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

Source: Applications in Plant Sciences, 4(12)

Published By: Botanical Society of America

URL: <https://doi.org/10.3732/apps.1600078>

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PRIMER NOTE

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* (Brassicaceae), 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 rosid 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* (Brassicaceae) was included because its genomic resources were used in primer design and can test the applicability of these primers to other closely related rosid lineages (Wang et al., 2009).

¹Manuscript received 25 June 2016; revision accepted 19 September 2016.

The authors thank Cíntia Silva-Luz for providing leaf material of *Schinus* and *Beiselia*. Research was supported in part by the National Science Foundation (grant no. 1403150 to M.R.G. and A.W.) and the Provost/COS/ESP Institutional Graduate Fellowship Research Award to E.S.C. from George Mason University. Any opinions, findings, and conclusions or recommendations expressed in this material are those of the authors and do not necessarily reflect the views of the National Science Foundation. Publication of this article was funded in part by the George Mason University Libraries Open Access Publishing Fund.

⁴Author for correspondence: ecoll11@masonlive.gmu.edu

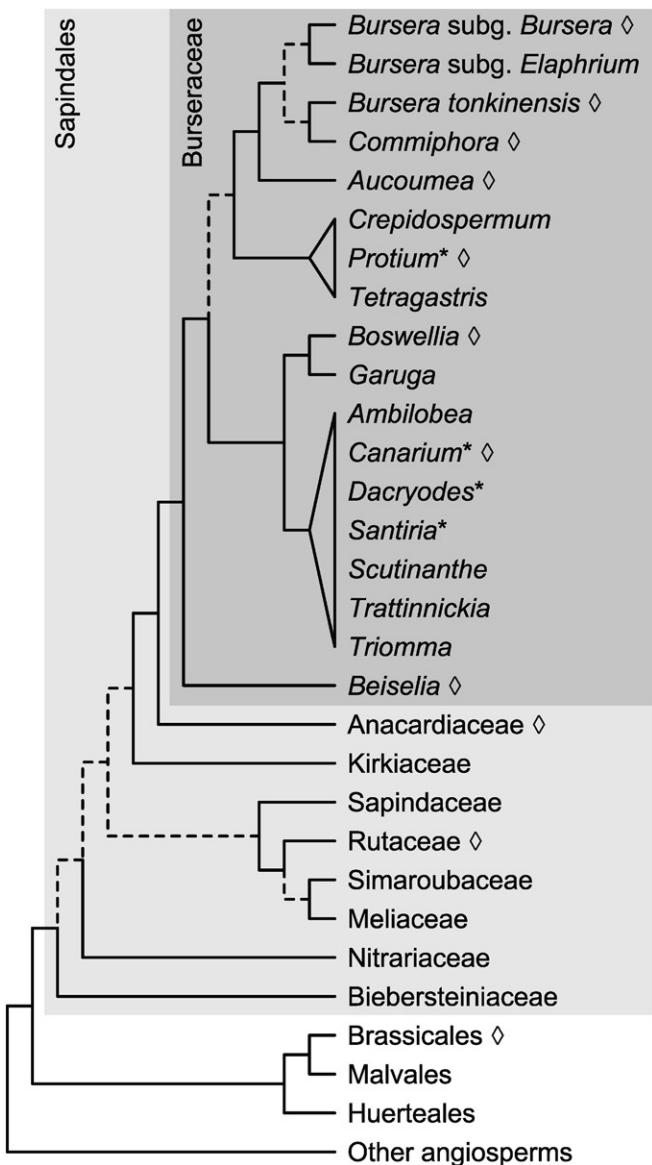


Fig. 1. Phylogeny of Sapindalean lineages condensed from Wang et al. (2009), Weeks et al. (2014), and Muellner-Riehl et al. (2016); nodes having low or conflicting support are indicated by dashed branches. Lineages sampled by the current study are noted by open diamonds. Generalized generic phylogeny 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 (10x 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_ot cycles (Mathieu-Daudé 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_ot 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_ot 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.

