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#### MOLDS IN LITTER, AIR AND IN-CONTACT WORKERS

#### IN POULTRY HOUSES.

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#### INTRODUCTION

Some mycotic diseases are quite common among birds. The possibility of the transmission of these diseases from birds to man can not be neglected. From these diseases, aspergillosis proved to be the most serious.

Asperoillosis occurs in crowded poultry houses

Aspergillosis occurs in crowded poultry houses and sometimes causes severe losses in the flock of young chickens. This mostly occurs after exposure of the birds to spores of the fungus from heavily contaminated litter, hau or other moldy feedstffs [Ross 1966 Ainsworth and Austwick 1973 and Hubbert et al. 1975].

Aspergilli are abundant in the environment and can grow under a wide range of conditions in litter as saprophytes and produce large numbers of spores which are distributed throughout the environment [Aleksandrov et al. 1974].

Maπ is also susceptible to pulmoπary aspergillosis which is an occupational disese of agriculture workers and others who are often exposed to high concentrations of spores [Buxton and Fraser, 1977].

The purpose of this study was to detect fungus contamination in litter, air and among workers in poultry houses.

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## Material and Methods

- Poultry Houses: The study was conducted at Zagazig Sharkia Governorate. Five broiler poulty houses, of a deep litter system, were investigated periodically for collecting specimens from litter and air.
- <u>Man</u> Fifteen persons in-contact with birds in poultry houses were examined.

## Collection of specimens:

- 1. Litter: Under complete aseptic conditions samples of litter from different poultry houses were collected. Ahandfull samples of the litter were collected randomly, in sterile polyethylene bags, from different places. The procedure recommended by Williams et al.[ 1975 ] was adopted.
- 2. Air:- This was carried out by the use of settling plate technique. Five sterile Petri-dishes of 10 cm diameter containing Sabouraud's dextrose agar were placed open for 30-60 seconds in different places in each poultry house that was previously subjected to litter sampling. These plates were placed at a hight corresponding to the poultry level. The plates were then covered and labelled.
- 3. Man: 15- workers attending the same poultry houses were examined and subjected to nose, ears, and hands swabing. Under aseptic conditions, 2 sterile cotton awabs were taken from inside the nares and external

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ears, they being thus moistened with their secretions.
Two sterile contton swabs were also streaked against the skin of worker's hands. All swabs were placed into sterile tubes containing Sabouraud's dextrose broth.

Collected samples were transferred to the laboratory within 2 hours.

## II. Mycological investigations :

- I. Litter: One gm. of thoroughly mixed litter sample was suspended in 9 cc. sterile saline solution. One ml. from the sample was transferred into 2 sterile Petri-dishes. Ten ml. of melted Sabouraud's dextrose agar (at 40 C) were aseptically poured into each dish and thoroughly mixed with the sample. After solidification, the plates were labelled, and incubated at 25 C for 3 5 days and were not discarded before 2 weeks.
- 2. Air Sabouraud's settling plates were incubated for 3-5 days at 25 C and they were not discarded before 2 weeks.
- 3. Man: Swabs from man were pressed against the sides of the tubes to expel the excess of broth into the Sabouraud's dextrose broth. The tubes were then centrifuged for 20 minutes at 3000 r.p.m. The supernatant fluid was discarded. Loopfulls from the sediment were inoculated into 4 plates containing Sabouraud's agar. Plates were incubated at 25 C for 3-5 days and were not discarded before 2 weeks.

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# 8III. Identification of various isolated Fungi: Fungal growth was identified according to Emmons et

al., (1977) and Treagan and Pulliam (1982).

Results are shown in Tables (1 & 2).

Table (1) Molds isolated from 50 samples of each litter and air in poultry houses.

ar ringa Southill	Litter									
Mold spp.										
	No.	of i	solates	%	No.	o f	isolat ,	es (	% 	
A.fumigatus		rair s	8	16	.27		-10		20	
A.flavus		10		20			6		12	
A.niger ·		<u></u>	el III en	4- <b>-</b>			3		6	
Penicillium	1	2	or verific d	4			4		8	
Alternaria .			· ma pl	etes			6		12	
Mucor		9,92	2 (18)	4			4		. 8	
Cephalosporium			gradines :	2_0			1		2	
Scopulariopsis			1	2			1		2	

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Table [2] Molds isolated from mose pars and

Table [2] Molds isolated from mose, ears and hands of 15 workers in poultry houses.

