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Key Words
AGEs • Androgen • Liver biochemistry • Immunostaining

Abstract
Background/Aims: Advanced glycation end products (AGEs) have been related to a wide range of liver disorders including hyperandrogenic states such as the Polycystic Ovary Syndrome (PCOS). The aim of the present study is to evaluate the potential impact of dietary glycotoxins exposure and androgen excess on hepatic histology and biochemistry in an androgenized female rat model. Methods: The study population consisted of 80 female Wistar rats, divided in 3 groups, a group of prepubertal (Group A, n=30) and adult rats (Group B, n=20) that were androgenized via subcutaneous implantation of dihydrotestosterone-containing pellets as well as a group of adult non-androgenized rodents (Group C, n=30). All groups were randomly assigned either to a high-AGE or low-AGE diet for 3 months. Results: Rats fed with a high-AGE diet exhibited significantly elevated levels of gamma-glutamyl transferase (γGT) (p≤0.0002) and indices of AGE immunostaining in liver tissue (p<0.01) when compared to the respective low-AGE group, while aspartate aminotransferase (AST) levels were affected only in non-androgenized animals (p=0.0002). Androgenization per se constitutes an aggravating factor as demonstrated by the elevated γGT levels in adult androgenized animals compared to non-androgenized, independent of diet content (p=0.0002) and by the elevated AST and alanine aminotransferase (ALT) levels in low-AGE subgroups (adult androgenized vs. non-androgenized, p=0.0002) followed by increased immunohistochemical AGE deposition in hepatocytes of the latter categories (p=0.0007). Conclusion: The present study suggests that androgens and glycotoxins may contribute synergistically to distort hepatic physiology and function as observed in hyperandrogenic conditions.

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Introduction

Advanced glycation end products (AGEs) constitute a complex and heterogeneous group of compounds derived from the non-enzymatic glycation of proteins, lipids and nucleic acids [1] produced not only endogenously but also from exogenous sources [2]. Westernized diets are considered a fundamental component of exogenously consumed glycotoxins generated under specific cooking methods [2] and are believed to contribute to total body burden of these compounds and to their tissue deposition [3, 4].

Crucial to the accumulation and biological action of AGEs is the process of their liver degradation (besides kidney) -and predominantly hepatic sinusoidal endothelial cells and to lesser extent Kupffer cells- being involved in their elimination through a scavenger receptor-mediated mechanism [5]. Consecutively, any disturbance in liver function could negatively affect hepatic clearance of AGE adducts whereas exposure to increased glycotoxins could adversely impact hepatic sinusoidal elimination and scavenger function [6].

Besides its pivotal role in compound degradation, liver has the potency to generate low levels of reactive oxygen species (ROS) and, in the case of AGEs, is presumed to contribute to inflammatory processes through the AGE-RAGE (Receptor for AGEs) axis expressed in hepatocytes and hepatic stellate cells [7]. In line with this, several human and experimental studies have shown an association between AGEs with a spectrum of liver disorders from simple steatosis and biochemical aberrations to non-alcoholic steatohepatitis and hepatic cirrhosis either as a result of disturbed liver removal [8-10] or as a possible confounding factor to pathophysiology and progression of these diseases through the induction of oxidative stress and inflammation [7]. Regarding exogenous AGEs, experimental evidence confirms their increased serum and liver tissue deposition and also their ability to promote hepatic oxidative stress both in vitro and possibly in vivo affecting liver biochemistry [11-14].

Increased serum AGEs are observed in hyperandrogenic states such as Polycystic Ovary Syndrome (PCOS) [15] in which higher prevalence of abnormal liver biochemistry is also documented [16, 17]. Although obesity and insulin resistance are the main pathogenetic mechanisms involved, hyperandrogenemia presents an additive adverse factor on liver function of these women [16] though the exact physiological mechanisms are not yet clarified. Interestingly, in hyperandrogenic states such as PCOS, AGEs are found to be positively correlated with testosterone levels implying an interaction between AGE proteins and hyperandrogenaemia [18]. A previous study from our group has shown the negative impact of androgens and nutritional AGEs in the ovarian tissue of female rats in terms of disturbed detoxification system [19].

