

Air drying technique for cytogenetic studies in freshwater planarians (Platyhelminthes)

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Cytogenetic studies in seawater or land planarians have been generally performed by means of the squash technique (BALL and DE VRIES 1983; CURINI-GALLETTI *et al.*, 1985, 1989a, b; MARTENS *et al.* 1989a, b; OKI *et al.* 1991; PUCCINELLI and CURINI-GALLETTI 1987; TAMURA *et al.* 1991). However, the quality of chromosome preparations obtained by squashing in terms of spreading and detritus background is not as good as that obtained by using the air drying technique in other taxa. In addition, few information is available about citogenetic techniques for freshwater invertebrates.

The air drying technique (ROTHFELDS and SIMINOVITCH 1958) was proposed for cytogenetic analysis of mammalian bone marrow cells and it included previous tissue disaggregation. For nearly forty years the technique became widely used, with several modifications, for cytogenetic studies even in plant cells.

This paper describes the modifications made in the technique for the analysis of planarians chromosomes.

Specimens were collected in Arroyo Zapata, a small water course in the Province of Buenos Aires, Argentina during the spring season.

Whole living animals — three or four each time — were washed in physiological solution NaCl 0.6% (GABE 1968) and then transferred to a clock glass with a solution of colchicine (0.3%) dissolved in Gabe's solution. The animals were cut with a scalpel under a magnifying glass until complete disaggregation. The disaggregated tissues were transferred to a centrifuge tube. After resuspending with a Pasteur pipette the cells were left into the tube for three hours. The material was centrifuged at 800 r.p.m. for 5 min and the supernatant was replaced with hypotonic solution (KCl 0.075 M). After incubation for 30 min at 20°C the hypotonic solution was replaced by fixative (methanol-acetic acid 3:1) and the tubes were stored at 4°C overnight. After

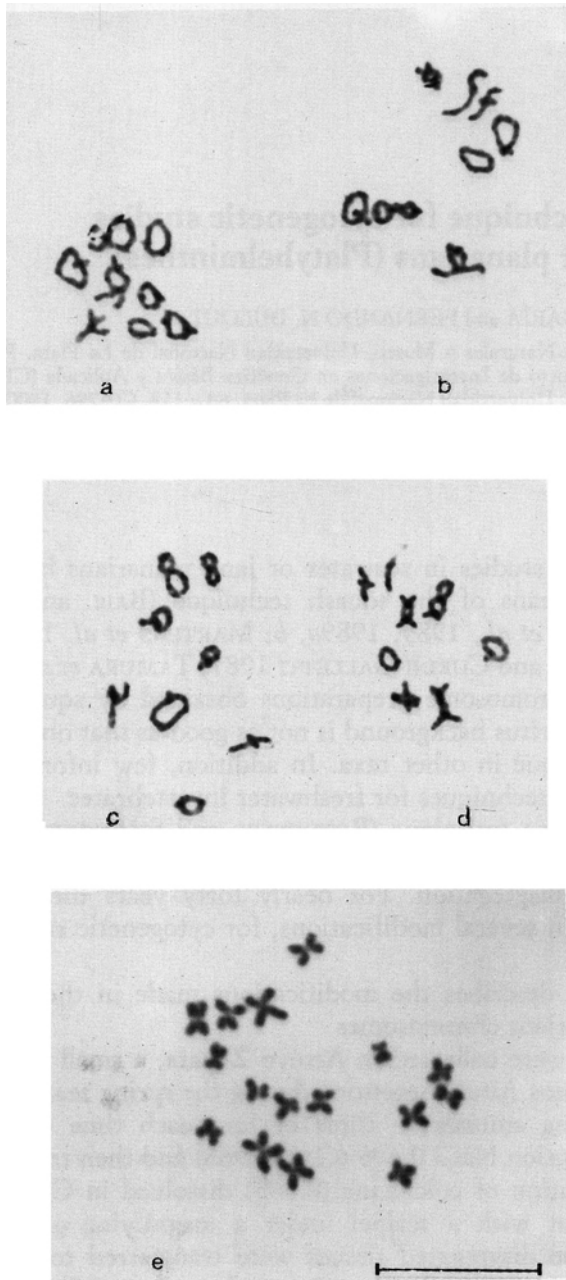


Fig. 1. — Micrographs of different plates: a. b. c. d. diacinesis — metaphase I; e. Mitotic metaphase. The bar indicates 10 μm.

one or two changes of fixative, chromosome preparations were made by dropping the cell suspension on frozen slides. Preparations were stained with Giemsa 5% for 2 minutes.

Using the whole specimens good plates of mitotic and meiotic cells in the same slide were obtained. As it can be seen in Fig. 1, chromosomes are well spread and the slide has not any detritus or cytoplasmic background. Consequently, morphological analysis of the chromosome complement is very easy and fast. On the other hand, the technique is very simple and reliable and can be used in other invertebrates.

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