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Authors: Collins, Elizabeth S., Gostel, Morgan R., and Weeks, Andrea

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AN EXPANDED NUCLEAR PHYLOGENOMIC PCR TOOLKIT FOR SAPINDALES¹

ELIZABETH S. COLLINS^{2,4}, MORGAN R. GOSTEL³, AND ANDREA WEEKS²

²George Mason University, 4400 University Drive, MSN 3E1, Fairfax, Virginia 22030-4444 USA; and ³Department of Botany, National Museum of Natural History, Smithsonian Institution, MRC 166, P.O. Box 37012, Washington, D.C. 20013-7012 USA

- **Premise of the study:** We tested PCR amplification of 91 low-copy nuclear gene loci in taxa from Sapindales using primers developed for *Bursera simaruba* (Burseraceae).
- **Methods and Results:** Cross-amplification of these markers among 10 taxa tested was related to their phylogenetic distance from *B. simaruba*. On average, each Sapindalean taxon yielded product for 53 gene regions (range: 16–90). *Arabidopsis thaliana* (Brassicales), by contrast, yielded product for two. Single representatives of Anacardiaceae and Rutaceae yielded 34 and 26 products, respectively. Twenty-six primer pairs worked for all Burseraceae species tested if highly divergent *Aucoumea klaineana* is excluded, and eight of these amplified product in every Sapindalean taxon.
- **Conclusions:** Our study demonstrates that customized primers for *Bursera* can amplify product in a range of Sapindalean taxa. This collection of primer pairs, therefore, is a valuable addition to the toolkit for nuclear phylogenomic analyses of Sapindales and warrants further investigation.

Key words: Anacardiaceae; Burseraceae; low-copy nuclear genes; microfluidic PCR; Rutaceae.

Low-copy nuclear gene regions offer increased phylogenetic utility for species- and population-level studies of plants as compared to chloroplast and nuclear ribosomal markers (Zimmer and Wen, 2012), yet sampling these regions remains challenging due to the dearth of universal primers and barriers to sequencing whole or partial nuclear genomes from multiple individuals. Consequently, assessing the phylogenetic limits of custom-designed target sequences or primers for low-copy nuclear gene regions is critical to fully realizing their broader impacts for advancing plant systematics. We report the results of a cross-amplification study incorporating primers for 91 low-copy nuclear gene loci created by Gostel et al. (2015) for species-level phylogenetics of Malagasy *Commiphora* Jacq. (Burseraceae). Primers for these markers were developed using genomic resources from two rosoid orders by mapping sequence data from a transcriptome of *Bursera simaruba* (L.) Sarg. (Burseraceae; Sapindales) (Matasci et al., 2014) to 950 putative low- or single-copy nuclear gene loci of *Arabidopsis thaliana* (L.) Heynh. (Brassicaceae; Brassicales) (Duarte et al., 2010). Gostel et al. (2015) further optimized the primers for microfluidic

PCR-based target enrichment, a method that allows simultaneous and cost-effective amplification of multiple loci (Blow, 2009; Uribe-Convers et al., 2016).

We tested cross-amplification of these primers using 10 taxa that have varying phylogenetic distances from *B. simaruba* within Sapindales and included *A. thaliana* as the outermost limit of the survey. Sapindales is a widespread group that includes ca. 6700 species within nine families (Angiosperm Phylogeny Group, 2016) (Fig. 1). Molecular phylogenies of this order often lack sufficient phylogenetic support along their backbone as well as at the species level (e.g., Fine et al., 2014; Grudinski et al., 2014), thus our understanding of Sapindalean systematics could benefit from an expanded phylogenetic toolkit such as that provided by the Gostel et al. (2015) primers.

