# TOMATO (Lycopersicon esculentum Mill.) BREEDING FOR YIELD

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### **Abstract**

Inheritance of yield has been researched for 12 hybrids originating from diallel crossing of six inbreed lines of tomato. The inheritance mode of fruit yield in tomato has been determined by the mean generation analysis. Eleven hybrids showed significant reaction of additive genes. Six crossing combinations showed interaction dominant x dominant genes, in 6 hybrid epistasis additive x dominant genes; and in 4 hybrids additive x additive genes. Most significant interactions (most often for researched genotypes) were dominant x dominant genes, and within this combination duplicate type of epistasis. This type of gene interaction has unfavourable effect to yield, and in process of selection for better yield it causes the decrease of total yield.

**Key words:** tomato, inheritance, yield, mean generation analysis

# Introduction

Yield is one of the most important selection program parameters. The aim of this paper is to determine and explain the differences in the yield of five tomato hybrids. In this way, it is possible to define which gene effects are more important for the yield expression. The obtained results would suggest that there is possibility for applying such research in plant breeding, which is aimed at selecting high-yield hybrids.

The average fruit weight per plant represent a yield component. In their research, Das et al.1988 pointed out the following facts: 10-20 genes affected the fruit weight, the additive variance was prominent in fruit weight inheritance, there was partial dominance in light fruit weight, and there was no interallele interaction. Conti et al. (1984) established that were significant epistatic gene effects on fruit weight inheritance. Singh and Singh (1984) achieved similar results in their researches. They established the fact apart from additive and dominance gene effects there were also epistatic ones. As for the inheritance of this characteristic, Singh and Singh (1985) concluded the following: additive genes prevailed over dominance ones in inheriting fruit weight, additive gene values were higher than dominance gene values, and (j) and (l) epistatic types were established. Dhaliwal and Nandpuri (1988) found out that one out of the three investigated combinations did not suggest the existence of the digenic interaction in inheriting this feature, but the other two combinations had additive x dominance gene effects. Applying the 6-parameter model, proved additive activity epistatic gene effects on fruit weight and earliness as their features.

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#### Material and method

Six parental tomato genotypes that differed in fruit yield have been selected. Their crossing has been performed by applying the method of diallele crossing. Four of them (D-150, S-49, S-35 and H-52) were pure lines, one was a cultivar (SP-109) and one was a population (Kgž). All of them belonged to the collection of tomato genotypes of the Institute for Vegetable Crops in Smederevska Palanka. The trial was set in 2005 by applying the method of random block system in three replications. The inheritance mode of fruit shape was determined on the basis of the mean generation of crossing the two parents (P1, P2, F1, F2, BC1, and BC2) employing the additive - dominance model (a three-parameter model). Gene effects were marked with the following symbols: mean value (m); additive gene effects (d); dominance gene effects (h); additive x additive (i); additive x dominance (j); dominance x dominance (l). On the basis of testing the adequacy of this model, it was concluded that there was either presence or absence of epistatic gene effects. The adequacy of the additive - dominance model was tested in two ways: by the general test ( $\chi$ 2-test). Applying the six-parameter model (Mather and Jinks, 1982), the estimation of the values of epistatic gene effects was performed. This investigation dealt with digenic epistasis between additive, dominance, and additive x dominance genes.

### **Results**

Variability of the average yield values has been expressed by variation coefficient (V), which was below 10. Variation coefficient in F<sub>1</sub> generation has been moving from 1,84% (H-52 x SP-109) to 6,32% (S-49 x H-52) and has been lower than for parents. In F2 generation the variation coefficient has been from 7,54% (S-35 x H-52) to 18,91 % (D-150 x S-49) (Tab 1). Common  $\chi^2$ -test proved that model with three parameters is adequate only for hybrids D-150 x SP-109 and S-49 x H-52. For crossings where no adequate three parameter mode has been established, the model with six parameters has been used in order to evaluate the gene effect. Combination D-150 x SP-109 had significant values of additive and dominant gene effect. The evaluated values of dominant genes have been higher than additive genes, so the dominant genes could be crucial in inheriting the yield component within this combination. Values of additive and dominant gene effect in hybrid S-49 x H-52 have been significant and values of dominant gene effect have been higher than additive. Values of  $\chi^2$ -test for both hybrids have been compatible and exclude the presence of non allele interactions (epystasis). Testing of other hybrids by using three parameters model proved the epistatic gene reaction. Based on significant  $\chi^2$ -tests, the presence of two significant genes, the presence of two-gene epistasis has been recorded (Table 2).