Taking all these into consideration, the aim of the present study was to explore whether liver per se could be another possible organ affected by the combined exposure to dietary glycotoxins and androgen excess considering that both parameters may have a negative liver impact.

Materials and Methods

Animals

Our study population consisted of 80 female Wistar rats, 30 of which were at the age of 4 weeks (prepubertal stage of life) while the others were at the age of 12-20 weeks (adult life) (Fig. 1). The initial population was divided into 3 main groups: Group A (n = 30) was submitted at 4th week of age (prepubertal) to androgenization, a process previously described by Mannerås et al. [20]. Specifically, 90-day continuous release pellets containing 7.5 mg dihydrotestosterone (DHT) were subcutaneously implanted to animals leading to a daily exposure of 83 μg. This experimental model was selected to mimic both the reproductive and metabolic characteristics of PCOS women, whose DHT levels are found ~1.7-fold higher than those of healthy women [20]. Group A was subsequently randomly divided into two subgroups based on food content; Subgroup A1 (n = 15) was fed commercial chow with low-AGE content (LA) while subgroup A2 (n = 15) was fed commercial chow with high-AGE content (HA).
Group B was composed of 20 older rodents, aged 12-20 weeks (adults), which were submitted to the same previously described process of androgenization. Similarly, a further random division into two subgroups based on the criteria of AGE content, followed androgenization. Subgroup B1 (n = 10) was fed commercial chow low in AGE content while subgroup B2 (n = 10) was exposed to a high-AGE content diet. Finally, group C was composed of 30 non-androgenized animals, which at 12-20 weeks of age, were equally subdivided in two groups. Subgroup C1 (n = 15) followed a low-AGE content diet while subgroup C2 (n = 15) was fed a high-AGE commercial chow.

The animals were housed four to five per cage under controlled conditions (21–22°C, 55–65% humidity, 12-h light/12-h dark cycle) and were given pelleted food and water ad libitum at ELPEN (Experimental-Research Centre, ELPEN Pharmaceuticals, Athens, Greece). Animal care and experimental procedures conformed to the “Guide for the Care and Use of Laboratory Animals” (Department of Health, Education and Welfare, Athens, Greece) and were approved by the Institutional Animal Care and Use Committee.

The diets used were derived from a single standard rat chow (AIN-93G) purchased from Bioserve (Frenchtown, NJ, USA), consisting of 18% protein, 58% carbohydrate, 7.5% fat and 3.73 kcal/g. Regular AIN-93G chow is normally prepared by heating at 190°C for 30 min. Analysis of this preparation was performed as previously described [3]. Briefly, it contained 436.9 ± 88.1 mg CML/100 g protein, 1,179.98 ± 127.90 mg fructoselysine/100 g protein and 302.78 ± 32.82 mg furosine/100 g protein and was considered as an HA diet.

The same rodent mix was also prepared without heating. This preparation was of equivalent macro- and micronutrient and energy content but contained 7.7 ± 2.2 mg CML/100 g protein, 601.01 ± 17.7 mg fructoselysine/100 g protein and 154.22 ± 4.54 mg furosine/100 g protein and was considered an LA diet.

Body weight was monitored weekly. The study was concluded after 3 months and rats were sacrificed with administration of 20 mg/mL xylazine hydrochloride and 100 mg/mL ketamine hydrochloride, under anesthesia with ether, allowing blood sample collection and tissue retrieval.