METHODS AND RESULTS

Taxonomic sampling and molecular methods—Appendix 1 contains accession information for the 11 taxa sampled; Fig. 1 displays their phylogenetic relationships. *Bursera simaruba* (*Bursera* Jacq. ex L. subgenus *Bursera*) and *C. grandifolia* Engl. were included as positive controls; prior work has shown that all or most of the custom-designed primers amplify PCR product in these two species (Gostel et al., 2015). For experimental taxa, we included *B. tonkinensis* Guillaumin, which is sister to *Commiphora* (Weeks and Simpson, 2007), as well as *Aucoumea* Pierre, the monotypic genus sister to *Bursera* and *Commiphora* (Weeks et al., 2014). One species from each of *Boswellia* Roxb. ex Colebr., *Canarium* L., and *Protium* Burm. f. were included, as well as *Beiselia* Forman, the monotypic genus sister to all other Burseraceae (Weeks et al., 2014). We included one species of Anacardiaceae, the family that is sister to Burseraceae (Weeks et al., 2014), and one species of Rutaceae, which represents the Sapindalean clade sister to Burseraceae–Anacardiaceae–Kirkiaceae (Muellner-Riehl et al., 2016). *Arabidopsis thaliana* (Brassicales) was included because its genomic resources were used in primer design and can test the applicability of these primers to other closely related rosoid lineages (Wang et al., 2009).

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⁴Author for correspondence: ecolli11@masonlive.gmu.edu

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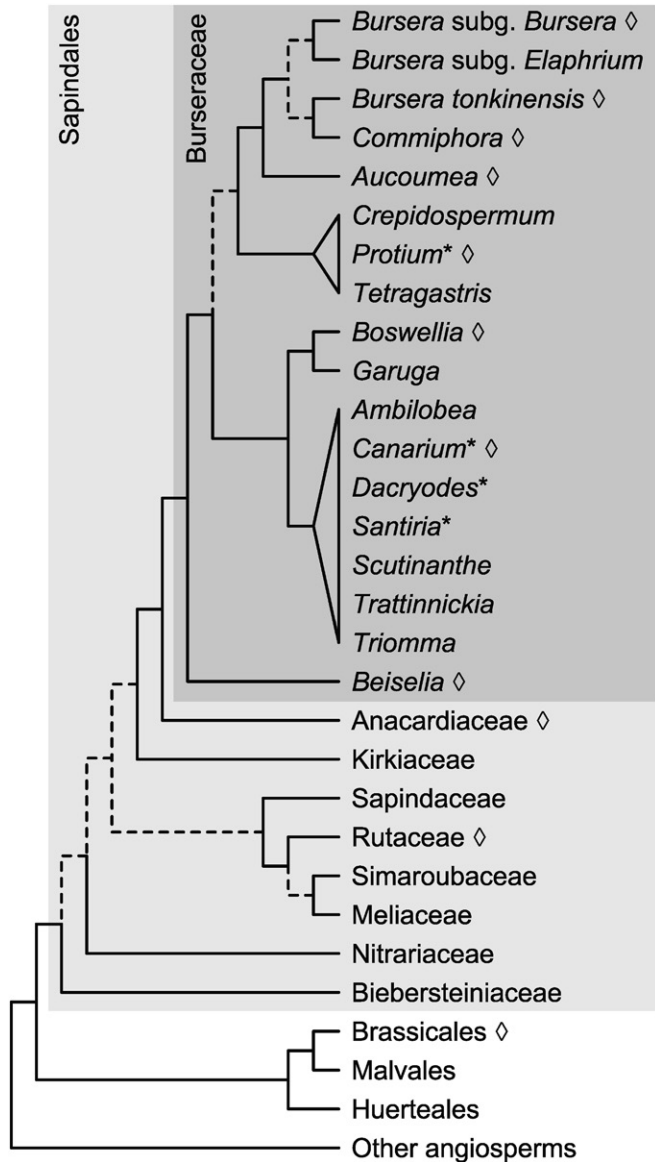


Fig. 1. Phylogeny of Sapindalean lineages condensed from Wang et al. (2009), Weeks et al. (2014), and Muellner-Riehl et al. (2016); nodes having low or conflicting support are indicated by dashed branches. Lineages sampled by the current study are noted by open diamonds. Generalized generic phylogeny of Burseraceae does not depict *Rosselia* or *Pseudodacryodes*, which have not been included in any molecular phylogenetic analysis; paraphyletic genera are indicated by asterisks.

