



The use of breast milk iodine concentration in the first week of lactation as a biomarker of iodine status in breast-feeding women

Shuchang Liu^{1,2}, Andrew Sharp^{2*}, Xiaoqin Luo³, Steven Lane⁴, Elmer V Villanueva^{5,6}, Zhiliang Lu^{1*} and Zheng Feei Ma^{5,7*}

¹Department of Biological Sciences, Xi'an Jiaotong-Liverpool University, Suzhou 215123, People's Republic of China

²Harris-Wellbeing Research Centre, Faculty of Health & Life Sciences, University of Liverpool, Liverpool L8 7SS, UK

³School of Public Health, Xi'an Jiaotong University, Xi'an 710006, People's Republic of China

⁴Health Data Science, University of Liverpool, Liverpool, UK

⁵School of Science, Xi'an Jiaotong-Liverpool University, Suzhou 215123, People's Republic of China

⁶Victoria University, Footscray Victoria Melbourne 3011, Australia

⁷Centre for Public Health and Wellbeing, School of Health and Social Wellbeing, College of Health, University of the West of England, Bristol BS16 1QY, UK

(Submitted 19 June 2023 – Final revision received 7 August 2023 – Accepted 16 August 2023 – First published online 29 August 2023)

Abstract

Breast milk iodine concentration (BMIC) is a promising indicator of iodine status in lactating women. However, there are limited data on its usefulness to reflect maternal iodine deficiency. Therefore, the aim of our study was to assess iodine concentration in breast milk and urine samples in exclusively breast-feeding women. Eligible pregnant women undergoing routine antenatal care in a large hospital in Shaanxi Province, China, were followed up from the third trimester of pregnancy until the first week of lactation. Urine samples (20 ml) were collected during pregnancy and lactation. Iodine concentration in samples was measured based on Sandell–Kolthoff reaction. Breast milk samples (5 ml) were provided during lactation. A receiver operating curve (ROC) was constructed to determine the diagnostic performance of BMIC. An iodine-specific FFQ was completed twice during pregnancy and lactation. A total of 200 women completed the study. The overall median BMIC was 89 µg/l, indicating iodine sufficiency (i.e. BMIC reference range between 60 and 465 µg/l). Women reported similar median urinary iodine concentration (UIC) during pregnancy and lactation (112 and 113 µg/l, respectively), but their iodine status differed – mild-to-moderate iodine deficiency during pregnancy and iodine sufficiency during lactation. The ROC for BMIC using UIC as a reference standard was 0.755 (95% CI: 0.644, 0.866). In conclusion, this study demonstrated that women were iodine sufficient in the first week of lactation as assessed by UIC, which was consistent with BMIC. These findings suggested that BMIC is a useful biomarker to assess iodine status in lactating women.

Keywords: Breast milk iodine concentration: Urinary iodine concentration: Maternal iodine status: Pregnancy: Lactation

Iodine is an essential micronutrient required for the production of thyroid hormones, which regulate metabolism, growth and development⁽¹⁾. Pregnant and lactating women have higher iodine requirements because they need to support the iodine needs of their fetus and baby⁽²⁾. A recent meta-analysis reported that the overall worldwide prevalence of maternal iodine deficiency was 53%⁽³⁾. As a result, insufficient iodine intake can cause many iodine deficiency disorders, which affect both mothers and infants, e.g. congenital hypothyroidism, intellectual disability, neonatal hypothyroidism, hyperthyrotropenemia and growth retardation⁽⁴⁾.

The WHO recommends assessing iodine status based on urinary iodine concentration (UIC) for non-pregnant women, defined as moderate-to-severe iodine deficiency (UIC 0–49 µg/l), mild iodine deficiency (UIC 50–99 µg/l), iodine sufficiency (UIC 100–199 µg/l) and more-than-adequate and excessive iodine status (UIC ≥ 200 µg/l)⁽⁵⁾. In non-pregnant women, UIC is used to determine iodine status based on the principle that approximately 90% of ingested iodine is excreted in the urine⁽⁶⁾. However, using UIC to assess iodine status has some limitations. First, UIC is a short-term biomarker, which can be easily affected by recent iodine intake from the diet⁽⁷⁾. Second, there is high

Abbreviations: BMIC, breast milk iodine concentration; UIC, urinary iodine concentration.

* **Corresponding authors:** Zheng Feei Ma, email zheng.ma@uwe.ac.uk; Andrew Sharp, email a.sharp@liverpool.ac.uk; Zhiliang Lu, email zhiliang.lu@xjtu.edu.cn



intra- and inter-individual variation in UIC^(7,8). In lactating women, there is little evidence that supports the recommendation of a median UIC cut-off of < 100 µg/l to indicate iodine deficiency because iodine is also excreted in breast milk and thus < 90 % of ingested iodine is excreted via urine^(4,9).

Breast milk iodine concentration (BMIC) has been reported to be a promising biomarker of iodine status in breast-feeding women⁽¹⁰⁾, although there is no scientific consensus on whether BMIC can accurately reflect iodine status in breast-feeding women⁽⁴⁾. This is because the iodine content in breast milk is influenced by the mother's iodine intake and overall iodine status⁽⁴⁾. In addition, BMIC is independent of maternal fluid intake⁽⁴⁾. A recent systematic literature review has suggested that there have been inconsistencies in the relationship between BMIC and UIC in lactating women, which may be due to the differences in lactation stages, maternal iodine status and sampling collection time⁽¹¹⁾. Most of these studies were cross-sectional in design, and there has been a lack of high-quality, well-designed studies⁽¹¹⁾. Therefore, WHO suggested a number of research priorities including the assessment of iodine status during pregnancy and early infancy, which include the usefulness of BMIC⁽⁹⁾.

To date, there is limited information regarding BMIC in breast-feeding women and on the changes of iodine status in pregnant women who are then followed until lactation. China has eliminated iodine deficiency disorders for more than 20 years and considered an iodine sufficient country based on the median UIC of studies involving school-aged children and non-pregnant adults⁽¹²⁾. However, these findings may not reflect the iodine status of women during pregnancy and lactation, who, as stated earlier, have a substantially higher iodine requirement⁽¹³⁾. Some women during pregnancy and lactation remain at risk of iodine deficiency⁽¹⁴⁾. Therefore, the aim of our study was to assess iodine concentration in breast milk and urine samples in Chinese breast-feeding women from Shaanxi province, an iodine sufficient region.

Methods

Study population

The Women and Iodine Nutrition study was designed as a prospective, longitudinal, observational cohort study spanning from the third trimester of pregnancy to the first week of lactation, with a follow-up period of up to 3 months, in Shaanxi Province (in the western part of China). Pregnant women were recruited between May 2021 and May 2022 at Xianyang Central Hospital Affiliated with the Medical Department of Xi'an Jiaotong University.

