## Quantitative and Qualitative Assessment of Aflatoxins Residues in Home and Industrial Broilers ChickenLivers

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#### Abstract:

Aflatoxins are a group of toxins mostly produced by Aspergillus flavus and Aspergillus parasiticus, two moulds that thrive in warm. humid climates on different crops which are used in manufacture of chicken rations. People can be exposed to aflatoxins which have a carcinogenic, mutagenic, teratogenic, immunosuppressive, and other adverse effects by consuming chicken livers from poultry that ate Aflatoxin contaminated feed. The result revealed that the mean residual values of 100 liver sample; home chicken broiler livers (n=50) were 27.61 $\pm$  6.60 (µg/kg) for AFB1; 0.92 $\pm$  0.35 (µg/kg) AFB2;  $0.10 \pm 0.05$  for AFG1 and  $44.30 \pm 14.50$  (µg/kg) for AFG2. While in industrial chicken liver sample (n=50) were 5.44  $\pm$  0.87  $(\mu g/kg)$  for B1; 0.61 ± 0.14  $(\mu g/kg)$  for AFB2 and 7.10±1.10  $(\mu g/kg)$  for AFG2. The prevalence of the mean  $\pm SE$  of the total AFLs/ µg/kg residues in home and industrial rearing chicken broiler liver samples were 72.47 $\pm$  0.12 µg/kg and 12.54 µg/kg and 12.78  $\pm$ 0.008 µg/kg respectively. The detected total aflatoxins were exceed the MLs (15-20ppb) in home rearing chicken livers samples, while lied with the limits in industrial rearing liver samples. Monitoring of aflatoxin residues must be implemented and control programs in all broiler meat products.

**Key words**: Aflatoxin, chicken liver, home rearing, Industrial rearing, human risk

#### Introduction

Aflatoxins are chemicals that are among the most prominent types of mycotoxins that affect poultry. These toxins are secreted by some mould species. The danger of aflatoxin lies in the fact that they infect birds without any symptoms appearing. That is why it is called the silent killer and It is thought to be the most potent naturally occurring liver carcinogen to play a part in up to 28% of liver cancer cases globally (*Liu and Felicia*, 2010) Broiler chicken livers is uniquely rich nutrition profile

makes it a worthy addition to any nourishing diet. Due to high content oxygenating iron, low in of calories, Folate, Biotin that perform a multitude of functions, rich in protein, B vitamins, ultra-high vitamin A. rich in selenium and vitamin C and vitality of internal (Bordoni and Danesi organs (2017). Aflatoxins discovered began shortly after an outbreak of an unknown etiologic disease in turkeys in 1960. The illness was dubbed Turkey "X" disease, and it later linked to a tainted was groundnut meal imported from Brazil (Blount, 1961). From there, a thorough investigation revealed that mould species known а as Aspergillus flavus and Aspergillus parasiticus were build up a secondary metabolites referred after that to Aflatoxins which were the real cause of Hepatotoxicity, (Liu and Felicia, 2010; Molaghi et al., 2017; WHO, 2018, and Alam et al., 2020).Chicken meat are distinguished by their flavour. affordability, and ability to offer an exceptional, well-balanced source of animal proteins for people of all ages (kralik et al., 2017). It's known that mycotoxins not only cause problems for the animal feed industry but also threaten the human's safety. The adverse health effects of mycotoxins are range from acute poisoning to long-term effects such as immune deficiency and cancer (Bryden, 2012 and WHO, 2018). Due to their toxicity, the majority of political authorities

limit the amounts of AFs that can be used in human and animal food. The permissible levels of AFs in human and animal foods depend on the perceived danger and the governing bodies' purview (*Gupta*, 2018).