Hormonal and biochemical assays

Testosterone was measured by enzyme-linked immunosorbent assay (ELISA) using commercially available kits (Calbiotech, CA, USA). Insulin was also quantified using ELISA immunoassays purchased by
Biovendor Laboratory Medicine (Czech Republic). Serum AGE levels (U/mL) were measured by CML-specific competitive ELISA, as described previously. Serum SGOT, SGPT, γGT, glucose, fructosamine and triglycerides were measured by spectrophotometry using commercially available kits (Bayer Hellas).

**Immunohistochemical analysis**

Paraffin-embedded sections of formalin-fixed liver tissue were deparaffinized by xylene and dehydrated in graded ethanol. Sections were treated in 3% hydrogen peroxide in phosphate-buffered saline (PBS) for 15 min and then rinsed in PBS. To increase the immunoreactivity of AGEs, the sections were placed in 500 mL of 0.01 mol/L citric acid–buffered solution (pH 7.0) and microwaved at 500 W for 5 min. After thorough washing, the sections were incubated with normal rabbit serum for 20 min at room temperature to avoid nonspecific binding of the antibodies.

The sections were then incubated overnight at 4°C with the anti-AGE monoclonal antibody 6D12 (0.25 mg/ml stock, dilution 1:50; Research Diagnostics, MA, USA) in PBS containing 1% bovine serum albumin. Immunoreactivity was detected by the streptavidin–biotin–peroxidase method according to the manufacturer’s protocol. The final reaction product was visualized with 3, 3′-diaminobenzidine tetrahydrochloride (LSAB detection kit; Dako, CA, USA). Lung tissue sections from diabetic rats were used as positive controls for AGE antibody. Negative controls (for example, liver tissue in which the primary antibody was substituted with non-immune mouse or goat serum) were also stained in each run.

The evaluation of the immunostained slides was performed blindly and independently by two pathologists. AGE expression was categorized in four levels according to the staining intensity: 1, very weak; 2, weak; 3, moderate; and 4, strong. AGE expression was examined in intrahepatic biliary cells, sinusoidal cells, Kupffer cells and hepatocytes (nuclear and cytoplasmic staining was evaluated separately for the latter cell type). The different cell types were identified by morphology and location by three experienced pathologists (SS, GL, PK). The extent of AGE expression was assessed as the percentage of positive cells in each cell type out of the total number of cells enumerated. Staining intensity was categorized in four levels: 1, faint; 2, weak; 3, moderate; and 4, strong. An H-score based on the percentage of stained cells multiplied by staining intensity was calculated.

**Statistical analysis**

Regarding the examined parameters (body weight at baseline, body weight at 3 months, serum AGEs, AST, ALT, γGT, testosterone, insulin, CRP, serum triglycerides, serum glucose, serum fructosamine, AGEs immunohistochemical score), descriptive statistics were calculated; values are presented as mean±standard error (SE).

Given the deviation from normality, as attested by the Shapiro-Wilk test, non-parametric statistical tests were performed. Specifically, the overall heterogeneity between the six groups was evaluated by the Kruskal-Wallis test. Five meaningful pairwise comparisons were a priori constructed in the study design, namely: A1 vs. A2, B1 vs. B2, C1 vs. C2 (to evaluate the high vs. low AGEs comparison), as well as B1 vs. C1, B2 vs. C2 groups (to address the effects of androgenization per se); all pairwise comparisons were assessed by the Mann-Whitney-Wilcoxon test for independent samples. The level of statistical significance regarding the seven pairwise comparisons was set at 0.05/5 = 0.01 (Bonferroni correction for multiple comparisons) as appropriate, whereas the level of statistical significance regarding the overall heterogeneity remained at the 0.05 level, given that multiple comparisons did not pertain to the latter overall notion. Statistical analysis was performed with STATA/SE version 13 (Stata Corp., College Station, TX, USA).