Inclusion criteria were as follows: pregnant women during their third trimester (gestation weeks of 28 and above); aged between 18 and 50 years; healthy, not had medically diagnosed thyroid disease, not taking thyroid medication; singleton birth; intended to breastfeed for at least 7 days; of Chinese nationality; must live in Shaanxi at least one year; able to read and write in Chinese; had a healthy, singleton, full-term birth (pregnancy weeks 38–42); infant exclusively breastfed. Infants with fetal abnormality were excluded.

Written informed consent was obtained from all eligible women. Infants were consented by their participating mothers. Ethical approval for this study was approved by the Xi'an Jiaotong-Liverpool University Ethics Committee (reference no. 20-01-09) and Xianyang Central Hospital Affiliated to the Medical Department of Xi'an Jiaotong University (reference no. 20200009). The results of the Women and Iodine Nutrition study were reported according to the STrengthening the Reporting of OBservational studies in Epidemiology (STROBE) guidelines for the cohort studies^(15,16).

Socio-demographic data collection and other maternal and neonatal data

Women were required to complete a fifty-seven-item questionnaire including a validated thirty-three-item iodine-specific FFQ (including milk products, yoghurt, eggs, seafoods and fish) and questions on socio-demographics twice (during pregnancy and lactation). The questionnaire has been previously validated and used in assessing maternal iodine status of the Chinese population⁽¹⁷⁾. Other socio-demographics of participants including age, marital status, ethnicity, education level, occupation and pre-pregnancy BMI were obtained during routine antenatal care. Women who graduated high school, junior high school and elementary school and below were categorised as 'below university level' and those who graduated university or junior college and above were 'university level and above'. The daily iodine doses during pregnancy and lactation were retrieved using the brand name of the supplement provided via the questionnaire used. Pre-pregnancy height and weight were recorded using a stadiometer to the closest 0.1 cm and 0.01 kg. Pre-pregnancy BMI was categorised according to the criteria by the Chinese adults proposed by the WHO's recommendation as follows: underweight, < 18.5 kg/m²; normal weight, 18.5–24.9 kg/m²; overweight, ≥ 25.0 kg/m² and obese, ≥ 30.0 kg/m²⁽¹⁸⁾.

Data on blood pressure, thyroid-stimulating hormone, free thyroxine (FT4) and thyroid peroxidase antibody (TPOAb) during pregnancy were retrieved from medical records. The normal reference ranges for thyroid-stimulating hormone, FT4 and TPOAb were as follows: 0.27–4.20 µIU/ml, 0.93–1.70 ng/dl and < 34.0 µg/ml. Thyroid-stimulating hormone, FT4 and TPOAb values were determined using electrochemiluminescence immunoassay on a Roche E602 immunochemistry analyser⁽¹⁹⁾. In addition, total gestation weeks at delivery, sex, type of delivery, birth weight and length and APGAR score (appearance, pulse, grimace, activity and respiration) at 1, 5 and 10 min were also assessed and obtained from neonatal records.

Urine sample collection

During the third trimester of pregnancy, participants were provided with instructions and equipment to collect one approximately 20 ml (non-fasting), mid-stream urine sample between 09.00 and 12.00. During the first week of lactation, participants were asked to provide two (non-fasting), mid-stream urine samples between 09.00 and 12.00 on two separate days (i.e. the 3rd and 4th day of lactation period), to control for the intra-individual and within-day UIC variation, while minimising participant burden⁽²⁰⁾. The internationally



recommended method to determine iodine status based on UIC is the collection of morning or spot urine samples in the non-fasting state⁽⁵⁾.

Breast milk sample collection

Women were asked to clean their breasts with water before collecting the breast milk sample manually. Approximately one 5 ml breast milk sample (non-fasting sample) was collected between 09.00 and 12.00 from women before their infants were fed at the 3rd day of lactation because not all women's milk would come in on the 1st or 2nd day of lactation^(20,21). Since the literature review does not show whether BMIC is affected by a recent meal, women were allowed to have their usual diet before breast milk collection⁽²²⁾. Currently, there is no evidence supporting the fact that BMIC differs with regard to the milk from the start and end of a feed, left or right breast or diurnal variation^(23,24).

Laboratory analysis

All urine and breast milk aliquots were kept frozen at -20°C from time of sampling in Xianyang Central Hospital Affiliated to the Medical Department of Xi'an Jiaotong University until analysis.

After thawing, samples were vortexed using a vortex mixer until homogenous. Samples from the same woman were analysed in the same batch.

Assessment of breast milk iodine concentration and urinary iodine concentration in lactating women. BMIC and UIC were measured colourimetrically based on the Sandell–Kolthoff reaction adapted for a 96-well microplate using a microplate reader (Denley Dragon, Wellscan MK 3, Thermo Fisher Scientific)⁽²⁵⁾. The equipment was calibrated following the manufacturer's instructions and quality control samples with known concentrations of iodine were included in the same run of the iodine analysis for both BMIC and UIC. A BMIC reference range of 60–465 $\mu\text{g/l}$ is used as indicative of sufficient iodine status in exclusively breast-feeding women residing in iodine sufficient regions⁽¹⁰⁾. The recommended UIC cut-offs during pregnancy and lactation to determine iodine sufficiency are ≥ 150 and ≥ 100 $\mu\text{g/l}$, respectively⁽⁵⁾.

Statistical analysis

SPSS statistical software package version 25.0 (IBM Corp.) was used to perform the statistical analysis. Parametric data were expressed as mean and standard deviation, and non-parametric data were presented as median (25th, 75th percentile). Categorical variables were reported as counts and percentages, χ^2 tests were used to assess differences in categorical variables. For related samples, comparisons of medians or means were performed using Wilcoxon sign rank tests for non-normally or paired t-test was used for normally distributed variables.

Receiver operating curves were constructed to determine the diagnostic performance of BMIC using a UIC cut-off of 100 $\mu\text{g/l}$ for lactating women. An optimal cut-off for BMIC was identified and sensitivity (proportion of cases correctly identified), specificity (proportion of non-cases correctly identified) along

with negative predictive value and positive predictive value were then calculated to assess the accuracy of the cut-off^(26,27). Sensitivity is defined as the proportion of those who are correctly identified as iodine deficient by BMIC (true positives), while specificity is defined as the proportion of those who are correctly identified as not iodine deficient by BMIC (true negatives)^(26,27). The negative predictive value is expressed as the proportion of those with negative test results who are correctly identified as not iodine deficient. The positive predictive value is reported as the proportion of those with positive test results who are correctly identified as iodine deficient^(26,28). As there is no gold standard to assess individual iodine status for lactating women, the mean of two spot urine samples collected from each lactating women was used as the reference standard for the determination of the sensitivity, specificity, negative predictive value and positive predictive value of BMIC. When the area under the receiver operating curve is ≥ 0.7 , it is considered to have the acceptable discrimination for distinguishing iodine deficiency from iodine sufficient⁽²⁹⁾.

