

Table 4. Species % composition by community.

| Species | Lowland forest | Upland forest | Sand Prairie |
|-----------------------|----------------|---------------|--------------|
| <i>D.affinis</i> | 56.3 | 48.5 | 56.1 |
| <i>D.algonquin</i> | 0.7 | 0.0 | 7.0 |
| <i>D.athabasca</i> | 0.4 | 0.0 | 0.0 |
| Undetermined | | | |
| aff.-alg.-ath.?? | 13.3 | 0.0 | 7.5 |
| <i>D.falleni</i> | 9.7 | 26.6 | 14.4 |
| <i>D.tripunctata</i> | 9.4 | 0.0 | 0.0 |
| <i>D.robusta</i> | 5.0 | 17.8 | 1.1 |
| <i>D.putrida</i> | 3.5 | 6.6 | 11.8 |
| <i>D.buskii</i> | 0.5 | 0.0 | 0.5 |
| <i>D.quinaria</i> | 0.4 | 0.0 | 1.6 |
| <i>D.duncani</i> | 0.2 | 0.0 | 0.0 |
| <i>D.testeca</i> | 0.2 | 0.0 | 0.0 |
| <i>D.immigrans</i> | 0.1 | 0.0 | 0.0 |
| <i>D.victoria</i> | 0.1 | 0.0 | 0.0 |
| <i>D.melanogaster</i> | 0.1 | 0.5 | 0.0 |
| Total collected | 5657 | 608 | 187 |

allow a more accurate comparison of the species compositions of the two forest communities.

Few members of the species commonly associated with humans (*D.buskii*, *D.immigrans* and *D.melanogaster*) were collected. We conclude that our samples represent natural *Drosophila* populations and not human-associated ones.

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Reference: Heim, W.G. 1978, DIS 53:216.

Silva, F.J. Universidad de Valencia, Espana. Partial inhibition of the effect of the mutant red malpighian tubules (red) by other eye colour mutations of *Drosophila melanogaster*.

A study of eye pigments and related metabolites in adult flies (9 days after emergence) of ten strains of double mutants of *D.melanogaster* has been carried out. All the strains carry two eye colour mutations, one of them being "red."

The separation of eye pigments and related metabolites in these strains was carried out

by means of two-dimensional thin-layer chromatography (TLC) on cellulose plates, using as extraction solvent methanol-acetic acid-water (4:1:5 by vol) and as elution solvent isopropanol-2%-ammonium acetate (1:1, v/v) for first dimension (3 hr) and 3% aqueous ammonium chloride for the second one (50 min). Quantification was made by measuring the fluorescence directly on the plate with a fluorescence spectrophotometer (PERKIN ELMER MPF 44B). The results of the ten double mutants strains are compared with the patterns of the single mutants (Ferre et al. 1983) in Table 1.

Although the effect of mutant "red" consists in a strong reduction of pteridines (except bipterin), some double mutants such as cn red, rb red, and cm red and to a lesser extent ltd red, cl red and v red, present a significantly higher quantity of drospterins, PDA and sepiapterin and a lower quantity of bipterin compared to the mutation red alone. This reduction may be interpreted as a partial inhibition of the effect of red mutant. Especially interesting is the case of cn red strain, since the gene *cn⁺* is known to be the structural gene of the enzyme Kynurenine hydroxylase and thus to affect only ommochrome synthesis.

The malpighian tubules of the wild type presents a light yellow colour produced by the accumulated 3-OH-kynurenine; however, the malpighian tubules of the mutant "red" are red-coloured (Oster 1954), due to the conversion of the accumulated 3-OH-kynurenine into ommochromes, of which a small quantity is xanthomatin and a larger quantity is ommin (Wessing & Bonse 1966). In addition this mutant, in the eyes, accumulates 31% of brown pigment (H_2 -xanthomatin) and around 5% of drospterins compared to the wild type (Ferre et al. 1983). Transport defects in malpighian tubules are the basis for the anomaly in some eye colour mutants of *D.melanogaster* that have reduced amounts of both pteridines and ommochrome (Sullivan et al. 1975; Howells et al. 1977; Sullivan et al. 1979). This fact strongly suggests that the gene *red⁺* is acting on the transport of pigments precursors, being unable to transport precursors of pteridines and ommochromes efficiently. For reasons not known it seems that certain mutations as cn, rb, cm, ltd, cl and v produce a partial inhibition of

Table I. Percentages of eye-pigments and related metabolites (Or-R has arbitrarily received the values of 100). NDP(Neodrosopterin), DP(Drosopterin), IDP(Isodrosopterin), ADP(Aurodrosopterin), PDA (2-amino-4-oxo-6-acetyl-7,8-dihydro-3H,9H-pyrimido [4,5-b] [1-4] diazepine), SP(Sepiapterin), Spot 7(unidentified pteridine), AHP(Pterin), BP(Eiopterin), IXP(Isoxanthopterine), XTC (Xanthurenic acid) and H₂-XTM (Dihydroxanthomatin). ND (not detected).

