Influence of Bisphenol A on Thyroid Volume and Structure Independent of Iodine in School Children

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Abstract

Background

Although several studies have evaluated the relationship between bisphenol A (BPA) and thyroid functions, their results are not entirely consistent. Little is known about BPA in relation to thyroid volume and structure.

Methods

We examined the association of BPA with thyroid volume and thyroid nodules using data from 718 Chinese children living in the East Coast of China in 2012. First morning urine samples were collected for the determination of urinary BPA, creatinine, and urinary iodine concentrations (UIC). Thyroid volume (TV) and nodules were assessed by thyroid ultrasonography.

Results

The median of TV was 3.14ml. 459 (63.9%) children took iodized salt at home and the median of UIC was 159 μg/l. BPA was detected in 99.9% of the urine samples and the medians for boys and girls were 2.64 and 2.35 μg/g creatinine, respectively. Of all participants 14.0% had thyroid nodules. Urinary BPA concentration was inversely associated with thyroid volume (β = -0.033, 95% CI: -0.053, -0.013) and the risk for multiple nodules (OR = 0.78; 95% CI: 0.63, 0.97). The associations above were similar for children who consumed iodized salt and those consumed non-iodized salt.
Conclusions

The data suggest that BPA may be one of the influencing factors for TV and thyroid nodules and its effects are independent of iodine nutrition status in children.

Introduction

Thyroid volume (TV) is usually measured to evaluate the goiter status and then to assess the degree of iodine deficiency in a population [1]. The influencing factors of TV in children include age, sex, body surface area (BSA), pubertal stage, and iodine nutritional status [2–4]. In addition to iodine nutrition status, other chemicals could also contribute to the development of non-toxic diffuse goiter [5]. Recent epidemiological studies have suggested that Bisphenol A (BPA) disrupts thyroid function in adults [6–9]. BPA is one of the highest production and consumption volume chemicals in the world [8]. Studies from different countries and areas have demonstrated the presence of urinary BPA in more than 90% of their study populations, suggesting a common exposure to BPA worldwide [7, 10–11]. Exposure to BPA is prevalent among children and adolescents [10], who may be more sensitive to its adverse effects including the effect on thyroid function.

The endocrine-disrupting effect of BPA might contribute to numerous complex outcomes including diabetes [12], obesity [13–15], and some other medical disorders [16] such as cognitive and behavioral disorders [17–19]. BPA and other environmental disrupting chemicals (EDCs) may influence the levels of thyroid and steroid hormones and result in increased risks of obesity and some other chronic conditions [20].

Either excess iodine intake or iodine deficiency was associated with an increased risk of developing thyroid nodules [21]. In order to reduce the iodine deficiency, the Universal Salt Iodization (USI) program was launched in China in October 1994 [22]. This program has been implemented for over 20 years and iodized salt remains the main dietary source of iodine. USI has significantly contributed to the prevention of iodine deficiency disorder (IDD) worldwide [23]. However, excess iodine consumption may also have adverse public health impacts [24]. It has been argued that the uniform iodized salt criterion might not have worked across China as expected [22]. The iodine status in children and adults is above an adequate level (100–200μg/L) in some areas and researchers have suggested to reduce the iodine intake accordingly [25]. It is not known if iodine nutrition status has an impact on the association between BPA thyroid measures.

In the current study, we explored the influence of BPA, in addition to iodine, on thyroid volume as well as thyroid nodules. Iodine intake is required for the production of thyroid hormone, and iodine nutrition would change thyroid volume and structure. Both thyroid volume and thyroid nodules [21] can be associated with abnormalities of thyroid function. We therefore also accessed the associations between BPA and thyroid outcomes in children according to their iodine nutrition status.

Materials and Methods

Study population

Three primary schools were selected from Minhang District in Shanghai, Haimen City in Jiangsu Province and Taizhou City in Zhejiang Province, respectively. Four classes in each grade from grade 3 to grade 5 in these schools were randomly selected and all students without
preexisting thyroid conditions in selected classes were enrolled into this study. Since thyroid
evaluations were not routinely conducted in children, no student reported preexisting thy-
roid abnormalities. Among 1267 children enrolled, 1248 provided first morning urine samples
and 1234 students completed routine physical examinations. An ultra-sound test for thyroid
gland volume was performed on 1064 students. BPA and urinary creatinine were measured for
803 urine samples. After excluding those with no urinary test or thyroid test, we included 718
students in the current analysis. Written consent from parents or guardians of all participants
were received and the study was approved by the Ethical Review Board of the School of Public
Health of Fudan University.

Anthropometric measurements

Anthropometric measurements, including standing height (cm), weight (kg), and circumfer-
ences of the waist, hip and chest (cm) were taken by trained health professionals according to a
standard protocol. The standing height was measured to the nearest 0.1cm without shoes.
Weight was measured to the nearest 0.1kg using a digital weight scale. Body mass index (BMI)
was calculated as weight in kilograms divided by the square of height in meters and over-
weight/obese status were assessed using the BMI growth reference values for Chinese children
[26].

