

Whittinghill, Maurice An aid in arranging flies for separating or counting them.

a line, which may be easily formed as follows. The flies are emptied from the etherizing bottle upon a card which has along its middle a flexible fold by which the card may be bent to form a trough. A few strokes of the brush distribute the pile of flies evenly along this groove. The card is now held flat with one hand while the other goes down the line casting each fly quickly to one side or the other with a brush handle or needle, thus sorting into two groups. If further classification is necessary, these two groups may be swept to opposite ends of the card, or to different cards, and each group easily arranged for the next sorting. After such a sorting each of the groups is found to lie in approximately linear order, which makes counting simple. Finally the card may be bent again to make a chute for dispensing the flies accurately into the morgue.

To facilitate sorting and counting flies it is advantageous to have them arranged approximately in

Stern, Curt Technique for the study of certain genetic constitutions in hypodermis spots.

based primarily on the occurrence of somatic segregation. A heterozygous cell  $Aa$  segregates into two cells  $AA$  and  $aa$ . If  $aa$  represents the genetic constitution to be tested and if the cell  $aa$  is viable and divides, a spot will appear. Often the genetic constitution to be tested in spots leads to inviability of the segregate thus not resulting in a visible aberrant area. The frequency of somatic segregation is rather variable so that conclusions as to inviability of certain constitutions can be drawn with reservation only, considering the possibility of absence of segregation. However, the following method furnishes reliable controls. Let  $a$  be the gene to be tested and  $b$  and  $c$  genes in the same linkage group effecting hypodermal characteristics in small spots. By mating flies of the constitution  $abC/ABc$  are produced. Somatic segregation will lead to  $abC/abC$  and  $ABc/ABc$  cells. The latter, known to be viable, will be able to produce a spot; the former, if viable will appear in direct contact with the  $ABc/ABc$  spot as an  $abC/abC$  twin-spot. If  $aa$  leads to inviability no twin spot will be formed. In case of sex-linked genes yellow ( $y$ ) may be used for  $b$  and singed<sup>3</sup> ( $sn^3$ ) for  $c$ , so that either only single  $sn^3$  spots or  $y$  next to  $sn^3$  twin-spots will be found (both  $y$  and  $sn^3$  being recognizable as characteristics of even-single setae). Somatic segregation is caused by four strand crossing-over. If it occurs to the right of all loci studied, the foregoing holds true completely except for developmental reasons which may make the mosaic areas so small as not to cover at least one seta per single spot. Somatic crossing-over between the genes studied will not lead to twin spots. However, the frequency of somatic crossing-over at the spindle fibre region is high enough to produce a sufficient number of potential twin

One tool in studies on gene action is the production of mosaic spots. Their appearance is