AN UNDESCRIBED SPECIES OF PHOMOPSIS AND ITS ASSOCIATED ASCIGEROUS STATE

V. G. RAO and B. R. DAYAKAR YADAV

Maharashtra Association for the Cultivation of Science, Research Institute, Pune 411004, India.

DURING the course of investigation into the fungal diseases of ornamental plants, an interesting leaf-spot disease was encountered by the authors on Aregelia spectabilis Mez. incited by a species of the form-genus Phomopsis (Sacc.) Sacc. The same spots later showed the development of a perithecial state which was identified as a species of Diaporthe nitschke¹. Since these two fungi were noted for the first time on one and the same host, a detailed study was undertaken in respect of their morphology, taxonomy and identification. A review of literature^{1,2} also showed no report of these two fungi on Aregelia spectabilis Mez. or its related hosts of the family Bromeliaceae so far. Hence, these two fungi have been described here as new taxa based on comparative morphology and hostrelationship² as under.

Phomopsis spectabilae sp. nov. (figure 1)

Pycnidia on leaves chiefly solitary, immersed, dark brown, sub-spherical to irregular, upto 151 μ m in diam; opening by an apical ostiole; pycnidial cavity being unilocular; conidiophores small, hyaline, simple, lining inside wall of the cavity; conidia of two types; alpha conidia (phialospores) hyaline, uni-celled, fusiform to elliptical, guttulate, $5.5-6\times2-2.7~\mu$ m, beta conidia (phialospores) hyaline, elongated, filiform, curved, measure $13.5-19.6\times2~\mu$ m.

Maculae saepe irregulares et oblonga, ad terminibus, coalitus, melleus ad medio, et avellaneus marginum; pycnidia punctiformae, hypophyblae, solitarieae, immersa, fuscaburnneae, sub-sphaerica vel irregulares, usque 151 μ m diam., ostiolata, unilocularia; pycnidiosporae alphae et betae; alpha (A) sporae hyalina, unicellularia, fusiformia vel elliptica, guttulata, magnit $5.5-6\times2-2.7~\mu$ m; beta (B) conidiae hyalina, elongata, filiformia, curvata, magnit; $13.5-19.6\times2~\mu$ m.

Diaporthe spectabilue sp. nov. (figure 2)

Ascostromata (ascomata) scattered, botryose, black with long neck, upto 300 μ m diam., narrowly elongated neck and opening by a narrow ostiole; asci 8-spored, fusiform, sessile, bitunicate, evanascent, measure $41.6-45\times8~\mu$ m; ascospores 1-septate, guttulate, acute at ends, measure $9.9-11.6\times3.3~\mu$ m.

