Original Article
Decrease in prosaposin in spermatozoon is associated with polychlorinated biphenyl exposure

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Received January 11, 2015; Accepted February 28, 2015; Epub March 1, 2015; Published March 15, 2015

Abstract: Polychlorinated biphenyls (PCBs) are a class of ubiquitous persistent organic pollutants and they have been associated with declining male fertility. In the present study, we aimed to determine the responsiveness of prosaposin (Psap) expression to PCB exposure. Male C57 mice were exposed to PCB mixture (Aroclor 1254) of environmental related doses by oral gavage. After exposure for 50 days, the expression of Psap was significantly decreased by PCB exposure in epididymides and epididymal spermatozoa, but not in testis. The Psap abundance in sperm was decreased in a dose-dependent manner. Benchmark dose modeling revealed the 95% lower confidence limit on the benchmark dose (BMDL) and Benchmark Dose (BMD) for Psap reduction were 1.25 and 8.89 μg/kg Aroclor 1254, and for sperm motility reduction were 11.85 and 61.9 μg/kg Aroclor 1254. The depressed Psap level also showed a significant correlation \( P<0.01, r=-0.531 \) with PCB accumulation in liver. In men with detectable PCB exposure in semen, Psap expression in sperm was significantly decreased whereas the semen parameters were unaffected. Linear regression showed that a negative association between total PCB level in seminal plasma and Psap level in ejaculated spermatozoa \( P<0.05, r=-0.396 \). In conclusion, our data suggested that the abundance of Psap in sperm sample may be a sensitive endpoint to predict PCB exposure.

Keywords: Polychlorinated biphenyls, prosaposin, sperm, human, mice

Introduction

Polychlorinated biphenyls (PCBs) are a class of legacy chemicals with high environmental persistence. Despite the ban on their production, PCBs remain to be a global environmental problem due to their high resistance against electrical, thermal, chemical, or biological breakdown and ongoing “leaking” to the environment from existing applications and waste. Accumulated evidence have associated exposure to PCBs with many adverse health effects, including effects on reproduction [30] and development [34], thyroid system [36], nervous system [27], immune system [28], cardiovascular system [22], and metabolism [18].

The effects of PCBs on male reproductive system have long been of concerned. PCB congeners have shown estrogenic, antiestrogenic, or anti-androgenic effects [16] which might disrupt the male reproductive functions and hypothyroidism induced by PCB exposure in early life also results in declines in reproductive endpoints. A number of animal studies have reported adverse effects in relation to PCBs following both developmental exposures and exposure in adulthood, including reduced organ weights, impaired spermatogenesis, and disrupted steroid hormone function [1, 5, 6, 11]. Epidemiological studies have associated exposure of human to PCBs with declining semen parameters, sperm DNA integrity, and circulating reproductive hormone levels [30]. Impaired male reproductive functions are undoubtedly impact final reproductive outcomes. A recent study has demonstrated that exposures of PCBs in male partner might contribute to decreased human fecundity, as measured by a prolonged time to pregnancy (TTP) [9]. The research reaffirmed the significance of PCB-related health consequences in human fertility.
In toxicological studies, biomarkers of effect are used to associate the xenobiotic exposures with their harmful downstream health consequences. They are typically physiological indicators which presence after physiologically relevant doses and response to xenobiotic of interest in a specific and sensitive manner. Furthermore, they should be facile in measurement and validatable in humans exposed to the toxin. Semen parameters are the most frequently used biomarkers to investigate the xenobiotic-related effects on male fertility. They are correlated directly to the fertility and can be measured objectively with a relative simple tool [33]. However, semen quality parameters provide less information about the mechanism of the action and the organ targeted. Toxicants can affect the male reproductive system at one of several sites, or at multiple sites, as well as physiological functions. There is so far no single all-encompassing biomarker of reproductive capacity in men. Identifying novel biomarkers reflecting have no doubt enabled us to better evaluate the effect of the exposure on the male reproductive system.

