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## Research Article

# Proteomics Approach (iTRAQ) to Test Subchronic and Chronic Toxicity Effects of Mice Exposed to Drinking Water Source from Yangtze River Delta

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### ABSTRACT

**Objective:** To investigate the combined effects of the pollutants in source of drinking water for local populations, mice models were used to test the subchronic toxicity for three months and chronic toxicity for six months.

**Methods and Results:** By using proteomics approach (iTRAQ), we had identified 29 differentially expressed proteins, which were related to amino acids metabolism, cell skeleton reorganization, apoptosis and cancer genesis. Moreover, 12 out of these 29 identified proteins are located in mitochondria, indicating drinking water source from Yangtze River Delta may exert oxidative damage stress to liver cells. All these cellular processes had been verified by hepatic histopathology, real time-PCR, flow cytometry test and our previous metabolic profile results.

**Conclusion:** All these tests indicated that drinking water source from Yangtze River Delta could accelerate aging, enhance apoptosis and carcinogenesis risk.

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## Introduction

The proteomics approach is increasingly relied on for identifying the critical cellular process and biomarkers relevant to health risk assessment and early disease detection, as well as for the monitoring of disease progression and molecular classification [1]. The current advances in the field of proteomics provide the opportunity to study the expression changes in global proteome level of cells and tissues at any given time point. Especially, the iTRAQ isobaric tags, one of the isotopic labeling approaches, allows to identify the relative and absolute abundance of proteins among up to eight samples simultaneously in one liquid chromatography-mass spectrometry run (LC-MS) [2].

The Yangtze River is the main source of drinking water for cities located in southeast of China. Many pollutants at low concentrations have been found in this river, such as organochlorine pesticides (OCPs), phthalates (PAEs), polycyclic aromatic hydrocarbons (PAHs), and heavy metals

(especially cadmium and arsenic) [3-6]. Not all these possible carcinogens are efficiently removed during the process of treatment of drinking water sources. And the treatments also introduce disinfection by-products and emerging contaminants in source of drinking water, which correlate with carcinogenesis [7]. In our previous work, we had found that pollutants and disinfection by-products in drinking water source from Yangtze River Delta may pose potential carcinogenic risk to local populations. And this conclusion had also been proved by other tests. Long-term exposure to drinking water source from Yangtze River Delta may lead to DNA damage and oncogenesis disorders in mouse models [3-6].

There are two important pictures which are still missing. Firstly, besides all the progresses we made in transcriptomic and metabolic profiles, the proteomic data is still unknown. Secondly, all the previous work we made is based on the fact that the mice exposed to drinking water source from Yangtze River Delta for three months, and we have no idea about

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the subchronic toxicity effects of the mice exposed for six months or even longer. To address these research gaps, we present proteomics test by comparing the mice fed with distilled water, drinking water source from Yangtze River Delta for three months and for six months.

## Materials and Methods

### I Water Sample

To investigate the temporal variations of pollutants in drinking water source from Yangtze River Delta. Water samples were tested in April, June and August 2017. The semi-volatile organic pollutants in samples were extracted by liquid-liquid method (EMC 2006) and DSQ II Single Quadrupole GC/MS (ThermoQuest, San Jose, CA, USA). Details about the chemical analysis were described in our previous work [6]. Recoveries of the detected organic pollutants were above 76.4%.

### II Animal Test

Sixty three-week-old male mice (C57) were got from the Experimental Animal Center, Academy of Military Medical Science (Beijing, China). After acclimation for seven days at the Nanjing General Hospital of Nanjing Military Command, all mice were randomly divided into three groups. Twenty mice each group, housed in two polypropylene cages, standard animal house conditions ( $22\pm 2^\circ\text{C}$ , 12 h light/dark cycle, and humidity of  $50\pm 5\%$ ). The test group mice were fed with drinking water source from Yangtze River Delta for three months (Sub chronic toxicity), six months (chronic toxicity) and the control group was fed with distilled water. The water for mice was monitored daily.

### III Hepatic Histopathology

After the mice were sacrificed, liver samples were harvested. Liver tissues were washed in normal saline, and then fixed in 10% formalin for more than 12 h, and sectioned at 5- $\mu\text{m}$  thickness, then mounted onto slides. Tissue slides were stained with hematoxylin and eosin (H&E), and then observed with optical microscope [6]. And tissue slides were also stained with phalloidin and DAPI, and observed under the laser confocal microscopy.

