MOLECULAR PHYLOGENY OF ROCHELIA (BORAGINACEAE) BASED ON NRDNA ITS AND CPDNA TRNL-F SEQUENCES

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Received 20 12 2009. Accepted for publication 12 05 2010


We here present molecular phylogeny of the genus Rochelia (Boraginaceae-Eritrichieae). A total of 8 species of Rochelia and 2 species of Lappula as outgroups were included in analyses using nrDNA ITS and cpDNA trnL-F separately and in combination. To examine evolutionary trend of morphological characters, we mapped six diagnostic characters on the combined tree using MacClade 4. The analyses revealed that sect. Rochelia due to inclusion of the monotypic section Cryptocarpa (Rochelia cardiosepala) is not monophyletic. Likewise, its subsections, Rochelia and Pedunculares are paraphyletic. Rochelia persica and R. disperma along with R. cancellata of the monospecific subgenus Neo-Rochelia, as unresolved branches, were sisters to the remaining species. One of six diagnostic characters examined (non-hamate tip of calyx hairs) had evolved as reversal in both R. persica and R. bungei and the other one (nutlets completely clasping the adaxial part of gynobase) had undergone parallel evolution between R. cancellata plus R. peduncularis and R. cardiosepala. Based on the present molecular analyses, the current infrageneric classification of Rochelia, at least at the sectional and subsectional level based upon traditional morphological characters is artificial.

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Key words. Boraginaceae, cpDNA trnL-F, Eritrichieae, Molecular Phylogeny, nrDNA ITS, Rochelia.

cpDNA trnL-F و nrDNA ITS فیلوزن مولکولی جنس Rochelia (Boraginaceae) بر اساس توالی های (Rochelia cancellata) تیلن، دانشگاه آزاد اسلامی، واحد قم، هدم در اثر خوش‌ساخترین، استادیار گروه زیست‌شناسی دانشگاه آزاد اسلامی، واحد قم. درک شارح کاوش پور ایسلاو، دانشیار گروه علوم گیاهی، دانشگاه تربیت مدرس. درک سارا سعادت‌نده، استادیار گروه زیست‌شناسی دانشگاه آزاد اسلامی، واحد علوم و تحقیقات. درک هدایت عظیم، دانشگاه تربیت مدرس.

کلید واژه‌ها. Boraginaceae، cpDNA trnL-F، Eritrichieae، مواد فیلوزنی، nrDNA ITS، Rochelia.

