HETEROSIS, INBREEDING DEPRESSION AND GENE ACTION OF SOME QUANTITATIVE CHARACTERS IN A DIALLEL CROSS AMONG FIVE WHITE LUPIN GENOTYPES

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ABSTRACT

The present investigation was carried out at Giza Research Station during 1999/2000, 2000/2001 and at Shandaweel Research Station in 2001/2002 growing seasons. All F_1 and F_2 Hybrids (including reciprocals) in a diallel mating design between five white lupin genotypes differing in origin (Giza-1, Dijon-2, Line 20, Line 37/3 and Kiev Mutant) were studied. Significant differences among genotypes and some crosses were detected for all studied characters, indicating genetic variability for all variables. Significant differences between some hybrids and their reciprocals were observed for all studied charcters, suggesting that earliness, seed yield and its components appeared to be influenced by the female parent. Negative heterosis percentage relative to mid parent was significant in seventeen crosses for days to maturity. Heterosis percentage was significantly positive in twelve, seven, eleven, thirteen, five and twelve crosses for plant height, number of branches, pods, seeds, 100-seed weight and seed yield per plant, respectively. Moreover, heterosis over the better parent was significant in some crosses for all studied characters except number of branches per plant. Eight, eleven, three, six, seven, five and eight crosses expressed significant inbreeding depression in F_2 for days to maturity, plant height, number of branches, pods, seeds, 100-seed weight and seed yield per plant. respectively. Additive component of genetic variability "D" was consistently significant or highly significant over generations for all studied characters. Dominance effects played an important role in the inheritance of all traits. The additive gene action was more important than the non-additive one and selection would be effective in early generations for improving these traits. Narrow sense heritability ranged from 29.0 % for 100-seed weight to 89.7 % for days to maturity. In general, position of array members on the Wr, Vr graph varied widely among generations. For P_1 (Giza-1) and P_4 (Line 37/3) had most recessive alleles for days to maturity. Moreover, P_5 (Kiev Mutant) contained most recessive genes for plant height, number of branches, pods, seeds, 100-seed weight and seed yield per plant.

Key words: White lupin, Diallel cross, Heterosis, Inbreeding depression, Heritability.

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INTRODUCTION

White lupin, Lupinus albus L, is an annual grain legume traditionally cultivated around the Mediterranean and along Nile Valley where it is used for human consumption, green manuring and as forage (Huyghe, 1997). The species (L. albus L) is diploid with 2 n = 50 and a generally self-pollinated crop with an average of 3.7 % outcrossing (Barbacki and Kapsa, 1960). The Egyptian genotypes have good productivity, but the high level of alkaloid in the seeds and their late maturity are considered the main problems of the genotypes. Cross breeding and selection in the segregating generations could improve these traits (Hoballah, 1991).

Diallel cross technique (El-Hady et al 1998; Abdalla et al 1999 and others) have been used to obtain considerable information on the magnitude of heterosis and to gain a better understanding of the nature of gene action involved in controlling quantitative characters. The exploitation of heterosis through synthetics and ultimately hybrids could pay off in improved yield potential. Several authors reported that the manifestations of heterotic effects in lupins ranged from significantly negative to significantly positive estimates for number of days to maturity, plant height and number of branches per plant as well as for seed yield and its components (Polkanova and Nagorskava, 1976; Lukashevich, 1981; Strautina, 1979; Okaz et al 1986; Agarkova et al 1991 and Sech & Huyghe, 1991). The studies of Savvicheva (1979) and Lukashevich (1981 & 1983) indicated that early maturity, plant height, number of pods and seeds per

plant as well as 1000 seed weight expressed intermediate inheritance or showed dominance or overdominance depending on the cross of F₁ intervarietal hybrids of L. luteus. On the other hand, Agarkova et al (1991) reported that heterosis for yield and stem length due to the additive effect of recessive genes could be greater than that resulting from heterozygosity in intraspecific hybrids of L. angustifolius. Debelvi and Derbenskii (1988) found that depression of lateral branching on the central raceme was predominant in hybrids of L. angustifolius when a determinate habit variety was the maternal parent.

The estimation of additive and dominance components of genetic variance is very important in evaluating the potential of any heterotic response. Narrow-sense heritability is the ratio of additive genetic variation to phenotypic variation and expresses the extent of resemblance between relatives and allows the determination of the response to selection. Yassen (1988) reported that genetic variances were due to dominance or additive effects of genes for flowering time, plant height, vield and some of its components and heritability estimates in narrow sense for 100-seed weight recorded 23.81, 24.61 or not calculated, depending on the cross in F₂ hybrids of L. albus. Sech and Huyghe (1991) stated that flowering time and reproductive characters showed high heritability and a mainly additive effects in F₁ hybrids of L. albus. Moreover, many of the yield characters were controlled by genes with additive effects, indicating that selection for these characters would be likely to be effective as reported by Klochko et al (1996) in L. angustifolius, and Khotyljova et al (1996) in L. luleus.

The present study was undertaken to estimate the magnitude of heterosis and inbreeding depression as well as understanding the nature of gene action and estimating heritability of different characteristics in white lupin hybrids. 2001/2002 using the diallel mating design including reciprocals among five widely diverse white lupin genotypes. Three varieties: Giza-1 (P₁), Dijon-2 (P₂) and Kiev Mutant (P₅) along with the two promising M₆ mutant lines: Line 20 (P₃) and Line 37/3 (P₄), which were assessed and selected from a previous study (**EI-Sayad**, 1997), were used as parents in the current investigation. Origin, pedigree and some characteristics of these genotypes are presented in Table (1).