Evaluated genetic parameters for these crossings have been calculated by analysing average generation by using six parameters model. Analysing the model of genetic parameters by using the model with six parameters only two-gene epystasis have been found (between additive, dominant and additive and additive x dominant genes – Table 3).

No significant value of epystatic gene effect has been found for hybrid D-150 x S-49, although  $\chi^2$  and scaling tests pointed to its presence. It is probable that these were higher gene interaction levels, while this method determinates only two gene epystasis. Heritability level in higher sense, within this hybrid group was from 87,52% (D-150 x Kg-ž) to 97, 56% (D-150 x S-49), which were high values of heritability (Table 2).

Hybrids with common parent S-49 had genetic parameters only for crossing S-49 x SP-109. Additive gene effects and epystasis effect dominant x dominant genes had significant values. Hybrids S-35 x SP-109 had significant interaction dominant x dominant but with opposite

pre-sign than dominant gene values, which proves the dominant type of epystasis. Crossing combination H-52 x Kg-ž had duplicate type of epistasis (Table 3).

### **Discussion**

From 12 researched hybrids including hybrids D-150 x SP-109 and S-49 x H-52, za koje je inheritance has been found by using model with three parameters, the significant level of hybrids has been found only in 7 hybrids. All crossing combinations had additive and dominant gene effect, where dominant values were higher than estimated values of additive genes, except for combination Kg-ž x SP-109 where established values of dominant genes have been negative. Dominant genes usually dominate over additive, which is compatible with Sonone et al. (1986) and Singh and Singh (1989), results.

In our research, the significant level of additive and dominant level has been proved for six hybrids. In S-35 x Sp-109 hybrid, dominant genes values have been higher than additive, while in hybrid Kg-ž x SP-109 value of additive genes have been higher than values of dominant genes. These conclusions are compatible with Singh et al (1988) who proved that both gene effects have significant values and that both gene effects impact the yield expression.

Significant level of additive genes has been found in five hybrids. These results are compatible with many research results proving that additive genes are dominant in yield inheriting: Kale et al. (1988), Sonone et al. (1986), Crill et al. (1987), Das et al. (1988), Singh and Singh (1993), Hegazi et al. (1995).

If genotypes were to be classified according to epiystasys expression frequency, in 12 researched hybrids the most common was (l) type of epistasys (6 times) and (j) type (6 times) and they could be the most significant which is compatible with Singh and Singh (1985), results who proved that (j) and (l) epystasis types are more significant than (i) epistasis in inheriting the yield impact per plant. Important notification in our research is that large number of hybrids had duplicate epistasis. In all combinations with significant values (h) and (l) this type of epistasis has been found, which proves that this type of interactions in stable in yield inheriting. This type of epistasis impacts the lower expression of the researched trait. Practically, this is a negative appearance in tomato breeding for increased yield. Our results have been compatible with Dhaliwal and Nandpuri (1988) who analysed three hybrid combinations with expended two-gene model and found duplicate type of epistasis. They considered that selection in the first generations after crossing would be slow, while in later generations much faster.

Results of interactive gene reaction have not been completely compatible with Singh et al. (1989). Our research proved interaction additive x dominant (j) type of epistasis in three hybrids (S-35 x Kg-ž, S-35 x SP-109 i Kg-ž x SP-109). These hybrids have been connected with the same mother or father components, which lead to a conclusion that this type of epistasic effect could be fixed by selection of parents. These hybrids have had second type of epistasis (duplicate type between dominant and dominant x dominant genes), so they have been the most important in analysing this trait.

### **Conclusion**

From 12 researched hybrids, 11 showed significant reaction of additive genes; in 7 significant reaction of dominant genes. In 6 crossing combinations the interaction dominant x dominant genes has been found in 6, in 6 hybrids epistasis additive x dominant genes; and in 4 hybrids

additive x additive genes. As the most significant interactions the most common for researched genotypes have been dominant x dominant genes and within this combination duplicate type of epistasis. This type of interactive gene effect has unfavourable impact on yield in selection process with the aim to increase yield, since it effects the decrease of total yield.

