Inheritance of Rind Color and Reverse Striping in a *Cucurbita pepo* (subsp. *texana*) **Cross**

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Introduction

Several loci determining exterior fruit color have been characterized in C. pepo (5), among which are the L-1 and *L-2* loci and the *D* gene. Interactions of multiple alleles at the L-1 and L-2 loci confer varying degrees of dark or light pigmentation, with the dominant alleles *L*-1 and *L*-2 conferring dark green coloration and their recessive counterparts *l*-1 and *l*-2 conferring a light green rind color (6). In addition, the $l-1^{BSt}$ allele causes fruit to exhibit broad, contiguous dark stripes in combination with the L-2 allele, and is recessive to L-1 but dominant to l-1 (3). A recently identified allele, $l-2^R$, reverses the stripes (a phenotype known as "reverse striping") in the presence of any striping allele—such as *l*-1^{BSt}—at the other locus, such that the broad stripes in between the fruit's vein tracts are lighter than the background color, which remains darker over the vein tracts (4). The *D* gene affects both fruit and stem color, with the dominant *D* allele conferring dark coloration to stem and fruit beginning at about 15-18 days after anthesis (6). Epistatic to *l*-1 and *l*-2, the *D* allele can cause fruits which would otherwise be light colored (i.e. *l*-1/l-1, l-2/l-2) to be dark.

Current evidence points to two possible systems of reverse striping inheritance in *C. pepo*, (1,4) In Paris' model (4), complete or partial RS phenotype in delicata x spaghetti squash results from the presence of at least one copy of the *l*-1^{BSt} allele and the *l*-2^R allele, independent of any other loci such as the *D* gene. Loy's more recently proposed system (1), based on crosses involving egg gourd, is the following: RS phenotype results from the presence of the dominant *D* allele, at least one copy of the *l*-1^{BSt} allele. The heterozygous genotype *L*-2 /*l*-2^R is responsible for "partial" RS in the first system and heterozygous *L*-2/*l*-2 for a similar "Type 2 RS" in the second.

Crosses between delicata and acorn squash are often used for the aesthetic of rind patterns on winter squash. The goal of this study was to initiate an evaluation of the two proposed genetic systems in these types of cultivars.

Materials and Methods

A cross between a delicata squash (*Cucurbita pepo* subsp. *texana* cv. Bush Delicata) expressing reverse striping and a dark green acorn squash (*Cucurbita pepo* subsp. *texana* cv. Sweet Reba) was performed. The F_1 was self-pollinated, and the F_2 population grown in the field in the summer of 2012 at the Homer C. Thompson Vegetable Research Farm in Freeville, NY, alongside the parents and F_1 . Fruits were harvested at maturity , and photographs of fruit from parents, F_1 and 135 F_2 plants were used for phenotyping.

Results and Discussion

The cross segregated for exterior fruit color (Figure 1). Fruit from 122 of the F_2 plants fell into easily identifiable RS ('Bush Delicata') or dark ('Sweet Reba') phenotypic categories, while 13 plants did not fit these categories and had fruit that showed light-green splotchiness or orange mottling. Of these 122 plants, 68 were RS and 54 were dark with respect to rind color. Fruit were not further separated into completely or partially reversed phenotypic classes due to difficulty in discerning differences.

The predictions of each of the afore-mentioned systems of RS inheritance with respect to the cross were compared against the obtained results. Acorn squash is known to have genotype DD, l-1/l-1, L-2/L-2 (2). Following the model of Paris, delicata has genotype DD, l-1^{BSt}/l-1^{BSt}, l- $2^{R}/l-2^{R}$ (4). Using the system of inheritance proposed by Loy, delicata is assigned the genotype DD, *l*-1^{BSt}/*l*-1^{BSt}, *l*-2/*l*-2. In either system the F_1 would be heterozygous at the L-2 locus and confer a partially reverse striped phenotype, which we observed (Figure 1). The expected F₂ phenotypic ratio for this cross predicted by both models is a 9:7 ratio of RS to dark exterior fruit color (Table 1). The *L*-1^{BSt}/__, *L*-2/L-2 genotype would ordinarily confer a broad, normal striping phenotype, but because of epistatic interaction in the presence of *D*, the stripes would not be apparent and the fruit are simply dark. Of the 122 classifiable plants, 68 were RS and 54 were dark—results consistent with either model ($\chi^2 = 0.013$, p= 0.9092).

The difference between the two systems involves the *D* gene and the existence of the $l-2^R$ allele. Loy, in his crosses, found that the dominant *D* allele was necessary to confer the RS phenotype (1), whereas Paris found RS exhibited even in crosses involving only light-stemmed (*dd*) plants (4). Differences in these systems may be a result of the different parents used for crossing (H. Paris, personal communication).

Because of the nature of our cross, especially its lack of segregation for the *D* gene, clarification regarding the inheritance of RS and the role of the *D* gene in reverse striping in *C. pepo* cannot be further elucidated using our current data. Additional crosses and plantings are planned.

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Literature Cited

Table 1. Expected genotypic frequencies and associated phenotypes of the F2 population given two systemsof reverse striping

Genotype:	Genotype:		
D-Independent Model	D-Dependent Model	Frequency	Phenotype
DD, L-1 ^{BSt} /, L-2/L-2	DD, L-1 ^{BSt} /, L-2/L-2	3/16	dark
DD, L-1 ^{BSt} /, L-2/l-2 ^R	DD, L-1 ^{BSt} /, L-2/l-2	6/16	RS (partial/Type 2)
DD, $L-1^{BSt}/$ _, $l-2^{R}/l-2^{R}$	DD, L-1 ^{BSt} /, l-2/l-2	3/16	RS (complete/Type 1)
DD, l-1/l-1, L-2/L-2	DD, l-1/l-1, L-2/L-2	1/16	dark
DD, l-1/1-1, L-2/l-2 ^R	DD, l-1/1-1, L-2/l-2	2/16	dark
$DD, l-1/l-1, l-2^R/l-2^R$	DD, l-1/l-1, l-2/l-2	1/16	dark



Figure 1. Delicata with complete reverse striping (left), acorn with dark green rind color (middle), and F₁ with intermediate/partial/"Type 2" reverse striping (right)