Whole genomic DNA was extracted from taxa using the FastPrep FastDNA Spin Kit (Bio101 Systems, La Jolla, California, USA) or the cetyltrimethylammonium bromide (CTAB) method (Weeks et al., 2005). Primer development for the 91 markers is detailed by Gostel et al. (2015); primer sequences are listed in Table 1. Markers were amplified via PCR in 15- μ L reactions including: 0.15 μ L of forward and reverse primers (50 μ M), 0.75 μ L spermidine (4 mM), 7.5 μ L GoTaq Green Master Mix (Promega Corporation, Madison, Wisconsin, USA), 5.6 μ L nuclease-free water, and 1 μ L genomic DNA (0.1–25.8 ng/ μ L). Markers that failed to amplify for *B. simaruba* and *C. grandifolia* were then trialed using reaction chemistry based on that recommended for microfluidic PCR-based target enrichment including: 0.15 μ L of forward and reverse primers (50 μ M); FastStart High Fidelity PCR System reagents (Roche Diagnostics, Mannheim, Germany), composed of 1.5 μ L FastStart High Fidelity Reaction Buffer without MgCl₂ (10 \times concentration), 2.7 μ L MgCl₂ (25 mM), 0.75 μ L DMSO, 1.2 μ L Nucleotide Mix (10 mM), 0.15 μ L FastStart High Fidelity Enzyme Blend

(5 U/ μ L); 0.75 μ L Loading Reagent (Fluidigm Corporation, San Francisco, California, USA); 6.8 μ L nuclease-free water; and 1 μ L genomic DNA.

The PCR thermocycler protocol followed that of Gostel et al. (2015) and included three alternating standard and C_t cycles (Mathieu-Daude et al., 1996), beginning with 2 min at 50°C, 20 min at 70°C, and 10 min at 95°C. The first set of 10 standard cycles included a denaturation step at 95°C for 15 s, annealing at 60°C for 30 s, and extension at 72°C for 1 min. Two C_t cycles followed, including four steps consisting of 95°C for 15 s, 80°C for 30 s, 60°C for 30 s, and 72°C for 1 min. Standard and C_t cycles alternated two more times with eight, two, eight, and five cycles, respectively. After 35 cycles, samples were held at 4°C prior to being visually verified via agarose gel electrophoresis (1% agarose; 94 V for 40 min). Low DNA mass ladder (Invitrogen, Carlsbad, California, USA) was included in the first and last wells of each gel to guide length estimation of PCR products.

Marker amplification results—Table 1 contains amplification results for the low-copy nuclear loci, including the range of amplicon lengths for all taxa and GenBank numbers for markers sequenced by Gostel et al. (2015) for *B. simaruba* and *C. grandifolia* that had ≥ 15 sequence reads mapped. Table 2 summarizes marker amplification success for each taxon. Ninety primer pairs amplified product in *B. simaruba* and, on average, 54 primer pairs worked for other Burseraceae taxa. The low number of markers amplified in *Aucoumea* (16) was unexpected given its close relationship to *Bursera*. This result may have been caused by primer mismatch due to increased genetic change within this monotypic genus, as evidenced by its long branch within Burseraceae phylogeny (Weeks et al., 2014). In total, nine primer pairs worked for every Burseraceae taxon tested, and if *Aucoumea* is excluded as an outlier, the panel of family-universal primer pairs increases to 26. Thirty-four and 26 primer pairs generated product in Anacardiaceae and Rutaceae, respectively, while only two primer pairs worked in *Arabidopsis*. Comparing the Burseraceae panel to that of Anacardiaceae and Rutaceae reveals 16 and 12 successfully amplified regions in common, respectively, with eight shared among the three families. PCR chemistry may have suppressed amplification of markers, as high-fidelity PCR reagents were not used due to their high cost. Among the positive controls, high fidelity as compared to standard PCR reagents increased amplification success by 8% (*Bursera*, 83 to 90 primer pairs) and 85% (*Commiphora*, 39 to 72 primer pairs). Thus, our experimental results report a conservative baseline for the cross-amplification success of these primer pairs.

