Effects of Carbohydrates on the Toxicity of \textit{p}-Aminophenyl Arsenoxide against \textit{S. cerevisiae}

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Received Date: November 25, 2016
Accepted Date: February 24, 2017
Published Date: March 03, 2017

Abstract

The effects of carbohydrates on the toxicity of \textit{p}-Aminophenyl arsensoxide (\textit{p}-APAO) against \textit{S. cerevisiae} were investigated by Microcalorimetry, including glucose, fructose, lactose, sucrose, N-acetyl glucosamine, 1,3,4,6-tetra-O-acetyl-D-glucosamine (OADG), and chito-oligomer. The toxicity of \textit{p}-APAO on \textit{S. cerevisiae} decreased in the presence of OADG or chito-oligomer. Among the test carbohydrates, the antagonism of OADG was better than those of other sugars. These results suggested that OADG and chito-oligomer were \textit{p}-APAO antagonists. Carbohydrates could form complex with \textit{p}-APAO to prohibit \textit{p}-APAO from polymerizing. This result was attributable to hexose carriers not recognizing the \textit{p}-APAO effectively.

Keywords: \textit{p}-Aminophenyl arsensoxide; Carbohydrates; Toxicity; Trimer; \textit{S.cerevisiae}; Microcalorimetry

Introduction

Since arsenic trioxide was used to treat acute promyelocytic leukemia remarkably effective, scientists have been working on it and its derivatives treatments to variety of other types of cancer\textsuperscript{[1-3]}. Though it is controversial in many applications with high toxicity, Food and Drug Administration of American (FDA) approved it as the treatment of acute promyelocytic leukemia in 2000. In order to overcome some of the shortcomings of arsenic therapeutics with low dose, high toxicity, especially limited dose to the solid tumor in clinic, scientists tried all kinds of ways to decrease the toxicity of arsenic chemicals\textsuperscript{[4-8]}. These methods included: 1) Designing new kinds of inorganic or organic arsenic chemicals; 2) Using other compounds or medicine with arsenic trioxide as combination therapy to both reduce the toxicity of arsenic chemicals and enhance the curative effects. Recently some new types of inorganic and organic arsenic (III, As) compounds with good biological activity have been synthesized. Most of them based on arsenic trioxide or \textit{p}-aminophenyl arsensoxide (\textit{p}-APAO)\textsuperscript{[4-7,9]} which could be prepared from the commercial goods 4-arsanilic (V, As) acid as the pig feed additive\textsuperscript{[10]}. Arsenic trioxide has been applied in clinic with other compounds to cut down its toxicity\textsuperscript{[11,12]}. If the acute toxicity of arsenite is reduced, its range of application would be enlarged\textsuperscript{[13]}. Our previous work indicated that chito-oligomer and glucosamine have significant effect to lower the toxicity of arsenic trioxide\textsuperscript{[13]}.

In fact organoarsenic (III, As) compounds are less toxic than inorganic arsenic compounds such as arsenic trioxide (III, As)\textsuperscript{[14]}. Some of them would be the candidate of new anticancer drugs\textsuperscript{[7]}. To find some substances with the function of lowering the acute toxicity of organic arsenic compounds and strengthening their therapeutic effect is a promising research topic. Based
on the above ideas and results of our previous studies we chose the carbohydrates as our objects to study the effects on the organoarsenic compound p-aminophenyl arsenoxide (p-APAO). These carbohydrates included: monosaccharide, disaccharide and chito-oligomer. Glucosamine could be commercially available through the hydrolysis of chitin, which composes the exoskeletons of crustaceans and other arthropods, cell walls in fungi and many higher organisms. Due to its safety and effectiveness, glucosamine has been approved very popularly as a dietary supplement in almost all countries \(^\text{15,16}\).

