‘Evergreen’ Peach, Its Inheritance and Dormant Behavior

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Abstract. The evergreen (EVG) peach, first described in Mexico, was used as a parent with deciduous (DE) peaches to develop F1 and F2 hybrid populations in Mexico, Florida, Georgia, and West Virginia. F1 trees were DE and F2 plants segregated 3 DE : 1 EVG. In West Virginia, the most temperate location, the heterozygous class could be distinguished in the first few years of growth by late leaf abscission in the fall. Segregation ratios suggest that the EVG trait is controlled by a single gene, evg, the EVG state being homozygous recessive. Evergreen trees were characterized by insensitivity of shoot tips to daylength and failure of terminal growth to cease growth until killed by low temperature. Lateral buds of EVG trees went dormant in the fall. Deep supercooling occurred in both EVG and DE trees, but it appeared later in EVG trees, was of shorter duration, and occurred to a lesser extent. Evergreen germplasm may be useful in developing peach cultivars for frost-free subtropic and tropical areas. It also presents a useful system for studying dormancy and cold hardiness.

The ‘Evergreen’ (EVG) is a feral, seed-propagated peach in central Mexico, growing in the area near lat. 18°53’ to 18°11’ N and long. 98°42’ W. It evidently descended from introductions by Spanish conquerors. This area of production varies from 900 to 2200 meters in elevation, is without killing frosts, has ≈85% of its rain (1250 mm) from early June to mid-October, and is without supplemental irrigation. Flowers and fruits on EVG trees occur year-round with one main crop ripening in November to early December and a second, lesser crop in June (Acosta and Barrios, 1987; Diaz, 1974). The second crop is usually not harvested due to small fruit induced by drought and low demand due to competition of other fruiting species.) Summer bloom sets fruit because night temperatures at this elevation are not above 15°C (Edwards, 1987; Edwards et al., 1990). Fruit require about 160 days to mature from full bloom. Thus, flowering in summer (July) gives a crop in winter and flowering in winter (January) gives a crop in summer (Diaz, 1974; Sanchez, 1975).

The insensitivity of the EVG peach to cessation of terminal growth under shortening photoperiods and lower temperatures (Mowrey and Sherman, 1986) and the ability to set flower buds throughout the year make it an interesting genotype. The ability of closely related deciduous (DE) and EVG germplasm may provide an interesting model system to study processes related to acclimation and dormancy where the two processes are not superimposed on each other. Our observations revealed that indicators of the onset of dormancy, such as leaf senescence and abscission, and adherence of bark tissue to the underlying xylem tissue did not develop in EVG trees except in climates where they were partially or fully defoliated by stress or leaf diseases. The objective of this study was to test the hypothesis (Lammerts, 1945) that expression of the EVG habit is under control of a single recessive gene and to develop comparative information on the ability of EVG trees to acclimate in different locations.

