Parallel Evolution of Leaf Pubescence in *Encelia* in Coastal Deserts of North and South America

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Summary. Members of the shrub genus *Encelia* occur in the arid coastal regions of western North America and South America along gradients of precipitation extending from less than 50 mm annually to somewhat over 350 mm. At moist ends of the gradient species possess pubescent, green leaves. Proceeding to drier regions, the species possess progressively more pubescent, white leaves. Leaf absorbance to solar radiation decreases with precipitation in an identical pattern along gradients in both North and South America.

Introduction

Along gradients of decreasing precipitation in arid lands of southwestern North America, there are numerous species that increase in leaf pubescence (hairiness) or are replaced by more pubescent-leaved species in response to increased aridity (Ehleringer 1981a). Recent studies of such species suggest that leaf pubescence results in increased reflectance to solar radiation, lower leaf temperatures and lower rates of water loss (Ehleringer et al. 1976; Ehleringer and Björkman 1978; Ehleringer 1981b). The most studied examples of pubescence increase are found in the genus *Encelia*. For the pubescent-leaved shrub *Encelia farinosa*, the presence of hairs is apparently an important feature which enables this species to successfully occupy hot, desert habitats (Ehleringer and Mooney 1978).

Members of the genus *Encelia* are common in the arid regions of southwestern North America (Munz 1959; Shreve and Wiggins 1964). They also occur in the coastal, arid regions of northern Chile, Peru, and the Galápagos Islands in South America (Blake 1914; Wiggins and Porter 1971). The taxonomic descriptions of *Encelia* species suggest that as in North America both glabrate and pubescent-leaved species occur in South America.

The purpose of this study was to compare the distributions of coastal *Encelia* species in North and South America to determine if parallel evolution of leaf pubescence and hence spectral properties was occurring within this genus. Specifically we wanted to know whether at points of homocline between North and South America were plants found with comparable degrees of leaf pubescence. If so, this would provide strong evidence for the role of climate, specifically aridity, in the selection of leaf morphological properties.

Methods

*Encelia* species were sampled along gradients of decreasing precipitation. In North America species were sampled along a coastal transect from Laguna Beach, California (33° 33' N, 117° 47' W) south to Magdalena Bay, Baja California Sur (24° 38' N, 112° 09' W) during August 1979. In Chile species were sampled along a coastal transect from Fray Jorge (30° 34' S, 71° 11' W) north to Antofagasta (23° 29' S, 70° 26' W). Climatic data for these regions are available from U.S. Weather Bureau Records, Hastings and Humphrey (1969), Almeida and Saxe (1958), and Hajek and di Castri (1975).

The species distributions were determined by examination of herbarium vouchers from the California Academy of Sciences, San Diego Natural History Museum, Stanford University, University of California, Berkeley and the University of Arizona Herbarium as well as by personal observations.

Leaf absorbance to solar radiation was measured using a 23 cm diameter Ulbricht integrating sphere. Light to the integrating sphere was either sunlight (via a heliostat) or monochromatic (Bausch and Lomb monochromator). The leaf absorbance is a percentage measurement and thus changes in leaf absorbance are changes in the percentage of light absorbed by the leaf. Further details of the theory and measurement of leaf absorbance using this technique are described in Ehleringer and Björkman (1978) and Ehleringer (1981b).

Results

The distributions of coastal *Encelia* species in North America is essential parapatric. The distribution boundaries are distinct and appear to be associated with changes in vegetation type. While the species or varieties typically do not occur together, there are rare regions of overlap, and in these zones hybrids between the two species are evident. As shown in Fig. 1 the distribution of *Encelia californica* Nutt. extends from Pt. Sal, California (34° 33' N, 120° 36' W) south to El Rosario, Baja California Norte (30° 03' N, 115° 43' W), where it is a dominant component of the coastal sage vegetation type. In the Valle El Rosario (30° 03' N, 115° 43' W) *Encelia asperfolia* (Blake) Clark and Kyhos first appears and extends south to the beginning of the Vizcaíno Desert at Villa Jesus Maria, Baja California Norte (28° 18' N, 114° 00' W). *Encelia palmeri* Vasey and Rose is quite common throughout the low elevation Vizcaíno Desert region from Villa Jesus Maria south to Punta Conejo, Baja California Sur (24° 04' N, 111° 01' W).

In Chile, the distributional boundaries of *Encelia* species are somewhat more difficult to establish because of the small number of collections which have been made. *Encelia canescens* var. *oblongifolia* (DC.) Blake occurs in the xerophytic vegetation from Termus del Socó (30° 43' S, 71° 38' W) north to about Choros Bajas (29° 17' S, 71° 44' W) (Fig. 2). Farther north in more arid and open shrubland, *Encelia canescens* Lam. occurs from Ca-chiyuyu (29° 05' S, 70° 54' W) to Caldera (27° 03' S, 70° 50' W). *Encelia canescens* var. *lanuginosa* is found just south of...
Chañaral (26° 23' S, 70° 37' W) extending north into the driest regions of the Atacama Desert.

