated based upon the number and position of mismatches between the primer and the target sequence. These data were then evaluated in the context of Shaw et al. (2014) to provide generalizations, by taxonomic group, for primer utility in conjunction with sequence variability.

Finally, a small number of plant species have sequences available for multiple accessions or different subspecific taxa including *Fragaria vesca* L. (Rosaceae,  $N = 2$ ), *Gossypium herbaceum* L. (Malvaceae,  $N = 2$ ), *Olea europaea* L. (Oleaceae,  $N = 4$ ), and *Oryza sativa* L. (Poaceae,  $N = 3$ ). Shaw et al. (2014) specifically excluded species pairs with very low and very high levels of sequence divergence. Very high levels of divergence made alignment difficult, and very low levels provide too few characters for reasonable comparison across all flowering plants. Here I compare the variation at the subspecific level to that of higher-level relationships to determine if the same regions are useful at multiple taxonomic levels.

## METHODS

**Primers designed here**—Sixteen chloroplast genomes, representing a diversity of flowering plants, were downloaded from GenBank (see Appendix 1). Homologous gene sequences were aligned in Se-AL version 2.0a11 (Rambaut, 1996). Primers were designed based on simultaneous viewing of the Se-AL file and an Oligo 4.02 (Rychlik, 2002) file, using a single sequence from the pool. Primers were anchored in coding regions and were designed to have a minimum number of hair-pins and primer-primer interactions, annealing temperatures between 50°C and 64°C, and a 3' GC clamp if possible, targeting regions 400–1800 bp in length. Primer details are provided in Table 1, and are provided in the order of appearance in the tobacco genome (*Nicotiana tabacum* L. [GenBank Z00044.1]). The tobacco genome was the genome of choice for describing the location of primers prior to the recent accumulation of genomic data. A total of three different *trnS* primers were designed, corresponding to the three *trnS* genes encoded by the chloroplast genome (*trnS*-GCU, *trnS*-UGA, and *trnS*-GGA). Gene order is highly conserved on the chloroplast genome of

flowering plants, but does vary and can be highly informative, for example, as in the 22-kb inversion in almost all Asteraceae (Jansen and Palmer, 1987a, 1987b) and the 78-kb inversion in Fabaceae subtribe Phaseolinae (Bruneau et al., 1990). Some primer combinations are not useful in particular groups of plants due to structural rearrangements. In some cases, the downloaded genomes differ in the identification of specific genes.

**Primer utility**—The chloroplast genomes for species of eight genera (*Acorus* L., *Amborella* Baill., *Canna* L., *Ceratophyllum* L., *Cymbidium* Sw., *Helianthus* L., *Magnolia* L., and *Nelumbo* Adans.) and for subspecies of *F. vesca*, *G. herbaceum*, *O. europaea*, and *O. sativa* were compared to 130 primer pairs published by Shaw et al. (2005, 2007), Scarcelli et al. (2011), Dong et al. (2012), and those designed here. Complete chloroplast genome sequences were downloaded from GenBank (accession numbers, taxonomic identity, and original publication information provided in Appendix 3) and aligned manually in Sequencher (Gene Codes Corporation, Ann Arbor, Michigan, USA). A separate file containing the primer sequences was imported and automatically assembled using the settings “dirty data” and 100% sequence similarity with a minimum overlap of 16 bp. Additional rounds of alignment were conducted with successively lower levels of sequence similarity. Primers that failed to align automatically, or that aligned incorrectly, were realigned manually whenever possible (guided by the GenBank annotations). Alignment of the two *Gossypium* sequences required inversion of a large region of one taxon (arbitrarily selected as *G. herbaceum* subsp. *africanum* (G. Watt) Vollesen) approximately corresponding to bases 115,132–135,355 in the final alignment. The *Oryza* alignment includes *O. nivara* Sharma & Shastry because it is a potential progenitor of *O. sativa* (Li et al., 2006; but see Huang et al., 2012 for an alternative view point).

As mentioned above, degenerate primers provide broader utility, but reduced amplification efficiency. If a mismatch was detected in the last five bases at the 3' end of the primer, the mismatch was inferred to be fatal (IDT, 2009). If more than three mismatches were detected within any given primer, amplification was inferred to be unsuccessful. These criteria are arbitrary but have worked for me personally and are probably more strict than necessary.

**Sequence variability within species**—The sequences of *F. vesca*, *G. herbaceum*, *O. europaea*, and *O. sativa* were examined manually to assess the variation of the 130 regions. Length of the inferred amplicon was noted along with the number of mismatched bases (aka inferred substitutions; excluding primer regions), the number of insertion/deletion (indel) events, and the number of inversions. These data provided an estimate of the utility of the regions for inferring phylogeny among closely related subspecies, and potential for application to phylogeographic studies. Shaw et al. (2014) specifically avoided these types of comparisons due to the very small number of parsimony informative characters. Sequence diversity was estimated using three criteria calculated as: (1) [(number of substitutions\*2)+(number of indels)+(number of inversions)]/amplicon length, (2) number of substitutions+indels+inversions, and (3) sequence diversity (number of substitutions/sequence length). The first criterion (criterion 1) is a weighted rank, and includes information on the number of inferred substitutions (weighted twice as heavily as the other two components), indels, and inversions. Substitutions were weighted more heavily because chloroplast indels may be more homoplasious (Kelchner and Clark, 1997), especially among closely related taxa. Inversions are often low in homoplasy (Graham et al., 2000) and thus could be weighted more heavily, but are relatively rare so weighting was not employed. The 10 most variable regions for each species were identified, as measured under each criterion. Frequency of any specific “top 10” primer pair was summed across the four species.

## RESULTS

**Primers designed here**—The 72 primers targeted noncoding regions of the chloroplast genome with amplicon sizes of 500–1800 bp. Degenerate primers were avoided because they were assumed to decrease priming efficiency, as were mismatches within the last five bases at the 3' end of the primer. Only two primers required degenerate bases: one primer with two degenerate bases and another primer with one degenerate base. None of these degeneracies were located within the last five bases. In

TABLE 1. Region, primer name, primer sequence, amplicon position, and amplicon length for plastid noncoding regions relative to the *Nicotiana tabacum* L. (GenBank Z00044.1) genome.