Mold spp	No	ose	Ears		Hands	
	No. of	%	No.of	%	No. of	%
A.fumigatus	5	33.3	3	20.0	3	20.0
A.flavus	7	46.7	4	26.7	6	40.0
A.niger	3	20.0	3	20.0	2	13.3
Penicillium	ari amoru	field Arrive	3	20.0	2	13.3
Alternaria	one julia i La La La Alla I	o do ET e los	5	33.3	(2 1 <del>-</del> 1 1 1 1	
Mucor autiliza	nga <del>-</del> na a				2	13.3
Cephalosporium	era – philip		the office a		3	20.0
Hormodendrium	na sanari		2	13.3	(20.54	
Stemphyllium	3	20.0	· L	26.7	town of	

#### Discussion

Air and litter play an important role in transmitting some mycotic diseases between man and birds [Emmons et al. 1977].

Tables [1 & 2 ] show that Aspergillus fumigatus was isolated from litter and air of poultry houses with percentage of 16 and 20, respectively, whereas the percentage of A . flavus was 20 in

litter and 12 in air. Moreover, in man A.fumigatus was recovered from nose, ears and hands of in-contact workers with percentage of 33.3, 20.0 and 20.0, respectively, while the percentages of A. flavus were 46.7 in nose, 26.7 in ears and 40.0 on hands.

From these results, it is evident that A .fumigatus and A.flavus are isolated from litter, air and man. A fact which may throw light on the occurrance of these species in the litter and man, and the possibility of the transmission of their spores from litter to man'through air. Buxton and Fraser [1977] and Emmons et al. [1977] mentioned that Aspergillus species grows frequently and abundantly as saprobes on decaying straw, and under certain circumstances man and chickens are exposed to the disease by inhaling a great number of the Aspergillus spores, especially Aspergillus fumigatus.

Otherwise, A. fumigatus and A. flavus were reported to invade and colonize masal sinuses in man [Emmons et al. 1977]. Buxton and Fraser [1977] stated that Aspergillus spp. may be associated with lesions in the skin, external ear, nasal sinuses, and occasionally in the bones and meninges, but the ear and nails are relatively the commonest sites of infection.

Concerning, A.niger it was isolated from air, nose, ears and hands of in-contact workers with percentage of 6,20, 20 and 13.3 respectively.

A. niger was reported to be an important etiologic agent of atomycosis [Emmons et al. 1977]. Koneman et al. [1979] indicated that A. niger may be associated with fungus ball infections of the masal sinuses or lung, and the

agent mostly commonly recovred from cases of otitis ext-

erna "Swimmers ear".

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Table [1] also shows that, besides Aspergillus spp., there are many other molds recovered from litter. There were Penicillium, Mucor and, Scopulariopsis with percentage of 4,4 and 2, respectively. Penicillium, Alternaria, Mucor, Cephalosporium and Scopulariopsis were also isolated from air with percentage of 8, 12, 8, 2and2 respectively. Penicillium sp., Mucor sp., Aspergillus fumigatus, A.niger, and A.flavus were isolated from air by Dye and Vernon [1952], Gupta et al.[1960]. Aleksandrov and Peev [1974] and Metwally & Aidaros [1979]. These molds are blamed for the cause of many diseases in man.

Regarding in-contact workers, (Table 2) Clarifies the isolation of molds, other than asperillus spp., from nose, stemphyllium was isolated with percentage of 20.0 pemicillium, alternaria, hormodendrium and stemphyllium were isolated from the ear with percentage of 20.0, 33.3,13.3 and 26.7, respectively. Molds isolated from hands were Penicillium, Mucor, and Cephalosporium with percentage of 13.33, 13.3. and 20.0 respectively. Also, it is noted that most persons out of 15, were multicarriers of A.fumigatus, A.flavus and stemphyliium in nose, Alteranaria, stemphyllium, and A.flavus in ears and A.flavus, cephalosporium and A.fumigatus on hands.