Test for thyroid gland volume and thyroid nodules

Students in Taizhou and Haimen received thyroid ultrasonography performed by experienced
examiners in local comprehensive hospitals. For students in Minhang, thyroid ultrasonography
was performed at school using a real-time sector scanner with a 7.5-MHz/40-mm probe linear
transducer. The ultrasonographic examination was carried out on the children lying on a desk
with the neck extended. The volume of each lobe was calculated by using the following formula:
volume of one lobe (mL) = 0.479*maximum thickness*maximum width*maximum length
(cm)[27]. The total thyroid volume was the sum of both lobes and the isthmus volume was not
included. Standardized thyroid ultrasound technique was adopted according to the method
described by Fuse et al [28]. According to the Chinese national criteria for thyroid measure-
ment, goiter was defined by age-specific thyroid volume. The upper limits of thyroid volume
for children aged 9, 10, and 11 years were 5.0, 6.0 and 7.0, respectively. If thyroid volume
exceeded the relevant limit, a child was judged as being goitrous [29]. Discrete lesion(s) within
the thyroid gland that is palpably and / or ultrasonographically distinct from the surrounding
thyroid parenchyma were defined as thyroid nodule(s) [30]. It was also detected by ultrasonog-
raphy in our study. In case of abnormality in the sonographic examination of the thyroid,
parents of the children would receive a written note describing the abnormal results of the
examination.

Urine and salt samples collection and iodine concentration analyses

The day prior to a physical examination, a 50ml BPA free tube was distributed to each student.
First morning urine samples were collected in children’s home and brought to school by stu-
dents themselves in the next morning. The collected urine samples were kept frozen at -80°C
until analysis. Students were also asked to bring a salt sample of more than 20g from home for
iodine measurement.

Urinary iodine concentrations (UICs) were determined following the method proposed by
the Ministry of Health of the People’s Republic of China (WS/T107. 2006, and GB/T13025.7–
1999) [27]. Salt iodine content was also measured using a national standard method with a
proper quality control [31](GB/T 13025.7–1999). 10% urine samples were assayed in duplicate.
Analyses of urine BPA

Details of the assays for the detection of BPA have been published previously [32]. Briefly, after an aliquot (1.0 mL) of urine sample was enzymatically hydrolyzed, BPA was purified and separated by reversed-phase and anion-exchange mixed-mode solid-phase extraction. It was analyzed by ultra-performance liquid chromatography coupled with tandem mass spectrometry. It was derivatized by dansyl chloride, and then resolved by the mobile phase of methanol and water both containing 10 mM of ammonium formate on reversed-phase ultra-performance liquid chromatographic C18 column and ionized in positive ion mode. BPA was quantified by using an isotope internal standard curve method.

All the analyses of urine samples were completed in twelve batches within two months. Four spiked urine samples with BPA at the concentration of 5 ng/mL and four solvent blanks were prepared along with each batch to monitor the background values of phthalate metabolites and the accuracy of analytical procedure. The inter-batch recoveries of BPA of spiked urine samples varied between 81% and 121% with the inter-batch relative standard deviations (RSDs) ranging from 6% to 16%.

Urinary creatinine concentration in each sample was also measured using an enzymatic method. BPA concentrations were then adjusted for urinary dilution by creatinine levels.

Table 1. Characteristics of participants in the analysis and non-participants.

| Characteristics | Non-participants | Participants in the analysis | χ² | P-value |
|-----------------|------------------|----------------------------|----|---------|
|                 | N %              | N %                        |    |         |
| Total           | 549 100.00       | 718 100.00                 |    |         |
| Sex             |                  |                            | 1.431 | 0.232 |
| Male            | 262 47.72        | 367 51.11                  |    |         |
| Female          | 287 52.28        | 351 48.89                  |    |         |
| Age             |                  |                            | 75.772 <0.0001 |        |
| 9               | 119 21.68        | 321 44.71                  |    |         |
| 10              | 234 42.62        | 228 31.75                  |    |         |
| 11              | 196 35.70        | 169 23.54                  |    |         |
| BMI a           |                  |                            | 7.063 | 0.029 |
| Normal          | 379 69.03        | 560 77.99                  |    |         |
| Overweight      | 89 16.21         | 93 12.95                   |    |         |
| Obese           | 56 10.20         | 57 7.94                    |    |         |
| Iodized salt consumption b |          |                            | 0.807 | 0.369 |
| No              | 172 31.33        | 259 36.07                  |    |         |
| Yes             | 340 61.93        | 459 63.93                  |    |         |
| Urinary iodine(μg/l) c |              |                            | 0.935 | 0.626 |
| <100            | 131 23.86        | 167 23.26                  |    |         |
| 100–200         | 225 40.98        | 297 41.36                  |    |         |
| >200            | 174 31.69        | 254 35.38                  |    |         |

a: 25 missing
b: 37 missing
c: 19 missing

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Statistical analysis

Values below the method limit of detection (LOD) for BPA (0.06ng/ml) were replaced with a value of the LOD divided by the square root of 2 \[33\]. Since data were not normally distributed, their logarithmically transformed values were therefore used, including thyroid volume, BPA, BSA (BSA = 0.007184 * weight^{0.425} / height^{0.725}) and BMI. One-way analysis of variance (ANOVA) was performed to test the difference in medians of thyroid volume and urinary BPA concentration between groups categorized by sex, age, BSA (in tertiles), iodized salt consumption status, urinary iodine level and study area. Simple linear regression analysis was used to examine the correlations between thyroid volume and its determinants. Multivariable linear regression models were used to adjust for age, sex, iodized salt consumption, BSA (log-transformed) and BMI (log-transformed). The median levels of log-transformed BPA across quintiles of TV were calculated and then compared using ANOVA. Due to a considerable day-to-day variation in iodine excretion, one-spot urinary iodine level was not a proper indicator of iodine status for individuals \[34\]. Therefore, in current analysis, iodized salt consumption instead of iodine concentration in urine, was included in the multivariate models. We also assessed the correlation between UIC and iodized salt consumption and observed a significantly higher level of UIC in children who consumed iodized salt at home, suggesting that iodized salt consumption status could be a good proxy for iodine nutrition at a population level.

