Abstract

Objective: To assess the iodine nutritional status and the prevalence of goitre during pregnancy in a region of Hungary that appeared to be iodine sufficient in previous studies.

Design: A cross-sectional voluntary screening study was organized in which 313 pregnant women participated.

Methods: Urine iodine concentration and the volume of the thyroid gland were measured in every woman. In the presence of low urinary iodine concentrations, goitre, or both, thyroid function tests were performed.

Results: Iodine deficiency was found in 57.1% of the pregnant women, and was severe in 15.6%. The volume of the thyroid gland was enlarged in 19.2% of individuals. Nodular goitre was found in 17 women (5.4%). The frequency of goitre and the mean thyroid volume were increased in the group of iodine-deficient women. In the 89 cases of iodine deficiency or goitre, thyrotrophin concentrations were in the normal range; however, the free triiodothyronine:free throxine ratio was increased in 97% of them, indicating that the thyroid gland was in a stimulated state in these individuals.

Conclusions: Iodine deficiency with high prevalence of goitre was recognized among pregnant women in an area that previously appeared to be iodine sufficient. An unexpected mild iodine deficiency was also noted in the non-pregnant control group. Reassessment and continuous monitoring of iodine nutritional status is warranted even in populations that are apparently considered to be ‘at no risk’ of iodine deficiency, especially in pregnant women. Regular administration of iodine, starting at preconception or in early pregnancy and continuing during the period of nursing, is recommended in these regions.

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Introduction

Iodine supply is inadequate in most parts of Hungary, except in small areas of the Great Plains (1–4). Several attempts have been made to eliminate this iodine deficiency (5). Since the 1970s, the issue of inadequate iodine intake has been publicized by several authors as a problem requiring immediate attention (1–3). Iodized salt and iodine-containing tablets can be purchased on a voluntary basis; however, as no state-organized program has been developed, inadequate iodine intake has remained a nation-wide health problem. Previous studies found the region of Debrecen (which is situated on the Great Plains) to be an area of adequate iodine intake as judged by the low frequency of goitre in the population, except for a few clusters of small villages (4–6).

Previous investigations confirmed that the most detrimental effects of inadequate nutritional iodine intake appear in pregnant women and in children (7–9). Hormonal changes and metabolic demands during pregnancy result in alterations in the biochemical parameters of thyroid function (10). The effects of iodine deficiency become manifest even in geographical areas of borderline iodine supply, because the thyroid gland cannot meet the demand for increased production of thyroid hormone (10–13). The main changes in thyroid metabolism in pregnancy are well characterized (10–15). Iodine deficiency triggers compensatory stimulating events affecting the thyroid gland of both the mother and the fetus. With respect to long-term effects, the development of goitre is only of secondary importance; the detrimental effect of mild thyroid insufficiency (defined by a moderate decrease in
maternal free thyroxine (FT₄) concentration) on the neuropsychological development of the child has been repeatedly emphasized (9, 16–22).

We assumed that, in a community with apparently sufficient iodine intake, the increased iodine demand during pregnancy might lead to manifest iodine deficiency in pregnant women. In the present study we show that the iodine nutritional status found in the general population may not reliably reflect the iodine supply of pregnant women living in the same region.

Participants and methods

Participants

We enrolled 313 pregnant women into a cross-sectional voluntary screening study, which was coordinated by the physicians and nurses of the Health Care System in Debrecen. The ages of the women were 26.8 ± 5.2 (range 17–47) years and the gestational ages were 23.6 ± 7.9 (6–39) weeks. Multivitamin products, which may contain iodine, are routinely offered to pregnant women by gynaecologists. The consumption of these drugs and the use of iodized salt were elicited by detailed history and four groups were established according to the form of iodine supplementation: group A, no iodine supplementation (n = 119); group B, regular use of iodized salt, no consumption of iodine-containing multivitamins (n = 50); group C, taking iodine-containing multivitamins, no use of iodized salt (n = 91); group D, regular use of iodized salt and iodine-containing multivitamin tablets (n = 53). Overall, 46% of women consumed iodine-containing multivitamin products. The gestation times of those in the non-multivitamin consuming groups (A and B: 24.6 ± 8.1 weeks) and the multivitamin consuming groups (C and D: 23.9 ± 7.4 weeks) did not differ significantly.

To assess the iodine supply in the general population, samples from 220 non-pregnant women were also analysed.