LITERATURE CITED

- ANGIOSPERM PHYLOGENY GROUP. 2016. An update of the Angiosperm Phylogeny Group classification for the orders and families of flowering plants: APG IV. *Botanical Journal of the Linnean Society* 181: 1–20.
- BLOW, N. 2009. Microfluidics: The great divide. *Nature Methods* 6: 683–686.
- DUARTE, J. M., P. K. WALL, P. P. EDGER, L. L. LANDERR, H. MA, J. C. PIRES, J. LEEBENS-MACK, AND C. W. DEPAMPHILIS. 2010. Identification of shared single copy nuclear genes in *Arabidopsis*, *Populus*, *Vitis*, and *Oryza* and their phylogenetic utility across various taxonomic levels. *BMC Evolutionary Biology* 10: 61.
- FINE, P. V. A., F. ZAPATA, AND D. C. DALY. 2014. Investigating processes of Neotropical rain forest tree diversification by examining the evolution and historical biogeography of the Proteae (Burseraceae). *Evolution* 68: 1988–2004.

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

Locus ID ^a		Primer sequences (5'-3') ^a	GenBank accession no. ^b	B. simaruba	C. grandifolia	Amplicon length range among all taxa	
AT3G54460 ^c	F:	GGACACACCCTGGCTCTAG	KX767983	X	X	270–290	
	R:	CTCCCATGACTTTGGTTCGTGTC	KX76792	X	X	420–520	
AT2G04620	F:	TCCACATATTGAGTGAAAGCAA	KX76792	X	X	X	
	R:	AAATGGGAGTGGGAATGAGATG	KX76800	X	X	X	
AT4G37510 ^c	F:	TTCATTTGACACCTCCATAGATGAC	KX76800	280	X	X	
	R:	GCTTAGCCGGATTAAATGGTCTGC	KX76794	X	X	X	
AT3G22660 ^c	F:	AGATGAGATGAAATTGGTGAAC	KX76794	450	X	X	
	R:	TTTCTGCTTAGCTCTCTTCTCATCT	KX76795	630–640	X	X	
AT1G21840 ^c	F:	TGGGGAGAACCTGGAGAGAGG	KX76794	X	X	X	
	R:	ACCAATTATCCTAACCTCTGAA	KX767930	X	X	X	
AT2G04740 ^d	F:	CAAACCTCCAAAACCTAACCGG	KX767931	460–590	X	X	
	R:	TCAAAAGCCCTCAAAAGCTCTCTC	KX767986	X	X	X	
AT4G14605 ^d	F:	CTTCTCACTTATAGCAGGAGAAG	KX767987	510–580	X	X	
	R:	CTTCTCACAGCTTATCAAGTCA	KX767990	X	X	X	
AT4G19900 ^c	F:	GTTCCTCTGAGCGATTGAGCTTGA	KX767991	350–420	X	X	
	R:	CTTGTAGAGAGGCAAGTGG	KX767994	490	X	X	
AT4G29590	F:	GAGCAATTCCCCTCAAGAGCA	KX767995	X	X	X	
	R:	GTGCTTGTAACTCTTTGGTAATGG	KX767996	X	X	X	
AT5G04910	F:	TAAGAGTCCACAGCAGCATGAGT	KX768005	260	X	X	
	R:	TAAAAGAATGATGTCACTAGCTTGC	KX768006	X	X	X	
AT3G15110 ^e	F:	CTCACCTGGCGCATATGCTGCT	KX767902	X	X	X	
	R:	ATTCCTCTGTAACCTTGGCTCTGGA	KX767903	740–930	X	X	
AT1G18060 ^d	F:	AAACAAGAAAAGTTGCAGTAAGGA	KX767926	X	X	X	
	R:	GCTGGGCTCTGTCACTTTTGT	KX767927	590	X	X	
AT2G03667 ^c	F:	CTAGTTGGCTGGTGTATGATG	KX767927	X	X	X	
	R:	CACAAAGGAAATAAACGAAAGCCT	KX768007	400	X	X	
AT2G40760	F:	GGTGTATCATGGAAAGGGG	KX767940	X	X	X	
	R:	CGCTCTGCCCTCTCTTCTTCT	KX767941	320–350	X	X	
AT2G20790 ^{c,g}	F:	CCATTGTCATGGTCTCAAGAGATG	KX767941	640–810	X	X	
	R:	COATGGTGCACATTTAACGTGTTCC	KX767958,	530–930	X	X	
AT2G36740	F:	AGTCACAAAGACTGCAGTAT	KX767959,	X	X	X	
	R:	CATCCCTTGAGAAATAACCGTATCTGT	KX767960	X	X	X	
AT3G01380 ^d	F:	AATCATCATATAAGGGCAGCG	KX767961	340	X	X	
	R:	COAAGAAATAATAGAAGTTAGTGGT	KX767966	X	X	X	
AT3G10400	F:	CGGTCTTGGTCTCTTAATATAAGC	KX767908,	450–510	X	X	X
	R:	AGAGAAAAGACTAACAGSACAGC	KX767910	X	X	X	
AT1G59990 ^d	F:	GCTACTTGGTCTCTTAATATAAGC	KX767944	610–780	X	X	X
	R:	TGACACCACGMAATAAACCTAACATG	KX767945	X	X	X	
AT2G22370B ^c	F:	AACCCACATGGFACTGTTAACATG	KX767904,	520–570	X	X	X
	R:	CATCAGACATAGAGATGCAAGCAG	KX767905,	380–832	X	X	X
AT1G31780 ^{c,g}	F:	CTTGTCCCTTGGTTACTTGTATCCA	KX767906	X	X	X	
	R:	GGACCCAAAAGTGTACTACAGAG	KX767907	X	X	X	