The association of urinary BPA concentration (in quintiles) with thyroid nodules (no, solitary nodule, and multiple nodules) was assessed using multinomial logistic regression analysis. All analyses were performed by using SAS, version 9.3 software (SAS Institute, Inc., Cary, NC, USA), and all tests of statistical significance were base on two-side probability.

Results

Participant characteristics

A total of 1267 students were recruited for this survey and 718 of them were included in the present analysis. The demographic characteristics as well as iodine nutrition status were summarized in Table 1. There were no statistically significant differences in sex and iodized nutrition status between children included in this analysis (N = 718) and those excluded (549). The median of UIC for children included in the analysis was 159μg/l.

Thyroid volume measurements

The median thyroid volume of participants was 3.14ml, and was similar for boys (3.05ml) and girls (3.21ml). Thyroid volume significantly increased with age, BSA and urine iodine concentration (Table 2). Children consuming iodized salt had a relatively larger thyroid gland volume as compared with those consuming non-iodized salt. The proportions of children consuming iodized salt in Minhang, Haimen and Taizhou were 75%, 94% and 25%, respectively, which partly explained a significantly lower level of TV in Taizhou. According to the thyroid volume-based criteria by age for screening of goiter in children, 6% (15 boys and 25 girls) had goiter.

Urinary BPA concentrations

BPA was detected in 99.9% of the urine samples. Median BPA concentrations were similar for boys (2.64μg/g creatinine) and girls (2.35μg/g creatinine), but increased with age (p-trend = 0.028) (Table 2). The urinary BPA concentration also showed a geographic difference. The median BPA concentration was the lowest in Minhang (1.74μg/g creatinine), the highest in Taizhou (3.95μg/g creatinine), and in between in Haimen (2.22μg/g creatinine) (Table 2).
There was a significant inverse association between BPA concentration and TV (Fig 1). A simple linear regression model for the association showed a regression coefficient (β) of -0.036 (95% CI: -0.056, -0.016). After adjustment for age, sex, (ln)BSA and iodized salt consumption status, the association persisted (Table 3). Each log unit increase in BPA level was associated with a 0.088 SD unit decrement in thyroid volume. When we used BMI instead of BSA, and used urinary iodine concentration instead of iodized salt consumption as covariates, the inverse association between thyroid volume and BPA remained significant. In addition, the inverse associations did not differ significantly by sex, or iodized salt consumption status (Table 3).

Urinary concentration of BPA was not significantly associated with the risk of goiter while iodized salt consumption was related to a reduced risk for goiter (Odds Ratio (OR): 0.34; 95% CI: 0.14–0.84) (data not shown).

Prevalence of thyroid nodules

Thyroid nodules were detected in 100 children, accounting for 14% of all participants. Most nodules were accompanied by hypoechoogenicity. The prevalence of multiple thyroid nodules

Table 2. Thyroid volume and urinary BPA concentration of participants characterized by sex, age, BSA, area and iodine nutrition status.

| Characteristics                   | Thyroid Volume Median (IQR) | P-value | Urinary BPA concentration a Median (IQR) | P-value |
|-----------------------------------|-----------------------------|---------|----------------------------------------|---------|
| All                               | 3.14(2.44–4.11)             | 0.263   | 2.45(1.09–5.97)                        | 0.126   |
| Sex                               |                             |         |                                        |         |
| Male                              | 3.05(2.43–4.03)             |         | 2.64(1.13–6.40)                       |         |
| Female                            | 3.21(2.45–4.25)             |         | 2.35(1.04–5.40)                       |         |
| Age (years)                       |                             | <0.0001 |                                        | 0.028   |
| 9                                 | 2.92(2.25–3.85)             |         | 2.24(1.04–5.34)                       |         |
| 10                                | 3.74(2.75–4.70)             |         | 2.54(0.98–6.41)                       |         |
| 11                                | 3.02(2.46–3.58)             |         | 2.89(1.27–6.34)                       |         |
| BSAb                              |                             | <0.0001 |                                        | 0.050   |
| T1 (<1.06 m²)                     | 2.57(2.05–3.45)             |         | 2.26(1.01–5.12)                       |         |
| T2 (1.06–1.19 m²)                 | 3.31(2.59–4.41)             |         | 2.49(1.05–5.58)                       |         |
| T3 (≥ 1.20 m²)                    | 3.42(2.70–4.54)             |         | 3.02(1.27–6.85)                       |         |
| Iodized salt consumption           |                             | <0.0001 |                                        | <0.0001 |
| No                                | 2.73(2.16–3.50)             |         | 3.44(1.74–7.02)                       |         |
| Yes                               | 3.35(2.61–4.43)             |         | 2.14(0.85–5.21)                       |         |
| Urinary iodine (μg/l)             |                             | 0.0003  |                                        | 0.783   |
| <100                              | 2.75(2.21–3.55)             |         | 2.29(1.04–4.58)                       |         |
| 100–200                           | 3.18(2.45–4.11)             |         | 2.44(1.02–6.35)                       |         |
| >200                              | 3.33(2.55–4.43)             |         | 2.49(1.15–6.36)                       |         |
| Area                              |                             | <0.0001 |                                        | <0.0001 |
| Minhang                           | 4.25(3.44–5.17)             |         | 1.74(0.94–3.43)                       |         |
| Haimen                            | 3.13(2.46–4.07)             |         | 2.22(0.78–5.20)                       |         |
| Taizhou                           | 2.52(2.02–3.07)             |         | 3.95(2.04–10.48)                      |         |

