

Endotoxins enhance hepatocarcinogenesis induced by oral intake of thioacetamide in rats *

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Abstract

AIM To clarify whether endotoxin is of pathogenic importance for hepatocarcinogenesis, or the increased cancer risk results solely from the cirrhotic process.

METHODS The rat model of hepatoma was treated by the intake of 0.03% thioacetamide in drinking water for six months. During induction of hepatoma, rats were additionally treated with splenectomy and/or lipopolysaccharide administration. The liver nuclear DNA index and proliferation index were quantitatively analyzed by flow cytometry. Hepatic histology was examined with light and electron microscopes. Plasmic endotoxin concentration and γ -glutamyl transpeptidase activity were measured, and hepatoma incidence was recorded.

RESULTS Thioacetamide induced cirrhosis and hepatoma in Wistar rats with histology or regenerative nodule, fibrosis and neoplastic foci were quite similar to the pathogenic process of human cirrhosis leading to hepatoma. In comparison with TAA controls (DNA index: 1.15 ± 0.21), exo-endotoxin increased the DNA index by 7.8% (1.24 ± 0.25 , $P < 0.02$) and hepatoma rate by 16.7. Splenectomy-induced enteric endotoxemia increased the DNA index by 25% (1.44 ± 0.15 , $P < 0.01$) and hepatoma rate by 33%. A summation of the effects of these two factors increased the DNA index by 36% ($P \leq 0.01$) and hepatoma incidence by 50%, moreover, the level of endotoxemia showed a close relation with DNA index ($r = 0.96$, $P < 0.01$), as well as with the occurrence rate of hepatoma ($r = 0.00$, $P < 0.01$). Histological findings further verified such alterations.

CONCLUSION Lipopolysaccharide administration and/or splenectomy-induced enterogenic endotoxemia may enhance rat hepatocarcinogenesis induced by oral intake of thioacetamide.

INTRODUCTION

Endotoxemia is often associated with hepatic injury and cirrhosis^[1]. Patients with hepatic cirrhosis or hepatocarcinoma often present a 3 to 8-fold high systemic endotoxemia above normal^[2]. If cirrhosis is established, there is a 200-fold increased risk for development of hepatocarcinoma^[3]. It is unknown whether endotoxin is of importance in hepatocarcinogenesis. Recognized etiologic or predisposing factors include chronic liver disease caused by hepatitis B, C and D; hepatotoxins such as aflatoxin and alcohol; and cirrhosis of any cause^[4]. The pathogenetic mechanisms involved in human hepatocarcinogenesis remain largely unsolved.

Numerous animal models of cirrhosis have been developed^[5,6]. Continuous administration of thioacetamide (TAA) in drinking water to rats produced cirrhosis in 4 months and hepatocarcinoma in 6 months^[7]. The histology of the TAA model was more similar to human cirrhosis than that of the CCl₄-treated rats^[8]. Wistar rats were used in the TAA model.

Using a TAA-induced rat model of cirrhosis leading to hepatocarcinoma, we tried to prove the hypothesis that lipopolysaccharide (LPS) administration and/or splenectomy-induced enterogenic endotoxemia may play an important role in the pathogenesis of liver tumors. The present study was designed to investigate the histologic changes of hepatocarcinoma in the TAA models treated with LPS and/or splenectomy; DNA aneuploidy and incidence of hepatocarcinoma in various groups; and the relationship between endotoxemia and hepatocarcinogenesis.

