# **Genetic Analysis of Fiber Traits in Cotton**

## Yüksel BÖLEK<sup>1</sup>, Hatice ÇOKKIZGIN<sup>2\*</sup>, Adem BARDAK<sup>1</sup>

<sup>1</sup>KSU, Faculty of Agriculture, Department of Agricultural Biotechnology, Kahramanmara <sup>2</sup>Gaziantep University, Nurda 1 Vocational School, Gaziantep

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**Abstract:** Diallel analysis has been used in many breeding programs to increase yield and quality. Eight cotton genotypes (Giza-45, A kabat-100, Is-4, 108-F, Acala Prema, Çukurova-1518, Nazilli-84S, Stoneville-45) were crossed in a complete diallel design in Kahramanmara in 2007. Genetic characteristics for fiber quality traits; length (Len), strength (Str), micronaire (Mic), uniformity (Unf), elongation (Elg), spinning consistency index (Sci) and short fiber index (Sfi) were determined through Hayman-Jinks diallel analysis methods. The analysis of variance revealed significant differences among the parents. Inheritances of most of the characters were due to dominant genes that were determined for Len, Mic, Sci, Str in Is-4; for Sfi and Unf in Giza-45 and for Elg in Nazilli 84S. **Key Words:** Fiber quality, gene action, cotton, diallel

Pamukta Lif Özelliklerinin Genetik Analizi

Özet: Diallel analiz yöntemi verim ve kaliteyi artırmak için birçok ıslah programında kullanılmaktadır. Sekiz pamuk genotipi (Giza-45, A kabat-100, Is-4, 108-F, Acala Prema, Çukurova-1518, Nazilli-84S, Stoneville-45) 2007 yılında Kahramanmara 'ta tam diallel dizayna uygun olarak melezlenmi tir. Lif kalite karakterleri; uzunluk (Len), dayanıklılık (Str), incelik (Mic), üniformite (Unf), uzama (Elg), iplik olabilirlik indeksi (Sci) ve kısa lif indeksi (Sfi) için genetik özellikler Hayman-Jinks diallel analiz yöntemine göre belirlenmi tir. Varyans analizi ebeveynler arasında önemli farklılıklar ortaya çıkartmı tır. Is-4'te Len, Mic, Sci, Str için; Giza-45'te Sfi ve Unf için; Nazilli 84S'de Elg için dominant genler belirlenmi tir.

Anahtar Kelimeler: Lif kalitesi, gen hareketi, pamuk, diallel

#### INTRODUCTION

The knowledge of genetic structure and mode of inheritance of complex traits help breeders and producers to establish suitable breeding methodology to improve economically important traits (Garg and Kalsy, 1988). Selection criteria have been used in conventional breeding programs are usually based on phenotypic characteristics. Recently, molecular markers started to employ in breeding programs. On the other hand, environmental conditions affect the phenotypic characteristics that have polygenic nature (Antoni et al., 1991; Rivera et al., 1999). Also, it is difficult to characterize with morphological markers and they don't show high levels of variation (Lukonge, 2005). Repeated crossing and intensive selection between a few genotypes with desirable traits, led to the narrowing of the cotton gene pool that resulted in low genetic variation of cultivated cotton genotypes. It is needed to include new and diverse progenitors in the breeding programs from other species such as G. barbadense and wild type accessions to increase variability.

