

Gene symbol for the erectoides-type, short-internode mutant Wt 11242

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The gene responsible for the short-internode phenotype of mutant line Wt11242 has now been assigned two different symbols. Recently, Świącicki and Wolko (8) designated the mutant allele *dnd* (*densinodosum*) and showed *Dnd* was located in the *M* region of linkage group III. Earlier, Kusnadi et al. (2) found the Wt11242 mutant had an erectoides-type phenotype and was allelic with the semi-erectoides mutant *lka* characterized by Reid and Ross (5). They designated the second mutant allele *lka*¹¹²⁴². However, in the terminology now adopted by the Hobart pea group, the two mutant alleles would be written as *lka1-1* (type line WL5865) and *lka1-2* (type line Wt11242).

There are five loci in the erectoides series: *Lk*, *Lka*, *Lkb*, *Lkc* and *Lkd* (1, 4-6). The erectoides phenotype is characterized by short internodes, stiff stems, short peduncles and petioles, dark green leaves, some epinasty, and an absence of basal branching. In the *lka* mutants, the shorter internodes result from a reduction in both cell length and cell number (2, 5). The erectoides phenotype appears to result from impaired brassinosteroid synthesis or reception. Mutants *lk* and *lkb* are brassinosteroid deficient: the *lkb* mutation blocks conversion of 24-methylenecholesterol to campesterol (3) and *lk* blocks conversion of campesterol to campestanol (9). The *lka1-1* mutant is brassinosteroid insensitive (3). Response of mutant *lka1-2* has not been reported. The *lka1-2* mutation appears less severe than *lka1-1* (2). The plant brassinosteroid growth regulators are related to animal steroidal hormones. The pea *Lkb* gene has been sequenced and the amino acid sequence showed 60% similarity with the human homologue (7).

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