MATERIAL AND METHODS

The present investigation was carried out during 1999/2000, 2000/2001 and

Genotype	Origin Pedigree		Date of maturity Growth habit		Alkaloid content
P ₁ (Giza 1)	Egypt	Land race (local variety)	Late	Indeterminate	High alka- loid (bitter lupin)
P ₂ (Dijon 2)	France	French variety	Late	Indeterminate	High alka- loid (bitter lupin)
P ₃ (Line 20)	Egypt	M6-induced mutant line derived from the Hun- garian cv. Kereskedelmi by 20 KR*	Late	Indeterminate	Moderate alkaloid (bitter lupin)
P ₄ (Line 37/3)	Egypt	M6- induced mutant line derived from cv. Giza 2 by 40 KR.	Early	Indeterminate .	Moderate alkaloid (bitter lupin)
P ₅ (Kiev Mutant)	Ukraine	Ukrainian variety derived from cv. White 70 by gamma-ray	Early	Determinate	Free alka- loid (sweet lupin)

Table 1. Origin, pedigree and some characteristics of the five white lupin genotypes.

* KR of gamma-rays.

In 1999/2000 growing season, all possible cross combinations among the five white lupin parents were made. In the second season (2000 / 2001) rehybridization were made to increase F_1 seeds as well as F_2 seeds were raised from the F_1 plants at Giza Research Sta-

tion. During 2001/2002 growing season, all the diallel mating progenies were evaluated in a randomized complete block design with three replications at Shandaweel Research Station. Seeds were sown 20 cm within rows, 60 cm apart. Sowing dates took place at early Novem-

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ber. Number of plants sampled per plot varied from 10 plants in parents and F₁'s to 50 plants in F_2 's. The following characters were recorded: number of days from sowing to 90 % maturity of plants per plot, plant height (cm), number of branches per plant, number of pods per plant, number of seeds per plant, 100seed weight (g) and seed yield per plant (g). Differences among genotypes were tested by conducting a regular analysis of variance of complete block design on plot mean basis. Heterosis and inbreeding depression determination were conducted as outlined by Foolad and Bassiri (1983). Appropriate "t" test were made for the significance of the F₁ crosses means from the mid and better parental values (Wynne et al 1970) and for the F2 crosses means from F1 values (Al-Rawi and Kohel, 1969). Values of Vr, Wr gcnetic components and ratios were estimated according to Hayman (1954 a & b).

RESULTS AND DISCUSSION

Statistical analysis presented in Table (2) revealed highly significant differences among tested genotypes, parents, crosses and parent vs. crosses for most studied characters, indicating wide genetic variability for studied variables.

The mean performance of parents, F_1 's and F_2 's including reciprocals for different characters are presented in Table (3). Kiev Mutant behaved as the earliest parent in maturity and recorded the lowest value of plant height, number of branches, pods, seeds and seed yield per plant as well as 100-seed weight. Giza-1 was the latest genotype and gave the highest number of branches per plant. Moreover, Dijon-2 had the highest value

of plant height, 100-seed weight and seed vield per plant. Line 20 recorded the highest number of pods and seeds per plant followed by Line 37/3. Regarding the test crosses in F_1 and F_2 generations, four crosses: Line 20 x Kiev Mutant, Line 37/3 x Kiev Mutant and reciprocal and Kiev Mutant x Giza-1 exhibited fewer number of days to maturity. None of the crosses exceeded significantly the tallest parent. The cross Dijon-2 x Giza-1 recorded the highest value of plant height, while crosses Dijon-2 x Kiev Mutant, Line 20 x Kiev Mutant and Line 37/3 x Kiev Mutant had shortest plant heights. Two crosses: Giza-1 x Line 37/3 and Kiev Mutant x Line 37/3 gave higher number of branches per plant. Concerning number of pods/plant, five crosses: Giza-1 x Dijon-2, Dijon-2 x Line 20, Kiev Mutant x Giza-1, Kiev Mutant x Dijon-2 and Kiev Mutant x Line 37/3 exhibited the highest number of pods/plant. Four crosses: Giza-1 x Dijon-2 and reciprocal, Dijon-2 x Line 37/3 and Kiev Mutant x Line 37/3 recorded the highest number of seeds/plant. None of the crosses exceeded the heaviest 100seed weight parent. Five crosses: Giza-1 x Dijon-2, Dijon-2 x Line 37/3, Line 20 x Dijon-2, Line 37/3 x Giza-1 and Line 37/3 x Line 20 showed higher estimates for 100-seed weight. With regard to seed vield/plant, five crosses: Giza-1 x Diion-2. Dijon-2 x Line 37/3, Line 20 x Dijon-Line 37/3 x Giza-1 and Kiev Mutant x Line 37/3 exhibited highest values for this trait. The previously mentioned crosses would prospect in white lupin breeding for improving the maturity date and productivity.

Regarding to reciprocal effects (Table 3), results revealed significant differences between some hybrids and their

	Source of	Df	Days to maturity		Plant height		No. of branches/plant		No. of pods/plant	
			F 1	F ₂	Fi	F ₂	F ₁	F ₂	Fı	F ₂
An	Replicates	2	0,50	0.63	34,06	4.69	0.38	1.15*	38.02**	2.70
Annals Agric	Genotypes	24	208.65**	104,08**	1408.59**	2333.91**	2.30**	2.47**	94.95**	84.17**
erric Sci	Parents	4	217.27**	217.27**	6031.60**	6031.60**	6.66**	6.66**	249.35**	250.01**
i 47(2)	Crosses	19	206.59**	84.30**	322.53**	1653.38**	1.29**	1.52**	55.59**	46.97**
2002	P. vs. C	ł	213.31**	26.44*	3551.72**	473.09**	4.01**	3.75**	225.19**	127.66**
	Error	48	4.47	4.07	22.28	27.33	0.93	0.32	5.13	3.81

Table 2. Mean squares of variance for number of days to maturity, plant height, number of branches, pods, seeds, 100-seed weight and seed yield per plant in F_1 and F_2 of 5 parental diallel cross.