### IV Protein Extraction and Digestion

To eliminate individual variations, ten random selected frozen livers from each group were ground in liquid nitrogen in a prechilled mortar with pestle. The fine powder was split into two equal parts and transferred to a 2 ml Eppendorf tube and lysis buffer (50 mM Tris 8.0, 1% SDS, 5 mM EDTA) supplemented with 1 mM PMSF was added. The samples were sonicated for 15min (2s-on, 4s-off, 200 w) on ice using a high intensity ultrasonic processor. The remaining debris and unbroken cells were removed by centrifugation at 30000 g at  $4^\circ\text{C}$  for 15 min. The supernatant was transferred to a new tube, reduced with 10 mM DTT for 1 h at  $56^\circ\text{C}$  and alkylated with 55 mM iodoacetamide for 45 min at room temperature in the dark. The protein was precipitated with 4 volumes of prechilled acetone for 30 min at  $-20^\circ\text{C}$ . After centrifugation, the pellet was then dissolved in 0.5 M TEAB and then sonicated for 5min. Repeat the centrifugation step and collect the supernatant. The centrifugation step was repeated and the supernatant was collected. Protein content was

determined with Bradford (Biorad, Hercules, CA, USA) using bovine serum albumin as the standard (Sigma, St. Louis, MO, USA). Approximately 100  $\mu\text{g}$  protein for each sample was digested with trypsin (Promega, Madison, WI, USA) overnight at  $37^\circ\text{C}$  in a 1:20 trypsin-to-protein mass ratio.

### V iTRAQ Labeling

After trypsin digestion, peptide was dried by vacuum centrifugation. Peptides was reconstituted in 0.5 M TEAB and processed according to the manufacturer's protocol for 8-plex iTRAQ (AB Sciex, Concord, Canada). Briefly, one unit of iTRAQ reagent (defined as the amount of reagent required to label 100  $\mu\text{g}$  peptides) was thawed and reconstituted in 70  $\mu\text{L}$  isopropanol. As mentioned above, two aliquots from each subgroup (three months, six months and control) were labeled with different iTRAQ tags respectively by incubation at room temperature for 2 h according to the design. The peptide mixtures were then pooled and dried by vacuum centrifugation. The peptide labeling efficiency was tested by Ultrafle Xtreme MALDI-TOF/TOF (Bruker Daltonics, Bremen GmbH). The pooled mixtures of iTRAQ-labeled peptides were fractionated by SCX chromatography.

### VI Fractionation by Strong Cationic Exchange (SCX)

For SCX chromatography using the LC-20AB HPLC Pump system (Shimadzu Co., Kyoto, Japan), the iTRAQ-labeled peptide mixture was reconstituted with 4 mL buffer A (25 mM  $\text{NaH}_2\text{PO}_4$  in 25% ACN, pH 3.0) and loaded into a  $4.6\times 250$  mm Ultramex SCX column containing 5- $\mu\text{m}$  particles (Phenomenex, Torrance, CA, USA). The peptides were eluted at a flow rate of 1 mL/min with a gradient of buffer A for 10 min, 5-35% buffer B (25mM  $\text{NaH}_2\text{PO}_4$ , 1M KCl in 25% ACN, pH 3.0) for 11 min, 35-80% buffer B for 1 min. The system was then maintained in 80% buffer B for 3 min before equilibrating with buffer A for 10 min prior to the next injection. Elution was monitored by measuring absorbance at 214 nm, and fractions were collected every 1 min. The eluted peptides were pooled as 10 fractions, desalted by Stratacx 18 column (Phenomenex, Torrance, CA, USA) and vacuum-dried.

### VII LC-ESI-MS/MS Analysis by LTQ-Orbitrap HCD

Each fraction was resuspended in certain volume of buffer A (5% ACN, 0.1% FA) and centrifuged at 20000 g for 10 min. In each fraction, the final concentration of peptide was about 0.25  $\mu\text{g}/\mu\text{l}$  on average. 16  $\mu\text{l}$  supernatant was loaded on an Shimadzu LC-20AD nano-HPLC by the autosampler onto a 2 cm C18 trap column (inner diameter 200  $\mu\text{m}$ ) and the peptides were eluted onto a resolving 10 cm analytical C18 column (inner diameter 75  $\mu\text{m}$ ) made in-house. The samples were loaded at 40  $\mu\text{L}/\text{min}$  for 4min, then the 44min gradient was run at 400 nL/min starting from 2 to 35% B (95% ACN, 0.1%FA), followed by 2 min linear gradient to 80%, and maintenance at 80% B for 4 min, and finally return to 2% in 1 min.

The peptides were subjected to nanoelectrospray ionization followed by tandem mass spectrometry (MS/MS) in an LTQ Orbitrap Velos (Thermo Fisher Scientific, Waltham, MA, USA) coupled online to the HPLC. The mass spectrometer was operated in the data-dependent mode and switched automatically between MS (max of  $1\times 10^6$  ions) and MS/MS

(target value of 50,000). Intact peptides were detected in the Orbitrap at a resolution of 60 000. Peptides were selected for MS/MS using high energy collision dissociation (HCD) operating mode with a normalized collision energy setting of 45%; ion fragments were detected in the Orbitrap at a resolution of 7500. A data-dependent procedure that alternated between one MS scan followed by a maximum of eight MS/MS scans was applied for the eight most abundant precursor ions above a threshold ion count of 5000 in the MS survey scan. Parent ions with a charge state from  $z=1$  and unassigned charge states were excluded for fragmentation. Dynamic Exclusion settings were as follows: repeat counts, 2; repeat duration, 30 s; and exclusion duration, 120 s. Ion source parameters were as follows: spray voltage 1.5 kV; no sheath and auxiliary gas flow; ion-transfer capillary temperature 200°C. For MS scans, the  $m/z$  scan range was 350 to 2000 Da.