### Acknowledgements

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### Literature cited:

- Conti S. Candilo M.; Franssoldati P. 1984. Analysis Aof fruit unifornity in two processing tomato. (abstract) Genetica Agraria, Vol.38, No 3, 327-328.
- Crill I.P.; Tomas P.; Brayan H.H.; Hawkins W.1987. Hybrid vigor in Florida fresh market tomato "Independence". Proceedings of the Florida State Horticurtural Society, Vol 99, 343-350.
- Das N.D.; Chosh S.S.; Chattopadhyay T.K. 1988. Genetics of cultitative characters in tomato (*Lycopersicon esculentum Mill.*). Indian Journal of Agriciltural Sciences, Vol 58, No1, 64-65.
- Dhaliwal M.S.; Nandpuri K.S. 1988. Genetics of yield and its components in tomato. Annals of Biology, Vol 4, No 1-2, 75-80.
- Hegazi H.H.; Hassan H. M.; Moussa A.G. 1995. Heterosis and heritability estimation for some characters of some tomato cultivars and their hybrid combinations. Alexandria Journal of Agricultural Research, Vol 40, Vol 2, 265-276.
- Kale P.B.; Dod V.N.; Supe V.S. 1988. Genetics variability and correlation studies in tomato. PKV Research Journal, Vol 12, No 2, 115-118.
- Mather K.; Jinks I.L 1982. Biometrical Genetics. Third Edition. Champan and Hall. London
- Singh R.K.; Singh S. 1984. A study of interaction of additive, dominance and epistatic gene effects with micro- and makro-environmens in two tomato triple test crosses. Journal of Agricultural Science, Vol 103, No 1, 53-57.
- Singh R.P.; Singht S. (1985): Detection and estimation of components of genetic variation for some metric traits in tomato (*Lycopersicon esculentum Mill.*). TAG, Vol 70, 80-84.
- Singh P.K.; Singh R.K., Saha B.C.; Rajesh K. 1988. Genetic variability in tomato (*Lycopersicon esculentum Mill.*). Indian Journal of Agricultural Sciences, Vol. 58, No 9, 718-720.
- Singh U.P.; Taanki I.; Singh R.K. 1989. Studies on order effect and epistatic components for yield in double-cross hybrids of tomato. Haryana Journal of Horticultural Sciences, Vol 18, No 3-4, 265-271.
- Singh R.K.; Singh V.K. 1993. Heterosis breeding in tomato (Lycopersicon esculentum Mill.). Annals of Agricultural Research, Vol 14, No 4, 416-420.
- Sonone A.H.; Yadar M.D.; Thombre M.D. 1986. Combining ability for yield and its components in tomato. Journal of Maharashtra Agricultivar Universities, Vol 11, No 3, 288-290.

Table 1: The average value and variability factors for yield per tomato plant (g)