The epididymis is known to play an important role in providing the microenvironment for sperm maturation and storage of sperm [13]. Numerous proteins synthesized and secreted by the epididymal epithelia cells are thought to be involved in male reproductive activities including the initiation of sperm maturation, sperm-oocyte recognition, the acrosome reaction, interaction with the zona pellucida, and binding to and fuse with plasma membrane of the oocyte. The caput epididymis is the most metabolically active region, accounting for 70-80% of the total overall protein secretion in the epididymal lumen [13]. Characterizing global transcript responses to PCBs in the caput epididymis might yield a clue concerning the potential mechanisms for their toxicological effects and might provide candidate markers for fertility and toxicologic evaluations.

In our previous studies, we have demonstrated that low doses of PCB exposure up to 500 μg/kg impaired sperm count, sperm motility and sperm morphology without significantly affecting the weight or morphology of the organs [10, 11]. The exposure also induced a global alternation in the gene expression of caput epididymis [10]. Amongst the dysregulated genes, the expression of prosaposin (Psap) mRNA is dose-dependently down-regulated in the microarray data. Psap is a 65-70 kilodalton glycoprotein that is targeted to the lysosome, as well as is secreted to extracellular fluids (Hiraiwa et al, 1992) and a role of Psap in sperm-oocyte interaction has been suggested (29) (Magargee et al. 2000). The goals of this study were to validate the effects of PCB exposure on the protein expression of Psap in male reproductive tract and to determine whether PCB-induced diminutions in Psap were present on epididymal sperm.

Materials and methods

Animals and treatment

All animal experiments were conducted according to the research protocols approved by the Xiamen University Institutional Animal Care and Use Committee. Male C57 mice, aged 21 days and weighing 12-14 g, were purchased from Fujian Medical University, China, housed at 24±1°C under a 12:12 h light-dark cycle, with free access to food and water. After a quarantine period, mice with adequate weight gain and without clinical signs were divided randomly into three experimental groups. Aroclor 1254 of analytical grade purity (lot LB38310; Supelco, Bellefonte, PA) was dissolved in a 10% ethanol saline solution as previously described [10, 11] and serially diluted with the same solution for exposure experiments. The mice in each treatment were administered with Aroclor 1254 by oral gavage once every three days (either 5 or 500 μg/kg doses); control mice received an equal volume of vehicle (5 μL/g). The mice were sacrificed after 50 days exposure under slight ether anesthesia. The epididymis was removed, cleared of adhering connective tissue and stored at -80°C for further analysis or placed in 4% formaldehyde for 24 h, rinsed in water, and stored in 70% ethanol until further processing for immunohistochemical examination.

The epididymis of each mouse was excised and rinsed with prewarmed PBS. The isolated Cauda epididymis was placed in 1 mL of human tuba fluid (Chemicon, Temecula, CA), cut finely with scissors, and incubated in 5% CO₂ atmosphere for 15 min at 37°C. An aliquot of the sperm suspension was placed on a clean glass slide to make a wet preparation for motility evaluation.
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RNA isolation and Real-time PCR

Total RNA was extracted with Trizol (Invitrogen, Carlsbad, CA) according to the manufacturer’s instructions. One milligram of RNA was reverse-transcribed using the Quantitect Reverse Transcription kit (QIAGEN, Valencia, CA). Quantitative RT-PCR was performed using SYBR green fluorescence and the Qiagen Rotor Gene 3000 Real-Time PCR cycler (QIAGEN, Valencia, CA). Each reaction was run in triplicate and consisted of 20 ng of cDNA, Fast Power SYBR Green PCR System (Applied Biosystems) and 300 nM of the gene-specific primers. The fold change in gene expression was calculated using the comparative cycle threshold method with the housekeeping genes. The expression of L32, RPL11 and Cyclophilin were used as internal control in the quantification.