A correlation between BMIC and maternal UIC was assessed using a Spearman correlation coefficient. Logistic regression analysis was used to assess the associations between the predictors of BMIC and maternal UIC with adjustment for covariates. The dependent variables were BMIC and maternal UIC. Covariates adjusted in regression models included variables of age, UIC pregnancy, UIC lactation, delivery type, occupation and education. A $P < 0.05$ was taken as level of significance.

For the calculation of 24-h breast milk iodine excretion, a breast milk volume of 0.8 l/d was used^(1,30). The estimated infant's iodine intake is calculated as follows: total volume of breast milk consumed by the infant multiplied by BMIC. Currently, there is no recommended iodine intake for infants aged < 1 month old⁽³¹⁾.

The primary outcome of this study was BMIC, and the sample size was calculated using G*Power 3.1 (Heinrich Heine University) based on data (mean and standard deviation) from a study of Chinese lactating women⁽³²⁾. Therefore, on the basis of the literature and using BMIC as the primary outcome, in order to detect an effect size of 0.5, with 90% power and two-sided alpha (0.05), this meant at least eighty women would be needed in each group to detect a significant difference between the iodine deficient and iodine sufficient groups. After accounting for 20% attrition, a final sample size of 192 women was needed for the whole study.

Results

Study population

Of the 227 women who were approached and invited to participate, 200 pregnant women fulfilled the study criteria and were enrolled in the study. Those who were excluded from the study were either ineligible for the study criteria ($n = 3$) or not interested in the study ($n = 24$). Table 1 summarises the basic characteristics of women and their infants included in the study. The mean age of the women was 29.0 ± 4.2 years. The study population consisted of pregnant women with gestational ages ranging from 29 weeks and 6 days to 40 weeks and 3 days, with a



Table 1. Socio-demographic characteristics of women and their infants

Characteristics of participants	Value*	
	Mean	SD
Sample size, <i>n</i>	200	
Age (years)	29.0	4.2
Gestational age (week)	37.0	2.4
Delivery age (week)	39.9	0.8
Height (cm)	161.0	4.8
Pre-pregnancy weight (kg)	55.9	8.4
Pre-pregnancy BMI (kg/m ²)	21.5	2.8
Weight gain	14.9	3.7
Blood pressure (mm Hg)		
Systolic pressure	115.1	10.5
Diastolic pressure	74.9	7.3
	<i>n</i>	%
Education level		
Elementary school and below	1	0.5
Junior high school	24	12.0
High school	44	22.0
University or junior college	126	63.0
Graduate and above	5	2.5
Occupation		
Local enterprise employees	67	33.5
Foreign-funded enterprises	0	0.0
Government professional units	7	3.5
Private SME owners and individual business owners	21	10.5
Agriculture	7	3.5
Freelance	62	31.0
Other	36	18.0
Delivery type		
Caesarean section	92	46.0
Vaginal birth	108	54.0
Smoking status		
Smoker	3	1.5
Non-smoker	197	98.5
Infants		
Sex		
Male	98	49.0
Female	102	51.0
Birth weight (kg)		
Mean	3.3	
SD	0.4	
Birth length (cm)		
Mean	51.1	
SD	1.4	
	Median	IQR
APGAR		
1 min	10.0	10.0, 10.0
5 min	10.0	10.0, 10.0
10 min	10.0	10.0, 10.0

SME, small and medium-size enterprise; IQR, interquartile range.

* Data are means \pm SD, median (IQR) or *n* (%).

mean gestational age of 37 weeks. All women were negative for TPOAb. The mean pre-pregnancy weight of the participants was 55.9 ± 8.4 kg, who had a mean weight gain of 14.9 ± 3.7 kg during pregnancy. Their mean systolic blood pressure was 115.1 ± 10.5 mm Hg, and their mean diastolic blood pressure was 74.9 ± 7.3 mm Hg. Only 1.5% of participants (*n* 3) were smokers. The study population consisted of similar equal numbers of male and female infants, with ninety-eight males (49.0%) and 102 females (51.0%). The mean birth weight was 3.3 ± 0.4 kg, and the mean birth length was 51.1 ± 1.4 cm. The median APGAR scores at 1 min, 5 min and 10 min were 10.0, 10.0 and 10.0, respectively. These scores indicate that the infants were

generally healthy at birth. Sixty-six participants (33.0%) reported that they use supplements (including vitamin complex, vitamin D, vitamin C, DHA, Ca, Runkang brand pregnancy supplements and Forceval brand pregnancy supplements), but only seven of them used supplements containing iodine. The daily iodine dose in these supplements (*n* 7) amounted to 150 μ g. No infants had a birth weight < 2500 g (i.e. low birth weight).

Iodine status

The iodine status of women is presented in Table 2. The overall median (interquartile range (IQR)) BMIC was 89 μ g/l (74, 117 μ g/l). The overall median UIC during pregnancy was 112 μ g/l (85, 134 μ g/l), which was indicative of iodine deficiency (median UIC < 150 μ g/l), while the overall median UIC during lactation was 113 μ g/l (90, 133 μ g/l), indicating iodine sufficiency (median UIC \geq 100 μ g/l). No significant (correlations) differences were found between UIC during pregnancy and lactation ($P = 0.784$). The prevalence of iodine deficiency (as assessed by UIC) was significantly during pregnancy than that of lactation (69.8% *v.* 30.2%) ($P < 0.001$). The overall mean dietary iodine intake calculated from FFQ for pregnancy and lactation was 231.89 ± 146.02 and 237.26 ± 156.20 μ g/d, respectively.

For those women who took iodine-containing supplements from pregnancy to lactation (*n* 7), all median UIC were below their respective median cut-off values: median UIC during pregnancy was 123 μ g/l, and median UIC during lactation was 99 μ g/l (median cut-off value 100 μ g/l). Besides, the median BMIC of women who took iodine-containing supplements (*n* 7) was 97 μ g/l.

Table 3 shows BMIC by socio-demographic features of women. There was no difference in BMIC in terms of different categories of age, BMI, delivery type, education, occupation and smoking status. We observed a positive significant correlation ($r = 0.369$, $P < 0.001$) between BMIC and UIC in breast-feeding women (Fig. 1).

Usefulness of breast milk iodine concentration in the assessment of iodine status. Figure 2 shows the area under the receiver operating curve for BMIC using UIC as a reference standard was 0.755 (95% CI: 0.644, 0.866), which was within the acceptable range (≥ 0.7)⁽²⁹⁾. This suggested a 75.5% chance that BMIC would correctly distinguish iodine-deficient breast-feeding women from iodine sufficient breast-feeding women. Therefore, BMIC could be used as a biomarker of iodine status in breast-feeding women.