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INTRODUCTION

*Rochelia* Reichenb. is a cosmopolitan genus with its dispersal centre in S.W. and central Asia, extending to the Mediterranean area and Australia, comprising 15-20 species (Nasir 1989, Mabberley 1990, Luque 1992). It is represented in Iran by 8 species (Khatamsaz 2002). *Rochelia* is characterized by 2-nutlet fruits mostly ornamented by stellate papillae (Khatamsaz 2002, Riedl 1967, Hilger 1984, Kazempour Osaloo 1993). De Candolle first (1846) classified *Rochelia* in the monotypic tribe *Rochelieae* with the two species *R. stellulata* Reichenb. and *R. leiocarpa* Ledeb. Popov (1953) while keeping the genus in the tribe, following Zakirov's treatment (1941, cited therein) divided it into two sections, *Eurochelia* Zak. and *Cryptocarpa* Zak. Section *Eurochelia* was further divided into two series: *Stellulatae* Zak. and *Pedunculares* Zak. *Cryptocarpa* is a monotypic section (*R. cardiosepala* Bge.). Later on, Riedl (1967) in the Flora Iranica reduced the tribe to the subtribal level, *Rocheliinae*, within the tribe *Eritrichieae*. Based on the number of sepals, he reclassified the genus into two subgenera, *Rochelia* (with 5 sepals) and the monotypic Neo-*Rochelia* H. Riedl (with 9-10 sepals). The former subgenus was, in turn, divided into two sections, *Rochelia* (=*Eurochelia*) and *Cryptocarpa*. The section *Rochelia*, as a multi-species taxon, is characterized by linear or lanceolate sepals converging/recurving fruit but not enclosing it. Whereas, *Cryptocarpa* is distinguished by its cordate calyx completely enclosing fruit. Riedl (1967) substituted series *Stellulatae* and *Pedunculares* of section *Eurochelia* as subsections *Rochelia* and *Pedunculares* Zak., respectively. Several works were performed on the genus using non-molecular data. On the basis of nutlet micromorphology, Hilger (1984) tried to group the species of the genus. Pollen morphological and karyological studies (Diez & Benito 1991, Kazempour Osaloo 1991, Kazempour Osaloo 2001, Luque 1992) implied that *Rochelia* is related to *Lappula* Gilib. Hiltbert, no relatively comprehensive study on the molecular phylogeny of the genus has been conducted. Out preliminary phylogenetic analyses using either nrDNA ITS or chloroplast trnL intron and trnL-trnF intergenic spacer (hereafter abbreviated as trnL-F) sequences for 40 species of the tribe *Eritrichieae* and related tribes showed that *Rochelia* with three species sampled, formed a monophyletic group as sister to *Lappula* (Khoshsokhan et al. 2008, Khoshshokhan & Kazempour Osaloo 2008). The internal transcribed spacer (ITS) is the region of the 18S-5.8S-26S nuclear ribosomal cistron. The spacers contain the signals needed to process the rRNA transcript (Baldwin 1992, Baldwin et al. 1995) and have often been used for inferring phylogeny at the
generic and infrageneric levels in plants (e.g., Baldwin 1992, Baldwin et al. 1995, Kazempour Osaloo et al. 2003 & 2005, Ahangarian et al. 2007). \(\text{trn}L-F\), is the chloroplast DNA (cpDNA) sequence that is now widely used to investigate interspecific/generic relationships among angiosperms and other plants using the universal primers of Taberlet et al. (Taberlet et al. 1991, Shaw et al. 2005).

In this study, we attempt to infer infra-generic relationships in \textit{Rochelia} and evaluate character evolution among its species in the context of the combined nrDNA ITS-\(\text{trn}L\)-F phylogeny.

MATERIALS AND METHODS
A total of seven species representing two sections of the subgenus \textit{Rochelia} and a single species of subgenus \textit{Neo-Rochelia} plus 2 \textit{Lappula} species as outgroups, were included in molecular studies (Table 1). Total genomic DNA was extracted from fresh and dried leaves, using a modification of the 2X CTAB protocol of Doyle and Doyle (1987). The nrDNA ITS region was amplified as a sharp single fragment using the primer pair ITS5/ITS5m and ITS4 in all cases (White et al. 1990, Sang et al. 1995). The \(\text{trn}L\)-F region was amplified using primers \textit{c} and \textit{f} as one fragment (Taberlet et al. 1991). Each fragment was directly sequenced using the Big dye terminator cycle sequencing ready reaction kit with the same primers. Sequencing of the fragments was done in an ABI Prism 3730xl DNA Analyzer (Applied Biosystems, USA).

Sequence alignment
Sequences were edited using BioEdit ver. 7.0.9.0 (Hall 1999) and aligned using ClustalX (Larkin et al. 2007) followed by manual adjustment. Alignment of each dataset required the introduction of several single and multiple-base indels (insertions/deletions). Positions of indels were treated as missing data for all datasets.

Phylogenetic analyses
Phylogenetic analyses were performed on the aligned nrDNA ITS and \(\text{trn}L\)-F data matrices separately and in combination. These datasets were analyzed using Maximum parsimony (MP) criterion as implemented in PAUP* (Swoford 2002). Heuristic searches were performed with 100 replicates of random addition sequence, tree-bisection-reconnection (TBR) branch-swapping with MulTrees on and steepest descent off. Bootstrap values (Felsenstein 1985) with 1000 replications were calculated using the heuristic search option, simple sequence addition and TBR branch swapping. To assess combinability of these datasets, incongruent length difference (ILD, Farris et al. 1995) test was conducted using PAUP*. ILD test suggested that the both datasets were not incongruent (P=1). To examine evolutionary trend of morphological characters, we mapped six diagnostic ones on the combined nrDNA ITS-\(\text{trn}L\)-F phylogeny.