### VIII Data Analysis and Interpretation

The instrument data file (raw) for each fraction was merged and transformed to a mgf file by Proteome Discoverer (ver. 1.2.0.208; Thermo Fisher Scientific, Waltham, MA, USA). Peptide and protein identifications were performed using the Mascot search engine (ver. 2.3.0; Matrix Science, London, United Kingdom). Database searching was restricted to tryptic peptides of mouse (IPI\_mouse\_v3.87). Carbamidomethyl (C), iTRAQ8plex (N-term), iTRAQ8plex (K) were selected as fixed, iTRAQ8plex (Y), Deamidated (NQ), Oxidation (M) as variable modifications, one missed cleavage allowed and precursor error tolerance at 20 ppm, fragment tolerance at 0.1 Da. Proteins were identified on the basis of having at least one peptide whose individual  $e$ -value wasn't more than 0.05 and also identified as the top-ranked matching sequence for that spectrum. We further restricted the number of identifications made by imposing spectrum and peptide-level nonredundancy such that any given peptide and any given MS/MS spectrum were linked to only one protein. The complete list of identified peptides was then housed in an Excel (Microsoft, Redmond, WA, USA) database for calculation of ratios and coefficient of variation.

**Table 1:** The primer sequences of gene.

Primer	Forward 5'-3'	Reverse 5'-3'
Fah	AGGCTTCTGCGACAATGC	TGCTGCCGAGAAGAGTAGAAG
UK114	TTGGCACTGTCAATGAGATCTATAA	GCAGCGACTTGGAAGCAG
USO1	GACTCCAGACGGAGAACTGC	CCTTCTACAGGAAGTATTTTGC
Pcbd1	TCTACCCTGACTCTTATTTGCTTG	TGGTCCCTTCCTAGGTTCC
GOT1	GCTGTGCTTCTCGCCTAGTT	AAGACTGCACCCCTCCAAC
ABAT	TTCCGGAAGCTGAGAGACA	GCACCTCGTCCACCAAGA
ANXA2	GGAAATATGGCAAGTCCCTGT	TCTGGTAGTCACCCCTGGTGT
Dstn	CACAGACTTTGCCGTCAGG	CTTCATCCGCAACCTGAACT
Adss1	AGGCAGAGAAGAACGAGAAGAA	CTGGTGGAAAGTCGAACACAA
Hint2	GAAGATGGCTGCTGCTGTG	TTCCTCCCGTCCGACAC
Eif4g1	TATCCAGGTGCAAGCCCTAC	ACACCTGGGCTGGGTAATA
Ctsb	GTGTCTGTGAAGACCTGCTT	GGGATAGCCACCATTACAGC
Cyp3a13	AATTCCGCCCTGAAAGGT	AAAGGGCAGGTACATGTAAGGA
Lonp2	CAATTATTGCAACAAGCAACAAA	TGGTCATCTTGAACCGATCC

**Table 2:** Concentrations of organic pollutants in drinking water source from Yangtze River Delta (ng/L), Concentration data = means ± Standard deviations.

Organic pollutants	Concentration	Organic pollutants	Concentration
isophorone	14±6	indeno(1,2,3-cd)pyrene	14±7
pentachlorophenol	17±9	phenanthrene	37±3

The meaningful cutoff of proteins for upregulated  $\geq 1.5$  fold and downregulated  $\leq 0.67$  fold was selected as the differential expressed proteins (DEPs). We also used CV value between technical replicates to estimate the quantitation confidence.

### IX Real time-PCR for Genes Transcriptions Quantification

Total RNA was extracted from liver by TRIzol Reagent (Invitrogen, Carlsbad, CA) and reverse-transcribed by RT Kit (Fermentas, Lithuania). cDNA was quantified by Taq DNA Polymerase (Fermentas, Lithuania) and Realtime PCR Master Mix (SYBR Green) (Toyobo, Japan).  $\beta$ -actin was used as an internal normalizing gene. The primer sequences used were as follows.

### X Liver Mononuclear Cell Preparation and Flow Cytometry Test

Mononuclear cells were prepared by using a 200-gauge stainless-steel mesh, and cells were stained by DCFH-DA (2',7'-Dichlorofluorescein diacetate, Sigma, St. Louis, MO) to test the Cellular oxidative damage [8]. We arranged four groups: negative control, three months group, six months group, positive group (negative control +10% H<sub>2</sub>O<sub>2</sub>).