Materials and Methods

Trees of EVG peach were propagated onto locally used rootstocks at Chapingo, Mexico, Gainesville, Fla., Byron, Ga., and Kearneysville, W.Va. Observations were made for more than 2 years on their growth and fruiting at Tetela (Mexico), Chapingo, Gainesville, Byron, and Kearneysville. The EVG peach was hybridized with ‘Maravilha’, ‘BRI, BR6’, ‘Capo Bosco’, ‘Coral’, ‘L72-4-20’, and ‘Empress’ DE peaches at four locations (Table 1). The nine seed parent genotypes ranged from 250 to 1000 chill units (cu). Chilling requirement (CR) of parents, F1, and F2 seedlings was recorded by comparing bloom and leafing dates of these seedlings with standard ‘key’ cultivars, as shown by Sherman et al. (1988). The sources of the pollen parent EVG peach in Florida and Mexico were seedlings of the ‘Tetela EVG’. The source of EVG in Georgia and West Virginia was PI442380, which was collected in 1979 as seed from a local market in Tupachula, Chiapas, Mexico (lat. 15° N, long. 92° W). The vendor reported that the fruit were grown in Mexico State. The four seedlings and their progenies grown at Byron had white flesh fruit with little exterior red color and ripened in August. These characteristics fit the descriptions of the EVG trees grown at Gainesville and Chapingo. At Kearneysville, open-pollinated (OP) seed were collected from an F1 tree of ‘Empress’ (dwarf) OP × EVG (PI442380). It is assumed that most OP seed resulted from selfing (Fogle, 1977). Since the F1 planting consisted of several
producing spring flowers that set a crop. Flower bud formation in EVG peach occurs continuously along the shoot as it matures. Flower bud differentiation is concentrated in two main periods as this peach is forced into short ecodormancy periods by drought and low temperatures in summer and winter, respectively. Lateral, mature leaves are lost before the July flowering, mainly due to drought but also to powdery mildew \textit{Sphaerotheca pannosa} (Wallr. ex Fr.) Lev. Similarly, leaves are lost before January flowering due mainly to powdery mildew and rust \textit{Transchelia pruni-spinosa} (Pers.) Diet species, but also due to drought. Following leaf loss, a new flush of shoot growth is produced in some lateral buds. Each main bloom and crop are produced by flower buds that developed on new growth made during the preceding fruit development period. Peaches of low chilling requirement from Florida, grown in the same conditions near Tetela, do not have these two main flowering periods and their terminals and lateral buds enter into endodormancy during the winter months.

At Gainesville and Chapingo (≈400 cu), seedlings of EVG (seed imported from Tetela) grow as in the Tetela region, except that winter temperatures below –7°C kill new fully expanded leaves and terminate growth. Lateral flower and leaf buds of EVG at Gainesville enter into endodormancy. Spring bloom and leafing indicate that these buds require 450 to 500 cu to initiate growth without exhibiting symptoms of inadequate winter chilling. At Byron, the mature EVG (PI 442380) leaves drop off due to drought and rust in the fall or are killed by freezing temperatures. Terminal growth continues intermittently throughout the winter, although the tips of the EVG in Byron may be killed. In the absence of such damage, winter growth continues. Mature trees at Byron survived record lows of –16°C in 1983 and –21°C in 1988, with damage only to the terminal growth. This pattern of winter growth does not appear to make the trunk particularly cold-tender, under the conditions at Byron, despite some lack of acclimatization. Bloom occurs over an extended period from early February through

rows of closely spaced trees, any seeds derived from outcrossing were likely to be the result of intercrossing between F1 trees. F1 seeds were germinated in the greenhouse in Winter 1986 and field-planted in June 1987. Dwarf and standard trees were segregated, but planted in adjacent rows. At all locations growth type was evaluated visually. Trees were classified as EVG if they maintained their leaves and terminal buds growing at the end of the fall, started terminal bud growth earlier in spring than lateral buds, or both.

Deep supercooling capacity (as low temperature exotherms) of xylem tissues and flower buds was determined at Byron on a monthly basis from Sept. 1989 through Jan. 1990 and at Kearneysville from Oct. through Dec. 1989. Freezing of deep supercooled water was characterized in three to five buds and 0.5 g of xylem segments from EVG and DE trees using differential thermal analysis as described by Ashworth (1984). A cold-hardiness screening of EVG and DE floral buds was conducted in Dec. 1989. Approximately 45 to 60 buds were evaluated at each 5°C temperature interval from –5 to –30°C. Death of primordia expressed as browning was assessed following an overnight thaw at 3°C and 24 h at 25°C (Ashworth, 1984).

At Chapingo, cold damage was evaluated in F1 (open-pollinated) populations after a natural freeze (–8°C) on 24 Dec. Cold damage was rated on a 0 to 5 scale, where 0 = no tree damage and 5 = severe tree damage to the tree.