It is necessary to characterize the genetic structure of every breeding material for effective breeding scheme. Diallel analysis method was developed to have an idea about the genetic mechanisms and structure of populations for different traits (Hayman, 1954). The breeding programs for many crops use diallel analysis because it offers genetic information on quantitative traits to the breeders (Viana et al., 2001; Peyman, 2012; Akbari et al., 2013). Diallel analysis also gives information about good combining inbred lines suitable for hybrid production (Ado et al., 2006; Townsend et al., 2013). It is a strong technique to explain differences of genotypes, phenotypic variance, additive gene effects, variation ratio of specific gene combinations (Danehloueipour et al., 2007; Saleem et al., 2013). It also provides useful data about genetic control of quantitative traits and their gene effects model (Baker, 1978; Yao et al., 2011). This way, dominant parents and hybrids can be defined (Goval and Kumar, 1991; Ahuja and Dhayal, 2007). On the other hand, diallel analysis has been used for genetic control study of quantitative traits of autogam or allogam populations (Jinks and Hayman, 1953), study of general and specific combining ability (Griffing, 1956) and analysis of heterosis (Gardner and Eberhart, 1966). Diallel crossing technique in cotton has also been used by cotton breeders (Zangi et al., 2010) and it explains practical and important information about breeder's materials (Ali et al., 2011). With this, it also reports equality and distribution of dominant and recessive alleles (Mohamed et al., 2009) as well as better genotypes for whichever yield trait (Ragsdale, 2003).

Cheatham et al. (2003) reported that Australian and wild cotton genotypes have the genes to improve fiber quality and fertility. While fiber fineness and length

<sup>\*</sup>Sorumlu yazar: Çokkızgın, H., hcokkizgin@gantep.edu.tr

primarily exhibit dominance gene effects, fiber percentage and fiber strength are controlled by additive gene effects. On the other hand, fiber yield and fiber elongation extension are controlled equally by additive and dominant gene effects. Thus there is a need an effective characterization of germplasm for successful cotton breeding. The purpose of this work is to identify gene action of some fiber quality traits in cotton cultivars mostly used in crosses to develop high yielding quality cotton.

### **MATERIAL and METHODS**

**Plant material:** Eight cotton genotypes were used in the experiment. They were belonging to *G. hirsutum* L. (Is-4, 108-F, Acala Prema, Çukurova-1518, Nazilli-84S, Stoneville-453) and *G. barbadense* L. (Giza-45, A kabat-100) species. Genotypes were selected for important fiber characteristics in addition to verticillium wilt resistance (Table 1).

Table 1. Characteristics of the cotton genotypes used in the experiment

| Genotypes      | Characters  |
|----------------|---|
| Giza-45        | Length (36.3 mm), strength (42.7 g/tex), uniformity (87.5 %), elongation (6.7 %), spinning consistency index (215.5), short fiber index (3.4) and verticillium wilt resistance.                                   |
| A kabat-100    | Length (35.6 mm), strength (43 g/tex), micronaire (3.4), uniformity (85.6 %), short fiber index (3.4) and verticillium wilt resistance.   |
| Is-4           | Length (36.4 mm), strength (45.1 g/tex), micronaire (3.8), uniformity (88.1 %), spinning consistency index (223), short fiber index (3.4) and verticillium wilt resistance.                                       |
| 108-F          | Length (32 mm), strength (40.4 g/tex), uniformity (87.7 %), spinning consistency index (197.5), short fiber index (3.4) and verticillium wilt resistance.   |
| Acala Prema    | Strength (40), uniformity (86.7 %), spinning consistency index (188), short fiber index (3.4). Acala lines are most widely used in hybridization programs in the world. Vertcillium resistance and high yielding. |
| Çukurova-1518  | Commercial cultivar developed at Cukurova Agricultural Research Institute. Adapted to Mediterranean Region of Turkey. High ginning turnout (41%).   |
| Nazilli-84S    | Commercial cultivar developed at Nazilli Agricultural Research Institute. Adapted to Aegean Region of Turkey. High ginning turnout (45%).   |
| Stoneville-453 | Commercial cultivar developed at Stoneville Pedigree in USA. Adapted to Southeast of Turkey. High ginning turnout (42%).  |

*Field evaluations*: The  $F_1$  material was developed by crossing eight cultivars in all possible combinations in the experimental fields of Kahramanmara Agricultural Research Institute in 2007. In the experiment, plot length was 12 m, spacing between and within rows was 70x20 cm. Completely randomized block design with 3 replications was applied. Standard cultural practices were applied as fertilizer (14 kg N and 12 kg P<sub>2</sub>O<sub>5</sub>) and irrigations (total 7-10 times in about 8-10 days intervals as needed during growing seasons). Field was received total 680.6 mm amount of rain during 2007 growing season. Bolls were sampled from the first position of the middle fruiting branches of each genotype. Fiber characteristics; length (Len) (mm) (2.5% Span Length), strength (Str) (g/tex) micronaire (Mic), uniformity (Unf) (%), elongation (Elg) (%), spinning consistency index (Sci) and short fiber index (Sfi) were determined by HVI (High Volume Instruments).