### Result

#### I Organic pollutants in Drinking Water Source from Yangtze River Delta

There were 23 organic pollutants detected in drinking water source. It was found that PAEs and PAHs contributed more than 95% to the total concentration of the 23 detected organic pollutants (Table 2).

acenaphthylene	6±2	pyrene	8±4
anthracene	63±26	butyl benzy phthalate	7±3
benzo(a)anthracene	14±6	bis(2-ethylhexyl)adipate	372±234
benzo(b)fluoranthene	87±34	bis(2-ethylhexyl)phthalate	1914±653
benzo(k)fluoranthene	22 ±9	dibutyl phthalate	4932±1043
benzo(a)pyrene	7±6	diethyl phthalate	29±7
chrysene	16±10	2,4-dinitrotoluene	36±19
dibenz(a,h)anthracene	40±14	2,6-dinitrotoluene	300±18
fluorene	107±33	hexachlorocyclopentadiene	15±3
hexachlorobenzene	11±4	dimethyl phthalate	107±7
TOC(wet process)	1.860-2.656mg (RSD<1%)		

**II iTRAQ Test**

1245 proteins had been identified and 1237 proteins were confirmed after filtering. 15 proteins were differentially expressed after exposed to drinking water source from Yangtze River Delta for three months, and 33 proteins for six months compared to the control group. Considering the expression behaviour of these proteins, 29 proteins had been selected to evaluate the toxicity effects of drinking water source from Yangtze River Delta.

Among all the 29 proteins, 22 were related to signaling pathways, and 14 of the 22 proteins are located in the metabolic pathways; several proteins were related to apoptosis and cytoskeleton; 26 of the differentially expressed proteins were associated with cancer, 7 proteins among 26 indicate liver cancer directly. Another important detail was that 12 of 29 proteins were located in mitochondrial. All details were shown in (Table 3).

**Table 3:** The 29 differentially expressed proteins profiles in mouse livers.

Ref.	Full Name	Symbol	Fold		Disease	local	Gene Ontology		Pathway
			3 m	6 m			Function	Process	
[9]	Fumarylacetoacetase	Fah	0.94	2.2	Live cancer; hepatic necrosis; implanted secondary tumor	Nu	fumarylacetoacetase activity	L-phenylalanine, tyrosine, phenylalanine, arginine catabolic process; metal ion binding; hydrolase	Metabolism of amino acids
[10]	Ribonuclease UK114	Uk114	1.3	2.1	Cancer	Mito	endonuclease activity	inhibition of the translation by cleaving mRNA; endonuclease; tyrosine catabolic process; regulation of translational termination;	
[11]	Betaine-homocysteine S-methyltransferase 1	Bhmt	0.99	2	Prostate cancer	Cyto	homocysteine S-methyltransferase activity	protein methylation; regulation of homocysteine metabolism; S-methyltransferase activity; Amino-acid biosynthesis	Metabolism of amino acids
[12]	Isoform 4 of General vesicular transport factor p115	Uso1	1.1	2	Liver cancer; Prostate cancer	Cyto	protein transporter activity	intracellular protein transport; apoptosis; actin and microtubules;	
[13]	Sulfite oxidase	Suox	1.4	1.8	Prostate cancer	Mito	Sulfite-oxidase activity; electron carrier activity	oxidation-reduction process; response to nutrient	Metabolism of sulfur
[14]	Aspartate aminotransferase	Got1	1.4	1.8	Liver cancer	Mito	L-aspartate:2-oxoglutarate aminotransferase activity;	amino acid metabolism; the urea and tricarboxylic acid (TCA) cycles	Metabolism of amino acids

