

Complex relationships between studied traits are telling us how much could be restricted adaptive biological variation, and which are developmental strategies to reduce enormous potentials of such a variation to efficient ontogenetic programs realized in eclosed adult individuals.

References: Cluster, P.D., D. Marinković, R.W. Allard, and F.J. Ayala 1988, PNAS 84: 610-614; Jovanovska, M., 1990, Ph.D. thesis at the University of Belgrade; Marinković, D., M. Milošević, and M. Milanović 1986, Genetica 70: 43-52; Milošević, M., and D. Marinković 1989, Genetique Selection Evolution 21: 17-32.

**Koryakov, Dmitry E.<sup>1</sup>, Elena S. Belyaeva<sup>2</sup>, and Igor F. Zhimulev<sup>1,2</sup>.** <sup>1</sup>Department of Cytology and Genetics, Novosibirsk State University, Novosibirsk 630090, Russia <sup>2</sup>Institute of Cytology and Genetics, Novosibirsk 630090, Russia. The new *Drosophila melanogaster* nonlethal inversion, arisen from the *In(2R)bw<sup>VD2</sup>*.

In the stock *In(2R)bw<sup>VD2</sup>/CyO*, received from the Umea *Drosophila* stock center, part of the flies spontaneously lost lethality in homozygous condition. The new stock, called *In(2R)bw<sup>VK</sup>* - brown Variegated of Koryakov, was constructed. Cytological analysis has shown that the new stock has the same inversion - 41A-B - 59D6-E1, as the *bw<sup>VD2</sup>*.

*bw<sup>VD2</sup>* and *bw<sup>VK</sup>* with other rearrangements, displaying *bw*-Variegation phenotype (*In(2R)bw<sup>VD1</sup>*, *In(2R)bw<sup>V1</sup>*, *T(2,3)bw<sup>VD4</sup>*, and *T(2,3)bw<sup>V4</sup>*), and with two lethals *uex<sup>1</sup>* and *rl<sup>9</sup>*, located according to Dimitri (1991) in heterochromatin distally and proximally to the break point of the inversion *bw<sup>VD2</sup>*, respectively (Table 1). In the case of *bw<sup>VK</sup>* the

expected ratio of the *Cy/Cy<sup>+</sup>* offspring is 1:1, in the case of *bw<sup>VD2</sup>* the ratio is 2:1, and the percentage of surviving flies was calculated from these ratios. It should be noted that the surviving flies *bw<sup>VK</sup>/bw<sup>VD2</sup>* slightly differ in reciprocal crosses. In comparison with *bw<sup>VD2</sup>*, viability of heterozygotes of *bw<sup>VK</sup>* with *bw<sup>VD1</sup>*, *bw<sup>V1</sup>*, and *bw<sup>VD4</sup>* is noticeable higher, but with *bw<sup>V4</sup>* it is lower. Viability of *bw<sup>VK</sup>/uex<sup>1</sup>* flies is greatly higher than expected.

In some crosses the flies with wing defects were found in small amounts (Table 2), which is supposedly due to position effect variegation of the *uex* locus, because a very similar phenotype was described for the *uex* homozygotes (Maeda, 1984).

So, losing the lethality for *bw<sup>VK</sup>* is accompanied by increasing viability with other *bw*-Variegated rearrangements.

Comparison of genetic inactivation as a result of position effect variegation of the *bw*-locus was performed with the inversions *bw<sup>VD1</sup>*, *bw<sup>VD2</sup>*, and *bw<sup>VK</sup>*.

Cis-effect of *bw<sup>+</sup>*-variegation in rearranged chromosomes can be observed in *R(bw<sup>+</sup>)/R<sup>+</sup>(bw)* heterozygotes where *R* is eu-heterochromatin rearrangement. Precise calculation of pigmented facets is possible if their number is not higher than 30 per eye. Therefore, we established five classes of variegation: 0, 1-10, 11-20, 21-30 and more than 30 pigmented facets per an eye. As a rule, both eyes of the fly

Table 1. The viability of heterozygotes of *bw<sup>VD2</sup>* and *bw<sup>VK</sup>* with other rearrangements, displaying *bw*-Variegation phenotype and with two lethals (at 25°C).

		<i>bw<sup>VK</sup></i>		<i>bw<sup>VD2</sup></i>	
		number of flies	Survived (in %)	number of flies	Survived (in %)
<i>bw<sup>VK</sup></i>	Cy			124	
	Cy <sup>+</sup>			116	96.7
<i>bw<sup>VD2</sup></i>	Cy	133			
	Cy <sup>+</sup>	107	89.2		
<i>bw<sup>V1</sup></i>	Cy	121		160	
	Cy <sup>+</sup>	94	87.4	38	57.6
<i>bw<sup>VD1</sup></i>	Cy	135		164	
	Cy <sup>+</sup>	19	24.7	0	0
<i>bw<sup>V4</sup></i>	Cy	102		110	
	Cy <sup>+</sup>	41	57.3	33	68.8
<i>bw<sup>VD4</sup></i>	Cy	82		134	
	Cy <sup>+</sup>	22	42.3	8	17.0
<i>uex<sup>1</sup></i>	Cy	82		157	
	Cy <sup>+</sup>	168	134.4	63	86.3
<i>rl<sup>9</sup></i>	Cy	122		153	
	Cy <sup>+</sup>	126	101.6	80	102.6