* and ** significant at 5 % and 1 % level of probability, respectively.

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Source of		No. of see	ds/plant	100-see	d weight	Seed yield / plant		
variance	df	F1	F ₂	F ₁	F ₂	F1	F ₂	
Replicates	2	15.13	38.56	2,77	2.48	32.22	1.73	
Genotypes	24	1305.59**	1408.58**	29.83**	31.54**	259.22**	211.64**	
Parents	4	2946.57**	2946.57**	52.23**	52.23**	546.73**	546.73**	
Crosses	19	803.90**	967.92**	25.83**	28.76**	169.10**	136.84**	
P. vs. C	1	4273.92**	3629.17**	16.04	1.49	821.54**	292.46**	
Error	48	78.85	42.23	7.18	3.05	21.17	9.44	

* and ** significant at 5% and 1% level of probability, respectively.

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Genotype	Days to	maturity	Plant (CI	-	No. of brar	iches/plant	No. of pods/plant	
51	F ₁	F_2	$\overline{F_1}$	F ₂	F ₁	F ₂	\mathbf{F}_1	F ₂
P ₁ (Giza-1)	177.5	·	160,4		6.8		33.0	
P_2 (Dijon -2)	172.5		177.1		6.1		28.7	
P_3 (Line 20)	171.0		154.3		6.0		35.3	
P_4 (Line 37/3)	161.8		166.5		6.4		30.9	
P ₅ (Kiev Mutant)	157.6		66.0		3.1		12.9	
$P_1 \times P_2$	171.3	170.6	163.3	162.6	5.9	6.9	38.1	35.2
$P_1 \times P_3$	175.0	175.0	164.6	175.3	6.2	7.1	34.1	29.0
$P_1 \times P_4$	186.3	170.0	163.7	147.2	7.6	6.1	29.3	26.5
$\mathbf{P}_1 \mathbf{x} \mathbf{P}_5$	164.0	161.3	154.3	96.4	5.7	5.2	27.6	25.8
$P_2 x P_1$	169.6	171.0	172.4	178.4	6,8	6.8	40,4	29.8
$P_2 \times P_3$	166.0	168.6	166.9	151.3	6.6	7.3	30.2	41.1
$P_2 \times P_4$	163.6	168.0	147.3	167.2	6.1	7.0	35.0	28.8
$P_2 \times P_5$	160.6	160.6	147.9	107.5	6.0	5.2	33.8	27.3
$P_3 \times P_1$	169.6	167.0	174.8	165.1	6,9	6.2	35.3	32.3
$P_3 \times P_2$	166.0	176.3	161.5	171.7	6.1	6.2	36.8	32.8
$P_3 \times P_4$	162.0	169.3	155.3	142.3	5.7	6.6	32.3	35.3
P ₃ x P ₅	159.0	161.0	138.9	125.4	6.0	5.7	19.8	31.5
$P_4 x P_1$	164.6	170.6	170.6	166.4	6.6	6.3	30.9	36.3
$P_4 \times P_2$	164.3	165.3	170.9	171.5	6.8	5.6	31.7	31.4
P ₄ x P ₃	166.6	166.0	168.8	153.1	5.8	6.3	32.2	31.5
P ₄ x P ₅	154.6	158.3	153.4	114.7	4.5	5.2	33.7	27.7
P ₅ x P ₁	153.3	163.0	168.2	158.7	6.7	5.7	35.4	28.0
P ₅ x P ₂	154.3	164.6	170.8	166.6	5.7	7.2	34.1	33.9
P ₅ x P ₃	153.3	168.0	156.5	160.4	6.4	6.7	32.1	34.4
$P_5 \times P_4$	153.6	157.0	160.2	139.4	6.8	<u> </u>	32.4	35.3
LSD 0.05	3.54	3.31	7.75	8.58	1.03	0.93	3.72	4.58
0.0}	4.72	4.41	10.33	11.44	1.37	1.24	4.95	6.11

Table 3. Mean performance of parents, F_1 and F_2 generations for the studied characters.

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Construe	No. of see	ds/plant	100-seed v	veight (g)	Seed yield/plant (g)	
Genotype	$\overline{F_1}$	F_2	F ₁	F ₂	$\overline{F_1}$	F ₂
P_1 (Giza-1)	117,0		37.0		42.1	
P_2 (Dijon -2)	115,3		44.3		49.0	
P ₃ (Line 20)	125.5		38.6		47.3	
P ₄ (Line 37/3)	123,3		37.5		46.2	
P ₅ (Kiev Mutant)	50.6		32.5		16.7	
$P_1 \times P_2$	152.0	133.6	38.2	40.4	55.1	51.4
$\mathbf{P}_1 \times \mathbf{P}_3$	115.0	108.9	38,5	36.3	44.0	39.4
$P_1 \times P_4$	111.1	108.2	39.2	38.4	43.4	41.6
$P_1 \times P_5$	100,8	99.7	40.6	36.3	40.5	36.2
$P_2 x P_1$	149.7	134.0	40.8	32.9	60.6	38.3
$P_2 \times P_3$	100.6	167.8	41.0	34.2	41.2	57.1
$P_2 \times P_4$	130,8	131.9	41.8	39.5	55.2	52.6
$P_2 \times P_5$	130.2	106.3	36.7	33.9	43.8	34.7
$P_3 \times P_1$	134.8	131.9	33.4	36.8	45.8	48.2
$P_3 \times P_2$	137.0	126.6	44.6	35,5	61.9	44.5
P ₃ x P ₄	132,3	125.4	37,4	34.7	49.5	42.1
$P_3 \times P_5$	91.3	108.2	35,5	40.4	30.0	42.2
$\mathbf{P}_4 \mathbf{x} \mathbf{P}_1$	119.0	149.8	41.4	38.2	49.4	56.9
$P_4 \times P_2$	136.0	115.5	39.6	39,9	53.9	43.6
$P_4 \times P_3$	107.4	120.7	42.5	40,4	45,7	48.2
$P_4 \times P_5$	134.1	90.6	39.2	40.4	51.7	36.2
$P_5 \times P_1$	143.0	113.1	36.0	37.3	52.5	43.1
$P_5 \times P_2$	114.0	137.8	35,9	39.2	40.7	54.0
$P_5 \times P_3$	139.2	125.9	35,8	34.6	49.4	42.7
$P_5 \times P_4$	129.7	132.9	42.8	36.2	51.8	46.7
LSD 0.05	14,57	10.66	4.40	2.86	7.55	5.04
0.01	19.43	14.22	5.86	3.82	10.07	6.72

