



Effect of Ajinomoto (MSG) on Weight & Length of Uterine Tubes of Female Albino Rats

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ABSTRACT

Introduction: Ajinomoto (MSG) is the most popular and frequently used food additive. It is used to enhance the taste of food and is also used for preservation of food. Its use has been there for a longer period of time & has now become an essential component of almost all prepared salty food items. This extensively used product has various health concerns so this study has been conducted. **Aims & Objectives:** To see the effects of ajinomoto (MSG) given to the female albino rats on weight and length of uterine tubes. **Place and duration of study:** This experimental study of 2 weeks duration was conducted in the Department of Anatomy, Shaikh Zayed Postgraduate Medical Institute, Lahore. **Material & Methods:** 45 female rats were divided into three equal groups in a random fashion i.e. one control group A, other two experimental groups B & C. Salt (ajinomoto-MSG) was given in a dose of 0.04 mg/kg body weight to animals of group B i.e. low dose group and 0.08 mg/kg body weight to group C i.e. high dose group for about two weeks after dissolution in 1 ml of distilled water. Animals of Group A i.e. control group were given only equal amount of distilled water without any salt for same days. They were sacrificed on fifteenth day of the experiment. The uterine tubes were approached and carefully dissected out of abdomen. Gross examination was done for any apparent abnormality. Tubes were then weighed and their lengths were taken. Tissue fixation was done & serial 3-5 μm thick sections were stained with hematoxylin/eosin stains for their detailed histological study. **Results:** Uterine tubes of both low dose and high dose experimental groups showed reduced weight and length as compared to tubes of animals of control group. **Conclusion:** Ajinomoto (MSG) causes decrease in weight and length of uterine tubes.

Key words: Monosodium Glutamate (MSG), albino rat, uterine tubes.

INTRODUCTION

Ajinomoto (MSG) is the most commonly used flavor enhancer. East-Asians were using it as a seaweed broth in their cooking for a long time. It was adding meaty taste to their meals. For these reasons it was very popular among them but its proper discovery was made by a Professor of Tokyo University Kikunae Ikeda. He found it as brown colored crystals left after the evaporation of an aqueous solution of this seaweed extract. These crystal were called as "umami" by him in 1907.¹ Later on a method was launched by scientists for large scale production of MSG under Ajinomoto Corporation in Japan.² Hence it got the name of Ajinomoto.

Its major component- glutamic acid is present naturally in protein rich foods such as nuts, legumes, mushrooms, meat, tomatoes, and many dairy products.³ Mostly it is bound with other amino acids but small amount of it is found as free form

enhancing the flavor of food. It is the high content of free glutamate in food which is basically adding or enhancing taste.⁴ Synthetically it was at first produced by fermentation of wheat gluten and then by fermentation of nitrogenous carbohydrates using bacteria.^{5,6} Nowadays its production is done on large scale by the fermentation of nitrogenous carbohydrates with genetically treated bacteria.^{6,7} Initially its use was limited to east- Asians only but later on it gets popularity throughout the world because of flavor enhancing capability. Along with its wide & extensive use multiple questions were being raised from time to time for its safe use. Various health concerns came forward from time to time but despite all, it remained as the most favorite flavor enhancing agent throughout the world.⁸ Its first side effect was observed by scientists in 1957 during conduction of an animal study on retina,⁹ but got unnoticed. The most eye-opening side effects were reported in 1968 by the New England Journal of Medicine as cascade of symptoms including

nausea, vomiting, headache, chest tightness, tingling, facial swellings, increased heart rate etc¹⁰ collectively labeled as “Chinese restaurant syndrome”. This observation directed all researchers and scientists of world to analyze it for its other effects. Multiple researches were carried out on various experimental animals and different organs of the body and it was proved as retinotoxic,¹¹ neurotoxic,¹² nephrotoxic,¹³ hepatotoxic¹⁴ etc. Research work on male reproductive organs also established it as a substance implicating in male infertility by causing testicular hemorrhage and oligoospermia.¹⁵ In female reproductive organs it was found to produce cystic degeneration & atretic follicles in ovaries and also uterine fibroids by interfering with estrogen levels.¹⁶ Present study was thus conducted to rule out its various morphological and histological effects on uterine tubes of female albino rats.

MATERIAL AND METHODS

This study was conducted in the Department of Anatomy, Shaikh Zayed Postgraduate Medical Institute, Lahore.

Study Design: Experimental Study

Study duration: 2 weeks

45 female albino rats were purchased for this study. They were acclimatized for about a week by keeping in animal house of Punjab Post Graduate Medical Institute, Lahore. Feed¹⁷ was provided to them and free access to food and water was confirmed. Their 12 hours light/dark cycles were maintained & then after two weeks animals were assigned into control (A) & experimental groups (B and C) in a random fashion.