Methods

Urine iodine concentration was determined from the first morning urine void, by the Sandell–Kolthoff colourimetric method (23). The iodine concentration of the urine was standardized against that of creatinine, urine creatinine concentrations being measured on the days of sampling. Urine specimens were stored at −20°C until required for analysis. The iodine deficiency was regarded to be mild with urinary iodine concentration between 50 and 100 µg iodine/g creatinine, moderate in the range 25–50 µg iodine/g creatinine and severe when values were less than 25 µg iodine/g creatinine.

Thyroid volume measurements were performed by ultrasound (Siemens Sonoline SL-1, 7.5 MHz linear transducer). Each lobe of the thyroid gland was assessed separately by measuring the three main diameters, and the total volume of the thyroid was calculated by the algorithm \( \pi / 6 \times \text{height} \times \text{width} \times \text{depth} \). The upper limit of normal thyroid volume was 18 ml.

In cases of urinary iodine less than 50 µg/g creatinine or diffuse or nodular goitre, further tests were offered to the expectant mothers; 89 of them were willing to continue the investigations. Thyroid-stimulating hormone (TSH) was assayed by a third-generation immunochemiluminometric assay and FT₄ by a competitive-binding, one-step chemiluminescent immunoassay (both from Byk-Sangtec Diagnostica GmbH & Co., Dietzenbach, Germany). Free tri-iodothyronine (FT₃) was assayed by a two-step saturation-competitive binding microparticulate enzyme immunoassay (Abbott Laboratories, Abbott Park, IL, USA).

Data analysis

The relationship between thyroid volume and iodine excretion was analysed statistically using ANOVA, and groups were compared using the Newman–Keuls test. The median test and the Pearson \( \chi^2 \) test were applied to analyse the effect of iodine supplementation on urinary iodine excretion. The level of significance in all statistical tests was set at \( P \leq 0.05 \).

Results

Urinary iodine excretion

Iodine deficiency among pregnant women in the city of Debrecen was much more frequent than we had
expected before the study (Fig. 1): normal urine iodine concentration was measured only in 42.9% of the women. The level of iodine deficiency was mild in 25.6%, moderate in 15.9% and severe in 15.6% of pregnant women. The median urinary iodine excretion was 57, 68, 130 and 115 μg/g creatinine in groups A, B, C and D respectively (Fig. 2); that in the control group of 220 non-pregnant woman was 82 μg iodine/g creatinine. The regular use of iodized salt did not influence the median urinary iodine excretion, but the consumption of multivitamin drug products increased the urinary iodine concentrations significantly. No significant difference in the median urinary iodine excretion was found between groups C and D (consumption of iodine-containing multivitamins respectively without and with iodized salt), therefore we combined the results of groups A and B, and those of groups C and D in further analyses. Although the frequency of iodine deficiency was lower in women taking iodine-containing multivitamins than in those without iodine supplementation, adequate iodine intake was measured only in 55% of women who were taking regular multivitamin iodine supplementation (Fig. 3). Iodine deficiency was diagnosed in 69% of expectant mothers without iodine-containing multivitamin consumption, and in 45% taking iodine-containing multivitamins.

**Maternal thyroid ultrasound**

The volume of the thyroid gland was in excess of the normal value of 18 ml in 19.2% of the women. Nodular goitre was diagnosed in 17 (5.4%). The mean thyroid volume of women with severe iodine deficiency was significantly larger than that in the group with adequate iodine intake, and the frequency of goitre was increased in all groups with iodine deficiency (Fig. 4).

**Maternal thyroid parameters**

In 89 women with low urine iodine excretion or goitre, further thyroid tests were performed. The mean TSH was 1.11 ± 0.54 mIU/l, which is within the normal range of 0.3–3.0 mIU/l. However, the FT3:FT4 ratio was remarkably increased at 0.45 ± 0.13, and was above the upper limit of normal (0.25) in 97% of the women.

**Discussion**

Considering that previous studies have concluded that the region of Debrecen was an area of sufficient iodine supply (1, 2, 4, 5), it is striking that, on the basis of urinary iodine excretion, we found mild iodine deficiency in the non-pregnant female population. More importantly, 57.1% of pregnant women in Debrecen were found to be iodine deficient and 19.2% had goitre. One possible explanation is a change in available iodine

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![Figure 2](Image)

**Figure 2** Urinary iodine excretion in women grouped according to their iodine supplementation. Group A: no iodine supplementation; group B: regular use of iodized salt; group C: consumption of iodine-containing multivitamins, no use of iodized salt; group D: regular use of iodine-containing multivitamins and iodized salt. Data are presented as median+upper quartile. The median urinary iodine excretion was significantly lower in groups A and B than in groups C and D (*P* < 0.01). The use of iodized salt did not influence the urinary iodine concentrations. No significant difference was found between groups C and D.

