

شبكة المعلومات الجامعية







شبكة المعلومات الجامعية التوثيق الالكتروني والميكروفيلم



شبكة المعلومات الجامعية

جامعة عين شمس

التوثيق الالكتروني والميكروفيلم

قسم

نقسم بالله العظيم أن المادة التي تم توثيقها وتسجيلها على هذه الأفلام قد أعدت دون أية تغيرات



يجب أن

تحفظ هذه الأفلام بعيدا عن الغبار في درجة حرارة من ١٥-٥٠ مئوية ورطوبة نسبية من ٢٠-٠٠% To be Kept away from Dust in Dry Cool place of 15-25- c and relative humidity 20-40%



بعض الوثائـــق الإصليــة تالفــة



بالرسالة صفحات لم ترد بالإصل

IN PLANTS AS A RESULT OF VIRUS INFECTION

B 5789

By

SABRY YOUNIS MOHAMED MAHMOUD

B.Sc. (Plant Pathology), Kafr El-Sheikh, Tanta University (1984) M.Sc. (Plant Pathology), Kafr El-Sheikh, Tanta University (1993)

A thesis submitted in partial fulfillment of the requirements for the degree of

DOCTOR OF PHILOSOPHY

in Agricultural Science (Agricultural Virology)

Department of Agric. Microbiology Faculty of Agriculture Ain Shams University

2000



APPROVAL SHEET

ANTIVIRAL SUBSTANCES INDUCED IN PLANTS AS A RESULT OF B5789 VIRUS INFECTION

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SABRY YOUNIS MOHAMED MAHMOUD

B.Sc. (Plant Pathology), Kafr El-Sheikh, Tanta University (1984) M.Sc. (Plant Pathology), Kafr El-Sheikh, Tanta University (1993)

This thesis for Ph.D. Degree has been approved by:

Prof. Dr. A.I. Abo El-Ghar

Prof. of Plant Pathology, Fac. of Agric., Minufiya University.

Prof. Dr. B.A. Othman

Prof. of Agricultural Virology, Department of Agricultural Microbiology, Faculty of Agric., Ain Shams University.

B-A-Othman

M. A. Aballman Prof. Dr. M.A. Abo El-Nasr

Prof. of Agricultural Virology, Department of Agricultural Microbiology, Faculty of Agric., Ain Shams University. (Supervisor)

Date of examination 30 /4 /2000

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B.Sc. (Plant Pathology), Kafr El-Sheikh, Tanta University (1984) M.Sc. (Plant Pathology), Kafr El-Sheikh, Tanta University (1993)

Under the Supervision of:

Prof. Dr. M.A. Abo El-Nasr

Professor of Agricultural Virology (Department of Agricultural Microbiology, Faculty of Agriculture, Ain Shams University).

Prof. Dr. K.A. El-Dougdoug

Professor of Agricultural Virology (Department of Agricultural Microbiology, Faculty of Agriculture, Ain Shams University).

Prof. Dr. F.M. Maklad

Professor of Plant Pathology (Plant Pathology Department, National Research Centre, Dukki, Giza, Egypt).



ABSTRACT

Sabry Younis Mohamed Mahmoud "Antiviral substances induced in plants as a result of virus infection", unpublished Doctor of Philosophy Dissertation, University of Ain Shams, Faculty of Agriculture, Department of Agric. Microbiology, 2000.

Bean yellow mosaic virus (BYMV) was isolated depending on symptoms, ELISA and electron microscope from different governorates during season 1996. Tomato mosaic virus (ToMV) isolate was identified by indicator hosts and electron microscopy. It was found that *C. amaranticolor* is the suitable hypersensitive host to both viruses, based on acquired resistance and SDS-PAGE.

In leaves of *Chenopodium amaranticolor* locally infected by Tomato mosaic virus (ToMV) or Bean Yellow Mosaic Virus (BYMV), antiviral substance (AVS) was formed after 5 and 6 days from inoculation respectively in inoculated and uninoculated leaves as inhibitor of virus biosynthesis. Crude AVS was extracted by hydrated calcium phosphate and it was purified by DEAE - column chromatography. AVS was acquired systemic resistance against a virus-challenge inoculation. AVS consists of protein and carbohydrate (phosphorylated - glucoprotein) with a molecular weight of 19,400 and 21,500 Kda.

AVS lost its antiviral activity when treated with alkaline phosphate and α -glucosidase, while it does not when treated with lipase as well as it is inhibited by actinomycin D. This lead to think that its formation depends on the same mechanism which is responsible for the synthesis of cellular RNA, DNA-dependent.

By non-denaturing PAGE, AVS is separated into one antiviral active band in both ToMV and BYMV. SDS-PAGE in denaturing conditions revealed at least two new bands with respect to compared extracts of healthy plants.

The AVS-product is production of non-specific antiviral substances in a general, defence response in plants to viral infections, which could be utilized for practical purposes, such as the use of AVS treatments under green-house conditions and in open field, or the production of transgenic plants producing AVS in the future.

Key Words: BYMV, ToMV, Induced resistance, ELISA, Hypersensitivity, Antiviral substances (AVSs), DEAE-Cellulose column chromatography, Gel filtration column chromatography and SDS-PAGE.

ACKNOWLEDGMENT

Praise and thanks be to ALLAH, the most merciful for assisting and directing me to the right way

I would like to thank my advisor **Prof. Dr. M.A. Abo El-Nasr**, Professor of Agriculture Virology, Faculty of Agriculture, Ain-Shams University for suggesting the problems. I like to state that without his inspiration, this work would not have come out.

My heartly thanks are due to **Prof. Dr. Kh.A. El-Dougdoug**, Prof. of Agric. Virology, Department of Agric. Microbilogy, Fac. of Agric., Ain Shams Univ. for his sincere, kind supervision, critical and valuable criticesm either throughout the present work or during writing the thesis which help in illustring the thesis as it offered now.

Thanks are due to **Prof. Dr. F.M. Maklad,** Prof. of Plant Pathology, National Research Center for their kind helping and valuable suggestions throughout the period of the present work.

Thanks are also contributed my advisor **Prof. Dr. Afaf S. Fahmy**, Professor of Biochemistry, Molecular Biology Department, National Research Centre for her continuous supervision, kind help encouragement through the course of this work and giving the full facilities for proper research work.

I would like to thank **Dr. Saleh A. Mohamed** and **Dr. Tarek M. Mostaffa**, Lecture of Molecular Biology, Molecular Biology Department, National Research Centre, for their valuable help and providing the materials for the various experiments.

I wish to thank all the staff menbers of the Plant Pathology Department, National Research Center, specially, **Prof. Dr. A.A. Morsy** and Department of Agric. Microbiol., Fac. of Agric., Ain Shams Univ., for their help during this study.