BPA: Bisphenol A; BSA: Body Surface Area; IDR: Inter-quartile range

a: Creatinine adjusted (μg/mg)
b: In tertiles

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Association between urinary BPA concentration and thyroid volume

There was a significant inverse association between BPA concentration and TV (Fig 1). A simple linear regression model for the association showed a regression coefficient (β) of -0.036 (95% CI: -0.056, -0.016). After adjustment for age, sex, (ln)BSA and iodized salt consumption status, the association persisted (Table 3). Each log unit increase in BPA level was associated with a 0.088 SD unit decrement in thyroid volume. When we used BMI instead of BSA, and used urinary iodine concentration instead of iodized salt consumption as covariates, the inverse association between thyroid volume and BPA remained significant. In addition, the inverse associations did not differ significantly by sex, or iodized salt consumption status (Table 3). Urinary concentration of BPA was not significantly associated with the risk of goiter while iodized salt consumption was related to a reduced risk for goiter (Odds Ratio (OR): 0.34; 95% CI: 0.14–0.84) (data not shown).
was 6.9%, compared with 7.1% for solitary nodules. The risk for thyroid nodules was negatively associated with age but showed no sex related difference. There was no association of thyroid nodules with urinary iodine level or iodized salt consumption status (Table 4).

Table 3. Linear regression analysis for association of thyroid volume with BPA (both log-transformed).

| Model | \( \beta \) | 95%CI | P-value |
|-------|-------------|-------|---------|
| All participants Univariate analysis Model 1\(^a\) | -0.036 | -0.056 | -0.016 | <0.0001 |
| Multivariate analysis Model 2\(^b\) | -0.038 | -0.058 | -0.018 | <0.0001 |
| Model 3\(^c\) | -0.033 | -0.053 | -0.013 | 0.001 |
| Model 4\(^d\) | -0.037 | -0.056 | -0.018 | <0.0001 |
| Sex Male Model 3\(^c\) | -0.039 | -0.064 | -0.015 | 0.002 |
| Female Model 3\(^c\) | -0.032 | -0.062 | -0.003 | 0.034 |
| Iodized salt consumption Yes Model 3\(^c\) | -0.029 | -0.062 | 0.003 | 0.074 |
| No Model 3\(^c\) | -0.037 | -0.061 | -0.014 | 0.002 |

\(^a\): Univariate regression analysis  
\(^b\): Adjusted for sex and age only  
\(^c\): Adjusted for sex, age, (ln)BSA, and iodized salt consumption status  
\(^d\): Adjusted for sex, age, (ln)BMI, and urinary iodine concentration (categorical variable: < 100µg/l, 100–200µg/l and >200µg/l)

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Associations between urinary BPA concentrations and thyroid nodules

Urinary BPA concentration was negatively associated with the risk of multiple nodules, but not with the risk of solitary nodules before and after controlling for sex, age, iodized salt consumption and BMI (in quintiles). There was a reduced risk for multiple nodules with increasing BPA, with an OR being 0.78 (95% CI: 0.63, 0.97). No notable variations in OR estimates were observed across sex, age, and iodized salt consumption categories (Table 5).

Table 4. Prevalence of thyroid nodules in participants categorized by sex, age, and iodine nutrition.