**Results**

**Comparisons pertaining to the effect of AGEs**

The descriptive statistics regarding body weight and measured serum parameters of the study sample are presented in Table 1. Compared with group A1, group A2 presented with higher AGEs serum levels (as expected, p<0.0001), lower body weight at 3 months (190.7 ± 7.9 vs. 253.7 ± 8.8 g), higher γGT (15.17 ± 1.01 vs. 6.01 ± 0.31 IU/L, p = 0.0001), higher serum testosterone (8.96 ± 0.68 vs. 5.41 ± 0.49 ng/mL, p = 0.0004) and higher CRP (437.4 ± 16.3 vs. 225.5 ± 10.3 μg/mL, p<0.0001). Trends towards higher AST (p =
0.024), ALT (p = 0.036), serum triglycerides (p = 0.014) in group A2 vs. group A1 did not reach formal statistical significance in view of the correction for multiple comparisons (Table 1). Moreover, AGE immunohistochemical H-score was higher in group A2 vs. A1 in sinusoidal (p<0.0001, 162 ± 78.03 vs. 57 ± 54.05; Fig. 2, 3) and Kupffer cells (p=0.0001, 169 ± 66.21 vs. 53 ± 43.16; Fig. 2, 3).

Compared with group B1, group B2 exhibited higher serum levels of AGEs (as expected, p = 0.0002), higher γGT (12.79 ± 0.70 vs. 6.59 ± 0.27 IU/L, p = 0.0002) and higher serum testosterone levels (16.27 ± 2.21 vs. 6.73 ± 0.44 ng/mL, p = 0.0002). Trends towards lower body weight (p = 0.025) in group B2 vs. group B1 did not reach significance given the Bonferroni correction (Table 1). Furthermore, group B2 exhibited higher serum fructosamine levels compared to group B1, with marginal, however, statistical significance (p = 0.010). Interestingly, AGEs H-score was higher in group B2 vs. B1 in all cell types. In detail, B2 cases showed higher AGE H-score in hepatocytes (regarding cytoplasmic staining, p = 0.0011, 87 ± 35.54 vs. 35 ± 21.72; Fig. 2, 3), intrahepatic biliary cells (p = 0.010, 272 ± 119.19 vs. 96 ± 67.05; Fig. 2, 3), sinusoidal cells (p = 0.0002, 191 ± 48.68 vs. 77 ± 50.15; Fig. 2, 3) Kupffer cells (p = 0.0002, 259 ± 77.15 vs. 60 ± 59.43; Fig. 2, 3).

Regarding the group C2 vs. C1 comparison, higher AGE levels (as expected, p<0.0001), higher AST (203.4 ± 12.2 vs. 131.4 ± 6.1 IU/L, p = 0.0002), γGT (10.52 ± 0.31 vs. 3.67 ± 0.17 IU/L, p<0.0001), testosterone (9.42 ± 0.98 vs. 4.39 ± 0.63 ng/mL, p<0.0001), CRP (356.3 ± 11.1 vs. 176.3 ± 10.9 μg/mL, p<0.0001) and serum triglyceride levels (54.6 ± 3.0 vs. 44.6 ± 1.3, p=0.004) were observed in group C2. Trends towards higher ALT (p = 0.044) in group C2 did not reach the Bonferroni-corrected threshold of significance. Once again, AGE immunohistochemical H-score was higher in group C2 vs. C1 in all cell populations and more specifically in hepatocytes (regarding cytoplasmic staining, p = 0.0006, 77 ± 24.17 vs. 38 ± 27.85 and regarding nuclear staining p = 0.0001, 212 ± 88.51 vs. 78 ± 54.68; Fig. 2, 3), intrahepatic biliary cells (p<0.0001, 230 ± 76 vs. 53 ± 44.35; Fig. 2, 3), sinusoidal cells