CONCLUSIONS

Our study demonstrates that 90 of 91 primer pairs for novel low-copy nuclear loci developed by Gostel et al. (2015) for *B. simaruba* successfully amplify product in a broad range of Sapindalean taxa and effectively expand the phylogenomic toolkit for this order. Twenty-six markers amplify all Burseraceae taxa (excluding *Aucoumea*) and eight amplify all Sapindalean groups tested. Our results present a new source for universal targets or primers for phylogenetic reconstruction of taxa within Sapindales. Future efforts will include sequencing amplicons to determine the number of phylogenetically informative characters for each locus.

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TABLE 1. Primer pair sequences and validation results by taxon.

Locus ID ^a	Primer sequences (5'–3') ^a	GenBank accession no. ^b		Amplicon length range among all taxa	Taxa																						
		<i>B. simaruba</i>	<i>C. grandifolia</i>		<i>Arbidiopsis thaliana</i>	<i>Aucoumea klaniana</i>	<i>Beilschia mexicana</i>	<i>Boswellia neglecta</i>	<i>Bursaria simaruba</i>	<i>Bursaria tonkinensis</i>	<i>Canarium pilosum</i>	<i>Commiphora grandifolia</i>	<i>Phellodendron amurense</i>	<i>Protium guianense</i>	<i>Schinus molle</i>	<i>Schinus fasciculatus</i>											
AT3G54460 ^c	F: GGACACACCCCTGGCTCTAG R: CTCACATGACTTTGGTTCTGTC	KX767982	KX767983	270–290	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X		
AT2G04620	F: TCCACCAFAATTTGAGTGAGGAA R: AATGGGAGTGGGAATGAAATGFG	KX76792	KX767929	420–520	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X
AT4G37510 ^c	F: TTCATTTTGAGACCTCCATTAGATGAC R: GCTAGCCGGATTATCGCTCC	KX768000	KX768000	280	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X
AT3G22660 ^c	F: AGATGAGATGTGAAAATGGTTGAACC R: TTTCTGCTTAGCTCTCTCTTTTCACTI	KX767974	KX767975	450	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X
AT1G21840 ^c	F: TGTGGAGAAGTTGAAGAGAGAGAG R: CACCAATTTCCCAACCCTCTGAA	KX767930	KX767931	630–640	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X
AT2G04740 ^d	F: CAAATCCAAAACCCTAAACCCGG R: TCAAAAAGCCTTCAAAAGCTTCTC	KX767986	KX767987	460–590	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X
AT4G14605 ^d	F: CTTCTACTCATAGCAGCAGAGAG R: TTTCTCACAGCCTTATCAAGTCA	KX767990	KX767991	510–580	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X