Figure 3 indicates the optimal cut-off, in terms of optimising sensitivity and specificity of BMIC is 117 μ g/l, with a sensitivity of 0.645 and a specificity of 0.828. This meant that if the cut-off value was used to test for iodine deficiency, it would correctly identify 64.5% of participants who were iodine deficient and correctly identified 82.8% of participants who were not iodine deficient. A negative predictive value of 92.6% meant that 92.6% of participants who tested negative for iodine deficiency actually did not have the deficiency, while a positive predictive value of 39.2% meant that 39.2% of participants who tested positive for iodine deficiency actually had the deficiency. These values indicated that the BMIC cut-off value of 117 μ g/l was a good

Table 2. Iodine status of women (*n* 200)

	Values*		P-value
	<i>n</i>	%	
BMIC (µg/l)			
Median	89	–	
IQR	74, 117		
Prevalence of iodine deficiency according to BMIC cut-off < 60 µg/l, <i>n</i> (%)	23	11.5	–
Median UIC (µg/l)			
Pregnancy			
Median	112		0.784
IQR	85, 134		
Lactation			
Median	113		
IQR	90, 133		
Prevalence of iodine deficiency based on the UIC cut-off			
Pregnancy (< 150 µg/l)	169	69.8	<0.001
Lactation (< 100 µg/l)	73	30.2	
	Mean	SD	
Thyroid function during pregnancy			
TSH (µIU/ml)	2.11	0.39	–
FT4 (ng/dl)	1.30	0.08	–
TPOAb (µg/ml)	8.00	3.80	–
Dietary iodine intake (µg/d)			
Pregnancy	231.89	146.02	0.762
Lactation	237.26	156.20	
BMIC/Infant iodine intake (µg/d)			
Median	71.60		–
IQR	58.60, 93.40		

BMIC, breast milk iodine concentration; IQR, interquartile range; UIC, urinary iodine concentration; TSH, thyroid-stimulating hormone

* Data are means ± SD or *n* (%) or median and interquartile ranges (IQR). A *P* < 0.05 was taken as level of significance. *P* values are shown for comparison between pregnancy and lactation.

balance between accurately identifying iodine deficiency and avoiding false positives.

Table 4 shows the regression analysis for the predictors of BMIC in breast-feeding women. The only significant predictor of BMIC was UIC during pregnancy. This suggested that if a woman had a UIC pregnancy level greater than the UIC cut-off value, the woman was eight times more likely to have a BMIC above the optimal cut-off (i.e. 117 µg/l) when compared with a woman with a UIC pregnancy level below cut-off. However, the 95% confidence level suggested that this difference may be as low as 3.5 times or as high as 18.6 times.

Discussion

Our study was one of the first studies to use the BMIC reference range proposed by Dold *et al.*, which is for exclusively breast-feeding women residing in iodine sufficient regions⁽¹⁰⁾. In our study, a number of biomarkers of iodine status including BMIC and UIC were employed to provide a comprehensive assessment of iodine status for women during their first week of lactation in an iodine sufficient region of China. This is because during lactation, about 45% of maternal iodine is redirected to meet the infant's iodine requirement, resulting in a decrease in fractional iodine excretion in the maternal urine⁽³³⁾.

Table 3. BMIC by socio-demographic characteristics of women

	BMIC*		P-value
	Median	IQR	
Age (years)			
< 30	87	72, 116	0.278
≥ 30	95	77, 117	
BMI (kg/m ²)			
Underweight < 18.5	103	77, 127	0.535
Normal weight 18.5–24.9	91	74, 117	
Overweight and obese ≥ 25	82	69, 112	
Delivery type			
Caesarean section	89	74, 115	0.656
Vaginal birth	89	72, 119	
Education level			
Below university level	91	75, 116	0.960
University level and above	89	72, 119	
Occupation			
Employed	90	74, 117	0.391
Non-employed	83	68, 118	
Smoking status			
Smoker	88	74, †	0.574
Non-smoker	91	73, 117	

BMIC, breast milk iodine concentration.

* Data are median and interquartile ranges (IQR).

† Only three smokers.

Currently, there is no consensus on an acceptable BMIC cut-off to categorise iodine sufficiency during lactation, which is primarily due to the uncertainty about infant iodine requirements. Several BMIC cut-offs of 50, 75, 80, 92 and 100 µg/l have been proposed to ensure iodine sufficiency during lactation⁽¹¹⁾. However, these proposed BMIC cut-offs did not specify if they could be applied to breast-feeding women residing in iodine sufficient regions because some of these cut-offs may have been derived from iodine deficient breast-feeding women and therefore not suitable for iodine sufficient regions. For example, higher BMIC was reported in goitrous areas of Detroit than non-goitrous areas of Boston⁽³⁴⁾. Furthermore, no difference in BMIC was reported between goitrous and non-goitrous areas of Italy and New Zealand^(35,36). On the other hand, in a large multi-centre study of lactating women, Dold *et al.* proposed a broad reference range of 60–465 µg/l to suggest iodine sufficiency in exclusively breast-feeding women from iodine sufficient regions⁽¹⁰⁾.

The overall median BMIC in the first week of lactation was 89 µg/l, which was indicative of iodine sufficiency based on the BMIC reference of 60–465 µg/l suggested by Dold *et al.*⁽¹⁰⁾. In addition, our overall median BMIC in the first week of lactation was within the range of BMIC (from 43 to 138 µg/l) in the first week of lactation as reported in previous studies^(36–42). Of these previous studies (*n* 7), only two of them were conducted in iodine sufficient lactating women^(37,40). Our overall median BMIC was higher than that of Böhles *et al.*⁽³⁷⁾ (55 µg/l), but lower than that of Kart *et al.*⁽⁴⁰⁾ (138 µg/l). This highlighted that there have been very few studies that have investigated BMIC in the first week of lactation and data particularly from iodine sufficient regions are limited.

WHO recommends that breastfed infants receive adequate amounts of iodine in their diet to ensure a normal growth and development. Breast milk is the only dietary source of iodine for breastfed infants. BMIC is primarily influenced by the maternal