MATERIALS AND METHODS

Animals and treatment

Female Wistar rats (Experimental Animals Center of Shanxi Medical University) weighing $125 \text{ g} \pm 9 \text{ g}$, were housed in wire-bottom cages under a 12-hour light/dark cycle and fed with a balanced pellet diet

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ad libitum. Animals were randomly assigned to five groups: sham group ($n = 5$) as an untreated control receiving tap water ad libitum; TAA control group ($n = 6$), given 0.03% w/v thioacetamide (TAA, purity >99%, Shanghai Central Chemical Factory) in drinking water; TAA+ST group ($n = 6$) splenectomized (ST) a week before the experiment; during the last two months, 0.8mg% lipopolysaccharide (LPS, from *-Escherichia coli*-sterotype 055:B5, Sigma) added to drinking water containing 0.03% TAA for both TAA+LPS group ($n = 6$); and TAA+ST+LPS group ($n = 6$). Without splenectomy, animals underwent midline laparotomy and spleen manipulation as a sham surgery. All operations were performed under light ether anesthesia and sterile conditions. At the termination of a 6-month course, rats were anesthetized with ethyl ether and blood from abdominal aorta was collected in heparinized tubes for measurement of plasmic endotoxin concentration and glutamyl transpeptidase (GGT) activity.

Analysis of endotoxin and γ -glutamyl transpeptidase activity in plasma

Endotoxin in plasma was determined by means of a chromogenic limulus amoebocyte lysate (LAL) assay^[9]. Limulus kit (Shanghai Yihua Clinical Medical Biotech Co.) was used for endotoxin-specific quantitative measurement in this study. Syringes, needles and glasswares were made pyrogen-free with dry heat (200°C for 4 hours). The plasma was pre-treated with perchloric acid (PCA) for removal of the possible inhibitors to LAL assay. Briefly, 0.1ml of 0.18mol/L NaOH was added to a 0.1ml of test plasma and incubated at 37°C for 5min. Then 0.1ml of 0.32mol/L PCA was added for an additional 10min incubation. The precipitate formed was dissolved by addition of 0.2ml of 0.18mol/L NaOH with vigorous mixing. The solution was centrifuged at 2000rpm for 10min. Then 0.05ml of the supernatant diluted with 0.05ml of 0.4mol/L Tris-HCl (pH 8.0) was added to 0.1ml of limulus amoebocyte lysate (LAL) in pyrogen-free cuvette, and incubated in a thermostatic bath at 37°C for exactly 30min. The synthetic chromogenic substrate 0.05ml, *n*-tert-butoxycarbonyl-*L*-leucyl-*L*-glycyl-*L*-arginine-*p*-nitroaniline (Boc-Leu-Gly-Arg-PNA) was added to the lysate and incubated for 5min. The reaction was stopped with 0.5% sodium nitrite. The amount of PNA released from the substrate was detected after diazo-coupling by adding 0.5ml of 0.05% ammonium sulfate and 0.5ml of 0.05% *n*-*l*-naphthylethylenediamine dihydrochloride. Absorbance was read at 545nm on 721A spectrophotometer, using distilled water as the 0 absorbance value. A standard curve was plotted with the known endotoxin concentration for each run.

Simultaneously, saline replacing plasma was as reagent control.

γ -glutamyl transpeptidase (GGT) activity in plasma was detected according to the method as described by Xu KC^[10]. In brief, 0.1ml of plasma was added to a 0.5ml of 10mmol/L substrate, γ -*L*-glutamyl- α -naphthylamide (Shanghai Third Reagent Factory), and incubated at 37°C for 120min. The amount of α -naphthylamide released from the substrate was detected after adding 10ml of 0.2% sulfanilic acid and 0.004% sodium nitrite. The absorbance was read at 500nm on a 721A spectrophotometer, using sample control of adding plasma after 120min incubation as the 0 absorbance value. A standard curve was plotted with the known α -naphthylamide concentration for each run.

Light and electron microscopy

One half of the left lobe of the liver was resected, fixed in 10% neutral-buffered formalin and embedded in paraffin. Paraffin sections were stained with hematoxylin-eosin (HE) for light microscopy. For electron microscopic study, the selected areas with hepatocarcinoma were cut into small pieces and fixed in 2% glutaraldehyde for 2 hours. After postfixation in 1% osmium tetroxide, the fragments were dehydrated in a graded alcohol series and embedded in Polybed. They were trimmed into ultrathin section and stained with uranyl acetate and lead citrate, and observed under an electron microscope at 100KV.

