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Phylogenic Relationships of *Rubus* Species Revealed by Randomly Amplified Polymorphic DNA Markers

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Abstract

Korean cultivated bramble, which is known as Bokbunja-ddal-gi is regarded as having originated from Korean native *Rubus core anus*. However, little scientific evidence and significant morphological differences between Korean cultivated bramble (KCB) and R. coreanus throws doubt on the ancestry of KCB. This study was carried out to obtain phylogenetic information on KCB by comparing its nuclear genomic background with those of *R. coreanus*, black (*R. occidentalis*) and red (*R. idaeus*) raspberry, blackberry (*R. lanciniatus*), and *R. crataegifolius*. A total of 99 random amplified polymorphic DNA (RAPD) markers were generated and used for phylogenetic analysis of 76 *Rubus* accessions. Accessions of each species were grouped into each distinct subclade by the RAPD markers at a similarity coefficient of about 0.59. The KCB subclade formed a clade with *R. occidentalis* and *R. crataegifolius* subclades at a similarity coefficient of 0.47. The *R. coreanus* subclade formed a clade with *R. idaeus*, *R. lanciniatus*, and *R. crataegi folius* subclades at a similar level of genetic similarity. Only one KCB accession from Hoengsung was included in the *R. coreanus* subclade. The accession shows leaf and flower characteristics differed from the rest of the KCB accessions which show morphological similarity to black raspberry is more closely related to black raspberry than to *R. coreanus*. This brings about the need for close scientific evaluations on the ancestry of KCB at both morphological and molecular levels.

Key words: Random amplified polymorphic DNA (RAPD), phylogenetic relationship, Rubus species

Introduction

Bramble refers to thorny plants of the genus *Rubus*, such as raspberry, blackberry, and loganberries. Wild brambles first attracted the attention of herbalists due to their medicinal properties. *R. coreanus*, the *Rubus* species native to Korea, has been used for various medicinal purposes. Well-known medicinal properties of *R. coreanus* fruit include remedies for palsy, imbecility, arthritis, and stomach diseases. It also has beneficial effects for sexual disorders and cancer treatments (Jeong and Sin 1996; Park et al. 2003). The use of bramble fruit has been diversified as a beneficial food resource as customers' interest in healthier food has grown with economic development and

* **To whom correspondence should be addressed** Song Joong Yun E-mail: sjyun@chonbuk.ac.kr Tel: +82-63-270-2508 changes in food industry. The demand for bramble fruit has increased as a variety of new foods and beverages containing bramble fruit or its extract as a major ingredient or supplements have been developed. The increased demand was followed by the cultivation of bramble plants in farm fields in South Korea. As the cultivation was initiated in Gochang-gun area in North Jeolla province, the cultivated bramble was collectively called Gochang Bokbunja-ddal-gi, or simply Bokbunja. The production of Bokbunja has increased drastically in the last five years and the cultivation area was estimated to be over 2,500 ha in 2006 (Korean Black Raspberry Experiment Station).

Rubus is one of the most diverse genera in the plant kingdom, and it contains 12 subgenera (Jennings 1988). With the efforts of plant breeders, species bearing edible fruits have been selected, and cultivars producing larger and more flavorsome fruits have been developed. Raspberries and blackberries are domesticated

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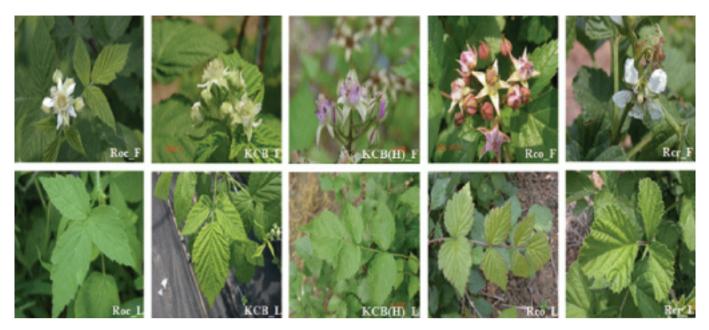


