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Recently (Hubby and Throckmorton, 1965) a study has been made, using vertical acrylamide electrophoresis, of the evolutionary relationships between adult soluble proteins within the virilis group of *Drosophila*. Presented here is an inter-species

survey of late third-instar larval lymph protein content, as shown by starch-gel electrophoresis, of the following seven species of the genus *Drosophila*: *melanogaster*, *simulans*, *immigrans*, *hydei*, *virilis*, *funnebris*, and *subobscura*. In addition, pooled samples of *melanogaster*, *funnebris* and *subobscura* species caught in the wild in various parts of Ireland, were examined for lymph protein content.

The results of the survey are shown in fig. 1. As expected, *melanogaster* and *simulans* had very similar patterns, with the complete absence of fraction 5 in the latter, constituting the major difference between the two. This protein band was found to be present in pooled samples of all twenty-six laboratory and wild stocks of *melanogaster* studied. It is possible that fractions 13 and 14 in *melanogaster* are very similar to fractions 3 and 13 in *simulans* (fig. 1) except that the electrophoretic mobilities are slightly altered. The protein patterns obtained for the other five species varied quite significantly from the above two species and from each other. *Drosophila immigrans* exhibited the least number of fractions having only two main bands not counting the "front" (fig. 1). The other four species all exhibited approximately six to nine fractions of differing concentration and composition.

The protein patterns exhibited by cultures of *melanogaster*, *funnebris*, and *subobscura* started from individuals caught in different parts of Ireland, were in close agreement with those of the laboratory bred stocks, and only minor differences were observed. In the case of *funnebris* for instance, protein fraction C of the laboratory stock (see fig. 1) was found to be split into two in the wild stocks. A survey of individual larvae has shown that the wild stock contained a mixture of individuals having the single and double band phenotypes.

The degree of difference and similarity between the larval lymph protein patterns of the different species can be correlated, to some extent, with their chromosomal arrangement. Those species with a primitive six chromosome pair configuration differ significantly in lymph protein pattern from *immigrans*, *simulans* and *melanogaster* which have four pairs. *D. simulans* and *D. melanogaster* are more evolved than *D. immigrans* and exhibit much more diversification of protein fractions. There is, however, quite a close resemblance between the pattern obtained from the Pacific 7 strain of *D. melanogaster* (see Duke, 1965) and that of *D. immigrans*. Four protein fractions of the *D. immigrans* pattern have similar electrophoretic mobility to fractions exhibited by *D. melanogaster* (fig. 1). By comparison, the lymph protein patterns from the four species containing six chromosome pairs are significantly different from those already described. *D. subobscura*, although having only two protein bands of exactly similar mobility to *melanogaster* fractions, has four others of very close mobility (see fig. 1). Four fractions in the lymph of *D. hydei* resemble those of *D. melanogaster* in electrophoretic mobilities. It may be significant that *D. hydei* alone of these four species contains the more evolved V-shaped chromosome. The patterns of *D. virilis* and *D. funnebris* are least like that of *D. melanogaster*. Generally speaking, therefore, the patterns of lymph protein content of the seven species studied, corresponded with the evolutionary trends within the genus as indicated by their characteristic chromosome configurations. These electrophoretic data should be firstly considered as an extra taxonomic aid in the systematics of *Drosophila*, and secondly as a basis for future work in a more refined biochemical approach to problems of evolution.

- References: Duke, E. J. (1965). Further studies on the inheritance of lymph proteins in *Drosophila*. *Gen. Res. Camb.* (in press).  
Hubby, J. L. and Throckmorton, L. H. 1965. Protein differences in *Drosophila*, II. Comparative species genetics and evolutionary problems. *Gen.* 52:203-215

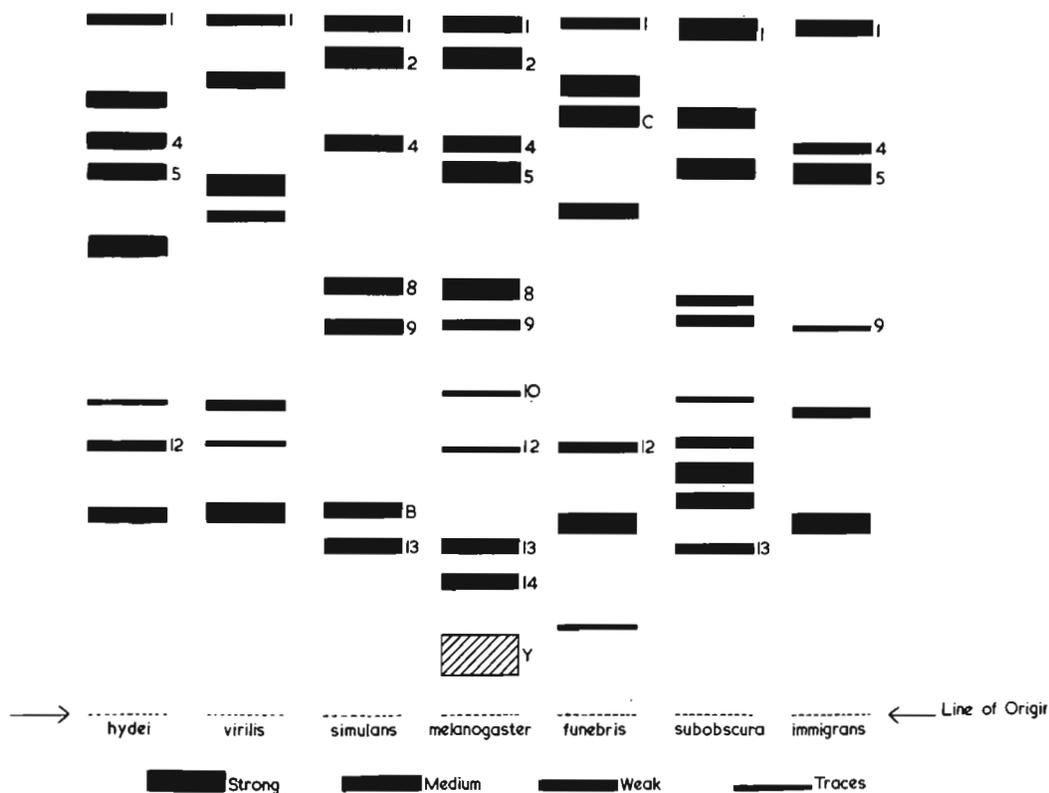


Fig. 1

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 The distribution of genetic potential for wing venation abnormalities in a natural population of *D. pseudoobscura*.

period beginning with the  $F_3$ . A variety of defects were detected, including missing posterior crossvein (cve), missing longitudinal vein (lv) and extra venation (ev). The potential for producing these defects is not restricted to only a few strains but is widespread throughout the population.

Egg samples from 25 strains derived from single females captured at Pinon Flats, Mount San Jacinto, California, were placed at 16°C and at 25°C for development. In each strain more than 1000 flies were sampled over a five generation

Table 1: Distribution of wing venation defects

<u>No. of Strains</u>	<u>lv</u>	<u>cve</u>	<u>ev</u>
6	+	+	+
1	+	+	0
6	0	+	+
3	+	0	+
0	+	0	0
1	0	+	0
7	0	0	+
1	0	0	0