# Curative Effect of Alpha Lipoic Acidon Parotid Gland Histopathological Changes in Adult Male Rats with Experimentally Induced Hypothyroidism

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#### **ABSTRACT**

Background:One of the most frequent chronic disorders is hypothyroidism. It affects various organs including the salivary glands. It causes pathophysiological changes including xerostomia. Alpha lipoic acid (ALA) can provide health benefits through its role as a potent antioxidant, metal chelator and reducing agent of the oxidized forms of other antioxidants such as vitamins C and E. Aim: To estimate the influence of hypothyroidism on rat's parotid gland and to illuminate the possible curative role of alpha lipoic acid(ALA). Methodology: Thirty-six adult male rats were haphazardlyarranged into four groups; nine animals in each as follows: Group 1 (control group): 4 rats were givenpurified water (vehicle of carbimazole) for threeweeks, the other 5 rats were given a daily oral dose of 2ml/kg corn oil only (vehicle of ALA) for four weeks after initiation of hypothyroidism. Group 2 (ALA group): were given a daily oral dose of 60 mg/kg ALA dissolved in its vehicle for 4 weeks. Group 3 (hypothyroidinduced group): were givena daily oral dose of 1.35 mg/kg of carbimazole liquified in its vehicle for 3 weeks to induce hypothyroidism. Group 4 (hypothyroid- ALA group): were given a daily oral dose of 1.35 mg/kg of carbimazole for 3 weeks followed by a daily oral dose of 60 mg /kg ALA treatment for the consecutive 4 weeks. The animals were anaesthetized and slaughtered at the end of the experiment. Parotid glands were collected, and the specimens were prepared for light microscopic examination using Hematoxylin and Eosin and Masson's trichrome stains were used, Morphometric analysis using ANOVA and Tukey's post hoc testswas performed. Results: Hypothyroidism caused histopathological changes in the form of general cellular distortion, cytoplasmic vacuolations, cellular infiltration, blood vessels congestion and fibrosis and morphometric changes in the form of a significant increase of collagen fibres in hypothyroid group. Alpha lipoic acid succeeded to improve those changes. Conclusion: Hypothyroidism has destructive effects on the Parotid gland structure. Fortunately, it is recommended to take Alpha Lipoic Acid in hypothyroid cases to help in decreasing such effects.

Keywords: adult rats, parotid gland, hypothyroidism, alpha lipoic acid.

## 1. INTRODUCTION

Hypothyroidism is the most prevalent thyroid disorder, which can be caused by a lack of iodine, thyroid gland tumours, autoimmune disorders, or decreased pituitary gland function <sup>(1)</sup>.

Salivary glands swelling is a common presentation in most of hypothyroid patients<sup>(2)</sup>. Many researchershave found a link between salivary gland dysfunction and autoimmune thyroiditis<sup>(3)</sup>. Moreover, impairment of thyroid function can lead to destructive aberrations in the salivary glands, so people with xerostomia should be evaluated for thyroid functions<sup>(4)</sup>.

Carbimazole which is a methimazole prodrug can inhibit thyroid peroxidase enzyme coupling and iodinating tyrosine deposits on thyroglobulin, lowering T3 and T4 production. The author also stated that after 2, 4, and 6 weeks of use of carbimazole, serum thyroxine, thyroid-stimulating hormone, and thyrotropin-binding inhibitory immunoglobulins were lowered <sup>(5)</sup>.carbimazolewaschosen to initiate experimental hypothyroidism <sup>(6)</sup>.

ALA is a compound of dithiol occurs naturally, which has a protective effect on radiation-induced thyroid gland, salivary gland and intestinal injury <sup>(6) (7)</sup>. ALA administration also decreases weight, thereby return blood TH levels to normal<sup>(8)</sup>. ALA is a co-factor for multi-enzyme complexes in mitochondria that increases glucose absorption and changes the action of numerous signaling molecules and transcript factors <sup>(9)</sup>.

Furthermore, ALA has a lipophilic nature that aids in crossing cell membranes, and it can effectively combat free radicals in both lipid and protein systems when combined with dihydrolipoic acid (DHLA)<sup>(10)</sup>. ALA can improve the body resistance to free radicals <sup>(11)</sup>. Other essential antioxidants, such as glutathione and vitamin C, can also be recycled by ALA <sup>(12)</sup>.

Up to the available literature, determining the specific curative role of ALA for the lethalproperties of carbimazole-induced hypothyroidism on salivary glands, particularly the parotid gland did not receive much condensation.