AT4G19900 ^c	F: GTTCTGTGAGACGATTTAGCTTGA R: CTTGTAGAGAGAGCAAGTCGG	KX767990	KX767991	350–420	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X
AT4G29590	F: GAGCAATCCCTTCAAAGAGGA R: GTCTTTGTATCTTTTGGTAATGG	KX767994, KX767996	KX767995, KX767997	490	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X
AT5G04910	F: TAAAGATGATGTCACTCAGCTTGG R: CTTCACTGGTGCATAATCTGTCTTC	KX768005	KX768006	260	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X
AT3G15110 ^c	F: ATTCTGTACCTTTGCTTCTGGA R: AACAAAGAAAGTTGCAGTAGAGGA	KX767902	KX767903	1560	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X
AT1G18060 ^d	F: GCTGGCTCTCTGCTCACTTTTGG R: CTAAGTCCCAAGAGATGAGTG	KX767926	KX767927	740–930	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X
AT2G03667 ^c	F: CACAAAGGAAATCAAGCAAGTCTT R: GGTGATATCATCTGGAAGGGGG	KX768007	KX768008	590	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X
AT2G40760	F: CGCTCTCGCCCTCTCTTTTC R: CCAATGTCAAATGGTCTCTGAAGATG	KX767940	KX767941	400	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X
AT2G20790 ^{o,e}	F: CCATGGTGCAAAATTAACGTCTTC R: AGTCCACAAGAACTGCAGTGAT	KX767940	KX767941	320–350	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X
AT2G36740	F: CATCTTTGAGAAATACCGTACTGT R: AATCATATAATAGGGCAGCCG	KX767958, KX767960	KX767959, KX767961	640–810	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X
AT3G01380 ^d	F: CCAAGAAATATAGAATTAGTCGGAC R: AGAAGAAAAGACTAACAGTACAGC	KX767966	KX767967	530–930	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X
AT3G10400	F: CCGTCTTTGAGCACGCTGA R: GTACTTGTCTCTTTAAATTTGATFAAGC	KX767966	KX767967	340	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X
AT1G59990 ^d	F: TGACACCACGAATAAATCCAAGC R: AACCCACATGGACTGTTAAACATG	KX767908, KX767910	KX767909, KX767911	450–510	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X
AT2G22370B ^c	F: CATCAGACATAAGAGATCCAGCAG R: CTTGTCTCTGGTTCATTGATCCA	KX767944	KX767945	610–780	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X
AT1G31780 ^{o,e}	F: TTTGGTCTCAATGATTTCAAGC R: CTTGTCTTGGTTCATTGATCCA	KX767904, KX767906	KX767905, KX767907	520–570	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X
AT1G31780 (INT) ^{o,e}	F: GGACCCAAAGTGTACTACAGAGAG R: GGACCCAAAGTGTACTACAGAGAG	KX767904, KX767906	KX767905, KX767907	380–832	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X

