

Supplementation of 10% calf serum was found to be the best among sera from various sources and at various concentrations examined.

Effects of silkworm blood on the growth of embryonic cells were also examined in the presence of 10% calf serum. The results are shown in Table 2.

Table 2. Effects of silkworm blood on the growth of *Drosophila* embryonic tissues in culture.

Source of blood	No. of explants tested	No. of explants in which growth was observed	Percent growth
Control	16	12	75
5th instar larvae, 5%	15	3	20
5th instar larvae, 10%	22	7	32
pupae, 10%	27	3	11

The addition of heat-treated blood collected from fifth instar larvae or pupae of silkworm exhibited no growth improvement in the culture of embryonic *Drosophila* tissues.

References: 1. Kuroda, Y., 1969 Japan. J. Genetics 44, Suppl. 1: 42; 2. Kuroda, Y., 1970 Exp. Cell Res. 59: 429.

David, J. and J. Bouletreau-Merle. University of Lyon, France. Two levels of egg retention in the genital tract of *Drosophila* females.

In *Drosophila*, egg production is controlled by vitellogenic activity, frequency of egg chamber resorptions and retentions. From the study of the relationship between daily egg production and frequency of females with an egg in the uterus, it is concluded that retentions may be

initiated at two levels of the female reproductive system.

The results obtained for normally fed, mated or virgin females are plotted on the figure. In spite of a rather significant variability, it appears that the points correspond to an increasing convex curve. So the time spent by each egg in the uterus is quite stable and the variations in fecundity correspond to variations of the duration of absence of intra-uterine egg. Of particular interest is the fact that results for virgin females appear to be

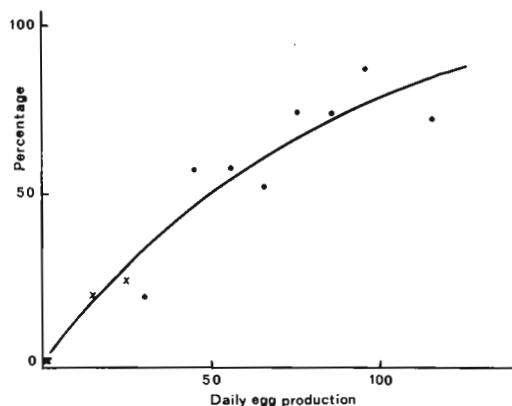


Figure: Relation between daily egg production and percentage of females with an egg in uterus (●: mated females; x: virgin females)

distributed on the same curve as those from mated females. From other studies (Merle and David, 1967) it is known that many stage 14 oocytes are observed in the ovarioles of virgin females. Therefore, this retention takes place in the ovarioles, not in the uterus.

Intra-uterine retention is observed when females are not offered a suitable site for egg laying. For example, properly fed, mated females were given only an agar medium for oviposition. In these conditions, the mean daily fecundity was only 2 eggs, but more than 80% of the females contained an egg in their uterus (data from Mrs. Van Herreweghe, 1970). Of course, in such eggs, embryonic development begins and cases of viviparity are observed (see also King, 1963).

In conclusion, retentions may be initiated at two levels. 1) In the uterus, when the conditions for egg laying are unsuitable. This retention involves the oviposition behavior and is probably controlled by the central nervous system.

2) In the ovarioles, when females are not mated. This retention corresponds probably to a lack of ovulation; that is, an absence of contraction of the wall of the ovarioles. Its determinism is not yet known.

References: King, R.C., 1963 D.I.S. 38: 96; Merle, J. and J. David, 1967 C.R. Acad. Sci. Paris 234: 2028-2030; Van Herreweghe, J., 1970 C.R. Acad. Sci. Paris 271: 108-110.