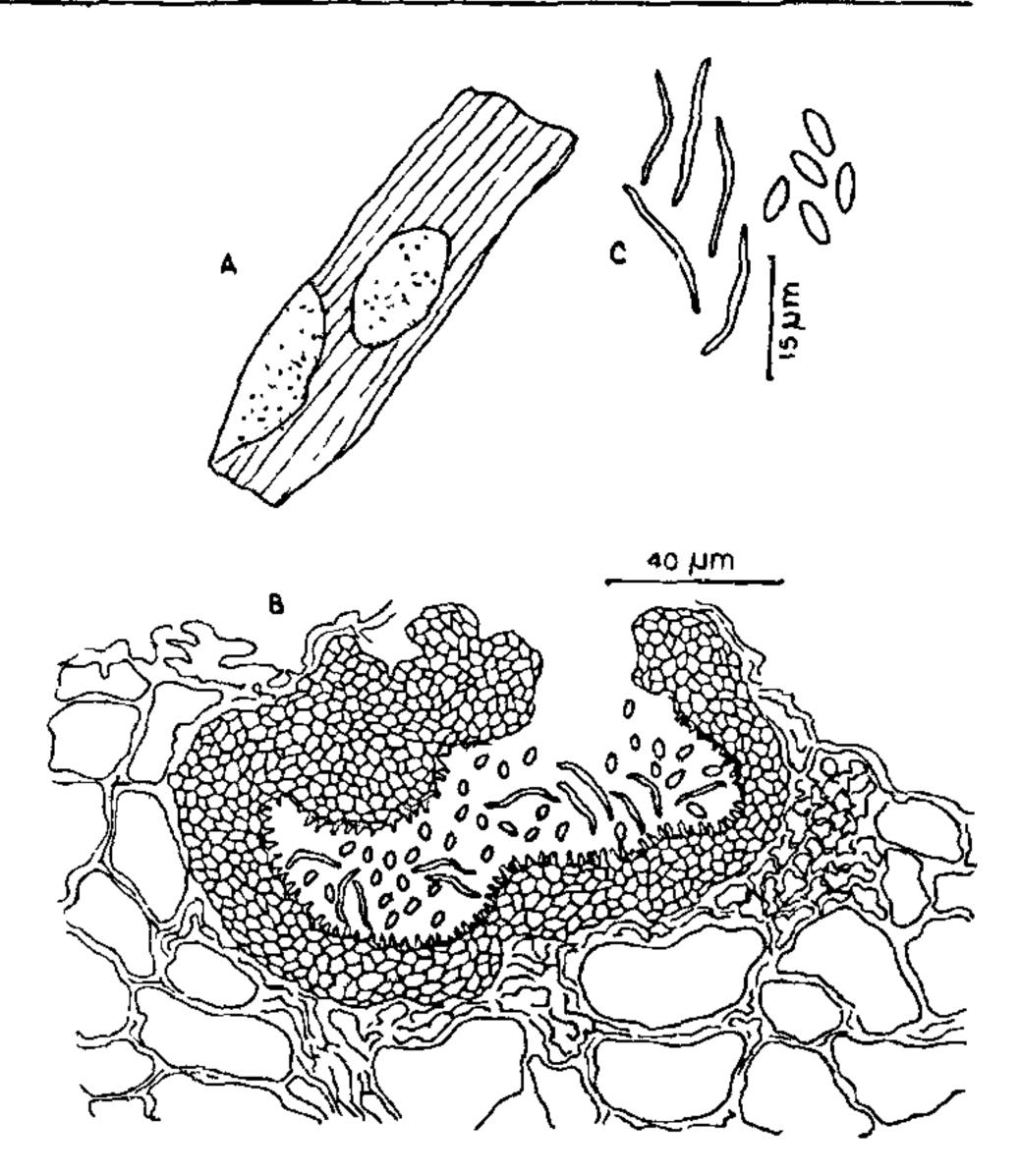


Figure 1. Phomopsis spectabilae sp. nov. A. Leaf spots (habit). B. V. S. of a pycnidium. C. Pycnidiospores (Conidia).