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>	
AT2G27760	F:	GAACCTTAACCCCTAACAACTGGAGAA			930
	R:	GGCGGTTCCTGGACCATAT			
AT2G27760 (INT)	F:	GAACCTTAAACCCCTAACAACTGGAGAA			160–470
	R:	CGAAATTCCCTAGCAGTGAACTCC			220–640
AT1G63160 (INT)	F:	GACGCTGTATCTAGGCTCCAG			
	R:	AAAATGTTGCAGTGTGAAGTTGGC			
AT1G63160	F:	GACGCTGTATCTAGGCTCCAG			1070–1490
	R:	CAACATGAGAACGAT			
AT1G65030	F:	CGGTTTCTGAAACTCGGGTACAG	KX767912	KX767913	340
	R:	CGGGGAAAAGAGGGTTTGG			
AT5G52180	F:	CTCGGAGAAATTGGTGGAAATGT	KX768003	KX768004	460
	R:	CATACAGAAAGCCGCTCGATA			
AT2G44760 ^d	F:	CAGCATGGAAATACGTGCTAGTA	KX767954	KX767955	530–900
	R:	TATCAACTGGAAACCCCTGGATAAG			
AT2G05320A ^c	F:	TGCCAAAGAACAGCTGGTTAAAGGA	KX767934	KX767935	440
	R:	TCTCCAAGAACAGCTGGTTAAAGGA			
AT4G31770 ^c	F:	GGGGTGAGAAATGAGAAATGACATG	KX767998	KX767999	580–780
	R:	ACAAGTTCTCCATCCAAATTCCAAA			
AT2G20330 ^{e,g}	F:	TCATTGAAGGTTGGGATTCAGC	KX767938	KX767939	610–750
	R:	ACGACTTGGCGATCTCTGATAAA			
AT1G66080	F:	CCTCTCTCATAGTGTGCT			
	R:	CCACAAAAGCACTGCATAAAGTT			
AT2G05170B ^c	F:	GCACAGTACATAAACCCATTGGT			
	R:	TGGCTTGTGGNCATATGAGAACTT			
AT1G65070	F:	CCTAATACTGGAGGGAAAATGCT	KX767914	KX767915	430–480
	R:	CAGTACTTCCCAGAGAAATCGAA			
AT5G67220	F:	CGGTAAAAAGCTCTCAGATCC			510–600
	R:	TATGGGAGGTTTACCTCT			
AT2G17265 ^{c,d,g}	F:	CTCTCCAGTGTCACTTTCAGTACAG	KX767936	KX767937	690
	R:	CTAGCACCAACTCTATCCACCTC			
AT2G46890B ^d	F:	TCTTTGCTGTCTCTGATATAGCCT			470–1690
	R:	CGATGTCGTCTCTGATATAGCCT			
AT2G31890B ^{c,g}	F:	TTATGGGAGGTTTACCTCT			570–780
	R:	CTTGAGAAATCTGTGGTCATCA			
AT2G46100 ^c	F:	TITAAGGACTTCGGCTTCAAA	KX767956	KX767957	310–370
	R:	GGCAGAAAAGATAAGCCCTCAG			
AT3G26580 ^{c,g}	F:	AGGTGAACGGTGGATTATGAT	KX767976,	KX767977,	660–920
	R:	GTGACGGTATTGCTCTGTAAG	KX767946	KX767947	
AT2G44660B	F:	GTTTTGCAGAAGGGATGATT			410
	R:	TGAAGGTTGGCTGGAGTATCT			
AT2G44660B (INT) ^c	F:	GTTTTGCAGAAGGGATGATT	KX767952	KX767953	590–1130
	R:	TGCTGAATCTGAACTCTAGTT			
AT3G49730	F:	COAAAATCTGAGATGGCTT	KX767953		520–900
	R:	AATCAACTCAGGCCTTCTCTC			140