Flow cytometry

According to Zuo LF^[11], with slight modification. The specimens for flow cytometry were prepared immediately after removal of liver. About 2g tissue was mechanically minced in 5ml of 0.05% collagenase (type IV, Sigma) and then through a 200 mesh filter. After washing with 0.1mol/L phosphate buffer solution (PBS), the well-dispersed hepatocytes were resuspended and fixed in 70% ethanol overnight, the cells were stained with propidium iodide reagents (propidium iodide 50mg/L, RNase 20mg/L, and Triton-X-100 1%) at 4°C for 30min. Chicken erythrocytes were used as internal controls. Flow cytometry was performed on a Fluorescence Activated Cell Sorter (FACS) 420 cytometer (USA). Data were analyzed on a compatible computer with HP-300 consort 30. A total of 10^4 cells were examined for each sample, DNA index (DI) expressed as a relative DNA content, was calculated by dividing mean channels of G0/1 peak in the treated rat by that of sham group. The coefficient of variation (CV) of G0/1 peak of the DNA ploid population was expressed by normalized standard deviation ($CV = \text{standard deviation} / \text{mean peak position} \times 100$). CV was always less than 5%. For the

DNA ploid criteria, it was classified as DNA-diploid if $DI=1.0\pm 2CV$, and DNA-aneuploid if $DI\neq 1.0\pm 2CV$. Cell cycle parameters were analyzed by DNA cell cycle analysis software Sum of Broadened Rectangles Model, and the cell proliferation index (PI) was expressed as the percentage of cells in S plus G2M phases.

Statistical analysis

All results were expressed as $\bar{x} \pm s$. Data were analyzed by Student's *t* test and multiple regression. *P* values <0.05 were considered statistically significant.

RESULTS

The oral intake of TAA in drinking water induced cirrhosis and, in some rats with hepatocarcinoma with null mortality. TAA-treated animals showed a progressive decline in body weight, particularly in the first month (data not shown). On the contrary, the ratio of liver weight to body weight were obviously higher in all TAA treated animals ($P<0.01$), compared with the sham group. The hepatocyte proliferation index was fairly high in the TAA+LPS or TAA+ST groups, and the highest in the TAA+ST+LPS group ($P<0.05$) compared with the TAA-model control (Table 1).

Table 1 Hepatocyte proliferation and liver weights

Groups	<i>n</i>	PI	Ratio of liver weight to 100g body weight
Sham	5	0.25 ± 0.032	4.32 ± 2.6
TAA control	6	0.33 ± 0.047	8.38 ± 2.0^b
TAA+LPS	6	0.35 ± 0.045	7.35 ± 1.3^b
TAA+ST	6	0.35 ± 0.077	7.40 ± 1.5^b
TAA+ST+LPS	6	0.45 ± 0.039^a	7.28 ± 1.5^b

^a $P<0.05$, vs TAA control, TAA+LPS and TAA+ST groups respectively. ^b $P<0.01$, vs sham group. PI, proliferation index. Rats were treated as described in Materials and Methods.

Endotoxemia level and GGT activity in systemic circulation

The TAA-treated rats with splenectomy or administration of 0.8mg% LPS (approximately corresponding to 50 μ g/100 g body weight daily) in drinking water during the last two months of the study showed increased endotoxemia level ($P<0.05$), compared with the sham group. The blood endotoxin level in the TAA+ST+LPS group further mounted up ($P<0.05$), compared with the sham group or the TAA control group. However GGT activity was markedly increased in the blood of the TAA-treated rats with splenectomy plus LPS administration ($P<0.05$), compared with solely TAA-treated group (Table 2).

Table 2 Endotoxemia levels and GGT activities in plasma

Groups	<i>n</i>	Endotoxin (EU/100ml)	GGT (U/100ml)
Sham	5	95 ± 15	17 ± 7
TAA control	6	108 ± 12	226 ± 33^d
TAA+LPS	6	118 ± 12^d	236 ± 44^d
TAA+ST	6	130 ± 31^d	266 ± 127^d
TAA+ST+LPS	6	130 ± 18^{ad}	442 ± 99^{abcd}

^a $P<0.05$ vs TAA control; ^b $P<0.05$, vs TAA+LPS group; ^c $P<0.05$, vs TAA+ST group; ^d $P<0.05$, vs sham group. Rats were treated as described in Materials and Methods.