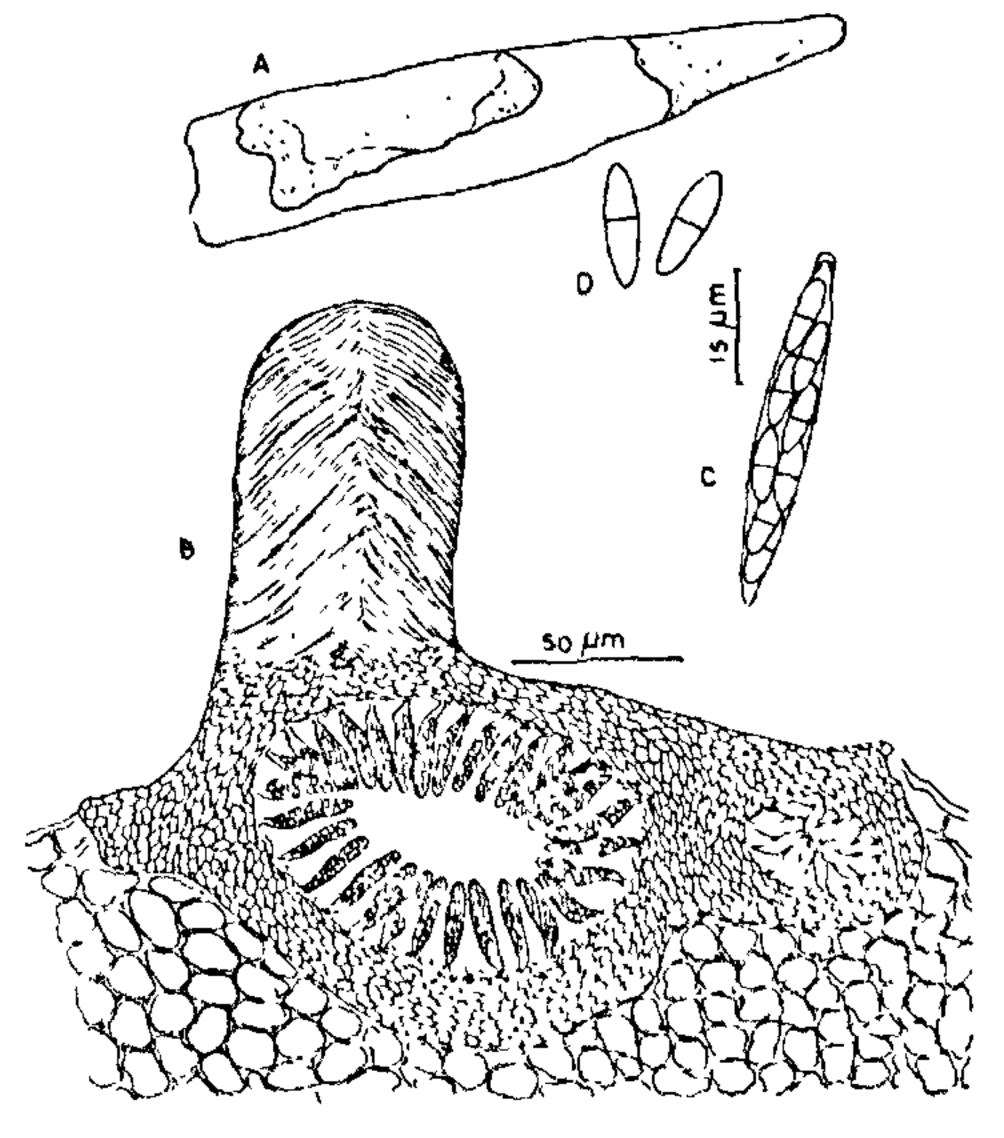


Figure 2. Diaporthe spectabilae sp. nov. A. Leaf spots (habit). B. V. S. of an ascocarp showing asci. C. Ascus. D. Ascospores.

Ascostramatae dispersae, botryosae, nigri, usque 300 μ m diam., neck elongata, leniter ostiolata; asci octo-sporae, fusiformia, sessilis, magnit. 41.6-45 × 8 μ m. bitunicatae, evanascentate; ascosporae uniseptata, guttulata, acutae ad apices, 9.9-11.6 × 3.3 μ m.

Matrix: În folus vivis Aregelia spectabilis Mez., (F. Bromeliaceae), leg. B. R. D. Yadav, in Oct. 1977, ad Pune, AMH 4082 (Holotypus).

Remarks: Both the perfect and imperfect stages were encountered in one and the same infection spot. This forms a new host record since both the fungi are reported herein for the first time on the said ornamental plant.

Grateful thanks are due to Dr V. P. Bhide for the interest in this study and to the Director of the Institute for facilities.

7 January 1985

- 1. Afx, von, J. A., The genera of fungi sporulating in pure culture, J. Cramer. Vaduz. (Germany, 1974, p. 315).
- 2. Sutton, B. C., The Coelomycetes, C.M.I. Publ., Kew, England, 1980, p. 696.