Chito-oligomer with non-toxicity and good water-solubility is the oligomer of glucosamine, which can be obtained by degradation of chitosan \(^\text{17}\). It has various biological activities such as antitumor and immune-enhancing effects \(^\text{18,19}\). Microcalorimetry currently is one of the most important techniques in thermochemistry and thermodynamic studies due to its high sensitivity, accuracy, and its non-invasive, non-destructive nature. It also allows real-time, on-line and automated analysis \(^\text{20-25}\). One of the most prominent characters of the growth of the microorganisms is the production of heat. Microbial activity could be quantified through detecting the heat output accompanied all the biochemical oxidation-reduction reaction. Based on detection of microbial cells’ metabolic heat microcalorimetry is an important tool to evaluate their growth activity in quantity and quality \(^\text{26}\). When the heat is detected and recorded by microcalorimeter as small as a microwatt, the useful information in the literature \(^\text{33}\). It was obtained as white solid in yield of 97% (29 g). \(^1\)H NMR (400 MHz, DMSO-\(d_6\)) \(\delta\) ppm 8.82 (brs, 1H), 7.92-7.88 (d, 2H), 5.94 (d, \(J = 8.8\)Hz, 2H), 5.38 (t, 1H), 4.96 (t, 1H), 4.24-4.20 (m, 1H), 4.09-4.01 (m, 2H), 3.59 (t, 1H), 2.20-1.89 (m, 12H). HRMS (ESI): [M+Cl]+ m/z calcd for: C\(_{14}\)H\(_{22}\)NO\(_9\), 348.1295; found: 348.1290. These data were agreement with the literature \(^\text{33}\).

### Synthesis of 1,3,4,6-tetra-O-acetyl-β-D-glucosamine hydrochloride

OADG hydrochloride was synthesized according to the literature (Scheme 2) \(^\text{33}\). It was obtained as white solid in yield of 97% (29 g). \(^1\)H NMR (400 MHz, DMSO-\(d_6\)) \(\delta\) ppm 8.82 (brs, 1H), 7.92-7.88 (d, \(J = 8.8\)Hz, 2H), 5.38 (t, 1H), 4.96 (t, 1H), 4.24-4.20 (m, 1H), 4.09-4.01 (m, 2H), 3.59 (t, 1H), 2.20-1.89 (m, 12H). HRMS (ESI): [M-Cl]+ m/z calcd for: C\(_{14}\)H\(_{22}\)NO\(_9\), 348.1295; found: 348.1290. These data were agreement with the literature \(^\text{33}\).

### Isothermal calorimetry detection

The effect of p-APAO on \(S. cererviae\) was investigat-ed by TAM air, which was an eight-channel isothermal batch calorimeter for heat flow measurements. The metabolic growth power-time curves of microorganism in 20 mL ampoules were obtained, and the experimental data were recorded by TAM Air Assistant. The 1.0 mg/mL As (III) bulk solution was obtained by adding 244 mg p-APAO into sodium hydroxide water solution, and diluted to 100.0 mL with volumetric flask. Before each batch experiment, the As (III) solution was adjusted to pH 7.0 by HCl. The inocula was homogeneous distributed into 50 mL of Luria-Bertani medium (LB medium) by gentle shaking to obtain the suspensions.

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Effects of carbohydrates on the bacteriostatic action of p-APAO

The 4.0 mL of the suspensions was added into each sterilized ampoules. The requested As (III) solutions were added by micropipette. Then the weighed carbohydrates were added into the sterilized ampoules, respectively. Sterilized water was used as a control instead of the sample. The pH of the mixture in each ampoule was adjusted to 7.0 by adding 1 mol/L HCl or NaOH. The liquor volume in each ampoule was adjusted to 5.0 mL by adding sterilized water. The ampoules were sealed tightly and shook for 3 min. The ampoules were placed in the calorimeter and signals obtained during growth were detected. The experiments were run at 28.00 ºC. Each batch experiment was carried out at least three times.

Results and Discussions

Power-time curve for growth of S. cerevisiae

Power-time curve for the growth of S. cerevisiae was shown in Figure 2a (A), which is a typical growth curve for S. cerevisiae and can be divided into four phases: lag phase, active recovery phase, stationary phase and decline phase. The calorimetric power (P), which reflects the multiplication of the cells, can be used as a parameter to characterize the growth of the bacteria. Since the area under the curve represents the total heat (Q) released during the experimental period. During the active recovery phase increases exponentially, the power-time curves obey the following equation: \( P_t = P_0 \exp (k t) \) or \( \ln P_t = \ln P_0 + kt \), where \( P_t \) is the power output at time t, \( P_0 \) is the initial power output, and k is the growth rate constant.

