Genetic mechanisms associated with determinate growth habit in cowpea (*Vigna unguiculata* (L.) Walp.)

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Motivation and Aim: Cowpea (*Vigna unguiculata* (L.) Walp.) is an important grain legume crop. It is known two types of growth habit that are classified as determinate and indeterminate. In wild species the growth habit is indeterminate and terminal shoot meristem of these plants remains in a vegetative state throughout the production of vegetative and reproductive structures. In plants with determinant growth habit, terminal shoot meristem switches from vegetative to reproductive state. It is known that *TFL1* is gene controlling determinate growth habit in many legume species. The aim of the present study was (1) to identify and characterize *TFL1* homologs in *Vigna* genome sequence and (2) to reveal allelic diversity of *TFL1* gene in *V. unguiculata* accessions with type of growth habit, sensitive and insensitive for different (dry and wet) environment conditions.

Methods and Algorithms: Based on the allelic differences of the *TFL1* gene described in the literature, DNA markers for distinguishing genotypes with different growth habit were developed. The homologous sequences of *TFL1* gene have been identified in *Vigna* Genome Server ('VigGS', http://viggs.dna.affrc.go.jp) using BLAST search. *In silico* analysis was performed with MEGA, FGENESH+ and PLACE software.

Results: Additional copies of the *TFL1* gene were found in *Vigna* genome. Phylogenetic analysis allowed to establish that the duplications occur in the common ancestor of legumes. 12 combinations of DNA markers were developed based on the known mutations of the *TFL1* gene and used for re-sequencing this gene in *V. unguiculata* accessions, sensitive and insensitive to changing type of growth habit in different environment conditions.

Conclusion: Comparison of *V. unguiculata* accessions with type of growth habit, sensitive and insensitive for different (dry and wet) environment conditions, suggest more complicate regulation of this trait than mechanism based on just *TFL1* allelic differences. Further comparative transcriptomic studies are needed.