| strain | NDP | DP | IDP | ADP | PDA | SP | Spot 7 | AHP | BP | IXP | XTC | H ₂ -XTM |
|---------|-------|-------|-------|-------|--------|----------|--------|--------|---------|--------|--------|---------------------|
| red | 6±2 | 4±2 | 4±2 | 9±2 | 79±4 | 43±10 | 66±10 | 24±5 | 301±30 | 51±5 | 20±7 | 31±1 |
| cn red | 70±14 | 36±5 | 39±7 | 80±5 | 175±33 | 84±14 | 69±18 | 52±2 | 174±46 | 30±2 | ND | -- |
| rb red | 41±10 | 16±2 | 9±1 | 32±5 | 145±26 | 128±30 | 101±9 | 22±2 | 323±42 | 26±3 | 37±4 | -- |
| cm red | 49±10 | 25±2 | 21±1 | 59±3 | 255±4 | 161±7 | 98±9 | 35±1 | 232±15 | 24±1 | ND | -- |
| v red | 12±1 | 6±2 | 7±2 | 35±2 | 169±27 | 47±5 | 61±8 | 29±3 | 122±3 | 23±2 | ND | -- |
| ltd red | 13±2 | 9±1 | 9±1 | 40±6 | 160±14 | 89±12 | 76±5 | 23±4 | 153±3 | 17±2 | ND | -- |
| cl red | 23±2 | 21±1 | 19±4 | 20±4 | ND | 1750±750 | 507±99 | 492±34 | 340±110 | 73±4 | ND | -- |
| pr red | 7±2 | 5±0 | 2±1 | 7±2 | 20±2 | 19±5 | 30±5 | 22±1 | 128±15 | 38±2 | 16±3 | -- |
| pn red | 5±2 | 3±0 | 2±1 | 4±2 | TRACE | 37±10 | 46±6 | 16±0 | 162±23 | 50±3 | ND | -- |
| dke red | 4±1 | 5±1 | 6±1 | 12±1 | 104±1 | 14±4 | 71±2 | 21±1 | 215±20 | 22±1 | 24±5 | -- |
| cho red | TRACE | TRACE | TRACE | TRACE | 33±4 | 39±1 | 61±3 | 25±2 | 308±28 | 35±1 | ND | -- |
| cn | 91±4 | 86±8 | 89±2 | 91±4 | 84±16 | 101±14 | 80±7 | 95±13 | 82±6 | 63±5 | ND | 4±0 |
| rb | 34±2 | 25±3 | 36±3 | 55±3 | 94±12 | 162±7 | 50±5 | 37±4 | 149±14 | 29±2 | 7±2 | 38±1 |
| cm | 22±4 | 19±2 | 21±2 | 51±6 | 202±21 | 199±15 | 66±4 | 34±6 | 182±30 | 20±6 | 6±3 | 43±1 |
| v | 93±2 | 80±4 | 79±2 | 93±4 | 108±11 | 146±9 | 100±9 | 102±6 | 103±6 | 107±4 | ND | 3±0 |
| ltd | 32±1 | 26±3 | 25±4 | 58±2 | 203±8 | 140±23 | 114±6 | 26±7 | 165±31 | 30±8 | ND | 8±1 |
| cl | 19±1 | 54±1 | 47±1 | 8±3 | ND | 1442±46 | 474±35 | 526±40 | 261±19 | 117±5 | 59±3 | 160±2 |
| pr | 45±3 | 33±9 | 48±11 | 33±3 | 35±4 | 64±6 | 58±11 | 39±10 | 105±17 | 109±10 | 22±7 | 122±0 |
| pn | 22±4 | 22±2 | 32±4 | 18±2 | 87±17 | 161±14 | 95±6 | 34±2 | 140±5 | 88±5 | 43±6 | 81±2 |
| dke | 20±3 | 51±13 | 55±17 | 86±9 | 109±15 | 128±34 | 61±4 | 38±4 | 119±7 | 66±10 | 108±15 | 153±1 |
| cho | 4±1 | 4±1 | 4±1 | 11±2 | 47±3 | 108±9 | 125±2 | 47±11 | 268±43 | 58±9 | TRACE | 66±1 |

this effect, causing these flies to recover the ability to transport and accumulate this substances in the eyes.

References: Ferre, J., F. J. Silva, M. D. Real & J. L. Mensua 1983, Chemistry and Biology of the Pteridines, de Gruyter: 669-673; Howells, A. J., K. M. Summers & R. L. Ryall 1977, Biochem. Genet. 15: 1040-1059; Oster, I. I. 1954, Dis 28: 77; Sullivan, D. T. & M. C. Sullivan 1975, Biochem. Genet. 13: 603-613; Sullivan, D. T., L. A. Bell, D. R. Paton & M. C. Sullivan 1979, Biochem. Genet. 17: 565-573; Wessing, A. & A. Bonse 1966, Z. Naturforsch. 21b: 1219-1223.