PCR was performed using the following primers: Psap: 5’-GCCAGAGGGCAGGAGCATT and 5’-CTGACCCAGGGACAGCAACA, L32: 5’-TCCACATTGTCAAGGAGCTG and 5’-ACTCATTTTCTCCTGGCCT, RPS11: 5’-CGTGACGAAGATGAAGATTGC and 5’-GCACATTGTGAATCGCACAGTC, Cyclophilin: 5’-ACACGCCATAATGGCACTGG and 5’-ATTTGCCATGGACAGATGCC.

Western blotting assay

Protein was extracted from frozen tissues using homogenization, and fraction samples (40 μg proteins) were mixed with 2x Coomassie brilliant blue, heated to 100°C in a water bath for 5 min. Next, 10% sodium dodecyl sulfate polyacrylamide gel electrophoresis (SDS-PAGE) was performed at a constant voltage of 100 V for 1.5 h. After SDS-PAGE, the proteins were transferred to a nitrocellulose membrane and blocked at room temperature for 2 h in PBS buffer containing 5% nonfat dried milk or 5% bovine serum albumin to prevent nonspecific binding of reagents, and then incubated overnight at 4°C with a primary antibody. Anti-Psap antibody was obtained from Abcam (Cambridge, MA). After this, the membrane was washed three times in PBST for 15 min and incubated with secondary antibody (1:10000 dilution; Pierce, USA) for 1 h at room temperature. Then the membrane was washed three times in PBST and chemiluminescence detection (Sigma, UK) was performed. The intensity of bands was quantified using Quantity One software (Bio-Rad).

Immunohistochemistry

Fixed tissues were embedded in paraffin and cut into 6 μm sections. The sections were deparaffinized, rehydrated, and stained following the previously described methods [11]. To optimize immunohistochemical staining, an antigen retrieval protocol was carried out by immersing the sections in 10 mM citrate buffer (pH 6.0) and heating in a microwave. The sections were incubated with anti-Psap antibodies overnight at 4°C in a humidified chamber and the negative control was incubated in the presence of irrelevant IgG instead of the primary antibodies.

Psap measurement in human spermatozoa and seminal plasma

A total of 36 male counterparts of women visiting the Affiliated Chenggong Hospital of Xiamen University for in vitro fertilization (IVF) treatment were enrolled in the study and donated the semen samples. The study was approved by the Institutional Review Board and informed consent was obtained from participants. All couples were diagnosed as female factor infertility and the cases with known causes of male infertility such as a history of orchitis, cryptorchidism, varicocele, vas deferens obstruction, and Y chromosome microdeletions were excluded. Semen parameters were analyzed according to WHO criteria (WHO 2010). Semen samples were centrifuged to separate the sperm from seminal plasma. For the resulting sperm, sperm proteins were isolated as previously described [7]. The sperm lysate and the seminal plasma were stored at -80°C for following Psap measurement.

Psap was determined using an ELISA kit (Uscn Life Science Inc., Wuhan, China) according to the manufacturer’s instructions. The enzyme-substrate reaction is terminated by the addition of a sulphuric acid solution and the color change is measured spectrophotometrically at a wavelength of 450 nm. The concentration of Psap in the samples is then determined by comparing the O.D. of the samples to the standard curve. The minimum detectable level of Psap detected is less than 8.2 pg/ml.

Determination of PCBs

PCB levels in mice livers and human seminal plasma were determined. For livers from exposed mice, a microwave-assisted extraction (MAE) procedure was used isolate PCBs [39].
Briefly, 1 g portion sample was weighted and 25 mL acetone-n-hexane (1:1, v/v) was used as the extractant. The extraction was carried out using MARS-X microwave-accelerated reaction system model (CEM, USA) under 100°C, 35 psi for 15 min at 300 W power output. The MAE solution were cooled to room temperature and dried through a short column packed with anhydrous Na$_2$SO$_4$. A cleanup on a SiO$_2$-Al$_2$O$_3$ column was followed and the columns were eluted with n-hexane. The eluate was finally concentrated to 1 ml under nitrogen for Gas chromatographic (GC) analysis.

