# The Construction of Genetic Map of Faba Bean (Vicia faba L.) Using RAPD Markers 

## Memen Surahman ${ }^{1)}$


#### Abstract

Fifty seven individuals of recombinant inbred lines (RILs) of faba bean (Vicia faba L.) descended from the cross between 34Morocco $x$ Kristall25 have been analyzed for RAPD markers. A total of 77 markers were assayed segregating in the 57 RI lines and grouped into 14 linkage groups, with total length of 973.2 cM .


Key words : Vicia faba, Recombinant inbred line, RAPDs, Linkage groups

## INTRODUCTION

Preliminary analysis of genetic linkage in faba bean includes the study of morphological traits (Cabrera and Martin, 1989) and was followed by the establishment of linkage maps based on morphological, isozyme, RFLP and RAPD markers (Van de Ven et al., 1991). Torres et al. (1993) reported, that when two F2 populations ( 20 F 2 individuals derived from the cross Vf6 x Vfl73 and 44 F2 plants from the cross Vf6 $x$ Vf 35 ) were analyzed using isozyme, RFLP and RAPD, 11 independently assorting linkage groups were identified in this population. Satovic et al. (1996) analysed 13 F2 families of faba bean descending from a trisomic plant. They have been analyzed for morphological, isozyme and RAPD markers. The study revealed a total of 10 linkage groups, six of which have been precisely assigned to specific chromosomes. The most recent map in faba bean has been reported by Vaz Patto et al. (1998). The map consist of 14 linkage groups with total length of 984.5 cM . The main objective of the present study was to construct genetic map.

## MATERIALS AND METHODS

For the construction of a linkage map a population of recombinant inbred lines (RILs) derived from the cross 34 Morocco $x$ Kristall25 was used. The 34Morocco is a bean pure line BPL228/ILB141 from ICARDA, Syria. Kristall25, developed at Hohenheim, Germany, is an inbred line from the German cultivar Kristall. These parents were chosen because of their large genetic and phenotypic difference (Schill et al., 1997).

The F2 plants from the cross 34 Morocco $x$ Kristall25 and from the cross Kristall25 x 34Morocco
were self-pollinated and advanced to the F7 generation using single-seed descent. The final population consists of 253 lines including reciprocal lines. A random sample of 57 lines was characterized based on RAPDs.

Segregation data were analyzed with MAPMAKER/EXP 3.0 computer program (Lander et al., 1987; Lincoln et al., 1993), with a LOD score 3.0 and with 0.4 theta ( 40 Haldane cM ; Haldane, 1919) as critical distance between two linked markers.

## ${ }^{*}$

## RESULTS

The 48 selected primers amplified a total of 115 polymorphic bands (data not shown) with an average of 2.3 polymorphic bands per primer. Fragment sizes ranged from about 550 bp to 2400 bp . With linkage criteria LOD $>3.0$ and a distance of less than $40 \mathrm{cM}, 77$ RAPD marker loci were classified into 14 groups (Figure 1). There were 38 unlinked markers. These markers were omitted.

The size of the 14 linkage groups ranged from 6.3 cM to 278.7 cM . The total length of map is 973.2 cM . The largest linkage group contains 16 loci (linkage group 3). The distance between markers ranges from 1.9 cM to 38.5 cM , with an average of 14.66 cM .

## DISCUSSION

A total of 77 markers were found segregating in 57 RI F7-sample lines (Figure 1). The total map distance defined among linked markers was 973.2 centiMorgan. A comparison of the present genetic map with the hitherto published maps shows that its total length resembles the map of Vaz Patto et al. (1998) with a map length of 984.5 cM . However these maps are still small when compared to the expected size of the Vicia faba

[^0]

Figure 1. Primary linkage map of faba bean (Vicia faba) with 77 RAPD-markers. Number to the left of a linkage group represents map distance in cM calculated with the Haldane mapping function. All linkages shown have a LOD score of at least 3.0. Designations to the right are locus names.
genome ( 1500 cM ; Vaz Patto et al., 1998). Besides the total map length a complete map is obtained only when the number of linkage groups corresponds to the haploid number of chromosomes. The haploid chromosome number of the faba bean is six $(x=6)$, therefore, six linkage groups, instead of the 14 linkage groups detected here, were expected. These results reflect problems associated with the use of relatively small populations (here: $\mathrm{N}=57$ ).