### Results

The EVG peach is indigenous to the area around Tetela, where it was observed to have terminal buds that continue to grow whenever temperatures are high enough. This growth habit results in long shoots and a weeping-type tree. In Tetela, the lowest CR (<250 estimated cu) peach genotypes from the Florida breeding program set terminal buds in winter and enter endodormancy, producing spring flowers that set a crop. Flower bud formation in EVG peach occurs continuously along the shoot as it matures. Flower bud differentiation is concentrated in two main periods as this peach is forced into short ecodormancy periods by drought and low temperatures in summer and winter, respectively. Lateral, mature leaves are lost before the July flowering, mainly due to drought but also to powdery mildew \textit{Sphaerotheca pannosa} (Wallr, ex Fr.) Lev. Similarly, leaves are lost before January flowering due mainly to powdery mildew and rust \textit{Transchelia pruni-spinosa} (Pers.) Diet species, but also due to drought. Following leaf loss, a new flush of shoot growth is produced in some lateral buds. Each main bloom and crop are produced by flower buds that developed on new growth made during the preceding fruit development period. Peaches of low chilling requirement from Florida, grown in the same conditions near Tetela, do not have these two main flowering periods and their terminals and lateral buds enter into endodormancy during the winter months.

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![Table 1. Progeny characteristics of nine deciduous (DE) seed parents crossed with evergreen (EVG). Homogeneity X² = 2.02; P = 0.95 based on 3:1 segregation.](image)
March, corresponding to genotypes with CRs of 450 to 700 cu. The extended bloom may be related to differences in bud maturity along the twig.

Lateral buds of EVG trees (PI442380) planted at Kearneysville appear to enter dormancy as fall progresses. Terminal growth continues until killed by low temperatures (~2C). Leaf drop does not occur until late in December. It is not clear if leaf drop is the result of cold temperatures, wind, or both.

F_1 hybrid trees of EVG × DE peaches exhibited a normal type of growth in all four locations of this study. However, leaf drop occurred later in the fall at Byron and Kearneysville than at Gainesville and Chapingo in most of the F_1 seedlings, probably more as a result of a lower CR than the effects of the EVG gene. The CR of the F_1 progeny ranged from 350 to 850 cu, about the mid-range, floral buds from the DE trees responded similarly to other

7 to –22C. In the freezing test, 11% of EVG buds survived –11C but lateral and terminal growth did not continue into the winter.

Fall for DE trees than for standard low-chill requirement cultivars, and lateral flower and leaf buds.

The CR of the EVG peach is probably the result of the effects of the EVG gene with either growth habit. In Dec. 1989, trees were again acclimated before the first killing freeze. Leaf drop was later in the fall for DE trees than for standard low-chill requirement cultivars, indicating that this trait is controlled by a single recessive gene with little environmental influence and is independent of the CR of lateral flower and leaf buds.

At Byron, F_2 seedlings segregated at a 3 DE : 1 EVG ratio (Table 1). However, recovery of yellow-fleshed EVG trees was less (59 white : 7 yellow) than the expected 3:1 (X^2 = 6.68, P < 0.01). The deciduous progeny approximated 3:1 (144 white : 52 yellow).

F_2 trees were observed for EVG in 1988 and 1989 at Kearneysville. In Dec. 1988, three classes of trees were readily observable in both the standard size and dwarf trees. Class I (DE) consisted of trees in which lateral and terminal buds were apparently dormant and all leaves had fallen from the trees at the time of normal leaf drop for DE trees in the area. Both lateral and terminal buds of Class II (semi-EVG) trees appeared dormant but most leaves were green and firmly attached to twigs at the time of evaluation in December, at least 1 month after normal leaf drop. Terminal shoots of Class III (EVG) trees continued growth, but lateral buds appeared dormant. Leaves were green and firmly attached. These trees segregated as Class I, 34 trees; Class II, 68 trees; and Class III, 34 trees, an exact 1:2:1 ratio. Chi squares for a 1:2:1 ratio (Class I, II, III) in standard and dwarf trees were .037 and .054, respectively (P = .95), indicating no association of the EVG gene with either growth habit. In Dec. 1989, trees were again evaluated, but only Classes I and III could be distinguished in a ratio of 102:34 (Table 1).

DE seedlings at Chapingo suffered less cold damage after a natural freeze than seedlings with the EVG habit (Table 2). Evergreen seedlings probably suffered the most damage because they had more prolonged growth in the fall and the tissues did not acclimate before the first killing freeze. Leaf drop was later in the fall for DE trees than for standard low-chill requirement cultivars, but lateral and terminal growth did not continue into the winter.