Table 2. A number of flies with wing defects from some crosses.

	number of flies	
	completely unexpanded wings	turbid, crumpled wings
<i>bw<sup>VK</sup>/bw<sup>V4</sup></i>	7*	23**
<i>bw<sup>VK</sup>/bw<sup>VD4</sup>***</i>	4*	5
<i>bw<sup>VD2</sup>/bw<sup>V1</sup></i>	0	3
<i>bw<sup>VD2</sup>/bw<sup>V4</sup></i>	14	18
<i>bw<sup>VD2</sup>/bw<sup>VD4</sup></i>	3	2

\* - Black mouthparts. \*\* - One wing may be smaller than another one, sometimes with bubbles. \*\*\* - In this cross a fly with notched wings planted apart (looking like Notch and Dichaete) was found

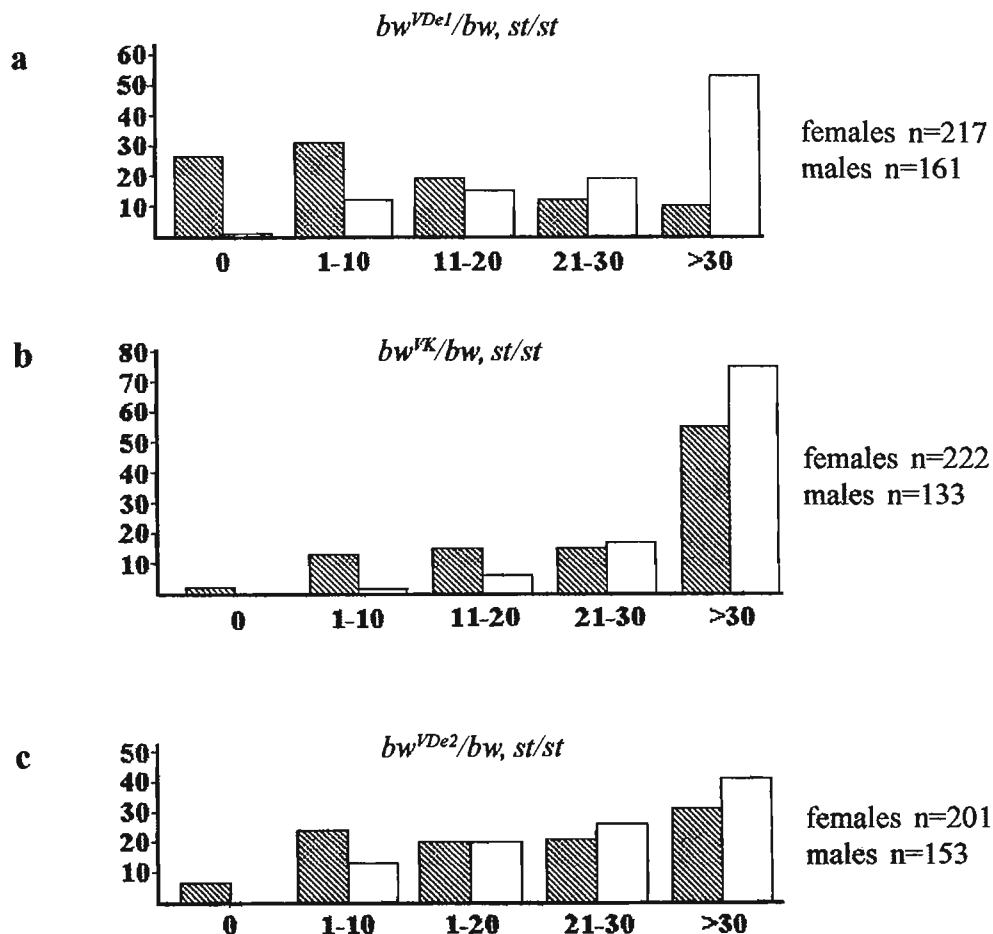


Figure 1. Comparison of the cis-inactivation of  $bw^+$  in different strains at temperature 18°C. Abscissa, classes with different number of pigmented facets; ordinate, number of eyes of every class (in%). Open columns are males, dashed are females.

belong to the same or to two neighboring classes.

Comparison of  $bw^+$  cis-effect in chromosomes with different inversions gives the following row:  $bw^{VDe1} > bw^{VDe2} > bw^{VK}$ . Cis-inactivation is stronger in females than in males in all these cases (Figure 1).

Cytological analysis shows that in  $X0; bw^{VK}/+$  males at 16°C (maximum enhancement of position effect variegation) only 11% of nuclei shows heterochromatization of the 59E region in rearranged homologue. In these cases the light chromosome region near 59D1-4 disappears and the bands 59D1-4 join to the chromocenter; they have diffuse morphology or are not seen at all. At the same conditions in  $bw^{VDe2}$  heterochromatization was found in 66% of nuclei (Belyaeva *et al.*, 1997).

So, the data show that  $bw^{VK}$  has noticeably decreased heterochromatization and frequent inactivation of the  $bw^+$  gene in comparison with  $bw^{VDe2}$ .

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