RESULTS
Phylogenetic analyses
Length of nrDNA ITS1-5.8S-ITS2 sequences for \textit{Rochelia} was 632 nucleotide sites, of which 22 sites were parsimony informative. MP analysis of the dataset resulted in 4 equally most parsimonious trees having a length of 35 steps and consistency index (CI)= 0.714 and a retention index (RI) = 0.697 (excluding uninformative characters). The aligned \(\text{trn}L\)-F dataset comprised of 866 nucleotide sites, of which 5 were informative. MP analysis of this dataset resulted in a single most parsimonious tree with a length of 5 steps (CI= 1, RI= 1). nrDNA ITS and \(\text{trn}L\)-F phylogenies are conflicting on the position of \textit{R. macrocalyx} Bge.; whereas, on the latter tree, it is a unresolved branch. The low resolution in \(\text{trn}L\)-F phylogeny is mainly due to the low number of parsimony informative characters. As noted in the material and methods section, ILD test suggested the incongruency of both datasets. The combined nrDNA ITS-\(\text{trn}L\)-F dataset was composed of 1498 nucleotide
Fig. 1. Strict consensus tree of 4 most parsimonious trees resulting from phylogenetic analysis of the combined nrDNA ITS - cpDNA trnL-F sequences for *Rochelia* and two outgroup taxa (Length= 40 steps, CI= 0.750, RI= 0.737). Numbers above branches are bootstrap values for 1000 replicates analyses; values < 50% are not indicated.

sites, of which 27 were parsimony informative. MP analysis of the dataset resulted in 4 equally most parsimonious trees with a length of 40 steps with a CI= 0.750 and an RI= 0.737. The strict consensus tree of these trees, the same as that of nrDNA ITS tree, with accompanying bootstrap values was presented in Fig. 1. On this tree, *Rochelia persica* Bge., *R. disperma* (L. f.) C. Koch. and *R. cancellata* are unresolved branches as sisters to a clade of the remaining five species examined. This clade was, in turn, composed of two subclades (A and B). The first subclade (A) contained well allied *R. bungei* Trautv. and *R. mirheydari* Riedl & Esfandiar (bootstrap value of 97%) and the second subclade (B) constituted a relatively weakly supported group (72% bootstrap) comprising *R. macrocalyx* and *R. cardiosepala*.

**Character evolution**

We here mapped six diagnostic characters on the strict consensus tree resulting from the combined nrDNA ITS-trnL-F dataset (Figs 2A-F). Character 1, tip of calyx hairs, has been undergone reversal evolution from hamate hairs to non-hamate ones in both *Rochelia persica* and *R. bungei* (Fig. 2A). The second character, fruit pedicel to sepal ratio [longer pedicel (>1)], is a synapomorphy for the clade comprising *R. bungei*, *R. mirheydari*, *R. macrocalyx*, *R. peduncularis* and *R. cardiosepala*. This character state is evolved from shorter pedicel (<1) (Fig. 2B). Character 3, shape of calyx lobes, have been changed from narrowly linear calyx through wide lanceolate/rectangular in *R. macrocalyx* and *R. peduncularis* to cordate only in *R. cardiosepala* (Fig. 2C). Sepal with prominent midrib is a synapomorphy for *R. macrocalyx*, *R. peduncularis* and *R. cardiosepala*, that has been derived from non-prominent midrib (character 4) (Fig. 2D). Character 5, medium-sized nutlets, > 3mm in length, is unique to *R. peduncularis* and *R. cardiosepala*, that is in fact a reversal from small-sized nutlets (≤ 3 mm) (Fig. 2E). Nutlets completely clasping the gynobase, character 6,
1. Tip of calyx hairs
- Non-hamate
- Hamate

2. Fruit pedicel to sepal ratio
- < 1
- = 1
- > 1
- equivocal

3. Shape of calyx lobes
- Narrowly linear
- Rectangular/ lanceolate
- Cordate

4. Sepal midrib
- Non-prominent
- Prominent
is evolved in parallel between *R. cancellata* plus *R. peduncularis* and *R. cardiosepala* (Fig. 2F).