For semen samples from human participants, 1 ml seminal plasma was fortified with 20 μl internal standard solution of 4, 4'-Dibromofluorobiphenyl (1000 ppb). The sample was thereafter mixed with 1 ml formic acid and equilibrated by ultrasonic treatment for 30 min. The resulting solution was extracted with solid phase extraction (SPE) cartridges. Prior to the sample application, the cartridges were washed with dichloromethane and activated with methanol and water. After conditioning, the cartridges were not allowed to dry. After sample loading, the SPE cartridge was rinsed with 1 ml water. The sorbent bed was dried by centrifugation (15 min, 3000 rotations/min). A SiO$_2$-Al$_2$O$_3$ column was placed under the SPE cartridges. The SPE cartridge was eluted with 2×500 μl n-hexane and 2×500 μl dichloromethane directly into the clean-up cartridge. Further, PCBs were eluted with 3 ml n-hexane and 3 ml dichloromethane from the clean-up cartridge. The final eluate was concentrated and stored as described above for liver samples.

An Agilent 6890N with electron capture detection (ECD) was used for PCB quantification (Agilent, USA). An aliquot of 1 μl of standard solution or sample solution was injected for each GC-ECD analysis. An aliquot of 1 μl of standard solution or sample solution was injected for each GC-ECD analysis. The number of determinations for each sample was three. Nitrogen was both the carrier gas at a flow-rate of 1.5 ml/min and the make-up gas at 60 ml/min. The temperature of the injector and the detector were maintained at 280 and 300°C, respectively. The column was programmed from 85°C, held for 2 min, to 180°C at 30°C/min, held for 2 min, to 200°C at 20°C/min, held for 3 min, to 230°C at 3°C/min, to 250°C at 5°C/min and held for 8 min.

Quantification of PCBs was performed using external standard calibration. To determine Aroclor 1254 accumulation in exposed mice, Aroclor 1254 was used as external standard and eleven major peaks were selected for quantification. To determine environmental exposure of PCBs in seminal plasma, twelve PCB congeners (PCB 8, PCB 18, PCB 28, PCB 44, PCB 66, PCB 101, PCB 103, PCB 153, PCB 138, PCB 187, PCB 128 and PCB 206) were used as standard. Peak area ratios (analyte response/internal standard response) were plotted against the concentration ratios (analyte concentration/internal standard concentration). Method limits of quantification (LOQ) were calculated as 3× standard deviation of the analyte values in procedural blanks. Method LOQs for individual PCBs ranged between were 2 to 6 ng/g LW. Recoveries ranged between 66% PCBs in semen.

Statistical analysis

All statistics were performed using SPSS 19.0. Results were reported as mean ± S.E. The data were statistically analyzed with one-way analysis of variance (ANOVA) and the significance level between data was examined using least significantly difference (Duncan; P=0.05) post hoc tests. Linear regression analysis was used to determine statistically if a measurement was dose-dependent or time-dependent, while a Pearson correlation was used to relate Psap expression to other endpoints.

Benchmark dose modeling

The United States EPA Benchmark Dose (BMD) Software Version 2.4 was utilized to compare benchmark doses (BMD). One standard deviation from the control mean was used to determine the benchmark dose response. For each endpoint, the best-fitting model was chosen using the methodology recommended in the Benchmark Dose Technical Guidance [15], which includes visual inspection of the model fit to the data and analysis of the chi-square value, p-value, Akaike's Information Criterion (AIC), and the 95% lower confidence limit on the benchmark dose (BMDL).

Results

PCBs dose-dependently decreased Psap in mouse epididymis and sperm

In the present study, mice were exposed to environmental related doses of PCBs for 50
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days, which decreased the sperm parameters in a dose-dependent manner whereas the body weight remained unchanged. Real-time PCR analysis at 5 μg/Kg group and 500 μg/Kg revealed a trend toward decreased Psap mRNA in epididymis (Figure 1A), which validated our previous finding in microarray [10]. The western blot analysis confirmed the trend (Figure 1B).