TABLE 1. Continued.

Locus ID ^a	Primer sequences (5'-3') ^a	<i>B. simaruba</i>	<i>C. grandifolia</i>	GenBank accession no. ^b	Amplicon length range among all taxa
AT2G44660A ^f	F: ATCGTATCACAGCACAGACATTGAR: R: GCAAAACAAACACCCATCAA	KX767950	KX767951	KX767950	790
AT2G21710 ^c	F: TTTCCTCCTTTACTAACATACAGCCTR: R: CTTGTCTGCACCTCTGTGATGAA	KX767942	KX767943 (5' only)	KX767942 (5' only)	1040-1360
AT2G21710 (INT) ^d	F: TTTCCTCCTTTACTAACATACAGCCTR: R: GCTGCCATCCCAGAGCTCTGG				750-860
AT2G22370A ^c	F: ATGGTGAGGCCCTGGAGATCTTC R: TAGGTGCTGTAACCAACAGATT	KX767924	KX767925		980-1320
AT1G77930A ^d	F: ACCCTAATTCTGTTCTGCATTG R: GAGCAGTTCATAGCAGCTTGAAT				580-740
AT1G77930A (INT)	F: ACCCTAATTCTGTTCTGCATTG R: GCATCCCTTAACCTCTGAATT				410-460
AT5G02250 ^d	F: CACTTATCCCTAATGTTCCAAGAAC R: GGATCTGCCCTGTTTCAAAATAT				1240-1680
AT2G31440	F: GTATGGAGGGTTCTTCCTTTG R: ATTCCTTGCAAGAGATGAACATACA				1000-1350
AT1G77550A ^d	F: TGTGAGCTTCTCTATATTGCGC R: TGATGCTTCAGAACAGAACAGA	KX767920	KX767921	KX767920	740-860
AT3G15290 ^c	F: GATGTTGTTGAGGCTATGTG R: ATCTGCAAGTCTAAAGGCCAT				1090
AT5G111980	F: TTCAACCCTGATCCCCAAATTAC R: GACAGAGATCCCCCTCAAGTATC			N/A	
AT5G14580 ^c	F: TATACTGTTGCGAGAATTCCGG R: TCCTGTGCAAACTTATCTAAGGCCT				1030-1750
AT2G31840 ^d	F: AGTGTATTGATGGTGTCCCTGATGT R: CATCTTGGTGAAGTAGCCCTACAG				480-1220
AT5G57655	F: TTGGTTATGCUAGTGTAAATCGA R: CTACAGTGCACATTGGAAACCAT				340-1340
AT2G47760	F: CAGCATGGAAATACGTTGCTAGTA R: TATCAACTGGACCCCTGGATAAG				620-1480
AT3G29130 ^d	F: TTGCCGAGGTCTGGTGAATT R: AAGTACTTCTCTGTTGATTTCCG				980-1720
AT3G13200 ^c	F: AACATTCGGCTTTCTCTCT R: GAATCATCAGAATCTACATCGT				1970
AT4G33030 ^d	F: GATGGTGTCTTGTACTGCTTTG R: COAGAAAACAGTGGCATATTCTG				770-1340
AT1G73180 ^d	F: AACTCCTGCCAGTGTCCAATATA R: AGAATGCCATTACCCAGTAGT				810-1000
AT2G31890A	F: AGATTGGAGGGAGCTACTTTATT R: CCTCCCTATACGTGCTGAAATCC	KX767948	KX767949	KX767948	450-620
AT3G46220 ^d	F: CAATTGAGGAGTGAATGGTGCCT R: TCCATTCTGCTGAAAGCTTGT				330-570
AT2G05120 ^e	F: TGTCAAAGCTCTGGTCTCATCAA R: CGAGGAAGAACACTGAAGCACTAG	KX767932	KX767933	KX767932	370-570
AT1G73740 ^d	F: TTGATATTGGAGGCTCTTGGG R: CACAGCTCTGAAACAAACAGAG				870-1230

TABLE 1. Continued.