							phosphatidylserine decarboxylase activity		
[15]	Pterin-4a-carbinolamine dehydratase	Pcbd1	1.3	1.8	dysplasia; early stages of cancer; malignant transformation;	Cyto	phenylalanine 4-monooxygenase activity	L-phenylalanine catabolic process; positive regulation of transcription; tetrahydrobiopterin biosynthetic process;	Metabolism of amino acids
[16]	Annexin A2	Anxa2	1.2	1.7	Liver cancer;	Cyto	calcium-dependent phospholipid binding	cytoskeletal protein binding; Rab-GTPasebinding; phosphatidylinositol-4,5-bisphosphate binding	Prostaglandin Synthesis and Regulation
[17]	4-aminobutyrate aminotransferase	Abat	1.2	1.7	Liver cancer	Mito	4-aminobutyrate transaminase activity	transaminase activity; gamma-aminobutyric acid catabolic; DNA-damaging product formation;	Metabolism of amino acids
[18]	4-hydroxy-2-oxoglutarate aldolase	Hoga1	1.2	1.6	engineered invasive esophageal squamous cell carcinoma	Mito	catalytic activity	metabolic pathway of hydroxyproline	Metabolism of hydroxyproline
[19]	Serine hydroxymethyl transferase 2	Shmt2	1.3	1.6	Prostate cancer	Mito	transferase activity	Regulation of metabolism; Serine+tetrahydrofolate=glycine +5,10-methylene tetrahydrofolate acid	Metabolism of cyanoamino acid
[20]	Microtubule-associated protein RP/EB family member 1	Mapre1	1.2	1.5	Gastric cancer	Cyto	microtubule plus-end binding	Cytoskeleton; mitosis; microtubule organizing center	Cell Cycle G2/M Transition
[21]	Coiled-coil-helix-coiled-coil-helix domain-containing protein 3	Chchd3	1.4	1.5	Anaplastic Cancer	Mito	phosphatase binding	mitochondrial fusion	
[22]	Destrin	Dstn	1.4	1.5	Prostate cancer	Cyto	actin binding	actin cytoskeleton; positive regulation of actin filament depolymerization; actin filament severing; catabolic process	
[23]	Isoform 1 of Phosphotriesterase-related protein	Pter	0.95	1.5	Colon cancer; Leukemia; Prostate cancer	Cyto	hydrolase activity; zinc ion binding		
[15]	Phosphoglycerate kinase 1	Pgk1	0.97	0.67		Cyto	Phosphoglycerate kinase activity	glycolysis; ATP binding; phosphoglycerate kinase activity; transferase activity;	Gluconeogenesis
[24]	Legumain	Lgmn	0.93	0.66	Leukemia; colorectal cancer	Cyto	peptidase activity	serine/threonine kinase activity; cysteine-type endopeptidase; negative regulation of growth	Lysosome
[25]	Isoform 1 of Core histone macro-H2A.1	H2afy	0.99	0.64	Colon cancer; leukemia	Nu	chromatin binding	nucleosome assembly; chromatin modification	Systemic lupus erythematosus

[26]	Succinate dehydrogenase [ubiquinone] iron-sulfur subunit,		0.98	0.61		Mito	protein binding	glycerol-3-phosphate dehydrogenase complex; glycerophosphate shuttle; oxidoreductase	TCA Cycle
[27]	Isoform 1 of Eukaryotic translation initiation factor 4 gamma 1	Eif4g1	0.84	0.59	Sarcoma; Leukemia; mitochondrial disorder; Hepatitis; Colorectal cancer	Cyto	translation initiation factor activity	regulation of translation	Metabolism of mRNA; translation; PI3K Cascade; eIF4F complex; mTOR signalling pathways
[28]	Histidine triad nucleotide-binding protein 2	Hint2	0.77	0.59	Prostate cancer; Colon cancer; leukemia	Mito	catalytic activity	Apoptosis; steroid biosynthesis; hydrolase	
[29]	Cytochrome P450 2F2	Cyp2f2	0.87	0.58	Growth suppression; insulin resistant;	ER	Monoxygenase; electron carrier; oxidoreductase activity	response to toxin; naphthalene metabolic process	Metabolism of xenobiotics by cytochrome P450
[30]	Cathepsin B	Ctsb	0.98	0.58	mitochondrial disorder; hepatitis; leukemia; prostate cancer	Mito Cyto	cysteine-type endopeptidase activity	response to wounding; endopeptidase; response to stress; negative regulation of apoptosis	Lysosome
[31]	Isoform 1 of Lon protease homolog 2	Lonp2	0.95	0.54	Liver cancer; leukemia	Pero	protein binding	signal peptide processing; ATPase; serine-type endopeptidase	Metabolism of proteins
[32]	ATP synthase subunit f	Atp5j2	1	0.51	hepatitis; leukemia; carcinoma in situ lesion	Mito	ATPase activity	transmembrane transporter; ATPase; mitochondrial proton-transporting ATP synthase complex	Oxidative phosphorylation; Electron Transport Chain
[33]	Majoy urinary protein 20	Mup20	0.62	0.37			pheromone binding	olfactory learning	
[34]	Isoform 1 of Adenylosuccinate synthetase isozyme 1	Adss1	0.89	0.63	Liver cancer; colorectal cancer; obesity; leukemia	Cyto	adenylosuccinate synthase activity	purine nucleotide biosynthetic; adenylosuccinate synthase; AMP biosynthetic	Metabolism of amino acids
[35]	Cytochrome P450 3A13	Cyp3a13	0.67	0.57	insulin resistant; liver cancer	ER	Monoxygenase; electron carrier; aromatase; iron ion binding	Metabolism of xenobiotics by cytochrome P451	Monoxygenase; electron carrier; aromatase; iron ion binding
[36]	Long-chain specific acyl-CoA dehydrogenase	Acadl	1.6	1.5	sarcoma	Mito Cyto		lipid catabolic; negative regulation of fatty acid oxidation; long-chain-acyl-CoA dehydrogenase	TCA Cycle

Nu: Nuclear; ER: Endoplasmic Reticulum; Cyto: Cytoplasm; Mito: Mitochondrial; Pero: Peroxisome.