TABLE I. Continued.

Locus ID ^a	Primer sequences (5'–3') ^a	GenBank accession no. ^b		Amplicon length range among all taxa
		<i>B. simaruba</i>	<i>C. grandifolia</i>	
AT2G27760	F: GAACCTTAAACCCCTAACAAATGGAGAA R: GGCGTTCCGGTGACCATAT	X		930
AT2G27760 (INT)	F: GAACCTTAAACCCCTAACAAATGGAGAA R: CGAAAATTCCTTAGCAGTGAACCTCC	X	X	160–470
AT1G63160 (INT)	F: GACGCTGTATCTAGGCCTCAAG R: AAAATGTTGCATGTGAAGTTTGGC	X	X	220–640
AT1G63160	F: GACGCTGTATCTAGGCCTCAAG R: CACCATGAGACGGCCCAAGTAT	X	X	1070–1490
AT1G65030	F: CGGTTTTCTGTAACCTCGGTACAG R: CGGGGAAAGAGAGGTTTTGG	X	X	340
AT5G52180	F: CTCGAGAAATTTGGTTGGAAATGT R: CATAAGAAAGCCGGTCGATA	X	X	460
AT2G44760 ^d	F: CAGCATGGAATACGTTTCGTAGTA R: TATCAACTGGACCCCTGGAATAAG	X	X	530–900
AT2G05320A ^e	F: TGCCAAGTGAACAGATAATTTGCT R: TCTCCAAAAGTCGTTTAAAGGA	X	X	440
AT4G31770 ^e	F: GCGGTGAGAAATGAGAAATGACATG R: AACAAATTCCTCCAAATCCCAA	X	X	580–780
AT2G20330 ^{e,g}	F: TCATTGAAGGTTGGGATTTACGC R: ACGACTTGGTGATCTGTAATAA	X	X	610–750
AT1G66080	F: CCTCTTCTTCCATAGTTGCT R: CCCAAAAACGACTGCATAAAGTT	X		900
AT2G05170B ^e	F: GCACAGTACATTAACACCAATTTGTT R: TGGCTTTGGTCTATGAGAAATCTT	X	X	430–480
AT1G65070	F: CCTAATACTGGAGGAAACTGCT R: CAGTACTTCCCAGAGAATTCGAA	X	X	510–600
AT5G67220	F: CGGTTAAAATGCTCTCAGGATCC R: CATCTGCCGAATGAGTAACCCTTCT	X		690
AT2G17265 ^{c,d,g}	F: TTATGGAGGTTTCGGTTTGATTCG R: CTAGCACCAACTCTATCCAACTC	X	X	470–1690
AT2G46890B ^d	F: TTCCTTGGCTGCTACCCTCTCTCAG R: CGATGCTGCTTCTGATATAGCCT	X	X	570–780
AT2G31890B ^{e,g}	F: CTCCTCAGTGCAGTTHAACAG R: CTTGAGAAATGTTGGTCCATCA	X	X	410
AT2G46100 ^e	F: TTTAAAGGACTTCGCCGTTTCAA R: GCGAAGAAATAGCCCTCCAG	X	X	310–370
AT3G26580 ^{e,g}	F: AGGTAAACGTTGGATATGATG R: GTGACGTTATTGCCCTGTAAG	X	X	660–920
AT2G44660B	F: GTTTTGGCAGGAAAGGATGATTT R: TGAAGGTTTGGCTGGAGTTATCT	X	X	590–1130
AT2G44660B (INT) ^e	F: GTTTTGGCAGGAAAGGATGATTT R: TGCCTGAATCTTGAACCTTAGTTT	X	X	520–900
AT3G49730	F: CCGAAACTGGAGATGGCTTTG R: AATCAACTCAGGCCTTCTTTTCTC	X	X	140

TABLE I. Continued.