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reciprocals. One cross (Line 20 x Giza-1) for days to maturity, two crosses (Dijon-2 x Giza-1 and Kiev Mutant x Line 37/3) for plant height, one cross (Kiev Mutant x Line 37/3) for number of branches/plant, two crosses (Line 20 x Giza-1 and Kiev Mutant x Giza-1) for number of seeds/plant, one cross (Line 37/3 x Line 20) for seed index and one cross (Kiev Mutant x Giza-1) for seed vield/plant in both of F_1 and F_2 as well as two crosses (Kiev Mutant x Dijon-2 and Kiev Mutant x Line 73/3) in F₂ expressed significant reciprocal effects. Results suggested that earliness, seed yield and its components appeared to be influenced by the female parent. Reciprocal differences may result from plasmon differences or plasmon gene interactions. Also, maternal effects showed possibility to pass from F_1 to F_2 . The fact that such effects were detected in F_2 indicated the importance of maternal and plasmon effects in inheritance of these characters.

Heterosis percentages relative to mid (\overline{MP}) and better (\overline{BP}) parent are presented in Table (4). Regarding number of days to maturity, data showed that seventeen crosses were significantly earlier than the means of mid-parents with heterosis percentage ranging from -1.7 to -8.5. Additionally, heterosis percentages relative to mid parent were significantly positive in twelve, seven, eleven, thirteen, five and twelve crosses with a range of 4.3-48.6; 15.2-42.2; 15.0-63.9; 9.6-70.6; 11.0-22.3 and 13.2-78.6 % for plant height, number of branches, pods, seeds, 100-seed weight and seed vield/plant, respectively. However heterosis percentages relative to better parent were significant in five, two, five, four, one and three crosses with a range of -2.5 to -2.9; 4.9-9.0; 13.3-22.4; 12.9-29.9; 14.1 and 23.7-26.3% for days

to maturity, plant height, number of pods, seeds, 100-seed weight and seed yield per plant, respectively. The high magnitudes of heterotic values found in these materials were expected due to the diversity of the parents. Therefore, improvement would be expected from selection in the advanced segregating generations. Yassen (1984) found significant heterosis over mid parent for days to flowering, plant height, pods/ plant, seeds / pod, 100-seed weight and seed yield/plant. These findings are in accordance with those reported by Lukashevich (1981); Strautina (1979) and Agarkova et al (1991).

Results of inbreeding depression in F₂ generation for studied traits are given in Table (4). Concerning number of days to maturity, eight crosses exhibited significant inbreeding depression ranging from -2.4 to -9.5 %. Moreover, eleven, three, six, seven, five and eight crosses expressed significant inbreeding depression in F_2 for plant height, number of branches, pods, 100-seed weight and seed yield per plant with a range of 5.6-37.5; 14.9-19.7; 14.9-20.9; 9.6-32.4; 10.6-20.4 and 13.6-36.8 %, respectively. On the other hand, significant inbreeding gain was observed in some F_2 crosses for some studied characters. These results are in general agreement with those reported by Abdalla et al (1999) in faba beans.

Estimates of the genetic and environmental components of variance and other statistics derived from these estimates are presented in Table (5). Additive components of genetic variability (D) were significant or highly significant for all traits, indicating that the additive gene action was more important than the non- additive in controlling the inheritance of the