**DISCUSSION**

The present data did not support the monophyly of the subgenus *Rochelia*. Likewise, its species rich section, *Rochelia*, due to inclusion of the monotypic section *Cryptocarpa*, is paraphyletic. Furthermore, the two subsections of *Rochelia*, viz, *Rochelia* and *Pedunculares* appeared to be non-monophyletic. The subsection *Rochelia* was represented herein by 4 species, two of which, *R. disperma* and *R. persica*, along with *R. cancellata* of the subgenus *Neo-Rochelia* are unresolved branches. The two others, *R. bungei* and *R. mirheydari* are the closest sister taxa (subclade A), as unified with a subclade of subsection *Pedunculares* and the monotypic section *Cryptocarpa* (subclade B, Fig.1). *Pedunculares* is a small subsection with 5 species, 2 of which, *R. peduncularis* and *R. macrocalyx*, were included herein. They were not united as sister species, but the former species is well allied with *R. cardiosepala* of the section *Cryptocarpa* (see also Hilger 1984). The derived position of *R. cardiosepala* within the section *Rochelia* indicates that this species should be classified within it, and thus, the sectional status of *Cryptocarpa* is no longer tenable. The species is characterized by specialized autapomorphies including cordate calyx lobes and completely invisible nutlets in calyx (Fig. 2C).

All six diagnostic characters examined herein but non-hamate tip of calyx hairs and nutlets completely clasping the gynobase, were not homoplasious for *Rochelia* species (see Fig. 2). The shorter pedicel is evolved once to longer pedicel in the clade of five species comprising the *R. mirheydari* through *R. cardiosepala* (Fig. 2B). The character 3, shape of calyx lobes underwent evolutionary changes twice terminating to cordate calyx in *R. cardiosepala*. These character states were evolved from narrowly linear calyx to wide lanceolate/rectangular one in both *R. macrocalyx* and *R. peduncularis* (Fig. 2C). The next character is sepal with prominent midrib which is a synapomorphy of the subclade of *R. macrocalyx*, *R. peduncularis* and *R. cardiosepala* (Fig. 2D). In the regional Floras, e.g., USSR, Iranica, Iran and Pakistan (Popov 1953, Khatamsaz 2002, Riedl 1967 and Nasir 1989), this feature along with the two later ones, were mostly used a key characters to separate them from other species. The two other characters, nutlets size, (medium) and attachment of nutlets to the adaxial part of gynobase (completely clasping) are shared by *R. peduncularis* and *R. cardiosepala*. The latter character state is also found in the other unrelated species, *R. cancellata* (Fig. 2F). In terms of this feature (nutlets
clasping the gynobase), Hilger (1984) suggested, however, that these three species are closely related.

CONCLUDING REMARKS

The present phylogenetic hypothesis showed that the infrageneric classification of *Rochelia*, at least at the sectional and subsectional level based upon traditional morphological characters is artificial. To treat a comprehensive circumscription of the genus at the infrageneric level, more taxa and other fast evolving DNA fragments are definitely necessary.

ACKNOWLEDGMENTS

We thank staff of Ferdowsi University of Mashhad Herbarium for providing a duplicate of *Rochelia mirheydari* and *Rochelia peduncularis* and our cordial thanks also due to S. Karaman and B. Bani of Gazi University, Ankara for providing some specimens of *R. cancellata* for DNA extraction.

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