Figure 1. Expression of Psap in the testes and epididymides of mice exposed to Aroclor 1254. A. The expression of Psap mRNA in the caput epididymides of mice exposed to Aroclor 1254. B. Effects of Aroclor 1254 exposure on the protein abundance Psap in mice epididymis. C. Effects of Aroclor 1254 exposure on the protein abundance of Psap in mice testis. Each column represents the mean ± SE of at least three separate experiments. "*" indicates significantly different to control at \( P<0.05 \). "#" indicates significantly different to 5 μg/Kg group at \( P<0.05 \).
and showed the protein expression responded to PCB exposure in a dose-dependent manner. In the 500 μg/Kg group, the protein expression decreased 70% over control.

Sertoli cells are important source of Psap in male reproductive tract and the expression of Psap in testes was also investigated by western blot. In contrast to the epididymides, the testes showed no sign of reduction of Psap abundance, even in the highest dose tested (Figure 1C).

We further investigate the alteration of Psap distribution in testes and epididymides after PCB exposure. In coincidence with previous reports, an intense staining was found in testes and a moderate staining present in the caput epididymides (Figure 2). Neither the intense nor the pattern of the staining was observably changed in testes after 500 μg/Kg Aroclor 1254 exposure. In the caput epididymis, Psap staining present in the supernuclear region of epithelia cells (arrowhead), as well as the apical surface of the epithelium (arrow). In the lumen, staining presented associated with epididymal sperm (curved arrow), which suggested a role of the protein in the epididymal maturation. As the mice exposed to 500 μg/Kg Aroclor 1254, the decrease in the staining was observed in the epithelium and the lumen.

Western blot was performed on the isolated epididymal spermatozoa to determine the responsiveness of Psap to PCB exposure. The decrease of Psap associated with epididymal sperm showed a dose-dependent manner (P<0.001, r=0.674). In mice exposed to 50 μg/
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Kg and 500 μg/Kg Aroclor 1254, sperm Psap was decreased by 42 percent and 58 percent relative to control value (Figure 3).

Benchmark dose modeling for Psap and sperm parameters

To evaluate dose-response effect of Aroclor 1254 on Psap expression, BMD modeling was utilized. The dose-response effect on Psap was compared with epididymal sperm concentration, sperm motility and sperm abnormality, which are frequently used as toxicological endpoints in animal and epidemiological studies [30]. In coincidence with our previous studies [10, 11], the sperm concentration and sperm motility were significantly reduced at 500 μg/Kg group, whereas the sperm abnormality was significantly increased in 50 μg/Kg group and 500 μg/Kg group. For Psap expression, sperm count and sperm motility, the Hill Model fit the data best based on global Goodness-of-Fit criteria, AIC, Chi-squared residuals and graphical verification (Figure 4). For sperm abnormality, however, all available models are a poor fit to the present data, which suggested that no certain estimation on BMD can be derived from the data.

Benchmark dose modeling indicated that BMD using percent reduction in sperm Psap was 8.89 μg/kg while BMD for sperm count and sperm motility was 164.67 μg/kg and 61.9 μg/kg.
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Figure 4. Comparison of dose-response effects of Aroclor 1254 on endpoint. The curve is calculated using the Hill Model. A. Decrease incidence of Psap expression in sperm in response to PCB exposure. B. Decrease incidence of epididymal sperm count in response to PCB exposure. C. Decrease incidence of epididymal sperm motility in response to PCB exposure. Data are presented as percent alternations relative to the respective control. The BMD is the concentration that elicits a response of one standard deviation over control mean. The BMDL is the concentration corresponding to the lower 95% confidence interval.

kg. The BMDLs for Psap expression, sperm count and sperm motility was 1.26 μg/kg, 15.73 μg/kg and 11.85 μg/kg, respectively.