![Figure 3](Image)

**Figure 3** The distribution of urinary iodine concentrations separately among pregnant women who were (open bars, n=144) or were not (solid bars, n=169) consuming iodine-containing multivitamins (150 μg/day). The prevalence of low urinary iodine excretion was significantly greater in the latter group (*P* < 0.01). Even with iodine replacement, only 65% of the women were iodine sufficient.
in this area. A decrease in iodine intake in the past 15 years has also been observed in the USA, resulting in low urinary iodine concentrations in 15% of women of childbearing age (24). In Hungary, where both iodized and non-iodized salts are available, iodized salt is not used during the industrial production of foodstuffs, and usually only a small part of a meal is prepared at home. In our study, the regular use of iodized salt at home before and during pregnancy did not result in an improvement in iodine nutritional status, as urinary iodine excretion in women supplementing their iodine intake did not differ significantly from that in women not taking any form of iodine replacement. Among pregnant women receiving no iodine supplementation or using only iodized salt at home, the prevalence of iodine deficiency was 69%.

Nearly 50% of pregnant women involved in this study received iodine supplementation in the form of multivitamin tablets containing 150 μg iodine, taken once a day. These products were offered as part of the routine obstetric regimen, and not with the specific purpose of counteracting iodine deficiency (as iodine deficiency had not been suspected). Even in pregnant women receiving this form of iodine supplementation, urinary iodine excretion was sufficient in only 55%. The reason for this may be multifold. The recent WHO recommendation is for the daily nutritional intake of 200 μg iodine during pregnancy and lactation (13). However, in goitre-endemic regions we would expect to encounter decreased iodine concentrations in the thyroid at the beginning of pregnancy (11). In such cases, favourable results are more likely to be achieved with 300–400 μg daily iodine. Compliance with the drug therapy is also an important factor. Finally, exact data are not available concerning the pharmacokinetic properties of the iodine-containing drug products. Our results demonstrated that occasional iodine supplementation in the form of multivitamins prescribed by gynaecologists was not adequate across the pregnant population as a whole. Conscious iodine supplementation therapy is necessary, with regular control of the compliance of women.

The urinary iodine excretion and the frequency of goitre in pregnant women in Debrecen were similar to those found by studies examining large populations around Brussels and in the southwest of France, which are marginally iodine-deficient regions (11, 25). These results indicate that a general survey of iodine nutritional status in pregnancy is recommended even in regions that are presumed to be iodine sufficient.

The causative role of iodine deficiency underlying the high frequency of goitre was strongly supported by the facts that the thyroid volume and the frequency of goitre were significantly larger in women with iodine deficiency than in the group with adequate iodine intake. As there are no methods available to estimate intrathyroidal iodine pools on an individual basis, biochemical markers such as increased FT3:FT4 ratio, decreased FT4, increased thyroglobulin concentration and increased thyroid gland volume are useful surrogate tools (10–13). The FT3:FT4 ratio was increased in 97% of iodine-deficient women. This can be interpreted as the result of the pathological stimulation of the thyroid gland by iodine deficiency.

The association of low FT4 with a normal TSH value seems to be typical for iodine-deficient pregnancies. In late pregnancies, decreased FT4 concentrations with normal TSH were found in 17% and 33% of women in the southwest of France and in Belgium, respectively (11, 25). The exact cause of this phenomenon is not clearly understood. One explanation is that the determination of free hormones is influenced by the high serum concentration of thyroxin-binding globulin, resulting in false-positive low FT4 concentrations (26). Others concluded that iodine-deficient pregnant women suffered from true hypothyroxinaemia (14). The latter theory is supported by the fact that, in a region of adequate iodine intake, the prevalence of low FT4 concentration was only 0.3% (27).

In conclusion, in the area of Debrecen, which previously appeared to be iodine sufficient, 57.1% of the pregnant women suffered from iodine deficiency and 19.2% had goitre. An unexpected mild iodine deficiency was also noted in the non-pregnant control group. The home usage of iodized salt does not seem to improve appropriately the iodine nutritional status in pregnancy, and occasional iodine supplementation in the form of multivitamin tablets resulted in partial success. Reassessment and continuous monitoring of iodine nutritional status in pregnancy is warranted even in populations that are apparently considered to be ‘at no risk’ of iodine deficiency. Regular administration of iodine, starting at preconception or in early pregnancy and continuing during the period of nursing, is recommended in these regions.
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References


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