

Postlethwait, J.H. University of California, Irvine, California. Effect of X-rays of the eye of heterozygous Antennapedia flies.

bearing bristles which appeared in the region of the vertex or orbital bristles. The morphology of the supernumary bristles and the ground pattern was characteristic of this region of the head. The bristles often encroached upon ommatidia, causing the eye to be smaller. Frequently, a sector in the dorsal anterior part of the eye also contained bristles.

The frequency at which defects occurred in Antp^R flies varied with the time of irradiation (Fig. 1), and

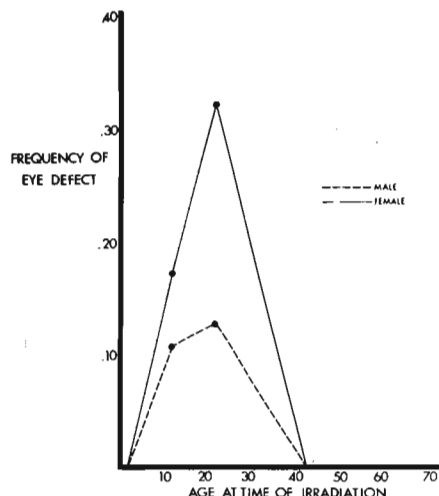


Fig. 1. Frequency of X-ray induced defects in the eye of Antp^R flies vs. time of irradiation.

there was a peak at 24 hours after egg deposition. The penetrance of the eye defect was greater in females than males. In irradiated progeny of the cross y;mwh Antp^R/mwh Sb Ubx X Df(1)sc⁸, w^a/Dp(1;3)sc^{J4} and the cross Antp^R/Sb Ubx X y sn³ f^{36a}, 72 of 383 Antp^R individuals irradiated prior to 45 hours had defective eyes, while only 1 of 944 of their Sb Ubx siblings irradiated at this time had a defective eye. This indicates that the factor responsible for the eye defect is on the Antp^R chromosome. Unirradiated Antp^R flies did not show such

irregularities in our experiments, nor did progeny of the cross y;mwh X Df(1)sc⁸, w^a/Dp(1;3)sc^{J4} and y sn³ f^{36a} X Oregon R irradiated with 1000r. Haskins and Enzmann¹, however, did obtain eye defects similar to the ones reported here after irradiation of an eosin stock.

Malformations of the head capsul without X-rays occur in *D. melanogaster* heterozygous for Antp^{LC} 2, *D. funebris* bearing aristapedia³, and *D. hydei* bearing ss^{Antp} 4. The eye effect in *D. funebris* is more pronounced in females than males³, as it was in our experiments. The gene erupt (er), also on the third chromosome, leads to a bristled structure protruding from the eye after irradiation of certain stocks at ten hours after egg laying. It is not known whether the eye defects reported here are due to the Antp^R gene itself, or to some other locus, such as er, on the Antp^R third chromosome.

References: 1. Haskins, D.P. and E.V. Enzmann, 1937 Amer. Nat. 71: 87-90; 2. Le Calvez, J., 1948 Bull. Biol. Fr. Belg. 82: 97-113; 3. Tiniakoff, G., 1939 DIS 11:52; 4. Gloor, H. and H. Kobel, 1966 Rev. Suisse Zool. 73: 229-252; 5. Glass, B., 1944 Genetics 29: 436-446.

Sanjeeva Rao, M. Osmania University, Hyderabad, India. The alteration of X-ray induced genetic damage by aflatoxin in *D. melanogaster*.

The treatment of *Drosophila* flies with certain chemicals, and antibiotics prior to irradiation has altered the genetic damage (Sobels 1961, 1963, 1964, 1965; Burdette 1961, Clark 1963 and M.S. Rao 1965) and one of the methods offered for explanation was the inhibition of protein

synthesis.

Aflatoxin is a collective name given to a group of highly toxic substances produced by certain strains of the mould *Aspergillus flavus*. The biological effects of this substance include: (i) inhibition of protein synthesis and also (ii) inhibition of m-RNA synthesis possibly through RNA polymerase.

With a view to find out whether aflatoxin would be able to alter the genetic damage akin to antibiotics experiments were undertaken to assess the alteration if any.

Oregon-K males of *D. melanogaster* were injected with 0.2 micro cc of saline solution containing 1 mg of aflatoxin dissolved in 1 cc of saline. The treated flies were exposed to