Correlation between PCB accumulation and endpoints

Internal exposure of PCBs in mice was estimated by determining the hepatic Aroclor 1254 accumulation and was correlated with sperm Psap expression by linear regression. For comparison purpose, epididymal sperm count, sperm motility and sperm abnormality were also correlated to hepatic PCB levels. The sperm Psap expression, epididymal sperm count, sperm motility and sperm abnormality were significantly correlated to hepatic PCB accumulation (Table 1). The correlation coefficient between Psap expression and PCB accumulation was higher than that between PCB accumulation and sperm motility or sperm count. In addition, correlations between alternations in endpoints were investigated. Relative expression of Psap in sperm was significantly associated with sperm abnormality (Table 1). However, no other significant correlation between Psap expression and other endpoints was observed.

Depressed Psap in human spermatozoon is associated with PCB exposure

PCBs were detected in 13 of total 36 samples collected from participants. The corresponding donors were considered as ‘exposed’. The mean concentration of total PCBs in seminal plasma of exposed participants was 0.141 ng/ml. The age, semen parameters and hormone levels of exposed and unexposed participants were comparable (Table 2). For exposed participants, the level of Psap in sperm was 0.45±0.16 ng/mg total protein, which was significantly different to the value of 0.65±0.24 ng/mg total protein in unexposed participants. In contrast, comparison of the Psap level in seminal plasma showed no significant difference between exposed and unexposed participants. Linear regression suggested that a negative association between total PCB level in seminal plasma and Psap associated with sperm (P<0.05, r=-0.396).
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Male reproductive endpoints, including sperm concentration, motility, and morphology, sperm DNA integrity and serum reproductive hormone levels have been associated with PCB exposure [30]. In the present study, we reported that Psap, a glycoprotein detectable in sperm, were dose-dependently reduced by PCBs. The exposure doses suppressing the Psap expression were comparable to the reported daily intakes in human [25, 41], when the dose translation from animal to human is taken into consideration [35]. Within the dose-range tested in the present study, the PCB induced repression of sperm Psap might precede other male reproductive endpoints, including sperm motility. In epidemiological studies, sperm motility is the most consistent reproductive endpoint associated with PCB exposure in men and data show a lack of exposure threshold for a PCB-related effect on sperm motility [30]. However, in the present study, the BMD and BMDL were lower for Psap reduction than percent reduction in sperm motility, which might suggested that the former might be a more sensitive predictor. This point was further fortified by the pilot study based on male participants enrolled from an IVF program, in which Psap levels in sperm were associated with semen PCB concentrations in semen.

Benchmark dose approach is use to derive a point of departure (POD) in estimating the dose-response effect. The method is less dependent on dose selection and spacing, and it takes into account the shape of the dose-response curve [14]). It provides a useful tool for analyzing data with limited dose setting and small sample size, because it is difficult for the No-Observed-Adverse-Effect-Level approach to distinguish differences in sensitivity to toxic exposure in such data set. In the present study, BMD modeling was used to allow a comparison of dose-response effects between endpoints. Although it might call for a more intense dose spacing and larger data set to determine the ‘true’ BMDs lie between 50 μg/kg and 500 μg/kg, it could be clearly distinguished from the response-curves that the response of Psap precedes sperm count and motility.

The correlation between Psap reduction and the increase of sperm abnormality might suggest a functional association. The role of Psap in male reproductive system has been suggested in knock-out mouse model. Ablation of Psap gene in mice results in several abnormalities, such as a decrease in testis size and an involution of the prostate, seminal vesicle, and epididymis, although levels of testosterone in blood remain normal. The spermatogenesis of Psap deficient mice is impaired and late spermatids were particularly affected. It appears that Psap is involved in the development and maintenance of the male reproductive organs, as well as, in cellular differentiation in male reproductive system [32].