Locus ID ^a	Primer sequences (5'-3') ^a	GenBank accession no. ^b		Amplicon length range among all taxa																		
		<i>B. simaruba</i>	<i>C. grandifolia</i>																			
AT2G44660A ^f	F: ATCGTATCAGACACAGACTTTGA R: GCAAAACAACCCACCACATCAAAA	KX767950	KX767951	790																		
AT2G21710 ^e	F: TTTCCTCCTTTACTAAACATACAGCCT R: CTTGTCTGCAACCTCTGATTTGAA	KX767942 (5' only)	KX767943 (5' only)	1040–1360																		
AT2G21710 (INT) ^d	F: TTTCCTCCTTTACTAAACATACAGCCT R: GTGCATCCCAAGAGCTCTGG			750–860																		
AT2G22370A ^e	F: ATGTTGAGCCCTTGAGATTCTTC R: TAGGTGCTGTTACTTTCAACCAGTT			980–1320																		
AT1G77930A ^d	F: ACCCTAATCTGTTCTGGATTG R: GACAGTTCAATAAGCAGCTTGAAT	KX767924	KX767925	580–740																		
AT1G77930A (INT)	F: ACCCTAATCTGTTCTGGATTG R: GCATCCCTCTTAACCTCTGCAATT			410–460																		
AT5G02250 ^d	F: CACTATCCCCATGTTTCCAGAGAAC R: GGATCTGCCCTGTTTCCAAATAT			1240–1680																		
AT2G31440	F: GTA TGGAGGGCTTTTCTTCTTTG R: ATTCCTGCAGCAAGATGAACTACA			1000–1350																		
AT1G77550A ^d	F: TGTGAGCTTTCTATAATGTGGCC R: TGATGCTTCATGACCAGACAAGA	KX767920	KX767921	740–860																		
AT3G15290 ^e	F: GATGTTGTAGTCGAGGCTATTGTG R: ATCTGCAAGTTCTAAGGACCCTAT			1090																		
AT5G11980	F: TTC AACCATGCCATCCCAAAATTAAC R: GACAGAGATCCGCTTCCAGTTATC			N/A																		
AT5G14580 ^e	F: TATACGTATGGCAGAAFTTCCGG R: TCTTGTGCAATCTTATCTAAGGCT			1030–1750																		
AT2G31840 ^d	F: AGTATGATTTGGTGCTGTATGT R: CATCTGGTGAGGTAGCCTACAG			480–1220																		
AT5G57655	F: TTGGTTATGCTCAGTAAATCCGA R: CTACAGTCAGATTTGAAAAGCAT			340–1340																		
AT2G47760	F: CAGCATGGAATACGTTTGCTAGTA R: TATCAACTGGACCCTGGAATAG			620–1480																		
AT3G29130 ^d	F: TTTCGCCGAGTTCTGGTGATT R: AACTACTTCTCTGTTGATTCATCCG		KX767980, KX767981	980–1720																		
AT3G13200 ^e	F: AACTCATCGGCTTTTTCTCTCT R: GAATCATCAGCATCTACATGGGT			1970																		
AT4G33030 ^d	F: GATGGTGTCTTTGGTACTGCTTTG R: CCAAGAAAACAGTGGCATTATCTG			770–1340																		
AT1G73180 ^d	F: AACTCTGCCAGTGTCAAAATAFA R: AGAATGCCATATCACCGTAAGT			810–1000																		
AT2G31890A	F: AGATTGGAGGGAGCTACTTTAT R: CCTCCCATATACTGTCTGAAATCC	KX767948	KX767949	450–620																		
AT3G46220 ^d	F: CAATTAGGAAATGAAATGGTGGGT R: TCCATTCTTTGAAAAGCTCTCTGT			330–570																		
AT2G05120 ^{e*}	F: TGTCAAAGCTCTGGTCTCATGAAA R: CGAAGGAAGAACTGAAGCATCTAG	KX767932	KX767933	370–570																		
AT1G73740 ^d	F: TTGATATTGGAGGCTTTTGGG R: CACCAGCTCTTGAAAACAACGAG			870–1230																		

TABLE 1. Continued.

Locus ID ^a	Primer sequences (5'-3') ^a	GenBank accession no. ^b		Amplicon length range among all taxa
		<i>B. simaruba</i>	<i>C. grandifolia</i>	
AT3G17170 ^d	F: GATGATGAACACTATTTTCCCTGAGGC R: TCTTGAACCTTCTCATTACACTGC			630-900
AT3G14910 ^e	F: GGAGCTATTTATCAAAAGTTGTGCC R: AAAGCAATATACGACCAAGAAATCTG	KX767968	KX767969	360-840
Total no. of primers amplified/taxon				

Note: INT = reverse primer is an internal primer for the locus.

^aPrimer originally developed by Gostel et al. (2015).

^bGenBank accession numbers from loci used in phylogenetic analysis in Gostel et al. (2015). GenBank numbers were only created for loci of *Bursera simaruba* and *Commiphora grandifolia* that were used in the phylogenetic analysis in Gostel et al. (2015). Some loci have two GenBank numbers for a species because sequence reads did not cover the full length of the locus. The first GenBank number corresponds to the read from the 5' end of the locus; the second GenBank number corresponds to the read from the 3' end of the locus.

^cUniversal Burseraceae primer (excluding *Aucoumea*).

^dPrimer for which high-fidelity TAQ increased amplification success for *Commiphora grandifolia*.

^ePrimer for which high-fidelity TAQ increased amplification success for *Bursera simaruba*.