### Table 1. Measurements of Body weight, serum biochemical and hormonal parameters in the six study groups. § derived from Mann-Whitney-Wilcoxon test for independent samples. † derived from Kruskal-Wallis test (level of statistical significance: 0.05). Values are presented as mean±S.E. P-values in bold denote statistically significant differences

| Parameters                      | Body weight at baseline (g) | Body weight at 3 months (g) | ALT (IU/L) | AST (IU/L) | γGT (IU/L) | Testosterone (ng/mL) | CRP (μg/mL) | Triglycerides (mg/dL) | Serum glucose (mg/dL) | Fructosamine (μM/L) |
|---------------------------------|-----------------------------|-----------------------------|-----------|------------|------------|----------------------|------------|----------------------|----------------------|---------------------|
| Group A1 (mean±SE)             | 97.7±4.7                    | 190.7±7.9                   | 1.0001    | 0.0001     | 0.0001     | 0.0001               | 0.0001     | 0.0001               | 0.0001               | 0.0001              |
| Group A2 (mean±SE)             | 147.4±4.9                   | 206.6±5.9                   | 0.0004    | 0.0004     | 0.0006     | 0.0004               | 0.0004     | 0.0004               | 0.0004               | 0.0004              |
| Group B1 (mean±SE)             | 138.1±4.2                   | 201.6±4.1                   | 0.0004    | 0.0004     | 0.0004     | 0.0004               | 0.0004     | 0.0004               | 0.0004               | 0.0004              |
| Group B2 (mean±SE)             | 148.2±4.2                   | 214.3±5.5                   | 0.0001    | 0.0001     | 0.0001     | 0.0001               | 0.0001     | 0.0001               | 0.0001               | 0.0001              |
| Group C1 (mean±SE)             | 158.2±4.1                   | 221.6±5.2                   | 0.0001    | 0.0001     | 0.0001     | 0.0001               | 0.0001     | 0.0001               | 0.0001               | 0.0001              |
| Group C2 (mean±SE)             | 165.2±4.1                   | 230.8±5.7                   | 0.0001    | 0.0001     | 0.0001     | 0.0001               | 0.0001     | 0.0001               | 0.0001               | 0.0001              |
Fig. 2. Immunohistochemical expression of AGEs in different cell types of female rat liver tissue. Higher AGE immunoreactivity is observed in sinusoidal and Kupffer cells of androgenized 4-week-old rats with a high AGE diet (A2) compared to androgenized following a low AGE diet (A1). Adult (12-20 weeks) androgenized rodents following a high-AGE diet (B2) exhibited stronger AGEs immunoreactivity in hepatocytes, intrahepatic biliary cells, sinusoidal and Kupffer cells compared to adult androgenized animals following a low AGE diet (B1). Non-androgenized high AGE animals (C2) displayed higher AGE staining in all cell populations compared to non-androgenized low AGE ones (C1). Magnification × 200.

(p<0.0001, 203 ± 55.18 vs. 92 ± 36.78; Fig. 2, 3), Kupffer cells (p<0.0001, 226 ± 63.91 vs. 78 ± 56.45; Fig. 2, 3).

**Comparisons pertaining to the effect of androgenization**

Table 2 presents the results of the a priori designed comparisons between the study groups pertaining to the effects of androgenization *per se*.

Groups C1 and C2 presented with significantly lower weight at three months than groups B1 and B2 (p = 0.0008 and p = 0.005, respectively), although the respective differences were not significant at baseline. Regarding γGT, lower levels were consistently noted in groups C1/C2 vs. B1/B2 (p<0.0001 and p = 0.006, respectively, Fig. 4A); AST and ALT levels were lower in C1 vs. B1 group (p = 0.002 for both comparisons, Fig. 4B, C). CRP levels in groups B1/B2 were steadily higher than the respective levels in groups C1/C2 (p<0.0001 and p = 0.002, Fig. 4D). Group B1 exhibited higher levels of serum triglycerides (p<0.0001) than group C1, but these difference was not reproduced at the C2 vs. B2 comparison. Trends towards higher serum fructosamine levels were observed in Group B2 vs. group C2 (p = 0.013). Finally, AGE immunohistochemical H-score was higher in group B1 in hepatocytes (regarding nuclear staining, p = 0.0007, 200 ± 81.64 vs. 78 ± 54.67; Fig. 3).