| Characteristics         | N   | No nodule (N (%)) | Thyroid nodules Solitary nodule (N (%)) | Multiple nodules (N (%)) | P-value |
|-------------------------|-----|-------------------|----------------------------------------|-------------------------|---------|
| All                     | 718 |                   |                                        |                         |         |
| Sex                     |     |                   |                                        |                         | 0.413   |
| Male                    | 367 | 321(87.47%)       | 21 (5.72%)                             | 25 (6.81%)              |         |
| Female                  | 351 | 297(84.61%)       | 28 (7.98%)                             | 26 (7.41%)              |         |
| Age                     |     |                   |                                        |                         | 0.004   |
| 9                       | 321 | 287(89.41%)       | 21 (6.54%)                             | 13 (4.05%)              |         |
| 10                      | 228 | 191(83.78%)       | 19 (8.33%)                             | 18 (7.89%)              |         |
| 11                      | 169 | 140(82.84%)       | 9 (5.33%)                              | 20 (11.83%)             |         |
| Iodized salt consumption|     |                   |                                        |                         | 0.510   |
| No                      | 259 | 228(88.03%)       | 16 (6.18%)                             | 15 (5.79%)              |         |
| Yes                     | 459 | 390(84.97%)       | 33 (7.19%)                             | 36 (7.84%)              |         |
| Urinary iodine(μg/l)    |     |                   |                                        |                         | 0.202   |
| <100                    | 167 | 144(86.22%)       | 11 (6.59%)                             | 12 (7.19%)              |         |
| 100–200                 | 297 | 247(83.17%)       | 23 (7.47%)                             | 27 (9.09%)              |         |
| >200                    | 254 | 228(89.77%)       | 14 (5.51%)                             | 12 (4.72%)              |         |
| Area                    |     |                   |                                        |                         | <0.0001 |
| Shanghai                | 200 | 150(75.00%)       | 25 (12.50%)                            | 25 (12.50%)             |         |
| Haimen                  | 261 | 230(88.12%)       | 13 (4.98%)                             | 18 (6.90%)              |         |
| Taizhou                 | 257 | 238(92.61%)       | 11 (4.28)                              | 8 (3.11%)               |         |

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Associations between urinary BPA concentrations and thyroid nodules

Table 5. Associations of urinary BPA concentration with thyroid nodules.

|          | Solitary nodule OR (95%CI) | Multiple nodules OR (95%CI) |
|----------|-----------------------------|----------------------------|
| All subjects | Univariate analysis         | 0.98 (0.79, 1.20)          | 0.77 (0.63, 0.95)       |
|          | Multivariate analysis b     | 0.98 (0.79, 1.20)          | 0.76 (0.61, 0.94)       |
| Sex b    | Male 1.07 (0.79, 1.47)      | 0.82 (0.61, 1.09)          |                          |
|          | Female 0.90 (0.68, 1.21)    | 0.69 (0.49, 0.96)          |                          |
| Age(years) b | 9 1.20 (0.87, 1.66)    | 0.68 (0.43, 1.09)          |                          |
|          | 10 0.90 (0.64, 1.26)       | 0.74 (0.52, 1.06)          |                          |
|          | 11 0.72 (0.43, 1.22)       | 0.80 (0.56, 1.15)          |                          |
| Iodized salt consumption b | Yes 1.01 (0.79, 1.30) | 0.74 (0.57, 0.96)          |                          |
|          | No 0.92 (0.62, 1.35)       | 0.79 (0.53, 1.19)          |                          |

a: Compared with participants with no thyroid nodules
b: The results of multiple analysis adjusting for age, sex, BSA (in quintiles), and iodized salt consumption state

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Discussion

In this study of school children conducted in east coast China, we observed diversity in thyroid volume with varied iodine nutrition status. Urinary BPA concentration was inversely associated with thyroid volume, but not with goiter. An increased BPA level was also associated with a reduced risk for multiple thyroid nodules.

Studies that measured thousands of individuals from several different countries overwhelmingly detected BPA in adults, adolescents, and children [35]. Considering the endocrine disruptor effect of BPA and its common exposure, many studies have focused on the potential effects of BPA on various chronic conditions, including diabetes, obesity [14–15], as well as thyroid functions [7].

Epidemiologic studies have revealed an association between BPA exposure and altered thyroid hormones. A cross-sectional study conducted in Shanghai examined this association in 3394 subjects and found increased free triiodothyronine (FT3) and decreased TSH with increased urinary BPA [7]. High urinary BPA level was associated with increased thyroid function (adjusted OR: 1.71, 95% CI: 1.26, 2.32)[7]. The NHANES study also reported a suggestive inverse relationship between urinary BPA and total T4 and TSH among adults [6]. Another survey in a Thai population observed a significantly negative correlation between serum BPA and FT4 level in males only, but BPA was not associated with TSH in either males or females [8]. The CHAMACOS study revealed that exposure to BPA during pregnancy was related to reduced T4 in pregnant women and decreased TSH in male neonates [36]. BPA was observed to have a direct effect on thyroid follicular cell and leads to an altered expression of the genes involved in thyroid hormones synthesis both in vitro and in vivo (zebrafish) models [37].

Other potential mechanisms for the association between BPA and thyroid hormones include inhibiting T3 pathways during metamorphosis [38] and thyroid hormone receptor (TR) transcription suppression [39].

Children are commonly exposed to BPA [10, 19], and those having higher BPA levels experience an elevated risk for being obese/overweight [14, 40]. Obesity is now becoming a great public health concern and is closely linked to numerous adverse health outcomes. Studies suggest that altered thyroid function may be one of the reasons for the relationship between BPA and obesity [41]. However, the association of BPA with abnormality in thyroid morphology and structure has not been well studied previously.

The main determinants of thyroid volume in both boys and girls have been shown to be age, urinary iodine, BSA, and pubertal stage. The findings of our study of Chinese children that thyroid volume was related with age, BSA, and iodine nutritional status are in a general agreement with those of previous studies conducted among other populations [2]. In another study conducted in Zhejiang Province, China, age, sex and BSA were the influencing factors for thyroid volume in children aged 6–12 years, whereas urinary iodine concentration had little effect on the TV [3]. In the study conducted in healthy Greek children by Kaloumenou et al [2], main determinants of thyroid volume in children were age, body surface area, and pubertal stage. The prevalence of thyroid nodule was also significantly increased with age and showed a female predominance [42]. Another study conducted in China indicated that excess iodine intake or iodine deficiency was also risk factors for thyroid nodules [21].