For ejaculated sperm, Psap might also be essential for fertilization. The seminal Psap

### Table 1. Correlation coefficient between Psap expression, hepatic Aroclor 1254 concentration, sperm motility and abnormality

<table>
<thead>
<tr>
<th>Endpoint</th>
<th>P</th>
<th>correlation</th>
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<tr>
<td>PCB accumulation versus sperm abnormality</td>
<td>0.001</td>
<td>0.557**</td>
</tr>
<tr>
<td>PCB accumulation versus Psap</td>
<td>0.003</td>
<td>-0.531**</td>
</tr>
<tr>
<td>PCB accumulation versus sperm motility</td>
<td>0.008</td>
<td>-0.478**</td>
</tr>
<tr>
<td>PCB accumulation versus sperm count</td>
<td>0.023</td>
<td>-0.413*</td>
</tr>
<tr>
<td>Psap versus abnormality</td>
<td>0.029</td>
<td>-0.4*</td>
</tr>
<tr>
<td>Psap versus sperm count</td>
<td>0.054</td>
<td>0.355</td>
</tr>
<tr>
<td>Psap versus sperm motility</td>
<td>0.13</td>
<td>0.283</td>
</tr>
</tbody>
</table>

**”indicates significant at P<0.05; “***”indicates significant at P<0.01.

### Table 2. Characteristics of participants exposed and unexposed to PCBs

<table>
<thead>
<tr>
<th></th>
<th>Exposed</th>
<th>Unexposed</th>
<th>P</th>
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</thead>
<tbody>
<tr>
<td>Age (yr)</td>
<td>32.62±5.80</td>
<td>33.30±5.35</td>
<td>0.72</td>
</tr>
<tr>
<td>Semen volume (ml)</td>
<td>2.46±0.80</td>
<td>2.40±0.92</td>
<td>0.85</td>
</tr>
<tr>
<td>Sperm concentration (10⁹/ml)</td>
<td>43.46±12.14</td>
<td>44.78±14.58</td>
<td>0.78</td>
</tr>
<tr>
<td>Motile sperm (%)</td>
<td>57.30±23.31</td>
<td>53.53±20.70</td>
<td>0.38</td>
</tr>
<tr>
<td>Normal form (%)</td>
<td>9.1±4.29</td>
<td>10.2±5.4</td>
<td>0.348</td>
</tr>
<tr>
<td>E2 (pg/ml)</td>
<td>22.73±7.44</td>
<td>27.91±9.52</td>
<td>0.966</td>
</tr>
<tr>
<td>T (ng/ml)</td>
<td>4.66±1.29</td>
<td>5.07±1.62</td>
<td>0.715</td>
</tr>
<tr>
<td>FSH (mIU/ml)</td>
<td>4.69±1.41</td>
<td>6.01±4.03</td>
<td>0.106</td>
</tr>
<tr>
<td>LH (mIU/ml)</td>
<td>3.83±1.07</td>
<td>4.63±2.01</td>
<td>0.359</td>
</tr>
<tr>
<td>PRL (ng/ml)</td>
<td>10.87±5.76</td>
<td>9.79±4.67</td>
<td>0.243</td>
</tr>
<tr>
<td>Fertilization rate (%)</td>
<td>79.79±22.28</td>
<td>81.26±18.92</td>
<td>0.83</td>
</tr>
</tbody>
</table>