### III Realtime PCR

Overall pattern of mRNA expression is similar to that of protein expression in mammalian systems, and integrated analyses of mRNA and protein expression data by concurrent measurement was observed

strong correlation in mammalian cells [37-38]. In order to complement the proteomics results, the real-time quantitative PCR was performed on 14 genes encoding differentiated proteins. The fold change of mRNA is in line with the proteins and indicates the high confidence of our proteomics result. Details were referred to (Table 4).

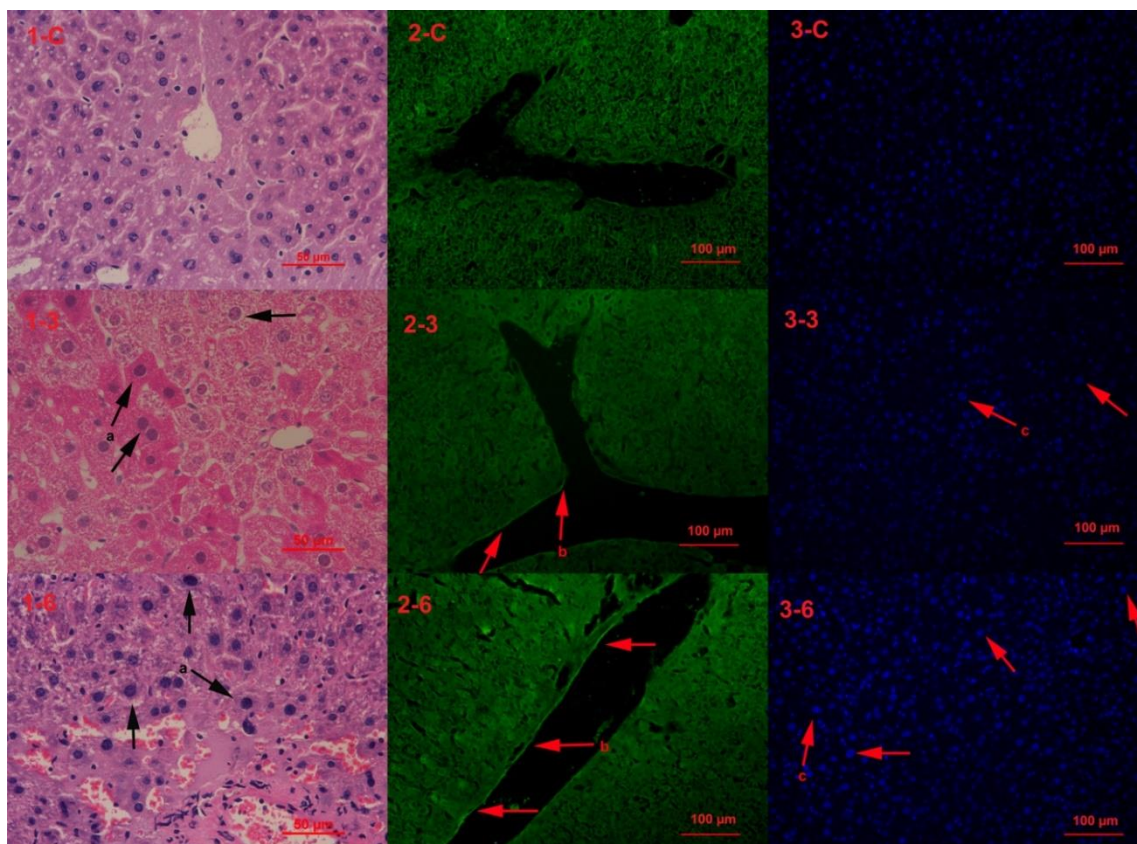
**Table 4:** Analysis by RT-qPCR of the expression of 14 oncogenes.

Gene Symbol	3 mon	6 mon	Gene Symbol	3 mon	6 mon
<i>Fah</i>	1.4	3.1	<i>Adssl1</i>	0.78	0.72
<i>Ukk114</i>	1.2	2.8	<i>Hint2</i>	0.49	0.47
<i>Uso1</i>	1.4	3.3	<i>Eif4g1</i>	0.84	0.59
<i>Pcbd1</i>	1.3	3.1	<i>Ctsb</i>	0.93	0.62
<i>Got1</i>	1.3	2.7	<i>Cyp3a13</i>	0.67	0.49
<i>Abat</i>	2.1	2.8	<i>Dstn</i>	3	3.2
<i>Anxa2</i>	2.1	2.8	<i>Lonp2</i>	0.89	0.47

### IV Hepatic Histopathology

The first line of the photos were control group (1-c, 2-c, 3-c), the second line were mice exposed to drinking water source from Yangtze River Delta for 3 months (1-3, 2-3, 3-3), and the third line were mice exposed

for six months (1-6, 2-6, 3-6). The tissue pathologic phenomenon was emerged in 3 months group and exacerbated in 6 months (Figure 1). All the pathology results indicate the tissue were in disorder under the stress and exacerbated with longer time.



