Phylogenetic Relationships of the Sweetpotato

[Ipomoea batatas (L.) Lam.]

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Abstract. Twenty-four accessions of Ipomoea, representing 13 species of section Batatas and the outgroup species I. gracilis and I. per-caprae were analyzed for restriction fragment length polymorphisms. Polymorphisms were detected by probing Southern blots of restriction enzyme-digested genomic DNA with 20 low or moderate copy number sequences isolated from an I. batatas cv. Georgia Red genomic library. Data were analyzed cladistically and phenetically.

Ipomoea trifida, I. tabascana, and collection K233 are, of the materials examined, the most closely related to sweetpotato (I. batatas). Ipomoea littoralis, the only Old World species in the section, is a sister species to I. tilaea. Ipomoea littoralis, I. umbratica, I. peruviana, I. cynanchifolia, and I. gracilis are shown to be diploid (2n = 2x = 30). In contrast, I. tabascana is tetraploid (2n = 4x = 60). The intrasectional relationships of section Batatas species and the role of tetraploid related species in the evolution of the cultivated I. batatas are discussed.

Sweetpotato production worldwide exceeded 1.3 x 10^6 t in 1989 (Food and Agriculture Organization, 1990), and the enormous potential of this crop as a carbohydrate source is widely recognized. To increase current production levels, new genes for resistance to various biotic and abiotic factors must be identified. Resistance to many important diseases and insects does not appear to be present in the I. batatas gene pool, which has focused attention on the use of exotic germplasm for sweetpotato improvement (Iwanaga, 1988; Orjeda et al., 1990). Nishiyama and Teramura (1962) were the first to use exotic germplasm in the form of the feral sweetpotato segregate K123 (Jones, 1967). Resistance to sweetpotato weevil (Cylas spp.), scab [Elsinoe batatas (Saw.) Viegas and Jenkins], and black rot (Cerato cystis fimbriata Ell. et Halst.) have been identified in plants reported as I. trifida and I. littoralis (Iwanaga, 1988). While the potential for the use of most section Batatas species in sweetpotato improvement has yet to be ascertained, other distantly related species contribute indirectly; I. setosa L. and I. carnea ssp. fistulosa Mart., ex Choisy are used as virus-indicator and flower-inducing rootstock, respectively. Increased awareness of the potential contribution of Ipomoea spp. to sweetpotato improvement is reflected in the recent efforts to collect and establish gene banks of these materials (de la Puente, 1988; Jarret et al., 1989).

Until recently, species boundaries in Ipomoea sect. Batatas were poorly understood. As a result, many species names have been misapplied in the taxonomic and agronomic literature, and different workers have often described similar and/or identical materials under different names, or material of distinct species under the same name. Austin (1978) provided a taxonomic revision of section Batatas based on the study of several hundred specimens, including type specimens of most of the species. He recognized 11 species and three other taxa believed to be of recent hybrid origin. This revision provides a soundly based outline of the taxonomy and nomenclature of Ipomoea sect. Batatas. Since 1978, three additional species have been recognized in the section [I. peruviana, formerly considered a synonym of I. batatas; I. umbratica, formerly placed in sect. Eriopsernum; and I. tabascana], a recently described endangered species from Mexico (Austin, 1988; Austin et al., 1991; McDonald and Austin, 1990)]

One unresolved taxonomic problem that is particularly relevant to the understanding of sweetpotato evolution involves the relationship of I. batatas to a series of wild tetraploid plants that closely resemble cultivated sweetpotato in the structure of their leaves, flowers, and fruits. Mexican accessions of such tetraploids have been identified in the literature as I. gracilis (Jones, 1970), I. littoralis (Nishiyama et al., 1975), I. trifida (Shiotani and Kawase, 1987), I. batatas (Austin, 1988), or descendants of recent hybrids between cultivated sweetpotatoes and wild diploid weeds such as I. trifida (Austin, 1977). K233 is representative of one form of this latter group of plants.

To understand the origins of the cultivated sweetpotato, the relationships of the wild species with one another and with the cultivated sweetpotato must be more fully understood. However, little is known about the relationships of the species in this group. Austin (1988) has presented several phenetic analyses based on morphological characters. However, the morphological similarity of the species and the great plasticity of morphological characters in this group make it difficult to use morphological characters alone for phylogenetic reconstruction. We have, therefore, used restriction fragment length polymorphisms (RFLPs) for our phylogenetic study of this section. RFLPs can provide many genetic markers of high heritability. They have proven valuable for phylogenetic studies in several crop genera, including Lycopersicon (Miller and Tanksley, 1990), Brassica (Song et al., 1988), Solanum (Debener et al., 1990), Lens (Havey and Muehlbauer, 1989), and Glycine (Menacio et al., 1990).