Genotypes	X	<u>+</u> S <sub>x</sub>	σ	V	Genotypes		X	<u>+</u> S <sub>x</sub>	σ	V
D-150	907,4	36,77	63,69	7,02	S-49 x KG-ž	BC <sub>2</sub>	2277,7	25,02	43,34	1,90
D-150 x S-49 F <sub>1</sub>	1839,9	27,29	47,27	2,57	S-49 x SP- 109	$F_1$	1073,5	17,42	30,17	2,81
$F_2$	1847,2	201,7	349,4	18,91		$F_2$	1685,0	105,9	183,50	10,89
$BC_1$	1778,0	157,1	272,2	15,31		$BC_1$	1872,2	129,9	225,08	12,02
$\mathrm{BC}_2$	1952,9	34,74	60,20	3,08		$BC_2$	1418,5	81,34	140,9	9,93
D-150 x S-35 F <sub>1</sub>	1768,4	29,70	51,45	2,91	S-35		1572,3	40,14	69,52	4,42
$F_2$	1463,6	124,38	215,44	14,72	S-35 x H-52	F1	1383,8	18,34	31,78	2,29
$BC_1$	1701,4	71,22	123,36	7,25		$F_2$	1254,0	54,56	94,49	7,54
$\mathrm{BC}_2$	1795,4	124,59	215,80	12,02		$BC_1$	1332,8	84,49	146,34	10,98
D-150 x H-52 F <sub>1</sub>	1679,8	29,16	47,05	2,80		$BC_2$	1484,7	87,58	151,69	10,22
$F_2$	1583,5	141,56	245,2	15,48	S-35 x Kg-ž	$F_1$	1672,2	21,39	37,05	2,22
$BC_1$	1779,3	63,55	110,08	6,19		$F_2$	1364,0	77,97	135,06	9,90
$\mathrm{BC}_2$	1255,7	36,37	63,00	5,02		$BC_1$	1303,6	78,49	135,15	10,43
D-150 x Kg-ž F <sub>1</sub>	2090,5	45,62	79,03	3,78		$BC_2$	1213,3	53,90	93,36	7,69
$F_2$	1494,0	112,21	194,36	13,01	S-35 x SP-109	$F_1$	1568,5	94,66	163,97	10,45
$BC_1$	1800,8	10,91	18,89	1,05		$F_2$	1191,2	65,41	113,3	9,45
$\mathrm{BC}_2$	1783,6	9,99	16,54	0,93		$BC_1$	1456,7	102,71	177,91	12,21
D-150 x SP-109 F <sub>1</sub>	1585,3	39,56	68,52	4,32		$BC_2$	1624,8	29,59	51,24	3,15
$F_2$	1204,2	123,27	213,51	17,73	H-52		1122,3	36,60	63,40	5,65
$BC_1$	1188,7	33,27	57,45	4,83	H-52 x Kg-ž	$F_1$	1698,7	21,39	37,05	2,18
$\mathrm{BC}_2$	1010,0	62,71	108,61	10,73		$F_2$	1738,6	77,97	135,06	7,78
S-49	1580,3	38,62	66,88	4,23		$BC_1$	1682,8	78,49	135,95	8,08
S-49 x S-35 F <sub>1</sub>	2195,1	45,48	78,77	3,59		$BC_2$	1213,3	53,9	93,36	7,69
$F_2$	1821,9	104,25	180,57	9,91	H-52 x SP-109	$F_1$	1471,6	15,62	27,05	1,84
$BC_1$	2107,2	25,68	44,48	8,11		$F_2$	1596,4	65,21	112,95	7,08
$\mathrm{BC}_2$	2190,1	198,29	341,46	16,20		$BC_1$	1780,3	32,89	56,96	3,20
S-49 x H-52 F <sub>1</sub>	2549,6	92,97	161,02	6,32		$BC_2$	1087,9	56,84	98,46	9,05
$F_2$	2048,7	218,82	368,62	17,99	Kg-ž		1528,6	35,87	62,14	4,07
$\mathrm{BC}_1$	1937,8	172,7	299,10	15,44	KG-ž x SP-109	$F_1$	16,54	29,74	51,51	3,11
$\mathrm{BC}_2$	1776,8	48,33	83,72	4,71		$F_2$	1534,0	99,93	173,08	11,28
S-49 x Kg-ž $F_1$	1987,7	42,82	74,17	3,73		$BC_1$	1303,7	14,37	250,06	19,18
F2	1878,1	85,21	147,58	7,86		$BC_2$	1078,3	34,22	59,28	5,50
$BC_1$	1732,8	27,48	47,60	8,75	SP-109		730,5	15,84	27,45	3,76

LSD<sub>(0,05)</sub> 77,15 (0,01) 109,76 Table 2: Genetic analysis of gene effect per plant using model with three parameters

Genotypes	(m)	t <sub>(m)</sub>	(d)	t <sub>(d)</sub>	(h )	$t_{(h)}$	$\chi^2$	$h^2$
D-150 x S-49	1311,43	51,76	-310,34	12,31	601,25	15,61	67,16**	97,58
D-150 x S-35	1264,23	46,65	-311,77	11,46	534,86	13,05	24,64**	91,99
D-150 x H-52	1021,88	40,76	-45,28	1,82	665,39	17,28	58,53**	93,95
D-150 x Kg-z	1291,76	57,03	-62,22	4,92	970,24	21,44	150,10**	87,52
D-150 x SP-109	798,05**	48,27	70,25**	4,32	723,58**	17,21	6,63	93,31
S-49 x H-52	1347,27**	51,33	231,87**	8,88	1145,67**	4,30	1,13	91,42
S-49 x SP-109	1179,91	58,40	440,07	21,73	-84,74	3,10	70,83**	93,96
S-35 x H-52	1337,97	50,29	210,37	7,71	41,27	1,25	10,75**	65,71
S-35 x SP-109	1131,99	52,67	379,15	17,78	963,04	17,12	122,81**	80,56
H-52 x Kg-z	1302,58	23,44	-232,88	9,43	300,34	10,56	173,34**	85,32
H-52 x SP-109	1048,43	57,63	315,361	17,28	466,91	18,67	113,34**	87,18
Kg-ž x SP-109	1125,42	61,02	399,78	21,64	495,68	14,70	14,86**	92,33
t <sub>0,05</sub>		2,228		2,776		2,179		
0,01		3,169		4,604		3,055		

