The incidence of congenital hypothyroidism (CH) in Guangxi, China and the predictors of permanent and transient CH

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Abstract

Background: The incidence of congenital hypothyroidism (CH) differs significantly among different ethnicities and regions, and early differentiation of transient CH is important to avoid unnecessary prolonged treatment with L-T₄.

Objective: To investigate the incidence of CH based on the newborn screening program in Guangxi Zhuang Autonomous Region, China, and to analyze the predictors that might allow for an early differentiation between permanent (P) and transient (T) CH.

Design and methods: Data from newborn screening program over a seven-year period (January 2009 to January 2016) at Guangxi Maternal and Child Health Hospital are analyzed. Blood samples were collected on filter paper between 3 and 7 days after birth, and TSH level was measured by time-resolved fluorescence assay. Individuals with increased TSH (TSH ≥ 8 IU/L) levels detected by newborn screening were recalled for further evaluation. Serum TSH, FT₃ and FT₄ were determined by electrochemiluminescence assay using venous blood samples. Diagnosis of CH is based on elevated TSH levels (>10 IU/L) and decreased FT₄ levels (<12 pmol/L). Patients with elevated TSH levels and normal FT₄ levels were diagnosed as hyperthyrotropinemia. Permanent or transient CH was determined by using the results of thyroid function tests after temporary withdrawal of L-T₄ therapy at approximately 2–3 years of age.

Results: Among 1,238,340 infants in the newborn screening program, 14,443 individuals were recalled for reevaluation (recall rate 1.18%), 911 and 731 individuals were subsequently determined to have hyperthyrotropinemia and CH respectively; thus, a prevalence of 1:1359 and 1:1694 for hyperthyrotropinemia and CH. Of the 731 patients with CH, 161 patients were diagnosed with permanent CH (PCH), and 159 patients were diagnosed with transient CH (TCH), the other 411 patients are too young to determine their subtypes. Patients with PCH required an increasing dose of L-T₄ during the first few years, whereas patients with TCH required a decreased dose of L-T₄. The TSH levels at diagnosis and the dose of L-T₄ used were significantly higher in PCH cases than in TCH cases.

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Key Words

- congenital hypothyroidism
- prevalence
- predictor
- L-T₄ dose
- Guangxi
- China
transient cases. The FT₄ levels at diagnosis were significantly lower in PCH cases than in TCH cases. The TSH levels at diagnosis, FT₄ levels at diagnosis and L-T₄ doses at 90 days were evaluated as predictors for differentiating PCH and TCH, and their accuracy at their respective optimal cutoffs were determined to be 60.6%, 66.7% and 93.9%, respectively. Conclusions: The CH incidence in Guangxi Zhuang Autonomous Region is slightly higher (1:1694) compared to the worldwide levels (1/2000–1/4000). The PCH and TCH ratio is close to 1; thus, the estimated PCH incidence is 1/3388, which is similar to reported worldwide average incidence (1/3000). The L-T₄ dose required at 90 days (>30 μg/day) has the highest predictive value for PCH. Earlier differentiation of PCH and TCH helps to determine appropriate treatment course.

Introduction

Congenital hypothyroidism (CH) is one of the most common causes of intellectual disability and growth retardation with worldwide incidence ranging from 1:2000 to 1:4000 newborns (1, 2, 3). Newborn screening allows for early detection and treatment of CH, thus effectively preventing intellectual disability and short stature due to lack of thyroid hormone.

The incidence of CH differs significantly among different ethnicities and regions. For example, Zeinalzadeh and coworkers reported a CH prevalence of 1:666 (TSH cutoff level 10IU/L) in East Azerbaijan, Iran (4), whereas a CH prevalence of 1:2591 (TSH cutoff level 10IU/L) was reported in the Republic of Macedonia (5). Two previous studies reported the CH prevalence based on newborn screening programs in Han Chinese, and the incidence of CH was reported as 1:1678 (TSH cutoff level 9IU/L) and 1:3009 (TSH cutoff level 10–15IU/L), respectively in Zhejiang (6) and Shanghai (7) province. Little is known about the incidence of CH in Guangxi Zhuang Autonomous Region, which is located in South-west China with a population of distinct ethnic composition.

CH can be classified into PCH and TCH according to the clinical course. PCH refers to a persistent deficiency of thyroid hormone that requires lifelong treatment. TCH refers to a temporary deficiency of thyroid hormone at the early stage of life, but thyroid hormone production recovers to normal afterwards. Recovery to euthyroidism typically occurs in the first few months or years of life. The current guidelines recommend that all CH children be re-evaluated at the age of 3 years (8). Early differentiation of TCH is important to avoid unnecessary prolonged treatment with L-T₄, and it is also helpful for predicting prognosis and alleviating families’ psychological burden. Thus, we investigated the CH incidence based on the newborn screening program in Guangxi Zhuang Autonomous Region, China and to analyze the factors that might allow an early differentiation between TCH and PCH.