Therefore, this research aims at the evaluation of ALA as a treatment for the adverse effects of hypothyroidism on the parotid gland.

## 2. MATERIAL AND METHODS

#### 2.1 Chemicals

Carbimazole: SML0931 (Sigma -Aldrich, St Louis Co., MO, USA),5 mg capsules.

Alpha Lipoic Acid: (Thiotacid or Thioctic acid), 300 mg capsules.

They were purchased from Eva-Pharm Pharmatheutical Company - Egypt.

## 2.2 Experimental animals

Thirty-six adult male albino Sprague-Dawley rats of 195 gm body weightwere bought for the experiment. The ratswere purchased from the Animal House at Zagazig University's Faculty of Veterinary Medicine. The animals were kept in a lab setting under supervision. All the trials were carried out in agreement with the strategies of the Institutional Animal Care and the norms of the Ethical Committee of Faculty of Medicine; Zagazig University(ZUIACUC/3/F/118/2020).

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## 2.3 Experimental Plan

The animals were distributed into four groups (nine animals in each) as follows:

**Control Group I:**where therats were further divided into 2 subgroups:

- ➤ Subgroup a:contained 4 rats were given a daily oraldosage of purified water only (the vehicle of Carbimazole) for three weeks through an intragastric tube.
- ➤ Subgroup b: contained5 rats were given a daily oraloraldosage of 2 ml/kg/day corn oil only (the vehicle of ALA) for fourweeks (13) by intragastric intubation.

**ALA group II:**contained the rats were given a daily oraldosage of 60mg/kg ALA dissolved in its vehicle for four weeks through intragastric tube.

**Hypothyroidism- inducedgroup III:** contained rats were given a daily oral dosage of 1.35 mg/kg Carbimazole dissolved in its vehicle for four weeks in order to induce hypothyroidism by intragastric intubation<sup>(14)</sup>.

**Fourth** (hypothyroid-ALA) group: where the rats were given a daily oraldosage of 1.35mg/kg carbimazole dissolved in purified water for three weeks followed by a daily oral dose of 60 mg/kg ALA liquefied in corn oil for the successive four weeks by using intragastric intubation.

## 2.4Methods

# I. Experimental Methodology:

At the beginning of the study, all the used rats were weighed and marked. In the trial termination, the animals were weighed again to be compared to their initial weights, then they were anaesthetized with thiopental (120 mg/kg intraperitoneal injection), and finally decapitated. The excised parotid glands were processed for histopathological examination.

## II. Light Microscopic Examination:

After fixing the samples in 10% Formalin, paraffin blocks were formed. Sections of  $4\mu m$  thickness were prepared and stained with Hematoxylin and Eosin (H & E) and Masson's trichrome stains.

## **I**II. Morphometric study and Statistical analysis:

Image J Analysis Software (Fiji image j; 1.51 n, NIH, USA) was used at the Human Anatomy and Embryology Department, faculty of medicine, Zagazig University. The Areapercentage of collagen fibres ( $\times$ 400) was measured. In all groups, measurements were taken in 10 non-overlapping fields per rat paraffin block. The obtained informationwas collected, charted, statistically analyzed and represented graphically. The Values were exhibited as mean and standard deviation (SD). Multiple evaluations of the groups were performed using one-way (ANOVA) and Tukey's post hoc Tests. The obvious level was set at p  $\leq$ 0.05. Statistical analysis was done by statistical analysis system SAS (Cary, NC, USA).

# 3. RESULTS

## 3.I. Light Microscopic Examination

## 3.1.1 Hematoxylin and Eosin Stain (H&E) Results:

As no obivious difference is observed among control and Alpha Lipoic Acid groups, so, both are represented as control groups on the description of histopathological results. Accordingly, they are named: negative and positive groups respectively.

The control groups showed that the parotid gland is formed of lobules; each lobule contained regular densely packed serous acini and striated ducts with a fine network of interlobular connective tissue (Fig.1a). One layer of high pyramidal epithelial cells around a lumen in each serous acinus, withamong thin fibrous connective tissue septa. The striated ducts among the acini were lined with a single layer of low columnar epithelial cells (Fig. 1b).

In contrast, the hypothyroidism-induced group exhibit that acinar cells lose their normal arrangement withpyknosis. The striated ducts and interlobar ducts are dilated. They become lined with more than one layer of epithelial cells (hyperplasia) and surrounded with wide areas of fibrosis (Fig.1c). Most of acinar cells shows large cytoplasmic vacuolations. Cellular infiltration is observed. There is massive congestion of thee blood vessels with periductal congestion. (Fig. 1d).

