MORPHOLOGICAL AND ANATOMICAL STUDIES OF THE NEWLY RECORDED RHUS CHINENSIS MILL. (ANACARDIACEAE) FROM TURKEY

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Abstract

Rhus chinensis Mill. (Anacardiaceae) has been reported as a new record for the flora of Turkey. Detailed morphological description and leaf anatomical properties are provided. Capitate glandular and nonglandular trichomes, and also epicuticle hairs have been observed in the leaf surfaces of R. chinensis. Fruit micromorphology and chromosome number of this species have also been evaluated.

Introduction

Anacardiaceae Lindl., the Cashew family, includes more than 700 species in the world belonging to 82 genera that are primarily distributed pantropically. The genus Rhus L., the sumac genus (Spondiadoideae, Rhoaeae), is one of the most widespread and recognizable genera in the north temperate zone, includes approximately 250 species from subtropical and warm temperate regions of the world. The genus is divided into two subgenera: Rhus L. and Lobadium (Raf.) Torr. & A.Gray (Barkley, 1937; Yi et al., 2007).

During field survey in Artvin Province (Turkey), some interesting specimens belonging to Anacardiaceae were collected. After critical study and consultation with relevant literature (Davis et al., 1967; Linchevskii, 1974; Davis, 1988; Güner et al., 2000; Hsu and Su, 2013; Eminağaoğlu, 2015), these specimens were identified as Rhus chinensis Mill. The species was not reported earlier from Turkey (Eminağaoğlu and Anşin, 2003; Eminağaoğlu and Anşin, 2004; Eminağaoğlu et al., 2008; Özhatay et al., 2011; Eminağaoğlu and Özcan, 2013, 2014; Yuksel and Eminağaoğlu, 2017). The number of species of Rhus in the flora of Turkey is increased to 2 with the addition of this species. In this study, we describe detailed morphological characters of Rhus chinensis, and investigate the anatomical and cytological properties of the species.

Material and Methods

Morphological analysis

Plant materials were collected from different parts of Artvin, Turkey, at different altitudes between 2013 and 2017. The collected materials were critically studied. The voucher specimens have been deposited at the Herbarium of Artvin Coruh University (ARTH), Artvin, Turkey.

Anatomical preparation

For anatomical investigation leaf samples were stored in 70% alcohol. Transverse sections of leaf, and paradermal sections of upper and lower epidermis of leaves were prepared manually using commercial razor blades and stained in Haematoxylin for about 15 min. To remove the excess stain, sections were washed in water several times (Algan, 1981). Semi-permanent slides were mounted in glycerin or permanent slides were covered with glycerin-gelatin (Vardar, 1987).

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Well stained sections were examined under a light microscope and photographed using an Olympus BX-53 microscope with digital camera attachment DP-73.

**Micromorphological analysis**

Micromorphological features of the fruits were studied using a stereomicroscope (Leica M60 with a digital camera attachment DFC 295) and a scanning electron microscope (Zeiss Evo LS 10, ACU-Biltekmer). For scanning electron microscopy, fruits and seed covered with endocarp were separately placed on stubs using double-sided adhesive tape, and coated with gold in a Cressington sputter coater 108 auto coating apparatus for 2 min. Fruits and endocarp were examined and photographed from the same region (from the middle part of the lateral region).

**Chromosome count**

For mitotic chromosome observation, root tips were cut off and pretreated with 1-Bromonaphthalen at 4°C for 16 h (Ozcan et al., 2011), then fixed in fresh Carnoy absolute alcohol-glacial acetic acid (3:1) for 24 h at 4ºC. For chromosome counts, root tips were hydrolyzed in 5N HCl for 3-5 min at room temperature and then rinsed with distilled water for 2-3 min. Staining was carried out in lacto-propionic orcein at least for 3 h at room temperature. Permanent slides were prepared from at least ten well-spread cells. The best metaphase plates were photographed with Olympus BX-53 microscope with digital camera attachment DP-73.