DNA index and hepatoma rate

The livers of rats in the TAA control group showed many superficial yellowish macronodules and one case of visible hepatocarcinoma out of six rats. With LPS administration and/or splenectomy, numerous micronodules appeared in livers and more cases of noticeable tumors. Moreover much more hepatocarcinomas were observed under the light microscope. DNA index as an indicator of the severity of malignant lesions were stepped up in the TAA models additionally treated with LPS and/or splenectomy. Therefore, they have significant differences between groups ($P<0.02$), particularly in the splenectomized TAA-treated rats with oral LPS administration (Table 3).

Table 3 DNA ploidy and incidence of hepatoma in various groups

Groups	<i>n</i>	DNA index	Rate of DNA aneuploid%	Incidence of hepatoma			
				Visible Under microscopy			
				(%)	<i>n</i>	(%)	<i>n</i>
Sham	5	1.0	0	0	(0/5)	0	(0/5)
TAA control	6	1.15 ± 0.21	50	17	(1/6)	50	(3/6)
TAA+LPS	6	1.24 ± 0.25^a	50	33	(2/6)	50	(3/6)
TAA+ST	6	1.44 ± 0.15^{ab}	100	50	(3/6)	67	(4/6)
TAA+ST+LPS	6	1.56 ± 0.07^{abc}	100	67	(4/6)	67	(4/6)

^a $P<0.02$, vs TAA control group; ^b $P<0.02$, vs TAA + LPS group; ^c $P<0.02$, vs TAA + ST group. DNA index=mean channels of G0/1 peak in the treated rat \div mean channels of G0/1 peaks in the normal controls. 10^4 cells of a rat liver were examined by flow cytometry. Rats were treated as described in Materials and Methods.

Histology

Under the light microscope, no noticeable changes were observed in untreated control rats (Figure 1). The livers of the TAA models with or without LPS and/or splenectomy had numerous micronodules of various sizes in all lobes. The nodules were separated by markedly proliferated collagen fiber, forming pseudolobules. Proliferation of spindle cells, probably fibroblasts, was recognized around such hyperplastic nodules (Figure 2). In the transforming

hepatocytes, atypical hyperplasia with nuclear heterochromatism and basophilic cytoplasm were seen mainly in perivenular and in periportal areas. Nuclear atypism was also seen in the hepatocytes. Moreover, disarrangement of the hepatic cords became apparent and formed considerable pseudoglandular (acinar) structures which were transformed from original liver parenchymal cells (Figure 3). Highly malignant lesions were observed in the TAA

+ST+LPS group, in which atypism was more apparent and formed a “muroid lake” with a float of small atypical cells (Figure 4).

Electron microscopic investigation clearly showed that the disarrangement of the altered hepatocytes and nuclear atypism were very apparent, together with less cytoplasm containing very few organelles (Figure 5). Collagen fibrils were markedly present around the neoplastic cells (Figure 6).