On the other hand, the parotid gland tissues in the hypothyroidism-ALA group, regain most of the distinctive general acinar configuration, and acinar epithelial cell vacuolations are greatly reduced. The ducts restored their normal epithelial lining. The fibrosis around the ducts and blood vessels is decreased (Fig. 1e, 1f).

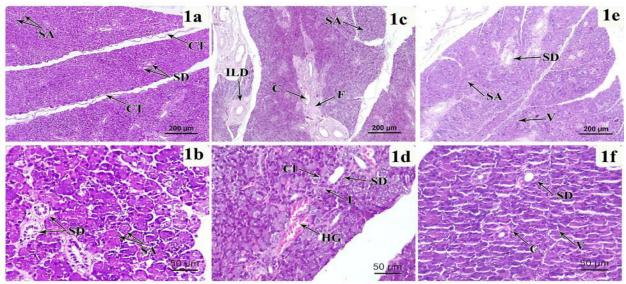


Figure (1):(1a) A Photomicrograph of a section in an adult albino rat parotid gland (control group) shows: the lobules of the gland which formed of serous acini (SA), striated ducts (SD) with a fine network of interlobular connective tissue (CT).(1b) Ahigher magnification of (1a) shows: normal serous acini (SA). Among the acini, striated ducts (SD) withone layer of columnar epithelium are observed. (1c): A Photomicrograph of a section in an adult albino rat parotid gland (hypothyroidism induced- group)shows:disturbed general distribution of the acini (SA), areas of widespread fibrosis (F), dilated interlobar duct (ILD) and periductal congestion (C). (1d): A magnified Photomicrograph of section in an adult albino rat parotid gland (hypothyroidism induced group) shows: striated ducts are dilated (SD) and lined by more than one layer of epithelial cells (L), cellular infiltration (CI) and hemorrhage (HG). (1e): A Photomicrograph of a section in an adult albino rat parotid gland (hypothyroid \_ alpha lipoic acid group)shows: most of the acini looknormally (SA),only limited acini show vacuoles in the cytoplasm (V) and most of the

striated ducts show normal character (SD). (1f): A magnified Photomicrograph of a section in an adult albino rat parotid gland (hypothyroidism -induced &alpha lipoic acid group) shows: few epithelial cytoplasmic vacuolations (V), a little periductal congestion (C) and the striated ducts appear normal shape (SD).

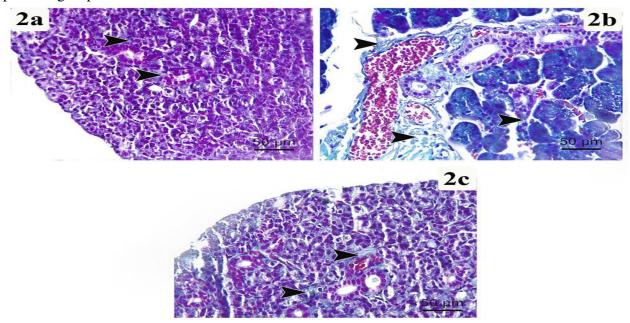
(H and E x 100, Scale bar 200 µm 1a, 1c & 1e)- (x 400, Scale bar 50 µm 1b, 1d & 1f).

#### 3.1.2 Masson's Trichrome Stain Results:

The Control groups revealsfew basophilic collagen fibres surrounding the ducts (Fig. 2a). In contrast, in hypothyroidism - induced group, abundant collagen fibres surrounding the ducts, acini and the blood vessels are observed (Fig. 2b). However, hypothyroid-ALA group exhibits few collagen fibers around the ducts and blood vessels (Fig. 2c).

## Statistical analysis:

Based on Masson's Trichrome results, these findings are proved by the statistical analysis of area percentage of collagen fibres in all experiment groups (Figure 3 & Table. 1). A significant increase is observed inhypothyroidism- induced group in comparison with the control negative and control positive groups. Furthermore, there is a significant decrease in hypothyroidism-induced + ALA group comparison with hypothyroidism- induced group. No significant differences are found comparing the hypothyroid induced + ALA with the control negative and positive groups.



<u>Figure 2:</u> Photomicrographs of Masson's Trichromestained sections of different experimental groups presenting the collagen fibres distribution. (2a)Control group, (2b) Hypothyroidism-induced group (2c) Hypothyroidism-induced + ALA groups. Arrow heads indicate theblue staining of the collagen fibres in the interstitium and around the blood vessels.