*Statistical analysis*: The data were analyzed using diallel type methods developed by Hayman-Jinks (1953). Parameters calculated are follows: E: Expected environmental component of variation, D: Additive genetic variance, F: Mean of Fr values over arrays, H1: Variation due to dominant effect of gene, H2: Variation

due to dominant effect of gene correlated with gene distribution, KD/KR: Ratio of dominance and recessive allels, h<sup>2</sup>: Net dominance over all loci in heterozygous phase, K: Effective genes, Wr: Co-variance, Vr: Variance, Hg: Broad sense heritability, Hd: Narrow sense heritability.

### **RESULTS and DISCUSSION**

The results for the analysis of variance revealed that the mean genetic differences among the hybrids and their parents in  $F_1$  generation were highly significant for all the fiber characters studied (Table 2).

Diallel analysis showed for all traits (H1/D)  $^{1/2}$  value greater than 1 (Table 3). Therefore, there is overdominance for the traits because in a diallel analysis, if (H1/D)  $^{1/2}$  value greater than 1, there is over-dominance for the trait (Balc1 and Turgut, 2006). Thus, in hybrid studies there may be a high chance of success in selecting superior genotypes. On the other hand, variation in environment can effect plant development and growth as well as quality of the product. If environmental variance is high then breeding can result unsuccessfully ( ener et al., 1999). Because of environmental variance is too high (30.539±148.093) for Sci in this experiment, breeding studies may be unsuccessful for this trait. Additionally, KD/KR ratio less than 1, meaning that recessive genes are excess for Sci (Table 3).

| Characters                 | Mean Squares | F       |  |
|----------------------------|--------------|---------|--|
| Micronaire                 | 0.318        | 38.585* |  |
| Length (mm)                | 22.028       | 23.451* |  |
| Strength (g/tex)           | 54.216       | 17.873* |  |
| Spinning consistency index | 1669.603     | 28.247* |  |
| Uniformity (%)             | 4.981        | 6.333*  |  |
| Short fiber index          | 0.454        | 7.546*  |  |
| Elongation (%)             | 0.832        | 7.754*  |  |