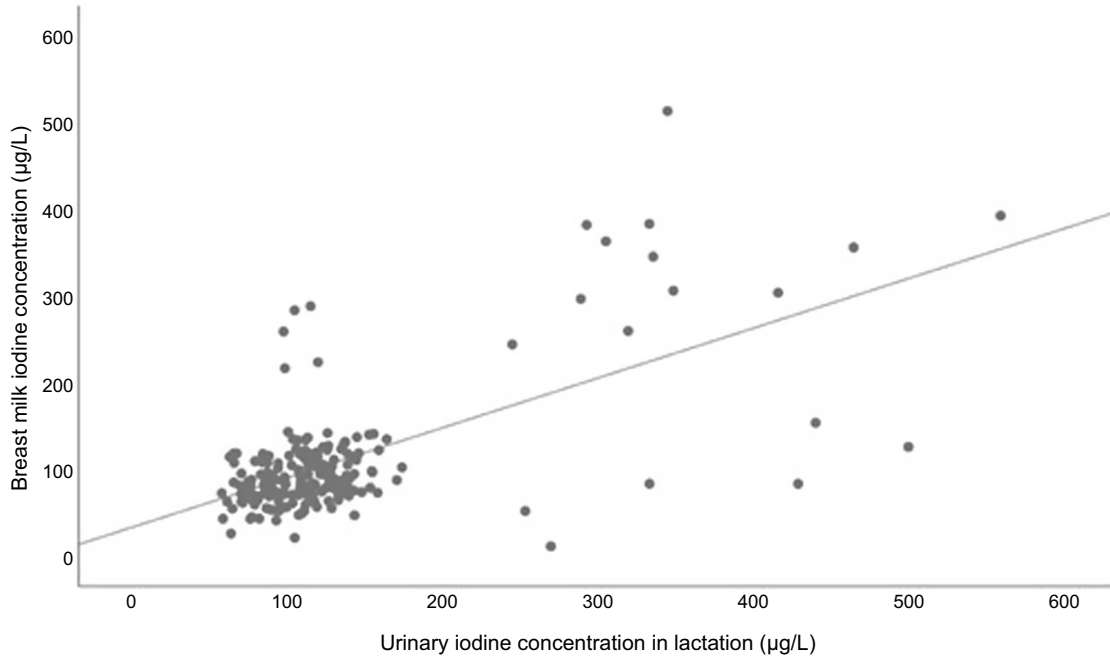


Fig. 1. Scatter plots of 200 samples illustrating the correlation between BMIC and UIC during lactation. BMIC, breast milk iodine concentration; UIC, urinary iodine concentration

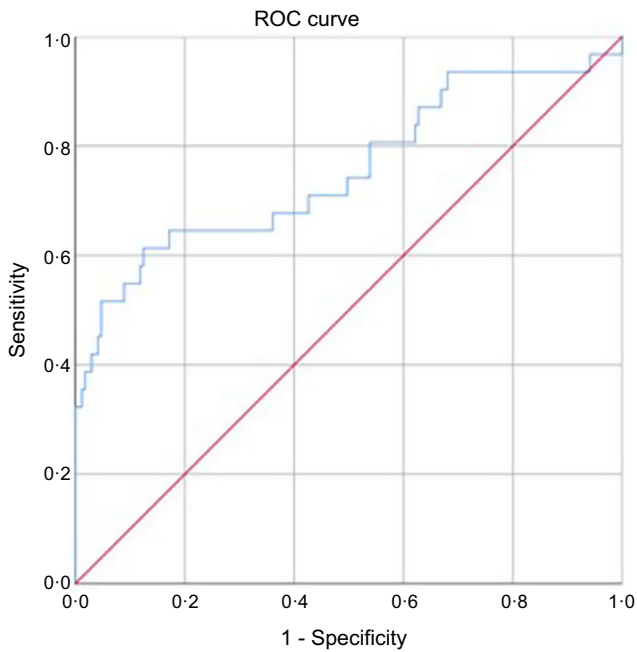


Fig. 2. The ROC curve for BMIC using UIC as a reference standard, diagonal line indicates chance (area = 0.5). ROC, receiver operating curve; BMIC, breast milk iodine concentration; UIC, urinary iodine concentration

dietary iodine intake⁽⁴³⁾. Women who had low iodine intake were reported to have a lower BMIC compared with women with sufficient iodine intake⁽¹¹⁾. Other factors such as stages of lactation and the geographical location of the women have been reported to affect BMIC. Women who lived in regions with naturally low soil iodine concentration were likely to have lower BMIC⁽⁴⁴⁾. Due to the collection of breast milk on the 3rd day postpartum, it is

possible that some women would still be producing colostrum, while others are likely to be producing transitional milk⁽⁴⁵⁾. In studies from the USA, Germany, Italy, China, New Zealand, Korea and Morocco, BMIC appeared to be highest in colostrum and decreasing throughout lactation, although not all studies have reported this pattern^(1,20,46–48). Therefore, it is assumed that BMIC will fall below the reference range for later mature milk with the prolongation of lactation stages.

This study reported that the area under the receiver operating curve for BMIC using UIC as a reference standard was 0.755, which was within the acceptable range. In addition, using the plot of sensitivity and specificity, our study reported an optimal BMIC cut-off of 117 µg/l might be used for categorising iodine sufficiency in lactating women residing in an iodine sufficient region. Although there have been many proposed BMIC cut-offs to determine iodine sufficiency in lactating women, the most commonly used BMIC cut-offs in the literature are 75 and 100 µg/l^(10,22,49–51). However, there is still no scientific consensus on the agreed BMIC cut-off. One of the possible reasons is because the iodine needs for infants are still inconclusive⁽¹⁾. In addition, the breastmilk samples from the published studies had different collection periods of lactation (varied from days/weeks to months), making the comparison of BMIC between studies with different breast milk collection periods of lactation difficult. More studies assessing BMIC along with UIC and thyroid function in mother–infant pairs are warranted to define a median BMIC cut-off for assessing iodine sufficiency in lactating women residing in iodine deficient and sufficient regions.

This study found that while women were iodine deficient (median UIC < 150 µg/l) during pregnancy, they were categorised as iodine sufficient (median UIC ≥ 100 µg/l) during lactation. Despite both the median UIC values of women during

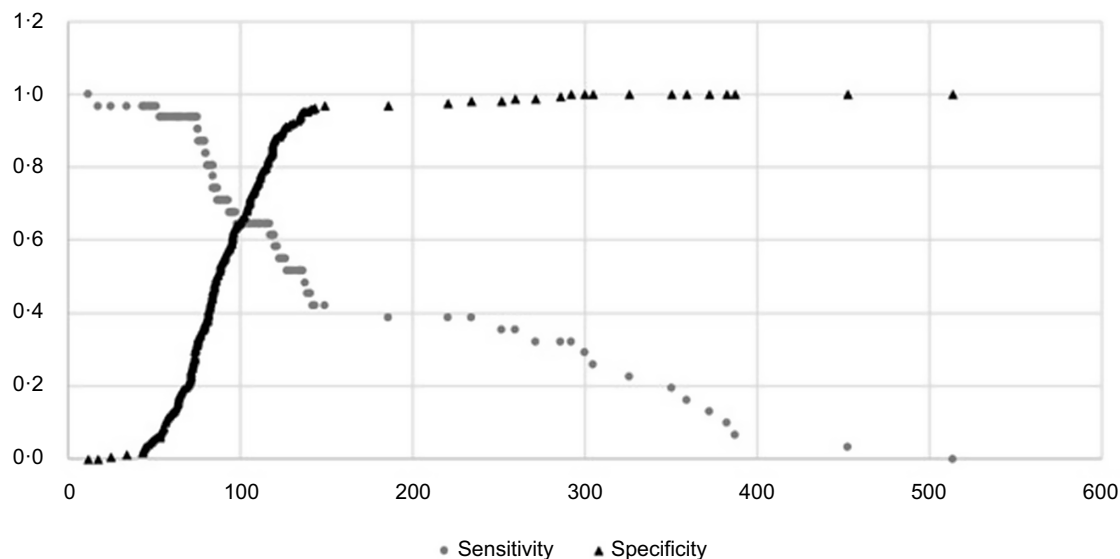


Fig. 3. Plots of sensitivity and specificity where lines cross are optimum value for classifying women as either being iodine sufficient or insufficient.