**Correlation analysis**

At the univariate analysis, serum AST, ALT and γGT levels correlated positively with AGEs (Spearman’s rho=+0.529, p<0.0001, rho=+0.285, p = 0.011 and rho=+0.711, p<0.0001, respectively). Similarly, AST, ALT and γGT levels correlated positively with serum testosterone (Spearman’s rho=+0.434, p = 0.0001, rho=+0.238, p = 0.034 and rho=+0.588, p<0.0001, respectively). Serum testosterone correlated positively with AGEs (Spearman’s rho=+0.687,
p<0.0001) and serum γGT correlated positively with CRP (Spearman’s rho=+0.648, p<0.0001).

Discussion

The main finding of this study relies on the aggravating impact of dietary glycotoxins and androgen excess on liver biochemistry of normal female rats. This was demonstrated by the increased immunohistochemical deposition of glycotoxins in hepatic cell types directly involved in AGE catabolism- such as Kupffer and sinusoidal cells- but also in hepatocytes and intrahepatic biliary cells. The findings are in accordance with the concept that liver is not only an organ of AGE metabolism and degradation [5] but also a site for accumulation of these molecules and probably AGE-related damage [8, 9]. Furthermore, as shown for the first time in the present study, androgen excess has an aggravating impact on several aspects of liver biochemistry parallel to AGE accumulation in hepatocytes of androgenized low-AGE fed animals.

The rats fed with high AGES for three months showed significantly elevated levels of gamma-glutamyl transferase (γGT) when compared to the respective low-AGE group.
an unchanged finding consistently noted in all under study populations- both androgenized and non-androgenized ones (Fig. 4) - possibly related to the increased glycotoxins deposition in intrahepatic biliary cells. Interestingly, this finding was also observed in previous studies indicating an important role of the biliary system in the disposition of AGE [21]. Furthermore, γGT levels were positively correlated with AGEs implying an aggravating impact of high AGE nutrition on this liver enzyme. According to our knowledge, this is the first study to present a possible association between dietary glycotoxins and γGT given that this parameter has not been determined in previous studies of rodents fed with diets high in AGEs [11, 13, 14].

The mechanism via which glycotoxins may exert their negative effect on γGT cannot be clarified from the present study though some assumptions can be proposed. A prominent speculation for elevated γGT levels is that this might reflect a defensive response /adaptation to oxidative stress or inflammatory reactions that allow cells access to increase intracellular glutathione, which is a main protective mechanism against oxidative stress [22]. Indeed, several studies have confirmed the response of γGT to experimentally-induced oxidative stress in several tissues [22]. As AGEs are potent triggers for ROS production in the liver [11, 12], it could be speculated that nutritional exposure to glycotoxins may represent an inflammatory environment, on the basis of elevated serum and hepatic AGEs deposition documented in the present experiment, to stimulate γGT production as defensive mechanism.

Regarding androgen exposure, the current study reveals that androgenized adult animals exhibited significantly elevated γGT levels compared to non-androgenized ones, independent of diet type suggesting that androgens may have an additional unfavorable effect on this enzyme. This finding is in accordance with human studies showing elevated levels of γGT in hyperandrogenic women [16] and furthermore, γGT levels were positively correlated with testosterone implying a possible pathogenetic association between these parameters.

The levels of other liver enzymes- aspartate (AST) and alanine aminotransferase (ALT) - signifying hepatocellular damage are also affected. Specifically, AST levels were found significantly higher in non-androgenized rats fed a high-AGE diet compared to low-AGE ones in accordance with the increased glycotoxins deposition in hepatocytes -both nuclear and cytoplasmic staining- of these animals. This finding agrees with previous studies showing
increased susceptibility of hepatocytes to the detrimental effects of AGEs from patients with varying degrees of liver impairment [10, 21]. Furthermore, a trend towards higher AST levels was observed in the group of androgenized prepubertal HA rodents compared to LA ones without reaching statistical significance (Fig. 4B).
Concerning the liver enzyme ALT, the impact of high-AGE diet seems to follow the same direction in the above populations though without statistical significance. The androgenized adult rats exhibited no difference for both enzymes which can be attributed to the relatively small number of animals in these subgroups (Fig. 4C).