Fig. 1. Morphological characteristics of leaf (L) and flower (F) of *Rubus* species used in this study. KCB: Korea cultivated bramble, KCB (H): Korea cultivated bramble in Hoengsung, Rco: bramble native to Korea (*R. coreanus* Mique), Roc: black raspberry (*R. occidentalis* L.), Rcr: *R. crataegifolius.*

forms of the wild *Rubus* species. Raspberries belong to the subgenus Idaeobatus, which contains about 200 species. Black, red, and purple raspberries are major domesticated species belonging to Idaeobatus. Black raspberries include the two main subspecies, *R. occidentalis* L. and *R. leucodermis* Dougl., which are indigenous to North America. *R. occidentalis* L. is more common in the eastern part of North America, and *R. leucodermis* Dougl. in the western part of North America. Black raspberry cultivars commercially cultivated in the USA originated from *R. occidentalis*, but not from *R. leucodermis* Dougl. (Jennings 1988). Red raspberries are widely distributed in the temperate regions of Europe, Asia, and North America. The two major subspecies in red raspberries are *R. idaeus* and *R. strigosus*. Blackberries belong to the subgenus Eubatus, which contains very large number of species (Jennings 1988).

The bramble cultivated widely in South Korea has been known as a domesticated form of the Korean native R. coreanus, which has been the traditional source of Bokbunja used for various medicinal purposes. Even though the cultivation history of the bramble in Korea is relatively new, as recent as the last five decades, little scientific documentation is available on the domestication or cultivation history. Therefore, the claim that KCB is a domesticated form of R. coreanus suffers from lack of supporting scientific evidence and documentation. Though R. coreanus bears black berries as KBC and black raspberry, flower color, and leaf shapes of KCB are rather more similar to those of black raspberry than to those of Korea native R. coreanus. KCB has white flowers and trifoliate leaves, whereas R. coreanus has purple flowers and penta- or nona-foliate leaves (Fig. 1). These morphological similarities between KCB and black raspberry have been the ground of speculation that KCB is

a derivative of black raspberry. Furthermore, cultivars of black raspberry and red raspberry were introduced to Korea from Canada and the USA in the 1960s (Ham et al. 1997). The random amplified polymorphic DNA (RAPD) markers generated using random primers and polymerase chain reaction (PCR) have been widely used for various polymorphism analyses including identification of cultivars, differentiation of plant species, and genetic analysis of phylogenetic relationships among strains or populations (Antonius and Klemola 1999; Charcosset and Moreau 2004; Mohan et al. 1997; Rafalski and Tingey 1993). Therefore, the objective of this study was to obtain basic phylogenetic information about the ancestry of KCB by comparing genomic background of KCB with those of *R. coreanus, R. crataegifolius*, black raspberry, red raspberry, and blackberry.

Materials and Methods

Plant materials

A total of 76 accessions of *Rubus* species were used in this study. Fourteen accessions of KCB were collected from the major cultivation areas in South Korea (Table 1). Black raspberry (Roc, *R. occidentalis* L., 15 cultivars) was obtained from National Clonal Germplasm Repository, Corvallis, Oregon, USA. *R. coreanus* (Rco, 21 species), red raspberry (Rid, *R. idaeus*, four species), blackberry (Rla, *R. lanciniatus*, four species), and *R. crataegifolius* (Rcr, 18 species) were obtained from the Korean Black Raspberry Experiment Station, Gochang, Korea.