**Figure 1:** Hepatic Histopathology. Note: a-nuclear bulging and chromosome condensing, key characteristic of aging (H&E staining); b-cytoskeleton reorganization in vascular endothelial cell (phalloidin staining); c-nuclear bulging (DAPI staining).

### V Oxidative Damage

The oxidative damage has been found in hepatic cells. Oxidative damage in six months group is twice more than in three months group, and both of these two groups are less than in the positive group. The peak in left

region means healthy cells, and the right regions represent oxidative damage cells. Details refer to (Figure 2).

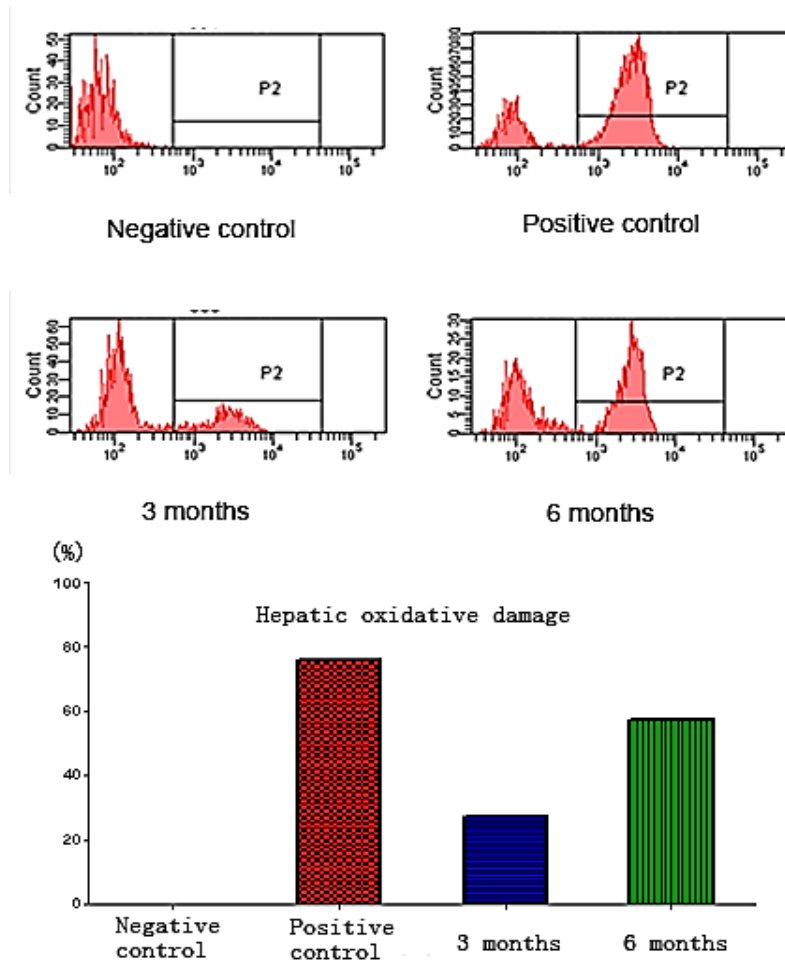


Figure 2: DCF (2',7'-Dichlorofluorescein diacetate) test the oxidative damage (P2 region).