**Results and Discussion**

**Rhus chinensis** Mill. Gard. Dict. ed. 8, n. 7 (1768).  

Shrub to small deciduous tree, 2–10 m tall; branchlets ferruginous pubescent, lenticellate. Leaves sessile, imparipinnately compound; rachis broadly winged, ferruginous pubescent; leaflets 9-13, leaflet blade ovate to oblong, 7–13×3–7 cm, increasing in size towards apex, adaxially dark-green, sparsely pubescent or glabrescent, abaxially pale-green, glaucous, and ferruginous pubescent, base rounded to cuneate in terminal leaflet, margin dentate, often crenate, lateral veins and reticulate venation impressed adaxially and prominent abaxially; petiole 4–9 cm long, densely pubescent. Flowers dioecious; inflorescence panicle, densely ferruginous pubescent. Staminate flowers 35–40 mm long, pedicel short, c. 1 mm, minutely pubescent; calyx pubescent, lobes ovate, c. 1 mm long, margin ciliate; corolla obovate-oblong, white, c. 2 mm long; filaments c. 2 mm long; anthers ovoid, c. 0.7 mm; disk annular, yellow; ovary very reduced or absent. Pistillate flowers: corolla elliptic-ovate, white, c. 1.6–2.0 mm long; calyx lobes c. 0.6 mm long; staminodes 4 or 5, disk annular, yellow; ovary ovoid, c. 1 mm long, densely white pubescent; styles 3; stigma capitrate. Drupe globose, slightly compressed, 4–5 mm in diam., mixed pilose and glandular-pubescent, red at maturity.

**Flowering period:** June to August; **fruiting period:** September to November.

**Specimens examined:** A8 Artvin, Hopa, Kemalpaşa, damp roadside, grassland, 15m, 27 July 2013, 41°30’33”N, 41°32’15”E, Ö. Emin. 18677 (ARTH 5250); Artvin, Hopa, Kemalpaşa, 36m, 29 September 2013, 41°27’52”N, 41°29’9”E, Ö. Emin. 19101 (ARTH 5251); Artvin, Hopa, Kemalpaşa, damp roadside, grassland, 10m, 8 May 2014, 41°28’22”N, 41°30’17”E, Ö. Emin. 19364 (ARTH 5269); Artvin, Arhavi, roadside, 40m, 5 June 2017, 41°20’47”N, 41°16’44”E, Ö. Emin. 22360 (ARTH 11342).
Fig. 1. *Rhus chinensis* Mill.: A. Winged rachis with pinnate leaves; B. Flowering branch; C. Pistillate flowers; D. Staminate flowers; E. Fruits. Scale bars: A=3 cm, B=5 mm, C=2 mm, D=8 mm, E=2 mm.

*Distribution:* China, Manchuria, India, Thailand, Korea, Japan, Tibet, Taiwan, North Korea, South Korea, Malaysia, India, Bhutan, Myanmar, Pakistan, Jammu and Kashmir, Laos, Thailand and Cambodia (Hassler, 2017). New record for Turkey.


*Uses:* Different parts of *R. chinensis* including root, bark, stem, leaf, flowers, fruit, seed and gall are used in the treatment of hemoptysis, inflammations, cough, dysentery, fever, jaundice, malaria, rheumatism, laryngitis, snakebite, stomach-ache and traumatic fractures. Galls on the
plant are also used for treatment of diarrhoea, hemorrhage, ulcer of mouth, diabetes, and rectal and intestinal cancer (Djakpo and Yao, 2010).

Leaf anatomical characteristics

Midrib: It is semi-circle and covers a large area. Under the upper and lower epidermis, several layers of collenchyma cells are observed. 12-15 collateral vascular bundles arranged in a circle. The larger bundles present near to upper epidermis are thin walled parenchymatous cells filled in the pith. At least one large resin duct is present in each vascular bundle. Sclerenchymatous sheath can be visible in phloem of vascular bundle as a cap (Fig. 2).