	D	ays to matur	ity		Plant height	<u> </u>	No. (of branches/	plant	N	o. of pods/pl	ant
Cross	Heterosis over MP	Heterosis over BP	Inbreeding depression ID									
$P_1 \times P_2$	-2.1**	-0,7	0.4	-3.3	-7.8*	0.4	-9.2	-13.2	-16.9*	23.3**	15.5**	7.6
$P_1 \times P_3$	0.3	2.3*	0.0	4.6*	2.6	-6.5**	-3.1	-8.8	-14.5	-0.3	-3.4	14.9*
$P_1 \times P_4$	9.8**	15.1**	8.8**	0.1	-1.7	10.1**	15.2*	11.8	19.7**	-8.4	-11.2	9.6
$P_1 \ge P_5$	-2.2**	4.1**	1.7	36.3**	-3.8	37.5**	14.0	-16.2*	8.8	20.0**	-16.4**	6.5
$P_2 x P_1$	-3.1**	-1.7	-0.8	2.1	-2.7	-3,5	4.6	0.0	0.0	30.7**	22.4**	26.2**
$P_2 \times P_3$	-3.4**	-2.9**	-1.6	0.7	-5.8*	9.4**	10.0	8.2	-10.6	-5.6	-14.4**	-36.1**
$P_2 \times P_4$	-2.2**	1.1	-2.7*	-14.3**	-16.8**	-13.5**	-3.1	-4.7	-14.8	17.5**	. 13.3*	17.7**
$P_2 \ge P_3$	-2,7**	1.9	0,0	21.6**	-16.5**	27.3**	30.4**	-1.6	13.3	62.5**	17.8**	19.2**
$P_3 \times P_1$	-2.8**	-0.8	-1.5	11.1**	9.00**	5.6*	7.8	1.5	10.1	3.2	0.0	8.5
$P_3 \times P_2$	-3.4**	-2.9**	-6.2**	-2.5	-8.8**	-6.3*	1.7	0.0	-1.6	15.0**	4.2	10.9
$P_3 \times P_4$	-2.6**	1.2	-4.5**	9.0*	-6.7**	8.4**	-8.1	-10.9	-15.8	-2.4	-8.5	-9.3
P ₃ x P ₅	-3.2**	0.9	-1.3	26.0**	-10.0**	9.7**	33.3**	0.0	5.0	-17.8**	-43.9**	-59.1**
$\mathbf{P}_{1} \mathbf{x} \mathbf{P}_{1}$	-3.0**	1.7	3.7**	4.3*	2.5	2.5	0,0	-2.9	4.6	-3.4	-6.4	-17.5*
$P_4 \ge P_2$	-1.7*	1.5	-0.6	-0.5	-3.5	-0.4	7.9	6.3	17.7*	6.4	2.6	0.9
P4 x P3	-0.1	3.0**	0.4	5.3*	1.4	9.3**	-6.5	-9.4	-8.6	-2.7	-8.8	2.2
P4 x P5	-3.1**	-1.9	-2.4*	31.9**	-7.9**	25.2**	-6.3	-29.7**	-15.6	53.9**	9.1	17.8**
$P_5 \times P_1$	-8.5**	-2.7*	-6.3**	48.6**	4.9*	5.7*	34.0**	-1.5	14.9*	53.9**	7.3	20.9**
P5 x P2	-6.5**	-2.1	-6.7**	40.5**	-3.4	2.5	23.9**	6.6	-26.3**	63.9**	18.8**	0.6
P5 x P3	-6.7**	-2.7*	-9.5**	42.0**	1.4	-2.5	42.2**	6.7	-4.7	33.2**	-9.1	-7.2
P5 x P4	-3.8**	-2.5*	-2.2	37.8**	-3.8	12.9**	41.7**	6.3	-4.4	47.9**	4.9	-8.9

Table 4. Percentage of heterosis over mid parent (\overline{MP}) and better parent (\overline{BP}) as well as percentage of inbreeding depression (ID) for various studied characters.

* and ** significant at 5 % and 1 % level of probability, respectively.

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		No. of seeds / p	lant		100-seed we	eight	Seed yield / plant		
Cross	Heterosis over MP	Heterosis over BP	Inbreeding depression ID	Heterosis over MP	Heterosis over BP	Inbreeding depression ID	Heterosis over MP	Heterosis over BP	Inbreeding depression ID
$P_1 \times P_2$	30.8**	29.9**	12.1**	-6.1	-13.8**	-5,8	20.8**	12.4	6.7
$P_1 \times P_3$	-5.2	-8.4	5.3	1.9	-0.3	5.7	-1.6	-7.0	10.5
$P_1 \times P_4$	-7.6	-9.9	2.6	5.1	4.5	2.0	-1.8	-6.1	4.2
$P_1 \times P_5$	20.3**	-13.8*	1.1	16.7**	9.7	10.6*	37.8**	-3.8	10.6
P ₂ xP ₃	28.8**	27.9**	10.5*	0.2	-7.9	19.4**	32.9**	23.7**	36.8**
$P_2 \times P_3$	-16.5**	-19.8**	-66.8**	-0.7	-7.5	16.6**	-14.5*	-15.9*	-38.6**
P2 x P4	9.6*	6.1	-0.8	2.2	-5.6	5.5	16.0**	12.7	4.7
P ₂ x P ₅	56.9**	12.9*	18.4**	-4.4	-17.2**	7.6	33.1**	-10.6	20.8**
P3 x P1	11.1*	7.4	2.2	-11.6*	-13.5*	-10.2	2.5	-3.2	-5.2
P ₃ x P ₂	13.8**	9.2	7.6	8.0	0.7	20.4**	28.4**	26.3**	28.1**
P3 x P4	6.4	5.4	5.2	-1.8	-3.1	7.2	5.8	4.7	14.9*
P3 x P5	3.6	-27.3**	-18.5**	-0.3	-8.0	-13.8*	-6.3	-36.6**	-40.7**
P4 xP1	-1.0	-3.5	-25.9**	11.0*	10.4	7.7	11.8	6.9	-15.2*
P4 x P2	14,0**	10.3	15.1**	-3.2	-10.6*	-0.8	13.2*	10.0	19.1**
P4 x P3	-13.7**	-14.4*	-12.4*	11.6*	10.1	4.9	-2.4	-3.4	-5.5
P4 x P5	54.1**	8.8	32.4**	12.0*	4.5	-3.1	64.1**	11.9	29.9**
P5 x P1	70.6**	22.2**	20.9**	3.5	-2.7	-3.6	78.6**	24.7**	17.8**
P5 x P2	37.4**	-1.1	-20.9**	-6.5	-19.0**	-8.4	23.7**	-16.9	-32.7**
P5 x P3	58.0**	10.9	9.6*	-4.0	-7.3	3.4	54.4**	4.4	13.6*
Ps x Pa	49.1**	5.2	-2.5	22.3**	14.1*	15.4**	64.4**	12.1	9.9

Table 4. Cont'd

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* and ** significant at 5 % and 1 % level of probability, respectively.