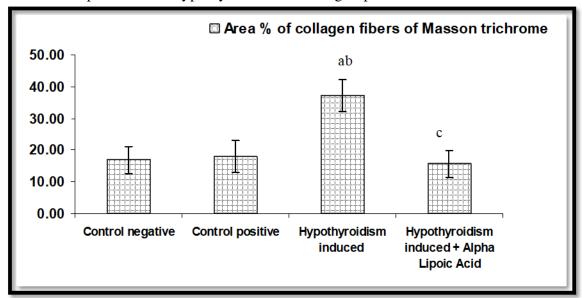
(Masson's Trichrome, scale bar x 50 µm, x400).

Table 1:Area percent of collagen fibers of Masson Trichrome stain for all experimental groups:

Parameter	control negative	control positive	Hypothyroidism induced	Hypothyroidism induced + Alpha Lipoic Acid	P-value
Area percent of collagen fibers	16.830±4.376	17.989±4.987	37.153±5.136 <sup>ab</sup>	15.679±4.258 <sup>c</sup>	0.0001

a-cMeans with different superscripts in the same row are significantly different (\* p<0.05, Tukey HSD test).

<sup>&</sup>lt;sup>c</sup>p<0.05 in comparison with Hypothyroidism induced groups



*Figure 3:* Abar chart demonstrating the analysis of the area % of collagen fibres of all groups.

## 4. Discussion

The goal of this research is to evaluate the curativeability of ALA supplement on the destructive changesproduced by carbimazole-induced hypothyroidism on parotid gland tissue of the experimental animal model.

Salivary glands are essential for maintaining a good oral sanitation and preserving the quantity and quality of saliva <sup>(15)</sup>. Thyroid hormonesare essential to maintain the function and histology of salivary glands <sup>(16)</sup>.

<sup>&</sup>lt;sup>a</sup>p<0.05 in comparison with control negative;

<sup>&</sup>lt;sup>b</sup>p<0.05 in comparison with control positive;

<sup>&</sup>lt;sup>a</sup>p<0.05 in comparison with control negative

<sup>&</sup>lt;sup>b</sup>p<0.05 in comparison with control positive

<sup>&</sup>lt;sup>c</sup>p<0.05 in comparison with Hypothyroidism induced groups

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Hypothyroidism can cause xerostomia; early diagnosis of xerostomia is critical for keeping and helping systemic and oral health <sup>(17)</sup>.

Regarding the light microscopic examination of the control groups, the study verified that the parotid gland was formed of lobules; each lobule contained densely packed serous acini and striated ducts with a fine network of interlobular connective tissue. Each serous acinus was found to be lined by one layer of high pyramidal epithelial cells surrounding a central lumen with thin fibrous connective tissue septa among the acini. Furthermore, the striated ducts between the acini were lined with a single layer of low columnar epithelial cells.

The results come in accordance with the previous researches, it was stated that the parotid gland was composed of lobules, each lobule was formed of regular serous acini, striated ducts and a fine network of interlobular Connective tissue. The authors observed that those acini had a narrow lumen surrounded by a one layer of pyramidal cells with basal round nuclei. In addition, a single layer of columnar epithelium lined the striated ducts (18) (19). Also, a previous study documented that the Histological Examination of sections of salivary gland of control rats revealed normal structure of its elements; acini were spherical, deeply stained and coalesced to each other. The acinar cells appeared pyramidal in shape with basophilic homogenous cytoplasm and clear cellular boundaries (20).

The results of the study did not show any difference in the histological pictures of the parotid glands between positive and negative control groups. This was in agreement Nasr El-Din and Abdel Fattahwho proved that there wasn't statistical change between control and Nigella Sativa oil (NSO) groups in all evaluated statistical parameters (19).

As regards the light microscopic examination of hypothyroidism-induced group, the study revealed that the acinar cells lost their normal arrangement and showed pyknosis. Most of the acinar cells showed large cytoplasmic vacuolations and cellular infiltration. The results revealed that the striated ducts and interlobar ducts were dilated, became lined with more than one layer of epithelial cells (hyperplasia) and surrounded with wide areas of fibrosis. The study also found that the blood vessels were massively congested. Periductal congestion is observed. They were surrounded by massive fibrosis.

In agreement with this study itwas documented that some acinar cells were opaque while other; were cloudy. They had deeply stained nuclei and highly vacuolated cytoplasm, with ductal hyperplasia, cellular infiltration and congested blood vessels<sup>(21)</sup>.