^fPrimer for which high-fidelity TAQ increased amplification success for *Bursera simaruba* and *Commiphora grandifolia*.

^gUniversal Sapindales primer (excluding *Aucoumea*).

^hFaint double band observed.

TABLE 2. Number of primer pairs amplified of the 91 primer pairs tested for each of the 11 taxa.

Species tested (Order; Family)	Primer pairs amplified/tested (%)
<i>Arabidopsis thaliana</i> (Brassicales; Brassicaceae)	2/91 (0.02)
<i>Aucoumea klaineana</i> (Sapindales; Burseraceae)	16/91 (17)
<i>Beiselia mexicana</i> (Sapindales; Burseraceae)	47/91 (52)
<i>Boswellia neglecta</i> (Sapindales; Burseraceae)	68/91 (75)
<i>Bursera simaruba</i> (Sapindales; Burseraceae)	90/91 (99)
<i>Bursera tonkinensis</i> (Sapindales; Burseraceae)	53/91 (58)
<i>Canarium pilosum</i> (Sapindales; Burseraceae)	71/91 (78)
<i>Commiphora grandifolia</i> (Sapindales; Burseraceae)	72/91 (79)
<i>Phellodendron amurense</i> (Sapindales; Rutaceae)	26/91 (28)
<i>Protium guianense</i> (Sapindales; Burseraceae)	54/91 (59)
<i>Schinus fasciculatus</i> (Sapindales; Anacardiaceae)	34/91 (37)

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APPENDIX 1. Accession information for taxa used in this study, including voucher information, country of origin, and latitude and longitude coordinate data, if available, and DNA extraction method.

Species	Voucher (Herbarium)	Country of origin	Geographic coordinates	DNA extraction method ^a
Sapindales				
Burseraeae				
<i>Aucoumea klaineana</i> Pierre	Walters <i>et al.</i> 466 (MO)	Gabon	00°07'12"S, 11°42'57"E	1
	McPherson 16293 (MO)	Gabon	00°27'S, 11°45'E	1
<i>Beiselia mexicana</i> Forman	Pell <i>s.n.</i> (TEX)	Mexico	NA	1, 2
<i>Boswellia neglecta</i> S. Moore	Weeks 00-VII-29-1 (TEX)	Ethiopia	NA	2
<i>Bursera simaruba</i> (L.) Sarg.	Weeks 16-VI-16-01 (GMUF)	USA	NA	1
	Goldman <i>s.n.</i> (BH)	USA	NA	2
<i>Bursera tonkinensis</i> Guillamin	Daly <i>et al.</i> 13929 (NY)	Vietnam	20°15'12.6"N, 105°43'2.5"E	1
<i>Canarium pilosum</i> A. W. Benn.	Bogler <i>s.n.</i> (TEX)	Malaysia	NA	2
<i>Commiphora grandifolia</i> Engl.	Gostel 121 (GMUF)	Madagascar	23°39'19.64"S, 44°37'44.36"E	1
	Weeks 10-I-09-10 (GMUF)	Madagascar	12°14'16.14"S, 49°22'12.906"E	1
<i>Protium guianense</i> (Aubl.) Marchand	Miller and Hawk 9391 (MO)	Suriname	04°45'22"N, 056°52'30"W	1
Anacardiaceae				
<i>Schinus fasciculatus</i> (Griseb.) I. M. Johnst.	Silva-Luz 287 (NY)	Argentina	24°52'05.4"S, 65°32'41.4"W	1
Rutaceae				
<i>Phellodendron amurense</i> Rupr.	Weeks 15-VII-13-01 (GMUF)	USA	38°49'53.76"N, 77°18'32.04"W	1
Brassicales				
Brassicaceae				
<i>Arabidopsis thaliana</i> (L.) Heynh.	Gostel <i>s.n.</i> (GMUF)	USA	NA	1

Note: NA = not available.

^a 1 = FastDNA, 2 = CTAB.