\*, Significant at 0.05

Table 3. Some genetic parameters for the fiber characters studied

| Source of             | NC .               |                    |                     | a · · · · · · · · · · ·    | <b>I</b> I :C : (0() | C1 ( C1 1 1       |                   |
|-----------------------|--------------------|--------------------|---------------------|----------------------------|----------------------|-------------------|-------------------|
| Variation             | Micronaire         | Length (mm)        | Strength (g/tex)    | Spinning consistency index | Uniformity (%)       | Short fiber index | Elongation (%)    |
| Е                     | $0.004 \pm 0.025$  | $0.472 \pm 0.845$  | $1.493 \pm 5.280$   | 30.539±148.093             | $0.445 \pm 0.738$    | $0.040 \pm 0.043$ | $0.058 \pm 0.079$ |
| D                     | 0.239±0.076        | $12.208 \pm 2.536$ | $16.402 \pm 15.840$ | 376.559±444.279            | 1.810±2.213          | 0.168±0.129       | 0.710±0.236       |
| F                     | 0.376±0.180        | -3.624±5.993       | 8.091±37.428        | -328.076±1,049.791         | 2.337±5.228          | 0.119±0.304       | 1.070±0.557       |
| H1                    | 0.771±0.175        | 18.899±5.831       | 103.062±36.413      | 2,213.187±1,021.332        | 10.128±5.086         | 0.744±0.296       | 1.945±0.542       |
| H2                    | 0.606±0.152        | 30.141±5.073       | 105.893±31.679      | 2,651.975±888.558          | 8.424±4.425          | 0.672±0.257       | 1.408±0.472       |
| D-H1                  | -0.533±0.150       | -6.691±5.003       | -86.659±31.243      | -1,836.628±876.326         | -8.318±4.364         | -0.577±0.254      | -1.235±0.465      |
| (H1/D) <sup>1/2</sup> | 1.798              | 1.244              | 2.507               | 2.424                      | 2.365                | 2.107             | 1.656             |
| H2/4H1                | 0.196              | 0.399              | 0.257               | 0.300                      | 0.208                | 0.226             | 0.181             |
| KD/KR                 | 2.561              | 0.787              | 1.218               | 0.695                      | 1.751                | 1.403             | 2.673             |
| $h^2$                 | $-0.002 \pm 0.102$ | $5.998 \pm 3.402$  | 33.899±21.246       | 922.617±595.905            | $1.219 \pm 2.968$    | 0.234±0.173       | 0.607±0.316       |
| К                     | -0.003             | 0.199              | 0.320               | 0.348                      | 0.145                | 0.348             | 0.431             |
| Yr, Wr+Vr for r       | -0.072             | -0.878             | 0.678               | -0.264                     | -0.814               | 0.787             | 0.214             |
| Hg                    | 0.773              | 0.829              | 0.647               | 0.813                      | 0.570                | 0.600             | 0.601             |
| Hd                    | 0.367              | 0.333              | 0.140               | 0.124                      | 0.159                | 0.176             | 0.390             |

E: Expected environmental component of variation, D: Additive genetic variance, F: Mean of Fr values over arrays, H1: Variation due to dominant effect of gene, H2: Variation due to dominant effect of gene correlated with gene distribution, KD/KR: Ratio of dominance and recessive alleles, h<sup>2</sup>: Net dominance over all loci in heterozygous phase, K: Effective genes, Wr: Co-variance, Vr: Variance, Hg: Broad sense heritability, Hd: Narrow sense heritability

In Wr-Vr graphs (Figure 1), co-variance and variance points show parents. If this points away from the regression line, there is epistatic gene effect otherwise if these points near origin and regression line, there are more dominant genes (Yıldırım, 2005). The inheritance of Sci is shown in a Wr-Vr graph in Figure 1. With regard to Sci, the parents; A kabat-100 (2) and Is-4 (3) had more dominant genes whereas Giza-45 (1), 108-F (4), Acala Prema (5), Çukurova-1518 (6), Nazilli-84S (7) and Stoneville-453 (8) carried more recessive genes. Stoneville-453 (8) for Elg, Len, Mic, Sfi and Unf; Çukurova-1518 (6) for Elg, Len, Mic; Giza-45 (1) for Sci and 108-F (4) for Str have more recessive genes than other parents. Moreover, Is-4 (3) for Len, Mic, Sci and Str; Nazilli-84-S (7) for Elg; A kabat-100 (2) for Sci and Sfi; Giza-45 (1), 108-F (4) and Cukurova-1518 (6) for Unf have more dominant genes than the other parents. In a

study, Ali et al. (2008) determined that parents have more dominant genes than recessive genes for the traits; Len, Str, Unf and Elg. When genotypes close to the regression line it could be postulated that there is no epistatic gene effect for the traits but genotypes away from the parabola indicates more recessive gene action. Analysis showed that Acala Prema was away from the regression line for all the traits. This result indicates Acala Prema have epistatic gene effect for all the traits. There is no epistatic gene effect for the genotypes near the regression line. If regression line cut the Wr axis above the origin, there is partial dominance and if regression line cut the Wr axis below the origin, there is over-dominance (Hayman, 1954). Griffing analysis describes partial-dominance for Str, but over-dominance was also present for all the traits except Sci and Str (Table 3).