In view of the relatively large size of the linkage group 3, it may putatively be assigned to chromosome number one of the faba bean. In the standard karyotype, this chromosome is the longest one ( Torres et al. 1993). It is a metacentric chromosome and about twice as long as each of the five subtelocentric chromosomes (Cabrera et al., 1989).

The average distance between markers in the present map is 14.66 cM when all 77 RAPD markers were used. Although the density of the map was lower than in the faba bean map constructed by Vaz Patto et al. (1989) which has an average distance of 12.95 cM , the map presented here can be considered as sufficiently dense for a QTL analysis. In general, a marker distance about of 20 cM is still adequate to detect QTL (Tanskley et al., 1993).

## REFERENCES

Cabrera, A., and A. Martin. 1989. Analysis of genetic linkage in faba bean (Vicia faba L.). FABIS Newslett 24:3-5.

Cabrera, A., J.I. Cubero, and A. Martin. 1989. Genetic mapping using trisomics in Vicia faba L. FABIS Newslett 23:5-7.

Haldane, J.B.S. 1919. The combination of linkage value and the calculation of distance between the loci of linked factors. J. Genet. 8:299-309.

Lander, E.S., P. Green, J. Abrahamson, A. Barlow, M.J. Daly, S.E. Lincoln, and L. Newburg. 1987. Mapmaker: An interactive computer package for constructing primary genetic linkage maps of
experimental and natural populations. Genomics I:174-181.

Lincoln, S.E., M.J. Daly, and E.S. Lander. 1993. Constructing genetic linkage maps with MAPMAKER/EXP version 3.0: A Tutorial and reference manual. Whitehead Institute for Biomedical Research. USA.

Satovic, Z., A.M. Torres, and J.I. Cubero. 1996. Genetic mapping of new morphological, isozyme and RAPD markers in Vicia faba L. using trisomics. Theor. Appl. Genet. 93:1130-1138.

Schill, B., A.E. Melchinger, E. von Kittlitz, and W. Link. 1997. Zächterische Brauchbarkeit von Intrapool-und Interpool Kreuzungen des mitteleuropäischen und mediterranen Genpools bei der Fababohne (Vicia faba L.). Vortr. Pflanzenzüchtung 38:127-145.

Tanksley, S.D., H. Medina-Filhoet, and C.M. Rick , Ganal., M.W., Prince, J.P., de Vicente, M.C., Bonierbale, M. W., Broun, P., Fulton, T.M., Giovannoni, J.J., Grandillo, S., Martin, G.B., Messeguer, R., Miller, J.C., Miller, L., Paterson, A.H., Pineda, O., Röder, M.S., Wing, R.A., Wu, W. and Young, N.D. 1993. High density molecular linkage maps of the tomato and potato genomes. Genetics 132:1141-1160.

Torres, A.M., Weeden, N.F., Martin, A. 1993. Linkage among isozyme, RFLP and RAPD markers in Vicia faba. Theor. Appl. Genet. 85:937-945.

Van de Ven, W.T.G., R. Waugh, N. Duncan, G. Ramsay, N. Dow, and W. Powel. 1991. Development of genetic linkage map in Vicia faba using molecular and biochemical techniques. Aspects Appl. Biol. 27:49-54.

Vaz Patto, M.C., A.M. Torres, and J.I. Cubero. 1998. Genetic mapping in Vicia faba L. using trisomics. $3^{\text {rd }}$ European Conference on Grain Legumes. Valladolid. p.64-65.


[^0]:    1) Department of Agronomy, Faculty of Agriculture,Bogor Agricultural University

    Kampus IPB Darmaga Bogor 16610, Bogor.Email:msurahm@lycos.com.Phone: (0251) 635141