Decreased secreting of thyroxin and subsequent rise in TSH may explain the association of goiter and iodine deficiency [25]. BPA suppresses TSH release from pituitary in a manner independent of both the thyroid hormone feedback mechanism and the estrogenic activity of BPA [6]. Therefore, the inverse association between thyroid volume and BPA observed in our study may partly due to the negative correlation between BPA and TSH. Children exposed to a higher level of BPA may have a reduced TSH and then prohibited the enlargement of thyroid gland.
The exposure of BPA may also alter thyroid functions, leading to thyroid abnormalities, such as subclinical hypothyroidism. The risk for developing thyroid nodules could therefore be changed [21], which might partly explain the association of BPA with thyroid nodules.

The sex difference in the BPA effect on the prevalence of thyroid nodules was consistent with findings from experimental studies and previous epidemiological studies on its effects on other chronic outcomes. BPA was associated with overweight/obesity among girls aged 9–12, but the association was not observed among boys [14]. A prospective birth cohort documented that BPA was detected in more than 97% of the gestational and childhood urine samples. The magnitude of the gestational BPA associations differed according to child’s sex, gestational BPA exposure affected behavioral and emotional regulation domains in children at 3 years of age, especially among girls [19]. Wang et al. conducted a cross-sectional study in children aged 8–15 and found a significant association between urinary BPA and BMI in females but not in males. [40] This sex-related discrepancy is possibly related to androgen-related differences in the metabolism of BPA [8]. It is also argued that BPA is estrogenic and capable of disrupting sex differentiation [38].

The effects of BPA on TV or thyroid volume were similar in participants consuming iodized salt and those consuming non-iodized salt, which suggested that the effects of BPA on TV and thyroid nodules were independent of iodine exposure.

The association of thyroid volume or thyroid echostructure with other chemicals in addition to iodine nutrition status, has been seldom explored [43]. To our knowledge, this is the first study on the potential effect of BPA on thyroid volume and thyroid nodules. Our study has some limitations. As a cross-sectional study, we did not have repeated measures of the urinary BPA concentration and thyroid volume overtime and did not evaluate long-term effects of BPA on TV. Information on the amount of salt consumed at home and outside of home was not collected and we can only use iodized salt consumption status as proxy of iodine nutrition, which was another limitation of our study. In addition, we had no information on thyroid hormones and were unable to evaluate whether these factors might explain the observed associations.

In conclusion, we observed an inverse association between urinary BPA with thyroid volume and risk for multiple thyroid nodules in children living in the East Coast of China. Further studies are needed to investigate the temporal relationship and potential public health implications of such an association.

**Author Contributions**

Conceived and designed the experiments: NW YZ CWF PXH HF QZ YC QWJ. Performed the experiments: NW HXW BW FJ QZ. Analyzed the data: NW YC CWF HXW BW QZ YC. Contributed reagents/materials/analysis tools: PXH MFS FJ HF QWJ. Wrote the paper: NW QZ YC.

**References**

1. Peterson S, Sanga A, Eklof H, Bunga B, Taube A, Gebre-Medhin M, et al. Classification of thyroid size by palpation and ultrasonography in field surveys. Lancet. 2000; 355(9198):106–10. PMID: 10675168
2. Kaloumenou I, Aleivizaki M, Ladopoulos C, Antoniou A, Duntas LH, Mastorakos G, et al. Thyroid volume and echostructure in schoolchildren living in an iodine-replete area: relation to age, pubertal stage, and body mass index. Thyroid. 2007; 17(9):875–81. PMID: 17956161
3. Zou Y, Ding G, Lou X, Zhu W, Mao G, Zhou J, et al. Factors influencing thyroid volume in Chinese children. Eur J Clin Nutr. 2013; 67(11):1138–41. doi: [10.1038/ejcn.2013.173](http://dx.doi.org/10.1038/ejcn.2013.173) PMID: 24065067
4. Liu P, Liu SJ, Su XH, Zhang SB, Ji XH. Relationship between urinary iodine and goiter prevalence: results of the Chinese national iodine deficiency disorders survey. J Endocrinol Invest. 2010; 33 (1):26–31. PMID: 19494707
5. Savchenko OV, Toupeleev PA. Lead, cadmium, manganese, cobalt, zinc and copper levels in whole blood of urban teenagers with non-toxic diffuse goiter. Int J Environ Health Res. 2012; 22(1):51–9. doi: 10.1080/09603033.2011.588324 PMID: 21660794

6. Meeker JD, Ferguson KK. Relationship between urinary phthalate and bisphenol A concentrations and serum thyroid measures in U.S. adults and adolescents from the National Health and Nutrition Examination Survey (NHANES) 2007–2008. Environ Health Perspect. 2011; 119(10):1396–402. doi: 10.1289/ehp.1103582 PMID: 21749963

7. Wang T, Lu J, Xu M, Xu Y, Li M, Liu Y, et al. Urinary bisphenol A concentration and thyroid function in Chinese adults. Epidemiology. 2013; 24(2):295–302. PMID: 23337242