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Genetic	Days to	maturity	Plant h	eight
components	F _l	F ₂	F ₁	F
D	7093**±6.64	71.11** ± 1.61	2002.95** ± 45.84	2001.73** ± 45.84
H	59.07 * ± 17.50	13.06± 4.36	1449.31** ± 123.82	512.19 ± 218.17
H ₂	44.99± 15.84	9.81± 3.95	944.73** ± 112.31	363.19±197.88
F h ² E	-21.84 ± 16.15	9.07 ± 4.04	1978.04 ** ± 114.53	1002.17* ± 201.8
h ²	44.56* ± 10.69	4.79 ± 2.67	752.79** ± 75.82	95.27 ± 133.60
E	1.49 ± 2.64	1.31 ± 0.66	7.58 ± 18.71	8.81 ± 32.98
$(H_1/D)^{1/2}$	0.913	0.429	0.851	0.506
$H_2/4H_1$	0,190	0.188	0.163	0.177
K _D /K _R	0.823	1.188	2.008	1.800
h ² /H ₂	0.990	0.489	0.797	0.262
Heritability	0.807	0.897	0.521	0.852
t^2	0.175	0.001	0.002	0.088

Table 5. Estimates of genetic and environmental components with mean of genetic estimates and heritability values for various studed characters in F_1 's and F_2 's diallel cross.

* and ** significant at 5 % and 1 % level of probability, respectively.

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Table 5. Cont.

Genetic	No. of brai	nches/plant	No. of po	ds/plant	No. of see	eds/plant
components	F	F ₁	Fi	F ₁	F1	F ₁
D	2.09 ** ± 0.11	2.10**±0.13	81.21** ± 10.38	80.82** ± 3.76	956.75 ** ± 82.33	968.16** ± 57.61
H ₁	1.20 * ± 0.29	1.23 * ± 0.36	92.76 * ± 28.05	49.15 ± 10.15	1253.04 * ± 222.36	723.83* ± 155.59
. H ₂	0.87 * ± 0.26	0.81 ± 0.33	67.09 ± 25.43	33.14 * ± 9.20	910.13*±201.68	560.08*±141.13
F	1.68 ** ± 0.27	1.75** ± 0.33	81.38*±25.94	67.15 ** ± 9.38	1088.36* ± 205.67	681.11 *± 14392
h ²	$0.77* \pm 0.17$	0.72*±0.22	45.56 ± 17.17	25.06 ± 6.21	895.44** ± 136.16	765.20**± 95.28
E	0.13 ± 4.38	0.12 ± 5.42	2.15 ± 4.23	2.53 ± 1.53	25.43 ± 33.61	14.03 ± 23.52
$ \begin{array}{c c} P \\ P \\$	0.759	0.765	1.069	0.779	1.144	0.865
5 H ₂ /4H ₁	0.181	0.164	0.181	0.169	0.182	0.193
K _D /K _R	1.879	1.914	1.742	1.888	1.805	1.614
h²/H2	0.888	0.897	0.679	0.756	0.984	1.366
Heritability	0.517	0.549	0.403	0.578	0.295	0.594
t ²	0.008	0.023	0.008	0.008	0.003	0.015

* and ** significant at 5 % and 1 % level of probability, respectively.

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Genetic components	100-seed weight (g)		Seed yield / plant (g)	
	F1	F2		F ₁
D	$15.08* \pm 2.65$	16.40* ± 2.84	175.04** ± 22.16	179.20** ± 0.86
H_1	14.51 ± 7.18	22.76 ± 7.69	177.15 ± 59.86	89.43** ± 2.33
H ₂	12.09 ± 6.51	13.66 ± 6.98	142.45 ± 54.29	63.19** ± 2.11
F	8.71 ± 6.46	21.90 ± 7.11	133.81 ± 55.36	136.47 ** ± 2.16
h^2	1.92 ± 4.39	-0.33 ± 4.71	170.70* ± 36.65	$60.44 ** \pm 1.43$
E	2.34 ± 1.08	1.01 ± 1.16	7.20 ± 9.04	3.04 ** ± 0.35
$(H_1/D)^{1/2}$	0.981	1,178	1.006	0.706
H ₂ /4H ₁	0.208	0.150	0.201	0.177
K _D /K _R	1.410	1.972	1.562	1,903
h^2/H_2	0.159	-0.024	1.198	0.956
Heritability	0.450	0.290	0.470	0.647
t ²	0,755	0.117	0.045	0.004

 \ast and $\ast\ast$ significant at 5 % and 1 % level of probability, respectively.

characters and that selection would be effective for these traits in segregating generations. In each case H_1 and H_2 (dominance components of genetic variance) were highly significantly different from zero and H_1 was greater than H_2 in all cases, indicating that the positive and negative alleles at the loci for these traits were not equal in proportion in the parents. Theoretically, H₂ should be equal to or less than H₁ (Hayman, 1954 b). H₁ vales were greater than D for 100-seed weight, suggesting that non-additive genetic variance is more important than the additive variance. These results are in general agreement with those obtained by Sech & Huyghe (1991) and Khotyliova et al (1996). The covariance of additive and dominance effects (F) was insignificantly negative in one case (number of days to maturity in F_1) and significantly positive in other traits. Negative estimates of F indicate excess of recessive alleles. while the positive estimates of F indicate excess of dominant ones. Overall dominance effects of heterozygous loci (h^2) was insignificantly negative in one case (100-seed weight in F₂), while these effects were positive and significant in other traits, indicating that the effect of dominance was due to heterozygosity and that direction of dominant was positive. 'All estimates of the environmental variance (E) were insignificant for all traits studied except seed yield per plant, indicating that this trait has been affected by environmental factors. Yassen (1988) and Sech & Huyghe (1991) found high estimates of environmental variance (E) in white lupin seed yield. Overall degree of dominance estimated by $(H_1/D)^{1/2}$ ranged from complete to over dominance in number of pods, seeds, seed vield/plant and 100-seed weight which suggested