Entry Number	Common Name (Taxon)	Collection location	Note (Variety)	Entry Number	Common Name (Taxon)	Collection location	Note (Variety)
1	South Korean	South Korea	Gochang, Jeonbuk	53	R. crataegifolius	South Korea	Gosung, Gyeongnam
2	cultivated bramble		Pyeongchang, Kangwon	54	-Rcr-		Kimhae, Gyeongnam
3	(Unknown) -KCB-		Hoengsung, Kangwon	55			Ulryungdo, Gyeongbuk
4			Sanchung, Gyeongnam	56			Goheung, Jeonnam
5			Yeonggwang, Jeonnam	57			Gokseong, Jeonnam
6			Jangsung, Jeonnam	58			Gurye, Jeonnam
7			Kimje, Jeonbuk	59			Youngam, Jeonnam
8			Sunchang, Jeonbuk	60			Jangheung, Jeonnam
9			Wanju, Jeonbuk	61			Gongju, Chungnam
10			Jangsu, Jeonbuk	62			Asan, Chungnam
11			Jeongeup, Jeonbuk	63			Taean, Chungnam
12			Jinan, Jeonbuk	64			Yeongwol, Kangwon
13			Taean, Chungnam	65			Gochang, Jeonbuk
14			Cheongwon, Chungbuk	66			Okcheon, Chungbuk
15	South Korean	South Korea	Taebak, Kangwon	67			Yeongwol, Kangwon
16	bramble		Hongcheon, Kangwon	68	R. crataegifolius	North Korea	Bacakdu Mt, NORTH KOR
17	(<i>R. coreanus</i> Mique)		Hoengsung, Kangwon		-Rcr-	North Rored	
18	-Rco-		Yangpyeong, Gyeonggi	69	Red raspberry	USA	Autum Bliss
19			Hamyang, Gyeongnam	70	(R. idaeus)		Golden Harvest
20			Sangju, Gyeongbuk	71	-Rid-		NOVA
21			Muan, Jeonnam	72			Canby
22			Bosung, Jeonnam	73	Blackberry	USA	Thorny
23			Wanju, Jeonbuk	74	(R. lanciniatus)		Thornless
24			Danyang, Chungnam	75	-Rla-		Creeping
25			Okcheon, Chungbuk	76			Ebano
26			Jecheon, Chungbuk				
27			Cheongwon, Chungbuk				
28			Cheongju, Chungbuk		_		
29			Gosung, Gyeongnam	Chemica	ls		
30			Goheung, Jeonnam	DNA	extraction kit	s were fro	om Gentra Syster
31			Gokseong, Jeonnam				asmid vector pGEM
32			Gurye, Jeonnam	· 1		-	adison, WI, USA).
33			Youngam, Jeonnam	•			from Sigma (St. Lou
34			Gongju, Chungnam		•	*	fioni Signia (St. Lot
				MO, USA), unless otherwis	e indicated.	
			Taean Chungnam				
35	Black raspherry	Δ	Taean, Chungnam				
36	Black raspberry (<i>R. occidentalis</i> L.)	USA	Black Hawk	Extractio	on of genomic I	DNA	
36 37		USA	Black Hawk Bristol		e		ung leaves using t
36 37 38	(R. occidentalis L.)	USA	Black Hawk Bristol Cumberland	DNA v	was isolated fro	om fresh yo	0 0
36 37 38 39	(R. occidentalis L.)	USA	Black Hawk Bristol Cumberland Plum Farmer	DNA v Puregene	was isolated fro DNA purificatio	om fresh yo n kit (Gentra	a System, Minneapo
36 37 38 39 40	(R. occidentalis L.)	USA	Black Hawk Bristol Cumberland Plum Farmer Shuttleworth	DNA Puregene MN, USA	was isolated fro DNA purificatio) following the in	om fresh yo n kit (Gentra nstruction pro	System, Minneapo by the manuf
36 37 38 39 40 41	(R. occidentalis L.)	USA	Black Hawk Bristol Cumberland Plum Farmer Shuttleworth New Logan	DNA Puregene MN, USA turer. Fres	was isolated fro DNA purificatio) following the is h leaf samples (30	om fresh yo n kit (Gentra nstruction pro 0 mg) were gr	a System, Minneapo povided by the manuf round with a mortar a
36 37 38 39 40 41 42	(R. occidentalis L.)	USA	Black Hawk Bristol Cumberland Plum Farmer Shuttleworth New Logan Black Knight	DNA v Puregene MN, USA turer. Fres pestle in l	was isolated fro DNA purificatio) following the in h leaf samples (30 liquid nitrogen. (30	om fresh yo n kit (Gentra nstruction pro 0 mg) were gr Cells of the g	a System, Minneapo ovided by the manuf round with a mortar a ground leaf tissue w