Elevated ALT and AST levels have been detected in PCOS women [17, 23]. In this study, the population of androgenized adult animals following a low-AGE, but not a high-AGE diet exhibits significantly higher levels of both aminotransferases when compared with the non-androgenized rats. Interestingly, this is in accordance with the immunohistochemical evidence of increased glycotoxins deposition in hepatocytes (nuclear staining) of androgenized animals shown for the first time in the present study. These findings clearly indicate that androgens per se play an important role in hepatic biochemistry and histology that is more pronounced in the low-AGE diet; nevertheless, in the high-AGE diet androgens do not seem to have an additive effect. This may indicate that the effect of diet on these liver enzymes prevails over the effect of androgenization or it may indicate a compensatory mechanism of action.

Furthermore, the observation of a positive correlation of ALT and AST with androgens seems to be compatible with human studies of hyperandrogenic women showing an association between biochemical and clinical hyperandrogenaemia with elevated ALT levels [16, 17]. The exact mechanism via which androgen may modify liver biochemistry is yet undetermined. Their role seems to be multifactorial and is believed to be implicated in multiple steps of hepatic physiology. Their impact may be apparent even in intrauterine stages of life by modifying liver metabolism and signaling [24, 25].

The present data of dietary AGEs on liver biochemical profile come in disagreement with recent studies [14]. Patel and colleagues have demonstrated higher levels of aminotransferases in rodents following a regular AGE diet compared to a high AGE one, probably as a consequence of adaptation of liver anti-oxidative mechanism(s) against AGEs. Of note, these measurements were observed at the 39th week of experiment while our findings were on the 12th week when defensive mechanisms could hypothetically be too early to be adapted. It is likely that due to the shorter duration of our study the compensative mechanisms against oxidative and inflammatory status provoked by AGE administration were not yet developed. Furthermore, our study provides evidence that androgen may be a contributing factor to sustain the oxidative stress leading to a further suppress of the anti-oxidative system.

Another interesting finding is the metabolic phenotype of rats with respect to dietary modification. Elevated fasting insulin levels with normal glucose and fructosamine levels were observed in non-androgenized and androgenized prepubertal HA-fed animals. In adult androgenized rats the impact of dietary AGEs in fasting insulin levels is less prominent, however, the marginal statistical significance of fructosamine levels may imply a distorted metabolic profile. Triglycerides seem to follow the same pattern with a trend towards elevated levels in androgenized prepubertal rats. These metabolic derangements may reflect an indirect impact of diet on liver biochemistry that is independent of the degree of obesity as the high-AGEs diet was related to a lower body weight. Furthermore, given that serum AGEs are known to be associated with insulin resistance [26] this abnormal metabolic profile could imply the development of disturbed insulin action that is known to be pathologically related to hepatic disorders and activation of inflammatory liver pathway [27].

