Pathogenic effect of bunt wheat on male reproductive system in Mice.

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Summary

This study was carried out to study the toxic effect of smut wheat on male reproductive system of mice. The experimental study was performed in the laboratory using 20 mice, which divided into two groups, one group feed on normal diet, the other fed on a mixture of 50% normal diet with 50% smut wheat. A histopathological study was done on the testes of Mice fed on smut wheat for 30 days. The histopathological study revealed thickness of basement membrane of seminiferous tubule of mice, and desquamation of the sertoli cells and spermatogonium from the basement membrane of seminiferous tubule and has adversely effect on male fertility.

Key words: bunt wheat, male mice, reproductive.

Introduction

Order ustilagenals, causes covered Kernal smut of cereals, it is considered as one of the most important pathogens on wheat and Barley and maize (1); The disease is quite common in southern and central parts of the country and responsible for losses each year (2), Very little work has been done on the toxic effect on man, poultry and other livestock animals (3). The serious disease caused by smuts which have been referred to us "ustilaginism", Debre and Nevot (4) had been described as an infantile disease similar to erythro – edema which may occur frequently in Central Europe in those infants fed on corn or bread containing spores of ustilago mydis , it is thought that consumption of ustilago. Smut of Tilletia carier Tilletia foetid (wallr.) Liro which belong to the same order ustilaginals have also been considered as toxic (5) ustilaginism, poisoning by smut spores which produces burning, itching, hyperemia, acrocyanosis and edema resemble ergotism, pellagra or infantile acrocynia (5and 6). It had been reported elsewhere that it provoke abortion in cattle and may cause contraction of uterine muscle and inhibit adrenaline. All the genera belong to same order have similar activities (7and 8). It was felt therefore that the histopathological study of female and male genetic systems of experimental animals fed on smut spore
of *Tilletia caries* was needed and it was hoped that this work may stimulate other workers in this field in order to expose the effect of these toxins on animals and human.

**Materials and methods**

Thirty adult male albino mice were used in this study at the age of 10 weeks and weight 24 gram ± 1 gram with optimum temperature incubation ranged between 22-25°C and fed on locally prepared diet. Tap water used as drinking water. Soft wood shaving used for bedding. 20 animals were divided into two groups, control group (T1) feed normal diet, and treated group (T2) fed on 50% normal locally prepared diet mixed with 50% smut wheat infected with *Tellitia caries* provided from the Ministry of Trade for 30 days.

**Histopathological Study:**

Male reproductive organs were dissected out after one month of treatment. The organs were fixed in small containers containing 20 ml of formalin 10% and dehydrated in progressively more concentrated alcohol, embedded in paraffin and cut into section 4-5 μm thickness stained with haematoxyline and eosin (H & E) for microscopical examination (9 and 10).

**Results and Discussion**

In the present study the result in the fig.1 and 2 show the normal structure of the testis. In this study the appearance of thickness of basement membrane of seminiferos tubules due to fibroses. The fibroses enclosed the tubules and this lead to decrease in the number of spermatid Fig. 3 and 4 may be due to the irritation by the trimethylamine and the destruction related to the polycyclic aromatic substance in the smut wheat. This results were in agreements with others (6,11,12,13 and 14) who found that there was a toxic, carcinogenic material polycyclic aromatic hydrocarbon (PAH), in addition to the trimethylamine which can cause irritation and damage of the living tissue in the body. This results showed the desquamation of the Sertoli cells and spermatogonium from the basement membrane, this spot the light on the irritative and destructive effect on male reproductive system which may lead consequently to infertility of male animals fed on smut wheat (15). and in some region completely degeneration were observed, this can be explained due to the irritation effect of the trimethylamine and polycyclic aromatic hydrocarbon (PAH) material in smut wheat (12, 13). Fig 5. some PAHs are carcinogens and may effect the male reproductive system (16 ). This result was disagree with the finding by (17) who state that the feeding of smut wheat for 45 days don’t cause toxic effect in chicken. Despite the effect of all PAH metabolism in Mice adrenal gland, testis and overy, hormonally elevated in each tissue by a variety of possible mechanism including induction and selective proliferation of cells (18). According to the histopathological results, it is recommended to make a regulation and standard limit for the use of smut wheat in feeding of animals to reduce the more economical losses and human health.
Fig. 1: Histopathological section of testis of the control male Mice shows spermatogonium (SG), primary spermatocyte (PS), Sertoli cells (SC), Spermatid (S), (H & E 40X).

Fig. 2: Histopathological section of control male Mice show normal structure of the testis (H & E 40X)
Fig. 3: Histopathological section of testis of the male Mice after 30 days post treatment with smut wheat show thickness of basement membrane of semineferous tubule due to fibroses (→), (H & E stain 40X).

Fig. 4: Histopathological section of testis of the male Mice 30 days post treatment with smut wheat show thickness (→) of basement membrane due to fibroses (H & E stain 10X).
Fig. 5: Histopathological section of testis of the male Mice after 30 days post treatment with smut wheat show desquamation of sertoli cells and spermatogonium from the basement membrane of semineferous tubules as well as fibrous interstitial tissue ( ——— ) & in some region completely degeneration were observed , (H & E 40X).

References


6- Stedman's medical spellchecker ©(2006). lippincott Williams & Wilkins.