Table 4. Predictors of BMIC in women

	Unadjusted Coef.			Adjusted Coef.†		
	β	OR	95% CI	β	OR	95% CI
Age	0.008	1.008	0.530, 1.915	0.273	1.313	0.634, 2.721
UIC pregnancy	2.091	8.094	3.521, 18.608 *	2.090	8.086	3.414, 19.154*
UIC lactation	0.550	1.733	0.863, 3.479	0.316	1.371	0.642, 2.930
Delivery type‡	-0.263	0.796	0.404, 1.464	-0.311	0.732	0.356, 1.506
Occupation§	0.143	1.154	0.513, 2.595	0.248	1.281	0.522, 3.144
Education	0.070	1.072	0.547, 2.101	-0.139	0.870	0.413, 1.833
BMI¶	0.165	1.179	0.391, 3.550	-	-	-
Pregnant history-Term	0.198	1.218	0.642, 2.314	-	-	-
Pregnant history-Preterm	-0.027	0.973	0.099, 9.572	-	-	-
Pregnant history-Abortuses	-0.088	0.916	0.604, 1.388	-	-	-
Pregnant history-Living	0.142	1.153	0.607, 2.187	-	-	-
Use of iodised salt**	0.216	1.241	0.889, 1.733	-	-	-
Use of supplement in pregnancy††	0.095	1.100	0.565, 2.141	-	-	-
Use of supplement in lactation‡‡	0.183	1.201	0.612, 2.357	-	-	-

BMIC, breast milk iodine concentration; UIC, urinary iodine concentration.

* Significant ($P < 0.05$).

† Adjusted for variables of age, UIC pregnancy, UIC lactation, delivery type, occupation and education in the model. Dependent variable: BMIC (cut-off of 117 $\mu\text{g/l}$).

‡ Categories for delivery type: 0 = natural birth, 1 = cesarean birth.

§ Categories for occupation: 0 = employed, 1 = non-employed.

|| Categories for education: 0 = completed high school or lower education, 1 = higher education.

¶ Categories for BMI: 0 = normal weight, 1 = under/overweight.

** Categories for use of iodised salt: 0 = use, 1 = not use/unknown.

†† Categories for use of supplement in pregnancy: 0 = use, 1 = not use.

‡‡ Categories for use of supplement in lactation: 0 = use, 1 = not use

pregnancy and lactation were being similar (i.e. 112 $\mu\text{g/l}$ and 113 $\mu\text{g/l}$, respectively), the categorisation of iodine status for pregnant and lactating women was different in this study. This suggested a change in the iodine profile of women through pregnancy and into lactation. The median UIC cut-off to determine iodine sufficiency in lactating women is 100 $\mu\text{g/l}$, which is lower than pregnant women because in lactating women, ingested iodine is excreted both in urine and breast milk⁽⁵⁾. Therefore, due to the variation in the partition of iodine between breast milk and urine, both BMIC and UIC are recommended to be included when assessing iodine status in breast-feeding women⁽⁴⁾. On the other hand, the median UIC

cut-off value for classifying iodine sufficiency during pregnancy is 150 $\mu\text{g/l}$, which was established on the basis of an average daily urine volume of 1.5 l⁽⁵⁾. However, during pregnancy, there is an increase in the glomerular filtration rate, which leads to increased daily urine volume and subsequently lowers UIC, and this may overestimate the prevalence of iodine deficiency in pregnant women^(5,9).

Although the median UIC remained similar throughout pregnancy and lactation, the median BMIC of breast-feeding women was within the BMIC reference range proposed by Dold *et al.*, which indicated iodine sufficiency⁽¹⁰⁾. This could be due to various factors such as a change in maternal iodine metabolism

and increased iodine absorption characterised by increased thyroid stimulation, which may have contributed to an increase in iodine availability for secretion into breastmilk⁽⁵²⁾. Another possible reason may be that the mammary gland has the ability to selectively accumulate and concentrate iodine, independent of iodine intake⁽⁵⁰⁾. Therefore, it can actively transport iodine from the maternal bloodstream into breastmilk, resulting in higher BMIC values, even if the UIC value remains constant throughout these critical stages. However, the increased iodine uptake of the mammary gland may take place at the expense of maternal iodine reserves if there is insufficient maternal iodine intake from the women's diet⁽⁵³⁾.

There are several challenges involved during the collection of breast milk samples from women in the early stages of lactation⁽⁵⁴⁾. New mothers, particularly first-time mothers, are unlikely to interrupt breast-feeding at the early stage of lactation because this is a critical period for the establishment of exclusive breast-feeding^(54,55). Our study had 57.5% of first-time mothers. Furthermore, breast milk is considered precious and beneficial for infants in Chinese culture. For these reasons, it can be difficult to recruit women to take part in this type of research. Robust data on iodine status, especially BMIC, are scarce⁽¹¹⁾. One of the challenges of this study was collecting breast milk samples from breast-feeding women due to limited volumes of breast milk produced. During the first week of lactation, only small volumes of breast milk are produced (i.e. mean volume on the first 24 h after birth and third day of lactation is 37.1 (range 7.0–122.5) g and 408 (range 98.3–775) g)⁽⁵⁶⁾. Breast-feeding women encountered difficulties expressing enough milk. Our data should be interpreted with caution because spot samples of breast milk and urine were used to assess iodine status in our study. Future studies should consider collecting multiple breast milk and urine samples over 24 h and wider time range to calculate the daily iodine excretion from lactating women with different iodine intake levels. This will provide a more comprehensive assessment of iodine status in lactating women.

This study has several strengths. First, two biomarkers of iodine status, BMIC and UIC were used in assessing iodine status of breast-feeding women. The UIC values during lactation were derived from spot urine samples collected from two consecutive days. Although it is suggested that at least ten spot urine samples are needed to reliably assess individual iodine status, it was not feasible to obtain in this study. It is suggested that two spot urine samples provide a better estimate of iodine status compared with a single spot urine sample⁽⁵⁷⁾. The participants in this study consisted of a rather homogenous group of women with similar age and BMI ranges. Second, breast milk samples were collected during the first week of lactation, which few studies have performed. Third, the urine and breast milk samples were collected during the same time period, which can reflect better comparability of the iodine status. In addition, to our knowledge, this was the first study to evaluate BMIC of breast-feeding women in China using the reference range of 60–465 µg/l proposed by Dold et al.⁽¹⁰⁾. The BMIC reported in our study was at the lower end of the reference range, which was likely influenced by the UIC of breast-feeding women, which was just above the UIC cut-off indicating iodine sufficiency. However, it is important to note that the reference range proposed by Dold

et al. has a broad range and might not be applicable to breast-feeding women living in iodine deficient areas⁽¹⁰⁾. Also, the reference proposed by Dold et al. does not identify the BMIC value that corresponds to severe, moderate, mild iodine deficiency, optimal, more-than-adequate and excessive iodine status in breast-feeding women in the early, mid and later stages of lactation⁽¹⁰⁾.