8. Sripraphradang C, Chailurkit LO, Aekplakorn W, Ongphiphadhanakul B. Association between bisphenol A and abnormal free thyroxine level in men. Endocrine. 2013; 44(2):441–7. doi:10.1007/s12020-013-9889-y PMID: 23377699

9. Meeker JD, Calafat AM, Hauser R. Urinary bisphenol A concentrations in relation to serum thyroid and reproductive hormone levels in men from an infertility clinic. Environ Sci Technol. 2010; 44(4):1458–63. PMID: 20030380

10. Wang B, Wang H, Zhou W, He Y, Zhou Y, Chen Y, et al. Exposure to bisphenol A among school children in eastern China: A multicenter cross-sectional study. J Sci Environ Epidemiol. 2014. PMID: 24866264

11. Calafat AM, Ye X, Wong LY, Reidy JA, Needham LL. Exposure of the U.S. population to bisphenol A and 4-tertiary-octylphenol: 2003–2004. Environ Health Perspect. 2008; 116(1):39–44. PMID: 18197297

12. Silver MK, O'Neill MS, Sowers MR, Park SK. Urinary bisphenol A and type-2 diabetes in U.S. adults: data from NHANES 2003–2008. PLoS One. 2011; 6(10):e26868. doi: 10.1371/journal.pone.0026868 PMID: 22046388

13. Trasande L, Attina TM, Blustein J. Association between urinary bisphenol A concentration and obesity prevalence in children and adolescents. JAMA. 2012; 308(11):1113–21. PMID: 22990270

14. Xu XH, Hong X, Xie LD, Li T, Yang YJ, Zhang Q, et al. Gestational and lactational exposure to bisphenol-A affects anxiety- and depression-like behaviors in mice. Horm Behav. 2012; 62(4):480–90. PMID: 23776476

15. Eng DS, Lee JM, Gebremariam A, Meeker JD, Peterson K, Padmanabhan V. Bisphenol A and chronic disease risk factors in US children. Pediatrics. 2013; 132(3):e637–45. doi: 10.1542/peds.2013-0106 PMID: 23958765

16. Lang IA, Galloway TS, Scarlett A, Henley WE, Depledge M, Wallace RB, et al. Association of urinary bisphenol A concentration with medical disorders and laboratory abnormalities in adults. JAMA. 2008; 300(11):1303–10. doi: 10.1001/jama.300.11.1303 PMID: 18799442

17. Xu XH, Hong X, Xie LD, Li T, Yang YJ, Zhang Q, et al. Gestational and lactational exposure to bisphenol-A affects anxiety- and depression-like behaviors in mice. Horm Behav. 2012; 62(4):480–90. PMID: 23240141

18. Tian YH, Baek JH, Lee SY, Jiang CG. Prenatal and Postnatal Exposure to Bisphenol A Induces Anxiolytic Behaviors and Cognitive Deficits in Mice. Synapse. 2010; 64(6):432–9. doi: 10.1002/syn.20746 PMID: 20169576

19. Braun JM, Kalkbrenner AE, Calafat AM, Yolton K, Ye X, Dietrich KN, et al. Impact of early-life bisphenol A exposure on behavior and executive function in children. Pediatrics. 2011; 128(5):873–82. doi: 10.1542/peds.2011-1335 PMID: 22025598

20. Hatch EE, Nelson JW, Stahlhut RW, Webster TF. Association of endocrine disruptors and obesity: perspectives from epidemiological studies. Int J Androl. 2010; 33(2):324–32. doi: 10.1111/j.1365-2605.2009.01035.x PMID: 20113374

21. Du Y, Gao Y, Meng F, Liu S, Fan Z, Wu J, et al. Iodine deficiency and excess coexist in china and induce thyroid dysfunction and disease: a cross-sectional study. PLoS One. 2014; 9(11):e111937. doi: 10.1371/journal.pone.0111937 PMID: 25375854

22. Meng F, Zhao R, Liu P, Liu L, Liu S. Assessment of iodine status in children, adults, pregnant women and lactating women in iodine-replete areas of China. PLoS One. 2013; 8(11):e81294. doi: 10.1371/journal.pone.0081294 PMID: 24282581

23. Pearce EN, Andersson M, Zimmermann MB. Global iodine nutrition: Where do we stand in 2013? Thyroid. 2013; 23(5):523–8. doi: 10.1089/thy.2013.0128 PMID: 23472655

24. World Health Organization/UNICEF/International Council for Control of Iodine Deficiency Disorders (2008) Elimination of iodine deficiency disorders: a manual for health workers. WHO Regional Office for Eastern Mediterranean.
25. Zou Y, Lou X, Ding G, Mo Z, Zhu W, Mao G, et al. An assessment of iodine nutritional status and thyroid hormone levels in children aged 8–10 years living in Zhejiang Province, China: a cross-sectional study. Eur J Pediatr. 2014; 173(7):929–34. doi:10.1007/s00431-014-2273-y PMID: 24500398

26. Hui L, Xin-nan Z, Cheng-ye J, Jie M. Body mass index cut-offs for overweight and obesity in Chinese children and adolescents aged 2–18 years. Chin J Epidemiol. 2010; 31(6):616–20.