Region	Primer name	$T_m$ (°C) <sup>a</sup>	Primer sequence	Amplicon position	Amplicon length (bp)
<i>trnQ</i> (UUG)– <i>psbK</i> IGS	<i>trnQ</i> -IGSR	62.7	ACCCGTTGCCTTACCGCTTGG	7457–8018	562
	<i>psbK</i> -IGSR	50.9	ATCGAAAACCTGCAGCAGCTTG		
<i>psbK</i> – <i>trnS</i> (GCU) IGS	<i>psbK</i> -IGSF	47.9	CCAATCGTAGATGTTATGCC	7937–8719	783
	<i>trnS</i> _GCU-IGSF	56.1	GGAGAGATGGCTGAGTGGA		
<i>trnG</i> (UCC)– <i>atpA</i> IGS	<i>trnG</i> _UCC-IGSF	56.3	CCTTCCAAGCTAACGATGCG	10,219–10,796	577
	<i>atpA</i> -IGSF	50.3	TGGACAGGTGAAGAAATTC		
<i>atpF</i> intron	<i>atpF</i> -E2R	47.3	CTCTGTTTTCGATTATCTAATAAAAT	12,582–13,372	791
	<i>atpF</i> -E1F	48.1	AGCAACAAATCCAATAAATCT		
<i>atpF</i> – <i>atpH</i> IGS	<i>atpF</i> -E1R	46.5	TAGATTTATTGGATTTGTGTC	13,352–13,927	575
	<i>atpH</i> -IGSF	48.5	CTTTTATGGAAGCTTTAACAATTTA		
<i>atpH</i> – <i>atpI</i> IGS	<i>atpH</i> -IGSR	56.9	CCAGCAGCAATAACGGGAAGC	14,059–15,400	1341
	<i>atpI</i> -IGSF	48.2	GTGTGTTGTTCTGTTTCTTTAG		
<i>rpoC1</i> intron	<i>rpoC1</i> -intR	49.9	AAGTGGGATGCTGTATTTTC	23,004–23,976	973
	<i>rpoC1</i> -intF	49.2	ACGAAGGTATCAAATGGG		
<i>trnS</i> (UGA)– <i>psbZ</i> IGS	<i>trnS</i> _UGA-IGSR	55.0	ATCAACCACTCGGCCATC	37,209–37,620	412
	<i>psbZ</i> -IGS	45.6	AATAGCCAATTGAAAAGC		
<i>psaA</i> – <i>ycf3</i> IGS	<i>psaA</i> -IGSR	50.2	CGGCGAACGAATAATCAT	43,469–44,295	827
	<i>ycf3</i> -E3F	48.4	CCCGGTAATTATATTGAAGC		
<i>ycf3</i> intron 2	<i>ycf3</i> -E3R	54.5	ATCTCCCTGTCGAATGGC	44,362–45,193	832
	<i>ycf3</i> -E2F	53.2	GGCCGTGATCTGTCATTAC		
<i>ycf3</i> intron 1	<i>ycf3</i> -E2R	50.0	TTCCGCGTAATTTCCCTTC	45,370–46,163	794
	<i>ycf3</i> -E1F	48.1	CATTTACTATTACAGAGATGG		
<i>ycf3</i> – <i>trnS</i> (GGA) IGS	<i>ycf3</i> -E1R	45.5	ACAATTGAAAAGGTCTTATC	46,214–47,174	961
	<i>trnS</i> _GGA-IGSR	47.9	CAAAGCCTACATAGCAG		
<i>rpS4</i> – <i>trnT</i> (UGU)	<i>rpS4</i> -IGSR1	56.2	TCCTCGGTAACGCGACAT	48,065–48,570	506 max.
	<i>rpS4</i> -IGSR2	45.9	GGCTTTTTATTAGTTAGTCC		
	<i>trnT</i> _UGU-IGSF1	53.0	AGGTTAGAGCATCGCATTTG		
	<i>trnT</i> _UGU-IGSF2	47.9	GAGCATCGCATTTGTAAT		
<i>trnF</i> (GAA)– <i>ndhJ</i> IGS	<i>trnF</i> -IGSF	56.4	ATCCTCGTGTCCACCAGTTCAAA	50,277–51,024	747
	<i>ndhJ</i> -IGSF	49.3	RCCCTAATTTTATGAAATACA		
<i>ndhC</i> – <i>trnV</i> (UAC) IGS	<i>ndhC</i> -IGSR	52.9	ATCATATTCGTGAAGCAGAAACAT	52,644–53,776	1132
	<i>trnV</i> _UAC-E2F	58.3	GGTTCGAGTCCGTATAGCCCT		
<i>trnV</i> (UAC) intron	<i>trnV</i> _UAC-E2R	57.1	GGGCTATACGGACTCGAACC	53,757–54,380	624
	<i>trnV</i> _UAC-E1F	52.8	GTAGAGCACCTCGTTTACAC		
<i>trnV</i> (UAC)– <i>atpE</i> IGS	<i>trnV</i> _UAC-E1R	52.8	GTGTAACGAGGTGCTCTAC	54,361–55,032	672
	<i>atpE</i> -IGSF	56.6	AGTGACATTGATCCRCAAGAAGC		
<i>atpB</i> – <i>rbcL</i> IGS	<i>atpB</i> -IGSR	48.4	AAGTAGTAGGATTTGATTTCTCAT	56,756–57,615	859
	<i>rbcL</i> -IGSR	53.9	AGTCTCTGTTTGTGGTGACAT		
<i>rbcL</i> – <i>accD</i> IGS	<i>rbcL</i> -IGSF	58.5	GCTGCTGCTTGTGAGGTATGG	58,960–59,865	905
	<i>accD</i> -IGSR	51.1	AATTGAACCCACATTTTCCATA		
<i>accD</i> – <i>psaI</i> IGS	<i>accD</i> -IGSF	48.2	GGTAAAAGAGTAATGAACAAC	61,143–62,161	1018
	<i>psaI</i> -IGSR	49.7	ATAAAGAAGCCATTGCAATTG		
<i>psaI</i> – <i>ycf4</i> IGS	<i>psaI</i> -IGSF	51.8	CCTAGTCTTTCCGGCAAT	62,127–62,682	556
	<i>ycf4</i> -IGSR	49.5	CCCCGTTATAAGTTCTATCC		
<i>ycf4</i> – <i>ycf10</i> IGS	<i>ycf4</i> -IGSF	47.0	ATTAGCCTATTTCTTGGG	63,153–63,541	389
	<i>ycf10</i> -IGSR	51.9	GCCAGTATTCACCAA		
<i>petA</i> – <i>psbJ</i> IGS	<i>petA</i> -IGSF	50.8	GAAACAGTTTGAGAAGGTCA	65,255–66,388	1133
	<i>psbJ</i> -IGSF	55.8	ATCCCGCATTGGGCTCATC		
<i>petL</i> – <i>psaJ</i> IGS	<i>petL</i> -IGSF	48.4	TCTATTAGCGGCTTTAACTATA	68,322–69,671	1350
	<i>psaJ</i> -IGSR	52.4	GCATCCGGGAATAACGA		
<i>psaJ</i> – <i>rpL20</i> IGS	<i>psaJ</i> -IGSF	46.5	ATGCGAGATCTAAAAACATA	69,565–71,404	1840
	<i>rpL20</i> -IGSF	46.6	CAGAATTAACGGGGATATA		
<i>rpL20</i> – <i>rpS12</i> IGS	<i>rpL20</i> -IGSR	51.3	CGTCTCCGAGCTATATATCC	71,372–72,319	947
	<i>rpS12</i> -IGSF	47.3	CAACTTATTAGAACAACAAGAC		
<i>clpP</i> intron 2	<i>clpP</i> -E3R	51.6	TTGCCTGTTCTTTGTACATAAAC	72,573–73,466	893
	<i>clpP</i> -E2F	50.9	GCTATTTATGAGCTATGCAA		
<i>clpP</i> intron 1	<i>clpP</i> -E2R	50.9	TTGCATAGCGCTATAAATAGC	73,446–74,451	1005
	<i>clpP</i> -E1F	54.9	TTGGGTTGACATATAGTCCGAC		
<i>clpP</i> – <i>psbB</i> IGS	<i>clpP</i> E1-IGSR	52.2	AGGGACTTTTGGAAACACC	74,481–74,970	490
	<i>psbB</i> -IGSR	51.5	ATACCAAGGCAACCCAT		
<i>psbH</i> – <i>petB</i> IGS	<i>psbH</i> -IGSF	48.5	AACTACTCCTTTGATGGG	77,214–78,377	1163
	<i>petB</i> -E2R	44.1	TAGTAAAAGTCATAGCAAA		
<i>petB</i> – <i>petD</i> IGS	<i>petBE2</i> -IGSF	50.8	ATGCACCTTCCAATGATACG	78,805–79,760	956
	<i>petD</i> -E2R	59.8	CCGAGGGAACCGGACAT		
<i>rpS3</i> – <i>rpS19</i> IGS	<i>rpS3</i> -IGSR	50.5	CAGTCTGAAACCAAGTGG	85,863–86,504	642
	<i>rpS19</i> -IGSF	45.9	TTTATATAACGGATAGTAGTGGT		
<i>ccsA</i> – <i>ndhD</i> IGS	<i>ccsA</i> -IGSF	45.5	ATGATATTTTCAACCTTAGA	116,344–117,614	1271
	<i>ndhD</i> -IGSF	43.6	CCGTAATAGGTATTGGTAT		



TABLE 1. Continued.

Region	Primer name	$T_m$ (°C) <sup>a</sup>	Primer sequence	Amplicon position	Amplicon length (bp)
<i>psaC-ndhE</i> IGS	psaC-IGSR	44.9	TCCTATACACGTATCATAAA	119,351–119,713	363
	ndhE-IGSF	42.4	TTCATCAATTTATCGTAAAC		
<i>ndhE-ndhI</i> IGS	ndhE-IGSR	45.6	GAAAATAAATAGGCACTCAA	119,912–121,251	1340
	ndhI-IGSF	46.9	CAATGACCGAAGAATATGA		
<i>rpS15-ycf1</i> IGS	rpS15-IGSR	47.7	GCAATTCTAAATGTGAAGTAAG	125,374–126,001	628
	ycf1-IGSR	45.6	ATTATCGATTAGAAGATTAGC		

<sup>a</sup>Melting temperature ( $T_m$ ) based on 50 mM NaCl solution.

contrast, 17 of the Scarcelli et al. (2011) primers have at least one degenerate base in the last five bases at the 3' end of the primer, and so are assumed to fail for at least some taxa.