Along the coastal transects in both North and South America Encelia taxa are replaced by increasingly more pubescent-leaved forms as precipitation levels decrease. Encelia californica and E. canescens var. oblongifolia have glabrate leaves and occur in the wettest regions (100–350 mm annually). When annual precipitation decreases to 50–100 mm, these taxa are replaced by the moderately pubescent leaved species E. asperifolia and E. canescens, respectively. On the driest sites where precipitation levels are below 50 mm annually, the heavily pubescent leaved species E. palmeri and E. canescens var. lanuginosa are found.

Climate diagrams were constructed for representative locations within the distributions of the six Encelia taxa to determine if there was homology in the macroclimates where they occurred. Figure 3 illustrates the climates for the three matched pairs, the South American climate diagrams being offset by six months. There is a strong similarity in macroclimates of morphologically matched pairs in North and South America, but not between adjacent taxa in the same continent, indicating that Encelia species with similar pubescence morphologies are found in similar climates.

The extent to which there is a functional parallel evolution of leaf pubescence in these Encelia species is indicated by the leaf absorbance to solar radiation. Changes in pubescence density should change the leaf absorbance and reflectance characteristics.

On both continents leaf absorbance over the visible wavelengths was measured on samples of all taxa in the field at the end of the growing season. In both the North and South American Encelia taxa, the leaf absorbance was high (84–85%) at the wet end of the aridity gradient (Fig. 4). Proceeding to drier sites in both continents, leaf absorbances decreased. The decreases in leaf absorbance were greatest at the driest sites where absorbances were 46 and 44% in North and South America, respectively. The changes in leaf absorbance with precipitation were identical for North and South American Encelia species.

The spectral characteristics of pubescent leaved species change with the extent of drought imposed upon the plant. Figure 5 shows the variations in spectral characteristics seen for leaves of E. canescens var. lanuginosa, a species from extremely arid lands, grown under well-watered and droughted conditions in the greenhouse. Under non-droughted conditions leaf pubescence is sparse, percent absorbance at all visible wavelengths is high, and the integrated 400–700 nm absorbance is 80%.
In contrast, under droughted greenhouse conditions the leaf pubescence becomes dense, the percent absorbance is decreased at all wavelengths, and the integrated 400-700 nm absorbance is 52%.

Discussion

Parallel and convergent evolution has been described extensively in the Mediterranean-type climate plant communities of California (North America) and Chile (South America) by di Castri and Mooney (1973), Mooney (1977), and Thrower and Bradbury (1977). Additionally, Mooney et al. (1974) have been able to demonstrate a similar convergence at the community level in the coastal desert regions which border the Mediterranean-type climate communities. At a finer level of resolution in the Mediterranean-type climates, Mooney et al. (1974) and Mooney (1977) have been able to show a strong degree of convergence in the morphology, phenology, and physiology of ecological pairs from the two continents. The current study extends these findings (at least for morphology) to species pairs in these desert regions.

What is the adaptive value of pubescence to these species? Recent studies suggest that pubescence or other reflectance increasing structures (such as waxes or salt glands) 1) reduce water loss and 2) are a necessary component of photosynthetic adaptation to arid habitats (Mooney et al. 1978; Ehleringer and Mooney 1978; Ehleringer 1981a; Ehleringer 1981c). For photosynthetic adaptation, it appears that perennial desert species either physiologically acclimate or morphologically acclimate to changes in the thermal environment (Ehleringer 1981c). Physiological acclimation involves a biochemical/physiological change in the photosynthetic system such that the leaf photosynthetic capacity is enhanced at the new temperature. Morphological acclimation occurs in those species which apparently do not physiologically acclimate. In morphological acclimation, there is a change in the leaf absorbance characteristics such that as air temperatures increase there is a decrease in leaf absorbance, and as a consequence leaf temperatures remain either constant or close to a fixed thermal optimum for photosynthesis. Morphological acclimation has been shown to occur in Encelia farinosa, an interior desert species (Ehleringer and Mooney 1978). The pubescence in the coastal desert Encelia may also prevent damaging high leaf temperatures. However, no data are available regarding the photosynthetic capacities of any of the coastal, pubescent species.

The leaf absorbance-precipitation response curve shown in Fig. 4 infers that coastal Encelia cannot physiologically acclimate their photosynthetic apparatus and therefore must morphologically acclimate. This morphological acclimation probably occurs with the onset of drought (Fig. 6). Thus, early in the growing season, after initial rains have fallen, the pubescence is minimal on all leaves along the precipitation gradient, resulting in high leaf absorbances and high photosynthetic potentials by all leaves. As drought develops and air temperatures increase, the plants at the driest end of the aridity transect respond first and produce lightly pubescent leaves with lowered leaf absorbances. This allows the leaves to remain close to the optimum temperature for photosynthesis. As the drought develops further and air temperatures increase more, the leaves produced are more pubescent and have progressively lower absorbances until at the end of the growing season the pattern observed in Fig. 4 is obtained.

The similarity in the extent of leaf pubescence development in Encelia in similar climates of North and South America and other California-Chile convergence studies suggests that by understanding how plants have adapted to the environment in one location it may allow us to a priori predict the adaptive modes expected to be found in similar environments elsewhere in the world.

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