Effect of concentrations of p-APAO on S. cerevisiae

Power-time curves for the growth of S. cerevisiae in the presence of As (III) were displayed in Figure 2. It shows the power-time curves for the growth of S. cerevisiae at 28 ºC in the presence of As (III) different concentration. As (III) had the ability to inhibit the growth of S. cerevisiae to a different extent. The inhibitory effects of As (III) increased at the concentrations ranging from 10.0 mg/L to 80.0 mg/L. With the increase concentration of the drug, the growth of S. cerevisiae became more slowly. When the concentration of As (III) increased to 80.0 mg/L, the power-time curve became a straight line, indicating that S. cerevisiae growth was completely inhibited.

Effects of OADG, COS on the toxicity of p-APAO

Due to the concentration of As (III) increased to 80.0 mg/L, the growth of S. cerevisiae was completely inhibited. So under the concentration of 80.0 mg/L As (III) all the power-time curves of the control C were almost straight lines in Figure 3 and Figure 4. The second control was that the power-time curves in the presence of 4 g/L OADG, 4 g/L COS, which had no obvious influence on the growth of S. cerevisiae in YPED medium. YPED medium is optimal medium for the growth of S. cerevisiae. To further illustrate the influence of different carbohydrates to p-APAO (80 mg/L) on the growth of S. cerevisiae, the power-time curves of the different concentrate about the same carbohydrate was tested (As shown in Figure 3 and Figure 4). In contrast to control A (which was the normal growth of S. cerevisiae).
Effects of Carbohydrates on the Toxicity of \(p\)-Aminophenyl Arsenoxide.

S. cerevisiae in YPED medium.) in the presence of 80.0 mg/L \(p\)-APAO different concentrations of OADG, COS were added to the solution. Power-time curve for growth of \(S.\) cerevisiae appeared again, which stands inhibitory effect of \(p\)-APAO on the growth of \(S.\) cerevisiae reducing. The power-time curve was closer to control A, the detoxification was better. The results showed that they reduced the inhibitory effect of \(p\)-APAO on the growth of \(S.\) cerevisiae. The inhibitory effect of \(p\)-APAO on the growth of \(S.\) cerevisiae decreased with the increased concentrations of OADG (Mw 347) and COS (Mw 1.4 \(\times\) 10\(^3\), if it is discounted to monosaccharide 177), respectively. These results suggested that OADG and COS could lower the toxicity of \(p\)-APAO on \(S.\) cerevisiae. 10 g/L of COS was almost equal to 20 g/L OADG, if these two concentrations are converted into molar concentrations, they are same molar concentrations. Actually the effect of 12 g/L OADG is almost the same of 10 g/L of COS (Figure 3 and Figure 4). This result indicated that OADG has the better capacity of detoxification to \(p\)-APAO than COS.

**Comparison analysis on carbohydrates to the toxicity of \(p\)-APAO**

The inhibitory effect of \(p\)-APAO (80 mg/L) on the growth of \(S.\) cerevisiae was influenced by different carbohydrates (12.0 g/L), and the power-time curves were shown in Figure 5. Sucrose, fructose, lactose, glucose and \(N\)-acetylglycosamine had slight influence on the inhibitory action of \(p\)-APAO on the growth of \(S.\) cerevisiae. But the power-time curve of OADG was the closest to blank control, which stands the best reduced arsenic toxicity effect on \(S.\) cerevisiae. The results demonstrated that OADG could dramatically promote the growth of \(S.\) cerevisiae, whereas other carbohydrates did not display remarkably effect on the growth of \(S.\) cerevisiae. So OADG had the greatest inhibitory effect on the \(p\)-APAO against the growth of \(S.\) cerevisiae among these six carbohydrates. The experimental results showed OADG and COS had clear effects on decreasing the toxicity of \(p\)-APAO on cells of \(S.\) cerevisiae.

