

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

Genetic fingerprint of some *Oreochromis* lines

B 5601

By

Yasser Mohamed Saad

B.Sc. Agriculture, Faculty of Agriculture, Genetics Department

Ain Shams University, 1996.

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

Master of Science

Agricultural Science (GENETICS)

Genetics Department Faculty of Agriculture Ain Shams University 2000



APPROVAL SHEET

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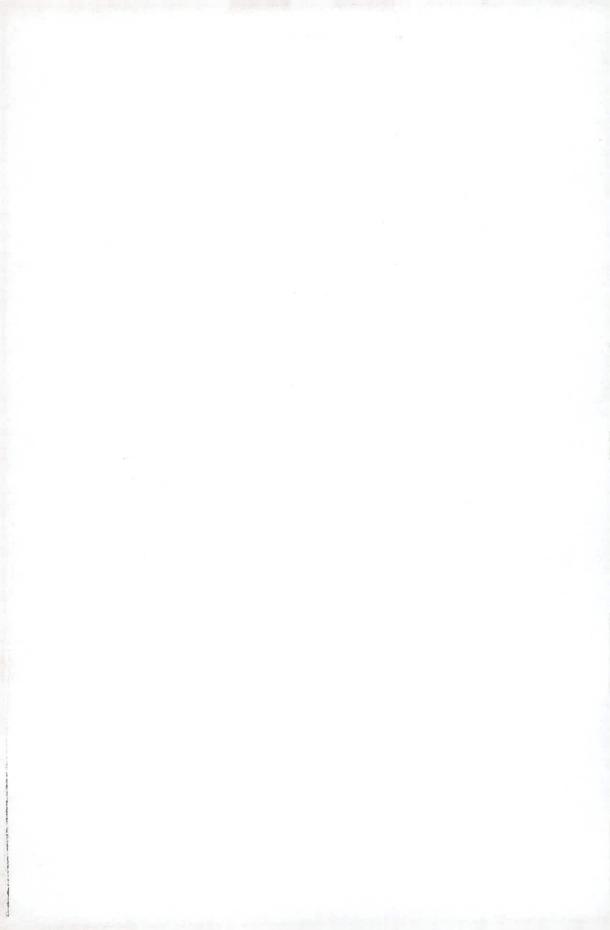
ABSTRACT

Yasser Mohamed Saad Genetic fingerprint of some Oreochromis lines. Master of Science, Genetics Department, Fac. of Agric., Ain Shams University, 1999

The genetic variations among three lines of Oreochromis niloticus collected frome different locations in Egypt were studied based on biochemical and molecular levels. The five isozyme systems, Esterase (*Est*), Lactate dehydrogenase (*Ldh*), Malate dehydrogenase (Mdh), Acide phosphatase (ACPH) and Superoxide dismutase (Sod) were studied in ten organs for each line. To discriminate isozymes among Oreochromis fish populations, different synthetic variations esterase substrates; α -naphthylbutyrate, α -naphthylpropionate and α naphthylvalerate were applied. Muscle and gills esterases had the value of line-substrate band specificity in comparison to both kidney and eye. SDS-PAGE protein electrophoretic patterns were employed to study inter and intra population variations of Oreochromis- lines. each population had a unique protien-banding pattern. RAPD markers were successfully used to discriminate nuclear genetic variations among the applied three lines. In this situation, thirteen different random primers (decamers) were used. Only, six primers (OpB-8, OpB-14, G12, A11, UBC-9 and UBC-4) succeeded in catching and amplifying genomic DNA of all lines.

Key word:

Fish, Nile Tilapia, *Oreochrom* is Electrophoresis, -protein, Isozymes, allozymes, DNA, RAPD-marker, Ontogenic variation.



Acknowledgement

I wish to express my sincere gratitude to prof. Dr. M. A. Rashed Professor of genetics Fac. Agric., Ain Shams Univ., for his supervision, valuable assistance, guidance and continious help during thesis preparation.

Also special thanks to Dr. A. A. El Gamal, associate professor of animal production, Fac. Agric., Ain Shams Univ., for his supervision and his valuable advise during thesis preparation.

I wish to express my deep thanks thanks to Dr. T. M. A. Tantawi, lecturer of genetics, Fac. Agric., Ain Shams Univ, for his supervision, valuable assistance, encouragement and unlimited guidance.

Deep thanks to all staff members of Ain Shams Center of Genetic Engineering and biotechnology (ACGB) for their helping and proceeding the gel documentation and statistical analysis.

I wish to express my thanks to all staff members of Aquaculture closed system laboratory Fac. Agric., Ain Shams Univ., for their great help during the collecting samples.

Additionally I wish to thank my family for their encouragement, assistance and support during my study.