27. Sang Z, Wang PP, Yao Z, Shen J, Halfyard B, Tan L, et al. Exploration of the safe upper level of iodine intake in euthyroid Chinese adults: a randomized double-blind trial. Am J Clin Nutr. 2012; 95(2):367–73. doi:10.3945/ajcn.111.028001 PMID: 22205314

28. Fuse Y, Saito N, Tsuchiya T, Shishiba Y, Irie M. Smaller thyroid gland volume with high urinary iodine excretion in Japanese schoolchildren: normative reference values in an iodine-sufficient area and comparison with the WHO/ICCIDD reference. Thyroid. 2007; 17(2):145–55. PMID: 17316117

29. Ministry of Health (2007) Diagnostic criterion for endemic goiter. MOH B, in Chinese.

30. Cooper DS, Doherty GM, Haugen BR, Kloos RT, Lee SL, Mandel SJ, et al. Revised American Thyroid Association management guidelines for patients with thyroid nodules and differentiated thyroid cancer. Thyroid. 2009; 19(11):1167–214. doi:10.1089/thy.2009.0110 PMID: 19860577

31. Tong YK HC, Li BQ, Li MH, Zhang NJ, Min SH, Fu SY General test method in salt industry: determination of iodide ion (GB/T 13025.7–1999). Beijing, China: China Criteria Publishing House; 1999. p. 1–5.

32. Wang HX, Zhou Y, Jiang QW. Rapid and sensitive analysis of phthalate metabolites, bisphenol A, and endogenous steroid hormones in human urine by mixed-mode solid-phase extraction, dansylation, and ultra-performance liquid chromatography coupled with triple quadrupole mass spectrometry. Anal Bioanal Chem. 2013; 405(12):4313–9. doi:10.1007/s00216-013-6779-3 PMID: 23430180

33. Braun JM, Smith KW, Williams PL, Calafat AM, Berry K, Ehrlich S, et al. Variability of urinary phthalate metabolite and bisphenol A concentrations before and during pregnancy. Environ Health Perspect. 2010; 120(5):739–45. doi:10.1289/ehp.1104139 PMID: 22262702

34. Jooste PL, Strydom E. Methods for determination of iodine in urine and salt. Best Pract Res Clin Endocrinol Metab. 2010; 24(1):77–88. doi:10.1016/j.beem.2009.08.006 PMID: 20172472

35. Vandenbroucke RN, Chahoud I, Heindel JJ, Padmanabhan V, Paumgartten FJ, Schoenfelder G, Urinary, circulating, and tissue biomonitoring studies indicate widespread exposure to bisphenol A. Environ Health Perspect. 2010; 118(8):1055–70. doi:10.1289/ehp.0901716 PMID: 20338858

36. Chevrier J, Gunier RB, Bradman A, Holland NT, Calafat AM, Eskenazi B, et al. Maternal urinary bisphenol A during pregnancy and maternal and neonatal thyroid function in the CHAMACOS study. Environ Health Perspect. 2013; 121(1):138–44. doi:10.1289/ehp.1205092 PMID: 23052180

37. Gentilcori D, Porreca I, Rizzo F, Ganbaatar E, Carchia E, Mallardo M, et al. Bisphenol A interferes with thyroid specific gene expression. Toxicology. 2013; 304:21–31. doi:10.1016/j.tox.2012.12.001 PMID: 23238275

38. Heimeier RA, Das B, Buchholz DR, Shi YB. The xenoestrogen bisphenol A inhibits postembryonic vertebrate development by antagonizing gene regulation by thyroid hormone. Endocrinology. 2009; 150 (6):2964–73. doi:10.1210/en.2008-1503 PMID: 19228888

39. Sheng ZG, Tang Y, Liu YX, Yuan Y, Zhao BQ, Chao XJ, et al. Low concentrations of bisphenol a suppress thyroid hormone receptor transcription through a nongenomic mechanism. Toxicol Appl Pharmacol. 2012; 259(1):133–42. doi:10.1016/j.taap.2011.12.018 PMID: 22227104

40. Wang HX, Zhou Y, Tang CX, Wu JG, Chen Y, Jiang QW. Association between bisphenol A exposure and body mass index in Chinese school children: a cross-sectional study. Environ Health. 2012; 11:79. PMID: 23083070. doi:10.1186/1476-069X-11-79

41. Hatch EE, Nelson JW, Stahlhut RW, Webster TF. Association of endocrine disruptors and obesity: perspectives from epidemiological studies. Int J Androl. 2010; 33(2):324–31. doi:10.1111/j.1365-2605.2009.01035.x PMID: 20113374

42. Hayashida N, Imazumi M, Shimura H, Okubo N, Asari Y, Nigawara T, et al. Thyroid ultrasound findings in children from three Japanese prefectures: aomori, yamanashi and nagasaki. PLoS One. 2013; 8(12):e83220. doi:10.1371/journal.pone.0083220 PMID: 24376666

43. Liu Y, Huang H, Zeng J, Sun C. Thyroid volume, goiter prevalence, and selenium levels in an iodine-sufficient area: a cross-sectional study. BMC Public Health. 2013; 13:1153. PMID: 24321191. doi: 10.1186/1471-2458-13-1153