Group A (Control):

Control group consisted of 15 rats that received equal amount of distilled water along with MSG free normal chick feed for 14 days.

Group B (Experimental):

Low dose experimental group comprised of 15 rats that were given MSG 0.04 mg/kg/day once a day for 14 days after dissolving in 1 ml of distilled water through orogastric tube.

Group C (Experimental):

High dose experimental group comprised of 15 rats that were given 0.08 mg/kg/day of MSG once a day for 14 days after dissolving in 1 ml of distilled water through orogastric tube.

Dissection was made on the fifteenth day of the experiment. Morphine was used for analgesia and sodium pentobarbitone for anesthesia.¹⁸ Rats were fixed one by one in supine position on dissection board. Abdominal cavities were opened by anterior

abdominal wall incision. Intestines were taken out & uterine tubes were identified (Fig-1). Bilateral uterine tubes were dissected out keeping ovaries and uterus intact. Their gross inspection was done & were later on weighed on electronic weighing machine. Lengths were taken after straightening them with the help of a steel ruler. These tubes were then fixed in formaldehyde for 24-36 hours & their embedded paraffin blocks were prepared after dehydration.¹⁹ Glass slides of 3-5µm thick sections were prepared & stained with Haematoxylin and Eosin (H&E) method²⁰ for histological examination.



Fig-1: Uterine tubes of albino rat attached to ovaries

Statistical analysis:

All observations were entered in computer and SPSS version 21.0 was applied for analysis. Both types of variables (qualitative & quantitative) were compared using Chi-square test & ANOVA.

RESULTS

After sacrifice, when uterine tubes were dissected out carefully and weighed, it was noted that animals of the control group had highest weight of tubes which was 43.07±10.89 mg, while in low dose experimental group B it was recorded 33.67±6.87 mg and in high dose experimental group C it was 28.80±12.34 mg (Table-1). There was statistically significant variation in weight of uterine tubes in the three groups (Table-2).

Groups	Mean weight (mg)	Standard Deviation	Min. weight	Max. weight
Group A (Control)	43.07	10.89	27	57.0
Group B (Low-dose experimental)	33.67	6.87	24.0	45.0
Group C (High-dose experimental)	28.80	12.34	14.0	54.0

Table-1: Mean weight of Uterine tubes of female albino rats

	Sum of squares	Degree of Freedom	Mean squares	Ratio of Variance	P-value
Inter Groups	1577.91	2	788.95	7.44	0.002**
Intra Groups	4452.66	42	106.02		
Total	6030.57	44			

Table-2: Various group comparisons for mean weight (mg) of uterine tubes of female albino rats.

Based on ANOVA

** Highly significant difference (p<0.01)

When weight of uterine tubes was compared group wise it showed that tubes of low dose & high dose experimental groups B and C had significantly more weight as compared to control group A but the difference in weight of both of experimental groups B and C was insignificant (Table-3, Fig-2).

Groups	Groups	Mean Difference	Std. error	P-value
Control Group A	Group B	9.40*	3.76	0.042*
	Group C	14.27*	3.76	0.001**
Experimental (low-dose) Group B	Group C	4.87	3.76	0.406 ⁺⁺

Table-3: Group wise comparison for weight of uterine tubes (mg) of female albino rats.

*The mean difference is significant at the .05 level.

Based on TUKEY'S Test

** Highly significant difference (p<0.01)

⁺⁺ Insignificant difference (p>0.05)

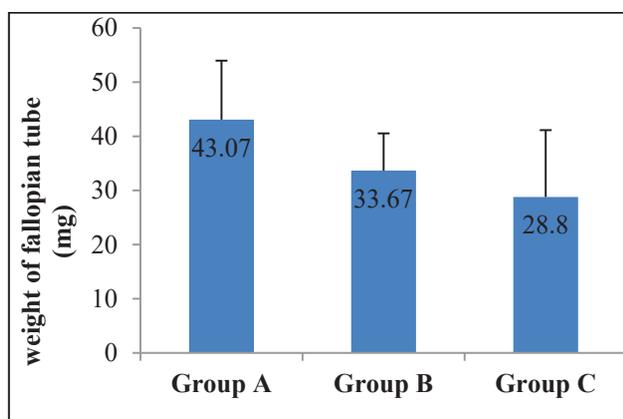


Fig-2: Graph showing comparison of mean uterine tube weight of female albino rats-control & both low & high dose experimental groups.

Similarly when length of uterine tubes was measured using steel ruler from ovarian to uterine ends after dissecting out, the animals of control group had maximum length of uterine tubes

2.28±0.35 cm, while in low dose experimental group B it was recorded 1.90±0.42 cm and 1.84±0.39 in high dose experimental group C (Table-4). The difference in length of tubes between these groups was significant (Table-5).