that the over dominance or espistasis are important in the inheritance of these traits. However, for number of days to maturity, plant height and number of branches/plant, the dominance relationship was suggested to be partial. Results reported by Yassen (1988); Sech and Huyghe (1991) and Klochko et al (1996) supported the evidence that both additive and dominance gene action are the most important components controlling variation in number of days to maturity, seed yield and its components. In general, the values of $H_2/4$ H_1 were slightly below the maximum value of 0.25. which arises when u=v=0.5 over all loci, indicating that positive and negative alleles were not equally distributed among the parents. The estimates of K_D/K_R were greater than one for all traits, indicating excess of dominant genes in the parents. The number of effective factors h^2/H_2 ranged from -0.024 to 1.366. It may be noted that this value is underestimated either when the dominance effects of all genes concerned are not equal in size and direction or when the distribution of the genes is correlated or when both conditions are fulfilled (Jinks, 1954). Narrow sense heritability estimates ranged from 0.290 for 100-seed weight to 0.897 for number of days to maturity. These results indicated that dominant genetic variance was more important for number of seeds/plant and 100-seed weight. However, days to maturity, plant height, number of branches, pods and seed vield/plant are mostly controlled by additive gene action and it is possible to breed for improving these traits. These results are in accordance with those reported by Yassen (1988); Sech and Huyghe (1991) and Klochko et al (1996) in lupins. Significance of t^2

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indicated the failure of the hypotheses postulated by Hayman (1954 b).

The Vr and Wr graphs are presented in Figs (1 to 14). The slope of the regression line in the Vr, Wr graph did not differ from unity for all traits except number of days to maturity in F_2 but was significantly different from zero for all characters except 100-seed weight in F_1 and F_2 generations. These results revealed the absence of genic interaction for all traits. However, days to maturity had complementary type of epistasis. The insignificant difference of the regression coefficient from zero indicated no real relationship among Vr and Wr and no further reliable interaction could be made.

The intercepts of the regression lines on the Wr axis deviated below the origin and showed over dominance for number of seeds / plant in F_1 . For the remaining traits partial dominance was observed as the regression lines deviated above the origin. The distribution of the parental arrays along the regression line for all the characters suggested wide distribution of dominant and recessive alleles among the parents. The scattering of parental arrays provided the information about the presence of different types of alleles in different parents for different characters

which is useful in practicing selection of parents for improving specific characters. However, the Vr, Wr graph varied among generations. This may be attributed to the large environmental variance. In the present study visual inspection of array points on the regression line of Vr. Wr graph will be restricted to all characters in F_1 and F_2 . For number of days to maturity array points for P₁ (Giza-1) and P₄ (Line 37/3) were farthest from the origin and had recessive alleles, while P5 (Kiev Mutant) appeared to possess most dominant alleles in this respect. On the other hand, P₅ (Kiev Mutant) was the farthest from the origin and had recessive alleles for plant height, number of branches, pods, seeds and seed yield per plant. With respect to 100-seed weight P₂ (Dijon-2) and P₅ (Kiev Mutant) were farthest from the origin and contained most recessive genes. In general, the graphic analysis indicated the inconsistency of array points over generations and that neither of the parents had the possible complete dominance or recessive alleles for any of the characters. It is, therefore, possible to accumulate the favourable genes the parents by using appropriate in procedures, i.e. recurrent sebreeding lection.



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REFERENCES

Abdalla, M.M.F.; D.S. Darwish; M.M. El-Hady and E.H. El-Harty (1999). Investigations on faba bans, *Vicia faba* L. 12- Diallel crossed materials grown under cages. *Egypt. J. Plant Breed. 3: 213-229.* Agarkova, S.N.; N.F. Pukhal and M.P. Miroshnikova (1991). Study of genetic aspects of heterosis in pea and *Lupinus* angustifolius. Genetika Moskva, 27 (10): 1799-1800.

Al-Rawi, K.M. and R.J. Kohel (1969). Diallel analysis of yield and other agronomic characters in *G. hirsutum* L. Crop Sci., 9: 779-783.

Barbacki, S. and E. Kapsa (1960). Variability in *Lupinus albus*. Genetica Polonica, 1: 61-101.

Debelyi, G.A. and V.L. Derbenskii (1988). Inheritance of the limited lateral branching character in hybrid populations of *L. angustifolius*. *Doklady Lenina*, 1: 22-24.

El-Sayad, Z.S. (1997). Breeding Studies on Lupin (L. albus) for some Important Economical Characters. pp. 89-102. Ph.D. Thesis. Fac. Agric., Al-Azhar Univ., Cairo, Egypt. El-Hady, M.M.; M.O. Omar; S.M. Nasr; K.A. Ali and M.S. Essa (1998). Gene action on seed yield and some yield components in F_1 and F_2 crosses among five faba bean (*Vicia faba* L.) genotypes. Bull. Fac. Agric. Cairo Univ., 37 (3): 369-388.

Foolad, M.R. and A. Bassiri (1983). Estimates of combining ability, reciprocal effects and heterosis for yield and yield components in a common bean diallel cross. J. Agrc. Sci., Camb., 100:103-108. Hayman, B.L (1954a). The analysis of variance of diallel tables. *Biometrics*, 10: 235-244.

Hayman, B.L (1954 b). The theory and analysis of diallel crosses. *Genetics*, 42: 336-355.