**Primer evaluation**—Three of the four sets of primers examined here were equally likely to amplify target chloroplast regions (81–85% should work; see Table 2). The Dong et al. (2012) primers were least likely to work based on the 12 species examined here (65% on average) and were particularly poorly matched to the *Oryza* genome (29% amplification success predicted), and only moderately suited for *Amborella* (52%), *Cymbidium* (52%), and *Helianthus* (57%). However, the Dong et al. (2012) primer pair *trnH-psbA* was not expected to work on any of the target species, possibly due, in part, to an extra “A” near the 3' end of the published sequence for the *trnH* primer. The primers designed here were poorly matched to three of the four monocots (*Cymbidium*, *Oryza*, and *Canna*; 61%, 64%, and 67%, respectively), despite being a good match for *Acorus* (81%). Scarcelli et al. (2011) primers were designed with monocots in mind and did an exceptional job matching the monocot genomes examined here, with amplification success ranging from 82–97%. They were almost equally good for the dicots examined here, with amplification success of 72–93%. The Shaw et al. (2005, 2007) primers were useful across the angiosperm phylogeny, with all anticipated amplification success percentages above 78%.

On average, the Shaw et al. (2005, 2007) and Scarcelli et al. (2011) primers are more degenerate, yet they were only slightly more likely to amplify the target sequences than the nondegenerate primers designed here, at least for nonmonocot taxa. With so many different primers available, most regions could be amplified in almost all target taxa provided an appropriate primer pair was selected. Indeed, many primer pairs should work in all 12 species examined here. Details of the inferred priming success are provided in Appendix S1, and species-specific notes on primer/sequence mismatches are provided in Appendix S2.

**Primer utility × sequence variability**—Shaw et al. (2014) conveniently summarized sequence variability across the chloroplast genome including the identification of the 13 fastest-evolving regions for six taxonomic groups (magnoliids, monocots, eurosids I, eurosids II, euasterids I, and euasterids II). Summing across these major groups, 28 different regions were identified as the most variable. Primers to amplify those 28 regions are detailed in Table 3, along with the Shaw et al. (2014) rank for each region (in bold typeface above each primer region), for each taxon examined here. Multiple primer pairs are available for each of the 28 regions except the *trnT-trnL* (Shaw et al., 2005 only), *ycf4-ycf10* (or *cemA*; current study only), and *ndhD-psaC* (none of the publications examined). The *ndhD-psaC* region was ranked 10th fastest for eurosids I, but as there are no

primers to be evaluated this region will not be discussed further. Primers are available for each of the remaining 27 regions.

Among the basal dicot grade (*Amborella* and *Magnolia*), successful primers are available for all 27 regions. Primer selection is more challenging for *Amborella* than for *Magnolia*. The top ranked region was the *rpl32-trnL* intergenic spacer (IGS). Shaw et al. (2007) primers will work for both taxa; Dong et al. (2012) primers will not. In contrast, *rps16-trnQ*, the second highest ranked region, has three sets of primers available (Shaw et al., 2007; Scarcelli et al., 2011; and Dong et al., 2012), all of which should work.

Among the monocots sampled (*Acorus*, *Cymbidium*, *Oryza*, and *Canna*), *Acorus* was the least difficult sequence to match and *Oryza* the most difficult. Structural rearrangements are the primary reason for failure to amplify across all available primers (e.g., *rbcL-accD* in *Oryza* and *petA-psbJ* in *Cymbidium*). One region cannot be amplified in *Acorus*—the *accD-psaI* IGS, despite the availability of four different primer pairs. In all, four regions cannot be amplified in *Cymbidium* with the primers studied here: *petN-psbM*, *psbM-trnD*, *atpB-rbcL*, and *petA-psbJ*. The *ndhA* region can be amplified in only some species of *Cymbidium* due to fatal substitutions in some species for all three primer pairs evaluated here. In *Oryza*, the *trnS[GCU]-trnG[GCC]*, *trnT-psbD*, *rbcL-accD*, *accD-psaI*, and *rps15-ycf1* cannot be amplified using any primer pair. In *Canna*, *ndhF-rpl32* will not amplify with either of the available primer pairs. Unfortunately, according to Shaw et al. (2014), *ndhF-rpl32* is the most variable and *psbM-trnD* is the third most variable region for monocots.

Basal eudicots were not evaluated by Shaw et al. (2014) in detail, so direct comparisons cannot be made here. Fortunately, at least one primer pair was successful for each of the 27 fastest-evolving regions, with the exception of the *ycf4-ycf10* region. The only available primers for this region were designed here, and they will not work for *Ceratophyllum*. In general, *Ceratophyllum* was more difficult to match than was *Nelumbo*.

Shaw et al. (2014) detailed variability of higher eudicots for four major groups: eurosids I, eurosids II, euasterids I, and euasterids II. Only a single species representing each group was included here. *Fragaria* (eurosids I) could not be amplified for a single region, the *ycf4-ycf10* IGS. According to Shaw et al. (2014), the fastest region for this clade was the *ndhA* intron. Both the Shaw et al. (2007) and Scarcelli et al. (2011) primers should work, but the Dong et al. (2012) primers will not. The second fastest region was the *trnS[GCU]-trnG[GCC]*, which should amplify with any of the primer pairs (Shaw et al., 2005; Scarcelli et al., 2011; or Dong et al., 2012).

The sole representative of eurosids II and euasterids I (*Gossypium* and *Olea*, respectively) could successfully be amplified by at least one pair of primers studied here. The fastest region

TABLE 2. Summary of amplification success probability for 130 pairs of chloroplast primers.

Publication <sup>a</sup>	No. of regions	Average % ampl.	Basal dicot grade/Magnoliids				Monocots			Basal eudicot grade			Eurosid I		Eurosid II		Euasterid I		Euasterid II	
			<i>Amborella</i>	<i>Magnolia</i>	<i>Acorus</i>	<i>Cymbidium</i>	<i>Oryza</i>	<i>Canna</i>	<i>Ceratophyllum</i>	<i>Nelumbo</i>	<i>Fragaria</i>	<i>Gossypium</i>	<i>Olea</i>	<i>Helianthus</i>						
Dong	21	65	11 (52%)	16 (76%)	14 (67%)	11 (52%)	6 (29%)	15 (71%)	17 (81%)	16 (76%)	14 (67%)	17 (81%)	12 (57%)							
Current study	36	81	31 (86%)	32 (89%)	29 (81%)	22 (61%)	23 (64%)	32 (89%)	32 (89%)	28 (78%)	33 (92%)	31 (86%)	32 (89%)							
Scarcelli	99	83	71 (72%)	92 (93%)	96 (97%)	92 (93%)	81 (82%)	71 (72%)	88 (89%)	73 (74%)	80 (81%)	79 (80%)	75 (76%)							
Shaw	33	85	27 (82%)	31 (94%)	29 (88%)	26 (79%)	29 (88%)	28 (85%)	28 (85%)	27 (82%)	27 (82%)	29 (88%)	28 (85%)							

<sup>a</sup>Dong et al., 2011; Scarcelli et al., 2011; Shaw et al., 2005, 2007.

for eurosids II was the *ndhF-rpl32* IGS. The Shaw et al. (2007) primer pair should work, but the Scarcelli et al. (2011) primer pair likely will not. The second most variable region was the *psbZ-trnG* IGS. For this region, both the Scarcelli et al. (2011) and Dong et al. (2012) primers should work, but the Shaw et al. (2005; as *trnfM-trnS*) primers will not. In euasterids I, the fastest region was the *rps16-trnQ* IGS. For *Olea*, the Shaw et al. (2007) and Scarcelli et al. (2011) primers should work, but not so the Dong et al. (2012) primers. The next-fastest region was the *rpl32-trnL* IGS. Both the Shaw et al. (2007) and Dong et al. (2012) primers should work.