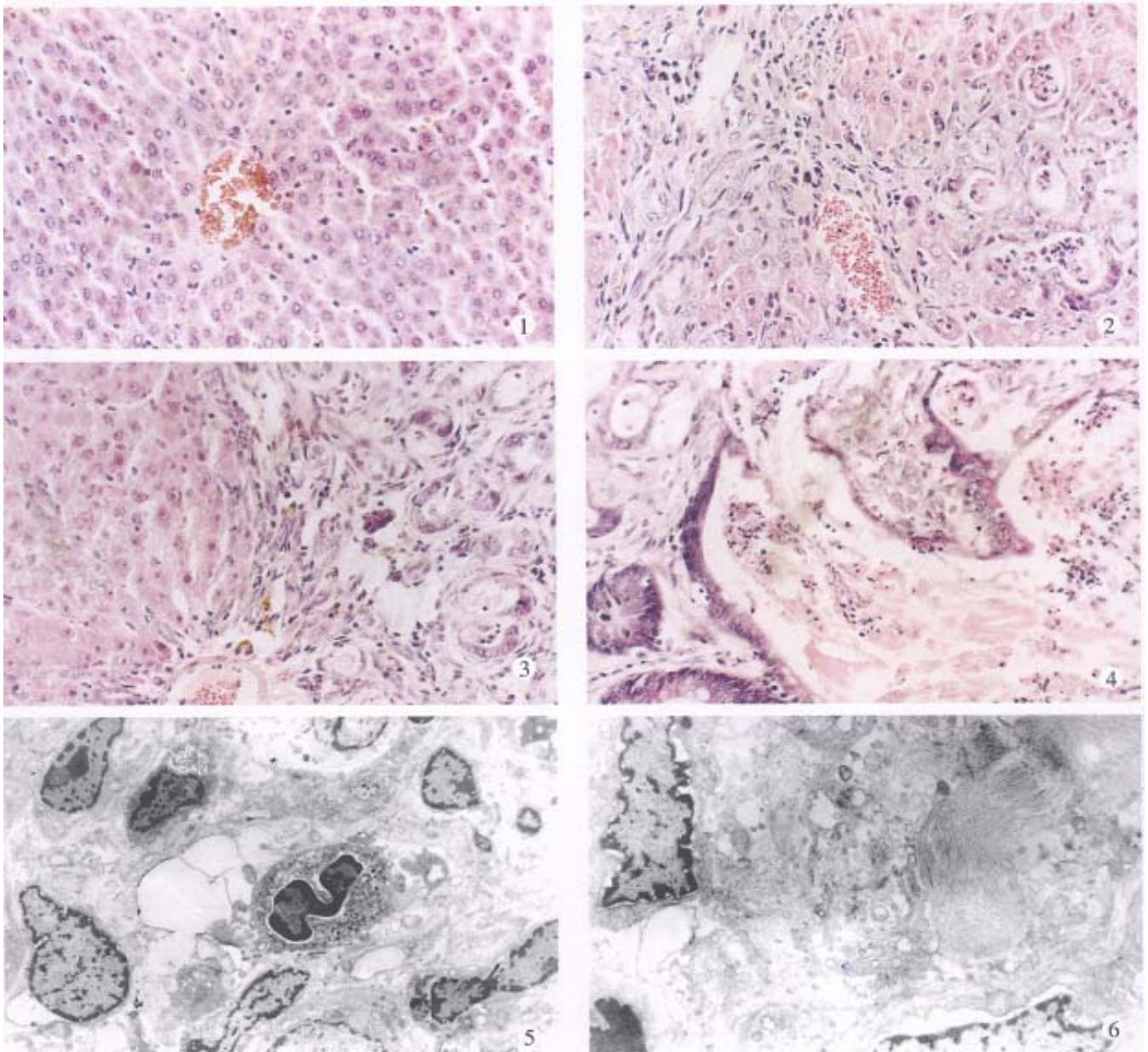


Figure 1 Section of liver from an untreated control (sham) rat. HE×20.

Figure 2 Section of liver from a TAA control rat treated as described in Materials and Methods. The nodules show the transforming hepatocytes and the hepatocarcinoma. The spindle fibroblasts are recognized around such hyperplastic nodules. HE×20.

Figure 3 Section of liver from a rat in TAA+ST group. Pseudoglandular structures and spindle cells are observed in perivenular areas. HE×20.

Figure 4 Section of liver from a rat in TAA+ST+LPS group. The disorganized and atypical hepatocytes formed a “muroid lake” with floating of small cells. HE×20.

Figure 5 Ultrastructure of hepatocellular carcinoma from a rat in TAA+ST+LPS group. Uranyl acetate and lead citrate staining ×400.