A statistically significant elevation of C-reactive protein (CRP) levels was observed in rats following a high-AGE diet compared to those exposed to a low-AGEs one (apart from androgenized adult group) (Fig. 4D). Furthermore, androgenization per se seems to have an aggravating impact as animals androgenized at adulthood compared to the non-androgenized control population show higher CRP levels independent of diet. AGEs are pro-inflammatory molecules shown to indirectly stimulate hepatic CRP production in vitro via stimulation of pro-inflammatory molecules (interleukin-6, interleukin-1 beta) by monocytes that upregulate CRP production in hepatocytes [28]. Furthermore, in an experimental study
of cultured Hep3B cells AGEs were shown to directly upregulate CRP mRNA levels via Rac-1 activation [29]. It is likely that their increased serum and liver deposition presented in this study may trigger and further deteriorate a similar biological effect *in vivo*. A similar pathogenic mechanism may underlie the aggravating impact of androgenization in inflammation as indicated by elevated CRP levels. It is known that in hyperandrogenic states such as in women with PCOS chronic low-grade inflammation is observed as reflected by increased circulating levels of inflammatory mediators [30]. Hyperandrogenemia can induce inflammation in reproductive-age women and one possible mechanism is through the release of cytokines such as tumor necrosis factor alpha (TNF-α) from mononuclear cells in a receptor-dependent fashion [31], or through enhanced leukocytic reactive oxygen species (ROS) generation [32]. ROS-induced oxidative stress, also triggered by AGEs [33], is a known activator of the nuclear factor-kappaB (NF-κB), a key regulator of inflammation [34]. Activation of this transcription factor presents an intersection point of both androgens and AGEs excess in the induction of inflammatory responses. Therefore, it is speculated that exogenously derived AGEs in combination with increased androgen exposure aggravate local inflammatory processes leading to augmented oxidative stress and release of inflammatory mediators including CRP release from liver.

Interestingly, in this experimental study, the inflammation marker CRP was also positively correlated with γGT. The elevated CRP levels in conjunction with the increased levels of γGT observed in the hyperandrogenic animals compared to the nonandrogenized ones may have implications beyond liver pathology given that these parameters are related to oxidative stress [35].

Another finding of the present study is that all rats fed with high AGEs for three months displayed significantly elevated levels of testosterone when compared to the respective low-AGE group. This was a consistent finding noted in the study populations - both androgenized and non-androgenized animals. This finding is confirmatory of our previous experimental studies presenting a distorted hormonal profile of elevated testosterone levels in normal female rats following a diet rich in glycoxins [3]. The pathogenic interference between AGEs and androgens is not yet clarified but may be related to the increased distribution of AGEs and RAGE in theca cells of female rats exposed to dietary glycoxins shown in previous studies [3]. Theca cells are responsible for the androgen production in the ovarian tissue and AGEs may modulate steroidogenesis, therefore a functional link between the increased deposition of AGEs in the ovary and the increased levels of serum testosterone cannot be excluded. Supportive of such interplay is the positive correlation between androgen and serum AGEs levels observed in PCOS women [18]. The differential effects on liver function between endogenous androgens (possibly stimulated by dietary AGEs) and exogenous androgens, is the aromatization of endogenous as opposed to non-aromatizable exogenous androgens. The experimental model of androgenization could indirectly favor the development of a PCOS phenotype exhibiting the hormonal characteristics (hyperandrogenic) of the syndrome. However, the observation of increased androgen levels in non-androgenized animals converges to an additive effect of glycoxins possibly through stimulation of ovarian androgen production.

This study has some limitations. Although the sample size was satisfactory some differences did not reach statistical significance especially in smaller size subgroups such as the androgenized in adult life population. Additionally, hepatic insulin resistance indexes indicative of the metabolic profile of the exposed animals were not evaluated. Finally, other parameters such as liver organ weight or body weight composition were not estimated.

**Conclusion**

The present study shows that high dietary AGEs induce apparent serum and liver tissue accumulation in all experimental groups. This is probably related to disturbed liver biochemistry that is observed throughout the study predominantly by elevation of γGT levels.
compatible with the immunohistochemical evidence of increased deposition in intrahepatic biliary cells. Androgenization per se is shown for the first time to have an additive negative liver impact that is more pronounced for aminotransferases in the low-AGE diet -parallel to the immunohistochemical evidence of increased hepatocyte AGE deposition- and more apparent for γGT levels, independent of diet.

Disclosure Statement

The authors declare no financial conflict of interest.

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