36 37 38 39 40 41 42 43	(R. occidentalis L.)	USA	Black Hawk Bristol Cumberland Plum Farmer Shuttleworth New Logan Black Knight Red fruit	DNA Puregene MN, USA turer. Fres pestle in I lysed by i	was isolated fro DNA purificatio) following the in h leaf samples (30 liquid nitrogen. C ncubating in lysis	om fresh yo n kit (Gentra nstruction pro 0 mg) were gr Cells of the g s solution at	ung leaves using to a System, Minneapo ovided by the manuf round with a mortar a ground leaf tissue we 65 °C for 60 min. C
36 37 38 39 40 41 42 43 44	(R. occidentalis L.)	USA	Black Hawk Bristol Cumberland Plum Farmer Shuttleworth New Logan Black Knight Red fruit NC 84-10-3	DNA Puregene MN, USA turer. Fres pestle in I lysed by i	was isolated fro DNA purificatio) following the in h leaf samples (30 liquid nitrogen. C ncubating in lysis	om fresh yo n kit (Gentra nstruction pro 0 mg) were gr Cells of the g s solution at	a System, Minneapo ovided by the manuf round with a mortar a ground leaf tissue we 65 °C for 60 min. C
36 37 38 39 40 41 42 43 44 45	(R. occidentalis L.)	USA	Black Hawk Bristol Cumberland Plum Farmer Shuttleworth New Logan Black Knight Red fruit NC 84-10-3 NC 84-10-7	DNA Puregene MN, USA turer. Fres pestle in I lysed by i lysate wa	was isolated fro DNA purificatio) following the in h leaf samples (30 liquid nitrogen. C ncubating in lysis s collected by co	om fresh yo n kit (Gentra nstruction pro 0 mg) were gu Cells of the g s solution at entrifugation	a System, Minneapo ovided by the manuf round with a mortar a ground leaf tissue we 65 °C for 60 min. C and treated with p
36 37 38 39 40 41 42 43 44 45 46	(R. occidentalis L.)	USA	Black Hawk Bristol Cumberland Plum Farmer Shuttleworth New Logan Black Knight Red fruit NC 84-10-3 NC 84-10-7 NC 84-10-2	DNA Puregene MN, USA turer. Fres pestle in I lysed by i lysate wa teinase K	was isolated fro DNA purificatio .) following the in h leaf samples (30 liquid nitrogen. Concubating in lysis s collected by co (6 mg/ml) for 60	om fresh yo n kit (Gentra nstruction pro 0 mg) were gr Cells of the g s solution at entrifugation min at 55 °C.	a System, Minneapo ovided by the manuf round with a mortar a ground leaf tissue w 65 °C for 60 min. C and treated with p . RNA was degraded
36 37 38 39 40 41 42 43 44 45 46 47	(R. occidentalis L.)	USA	Black Hawk Bristol Cumberland Plum Farmer Shuttleworth New Logan Black Knight Red fruit NC 84-10-3 NC 84-10-7 NC 84-10-2 Haut	DNA Puregene MN, USA turer. Fres pestle in I lysed by i lysate wa teinase K adding RN	was isolated fro DNA purificatio) following the in h leaf samples (30 liquid nitrogen. C ncubating in lysis s collected by co (6 mg/ml) for 60 Nase A (1.5 mg/m	om fresh yo n kit (Gentra nstruction pro 0 mg) were gr Cells of the g s solution at entrifugation min at 55 °C. nl) in the cel	a System, Minneapo ovided by the manuf round with a mortar a ground leaf tissue w 65 °C for 60 min. C and treated with p . RNA was degraded Il lysate and incubat