Table 3 : Genetic analysis of gene effect of yield per plant by using a model with six parameters

parameters / hybrids		-		-		-							
		[m]	$\mathbf{t}_{(\mathbf{m})}$	[d]	$\mathbf{t}_{(\mathbf{d})}$	[h]	$\mathbf{t}_{(\mathbf{h})}$	[i]	$\mathbf{t_{(i)}}$	[j]	$\mathbf{t_{(j)}}$	[1]	$\mathbf{t}_{(\mathbf{l})}$
D-150 x S-49		1168,5	1,28	-336,4**	12,59	2043,5	1,05	75,5	0,08	323,6	1,99	-1371,8	1,45
D-150 x S-35		100,8	0,69	-332,6**	11,95	3784,0 *	2,78	1139,1	1,92	475,8	1,59	-2116,4 *	2,69
D-150 x H-52		1278,8	2,17	-107,4 *	4,02	817,3	0,66	-263,1	0,45	1261,9**	7,47	-416,5	0,64
D-150 x KG-ž		2587,0	0,06	-313,3**	12,12	3807,0**	4,21	1191,5 *	2,66	658,7**	11,11	-1744,0**	3,77
S-49 x SP-109		1314,8 *	2,46	424,9**	20,74	1723,4	1,33	-159,4	0,30	57,5	0,17	-1964,3 *	2,52
S-35 x H-52		728,7	1,99	223,9**	8,05	1448,0	1,49	618,8	1,69	-732,4 *	2,54	-793,3	1,29
S-35 x SP-109		-246,8	0,69	420,9**	19,23	3936,3**	4,50	1398,1**	3,91	-1178,1**	5,27	-2121,3 *	3,78
H-52 x KG-ž		2540,4**	8,67	21,871	0,84	-3838,0**	4,68	-990,0 *	3,39	-4509,9**	17,31	2969,0**	29,69
H-52 x SP-109		1575,8**	5,03	195,9**	9,56	186,9	0,27	-649,5	2,08	992,9**	7,12	-290,9	0,74
KG-ž x SP-109		2501,6**	5,05	399,1**	21,39	-3023,0 *	2,55	-1371,9 *	2,77	-347,34	1,18	2176,3**	3,06
	t <sub>0,05</sub>		2,228		2,776		2,179		2,447		2,306		2,179
	$t_{0,01}$		3,169		4,604		3,055		3,707		3,350		3,055

## СЕЛЕКЦИЈА НА ПРИНОС ПАРАДАЈЗА (Lycopersicon esculentum Mill.)

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### Резиме

Парадајз спада у једну од напрофитабилнијих повртарских врста. Величина и квалитет приноса, поред раностасности, односно динамике сазревања плодова, дају привредни значај парадајзу. Познавање начина наслеђивања у селекцији ове врсте може да представља умногоме скраћивање времена које је потребно да се селекционише комерцијални хибрид и осигурати ми стабилност у његовој експресији. Наслеђивање приноса испитивано је код 12 хибрида пореклом из диалелног укрштања шест инбред линија парадајза (Д-150, С-49, С-35 и Х-52). Код 11 хибрида се испољило сигнификантно деловање адитивних гена; у 7 сигнификантно деловање доминантних гена на принос плодова парадајза. Код 6 комбинација укрштања утврђена је интеракција доминантни х доминатни гени, у 6 хибрида епистаза адитивни х доминантни гени; и у 4 хибрида адитивни х адитивни гени. Као најзначајније интеракције (најчешће се појављују код испитиваних генотипова), могу се издвојити доминантни х доминантни гени, и у оквиру њих дупликатни тип епистаза. У прктичној селекцији парадајза треба одабрати линије које усливљавају већи број позитивних генетских ефеката у потомству, а који ће у комерцијалним комбинацијама дати перспективне хибриде.

Кључне речи: парадајз, принос, наслеђивање, анализа просека генерација

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