In Egypt, the poultry industry is exceedingly based on the imported ingredients for feed formulation, in this way, the defilement with fungi are exceptionally tall either amid the stages of generation or amid transportation period or may happen amid capacity stages within the markets (Lizárraga-Paulínet al., 2011 and Bryden, 2012). Aflatoxins are considered the most important form of mycotoxins, most dangerous for their serious pathogenic effects and economic losses caused by the destruction of polluted crops; (Agag, 2004 and Kumar et al., 2020). The greatest accumulation of Aflatoxins was observed organs which in responsible for biotransformation, such as the liver and kidney (Del et al., 2005). According to the type of chicken broiler rearing, age and ration performed in addition to degree of mould contamination aflatoxin residues were dependant (Hussain et al., 2010 and Shareef, 2010). Therefore, this study was carried out to assess the prevalence and the content of aflatoxin residues in rearing chicken broiler liver samples which collected from both home rearing and industrial rearing broilers which having different age;

feeding and rearing systems located in Alsharqia governorate.

### Materials and Methods Standard Solutions:

The stock aflatoxin standard solutions were prepared according to the A O A C (2000).

# Extraction of Aflatoxins from the sample: *A O A C (2000)*

One hundred (100) gram liver sample was homogenized. 10 ml of 20% citric acid was added and mixed well 200mL of dichloromethane was added and held on the automatic shaker for 30 minutes. The mixture was filtered filtrate evaporated and the in vacuum. Add hexane and redetect the extracted material.

### Clean-Up:

The procedure was carried out using Solid-Phase Extraction columns.

#### **HPLC** analysis

Analysis of Aflatoxins was done by Agilent HPLC apparatus, Faculty of Veterinary Medicine, Zagazig University

#### **Results and Discussion**

The results showed in Table (1) the minimum, maximum and mean values  $\pm$  SE of Aflatoxins residual level (µg/kg) in the examined liver samples of home chicken broiler rearing livers were 1.2, 67.97 and 27.61 $\pm$  6.60 (µg/kg) respectively for AFB1;  $\leq$  1.0, 3.30 and 0.92 $\pm$  0.35 (µg/kg) respectively for AFB2;  $\leq$  1, 0.50 and 0.10 $\pm$  0.05; respectively for AFG1 and 2.40, 150.2 and 44.30  $\pm$  14.50 (µg/kg), respectively for AFG2. The obtained results were

nearly similar to those reported by Nashwa et al. (2017) and Ibrahim et al. (2018) but lower than those reported by El-Nabarawy et al. (2021), while in industrial chicken broiler rearing livers (Table2) were 2.20, 9.50 and 5.44  $\pm$  0.87 (µg/kg) respectively for AFB1;  $\leq 1, 1.35$ and  $0.61 \pm 0.14$  (µg/kg) respectively for AFB2 and 2.70, 12.50 and  $7.10 \pm 1.10$  $(\mu g/kg)$ for G2 respectively. The obtained results were nearly similar to those reported by Nashwa et al. (2017) and Abd El-Tawaab et al. (2019) and higher than those mentioned by Ebeed et al. (2015) and Sineque et (2017) and Gathumbi and al. Bebora (2000).

frequency The percentages of Aflatoxins residues in examined liver samples as given in Table (3) were 15(36.6%) and 15 (41.7%) for AFB1, 9(21.9%) and 6(16.6%) for AFB2, 2(4.9%) and Zero for AFG1 and 15(36.6%) and 15(41.7%) for AFG2 in examined liver samples of home and industrial rearing chicken broiler livers respectively. These results showed that AFs were arranged according to its prevalence the examined samples in as AFB1>AFG2>AFB2 > AFG1. The results obtained were lower to the result obtained by Sineque et al. (2017) in case of industrial samples while in home sample was higher. AFB1 were considered the most common and dangerous to the human liver because of their carcinogenic. mutagenic and teratogenic effects and

contaminated the chicken livers through the ration which performed to the reared chicken. The results were higher than those obtained by *Ebeed et al. (2015)* and arranged as AFB1>AFG1> AFG2 while AFB2 was not detected, while they were nearly similar to those reported by *El-Desouky et al. (2014), Faten et al. (2016),* and *Nashwa et al.* (2017).