36 37 38 39 40 41 42 43 44 45 46 47 48	(R. occidentalis L.)	USA	Black Hawk Bristol Cumberland Plum Farmer Shuttleworth New Logan Black Knight Red fruit NC 84-10-3 NC 84-10-7 NC 84-10-2 Haut Dundee	DNA Puregene MN, USA turer. Fres pestle in I lysed by i lysate wa teinase K adding RN the lysate	was isolated fro DNA purificatio) following the in h leaf samples (30 liquid nitrogen. C ncubating in lysis s collected by co (6 mg/ml) for 60 Nase A (1.5 mg/r at 37 °C for 15	om fresh yo n kit (Gentra nstruction pro 0 mg) were gu Cells of the g s solution at entrifugation min at 55 °C. nl) in the cel min. Proteir	a System, Minneapo ovided by the manuf round with a mortar a ground leaf tissue w 65 °C for 60 min. C and treated with p . RNA was degraded Il lysate and incubat as were precipitated
36 37 38 39 40 41 42 43 44 45 46 47 48 49	(R. occidentalis L.)	USA	Black Hawk Bristol Cumberland Plum Farmer Shuttleworth New Logan Black Knight Red fruit NC 84-10-3 NC 84-10-7 NC 84-10-2 Haut Dundee John Robertson	DNA Puregene MN, USA turer. Fres pestle in I lysed by i lysate wa teinase K adding RN the lysate adding the	was isolated fro DNA purificatio) following the in h leaf samples (30 iquid nitrogen. C ncubating in lysis s collected by co (6 mg/ml) for 60 Nase A (1.5 mg/m at 37 °C for 15 e protein precipit	om fresh yo n kit (Gentra nstruction pro 0 mg) were go Cells of the g s solution at entrifugation min at 55 °C. nl) in the cel min. Proteir ation solution	a System, Minneapo ovided by the manuf round with a mortar a ground leaf tissue w 65 °C for 60 min. C and treated with p . RNA was degraded Il lysate and incubat as were precipitated n to the cell lysate f
36 37 38 39 40 41 42 43 44 45 46 47 48 49 50	(<i>R. occidentalis</i> L.) -Roc-		Black Hawk Bristol Cumberland Plum Farmer Shuttleworth New Logan Black Knight Red fruit NC 84-10-3 NC 84-10-3 NC 84-10-2 Haut Dundee John Robertson Mac Black	DNA Puregene MN, USA turer. Fres pestle in I lysed by i lysate wa teinase K adding RN the lysate adding the lowed by i	was isolated fro DNA purificatio) following the in h leaf samples (30 liquid nitrogen. Concubating in lysis s collected by co (6 mg/ml) for 60 Nase A (1.5 mg/r at 37 °C for 15 e protein precipit inverting racks co	om fresh yo n kit (Gentra nstruction pro 0 mg) were gr Cells of the g s solution at entrifugation min at 55 °C. nl) in the cel min. Proteir ation solution ntaining the s	a System, Minneapo ovided by the manuf round with a mortar a ground leaf tissue w 65 °C for 60 min. C and treated with p . RNA was degraded Il lysate and incubat as were precipitated in to the cell lysate f samples for about 2 m
36 37 38 39 40 41 42 43 44 45 46 47 48 49	(R. occidentalis L.)	USA South Korea	Black Hawk Bristol Cumberland Plum Farmer Shuttleworth New Logan Black Knight Red fruit NC 84-10-3 NC 84-10-7 NC 84-10-2 Haut Dundee John Robertson	DNA v Puregene MN, USA turer. Fres pestle in I lysed by i lysate wa teinase K adding RN the lysate adding the lowed by i and centri	was isolated fro DNA purificatio) following the in h leaf samples (30 liquid nitrogen. C ncubating in lysis s collected by co (6 mg/ml) for 60 Nase A (1.5 mg/m at 37 °C for 15 e protein precipit inverting racks co fuging at 13,000	om fresh yo n kit (Gentra nstruction pro 0 mg) were gr Cells of the g s solution at entrifugation min at 55 °C. nl) in the cel min. Proteir ation solution ontaining the s -16,000 x g	a System, Minneapo ovided by the manuf round with a mortar a ground leaf tissue we 65 °C for 60 min. C and treated with p