In late December the EVG trees exhibited LTEs in the range of –7 to –1C, whereas LTEs in the DE trees were in the range of –7 to –22C. In the freezing test, 11% of EVG buds survived –11C compared with 80% of DE at the same temperature (Fig. 1). In general, floral buds from the DE trees responded similarly to other

Table 2. Cold-damage evaluation in three F_2 seedling populations of deciduous (DE) and evergreen (EVG) habit during a natural freeze (~8C) on 24 Dec. 1990 at Chapingo, Mexico.

| CR (cu) | BRI × EVG | Capo Bosco × EVG | Coral × EVG |
|--------|-----------|------------------|-------------|
| 150    | ---       | 1.7              | ---         |
| 250    | 0.7       | 2.4              | 1.2         |
| 350    | 0.3       | 1.3              | 0.6         |
| 450    | 0.5       | 1.4              | 1.1         |

CR = chilling requirement (cu) classes in F_2 lateral buds.

Table 3. Progressive change in the extent of low-temperature exotherm of current year shoots of deciduous (DE) and evergreen (EVG) germplasm. Low-temperature exotherm midpoint temperatures were determined using differential thermal analysis.

| Date          | DE | EVG |
|---------------|----|-----|
| Byron, Ga.    |    |     |
| 29 Sept. 1989 | –18| –17 |
| 12 Oct. 1989  | –17| –15 |
| 13 Nov. 1989  | –18| –15 |
| 13 Dec. 1989  | –24| –24 |
| 4 Jan. 1990   | –30| –21 |
| Kearneysville, W.V. |    |     |
| 18 Sept. 1989 | –20| –16 |
| 25 Sept. 1989 | –21| –16 |
| 11 Oct. 1989  | –23| –16 |
| 30 Oct. 1989  | –27| –18 |
| 9 Nov. 1989   | –27| –18 |
| 3 Dec. 1989   | –31| –25 |

Discussion

“Evergreen” growth was studied by Lammerts (1945) on two F_2 populations (16 and 70 trees) of ‘Chinese Dwarf Evergreen’ × ‘Rio Oso Gem’. These populations segregated 5 DE : 6 semi-EVG : 5 peach floral buds (Quamme, 1974).

Deep supercooling in DE flower buds began to appear in mid-October and the low temperature exotherm (LTE) appeared at increasingly lower temperatures as acclimation increased. In contrast, deep supercooling in the EVG trees did not appear until late November. Sporadic flowering in the EVG trees began in Georgia during the first week of January.

The LTE of xylem tissues in current year shoots in DE and EVG trees from September through January at Kearneysville and Byron (Table 3) followed trends similar to previous reports (Wisniewski and Ashworth, 1986). The ability to deep supercool increased over the fall period in both DE and EVG, but the extent of deep supercooling was much greater in the DE. A detailed study of cold acclimation and polypeptide changes in current year shoots of DE and EVG genotypes has been reported by Arora et al. (1992).

Further evaluation of samples from Kearneysville after 3 Dec. 1989 are not included due to severe cold temperatures that resulted in dieback of the EVG trees to ground level. Evaluation of samples from Byron were discontinued after the onset of flowering in the first week of January. A significant level of deep supercooling was still present in the xylem tissues of EVG samples at the time of flowering.

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EVG as opposed to previous beliefs that it had no CR. The gene that conveys the EVG character may be of special interest in the breeding of peaches in the frost-free subtropics and tropical highlands. This peach is interesting germplasm for acclimation response studies due to its lack of response to shortening day lengths in terminal growth cessation and terminal bud dormancy.

In field tests at Byron, unbudded EVG (PI442380) seedlings are surviving well on a peach tree short-life site. We don’t know if this superior survival relates to root reaction to ring nematode or to a unique trunk physiology that makes the tree less prone to spring cold damage.

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Fig. 1. Percent survival of floral buds from deciduous (●) and evergreen germplasm (○) collected on 29 Dec. 1989 in Byron, Ga.