The same resultwas obtained by other authors; the acini in hypothyroidism- induced rats showed indistinct outlines with an increase in the amount of inter acinar & interlobular connective tissue <sup>(2)</sup>. It was noticedthatgiving exogenous T3 to the developing rats cause separation of regular acini in salivary glands<sup>(22)</sup>.

Consistent with those findings, it wasnoted that the administration of propylthiouracil and methimazoleled toreduction in the number and size of the acini of the salivary glands (23) (24) (25). In addition, a previous study stated that pancreatic tissue of hypothyroid rats showed shrunken acini and cytoplasmic vacuolation (26). It wasnoticed that the nuclei of the hypothyroidism-inducedgroup in both parotid and submandibular glands appeared pyknotic, degenerated and

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some show pleomorphism, the cytoplasm was darkly stained  $^{(24)(20)}$ . It was found that the atrophy in serous acinar structure increased with the increase in the duration of hypothyroidism  $(p<0.05)^{(27)}$ .

These degenerative changes observed in the hypothyroid group may be due to the free radicals resulting from hypothyroidism, which have a damaging effect <sup>(23)</sup>. The study come in agreement with another study which stated that the degenerative alterations could be linked to elevated blood endotoxins and oxidative stress, which resulted in capillary damage, exudation, and apoptosis <sup>(28)</sup>. Another work found that those changes might be a result of an inflammatory response, which might restrict the diffusion of nutrients and oxygen to parotid cells <sup>(21)</sup>. Some investigation found that hypothyroidism caused arise in the formation of plasma Malondialdehyde (MDA) (a product of lipid peroxidation) and a reduction in the activity of plasma superoxide dismutase (SOD) (an antioxidant enzyme) this led to the development of oxidative stress in hypothyroidism resulting in cell destruction <sup>(29)</sup> <sup>(30)</sup>.

Regarding the light microscopic examination of hypothyroidism-induced + ALA group, this work showed that parotid tissue regained the normal shape most of the acini and the vacuoles in the cytoplasm were obviously decreased. In addition, the ducts restored their normal epithelial lining with decreased fibrosis around the ducts and blood vessels.

In line with the study results, it wasreported that ALA hadatherapeuticeffectin hypothyroidism-induced toxicity<sup>(31)</sup>. It wasnoticed that ALA causedenhancement the in tubular diameter and germinal epithelial thickness in the testis of hypothyroid rats<sup>(28)</sup>. It wasconveyed that ALA acted as a strong antioxidant, which couldreduce the histological alterations induced by Carbimazole in the tissue of adult male rats' testis<sup>(32)</sup>. It was discovered that taking ALA reduced obesity, restoring blood TH levels and lowering oxidative stress <sup>(8)</sup>. In addition, it was reported that ALA improved human micro vascular endothelial cells dysfunction induced by oxidative stress<sup>(33)</sup>. Another study found that postweaning administration of ALA might be useful in alleviating the developmental hypothyroidism-induced disrupted neurogenesis in Propylthiouracil (PTU)-exposed offspring<sup>(34)</sup>.

These encouraging findings could be attributable to the lipophilic nature of the ALA molecule, which easily crosses biological membranes and so reaches all cellular compartments, improving oxidative stress-related diseases such as atherosclerosis, metabolic syndrome, and diabetes mellitus <sup>(8)</sup>. In addition, ALA has an anti-inflammatory effect, which counteract the inflammatory signs like cellular infiltration and fibrosis <sup>(35)</sup>.

Regarding the area percentage of collagen fibres, the results of this study documented that there was an obvious increase in hypothyroidism- induced group related to the control negative and control positive groups. Furthermore, there was a significant decrease in hypothyroidism-induced + ALA group in comparison with Hypothyroidism- induced group without any statistical change between control negative and control positive groups.

These findings were in line with previous researches which confirmed that Carbimazole-induced hypothyroidism hadharmfulproperties on the parotid tissue of the animal model which include loss of normal acinar distribution, inter-acinar spaces became wide, cytoplasmic vacuoles, congested

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blood vessels, dilated ducts, and excess deposition of collagenous fibres around the blood vessels and inter-lobular ducts (3,4&24)

## 5. Conclusion

The study clarify a great role of ALA as a therapeutic curative agent to the toxic parotid gland histopathological effects that occurred due tocarbimazole-induced hypothyroidism in rats.

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