Primer failure in *Helianthus* (euasterids II) was primarily due to structural rearrangements (e.g., *trnS*[GCU]-*trnG*[GCC], *rpoB-trnC*, *trnE-trnT*, *rbcL-accD*). *rpl32-trnL* IGS was the fastest region according to Shaw et al. (2014), and either the Shaw et al. (2007) or Dong et al. (2012) primers should successfully amplify this region. The adjacent *ndhF-rpl32* IGS was the second most variable region. Both the Shaw et al. (2007) or the Scarcelli et al. (2011) primers should work.

**Subspecific sequence variability**—Intraspecific sequence variation was evaluated in four species: *F. vesca*, *G. herbaceum*, *O. europaea*, and *O. sativa*. This represents a tiny fraction of angiosperm diversity, but is the first analysis of subspecific diversity across the entire chloroplast genome for multiple species, in the context of available primer resources. Appendix S3 identifies the fastest-evolving regions among the four species, under three different criteria. On average, only five inversions per chloroplast genome were detected here and the distribution across species was very different. *Gossypium* and *Oryza* each had 10 inversions, *Fragaria* none, and *Olea* only one. Details of subspecific comparisons for all regions are provided in Appendix S2.

No single genic region was identified as the top 10 fastest for all four species. Pooling data across all three criteria, the most frequently identified genic region was the *psbZ-trnfM* IGS with eight occurrences out of a maximum of 12 possible, followed by the *trnS*(GCU)-*trnG*(GCC) IGS, with six occurrences, *rps16-trnQ* IGS and *trnT*(GGU)-*psbD* IGS each with five, and *rps12-psbB* IGS and *rps4-trnT*(UGU) IGS each with four occurrences. Data for individual species have limited general application, but are provided below.

*Oryza sativa*, the only monocot in this comparison, showed highest variation, based on rank, for *clpP-psbB* (0.0195, 924 bp), *atpB-rbcL* (0.0168, 1070 bp), and *psbM-trnD*(GUC) (0.0150, 523 bp). Two of the same regions were identified as fastest under criterion 2, *atpB-rbcL* (12 characters, 1070 bp) and *clpP-psbB* (11 characters, 924 bp), plus *rbcL-accD* (13 characters, 1824 bp). Sequence divergence was highest in and around the *clpP* region including what would be the *clpP* intron 2 (1.9455%, 257 bp), *clpP* intron 1 (1.0050%, 199 bp), and *clpP-psbB* (0.7576%, 924 bp). In contrast, the three fastest regions per Shaw et al. (2014) for monocots were *ndhF-rpl32* (rank 1), *ndhC-trnV* (rank 2), and *psbM-trnD* (rank 3).

The highest variation for *Fragaria* under criterion 1 was for *trnW*(CCA)-*psaJ* (0.0101, 789 bp), *trnT*(GGU)-*psbD* (0.0098, 1527 bp), and *trnP*(UGG)-*rps18* (0.0090, 1563 bp). Under criterion 2: *trnT*(GGU)-*psbD* (eight characters; 1527 bp), *trnP*(UGG)-*rps18* (eight characters, 1563 bp), and *petN-trnD* (seven characters, 2504 bp). Under criterion 3, the top three regions were *trnT*(GGU)-*psbD* (0.4584%, 1527 bp), *psbB-psbH* (0.4451%, 674 bp), and *rps4-trnT*(UGU) (0.4435%, 451 bp). Shaw et al. (2014) eurosids I top three regions were *ndhA* intron

TABLE 3. Amplification success prediction for the 28 fastest Shaw et al. (2014) regions.<sup>a</sup>

Approx. <i>Nicotiana</i> order	Genomic region	Publication <sup>b</sup>	Basal dicot grade/Magnoliids										Monocots		Basal eudicot grade		Eurosid I		Eurosid II		Euasterids I		Euasterids II		Average	
			<i>Amborella</i>	<i>Magnolia</i>	<i>Acorus</i>	<i>Cymbidium</i>	<i>Oryza</i>	<i>Canna</i>	<i>Ceratophyllum</i>	<i>Nelumbo</i>	<i>Fragaria</i>	<i>Gossypium</i>	<i>Olea</i>	<i>Helianthus</i>	<i>Olea</i>	<i>Helianthus</i>										
1	<i>trnH-psbA</i> IGS	Dong et al.	NO**	NO**	NO	NO**	NO	NO	NO**	NO	NO	NO	NO	NO	NO	NO	NO	NO	NO	NO	NO	NO	NO	NO	NO	0%
	<i>trnH-psbA</i> IGS	Scarcelli et al.	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	83%
	<i>trnH-psbA</i> IGS	Shaw et al.	YES	YES	YES	NO	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	92%
5	<i>matK</i> exon	Dong et al.	YES	YES	YES	NO	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	92%
	<i>trnK</i> (including <i>matK</i> )	Scarcelli et al.	YES	YES	YES	YES*	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	100%
7	<i>trnK-rps16</i> IGS	Scarcelli et al.	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	92%
	<i>trnK-rps16</i>	Shaw et al.	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	100%
8	<i>trnK-3' rps16</i>	Dong et al.	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	83%
	<i>rps16</i> intron	Shaw et al.	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	100%
9	<i>rps16</i> intron	Scarcelli et al.	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	83%
	<i>rps16</i> intron	Shaw et al.	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	100%
12	<i>rps16-trnQ</i> IGS	Dong et al.	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	58%
	<i>rps16-trnQ</i>	Scarcelli et al.	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	100%
16	<i>5' rps16-trnQ</i>	Shaw et al.	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	100%
	<i>trnS-trnG</i> IGS	Dong et al.	NO	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	75%
18	<i>trnS-trnG</i> (and intron)	Scarcelli et al.	NO	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	75%
	<i>trnS-trnG</i>	Shaw et al.	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	83%
26	<i>atpF</i> intron	Prince (here)	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	100%
	<i>atpH</i> intron/exon	Scarcelli et al.	NO	YES	YES	NO	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	67%
29-31	<i>atpH-atpI</i> IGS	Dong et al.	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	92%
	<i>atpH-atpI</i>	Prince (here)	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	100%
32	<i>atpH-atpI</i>	Shaw et al.	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	100%
	<i>rpoB-trnC</i> IGS	Dong et al.	YES	YES	YES	NO	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	67%
33	<i>rpoB-trnC</i>	Scarcelli et al.	NO	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	75%
	<i>rpoB-trnC</i>	Shaw et al.	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	83%
34	<i>petN-psbM</i> IGS	Scarcelli et al.	YES	YES	YES	NO	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	83%
	<i>petN-trnD</i>	Dong et al.	NO	NO	NO	NO	NO	NO	NO	NO	NO	NO	NO	NO	NO	NO	NO	NO	NO	NO	NO	NO	NO	NO	NO	17%
38-41	<i>ycf6-psbM</i>	Shaw et al.	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	75%
	<i>psbM-trnD</i> IGS	Dong et al.	YES	YES	YES	NO	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	83%
34	<i>psbM-trnD</i>	Shaw et al.	NO	NO	YES	NO	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	67%
	<i>trnE-trnT</i> IGS	Scarcelli et al.	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	83%
34	<i>trnD-trnT</i>	Shaw et al.	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	92%
	<i>trnT-psbD</i> IGS	Dong et al.	NO	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	67%
38-41	<i>trnT-psbD</i>	Scarcelli et al.	NO	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	83%
	<i>trnT-psbD</i>	Shaw et al.	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	92%
38-41	<i>psbZ-trnG</i> IGS	Dong et al.	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	83%
	<i>trnS-trnG</i>	Shaw et al.	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	67%

TABLE 3. Continued.

