

Takamura, T. & Y. Fuyama 1980, *Behav. Genet.* 10:105-120; Weisbrot, R.D. 1966, *Genetics* 53:427-435.

Sondergaard, L. University of Copenhagen, Denmark. Mating capacity of e/e and e/+ males under non-competitive conditions.

It is well-known that the mutant ebony (e) has several pleiotropic behavioural effects. Some of these have been thought to be the reason why the e gene, in contrast to most other mutant genes, stabilizes at a certain level in population cage experiments. One factor which is rarely considered is what one might call the "Don-Juan" factor, i.e., the number of females a male can mate within a given period. A male with a very efficient courtship could be at a selective disadvantage if he needs too long a recovery period after copulation compared to a male with a less effective courtship, but with a very short recovery period. To test the mating capacity, single unexperienced  $\sigma\sigma$  (12-24 hrs of age) were confined for 24 hrs with 12 one-week-old  $\text{♀♀}$  in light or complete darkness. e/e, e/+ and +/+  $\sigma\sigma$  were mated with e/e; e/+ and +/+  $\text{♀♀}$  also to test the effect of the female genotype on male performance. Results are shown in Table 1. The overall tendency is that e/+ and +/+  $\sigma\sigma$  perform better in light, whereas e/e  $\sigma\sigma$  perform equally well in light and darkness when mated to e/+ and e/e  $\text{♀♀}$ . In the light the order of the D.J. factor is e/+ > e/e > +/+, indicating overdominance for this trait.

Table 1. D.J. factor  $\pm$  s.d. (see text) for males confined for 24 hrs with 12 of the indicated genotype; experiments were performed in 24 hrs light and 24 hrs of darkness. In each experiment 75-100 were tested individually.

	darkness	light
+/+ $\text{♀}$ x e/+ $\sigma$	3.3 $\pm$ 1.5	4.8 $\pm$ 2.0
e/e $\text{♀}$ x e/+ $\sigma$	3.6 $\pm$ 1.7	5.9 $\pm$ 2.0
e/+ $\text{♀}$ x e/+ $\sigma$	3.4 $\pm$ 1.9	5.7 $\pm$ 2.4
+/+ $\text{♀}$ x e/e $\sigma$	2.9 $\pm$ 1.3	3.7 $\pm$ 1.9
e/e $\text{♀}$ x e/e $\sigma$	4.3 $\pm$ 2.1	4.5 $\pm$ 2.0
e/+ $\text{♀}$ x e/e $\sigma$	4.3 $\pm$ 1.9	4.4 $\pm$ 2.1
+/+ $\text{♀}$ x +/+ $\sigma$	1.8 $\pm$ 1.1	2.4 $\pm$ 1.6
e/e $\text{♀}$ x +/+ $\sigma$	2.5 $\pm$ 1.3	3.1 $\pm$ 1.5
e/+ $\text{♀}$ x +/+ $\sigma$	1.7 $\pm$ 1.5	3.2 $\pm$ 1.5

These observations are explainable by the fact that e/e flies are blind and that e/+ and e/e have a more efficient courtship behaviour (Kyriacou et al. 1978). However, this does not explain the observed differences between different females when tested to the same male genotype: in the light the scores are lower with +/+  $\text{♀♀}$ . In darkness the results are more complex: no differences were observed between  $\text{♀♀}$  mated to e/+  $\sigma\sigma$ ; e/e  $\sigma\sigma$  show lower scores with +/+  $\text{♀♀}$ ; +/+  $\sigma\sigma$  have a higher mating frequency with e/e  $\text{♀♀}$ . These differences could be explained by differences in female heat. A more possible explanation is a difference in the activity levels of both males and females. That is, increasing spontaneous activity in the order +/+; e/+; e/e. In the light +/+  $\text{♀♀}$  do not move around as much and therefore rarely meet a male; in the darkness they move around even less. However, with e/+ males this is compensated for by the higher activity of these males also in the dark. In the experiment with +/+  $\sigma\sigma$  sluggishness is only compensated for by the high activity of the e/e  $\text{♀♀}$  in darkness. In the dark the high activity of e/e  $\sigma\sigma$  compensates for differences between e/e  $\text{♀♀}$  and e/+  $\text{♀♀}$  activity.

Reference: Kyriacou, C.P., B. Burnet & K.J. Connolly 1978, *Anim. Behav.* 26:1195.

Spieß, E.B. University of Illinois, Chicago, Illinois. Discrete generation populations of *D. persimilis* selected for female receptivity and frequencies of KL-MD karyotypes.

Population box experiments were designed in 1979 with selection for early maturation of females (*D. persimilis*) in order to substantiate the relative frequency changes expected of KL and MD arrangements that had been characterized for female "switch-on" of receptivity by Yu & Spieß (1978). Three strains of KL (4,11,17) with amylase variant amy-1.09 and 3 strains of MD (7,16,35) with amy-1.00 derived from a McDonald Ranch, CA, population were intercrossed within homokaryotypes and introduced into plastic refrigeration boxes ("Bennett cages") with 8 holes for as many food vials to provide oviposition area for 200 initial pairs of flies. Females were virgins of 1-2 days past eclosion while males were as old or slightly older. Initial frequencies were approximately 90%: 10% of either arrangement and four populations were monitored by electrophoresing a sample of

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