

en the small increase in yield strength caused by dislocation locking might be getting offset by a decrease in strength caused by dynamic recovery.

Since stress relaxation in the plastic strain range occurs by a process of thermally activated motion of dislocations across the barriers,^{12,13} it is possible to calculate one of the activation parameters, viz., the activation volume from the relaxation curves. Activation volume, the product of the length of the dislocation getting activated, the Burgers vector and the width of the barrier, characterises the obstacle to dislocation motion. From the relation between creep and low temperature relaxation, it has been shown^{13,14} that the slope 'S' of the curve $\sigma_0 - \sigma$ versus $\ln t$ is given by

$$S = kT/v \quad (2)$$

where 'v' is the activation volume, k the Boltzmann's constant and T the absolute temperature. The slopes of the straight lines corresponding to 3.9 and 4.8 kg./mm.² are utilised to calculate the activation volume from equation (2). The estimated activation volume ($\approx 2.5 \times 10^{-21}$ cm.³) is in good agreement with that reported earlier from creep experiments.¹ While the estimated activation volume suggests that either intersection of glide and forest dislocations or the non-conservative motion of jogs is the rate-controlling mechanism in Indian commercial aluminum, our earlier results based on the combined data of creep and tensile testing indicated that the non-conservative motion of jogs producing point defects might be the most probable one.

CONCLUSIONS

Strain ageing experiments on Indian commercial aluminum showed the occurrence of sharp yield point associated with an yield drop. The increase or decrease in the yield strength after ageing depended on the stress level. The analysis of the relaxation curves yielded a value of 2.5×10^{-21} cm.³ for the activation volume.

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1. Prasad, Y. V. R. K., Sastry, D. H. and Vasu, K. I., To be published in *Trans. Ind. Inst. Metals*.
2. —, — and —, communicated to *J. Ind. Inst. Science*.
3. Mitra, S. K., Osbrone, P. W. and Dorn, J. E., *Trans. AIME*, 1961, **221**, 1206.
4. — and Dorn, J. E., *Ibid.*, 1963, **227**, 1015.
5. Nunes, A. C., Rosen, A. and Dorn, J. E., *Trans. ASM*, 1965, **58**, 38.
6. Basinski, Z. S., *Phil. Mag.*, 1959, **4**, 393.
7. Cottrell, A. H., *Dislocations and Plastic Flow in Crystals*, Oxford, London, 1953, p. 139.
8. —, *Trans. AIME*, 1958, **212**, 192.
9. Johnston, W. G., *J. Appl. Phys.*, 1962, **10**, 727.
10. Hahn, G. T., *Acta Met.*, 1962, **10**, 727.
11. Cottrell, A. H., *Conf. on the Relation between the Structure and Mechanical Properties of Metals*, H.M.S.O., 1963, **2**, 456.
12. Feltham, P., *J. Inst. Metals*, 1960-61, **89**, 210.
13. —, *Phil. Mag.* 1961, **6**, 259.
14. Sargent, G. A., *Acta Met.*, 1965, **13**, 663.

TEST-TUBE FERTILIZATION IN DICRANOSTIGMA FRANCHETIANUM (PRAIN) FEDDE

N. S. RANGASWAMY AND K. R. SHIVANNA

Department of Botany, University of Delhi, Delhi-7

THE technique of test-tube fertilization devised by Kanta *et al.*¹ helps to eliminate the path of pollen tubes through the stigma and style. It is therefore promising in studies on plant breeding and genetics, and has been thus far applied to a few systems.²⁻⁵ This paper reports our successful application of the technique to *Dicranostigma franchetianum* (Papaveraceae). Plants of this species were raised from seeds obtained from the Direktor, Institute für Kulturrepflanzforschung, Gatersleben, East Germany.

As a prerequisite to our work, anthesis, dehiscence of anthers, pollination, fertilization, and seed development were studied from fresh material and from material fixed in formalin-acetic-alcohol (40% formaldehyde solution 5 ml + glacial acetic acid 5 ml + 70% ethyl alcohol 90 ml). The fixed material was embedded in paraffin following the customary method and microtomed (10-15 μ). The sections were stained in iron hæmatoxylin and erythrosin, and mounted in Canada balsam. Pollen germination and pollen tube growth

were studied also from whole mounts and free-hand sections prepared in 1% iron acetocarmine.