Approx. <i>Nicotiana</i> order	Genomic region	Publication <sup>b</sup>	Basal dicot grade/Magnoliids				Monocots			Basal eudicot grade			Eurosid I		Eurosid II		Average	
			<i>Amborella</i>	<i>Magnolia</i>	<i>Acorus</i>	<i>Cymbidium</i>	<i>Oryza</i>	<i>Canna</i>	<i>Ceratophyllum</i>	<i>Nelumbo</i>	<i>Fragaria</i>	<i>Gossypium</i>	<i>Olea</i>	<i>Helianthus</i>				
50	<i>psbZ-trnFM</i> <i>trnT-trnL</i> IGS	Scarcelli et al.	YES 11 <sup>c</sup>	YES	YES 9 <sup>c</sup>	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	100%
55	<i>trnT-trnL</i> <i>ndhC-trnV</i> <i>ndhC-trnV</i> <i>ndhC-trnV</i> <i>ndhC-trnV</i> <i>ndhC-trnV</i>	Shaw et al. Dong et al. Prince (here) Scarcelli et al. Shaw et al.	YES 5 <sup>c</sup> YES YES YES YES	YES YES YES YES YES YES	YES 2 <sup>c</sup> YES YES YES YES	YES YES YES YES YES YES	YES YES YES YES YES YES	YES YES YES YES YES YES	YES YES YES YES YES YES	YES YES YES YES YES YES	YES YES YES YES YES YES	YES YES YES YES YES YES	YES YES YES YES YES YES	YES YES YES YES YES YES	YES YES YES YES YES YES	YES YES YES YES YES YES	YES YES YES YES YES YES	100% 100% 100% 92% 100% 92%
60	<i>atpB-rbcL</i> IGS	Prince (here)	YES	YES	YES	NO	YES	YES	YES	YES	YES	YES	YES	NO	YES	YES	YES	75%
62	<i>atpB-rbcL</i> <i>rbcL-accD</i> IGS	Scarcelli et al.	NO	YES	YES 12 <sup>c</sup>	NO	YES	YES	YES*	YES	YES	YES	YES	YES	YES	NO	YES	67%
64	<i>rbcL-accD</i> <i>rbcL-accD</i> <i>accD-psal</i> IGS	Dong et al. Prince (here) Scarcelli et al.	NO YES NO	YES YES NO	YES NO NO	YES NO NO	YES NO YES	YES NO NO	YES NO NO	YES NO NO	YES NO NO	YES NO NO	YES NO NO	YES NO NO	YES NO NO	YES NO NO	YES NO NO	75% 42% 8%
67	<i>accD-psal</i> <i>accD-psal</i> <i>accD-psal</i> <i>accD-psal</i> <i>yef4-cema</i> ( <i>yef10</i> ) IGS	Dong et al. Prince (here) Scarcelli et al. Shaw et al.	NO NO NO YES	YES YES YES YES	NO NO NO NO	NO NO NO YES	NO NO NO YES	NO NO NO YES	NO NO NO YES	NO NO NO YES	NO NO NO YES	NO NO NO YES	NO NO NO YES	NO NO NO YES	NO NO NO YES	NO NO NO YES	NO NO NO YES	67% 67% 58% 75%
70	<i>yef4-yef10</i> <i>petA-psbJ</i> IGS	Prince (here)	YES 6 <sup>c</sup>	YES	YES 6 <sup>c</sup>	YES	YES	YES	NO	NO	YES	YES	YES	YES	YES	YES	YES	75%
72	<i>petA-psbJ</i> <i>petA-psbJ</i> <i>psbE-petL</i> IGS	Dong et al. Prince (here) Shaw et al.	YES YES YES	YES YES YES	YES YES YES	NO NO NO	YES YES YES	YES YES YES	YES YES YES	YES YES YES	YES YES YES	YES YES YES	YES YES YES	YES YES YES	YES YES YES	YES YES YES	YES YES YES	75% 67% 75%
76, 77	<i>psbE-petL</i> <i>psa-rpl33</i> IGS	Dong et al. Shaw et al.	NO YES	NO YES	NO YES	NO YES	NO YES	NO YES	YES YES	YES YES	YES YES	YES YES	YES YES	YES YES	YES YES	YES YES	YES YES	50% 100%
116	<i>trnP-rps18</i> <i>psa-rpl20</i> <i>ndhF-rpl32</i> IGS	Scarcelli et al. Prince (here)	YES NO	YES YES	YES NO	YES NO	YES NO	YES NO	YES YES	YES YES	YES YES	YES YES	YES YES	YES YES	YES YES	YES YES	YES YES	92% 58%
118	<i>ndhF-rpl32</i> <i>ndhF-rpl32</i> <i>rpl32-trnL</i> IGS	Scarcelli et al. Shaw et al.	YES NO	YES YES	YES YES	YES YES	YES YES	YES YES	YES NO	YES NO	YES YES	YES YES	YES YES	YES YES	YES YES	YES YES	YES YES	83% 67%
121, 5 127	<i>rpl32-trnL</i> IGS <i>rpl32-trnL</i> <i>ndhD-psaC</i> IGS	Dong et al. Shaw et al.	NO YES	YES YES	YES YES	NO YES	YES YES	YES YES	YES YES	YES YES	YES YES	YES YES	YES YES	YES YES	YES YES	YES YES	YES YES	83% 92%
129	<i>ndhA</i> intron <i>ndhA</i> intron <i>ndhA</i> intron <i>ndhA</i> intron <i>rps15-ycf1</i> IGS <i>rps15-yef1</i> <i>rps15-yef1</i>	Dong et al. Scarcelli et al. Shaw et al. Prince (here) Scarcelli et al.	NO YES YES YES	NO YES YES YES	NO YES YES YES	YES* YES* YES* YES	YES YES YES YES	YES YES YES YES	NO YES YES YES	NO YES YES YES	NO YES YES YES	NO YES YES YES	NO YES YES YES	NO YES YES YES	NO YES YES YES	NO YES YES YES	NO YES YES YES	25% 100% 92% 83% 67%

<sup>a</sup> YES\* = will not work for at least one species in the genus; NO\*\* = will work if psbA primer is synthesized with one fewer A at the 3' end.  
<sup>b</sup> Shaw et al., 2005, 2007; Scarcelli et al., 2011; Dong et al., 2012.  
<sup>c</sup> Shaw et al. (2014) rank for the region within the specified taxonomic group.



(rank 1), *trnS*(GCU)-*trnG*(GCC) (rank 2), and *rps16* intron (rank 3).

In *Gossypium*, the most informative regions under criterion 1 were *psbZ-trnfM*(CAU) (0.0534, 1179 bp), *trnH*(GUG)-*psbA* (0.0444, 496 bp), and *rps4-trnT*(UGU) (0.0425, 635 bp). Criterion 2 fastest regions were *trnS*(UGA)-*trnG*(GCC) with 39 variable characters over 1673 bp, followed by *psbZ-trnfM*(CAU) with 37 characters for 1179 bp, and *trnT*(UGU)-*trnL*(UAA) with 33 characters over 1470 bp. Sequence divergence (criterion 3) was highest for *psbZ-trnfM*(CAU) (2.2053%, 1179 bp), then *trnS*(UGA)-*trnG*(GCC) (1.6736%, 1673 bp), and finally the *rps16* intron (1.6181%, 927 bp). Eurosids II top three regions for Shaw et al. (2014) were *ndhF-rpl32* (rank 1), *psbZ-trnG* (rank 2), and *trnT-trnL* (rank 3).

For *Olea*, the most informative regions under criterion 1 were *psbC-psbZ* (0.0411, 1045 bp), *trnS*(UGA)-*trnfM* (0.0333, 1203 bp), and *clpP* intron 2 (0.0313, 702 bp). The highest number of variable characters (criterion 2) were found in *rps16-trnQ* (29 characters, 2739 bp), *psbC-psbZ* (22 characters, 1045 bp), and *trnS*(UGA)-*trnfM* (21 characters, 1203 bp). Criterion 3 (percent sequence divergence) was highest in the same three regions as under criterion 1: *psbC-psbZ* (2.0096%, 1045 bp), *trnS*(UGA)-*trnfM* (1.5794%, 1203 bp), and *clpP* intron 2 (1.4245%, 702 bp). Shaw et al. (2014) euasterids I top three included *rps16-trnQ* (rank 1), *rpl32-trnL* (rank 2), and *ndhC-trnV* (rank 3).