Subjects and methods

1,238,340 newborns were screened for CH at the duration from January 2009 to January 2016 at the Newborn Screening Center of Guangxi, China. This center is responsible for half of the total population in Guangxi province. The coverage for newborn screening is 95.2% for this part of the country. The screening follows a standard screening protocol. Between 3 and 7 days after birth, 3 drops of heel blood were collected on a filter paper (S&S 903). The thyroid-stimulating hormone (TSH) level was measured by time-resolved fluorescence assay (Perkin Elmer).

The analytical sensitivity of TSH test was typically better than 0.002IU/L. The accuracy of the TSH test was assessed using both intra- and inter-assay. Intra-assay coefficients of variation at TSH concentrations of 14.6, 24.0 and 58.6IU/L were 7.1, 6.3 and 7.0%, respectively, while the inter-assay coefficients of variation at the same TSH concentrations were 8.1, 9.0 and 7.8%, respectively. Subjects with increased TSH (TSH≥8IU/L) levels were contacted and re-evaluated. Pre-term (<37 weeks gestation) and low-birth-weight infants (<2000g) underwent a second mass screening one month after birth, when their body weight reaches 2500g, even if data in the first screening at age 3–7 days was within the normal range.

The confirmatory test was carried out between 7 and 28 days after birth by electrochemiluminescence
assay (Cobas e601, Roche Diagnostics) with venous blood samples. The test results were compared to age-specific reference as included in Supplementary Table 1 (see section on supplementary data given at the end of this article). Diagnosis of CH is based on elevated TSH levels (>10 IU/L) and decreased FT₄ levels (<12 pmol/L). Patients with elevated TSH levels and normal FT₄ levels were diagnosed as hyperthyrotropinemia (HT). Thyroid ultrasonography and ⁹⁹mTc scintigraphy were performed during the neonatal period before treatment. L-T₄ replacement therapy started after the diagnosis of CH, and the dose was adjusted according to the serum TSH and FT₄ levels. The target serum FT₄ level should be >50% of the normal range by age. The target for TSH level should be within the normal range by age. Permanent CH was determined by temporary withdrawal of L-T₄ therapy at approximately 2–3 years of age. After one month of discontinuation of L-T₄ treatment, TSH and FT₄ levels were measured in venous blood sample. Individuals who showed contiguous dependency on L-T₄ were diagnosed with permanent CH. These children were then repeatedly evaluated at regular intervals for 1–1.5 years to monitor thyroid function. Those who did not need contiguous L-T₄ therapy were diagnosed with transient CH. The study was approved by the Medical Ethics Committee of Guangxi Maternal and Child Health Hospital. Informed consent was obtained from the parents of the patients.

### Statistical analysis

The data were analyzed using R software. Qualitative variables were compared using the Fisher test and quantitative variables were compared by t-test. All P values were two-sided with P<0.05 being considered as significant. A receiver-operating characteristic (ROC) curve analysis was performed to identify the optimal cutoff values of predictor, which were able to distinguish between PCH and TCH. The best cutoffs were defined by the highest Youden index ((specificity + sensitivity)−1).

### Results

During the study period, 1,238,340 newborns were screened. A skewed gender ratio was noticed in this population: 667,676 (53.9%) were male and 570,664 (46.1%) were female. The female:male ratio was 1:1.17. 80% of samples were taken from neonates at 3–5 days

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**Table 1** Distribution of serum TSH levels among screened newborns (n=12,38,340).

<table>
<thead>
<tr>
<th>Distribution of TSH levels (IU/L)</th>
<th>N</th>
<th>%</th>
</tr>
</thead>
<tbody>
<tr>
<td>0–4.9</td>
<td>11,40,493</td>
<td>92.1</td>
</tr>
<tr>
<td>5–7.9</td>
<td>81,062</td>
<td>6.5</td>
</tr>
<tr>
<td>8–9.9</td>
<td>9805</td>
<td>0.8</td>
</tr>
<tr>
<td>10–19.9</td>
<td>6109</td>
<td>0.5</td>
</tr>
<tr>
<td>≥20</td>
<td>871</td>
<td>0.1</td>
</tr>
<tr>
<td>Total</td>
<td>12,38,340</td>
<td>100</td>
</tr>
</tbody>
</table>

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**Figure 1** Variation of the TSH levels and positive rate based on the timing of newborn screening sampling.
after birth. 97% of the neonates were screened within the first two weeks after birth. The distribution of serum TSH levels among screened newborns is shown in Table 1. We plotted the mean TSH levels and the positive rates from days 3 to day 14 and observed a concomitant decline of TSH levels and screening positive rates, but a significant improvement in confirmed positive rates (Fig. 1).