ULTRASTRUCTURE OF SPERMS OF HEAT STERILIZED DYSDERCUS KOENIGII F

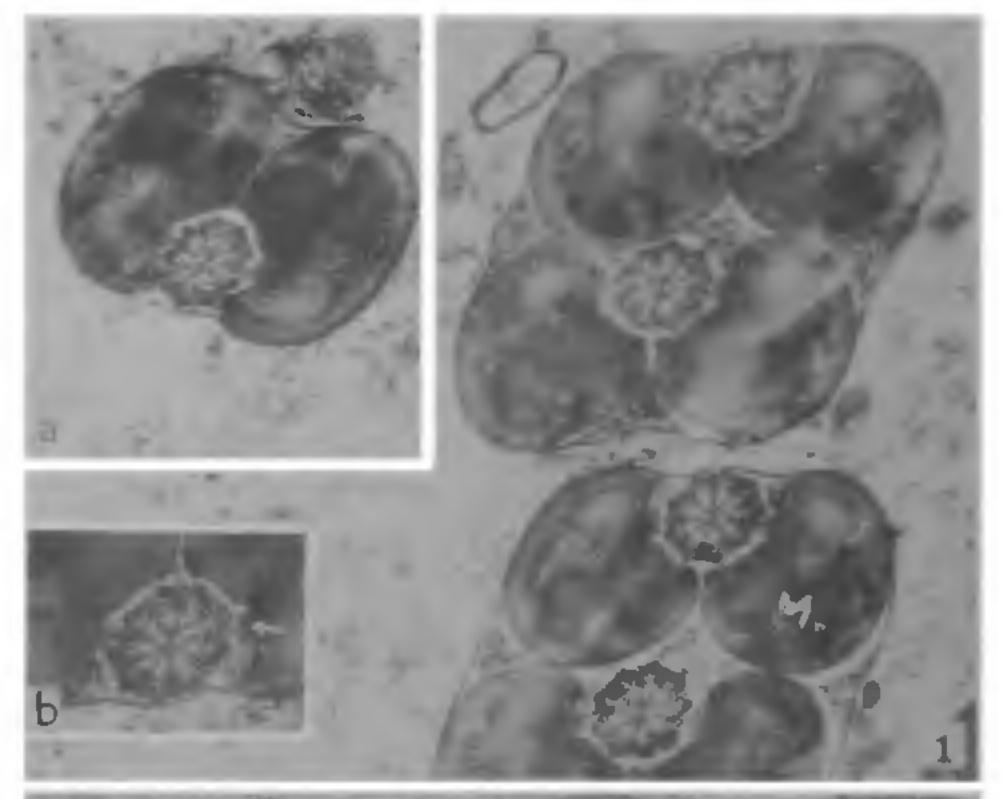
K. TIKKU and B. P. SAXENA*

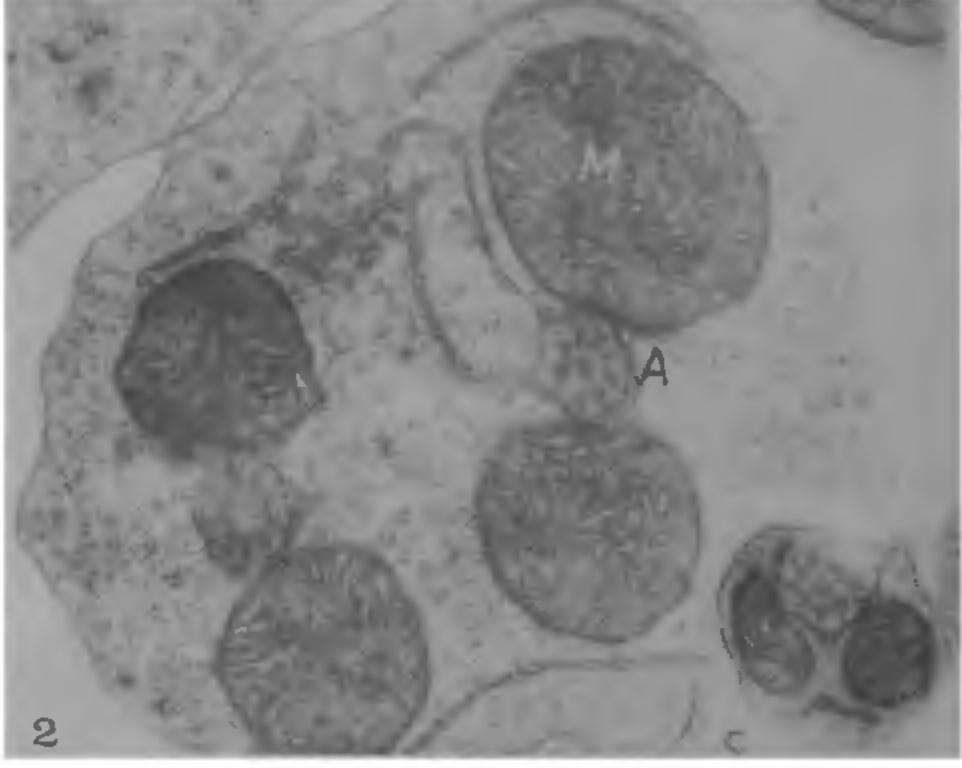
Division of Insecticides and Insect Physiology, Regional Research Laboratory, Jammu 180001, India.

In laboratory cultures of the Red Cotton Bug (Dysdercus koenigii F) almost 100% sterility was noticed when temperature of the culture room was increased to between 35° and 40°C. Females laid what looked like normal eggs after normal copulation, but the eggs failed to hatch. Their colour also did not change from pale yellow to orange which normally indicates the developmental process of normal eggs. To determine the cause of sterility, the sperms produced by these sterile males were examined using electron microscope.

Testes were dissected out and fixed in 2.5% glutaraldehyde in phosphate buffer (pH 7.2) along with the vas deferens (testis cut into 3 portions: upper, middle & lower; vas deferens kept as whole), post-fixed in 1% osmium tetroxide in the same buffer, dehydrated in acetone grades and embedded in Epon 812. Ultrathin sections were stained with uranyl acetate and lead citrate and examined in electron microscope (JEOL 100 CX-II).

Figure 1 shows a T.S. of mature sperms in the vas deferens of sterile adult male. Figure 2 shows the T.S.





Figures 1-2. 1. Two mature sperms (in vas deferens) having common plasma membrane ensheathment. Mitochondrial derivatives (M), each with 3 crystalline bodies, are seen surrounding the axoneme; the background has precipitated semen, × 29000. 2. Two spermatids developing in the cytoplasmic matrix. Mitochondrial derivatives (M) with prominent cristae and axoneme (A) getting formed. Initial formation of plasma membrane and the free ribosomes (dark granules) can also be seen, × 19000. [Insets: (a) Normal mature sperm, × 29000 (b) Curved end feet (arrow) of mitochondrial bridges, × 36000 (c) Tail end of a normal spermatid, × 19000