## DISCUSSION

A large number of “universal” primers have been published for amplification of various chloroplast regions. Some are more degenerate than others, presumably to be more widely applicable. Degeneracy is not required, however, and may not lead to greater success in the laboratory. On the other hand, nondegenerate primers with poor fit are likely to fail, and some primers published as “universal” are not necessarily so. The universal barcoding primers of Dong et al. (2012) were the least likely to be useful across the 12 taxa examined here, with an average success rate of 65%, and a very poor 29% success rate in *Oryza*. In contrast, the primers designed by Scarcelli et al. (2011) specifically for monocots were exceedingly well-matched to the monocots sampled (97% in *Acorus*, 93% in *Cymbidium*, 92% in *Oryza*, and 88% in *Canna*), and a good match across all angiosperms.

Unlike previous analyses, this study used published genomes and primer sequences to infer the likelihood of amplification success. Only a small number of published primers were evaluated, and additional primers will be added to future analyses. Indeed, as mentioned in the introduction, Ebert and Peakall (2009) and Dong et al. (2013) have primers that could be evaluated as well as those of Doorduyn et al. (2011) designed for species of Asteraceae. The evaluation conducted here shows parallels to prior studies in that general conclusions or recommendations are difficult to distill. For each region, there may be a number of primer pair options. Which primer pair is best is highly variable and depends upon the taxon being investigated. Scarcelli et al. (2011) primers are the best option for monocots in general, but will fail in specific combinations (e.g., *trnH-psbA* for *Canna*, *atpF* intron/exon for *Cymbidium*, and *trnD-trnT* for *Oryza*). Dong et al. (2012) primers are generally less successful, but they are the only primers that will work for *psbM-trnD* in *Amborella* and *Magnolia*. In several instances, a primer will work for some, but not all species in a genus, like

the Scarcelli et al. (2011) *matK* primers in *Cymbidium* or the *trnK-rps16* primers in *Helianthus*. Table 3 provides a quick summary of primer match for the top regions according to Shaw et al. (2014).

Prior studies have done an excellent job assessing variability of various noncoding regions across a diversity of angiosperms, particularly the recent work of Shaw et al. (2014). Those studies focused on infrageneric or even intergeneric comparisons. Here I compare sequence variability within species to see if the same markers are identified as the most variable, under slightly different criteria. This comparison was specifically avoided by Shaw et al. (2014) due to the small number of variable characters. The fastest regions identified here for *Oryza* were (depending upon criterion) *clpP-psbB*, *atpB-rbcL*, *psbM-trnD*, and *rbcL-accD*. In contrast, Shaw identified *ndhF-rpl32*, *ndhC-trnV*, and *psbM-trnD* as the fastest regions for monocots, with only one region of overlap between the two. For *Fragaria* (euasterids I), the list has no overlap at all. *Olea* (euasterids II) and *Gossypium* (euasterids I) each only overlap for a single region between the two studies. The lack of consensus over which region is the most variable at lower taxonomic levels has been pointed out by a number of papers including Särkinen and George (2013) for *Solanum*, and for 19 species pairs as demonstrated by Shaw et al. (2014). The comparison made here only adds to the argument that there is an acute need for additional comparative information.

Shaw et al. (2014) provided a solid foundation for which markers evolve the most quickly in major angiosperm clades, yet the fastest regions identified here for subspecies comparisons share little overlap with Shaw’s regions. This finding suggests the need for a thorough exploration of markers prior to undertaking a large comparative sequencing project. The methods employed here to examine expected primer utility can easily be applied to any taxon, provided complete chloroplast genomic data are available. When complete genome data are lacking, the results presented here can provide a rough estimate of the “best primers,” but this remains a work in progress.

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APPENDIX 1. Complete chloroplast genome sequences used to design universal flowering plant primers for 36 plastid noncoding regions. Format: Organism; GenBank number and version; publication.

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**Basal Dicot Grade:**

1. *Amborella trichopoda* Baill.; NC\_005086.1; Goremykin et al., 2003.

**Monocots:**

2. *Oryza nivara* Sharma & Shastry; NC\_005973.1; Shahid Masood et al., 2004.
3. *Oryza sativa* L.; NC\_001320.1; Hiratsuka et al., 1989.
4. *Saccharum* hybrid; NC\_005878.2; Calsa et al., 2004.
5. *Saccharum officinarum* L.; NC\_006084.1; Asano et al., 2004.
6. *Triticum aestivum* L.; NC\_002762.1; Ogihara et al., 2002.
7. *Zea mays* L.; NC\_001666.2; Maier et al., 1995.

**Eudicots:**

8. *Arabidopsis thaliana* (L.) Heynh.; NC\_000932.1; Sato et al., 1999.
  9. *Atropa belladonna* L.; NC\_004561.1; Schmitz-Linneweber et al., 2002.
  10. *Calycanthus floridus* L. var. *glaucus* (Willd.) Torr. & A. Gray; NC\_004993.1; Goremykin et al., unpublished (Goremykin, V., K. Hirsch-Ernst, S. Wolf, and F. Hellwig. Complete structure of the chloroplast genome of *Calycanthus fertilis*. Direct GenBank submission 9 July 2003).
  11. *Lotus japonicus* (Regel) K. Larsen; AP002983.1; Kato et al., 2000.
  12. *Medicago truncatula* Gaertn.; NC\_003119.6; Lin et al., unpublished (Lin, S., H. Wu, H. Jia, P. Zhang, R. Dixon, G. May, R. Gonzales, and B. A. Roe. *Medicago truncatula* variety Jema Long A-17 chloroplast, complete sequence. Direct GenBank submission 31 August 2001).
  13. *Nicotiana tabacum* L.; Z00044.1; Shinozaki et al., 1986. Note: this sequence has been updated since this article was published.
  14. *Nymphaea alba* L.; NC\_006050.1; Goremykin et al., 2004.
  15. *Oenothera elata* Kunth subsp. *hookeri* (Torr. & A. Gray) W. Dietr. & W. L. Wagner; NC\_002693.1; Hupfer et al., 2000. Note: this sequence has been updated since this article was published.
  16. *Spinacia oleracea* L.; NC\_002202.1; Schmitz-Linneweber et al., 2001.
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APPENDIX 2. Comparison of chloroplast regions with published primer pairs.