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Accordingly, contaminated litter [moldy straw] and air, other than infected person or birds, may a role in transmitting mycotic diseases to man and birds in poultry houses. A fact which throws light on the importance of environmental samitation in poultry houses, Particulary avoiding damp litter, cleaning food troughs and efficient ventilation.

#### Summary

Litter, air, and in-contact workers of five poultry houses were examined mycologically.

The isolated fungus species from 50 air samples were A.fumigatus, A.flavus, A.niger, Penicillium, Alternaria, Mucor, Cephalosporium and Scopulariopsis.

The fungus-species isolated from 50 litter samples were A.fumigatus, A. flavus, Penicillium and Scopulariopsis.

15 in-contact workers were found to be nose, ear and hand carriers of nine strains of which seven were previously isolated from air and litter of the poultry houses.

#### References

Ainsworth G.C. and Austwick P.K.C.[1973].Fungal diseases of animals.2gd Ed.Commonwealth Agriculture Bureaux, Farnham Royal, Slough, England.

- Zagazig Veterinary Journal 62
- Aleksandrov M. and Peev N.[1974].Fungi in the the air of poultry houses . Veternarno meditsinski Nauki, Bulgaria 11, No. 2,43-50.
- Buxton A. and Fraser G. [1977]. Animal Microbiology.Vol.

  1. Immunology, Bacteriology, mycology.Blackwell
  Scientific Publications LTD, 1st Published,
  London.
- Dye M.N. and Vernon T.R. [1952]. Air borne moulds spores
  New Zeel. J., Sc. and Tach. B.34, 118-127.
- Emmons C.W., Binford C.H., UTz J.P. and Kwon-Chung K.J.
  [1977]. Medical Mycology, 3rd Ed., Lea and
  Febiger, Philadelphia.
- Gupta K.D., Segani I.C. and Kasliwa R.M. [1960]. Survey of the allergic diseases caused by the air borne spores in Juiper. Ind. J.Chest. Dis.2,273.
- Hubbert W.T, McCuloch W.F., and Schnurrenberger P.R.[1975]

  Diseases transmitted from animals to man. 6 th
  Ed. Charless, C.; Thomas, Publisher, Spring field, Illinois, U.S.A.
- Koneman E.W., Allen S.D., Dowell V.R. and Sommers H.M.
  [1979]. Color Atlas and Textbook of Diagnostic
  Microbiology. J.B. Lippincott Co., Philadelphia, Toronto.
- Metwally M.Abd EL-R.and Aidaros H. [1979]. Air pollution in poultry houses in relation to various environmental conditions. Res. Bull. No. 39. Zagazig Journal of Agri. Research.

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- Ross E. [1966]. An aspergillosis outbreak in a broiler flock on bagasse litter. Hawaii Fm Sci.14, No. 3 pp. 5-6.
- Treagan L. and Pulliam L. [1982]. Medical Microbiology
  Laboratory Procedures. W.B. Saunders Co.
  PA, USA.
- Williams J.E., Mallenson and Soneyenbos G.H. [1975].
  Isolation and Identification of avian pathogens, American Association of avian pathologists"

ملخـــــ البحـــــ

# العفن في الفرشة والهوا عوالعمال المخالطين داخل مساكن الدواجن

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بدراسة خمسون عينه من الفرغه في طمين مزارع لانتاج دجاج التصحيسين ثم عسزل الانواع الاتيسه من العفن ، اسبر جيلوس فيوميجتيس ، اسبر جيلوس فيلافس، بنسليسوم ، ميسوكسر ، سكا بيولاريوبسس، كما تم عسسزل الانسواع الاتيسه من العفن ، اسبر جيلوس فيوميجيتس ، اسبر حيلوس فلافسس اسبر جيلوس نيجر ، بنسليسوم ، الترناريا ، ميوكسر ، سيفالوسبوريسس سكا بيولاريوبسسس، من خمسون عينه من الهوا اداخل هذه المساكن الدواجن،

فحم خمسة عشر من العامليين في هذه المساكن بأخذ عينيا من الانف ، والاذن والايدى بأمكين عزل نفس المعفق المعزولة سابقا من الغرشة والهوا ولاسيما فطر اسبر جليون فيوميجيس ، اسبر جلون فعلاقس ، اسبر جليون نيجسر ، بنسليسوم .