Under the climate of Delhi *Dicranostigma franchetianum* flowers during February-April. Anthesis occurs between 8 and 9 a.m. Anthers begin to dehisce toward the evening the day

before anthesis and continue to shed pollen until two hours after anthesis by which time pollination is also accomplished. Pollen germination occurs nearly 30 minutes after pollination and fertilization 24-36 hours thereafter. Six days after pollination a 2- to 4-celled

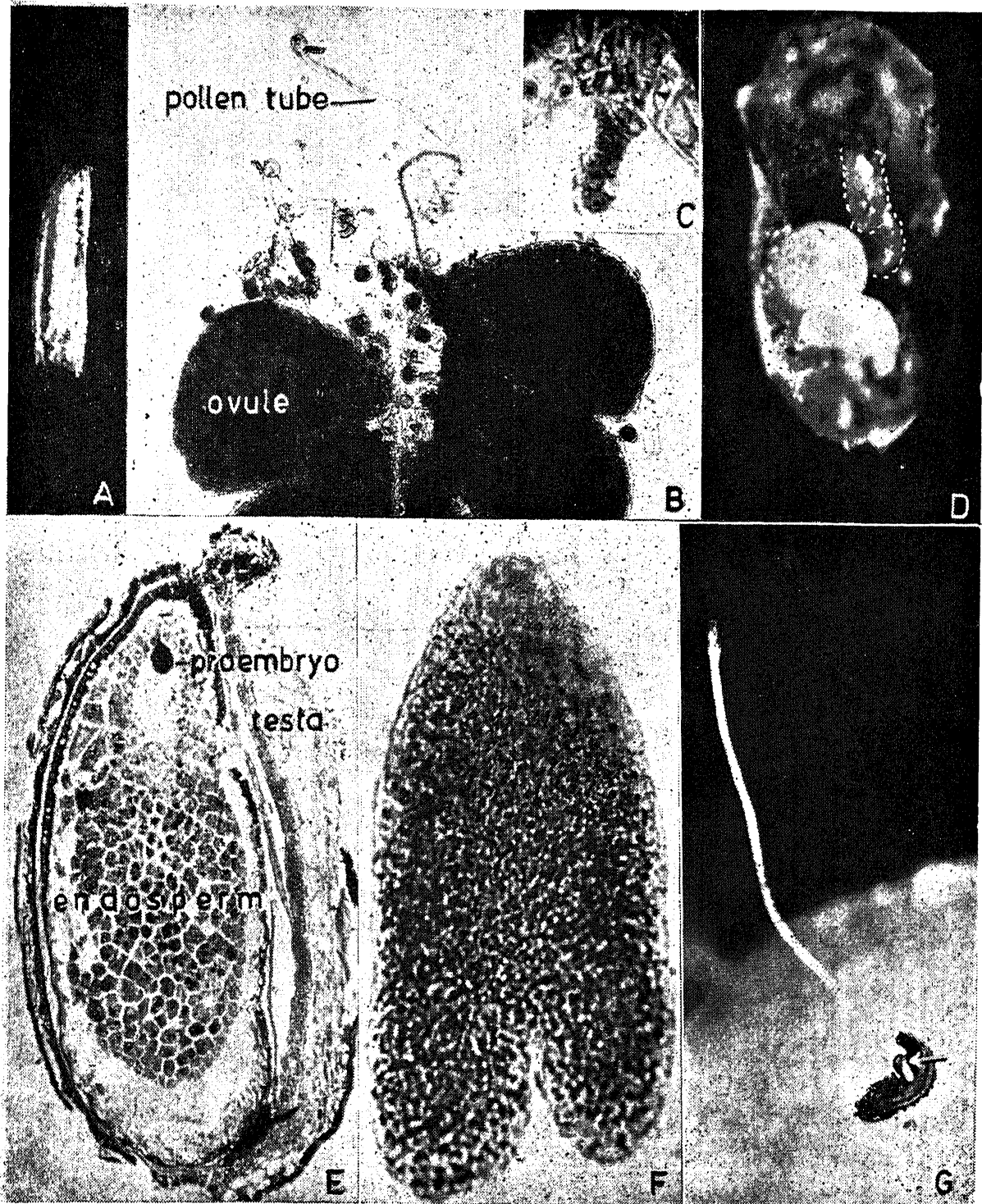


FIG. 1, A-G. Test tube fertilization in *Dicranostigma franchetianum*. A. An explant showing ovary wall, placenta, and ovules, $\times 6$. B. Whole mount of a few ovules removed from a culture 24 hours after pollination; note pollen germination, $\times 121$. C. Longisection of micropylar part of a fertilized ovule from a culture 4 days after pollination. The filamentous proembryo is obvious, $\times 102$. D. 7-day-old pollinated culture showing 4 developing seeds; the 2 that are in profile are highlighted by broken lines; unfertilized ovules are not in focus, $\times 22$. E. Longisection of a young seed collected from a culture 7 days after pollination. Globular embryo and massive endosperm are evident, $\times 110$. F. Whole mount of embryo dissected from seed obtained from 18-day-old pollinated culture, $\times 345$. G. 21-day-old pollinated culture showing seed germination *in situ*. In this view, the seedling does not show the root; the arrow-marked is a recently germinated seed which shows only the emergence of the radicle, $\times 4$.