Table 1. A total 76 Rubus accessions of Korean	cultivated bramble <i>R</i> coreanus <i>R</i> crat	aegifolius R occidentalis R idaeu	s and R lanciniatus
Table 1. A total 70 habas accessions of Rolean	cultivated bramble, n. coreanas, n. crat	acynonius, n. occiaentans L., n. iaaca	s, and n. ianciniatus.

washed with 70% ethanol. DNA pellet was dried and hydrated in 50 μl DNA hydration solution by incubating the DNA sample overnight at room temperature. DNA concentration was measured by both spectrophotometric assay and gel electrophoresis. DNA samples were used for PCR for nuclear genomes.

Random amplified polymorphic DNA analysis

Primers for the analysis of RAPD were purchased from SeouLin Bioscience (SRILS uniprimer kit, Seoul, Korea). RAPD analysis of nuclear genomic DNA was carried out with the DNA template prepared from leaf samples. PCR amplification was carried out in a 30 µl reaction mixture, containing ~0.01-0.1 ng of template DNA, 10 pmol primer (SRILS uniprimer kit), dNTPs (0.2 mM each), and Taq polymerase (1 unit, Ex Taq PCR, TaKaRa). The thermal cycles consisted of an initial denaturation (5 min at 94 °C), 40 cycles of amplification (repeated cycles of denaturation (1 min at 94 °C), annealing (1 min at 55 °C) and extension (1 min at 72 °C)), and a last extension (10 min at 72 °C). The amplified products were separated in 1% agarose gel at 100V for 30 min and visualized by staining the gel in ethidium bromide solution and photographed under UV light. RAPD markers were analyzed by UPGMA (unweighted pair-group method with arithmetic average) method using NTSYS (numerical taxonomy and multivariate analysis system) program (Sneath and Sokal 1973).

Results and Discussion

A total of 99 RAPD markers including 84 major bands and 15 minor bands were generated using the 12 SRILS primers.

Typical markers are shown in Fig. 2.

Profiles of RAPD markers were scored and analyzed by the UPGMA method using a NTSYS program. Accessions of each species were grouped into a distinct separate subclade based on the RAPD markers (Fig. 3). Some of the accessions in a species were very similar to each other and the accessions of wild *R. crataegifolius* were clearly distinct both morphologically and phenogenetically.

Accessions of each species were grouped into a distinct separate subclade by the species at the similarity coefficient of about 0.59. The RAPD markers revealed that KCB forms a distinct subclade separately from that of each Rubus species. The KCB subclade formed a clade with R. occidentalis and R. crataegifoliu subclade at the similarity coefficient of 0.47. The R. coreanus subclade formed a clade with R. idaeus and R. lanciniatus subclades at the similarity coefficient of 0.45. The clade consisting of R. coreanus and R. lanciniatus subclades is remotely but more closely related to R. idaeus subclade than the clade containing KCB and R. occidentalis (Fig. 3). This result indicates that KCB has a genetic constitution significantly different from other *Rubus* species. Among the *Rubus* species tested in this study, nuclear genomic background of KCB accessions shows relatively closer relatedness to that of R. occidentalis than to that of R. coreanus. This strongly suggests that the widely accepted ancestry of KCB to R. coreanus might have originated from legendary assumptions rather than sound scientific evidence.