Figure 6 Ultrastructure of cirrhosis from a rat in TAA+ST+LPS group. The bundles of collagen fibrils were present with atypical hepatocytes. Uranyl acetate and lead citrate staining ×800.

DISCUSSION

TAA-induced cirrhosis and hepatocarcinoma has been observed in Wistar rats. The results of the present study demonstrated for the first time that endotoxin may play an important role in the development of hepatocarcinoma in rats induced by continuous administration of 0.03% TAA in drinking water.

TAA is hepatotoxic owing to effects on DNA, RNA, protein synthesis and GGT activity, through which it induces cirrhosis and hepatocarcinoma^[1,8]. After 6 months of chronic TAA administration, the liver exhibited an external appearance of cirrhosis characterized by the presence of numerous yellowish nodules and whitish hepatocarcinoma. The microscopic study showed that micronodular cirrhosis with hepatocarcinoma resembles human cirrhosis with hepatocarcinoma in both its biological and morphological aspects in close agreement with other studies^[5-7]. Table 3 shows that TAA increases nuclear DNA content by 15% in rat liver, producing 50% cases with aneuploidy and 50% cases of hepatocarcinoma seen under microscopy was a suitable model for evaluating the effects of endotoxin on hepatocarcinogenesis.

DNA index representing a ratio of the DNA content in the treated rat to the mean DNA content of the diploid normal rats indicate precise malignancy of hepatocarcinoma, together with aneuploidy rates in various groups, which were highly correlative to the incidence of hepatocarcinoma (the former $r = 0.98$, $P < 0.01$; the latter $r = 0.96$, $P < 0.01$). Moreover, the histologic studies with light and electron microscopes well confirmed the interesting findings. In the hepatocyte, clonal perpetuation of accumulated DNA aneuploid affecting growth regulatory genes eventually leads to a continuation of severe progressive abnormalities characterized by dysplasia, formation of adenoma and, finally, cancer^[13]. In addition, plasma GGT activity, an enzyme of fetal hepatocytes, was a sign of poor differentiation and correlated with cancerous change ($r = 0.93$, $P < 0.05$) and the degree of endotoxemia ($r = 0.88$, $P < 0.05$).

Like Kupffer cells, splenic macrophages have a potential for endotoxin clearance by way of phagocytosis. In this study, rats undergoing splenectomy showed reduced endotoxin clearance and increased enterogenic endotoxin levels in systemic circulation by 37% ($P < 0.05$). And rats with cirrhosis after LPS administration at a dose of 50 µg/100g body weight daily for two months increased their endotoxemia by 24% ($P < 0.05$), compared with the normal controls. When the rat endotoxemia increased, inhibition of hepatic DNA synthetic enzymes by endotoxin^[12] resulted in a decrease of liver weights and formation of micronodules in comparison with the TAA control rats with

macronodules. In addition, splenic macrophages have an ability to generate tumor necrosis factor- α and play an important role in immunosurveillance^[13,14]. Therefore the down-regulation of Kupffer cell activity after splenectomy may have deleterious effects on host defenses^[15].

This study shows that endotoxemia from orally administered LPS, during the last two months, and/or from splenectomy before commencement of the experiment, significantly enhanced hepatocarcinogenesis ($P < 0.02$), based on the alterations of the DNA index. It seemed that the TAA model with splenectomy combining LPS administration was more deleterious to the development of hepatocarcinoma in rats than the TAA model treated with ST or LPS solely. In addition, it was noticed that the blood endotoxin levels in rats were significantly correlated with malignancy ($r = 0.96$, $P < 0.01$) and the incidence ($r = 0.99$, $P < 0.01$) of hepatocarcinoma. If so, it may be of importance in the pathogenetic process of hepatocarcinoma and in clinical therapeutic strategy. However, patients with cirrhosis presenting relatively high endotoxemia^[2] may have higher risk for development of hepatocarcinoma epidemiologically^[3], but what are the possible mechanisms involved in the pathogenesis and whether this result may be extrapolated into humans await further studies.

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