### Discussion

Male reproductive endpoints, including sperm concentration, motility, and morphology, sperm DNA integrity and serum reproductive hormone levels have been associated with PCB exposure [30].
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facilitates the sperm-egg interaction through a 60-amino acid sequence fragment, which is conserved among human and animals [2-4, 17, 29]. It is demonstrated that the pregnancy rate for thawed sperm from bulls is increased when the sperm is exposed to the fragment [3]. It appears that Psap itself is equally as good as the fragment in restoring sperm binding capability [17]. Although the mechanism of action is unknown, these studies suggest that the role of Psap was increasing the number of sperm binding to the outer egg layer, zona pellucida. It is postulated that insufficient amount of the protein essential for the initial binding of a spermatozoon to an egg investment might result in poor fertilizing potential of sperm, representing either a substantial proportion of the cells in a semen sample from certain subfertile individuals or most samples of frozen-thawed sperm [4]. Previous study has demonstrated that sperm-oocyte interaction is significantly reduced in PCB treated male rats [21]. If the Psap levels in sperm are sensitive to environmental exposure, would it possible that depressed Psap levels contribute to the prolonged TTP associated with PCB exposure? Our data failed to show any significant difference between exposed and unexposed participants in fertilization rate. However, due to the limited size of study cohort and incapability to adjust for potential confounders, it was unable to conclude that depressed Psap levels in sperm without major changes in semen parameters have no effect on fertilization rate.

Psap in male reproductive tract is secreted by Sertoli cells as well as epididymal principal cells [31]. In testis, Psap exists as a secretory protein as well as a lysosomal protein [24]. The Psap secreted by Sertoli cells binds to the tail of late spermatids and spermatozoa and is present in a relatively high concentration in luminal fluids from the rat rete testis, efferent ducts [23]. On the other hand, the fact that Psap is also synthesized in the epithelial cells in the extra testicular tract suggest that this protein might play an integral role through the process of sperm maturation [20]. In the efferent ducts, the Psap from testes is endocytosed by the nonciliated cells and the abundance decreased dramatically [23], which might suggest the differential role of Psap in testis and epididymis. In the present study, the expression of Psap in testis and epididymis is differentially regulated. Within the dose-range tested in the present study, no significant alteration of Psap expression is observed in testes. In contrast, the caput epididymides showed a decrease in Psap expression at both mRNA and protein levels after PCB exposure, which might suggest dysfunction in epididymides might be response for the declining Psap levels in sperm. Once again, it suggested that epididymis is a sensitive target organ contributing to PCB induced male reproductive toxicity, as it have been indicated in epidemiological studies [8].

The Psap level is regulated in vivo and in vitro as a result of steroid hormone actions. In the...
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endometrial epithelium, Psap expression increased in response to elevated levels of estrogen during the oestrous cycle [37]. Supplement of 17β-estradiol in breast cancer culture dose-dependently increased secretion of Psap into medium [12]. Using prostate cell line as an in vitro model, Koochekpour and colleagues show that the expression of Psap is under regulation of androgen receptor [26]. The toxic effects of PCBs on male reproductive system are associated with their ability to disrupt both the expression/activity of enzymes required for sex steroid synthesis/catabolism, and the expression/ability of hormone receptors to bind endogenous ligands. Thus, it is reasonable to postulate that the expression of Psap is responsive to endocrine disrupting activities of PCBs. However, there is a report indicating that the immunohistological staining of Psap in epididymal principal cells, as well as in narrow, clear, and basal cells, remain unchanged after orchidectomy with or without testosterone supplementation, efferent duct ligation, or hypophysectomy [19], which suggests an alternative mode of regulation on Psap. To clarify the specific effects of PCBs on Psap expression, additional experiments with PCB congeners in different mode of action are still required.

In summary, we have described the dose-dependent reduction of Psap in male reproductive tract after PCB exposure. The Psap levels in sperm were adversely associated with external and internal PCB exposure, in mice and men. The PCB induced depression of Psap preceded significant effects on sperm motility and concentration. Adversely association between Psap relative expression and sperm abnormality suggested a correlation between Psap levels and sperm function. Taken together, our data suggested that determining the expression of Psap in sperm might provide a link between PCB exposure and exposure-mediated damage to target organs, and have the potential to become a useful predictor of PCB related decrease in fertility.

Acknowledgements

The study was supported by and National Key Technology R&D Program of China (2007BAC27B02), the National Funds for the Natural Sciences (81302454) and the Major Science and Technology Platform in Xiamen City (No. 3502z20111006).

Disclosure of conflict of interest

None.

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