Hoballah, A.A. (1991). Breeding for Seed Yield, Its Components and Quality Traits in White Lupin (L. albus). pp. 136-140. Ph.D. Thesis, Fac. Agric., Cairo Univ., Egypt.

Huyghe, C. (1997). White lupin (L. albus L.). Field Crops Research, 53: 147-160.

Jinks, J.L. (1954). The analysis of continuous variation in a diallel cross of *Nicotiana rustica* varieties. *Genetics*, 39: 767-788.

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Khotyljova, L.V.; L.A. Tarutina; V. N. Kavtsevich and A.P. Savchenko (1996). Genetic control of quantitative characters in yellow lupin. *Abst. Book of 8 th Int. Lupin Conf.*, Pacific, California, USA.

Klochko, N.A.; L. Narvaes and N.F. Anikeeva (1996). Combining ability of *L.* angostifolius varieties in a diallel crossing scheme. *Plant Breed. Abst.*, 66 (4): 3808. Lukashevich, M.I. (1981). Breeding and genetical study of intervarietal hybrids of yellow fodder lupin. *Plant Breed. Abst.* 51 (5): 4306.

Lukashevich, M.I. (1983). Inheritance of some economical useful characters in hybrids of *L. luteus. Plant Breed. Abst.*, 53: 567.

Okaz, A.M.; E.A. El-Gharib; A.F. Abdalla; S.B. Mourad and H.E. Yassen (1986). Heterosis in some crosses of the genus Lupinus. Al-Azhar J. Agric. Res., 5: 296-306.

Polkanova, T.P. and M.D. Nagorskaya (1976). The breeding of L. luteus for rapid maturation. *Plant Breed. Abst.*, 46: 2537.

Savvicheva, K.L. (1979). Inheritance of growth period duration and yield in *L. leteus. Plant Breed. Abst. 49: 9260.* Sech, L. and C. Huyghe (1991). Diallel analysis in white lupin: consequences for breeding. *Agronomie.* 11: 719 - 726.

Strautina, E. (1979). Features of F_1 hybrids between mutants of L. albus L. Plant Breed. Abst., 49: 4297.

Wynne, J.C.; D.A. Emery and P.W. Rice (1970). Combining ability estimates in Arachis hypogaea L. II. Field performance of F_1 hybrids. Crop. Sci., 10: 713-715.

Yassen, H.E. (1984). Breeding Studies on Lupinus termis Forsk. pp. 61-72. M.Sc. Thesis, Fac. Agric., Al-Azhar Univ., Cairo, Egypt.

Yassen, H.E. (1988). Breeding Studies on Lupin for High Yield and Quality. pp. 94-97. Ph.D. Thesis, Fac. Agric., Al-Azhar Univ., Cairo, Egypt.

اجريت هذه الدراسة على خمسـة ابـاء فى الجيل الاول والتـاني خــلال المواسم (جيزة-١ ، ديجون-٢، السلالة ٢٠، السلالة ١٩٩٩ / ٢٠٠٠ ، ٢٠٠٠ / ٢٠٠١ ، ٣/٣٧، طفرة كييف) من الـترمس وجميـع ٢٠٠١ / ٢٠٠٢ بهدف تقدير قـوة الـهجين الهجن التبادلية بينها متضمنة الهجن العكسـية والتدهور نتيجة للتربية الذاتية وكذلك لدراسة

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عـدد القرون للنبات-عدد البذور للنسات- الانعز البة المبكرة. وزن ١٠٠ بذرة – وزن محصول النبات. تراوحت كفاءة التوريث بالمعنى الخلص

والأب الأحس، كذلك حدث تدهور معنوى حتى النضبج.

الأثر التجميعي كـانت معنويـة أو عاليـة كييف) محتويا على معظم العوامل المتنحيـة المعنوية وثابتة باختلاف الأجيال لجميع لصفة طول النبات وعدد الفروع الصفات المدروسة وكان هناك أيضه أدوراً للنبات وعدد القرون للنبات وعدد البذور لتأثير السيادة لجميع الصفات. وكان تـــأثير للنبات ووزن ١٠٠ بــنرة ووزن محصــول العوامل ذات الأثر التجميعي أكثر أهمية من

فعل الجين وتقدير نسبة التوريست والتوقسع دور العوامل ذات الأثر غير التجميعي فسي بإمكانية تحسين صفات عدد الأيام حتبى وراثة الصفات المدروسة مما يؤكد جدوى النضج- طول النبات- عدد الفروع للنبات- الإنتخاب لهذه الصفات خلال الأجيال

أظهرت النتائج تفوق بعض هجن الجيل من ٢٩ % لصفة وزن ١٠٠ بذرة السي الأول تفوقاً معنوياً مقارنة بمتوسط الأبوين ٨٩.٧ % لصفة عدد الأيام مسن الزر اعسة

في نباتات الجيل الثاني لبعض الهجن نتيجة أظهرت الأشكال البيانية لتوزيع الأباء للتربية الذاتية لجميع الصفات المدروسة. حول خط الانحدار انه كان مختلفا من كان هناك تأثيرات معنوبة وثابتة خــلال جبل الى آخر ولكن بوجـه عـام كان الأب الجيلين لبعسض الــهجن العكسية لجميــع الأول (جيزة-١) والأب الرابـــع (الســـلالة الصفات المدروسية مميا يوضيح أهمية (٣/٣٧) يحتوى كل منها عليه معظم التأثير ات الأمية في وراثة هذه الصفات. العوامل المتتحية لصفة عدد الأيام حتبي أظهر تحليل النتهائج أن العوامل ذات النضج بينما كهان الأب الخهامس (طفرة النبات.

> تحکيم: ا.د محمد يسن عبد الفتــــاح ا.د مظهر محمد فوزی عبد الله