The prevalence of the minimum, maximum and mean ±SE of the total AFLs/ µg/kg residues as given in Table (4) in home and industrial rearing chicken broiler liver samples were 71.91, 72.93 and  $72.47 \pm 0.12 \ \mu g/kg$  and  $12.54 \ \mu g/kg$  $,13.15 \text{ }\mu\text{g/kg}$  and  $12.78 \pm 0.008$ µg/kg respectively. The obtained results in industrial rearing broiler liver samples were nearly agreed with those mentioned by Abd El-Tawaab et al. (2019), Abdalla et al. (2021), and Morshdy et al. (2021) but higher than those mentioned by Ebeed et al. (2015) and Nashwa et

al., (2017). The obtained results in home rearing broiler livers samples were higher than those mentioned by Ebeed et al., (2015), Abd El-Tawaab et al. (2019), Abdalla et al. (2021), and Morshdy et al. (2021). The estimated total aflatoxins were exceeding the MLs (15-20ppb) inhome rearing chicken livers samples, while lied with the limits in industrial rearing liver samples, this was attributed to the type of ration performed, age of the chicken addition and of anti-aflatoxin additives. The accepted and none accepted liver samples according to MLs standards were illustrated, it is clear that all samples of livers collected from home rearing were rejected. On the contrary. all samples from industrial rearing were accepted because they were in line with the permissible limits for the presence of mycotoxins in general and Aflatoxins residues in particular in chicken livers in food stuff

**Table (1):** Incidence of Aflatoxins residues  $(\mu g/kg)$  in examined samples of home rearing chicken broiler liver

<b>C</b>	AFB1	AFB2	AFG1	AFG2
Min	1.2	$\leq 1.0$	$\leq 1$	2.40
Max	67.97	3.30	0.50	150.2
Mean ±SE	27.61 ±06.60	$0.92 \pm 0.35$	$0.10 \pm 0.05$	$44.30 \pm 14.50$

Table (2):	Incidence of Aflatox	insresidues (	(µg/kg) in e	examined sat	mples of
Industrial i	rearing chicken broile	er liver			

	AFB1	AFB2	AFG2
Min	2.20	$\leq 1$	2.70
Max	9.50	1.35	12.50
Mean ±SE	$5.44 \pm 0.87$	0.61±0.14	7.10±1.10

	Home rearing		Industrial rearing			
AFs	broiler liver		broiler liver			
	No	%	N0	%		
AFB1	15	36.6	15	41.7		
AFB2	9	21.9	6	16.6		
AFG1	2	4.9	0	0.0		
AFG2	15	36.6	15	41.7		
Total	41	100	36	100		

Table (3): Frequency % of AFs detection in Home and Industrial rearing chicken broiler liver samples

Table (4): Prevalence of total AFLs/  $\mu$ g/kg in Home and Industrial rearing chicken broiler liver

	AFLs/ µg/kg /Home	AFLs/ µg/kg /Industrial
Min.	71.91	12.54
Max.	72.93	13.15
Mean ±SE	$72.47 \pm 0.12$	$12.78\pm0.008$

Table (5): Accepted and none accepted liver samples according to MLs\* standards

AFLs in home rearing			AFLs in industrial rearing				
broiler livers*		broiler livers					
Acce	epted	Non	accepted	Accepted		Non accepted	
No	%	No	%	No	%	No	%
0	0	15	100	15	100	0	0

\*Permissible limit according to US department of Agriculture (USDA) and Food and Drug Administration (FDA) set a tolerance limit of 20 ppb for aflatoxins in view of their toxic effects on foods (Sarma et al., 2017).

# Conclusion and recommendations:

The Aflatoxin levels either single or total residue recorded in this study were higher than the allowed limits in home rearing liver samples, suggesting that the chicken liver from this sources represents a high risk to consumers and aflatoxin

monitoring should residue be performed and control programs in all chicken broiler rearing meat products. The encouragement of national authorities was recommended to monitor and ensure that the levels of mycotoxins in chicken broilers livers should be low as possible and comply with the

national and international MLs, legislation.