Despite its considerable strengths, this study was limited by the failure to follow-up for a longer period of time to obtain subsequent samples. This is because the pandemic of coronavirus disease 2019 (COVID-19) affected some aspects of this study, including recruitment of participants because of the city lockdown, travel restrictions and hospital guidelines to prevent the spread of the COVID-19 pandemic. In addition, women usually left the hospital and return home on the 4th day of postpartum, making continued follow-up and sample collection difficult.

Conclusions

This study demonstrated that women were iodine sufficient in the first week of lactation. In addition, these findings supported the proposed BMIC reference range of 60–465 µg/l for a group of exclusively breast-feeding women in our region. Given the lack of conclusive evidence, more studies on the usefulness of BMIC as a biomarker of iodine status are warranted in breast-feeding women and infants with varying iodine status and lactation stages. Only then can reasonable recommendations be made regarding the usefulness of BMIC to assess their iodine status.

Acknowledgements

We were extremely grateful to all the participants who took part in the cohort study. We would also like to thank Sheila Skeaff for providing the intellectual input to the study.

This work was supported by the Research Development Fund (RDF) (reference no. RDF-18-01-15) from Xi'an Jiaotong-Liverpool University.

S. L. (Shuchang Liu), A. S, X. L., E. V. and Z. F. M. were responsible for the study design. S. L. (Shuchang Liu) was responsible for research tool development, participant recruitment, field investigation, data collection, statistical analysis and wrote the first draft of the manuscript. S. L. (Steven Lane) assisted with the statistical analysis. A. S, X. L., E. V., Z. L. and Z. F. M. provided oversight and leadership responsibility for the research activity planning and execution, including the mentorship for PhD candidate S. L. (Shuchang Liu). All authors read and approved the final manuscript. All authors reviewed the manuscript.

The authors declare that they have no conflict of interest.

All procedures performed in the retrospective study involving human participants were in accordance with the ethical standards of Xi'an Jiaotong-Liverpool University Ethics Committee (reference no. 20-01-09) and Xianyang Central Hospital Affiliated to the Medical Department of Xi'an Jiaotong University (reference no. 20200009). The procedures were complied with the 1964 Helsinki declaration and its later amendments or comparable ethical standards.



References

- Andersson M & Braegger CP (2022) The role of iodine for thyroid function in lactating women and infants. *Endocr Rev* **43**, 469–506.
- Bath SC, Verkaik-Kloosterman J, Sabatier M, *et al.* (2022) A systematic review of iodine intake in children, adults, and pregnant women in Europe—comparison against dietary recommendations and evaluation of dietary iodine sources. *Nutr Rev* **80**, 2154–2177.
- Patriota ES, Lima IC, Nilson EA, *et al.* (2022) Prevalence of insufficient iodine intake in pregnancy worldwide: a systematic review and meta-analysis. *Eur J Clin Nutr* **76**, 703–715.
- Brough L (2022) Improving iodine status in lactating women: what works? *Curr Nutr Rep* **11**, 592–599.
- World Health Organization (WHO) (2007) Assessment of Iodine Deficiency Disorders and Monitoring their Elimination: A Guide for Programme Managers. no. 9241595825. Geneva: WHO.
- Zimmermann MB & Andersson M (2012) Assessment of iodine nutrition in populations: past, present, and future. *Nutr Rev* **70**, 553–570.
- Zimmermann MB (2008) Methods to assess iron and iodine status. *Br J Nutr* **99**, S2–S9.
- Als C, Helbling A, Peter K, *et al.* (2000) Urinary iodine concentration follows a circadian rhythm: a study with 3023 spot urine samples in adults and children. *J Clin Endocrinol Metab* **85**, 1367–1369.
- Andersson M, De Benoist B, Delange F, *et al.* (2007) Prevention and control of iodine deficiency in pregnant and lactating women and in children less than 2-years-old: conclusions and recommendations of the Technical Consultation. *Public Health Nutr* **10**, 1606–1611.
- Dold S, Zimmermann MB, Aboussad A, *et al.* (2017) Breast milk iodine concentration is a more accurate biomarker of iodine status than urinary iodine concentration in exclusively breastfeeding women. *J Nutr* **147**, 528–537.
- Liu S, Sharp A, Villanueva E, *et al.* (2022) Breast milk iodine concentration (BMIC) as a biomarker of iodine status in lactating women and children < 2 years of age: a systematic review. *Nutrients* **14**, 1691.
- Sun D, Codling K, Chang S, *et al.* (2017) Eliminating iodine deficiency in China: achievements, challenges and global implications. *Nutrients* **9**, 361.
- Zimmermann MB (2020) Iodine and the iodine deficiency disorders. In *Present Knowledge in Nutrition*, pp. 429–441 [BP Marriott, DF Birt, VA Stallings, *et al.*, editors]. Amsterdam, The Netherlands: Elsevier.
- Ding Y, Indayati W, Basnet TB, *et al.* (2020) Dietary intake in lactating mothers in China 2018: report of a survey. *Nutr J* **19**, 1–13.
- Vandenbroucke JP, von Elm E, Altman DG, *et al.* (2014) Strengthening the Reporting of Observational Studies in Epidemiology (STROBE): explanation and elaboration. *Int J Surg* **12**, 1500–1524.
- Von Elm E, Altman DG, Egger M, *et al.* (2014) The Strengthening the Reporting of Observational Studies in Epidemiology (STROBE) Statement: guidelines for reporting observational studies. *Int J Surg* **12**, 1495–1499.
- Yu Z, Zheng C, Zheng W, *et al.* (2020) Mild-to-moderate iodine deficiency in a sample of pregnant women and salt iodine concentration from Zhejiang province, China. *Environ Geochem Health* **42**, 3811–3818.
- NCD Risk Factor Collaboration (2016) Trends in adult body-mass index in 200 countries from 1975 to 2014: a pooled analysis of 1698 population-based measurement studies with 19.2 million participants. *Lancet* **387**, 1377–1396.
- Li N, Lu Y, Si P, *et al.* (2022) The impact of moderately high preconception thyrotropin levels on ovarian reserve among euthyroid infertile women undergoing assisted reproductive technology. *Thyroid* **32**, 841–848.
- Mulrine HM, Skeaff SA, Ferguson EL, *et al.* (2010) Breast-milk iodine concentration declines over the first 6 mo postpartum in iodine-deficient women. *Am J Clin Nutr* **92**, 849–856.
- Jorgensen A, O'Leary P, James I, *et al.* (2016) Assessment of breast milk iodine concentrations in lactating women in Western Australia. *Nutrients* **8**, 699.
- Semba RD & Delange F (2001) Iodine in human milk: perspectives for infant health. *Nutr Rev* **59**, 269–278.
- Andersen SL, Møller M & Laurberg P (2014) Iodine concentrations in milk and in urine during breastfeeding are differently affected by maternal fluid intake. *Thyroid* **24**, 764–772.
- Dorea JG (2002) Iodine nutrition and breast feeding. *J Trace Elem Med Biol* **16**, 207–220.
- Yan Y, Zhang Y, Liu L, *et al.* (2006) *Method for determination of iodine in urine by As³⁺-Ce⁴⁺ catalytic spectrophotometry. Health Standard of China, WS/T107–2006*. Beijing: China Standard Press.
- Li F & He H (2018) Assessing the accuracy of diagnostic tests. *Shanghai Arch Psychiatry* **30**, 207.
- Parikh R, Mathai A, Parikh S, *et al.* (2008) Understanding and using sensitivity, specificity and predictive values. *Indian J Ophthalmol* **56**, 45.
- Maxim LD, Niebo R & Utell MJ (2014) Screening tests: a review with examples. *Inhal Toxicol* **26**, 811–828.
- Hosmer DW Jr, Lemeshow S & Sturdivant RX (2013) *Applied Logistic Regression*, vol 398. Amsterdam, The Netherlands: John Wiley & Sons.
- Russell R, Beard JL, Cousins RJ, *et al.* (2001) *Dietary Reference Intakes for Vitamin A, Vitamin K, Arsenic, Boron, Chromium, Copper, Iodine, Iron, Manganese, Molybdenum, Nickel, Silicon, Vanadium, and Zinc*. Washington, DC: Institute of Medicine (US) Panel on Micronutrients.
- Dold S, Zimmermann MB, Baumgartner J, *et al.* (2016) A dose-response crossover iodine balance study to determine iodine requirements in early infancy. *Am J Clin Nutr* **104**, 620–628.
- Wang W, Sun Y, Zhang M, *et al.* (2018) Breast milk and infant iodine status during the first 12 weeks of lactation in Tianjin City, China. *Asia Pac J Clin Nutr* **27**, 393–398.
- Laurberg P & Andersen SL (2014) Breast milk—a gateway to iodine-dependent brain development. *Nat Rev Endocrinol* **10**, 134–135.
- Turner R (1933) Iodine content of human skim milk from goitrous and non-goitrous regions. *Proc Soc Exp Biol Med* **30**, 1401–1403.
- Hercus CE & Roberts K (1927) The iodine content of foods, manures and animal products in relation to the prophylaxis of endemic goitre in New Zealand: studies from the University of Otago, New Zealand. *Epidemiol Infect* **26**, 49–83.
- Vermiglio F, Presti VL, Finocchiaro M, *et al.* (1992) Enhanced iodine concentrating capacity by the mammary gland in iodine deficient lactating women of an endemic goiter region in Sicily. *J Endocrinol Invest* **15**, 137–142.
- Böhles H, Aschenbrenner M, Roth M, *et al.* (1993) Development of thyroid gland volume during the first 3 months of life in breast-fed *v.* iodine-supplemented and iodine-free formula-fed infants. *Clin Invest* **71**, 13.
- Chan SS, Hams G, Wiley V, *et al.* (2003) Postpartum maternal iodine status and the relationship to neonatal thyroid function. *Thyroid* **13**, 873–876.
- Costeira MJ, Oliveira P, Ares S, *et al.* (2009) Iodine status of pregnant women and their progeny in the Minho Region of Portugal. *Thyroid* **19**, 157–163.