Materials and Methods

Plant material

Plant material (Table 1) was, unless noted otherwise, obtained from the U.S. Sweetpotato Germplasm Repository (Jar-
Leaf tissue was collected from individual plants started from seed, frozen at –135°C, freeze-dried, ground to a fine powder in liquid N, and stored desiccated at –20°C until used. Accessions of *Ipomoea batatas* were obtained from an in vitro collection (Jarret, 1989), acclimated to soil in a quarantine greenhouse, and leaf tissue was harvested from individual plants. Leaf tissue of *I. littoralis* was harvested from a single accession of this species, collected in Queensland, Australia, in 1989, and maintained in quarantine in Griffin. Herbarium specimens of all plant material used in this study are available from the Southern Regional Plant Introduction Station.

DNA isolation, digestion, electrophoresis, and blotting

DNA was extracted from lyophilized leaf tissue following a procedure modified from that of Murray and Thompson (1980) and G. King (NPI, Salt Lake City, Utah, personal communication) as described by Gawel and Jarret (1991). Precipitated DNA was collected by centrifugation, resuspended in 400 µl of 1 M NaCl with heat (65°C for 30 min), and transferred to microtubes. Undissolved material was removed by centrifugation at 16,000 × g for 2 min, the supernatant was transferred to a new tube, and the DNA was ethanol-precipitated (Maniatis et al., 1982). DNA (3 to 5 µg) was digested with 9 to 15 U (U = unit of enzyme activity) of *Eco* RI, *Bam* HI, or *Msp* I for 8 h at 37°C. Fragments were separated on 0.8% agarose (BioRad) gels at 50 V for 20 h in TEA buffer. Following electrophoresis, the gels were denatured, neutralized, and Southern-blotted to nylon membrane following the membrane manufacturer’s recommendations (BioTrans-ICN, Irvine, Calif.). DNA was bound to the membranes by exposure to ultraviolet radiation (Stratagene, La Jolla, Calif.).

Library construction

Total genomic DNA, isolated as described above from ‘Georgia Red’, was further purified on cesium chloride (Maniatis et al., 1982), ethanol-precipitated, and digested with *Eco* RI. *Eco* RI-digested genomic DNA was ligated into dephosphorylated *Eco* RI-digested pUC 18 and used to transform *Escherichia coli* strain LL308. Recombinant plasmids were screened on X-gal and their insert size determined. Plasmids, with insert sizes from 0.5 to 1.0 kb, were cloned and their DNA isolated (Maniatis et al., 1982). Approximate copy number was determined by probing dot blots of plasmid DNA with *32*P-labeled *Ipomoea batatas* cv. Georgia Red genomic DNA. Inserts were isolated from 20 plasmids, bearing low copy number sequences, on low melting-point agarose and were random primer-labeled (BRL) with 50 µCi (1 Ci = 37 GBq) of *32*P-dCTP (NEN, Dupont, Wilmington, Del.). Unincorporated nucleotides were removed by chromatography on Sephadex G50 (Maniatis et al., 1982).

Hybridizations and autoradiography

Membranes were prehybridized for 4 to 6 h in a prehybridization solution containing 6 × SSC, 0.001% sonicated denatured salmon sperm (SS), 5 × Denhardt’s solution, and 0.1% SDS at 65°C. Hybridizations were carried out in 6 × SSC, 0.001% SS, 0.1% SDS, and denatured labeled plasmid DNA. Membranes were hybridized overnight at 65°C and washed successively in 2 × SSC, 0.1% SDS; 1 × SSC, 0.1% SDS; and 0.1 × SSC, 0.1% SDS for 30 min each at 65°C. Membranes were