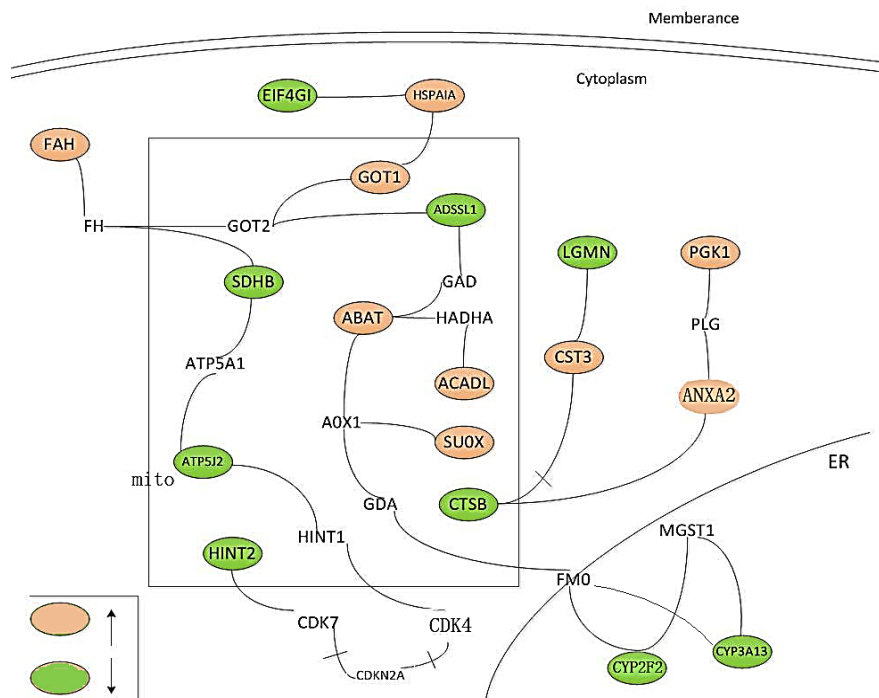


Figure 3: The interactions between the differently expressed proteins (proteins colored orange or green).



## Discussion

In an attempt to better understanding toxicity mechanisms and disease progression of drinking water source, we classified the cellular process and function into four sections: amino acids metabolism; cell skeleton (actins and microtubules) reorganization; apoptosis and cancer. But in the hepatic histopathology, we found the phenomenon that aging existed in liver tissue and actins organization in vascular endothelial cell. Aging is a bridge that adjoins oxidative stress, mitochondrial disorder to apoptosis and cancer [39, 40].

Amino acids catabolic process indicates the liver injury, which may cause rapid hepatocyte death [9, 10]. The actin and microtubules decide the direction of apoptotic cells [12, 16]. However, stress-induced failure of cell death programmes may lead to an accumulation of damaged cells and therefore enhance the risk of carcinogenesis [9, 22]. On the other hand, the aging can enhance the apoptosis risk; and as a defense mechanism, dealing with the oncogenes disorder, aging is an effective way to inhibit the carcinogenesis [39, 41].

It is found that the differential expression of mRNA and proteins was enlarged with longer time. The vascular endothelial cell injury was more serious and number of aging cells was also increased in six months group compared to three months. And the closely related differentially expressed proteins were presented in (Figure 3).

### I Metabolism of Amino Acids

In our previous work, the NMR-based metabolic profile of mice exposed to drinking water source for 90 days had shown that several amino acids had disorder in serum [6, 42]. And in our proteomic profile, we found that FAH, BHMT, UK114, GOT1, PCBD1, ADSSL1 were related to amino acid catabolic process.

### II Cellular Stress Responses

The flow cytometry test has showed the oxidative damage of liver. In the proteomic profile, we had found that 12 proteins located in the mitochondrial, 13 proteins in the cytoplasm, 2 in the endoplasmic reticulum (ER), 2 in the nuclear, and 1 in peroxisomal. The chemical stress may cause the disorder of mitochondrial, ER, peroxisomal, and inter-organelle crosstalk during cellular stress, like mitochondrial–nuclear crosstalk [43, 44]. ER Stress responses have been found playing important roles in maintaining cellular functions. Creld2 is a novel mediator in regulating the progression of various ER stress-associated diseases and CYP2F2 response to wounding [45].

Several proteins located in the mitochondrial are related to cellular stress response. One of them is CTSB, which is an important indicator of autophagy, apoptosis and response to stress. ACADL, SDHB, GOT1 are in the tricarboxylic acid (TCA) cycle, which is determined in the energy and biomolecules carbon skeleton in cell. SUOX is related to electron transfer, while ABAT may indicate the DNA damaging in mitochondrial. All the disorder of these proteins in mitochondrial may causes oxidative damage to cells, and leads to aging and apoptosis [40, 46-48].

### III Cytoskeleton Reorganization

On the other side of this text, it has been found that USO1, ANXA2, MAPRE1, DSTN were related to cell skeleton. The disorders of USO1 (p115), ANXA2 (P36) and DSTN which are associated with actin and microtubules decide the direction of apoptotic cells [12, 16]. After the containing in source of drinking water transferring into blood and exerting stress to the vascular, it had been observed that the cytoskeleton reorganization in vascular wall cells and reflected cells injury under stress [49].

### IV Aging, Apoptosis and Cancer

In the hepatic histopathology, we found the number of aging cells was increased in six months group comparing to three months group and control group. It also verified the accumulations of oxidative damage. The mitochondria disorder directly caused cell aging and raised up the apoptosis risk. While facing with the oncogenes disorders, aging is an effective way to restrict the carcinogenesis. However, the accumulation of cells damaging can also enhance the risk for cancer [9, 22, 50].

## Conclusion

The drinking water source from Yangtze River Delta may exert oxidative damage stress to hepatic cells and also causes the amino acids metabolism disorder, cell skeleton reorganization, oncogene differentially expression. All these cellular processes directly lead to cell aging. Comparing with the subchronic and chronic toxicity effects, we found that the water could accelerate aging, enhance apoptosis and increase carcinogenesis risk.

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