**Plausible mechanism for detoxification to the toxicity of \(p\)-APAO**

About 80% of As (III) was transported by the hexose carriers in \(S.\) cerevisiae in form of trimer (Figure 1) in aqueous. Large hinder impeded \(p\)-APAO to polymerize. Trimer of As (III) compounds had been predicted to form a six-membered ring, which is very similar to hexose. This structure could be recognized by the hexose carriers. It is also the main reason intoxication of As (III) in the cell\(^{[29]}\). OADG has stronger effect than the other carbohydrates. This result was due to the following factors:

1. Intermolecular hydrogen bond. 2-position naked amino group in OADG acts with conjugative \(\pi\) bond of benzene ring of \(p\)-APAO to form hydrogen bonds with amino and hydroxyl of \(p\)-APAO. The number of hydrogen bonds also was related with effect of action;

2. Intermolecular \(\pi\)-\(\pi\) interaction. Carbonyl in acetyl group interacts with conjugative \(\pi\) bond of benzene ring of \(p\)-APAO to form intermolecular \(\pi\)-\(\pi\) interaction, which plays an important role. In OADG there are four acetyl which could strengthen the \(\pi\)-\(\pi\) in-

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**Figure 3:** Effect of different concentrations of OADG on \(p\)-APAO influence of the growth of \(S.\) cerevisiae. A: Control; B: 4 g/L OADG; C: As; D: As + 4 g/L OADG; E: As + 6 g/L OADG; F: As + 8 g/L OADG; G: As + 10 g/L OADG; H: As + 12 g/L OADG

**Figure 4:** Effect of different concentrations of COS on \(p\)-APAO influence of the growth of \(S.\) cerevisiae. A: Control; B: 4 g/L COS; C: As; D: As + 4 g/L COS; E: As + 6 g/L COS; F: As + 8 g/L COS; G: As + 10 g/L COS

**Figure 5:** Effect of carbohydrates on \(p\)-APAO influence of the growth of \(S.\) cerevisiae: A: Control; B: As; C: As + OADG; D: As + glucose; E: As + fructose; F: As + \(N\)-acetylglycosamine; G: As + lactose; H: As + sucrose. Concentration of As: 80 mg/L, concentration of different carbohydrate: 12 g/L.
teraction with benzene ring of p-APAO, while there is none π-π interaction in the complex of other carbohydrate and p-APAO except N-acetyl glucosamine with only one acetyl. Due to Intermolecular π-π interaction and Intermolecular hydrogen bonds, these interactions resulted in large hinders to inhibit p-APAO to form trimers (Figure 1). Trimer of p-APAO could be recognized by hexose carriers effectively. Intermolecular interactions of carbohydrates with p-APAO actually made R become larger hinder, which led p-APAO more difficult to be transported by hexose carriers. This resulted in decreasing heap of arsenic compound in cell. So the cytotoxicity of arsenic could be detoxified correspondently. OADG has four acetyl and one naked amino. Acetyl could form π-π interaction with benzene ring of p-APAO. Acetyl also could form hydrogen bonds with amine and hydroxyl groups in p-APAO. Naked amine in OADG could form the hydrogen bond with p-APAO, too. All the facts could explain the strong detoxification of OADG. There are other factors such as the interaction of naked amino with arsenic (III, As) acid. Naked amino group with stronger electron-donating ability than OH can form more stable complex. So glucosamine can form more stable complex than glucose and fructose. COS with amine and hydroxyl could form hydrogen bond with p-APAO, which could also reduce the heap of arsenic compound in cells of S. cerevisiae. COS had the ability of detoxification, too. Acetyl-modified OADG could strengthen its lipophilicity to penetrate cell wall more easily. According to the literatures in the absence of an enzyme the hydroxymethyl group of compounds, such as glucose, fructose can react with arsenate in neutral aqueous solutions to form esters. It is easy for monosaccharides to form complexes with arsenite while not easy for disaccharides. Therefore these disaccharides had unobvious detoxification.

Conclusions

In conclusion, we obtained the general information about S. cerevisiae with different concentrations of p-APAO (III, As) by microcalorimetry in our experiments. OADG and COS were found to have significant effects to lower the toxicity of p-aminophenyl arsenoxide on cells of S. cerevisiae. The effect of OADG was stronger than those of other carbohydrates. These results suggested that OADG and COS may be applied as the assistant antidote for p-APAO. These carbohydrates could interact with p-APAO via intermolecular hydrogen bonds or π-π interaction to form complex to inhibit p-APAO’s polymerization. This led p-APAO not easily to form the similar structure of hexose. So p-APAO couldn’t be recognized by hexose carriers effectively. OADG and COS had the capability of detoxification to p-APAO.

Acknowledgments: This work was supported by National Natural Science Foundation of China (31371750) and National Natural Science Foundation of Hubei Province, China (2014CFB570)

References


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