40. Kart P, Türkmen MK, Anik A, *et al.* (2021) The association of lactating mothers' urinary and breast milk iodine levels with iodine nutrition status and thyroid hormone levels of newborns. *Turk Arch Pediatr* **56**, 207–212.
41. Laurberg P, Nøhr SB, Pedersen KM, *et al.* (2004) Iodine nutrition in breast-fed infants is impaired by maternal smoking. *J Clin Endocrinol Metab* **89**, 181–187.
42. Kurtoglu S, Akcakus M, Kocaoglu C, *et al.* (2004) Iodine status remains critical in mother and infant in Central Anatolia (Kayseri) of Turkey. *Eur J Nutr* **43**, 297–303.
43. Kirk AB, Kroll M, Dyke JV, *et al.* (2012) Perchlorate, iodine supplements, iodized salt and breast milk iodine content. *Sci Total Environ* **420**, 73–78.
44. Zimmermann MB (2009) Iodine deficiency. *Endocr Rev* **30**, 376–408.
45. Tudehope DI (2013) Human milk and the nutritional needs of preterm infants. *J Pediatr* **162**, S17–S25.
46. Dror DK & Allen LH (2018) Iodine in human milk: a systematic review. *Adv Nutr* **9**, 347S–357S.
47. Andersson M, Karumbunathan V & Zimmermann MB (2012) Global iodine status in 2011 and trends over the past decade. *J Nutr* **142**, 744–750.
48. Zhang Y, Zhao X, Shan L, *et al.* (2023) Variations in breast milk iodine concentration over 24 h among lactating women in Northern China. *J Nutr* **153**, 208–214.
49. Bazrafshan HR, Mohammadian S, Ordookhani A, *et al.* (2005) An assessment of urinary and breast milk iodine concentrations in lactating mothers from Gorgan, Iran, 2003. *Thyroid* **15**, 1165–1168.
50. Azizi F & Smyth P (2009) Breastfeeding and maternal and infant iodine nutrition. *Clin Endocrinol* **70**, 803–809.
51. Delange F (2007) Iodine requirements during pregnancy, lactation and the neonatal period and indicators of optimal iodine nutrition. *Public Health Nutr* **10**, 1571–1580.
52. Eltom A, Eltom M, Elnagar B, *et al.* (2000) Changes in iodine metabolism during late pregnancy and lactation: a longitudinal study among Sudanese women. *Eur J Clin Nutr* **54**, 429–433.
53. Fu M, Gao Y, Guo W, *et al.* (2022) Mechanisms of sodium/iodide symporter-mediated mammary gland iodine compensation during lactation. *Nutrients* **14**, 3592.
54. Zhao Y, Ding A, Arya R, *et al.* (2018) Factors influencing the recruitment of lactating women in a clinical trial involving direct oral anticoagulants: a qualitative study. *Int J Clin Pharm* **40**, 1511–1518.
55. Agostoni C, Braegger C, Decsi T, *et al.* (2009) Breast-feeding: a commentary by the ESPGHAN Committee on Nutrition. *J Pediatr Gastroenterol Nutr* **49**, 112–125.
56. Saint L, Smith M & Hartmann P (1984) The yield and nutrient content of colostrum and milk of women from giving birth to 1 month post-partum. *Br J Nutr* **52**, 87–95.
57. König F, Andersson M, Hotz K, *et al.* (2011) Ten repeat collections for urinary iodine from spot samples or 24-hour samples are needed to reliably estimate individual iodine status in women. *J Nutr* **141**, 2049–2054.