#### **Conflict of interest statement**

The authors declare that there are no conflicts of interest.

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## الملخص العربي التقييم الكمي والنوعي لمتبقيات الأفلاتوكسينات في كبد دجاج التسمين المنزلي والتجاري

الأفلاتوكسينات هي عائلة من السموم التي تنتج بشكل رئيسي من فطر اسبر جيللاس فلافس و اسبر جيللاس بار ازيتاكاس ، و هي وفيرة في البيئات الدافئة والرطبة على محاصيل مختلفة والتي تستخدم في تغذية الدجاج اللاحم. يمكن أن يتعرض الناس للأفلاتوكسينات التي لها تأثيرات مسرطنة ومثبطة للمناعة وغيرها من الأثار الضارة عن طريق تناول كبد الدجاج التي تأكل علفًا ملوئًا ومثبطة للمناعة وغيرها من الأثار الضارة عن طريق تناول كبد الدجاج التي تكل علفًا ملوئًا المنزلي (ن = 50) كان 16.1 في متوسط القيم المتبقية لـ 100 عينة كبد ؛ كبد دجاج التسمين المنزلي (ن = 50) كان 20.1 في متوسط القيم المتبقية لـ 100 عينة كبد ؛ كبد دجاج التسمين المنزلي (ن = 50) كان 20.1 في مقوسط القيم المتبقية لـ 100 عينة كبد ؛ كبد دجاج التسمين المنزلي (ن = 50) كان 20.1 في 50.0 (ميكرو غرام / كغ) لـ 81 ؛ 20.0 ± 20.0 (ميكرو غرام / كغ) كافئ كافئ كفئ كو غرام / كبن 10.0 ± 20.0 له موكو غرام / كغ) لـ 11 ؛ 20.0 ± 20.0 (ميكرو غرام / كغ) كافئ كافئ كافئ كافئ كو غرام / كبن 10.0 ± 20.0 له 20.0 له 20.0 الماذلي (ن = 50) كان 20.1 له 20.0 الماذلي عنينات كبد الدجاج المربي علي نطاق تجاري (ن = 50) كانت 20.4 ± 5.4 (ميكرو غرام / كغ) له 18 ؛ 20.0 الماذلي كافئ الماذلي كافئ الماذ عن معرد وغرام / كغ الماد عام الماذلي كبد الدجاج المربي علي نطاق تجاري (ن = 50) كانت 20.4 ± 5.4 (ميكرو غرام / كغ) له 20.0 معدل معدل عداد الدو عرام / كغ في عينات كبد دجاج التسمين المنزلي والتشار متوسط ± 32 لمجموع دام / كغ و 20.1 ± 10.1 (ميكروغرام / كغ و 20.5 ± 20.00 له 20.0 معدل والتجاري روغرام / كغ و 20.5 ± 20.00 انتشار مولي معدل مولي معدل معدل ميكروغرام / كغ في عينات كبد دجاج التسمين المنزلي معدل والتجاري روغرام / كغ و 20.5 ± 20.00 معدل معدل معدا والتجاري روغرام / كغ و 20.5 ± 20.00 معدل معدل معدل مولي مولي معدل والتجاري مولي مولي معدل ميكروغرام / كغ و 20.5 ± 20.00 معدل معدل والتجاري معدل معدو مام ميكروغرام / كغ و 20.5 ± 20.00 معدل معدل معدو والتجاري ميكروغرام / كغ و 20.5 ± 20.00 معدو عدا مركو عر م / كغ و 20.5 ± 20.00 معد ما والتجاري مي ميكروغرام / كغ و 20.5 ± 20.00 معدو مع ماندوان معدل معدو والتحالي الماذلي ما ميذلي ما دود الممو مي ما ميكرو غرام / كغ و 20.5 ± 20.00 معد ما ولي