Ara tırma Makalesi Research Article

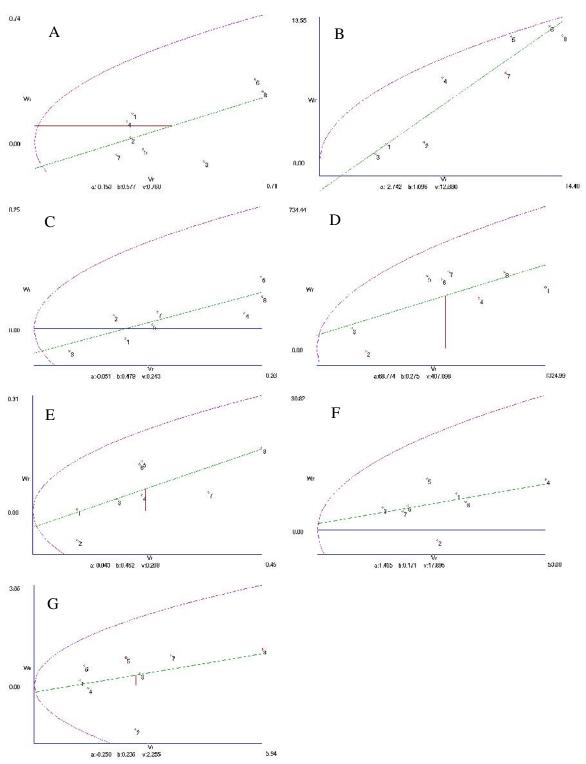


Figure 1. Vr-Wr graphs for fiber elongation (A), length (B), micronaire (C), spinning consistency index (D), short fiber index (E), strength (F), and uniformity (G). Parents were shown in numbers (1: Giza-45, 2: A kabat-100, 3: Is-4, 4: 108-F, 5: Acala Prema, 6: Çukurova-1518, 7: Nazilli-84S, 8: Stoneville-453)

D-H1 had negative values for all the traits (Table 3). This shows that dominant gene effect have more important role than additive effect in the formation of the traits. This result is parallel with additive and non-additive genes are important (Pole et al., 2008) for Len,

Str, Mic traits (Rauf et al., 2006). Aguiar et al. (2007) indicated most of micronaire index, fiber strength, fiber length, uniformity index, short-fiber index and fiber elongation traits were affected by additive genes. H2/4H1 unequal 0.25 for all traits except Str (Table 3)

indicating that dominant and recessive genes unequal ( ener et al., 2000). Str was near to 0.25 meaning it has equal allel frequency. Therefore, breeding will have a high success for Str (Ragsdale, 2003). Additionally, H1>H2 indicates that allelic frequency unequal for Mic, Unf, Sfi, Elg (Table 3). This result was also supported by H2/4H1 ratio that was less than 0.25. KD/KR ratio was higher than 1 for all traits except Len and Sci (Table 3). Therefore, there are dominant genes more than recessive genes in population for the traits. Effective gene was not determined because of K (effective gene) less than 1 for all traits.

Karademir and Gençer (2010) reported that ginning percentage, fiber length, fiber fineness and fiber elongation properties were influenced by additive while seed cotton yield, fiber strength and fiber uniformity ratio were influenced by non-additive gene effects in the experimental populations. Positive r for Yr, Wr+Vr indicates that parents with higher values for Str, Sfi and Elg have recessive alleles but dominant alleles have negative effect for the same characters. On the other hand, negative r for Yr, Wr+Vr indicates that parents with higher values for Mic, Len, Sci, Unf have dominant genes and the alleles increasing these values are also dominant. H2/4H1 ratios less than 1 and KD/KR ratios bigger than 1 except Len and Sci shows that parents have dominant gene frequency more than recessive gene frequency for all the traits (Table 3). Therefore, we can conclude that dominant alleles play an important role in the formation of these traits.

Broad sense heritability value was higher than narrow sense heritability value for all traits meaning that non-additive gene effect was effective over all traits (Mohamed et al., 2009).

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