Fig. 2. Leaf anatomy of *Rhus chinensis*: A-B. Midrib; C. Vascular bundles; D. Trichomes; E. Veinlet; F. Lamina. cl: collenchyma, ct: capitate trichome, eh: epicuticular hair, pc: pith cell, ph: phloem, pp: palisade parenchyma, sc: secretory canal (duct), dc: druse crystals, sp: spongy parenchyma, str: simple trichome, xy: xylem. Scale bars: A = 200 μm; B,C,E = 100 μm; D,F = 50 μm.
Lamina: The leaves show dorsiventral mesophyll. It is composed of single palisade layer and 3-4 spongy layers. The palisade tissue covers in equal areas to spongy parenchyma. Epidermis is covered with a thick cuticle. Upper epidermal cells with straight walls are distinctly larger than the lower ones (Fig. 2F). Non-glandular and capitate glandular trichomes are sparsely observed in the adaxial surface, but abaxial one included densely non-glandular multicellular and capitate trichomes with 4-8 head cells and epicuticle hairs. It bears hypostomatic type stomata which are found only in abaxial side of leaf (Fig. 3). They are sunken in lower epidermis (Fig. 2C, D).

Crystals: Druse crystal compounds are present in some collenchymatous cells of midrib and inside of the palisade cells of lamina. They are much bigger (Fig. 2F)

*R. chinensis* has dorsiventral mesophyll with single layer of palisade cells and 3-4 layers of spongy cells. Yanping *et al.* (2001) investigated formation of gall in this species and showed the differences in the ratio of palisade tissue and spongy tissue. In the present study, we observed three types of trichomes in the leaf of *R. chinensis* (Fig. 3). Mobius (1899), and Rost and Gilg (1912) reported two different types of trichomes (thick-walled bristle hairs and club-shaped trichomes) in *R. vernicifera* L. and *R. toxicodendron* L., respectively. Liu *et al.* (2008) found epicuticular hairs in *R. chinensis*. In the present study, crystal compounds were observed in *R. chinensis* (Fig. 2). McNair (1921) reported these types of crystals in *R. diversiloba* Torr. & A. Gray [*Toxicodendron diversilobum* (Torr. & A.Gray) Greene]. McNair (*l.c.*) also reported resin
ducts in *R. diversiloba*. Harada (1932) reported resin ducts in the petiole and veinlets of *R. succedanea* L. In the present study similar results were found for *R. chinensis* as like in *R. diversiloba*. Therefore our results are in accordance with previous studies.

**Fruit micromorphology**

Fleshy fruit has reddish colour with round shape, and endocarp is laterally compressed. Fruit length ranged from 4.5 to 4.7 mm. Results of fruit micromorphology are shown in Fig. 4. Exocarp cells are undulate and more or less inflated with striate surface. Epidermal surface was covered with glandular trichomes. Endocarp has regular sclerified cells.

![Fig. 4. SEM micrography of fruit of *Rhus chinensis*. A-C. Fruit; D-F. Endocarp.](image)

**Chromosome number**

Somatic chromosome number of *R. chinensis* has been determined as $2n=2x =30+0-2B$ (Fig.5).

![Fig. 5. Somatic metaphase of *Rhus chinensis*.](image)
Two B chromosome have also been found. In a previous report, Shang et al. (1990) documented chromosomes of four *Rhus* species, and reported chromosome number of *R. chinensis* as 2n = 30 +0-5B and chromosome numbers of other three taxa as 2n = 30. However, triploid cultivar was also detected in their study for *R. verniciflua* as 2n = 3x = 45. Parfitt et al. (1990) found gametic chromosome number of *R. aromatica* Aiton var. *pilosissima* (Engl.) Shinners as n = 15. Our results are in accordance with those previous reports and only diploid number was observed for *R. chinensis* in this study.

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**References**


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