

First report of leaf blight disease of *Gloriosa superba* L. caused by *Alternaria alternata* (Fr.) Keissler in India

Chandan Kumar Maiti · Surjit Sen · Amal Kanti Paul · Krishnendu Acharya

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Abstract Leaf blight disease was found on *Gloriosa superba* L. (Liliaceae), an endangered, herbaceous, perennial, climbing lily that produces colchicine, in West Bengal, India in 2004. Small brownish spots on leaves developed into concentric rings, which eventually darkened and coalesced to blight the entire leaf. The causal fungus was morphologically identified as *Alternaria alternata* (Fr.) Keissler. This is the first record of *A. alternata* on *G. superba*.

Keywords *Gloriosa superba* · Leaf blight · *Alternaria alternata*

Introduction

Gloriosa superba L. (glory lily, Liliacea) is an endangered, herbaceous, perennial, climbing medicinal plant found throughout India. Colchicine is the major compound isolated from the seed and rhizome of this plant (Sarin et al. 1974). It has abortifacient, anti-pyretic, anti-inflammatory and anti-leprotic properties (Guhabakshi et al. 2001). A leaf blight disease of this plant was observed during 2004–2006 in different areas of lower Gangetic plains of West Bengal, India. Recorded losses were estimated to be 65–80% per annum.

Symptoms

A leaf blight disease of this plant was observed during June to September (temperature $31 \pm 2^\circ\text{C}$). Early symptoms appeared as small, circular to oval, light brownish spots (25–38 mm), 2–6 per leaf, scattered at the tip, margin, and midrib of the leaves. Subsequently, the spots enlarged and usually developed into a concentric ring. At an advanced stage, the spots became dark brown to blackish in colour, gradually coalesced and became irregular in shape, then the affected leaf blighted completely (Fig. 1a–c).

Identification of the pathogen

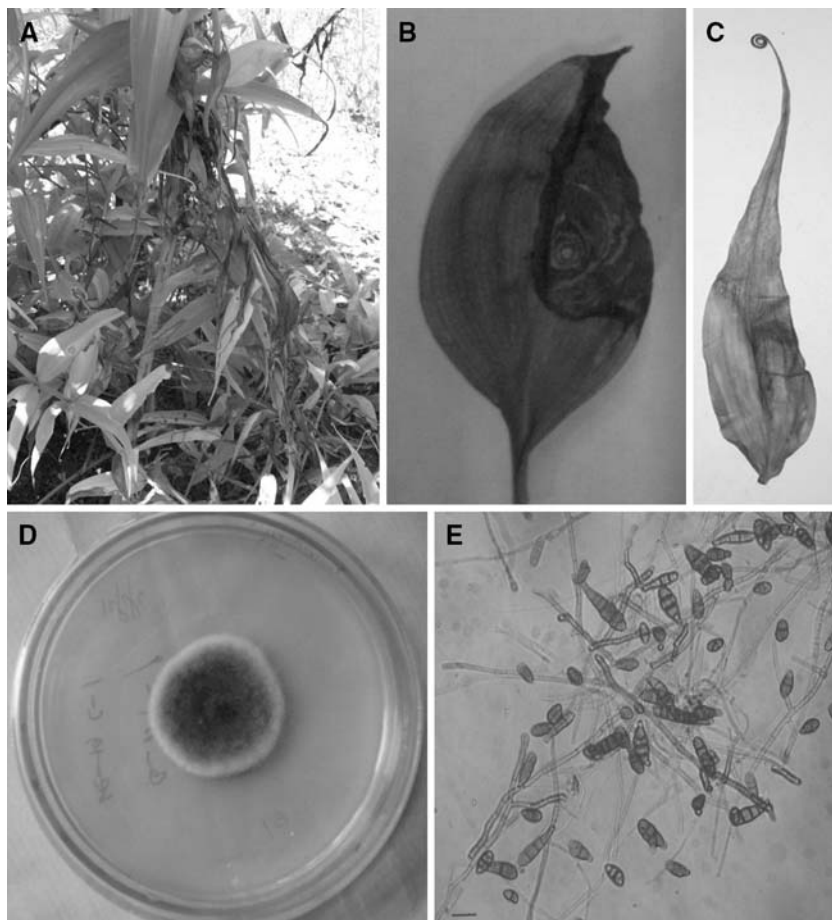
A fungus was consistently isolated from the blighted leaf on potato dextrose agar (PDA) as a pure culture (Fig. 1d). The fungus produced abundant branched, brownish, septate mycelia; conidia were found on leaf peels and scrapings, in cleared leaves and in vertical sections of leaves. Conidial size varied from $17.5\text{--}52.5 \times 8.5\text{--}16 \mu\text{m}$, with 3–5 transverse septa (Fig. 1e). Characteristics of conidia from cultures were similar to those of conidia isolated from infected plants. Based on the morphological characters, the organism was identified as *Alternaria alternata* (Fr.) Keissler (Ellis 1971), and the identification was confirmed by the Agharkar Research Institute, Pune, India.

Pathogenicity test

Leaf disks (14 mm) were surface sterilized (0.5% sodium hypochlorite for 2 min), then placed on 0.5% water agar containing $1 \mu\text{g/ml}$ of benzylaminopurine (Ash and Lanoiselet 2001). They were then inoculated with $10 \mu\text{l}$ of

C. K. Maiti · S. Sen · A. K. Paul · K. Acharya (✉)
Molecular and Applied Mycology and Plant Pathology
Laboratory, Department of Botany, University of Calcutta,
35, Ballygunge Circular Road, Kolkata 700 019, India
e-mail: krish_paper@yahoo.com

Fig. 1 **a** Leaf blight disease of Glory lily (*Gloriosa superba* L.) at an advanced stage, naturally infected with *Alternaria alternata*. **b** Close-up of a naturally infected leaf with clear concentric rings. **c** Blighted leaf at advanced stage. **d** 72 h old colony of *A. alternata* on PDA. **e** Mycelia and conidia of *A. alternata* isolated from a leaf lesion. Bar = 25 μ m



a conidial suspension (10^4 conidia/ml sterile distilled water). A control leaf disk was inoculated with sterile distilled water. The leaves were incubated at 25°C for 4–5 days. After 5 days, typical target spot symptoms developed on the inoculated disks. *A. alternata* was reisolated from such disks.

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