Locus ID ^a		GenBank accession no. ^b	Amplicon length range among all taxa		
			<i>B. simaruba</i>	<i>C. grandifolia</i>	
AT4G31790 ^d	F: ATTTGGTTGTTGAGGCCAAAGAAA R: GTCCAAAATGACCATCTGGAGTT		1620–2180		X X X X X X
AT5G10460	F: TGGTCATCAATTAGCAATTCTCAGC R: GCTCTTCAAAATCTCCAACT		1320–1800	X X X X X X	X X X X
AT4G26980 ^d	F: CTGCTAGTGGGTTCTGAATTGG R: ACTTCCTCAAGCATTGACAACCTCAT		940–1170	X X X X X X	X X X X
AT5G48790 ^d	F: GAGGATTGGTTGACTGAAGAG R: TCGGACCTTAAATGTGAATGTT		680–1250	X X X X X X	X X X X
AT5G15680A	F: TTCTCATCAAAACATCTGGGCC R: GAGGAATTGCAATCAGATTCTGGTC	KX768001	560	X X X X X X	X X X X
AT3G04650	F: CAAATCGCTTGTTGATGTGTTTC R: CTGGGGCAGTGGGATGTTTTC	KX767962, KX767964	490–660	X X X X X X	X X X X
AT2G25570 ^d	F: GACAACCTCAAAATCACAGCCAG R: GTCCCTCTCATGCCCCATG	KX767965	660–1160	X X X X X X	X X X X
AT2G31040 ^d	F: AAGTACTGGGGAGAAAGAG R: CCAAATGAGGATTGCACTTC		1230–1690	X X X X X X	X X X X
AT4G04955	F: GAACAGATACTGTTACAGCCAG R: TGAGCTTTAGTCCCTGAAAG		440–1170	X X X X X X	X X X X
AT3G21540 ^c	F: GTTGCTATTAGCTGATGCCAAA R: AATGGTTCTTGTAGCATGCCAA	KX767970, KX767972	730–1020	X X X X X X	X X X X
AT2G05170A ^c	F: GAAGGAAATGTTACAGTGAGGA R: TGAGAAGAATGGGGAGCTCTT	KX767973	650–880	X X X X X X	X X X X
AT2G28450 ^f	F: TTCTGAGATAATGCTTATGTCAGG R: CGCCAATTGTCAGTACCA		1330	X X X X X X	X X X X
AT3G07750 ^d	F: GCTATATTGTTGATTGAGGCCT R: TGGTGTCTCACGTTTAATGATC		940–1330	X X X X X X	X X X X
AT1G76450 ^d	F: CGTCGGCACAAAATTACAAGAATGGA R: TCAAATTCCCAGTCCAAAGTCCCAATC	KX767916, KX767918	1330–1570	X X X X X X	X X X X
AT3G10530 ^{c,d}	F: ATTCCCCATCAAATTCCACTCG R: ACTCATGGATCTCCAGTACTAA	KX767919	700–1960	X X X X X X	X X X X
AT3G61620 ^c	F: TTAGATATTGAGGCAAGAACGG R: CCTAAAAGGTATGGTCAAGGTT		1330	X X X X X X	X X X X
AT4G21170 ^d	F: TGGAGCTGTTATTATGCCCTTGT R: TAGTCCTAGTAAACAACAGC	KX767992	730–1130	X X X X X X	X X X X
AT3G22990 ^d	F: TCTCCCTTCACTGAACTGAGA R: ACAGTTCAAGGGCACATGATC		900–1130	X X X X X X	X X X X
AT4G18810B	F: TTATGATATTGAGGCAAGAACGG R: CCTAAAAGGTATGGTCAAGGTT	KX767988	540–640	X X X X X X	X X X X
AT1G77550B ^d	F: TCTAGGTCCATCTCAAGTGCAGA R: CTGGTGTGTTATGTGATTGATGTC	KX767922	760–820	X X X X X X	X X X X
AT5G16690 ^d	F: TGTTCCCTGAGAACATTGTCACT R: TCAAAGAACGGCTGAACTGTT		760–820	X X X X X X	X X X X
AT4G00560	F: CTGCTATGTCATAAACGCTCTCC R: GTCCACCAACATCAACAGTAAC	KX767984 (5' only)	900–1180	X X X X X X	X X X X

TABLE 1. Continued.