Approx. <i>Nicotiana</i> order <sup>a</sup>	Primary type	Location <sup>b</sup>	Genomic region	Shaw et al., 2005, 2007	Ebert and Peakall, 2009	Scarcelli et al., 2011	Dong et al., 2012	Dong et al., 2013	Current study
1	IGS	LSC	<i>trnH</i> (GUG)- <i>psbA</i>	✓		✓	✓	✓	
2	Exon	LSC	<i>psbA</i> exon			✓		✓	
3	IGS	LSC	<i>psbA-trnK</i> (UUU)	✓		✓		✓	
4	IGS	LSC	3' <i>trnK</i> (UUU)- <i>matK</i>	✓	✓				
5	Exon	LSC	<i>matK</i> exon			✓	*	✓	
6	IGS	LSC	<i>matK-trnK5'</i>	✓	✓			✓	
7	IGS	LSC	<i>trnK</i> (UUU)- <i>rps16</i>	✓	✓	✓		✓	
8	Intron	LSC	<i>rps16</i> intron	✓	✓	✓		✓	
9	IGS	LSC	<i>rps16-trnQ</i> (UUG)	✓	✓	✓	✓	✓	
10	IGS	LSC	<i>trnQ</i> (UUG)- <i>psbK</i>		✓	✓		*	✓
11	IGS	LSC	<i>psbK-trnS</i> (GCU)		✓	✓		*	✓
12	IGS	LSC	<i>trnS</i> (GCU)- <i>trnG</i> (UCC) and intron	✓	✓	✓	✓	*	
13	Intron	LSC	<i>trnG</i> (UCC) intron	✓	✓	✓			
14	IGS	LSC	<i>trnG</i> (UCC)- <i>atpA</i>		*	✓		✓	✓
15	Exon	LSC	<i>atpA</i> exon			✓		✓	
16	IGS	LSC	<i>atpA-atpF</i>		✓			✓	
17	Intron	LSC	<i>atpF</i> intron		✓	✓		✓	✓
18	IGS	LSC	<i>atpF-atpH</i>		✓	✓		✓	✓
19	IGS	LSC	<i>atpH-atpI</i>	✓	✓	✓	✓	✓	
20	Exon	LSC	<i>atpI</i> exon			✓		✓	
21	IGS	LSC	<i>atpI-rps2</i>		✓	✓		✓	
22	Exon	LSC	<i>rps2</i> exon			✓		*	
23	IGS	LSC	<i>rps2-rpoC2</i>		✓	✓			
24	IGS	LSC	<i>rpoC2-rpoC1</i>			✓		*	
25	Intron	LSC	<i>rpoC1</i> intron/exon 1		✓	✓		✓	✓
26	Exon	LSC	<i>rpoC1</i> exon 2			✓		✓	
27	Exon	LSC	<i>rpoB2</i> exon					✓	
28	IGS	LSC	<i>rpoB-trnC</i> (GCU)	✓	✓	✓	✓	✓	
29	IGS	LSC	<i>trnC</i> (GCU)- <i>ycf6</i>	✓					
30	IGS	LSC	<i>trnC</i> (GCU)- <i>petN</i>		✓	✓		✓	
31	IGS	LSC	<i>petN-trnD</i>			✓			
32	IGS	LSC	<i>petN-psbM</i>		✓		✓	✓	
33	IGS	LSC	<i>ycf6-psbM</i>	✓					
34	IGS	LSC	<i>psbM-trnD</i> (GUC)	✓	✓		✓	✓	
35	IGS	LSC	<i>trnD</i> (GUC)- <i>trnT</i> (GGU)		✓	✓		✓	
36	IGS	LSC	<i>trnT</i> (GGU)- <i>psbD</i>	✓	✓	✓	✓	✓	
37	Exon	LSC	<i>psbD</i> exon			✓		✓	
38	Exon	LSC	<i>psbC</i> exon			✓		✓	
39	IGS	LSC	<i>psbC-psbZ</i>		✓	✓		*	
40	IGS	LSC	<i>trnS</i> (UGA)- <i>trnG</i> (GCC)				✓		
41	IGS	LSC	<i>trnG</i> (GCC)- <i>rpS14</i>		✓				
42	IGS	LSC	<i>trnS</i> (UGA)- <i>trnJ</i> M	✓					
43	IGS	LSC	<i>trnS</i> (UGA)- <i>psbZ</i>						✓
44	IGS	LSC	<i>psbZ-trnJ</i> M(CAU)			✓			
45	IGS	LSC	<i>trnJ</i> M(CAU)- <i>psaB</i>			✓			
46	Exon	LSC	<i>psaB</i> exon					✓	
47	Exon	LSC	<i>psaA</i> exon					✓	
48	IGS	LSC	<i>psaA-ycf3</i>		✓	✓		✓	✓
49	Intron	LSC	<i>ycf3</i> intron 2		✓	✓		✓	✓
50	Intron	LSC	<i>ycf3</i> intron 1		✓	✓		✓	✓
51	IGS	LSC	<i>ycf3-trnS</i> (GGA)		✓				✓
52	IGS	LSC	<i>ycf3-rps4</i>			✓		✓	
53	IGS	LSC	<i>trnS</i> (GGA)- <i>rpS4-trnT</i> (UGU)	✓					
54	IGS	LSC	<i>rpS4-trnT</i> (UGU)					*	✓
55	IGS	LSC	<i>trnT</i> (UGU)- <i>trnL</i> (UAA)	✓	✓			*	
56	Intron	LSC	<i>trnL</i> (UAA) intron	✓		✓		*	
57	IGS	LSC	<i>trnL</i> (UAA)- <i>trnF</i> (GAA)	✓				*	
58	IGS	LSC	<i>trnL</i> (UAA)- <i>ndhJ</i>	✓		✓		✓	
59	IGS	LSC	<i>trnF</i> (GAA)- <i>ndhJ</i>		✓				✓
60	IGS	LSC	<i>ndhJ-ndhC</i>					✓	
61	IGS	LSC	<i>ndhC-trnV</i> (UAC)	✓	✓	✓	✓	✓	✓
62	Intron	LSC	<i>trnV</i> (UAC) intron		✓	✓		✓	✓
63	IGS	LSC	<i>trnV</i> (UAC)- <i>atpE</i>					✓	✓
64	IGS	LSC	<i>trnV</i> (UAC)- <i>atpB</i>			✓		✓	
65	Exon	LSC	<i>atpB</i> exon			✓		✓	
66	IGS	LSC	<i>atpB-rbcL</i>		✓	✓		✓	✓
67	Exon	LSC	<i>rbcL</i> exon			✓		✓	
68	IGS	LSC	<i>rbcL-accD</i>		✓	✓		✓	✓



APPENDIX 2. Continued.

Approx. <i>Nicotiana</i> order <sup>a</sup>	Primary type	Location <sup>b</sup>	Genomic region	Shaw et al., 2005, 2007	Ebert and Peakall, 2009	Scarcelli et al., 2011	Dong et al., 2012	Dong et al., 2013	Current study
69	Exon	LSC	<i>accD</i> exon			✓		✓	
70	IGS	LSC	<i>accD-psaI</i>	✓	✓	✓	✓	*	✓
71	IGS	LSC	<i>psaI-ycf4</i>		✓	✓		*	✓
72	Exon	LSC	<i>ycf4</i> exon			✓		✓	
73	IGS	LSC	<i>ycf4-ycf10(cemA)</i>		*			✓	✓
74	Exon	LSC	<i>cemA</i>					✓	
75	IGS	LSC	<i>ycf4-petA</i>			✓		*	
76	Exon	LSC	<i>petA</i> exon			✓		✓	
77	IGS	LSC	<i>petA-psbJ</i>	✓	✓		✓	✓	✓
78	IGS	LSC	<i>psbJ-psbE</i>					✓	
79	IGS	LSC	<i>petA-psbL</i>			✓		✓	
80	IGS	LSC	<i>psbE-petL</i>	✓	✓		✓	✓	
81	IGS	LSC	<i>petL-psaJ</i>					✓	✓
82	IGS	LSC	<i>petL-trnP(UGG)</i>			✓		✓	
83	IGS	LSC	<i>trnW(CCA)-psaJ</i>				✓	✓	
84	IGS	LSC	<i>trnP(UGG)-rps18</i>		*	✓			
85	IGS	LSC	<i>psaJ-rpl20</i>		*			*	✓
86	IGS	LSC	<i>rps18-rps12</i>			✓		*	
87	IGS	LSC	<i>rpl20-rps12</i>	✓				*	✓
88	IGS	LSC	<i>rps12-psbB</i>			✓			
89	IGS	LSC	<i>rps12-clpP</i>		✓	✓		*	
90	Intron	LSC	<i>clpP</i> intron 2		✓	✓	✓	✓	✓
91	Intron	LSC	<i>clpP</i> intron 1		✓	✓	✓	✓	✓
92	IGS	LSC	<i>clpP-psbB</i>		✓	✓		✓	✓
93	Exon	LSC	<i>psbB</i> exon			✓		✓	
94	IGS	LSC	<i>psbB-psbH</i>	✓				✓	
95	IGS	LSC	<i>psbH-petBE2</i>		✓			✓	✓
96	Intron	LSC	<i>petB</i> intron/exon 2			✓		✓	
97	IGS	LSC	<i>petBE2-petDE2</i>		✓	✓	✓	✓	✓
98	Intron	LSC	<i>petD</i> intron/exon 2			✓		✓	
99	IGS	LSC	<i>petD-rpoA</i>			✓		✓	
100	Exon	LSC	<i>rpoA</i> exon					✓	
101	IGS	LSC	<i>rpoA-rps11</i>					✓	
102	IGS	LSC	<i>rps11-rps8</i>		✓	✓		✓	
103	Exon	LSC	<i>rps8</i> exon					✓	
104	IGS	LSC	<i>rpl36-rpl14</i>	✓					
105	IGS	LSC	<i>rps8-rpl16</i>		✓			✓	
106	Intron	LSC	<i>rpl16</i> intron	✓		✓		✓	
107	IGS	LSC	<i>rpl16-rps3</i>		✓	✓		✓	
108	Exon	LSC	<i>rps3</i> exon			✓		✓	
109	IGS	LSC	<i>rps3-rps19</i>		✓			*	✓
110	IGS	LSC	<i>rpl22-rpl2</i>			✓		*	
111	Intron	IRb	<i>rpl2</i> intron/exon 1-2			✓		✓	
112	IGS	IRb	<i>rpl23-ycf2</i>			✓		*	
113	Exon	IRb	<i>ycf2</i> exon					✓	
114	IGS	IRb	<i>ycf2-ndhB</i>			✓		✓	
115	Exon	IRb	<i>ndhB</i> exon 2			✓		✓	
116	Intron	IRb	<i>ndhB</i> intron/exon 1			✓		✓	
117	IGS	IRb	<i>ndhB-rps7</i>			✓		✓	
118	IGS	IRb	<i>rps7-rps12</i>					✓	
119	Intron	IRb	<i>rps12</i> intron/exon			✓			
120	IGS	IRb	<i>rps12-trnV(GAC)</i>			✓		✓	
121	IGS	IRb	<i>trnV(GAC)-rrn16</i>			✓		✓	
122	Exon	IRb	<i>rrn16</i> exon			✓		✓	
123	IGS	IRb	<i>rrn16-trnI(GAU)</i>			✓		✓	
124	Intron	IRb	<i>trnI(GAU)</i> intron			✓		*	
125	Intron	IRb	<i>trnA(UGC)</i> intron			✓		*	
126	IGS	IRb	<i>trnA(UGC)-rrn23</i>			✓		*	
127	Exon	IRb	<i>rrn23</i> exon					✓	
128	IGS	IRb	<i>rrn4,5-trnN(GUU)</i>			✓		✓	
129	IGS	IRb	<i>trnN(GUU)-ycf1</i>					✓	
130	IGS	IRb/SSC	<i>ycf1-ndhF</i>					✓	
131	Exon	SSC	<i>ndhF</i> exon				✓	✓	
132	IGS	SSC	<i>ndhF-rpl32</i>	✓		✓		✓	
133	IGS	SSC	<i>rpl32-ccsA</i>			✓		✓	
134	IGS	SSC	<i>rpl32-trnL(UAG)</i>	✓			✓	✓	
135	Exon	SSC	<i>ccsA</i> exon			✓		✓	
136	IGS	SSC	<i>ccsA-ndhD</i>			✓		✓	✓