Nevertheless, scientific studies which used KCB as an experimental material referred to the scientific name of the material as *R. coreanus* rather than *R. occidentalis* (Cha et. at. 2007; Yoon et al. 2002). Furthermore, nearly all advertisements of the products made of KCB fruits refer KCB as *R. coreanus*. *R. occidentalis* is not included in the introduction to Bokbunja produced in

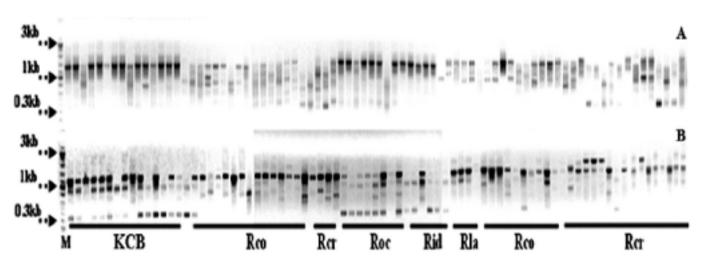


Fig. 2. Typical random amplified polymorphic DNAs generated for the bramble species. M: DNA ladder, KCB: Korea cultivated bramble, Rco: Korea native bramble (*R. coreanus* Mique), Roc: black raspberry (*R. occidentalis* L.), Rcr: *R. crataegifolius*, Rla: blackberry (*R. lanciniatus*), and Rid: red raspberry (*R. idaeus*). A (URP primer11), B (URP primer12).

Gochang-gun (http://www. gochang. jeonbuk.kr/farm/).

Only one KCB accession from Hoengsung was included in R. coreanus subclade. Hoengsung accession has unique morphological characteristics different from all the other KCB accessions. Its leaf and flower characteristics are similar to R. coreanus rather than to R. occidentalis to which all the rest of the KCB accessions share a close similarity. A few Korean black raspberry breeders have used wild R. coreanus germplasms collected from mountains of the southern part of the Korean peninsula (Kim et al. 2002a; Kim et al. 2002b). They assumed that KCB is R. occidentalis rather than R. coreanus (Kim et al. 2002a). Accessions in the KCB subclade show a diverse genetic distance. Especially, the KCB accession from Sunchang (accession #8) is included in the subclade at the similarity coefficient of 0.59. This result indicates that there is a considerable genetic variation among the local KCB accessions. Also, the KCB accession from Hoengsung is most likely domesticated from R. coreanus.

In conclusion, the genetic background inferred from the RAPD fragments shows that all KCB accessions, except one from Hoengsung, exhibit the closest relatedness to *R. occidentalis*

among the *Rubus* species used in this study such as *R. coreanus* Mique, *R. crataegifolius*, *R. occidentalis*, *R. idaeus*, and *R. lanciniatus*. This brings about the need for close scientific investigations on the ancestry of KCB at both morphological and molecular levels.

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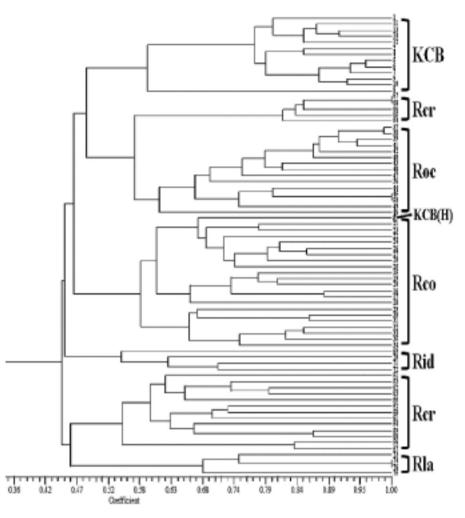


Fig. 3. A phylogram generated from the 99 nuclear RAPD markers for the *Rubus* accessions. KCB: Korea cultivated bramble, KCB (H): Korea cultivated bramble in Hoengsung, Rco: Korea native bramble (*R. coreanus* Mique), Roc: black raspberry (*R. occidentalis* L.), Rcr: *R. crataegifolius*, Rla: blackberry (*R. lanciniatus*), and Rid: red raspberry (*R. idaeus*).

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