| Species                          | Identifier  | Sample no. | Country of origin |
|---------------------------------|-------------|------------|-------------------|
| *Ipomoea batatas* (L.) Lam. (2n = 90) | P1538295    | 97         | Peru              |
|                                 | P1538300    | 98         | Peru              |
|                                 | Q27990      | 99         | New Guinea        |
| *I. cordato-triloba* Dennstedt   | P1518495    | 14         | Mexico            |
|                                 | P1540710    | 43         | Colombia          |
| *I. cynanchifolia* Meinsh.       | CIP460149   | 15         | Brazil            |
| *I. gracios R. Brown*            | P1538370    | 01         | Australia         |
| *I. lacunosa* Blume              | 67.36º      | 11         | United States     |
| *I. littoralis* Blume            | 67.36º      | 22         | Australia         |
| *I. peruviana* O’Donell          | CIP46025º   | 19         | Peru              |
| *I. pes-caprae* (L.) R. Brown    | P1518492    | 21         | Mexico            |
| *I. ramosissima* (Poir.) Choisy   | CIP460005º  | 18         | Peru              |
|                                 | CIP460036º  | 41         | Bolivia           |
| *Ipomoea sp.*                    | K233º       | 24         | Mexico            |
| *I. tabascana* McDonald & Austin | P1518473    | 90         | Mexico            |
| *I. tenuissima* Choisy*          | DLP2925º    | 12         | Mexico            |
| *I. tilliae* (Wild.) Choisy      | P1530994    | 51         | Dominican Republic|
|                                 | P1540731    | 70         | Colombia          |
|                                 | P1530998    | 53         | Dominican Republic|
|                                 | P1540722    | 17         | Colombia          |
|                                 | P1540724    | 80         | Colombia          |
|                                 | P1543818    | 81         | Costa Rica        |
| *I. umbraticola* House°          | 67.36º      | 74         | Mexico            |

*°Material obtained from J.A. McDonald, Univ. of Texas at Austin.
*From D.F. Austin, Florida Atlantic Univ., Boca Raton.
*From A. Jones, U.S. Dept. of Agriculture, Agricultural Research Service, 2875 Savannah Highway, Charleston, S.C.
°International Potato Center, Lima, Peru.
Root-tip chromosome counts

Procedures for examination of somatic chromosomes were essentially as described by Jones and Kobayashi (1968). Fixed root tips were hydrolyzed in 0.2 N HCl for 30 min at 65°C, allowed to cool for 30 min at room temperature, and stained with aceticorcin. Chromosomes were counted in a minimum of 10 cells per species examined.

Data analysis

Data were compiled into a 0-1 matrix and subjected to cladistic analysis by Wagner parsimony using the SWAP= GLOBAL and MULPARS = ON options of PAUP version 2.4 (Swofford, 1985). To evaluate the strength of the resulting cladase, the data were analyzed by the bootstrap method of Felsenstein (1986). Using the BOOT routine in PHYLIP (version 3.4), 100 bootstrap samples were generated by random resampling of the data set (Felsenstein, 1985) and separately subjected to Wagner parsimony analysis. Since bands visualized using the same probe may not be genetically independent (Gawel et al., 1992), they were linked using the FACTOR option of BOOT. The confidence level of each clade (the bootstrap statistic) is equal to the percentage of bootstrap trees in which that clade appears. The bootstrap values are plotted on the majority-rule consensus tree that the statistical support for different clades varied from 29% to 100%. Three clades showed significant (95% or better) support (bootstrap statistic = 95%) (Fig. 1). This result supports the traditional view that I. trifida and the wild Mexican tetraploids are the most likely ancestors of cultivated sweetpotato (Austin, 1988) and indicates that the recently described species I. tabascana is also a very close relative of the crop plant. The species included in this group are classified genome B (Nishiyama et al., 1975). Other divisions within the section clearly transcend the boundaries of ploidy, compatibility, and genome type (Table 2).

The closest relatives of cultivated sweetpotato seem to be the wild Mexican tetraploids (bootstrap statistic = 96%). The most strongly supported clade in the tree (bootstrap statistic = 100%) consists of the cultivated sweetpotato, the tetraploid K233, and seven other species: I. cordata-triloba, I. cynanchifolia, I. lacunosa, I. tabascana, I. tenuissima, I. trifida, and I. triloba. This group contains all the closest relatives of the cultivated sweetpotato; all other taxa share fewer than half of the DNA fragments examined with I. batatas and must be considered more distantly related. Studies of the evolution of the sweetpotato should concentrate on a closer examination of the species in this group.

The closest relatives of cultivated sweetpotato seem to be the wild species I. trifida and I. tabascana, and the wild Mexican tetraploid represented by accession K233 (Fig. 2). The close relationship of these four taxa is strongly supported by our data (bootstrap statistic = 95%) (Fig. 1). This result supports the
accessions in our collection from Veracruz, Mexico, that are morphologically very similar to K233 and identical to accessions identified by McDonald as *I. batatas* var. *apiculata*. We suggest that K233 is a segregate of *I. batatas* var. *apiculata*, which explains its placement close to *I. batatas*. Chromosome counts (data not shown) indicate that *I. batatas* var. *apiculata* is tetraploid (2n = 4x = 60).