APPENDIX 2. Continued.

Approx. <i>Nicotiana</i> order <sup>a</sup>	Primary type	Location <sup>b</sup>	Genomic region	Shaw et al., 2005, 2007	Ebert and Peakall, 2009	Scarcelli et al., 2011	Dong et al., 2012	Dong et al., 2013	Current study
137	Exon	SSC	<i>ndhD</i> exon			✓		✓	
138	IGS	SSC	<i>ndhD-ndhE</i>					✓	
139	IGS	SSC	<i>psaC-ndhE</i>						✓
140	IGS	SSC	<i>psaC-ndhG</i>			✓			
141	IGS	SSC	<i>ndhE-ndhI</i>					✓	✓
142	Exon	SSC	<i>ndhG</i> exon			✓		*	
143	IGS	SSC	<i>ndhG-ndhI</i>			✓		*	
144	Intron	SSC	<i>ndhA</i> intron	✓		✓	✓	✓	
145	IGS	SSC	<i>ndhA-ndhH</i>					✓	
146	Exon	SSC	<i>ndhH</i> exon			✓		✓	
147	IGS	SSC	<i>ndhH-rps15</i>					✓	
148	IGS	SSC/IRa	<i>rps15-ycf1</i>			✓			✓
149	IGS	IRa	<i>ycf1-rrn5</i>			✓			
Bonus	IGS	LSC	<i>rbcL-psaI</i>						✓
Bonus	IGS	LSC	<i>trnS-psbD</i>				✓		

<sup>a</sup> Several regions overlap.

<sup>b</sup> IR = inverted repeat; LSC = large single-copy region; SSC = small single-copy region.

\* Slightly different region from that listed.

APPENDIX 3. Complete chloroplast genome sequences used to assess primer utility. Format: Organism; GenBank number and version; publication.

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**Basal Dicot Grade:**

1. *Amborella trichopoda* Baill.; NC\_005086.1; Goremykin et al., 2003.
2. *Magnolia grandiflora* L.; NC\_020318.1; Li et al., unpublished (direct GenBank submission dated 22 February 2013).

**Monocots:**

3. *Acorus calamus* L.; AJ879453.1; Goremykin et al., 2005.
4. *Cymbidium aloifolium* (L.) Sw.; NC\_021429.1; Yang et al., 2013.
5. *Cymbidium mannii* Rehb. f.; NC\_021433.1; Yang et al., 2013.
6. *Cymbidium sinense* (Jacks. ex Andrews) Willd.; NC\_021430.1; Yang et al., 2013.
7. *Cymbidium tortisepalum* Fukuy.; NC\_021431.1; Yang et al., 2013.
8. *Cymbidium tracyanum* Rolfe; NC\_021432.1; Yang et al., 2013.
9. *Oryza nivara* Sharma & Shastry; NC\_005973.1; Shahid Masood et al., 2004.
10. *Oryza sativa* L. Indica group; NC\_008155.1; Tang et al., 2004.
11. *Oryza sativa* L. Japonica group; NC\_001320.1; Hiratsuka et al., 1989.
12. *Canna indica* L.; KF601570.1; Barrett et al., 2014.

**Basal Eudicot Grade:**

13. *Ceratophyllum demersum* L.; NC\_009962.1; Moore et al., 2007.
14. *Nelumbo lutea* Willd.; NC\_015605.1; Quan and Ding, unpublished (direct GenBank submission dated 16 February 2009).
15. *Nelumbo nucifera* Gaertn.; NC\_015610; Quan and Ding, unpublished (direct GenBank submission dated 16 February 2009).

**Eurosids I:**

16. *Fragaria vesca* L. subsp. *bracteata* (A. Heller) Staudt; NC\_018766.1; Njuguna et al., 2013.
17. *Fragaria vesca* L. subsp. *vesca*; NC\_015206.1; Shulaev et al., 2011.

**Eurosids II:**

18. *Gossypium herbaceum* L.; NC\_023215.1; Shang et al., unpublished (Shang, M., K. Wang, J. Hua, F. Liu, C. Wang, X. Zhang, Y. Wang, and S. Li. *Gossypium herbaceum* chloroplast, complete genome. Direct GenBank submission 11 February 2011).
19. *Gossypium herbaceum* L. subsp. *africanum* (G. Watt) Vollesen; NC\_016692.1; Xu et al., 2012.

**Euasterids I:**

20. *Olea europaea* L.; NC\_013707.2; Messina, unpublished (Messina, R. *Olea europaea* chloroplast, complete genome. Direct GenBank submission 3 March 2007).
21. *Olea europaea* L. subsp. *cuspidata* (Wall. ex G. Don) Cif.; NC\_015604.1; Besnard et al., 2011.
22. *Olea europaea* L. subsp. *europaea*; NC\_015401.1; Besnard et al., 2011.
23. *Olea europaea* L. subsp. *maroccana* (Greuter & Burdet) P. Vargas; NC\_015623.1; Besnard et al., 2011.

**Euasterids II:**

24. *Helianthus annuus* L.; NC\_007977.1; Timme et al., 2007.
  25. *Helianthus decapetalus* L.; NC\_023110.1; Bock et al., 2014.
  26. *Helianthus divaricatus* L.; NC\_023109.1; Bock et al., 2014.
  27. *Helianthus giganteus* L.; NC\_023107.1; Bock et al., 2014.
  28. *Helianthus grosseserratus* M. Martens; NC\_023108.1; Bock et al., 2014.
  29. *Helianthus hirsutus* Raf.; NC\_023111.1; Bock et al., 2014.
  30. *Helianthus maximiliani* Schrad.; NC\_023114.1; Bock et al., 2014.
  31. *Helianthus strumosus* L.; NC\_023113.1; Bock et al., 2014.
  32. *Helianthus tuberosus* L.; NC\_023112.1; Bock et al., 2014.
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