Groups	Mean lengths -cm	Standard Deviation	Min. length	Max. length
Control Group A	2.28	0.35	1.55	2.85
Experimental Group B (low dose)	1.90	0.42	1.15	2.85
Experimental Group C (high dose)	1.84	0.39	1.15	2.65

Table-4: Mean length (cm) of uterine tubes of female albino rats

	Sum of squares	Degree of Freedom	Mean squares	Ratio of variance	P-value
Inter Groups	1.708	2	0.854	5.693	0.006**
Intra Groups	6.300	42	0.150		
Total	8.008	44			

Table-5: Various group comparison for lengths of uterine tubes

Based on ANOVA

** Highly significant difference (p<0.01)

When individual group comparison for mean length of uterine tubes of albino rats was done, results showed that both experimental groups (low & high dose) B and C had significantly short length of tubes as compared to animals of control group A. However the difference of length between both experimental groups i.e. B and C was insignificant (Table-6, Fig-3).

Groups	Groups	Mean Difference	Std. error	P-value
Group A	Group B	0.38(*)	0.14	0.027**
	Group C	0.44(*)	0.14	0.009**
Group B	Group C	0.06	0.14	0.906 ⁺⁺

Table-6: Various groups comparison for mean length (cm) of uterine tubes of female albino rats

* The mean difference is significant at the .05 level.

Based on TUKEY'S Test

** Highly significant difference (p<0.01)

⁺⁺ Insignificant difference (p>0.05)

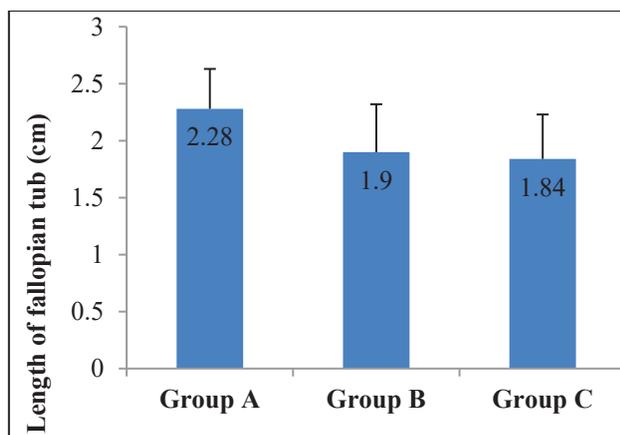


Fig-3: Graph showing comparison of mean length of uterine tubes of female albino rats-control & experimental groups

DISCUSSION

The use of synthetic salt is getting popular in our society. This improves the flavor and makes the food tasty and easily palatable. Ajinomoto is one of the most commonly used such salt; previously associated with Chinese food only but now is a choice of every food company for its products. It is also naturally present in our food as proteins. Although it is in extensive usage but still questions are there for its safe use. Research work is being continuously carried on to observe its effects on various organs.²¹ Present study was also planned to see any effects on morphology and histology of uterine tubes.

When weight of organ was recorded it was noticed that weight of uterine tubes in experimental groups both low & high dose B and C were significantly less than that of control group A with p-values 0.042 and 0.001 respectively (Table-3). This observation of weight reduction of uterine tubes is identical with weight loss of testis in an animal study conducted by Nayantara et al. in 2008. The study was performed on rats to see the effects of ascorbic acid on MSG induced changes in testis & it was observed that there was a loss of testicular weight in animals who received MSG on both short term and long term basis.²² In the present study length of uterine tubes were also taken after dissection and it was observed that the tubes of both experimental (low & high dose) groups B and C were of shorter length as compared to that of control group A with p-value 0.027 and 0.009 (Table-6).

MSG is known to cause reactive oxidative damage by the release of free radicals & both results of the present study i.e. reduction in weight and length of organ could be due to reactive oxidative damage created by free radical release by MSG. It is known

that a cell responds to any inner or outer insult by different responses which include permeability change in membrane, ATP depletion, mitochondrial loss, calcium ions influx or intra cellular accumulation of oxygen derived free radicals. These responses vary according to severity and duration of insulting factors & progress to degeneration.²³ In the present study there could be increased oxidative stress because of release of free radicals leading to irreversible cell injury. This is a possible explanation for the results of present study pertaining to reduction in weight and length of uterine tubes in both low and high dose experimental groups.

CONCLUSION

The results of the present research work indicate that oral administration of ajinomoto for two weeks caused reduction in weight and length of fallopian tubes.

However if this is given in food for a longer period of time it could cause severe effects on major organs and could be a cause of female infertility.

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