A role for the various tetraploid forms of *I. batatas* in the evolution of the cultivated hexaploid sweetpotato is unclear. Tetraploid materials examined to date, including *I. tabascana* (Table 2), have not demonstrated an ability to form storage roots. However, this failure has not yet been systematically examined. According to Martin et al. (1974), roots of their tetraploid accessions, when cut from the parent plant, were able to sprout. We have not verified this characteristic with K233 or *I. tabascana*. *Ipomoea trifida* (2n = 2x = 30) has a distinct perennial rooting habit (R.L.J. and A.W., unpublished data).

The hybrid members of the section, *I. × leucantha* Jacquin (Abel and Austin, 1973, 1980) and *I. × grandifolia* (Dammer O’Donnell (Austin, 1978), were not included in this analysis. Plant material of *I. × grandifolia* was not available. A hybrid origin for *I. × grandifolia* was proposed by Austin (1978). However, recent examination of newly acquired materials indicate that *I. × grandifolia* is a distinct species (D.F. Austin, personal communication) and should be included in future analyses.

Although less strongly supported by the bootstrap statistics, Figs. 1 and 2 illustrate a relationship between the North American species *I. lacunosa*, *I. tenuissima*, and *I. cordato-triloba*. Closely associated with these is *I. trifida* and, more distantly, *I. cynanchifolia*. *Ipomoea trifida*, endemic to the Caribbean and southern Florida, has also been suggested as a close relative to the sweetpotato (Austin, 1988). The South American species *I. peruviana*, *I. tiliacea*, and *I. ramosissima*; the Central American/Mexican species *I. umbraticola*, and the Old World species *I. littoralis* and *I. gracilis* appear to be more distantly related to *I. batatas* (Fig. 1).

Root-tip chromosome counts indicate that *I. peruviana*, *I. umbraticola*, *I. cynanchifolia*, *I. littoralis*, and *I. gracilis* are diploid, 2n = 2x = 30 (Table 2).

Although the three major clades in Fig. 1 are well supported, others parts of the cladogram are less well resolved (bootstrap statistics < 80%). These low bootstrap values reflect the relatively few unique character state changes supporting these clades. The level of within-species morphological and genetic (RFLP) variation in many of these species is high (A.W. and R.L.J., unpublished data on *I. trifida*; R.L.J., unpublished data on *I. batatas*). The topology of the cladogram is determined partly.

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**Table 2.** Ploidy, compatibility, and genome group of *Ipomoea* spp. and accessions. Chromosome counts, unless noted otherwise, are from Jones (1974).

| Species       | Ploidy | Self-compatible | Genome |
|---------------|--------|-----------------|--------|
| *I. batatas*  | 2n = 90| No              | B      |
| *I. cordato-triloba* | 2n = 30| Yes             | A      |
| *I. cyananchifolia* | 2n = 30| Yes             | A      |
| *I. gracilis* | 2n = 30| No              | B      |
| *I. lacunosa* | 2n = 30| Yes             | A      |
| *I. littoralis* | 2n = 30| No              | B      |
| *I. peruviana* | 2n = 30| No              | B      |
| *I. pes-caprae* | 2n = 30| Yes             | A      |
| *I. ramosissima* | 2n = 30| Yes             | A      |
| *I. sp. (K233)* | 2n = 60| No              | B      |
| *I. tabascana* | 2n = 60| No              | B      |
| *I. tenuissima* | 2n = 30| Yes             | A      |
| *I. tiliacea* | 2n = 60| No              | B      |
| *I. trifida* | 2n = 60| No              | B      |
| *I. triloba* | 2n = 60| Yes             | A      |
| *I. umbraticola* | 2n = 60| Yes             | A      |

* Nishiya et al. (1975).
* This paper.

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by the chance selection of particular genotypes from the complex array of genotypes that comprise each species. Further, natural hybridization is known to occur between species of this group (Abel and Austin, 1980; Austin, 1978). Intrgressive gene flow between clades decreases the amount of genetic differentiation between species and weakens the historical association between the characters and the original phylogeny, thus decreasing the statistical confidence of the phylogenetic reconstruction. To improve the analysis, more individuals of these taxa, in combination with additional characters, are needed.

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