Locus ID ^a		GenBank accession no. ^b			Amplicon length range among all taxa
		B. simaruba	C. grandifolia		
AT3G17170 ^d	F: GATGATGAACATTATTTCTTGAGGC R: TCTTGAACCTTCTCATTCACACTGC				630–900
AT3G14910 ^e	F: GGAGCTATTATCAAAGTTGTGCC R: AAAGCAATAATACGACCAAAGAACTCTG	KX767968	KX767969	360–840	
Total no. of primers amplified/taxon				2 16 47 68 90 53 71 72 72 26 54 34	

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). Some loci 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> (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)

GOSTEL, M. R., K. A. COY, AND A. WEEKS. 2015. Microfluidic PCR-based target enrichment: A case study in two rapid radiations of *Commiphora* (Burseraceae) from Madagascar. *Journal of Systematics and Evolution* 53: 411–431.

GRUDINSKI, M., C. M. PANNELL, M. W. CHASE, J. A. AHMAD, AND A. N. MUELLNER-RIEHL. 2014. An evaluation of taxonomic concepts of the widespread plant genus *Aglaja* and its allies across Wallace's Line (tribe Aglaeae, Meliaceae). *Molecular Phylogenetics and Evolution* 73: 65–76.

MATASCI, N., L.-H. HUNG, Z. YAN, E. J. CARPENTER, N. J. WICKETT, S. MIRARAB, N. NGUYEN, ET AL. 2014. Data access for the 1,000 plants (1KP) project. *GigaScience* 3: 17.

MATHIEU-DAUDE, F., J. WELSH, T. VOGT, AND M. MCCLELLAND. 1996. DNA rehybridization during PCR: The 'C_{ot} effect' and its consequences. *Nucleic Acids Research* 24: 2080–2086.

MUELLNER-RIEHL, A. N., A. WEEKS, J. W. CLAYTON, S. BUERKI, L. NAUHEIMER, Y.-C. CHIANG, S. CODY, AND S. K. PELL. 2016. Molecular phylogenetics and molecular clock dating of Sapindales based on plastid *rbcL*, *atpB* and *trnL-trnF* DNA sequences. *Taxon* 65: 1019–1036.

URIBE-CONVERS, S., M. L. SETTLES, AND D. C. TANK. 2016. A phylogenomic approach based on PCR target enrichment and high throughput sequencing: Resolving the diversity within the South American species of *Bartsia* L. (Orobanchaceae). *PLoS ONE* 11: e0148203.

WANG, H., M. J. MOORE, P. S. SOLTIS, C. D. BELL, S. F. BROCKINGTON, R. ALEXANDRE, C. C. DAVIS, ET AL. 2009. Rosid radiation and the rapid rise of angiosperm-dominated forests. *Proceedings of the National Academies of Science, USA* 106: 3853–3858.

WEEKS, A., D. C. DALY, AND B. B. SIMPSON. 2005. The phylogenetic history and biogeography of the frankincense and myrrh family (Burseraceae) based on nuclear and chloroplast sequence data. *Molecular Phylogenetics and Evolution* 35: 85–101.

WEEKS, A., AND B. B. SIMPSON. 2007. Molecular phylogenetic analysis of *Commiphora* (Burseraceae) yields insight on the evolution and historical biogeography of an "impossible" genus. *Molecular Phylogenetics and Evolution* 42: 62–79.

WEEKS, A., F. ZAPATA, S. K. PELL, D. C. DALY, J. D. MITCHELL, AND P. V. A. FINE. 2014. To move or to evolve: Contrasting patterns of intercontinental connectivity and climatic niche evolution in "Terebinthaceae" (Anacardiaceae and Burseraceae). *Frontiers in Genetics* 5: 409.

ZIMMER, E. A., AND J. WEN. 2012. Using nuclear gene data for plant phylogenetics: Progress and prospects. *Molecular Phylogenetics and Evolution* 65: 774–785.

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

Note: NA = not available.

^a1 = FastDNA, 2 = CTAB.