The mean TSH level at newborn screening in all measured samples was 2.3±3.2IU/L. The mean birth weight was 3113.1±438.9 g and the mean gestation was 38.8±1.6 weeks. Overall, 14,443 neonates (1.18%) were detected to have high TSH levels in the first and second dry samples and were recalled for confirmatory test. The confirmatory test revealed normal FT₄ levels and elevation of TSH in 911 neonates. These newborns were diagnosed as HT. 731 were diagnosed as CH with elevation of TSH levels and decreased FT₄ levels. The incidence of CH and hyperthyrotropinemia was 1/1694 and 1/1359, respectively. The significant differences between HT and CH regarding TSH (Fig. 2A) and FT₄ levels at diagnosis (Fig. 2B) are expected based on the definition. The analysis also revealed that more females are diagnosed with CH and more males are diagnosed with HT (Fig. 2E). Compared with individuals with HT, significantly more individuals with CH had abnormal thyroid morphology (Fig. 2F). There were no statistical differences between HT and CH regarding weight (Fig. 2C) and gestational weeks (Fig. 2D).

Among neonates with CH, 360 (49.2) were male and 371 (50.8) were female (female: male ratio 1.03:1). The prevalence of CH among male and female neonates was 1:1855 and 1:1538, respectively, and the difference was statistically significant (P=0.012). The mean TSH level at diagnosis of CH neonates was 83.51±26.84IU/L. The mean birth weight of CH neonates was 3158.6±775.6 g and the mean gestation was 38.9±1.6 weeks. The birth weight and gestation weeks are not different in CH neonates in comparison to non-CH neonates. 57,981 of all newborns and 52 of the 731 CH cases were pre-term babies, and 10,509 of all newborns and 12 of the 731 CH had low birth weight. The CH prevalence are significantly higher among pre-term (P<0.002) and small for gestational age newborns (P<0.019) compared with full-term newborns and norm birth weight newborns.

Of the 731 patients with CH, 161 patients were diagnosed as PCH and 159 patients were diagnosed as TCH, the PCH-to-TCH rate was about 1:1. The other 411 patients are too young to withdraw L-T₄ therapy, so we cannot determine their clinical phenotype. There were statistical differences between PCH and TCH regarding TSH levels at diagnosis (Fig. 3A), FT₄ levels at diagnosis (Fig. 3B) and thyroid morphology composition (Fig. 3F). There were no statistical differences between PCH and TCH regarding gestation weeks (Fig. 3D), weight (Fig. 3C) and gender composition (Fig. 3E).
The individual doses and average dose of L-T₄ at different age of newborns with PCH and TCH were shown in Fig. 4A and B, respectively. It is obvious that the PCH patients require a higher dosage and longer treatment duration. At the age of 900 days, 29.6% of TCH patients were treated with an average L-T₄ dose of 7.8μg/day, whereas 100% of PCH patients were treated with an average dose of 38.9μg/day.

The dependence on L-T₄ showed a significant difference between PCH and TCH. To further evaluate the effectiveness of using L-T₄ as a predictor, we performed ROC curve analysis at different times after diagnosis. The differentiating power increases at a later time point (Fig. 5A). The L-T₄ doses used at 90 days is a relatively better predictor taking into account of early diagnosis, the area under the curve (AUC), sensitivity and specificity (Supplementary Table 2).

We also evaluated the effectiveness for differentiating PCH and TCH using TSH levels at diagnosis, FT₄ levels at diagnosis and L-T₄ doses at 90 days (Fig. 5B). Based on the ROC curve analysis, we determined the optimal cutoff values for predicting CH subtypes: The TSH levels at diagnosis was set at 97.1 IU/L, which yielded a sensitivity of 84.56% and a specificity of 52.94%, with the AUC at 0.6889; the L-T₄ doses at 90 days was set at 30μg/day, which yielded a sensitivity of 58.09% and a specificity of 78.68%, with the AUC at 0.7220 and the value of FT₄ levels at diagnosis was set at 4.6 pmol/L, which yielded a sensitivity of 60.29% and a specificity of 69.85%, with the AUC at 0.6690.

To further assess the accuracy of the cutoffs for differentiating PCH and TCH determined by retrospective patient cohort, we used 33 prospective patients subsequently (Supplementary Table 3), who were identified with PCH or TCH from the total 731 CH cases during the process of manuscript preparation. As a result, the accuracy of predicting the CH subtypes are 60.6, 66.7 and 93.9% using the cutoff values of TSH levels at diagnosis, FT₄ levels at diagnosis and L-T₄ doses at 90 days respectively. The L-T₄ doses at 90 days is the best predictor of high accuracy.

**Discussion**

In this study, over 1.2 million newborns from Guangxi Zhuang Autonomous Region, China were screened for CH using a standard procedure with 8 IU/L as a cutoff value for TSH. As a result, 1.18% neonates were recalled for confirmatory test. This re-rate is higher than that of Mexico (0.15%) (9), Brazil (0.21%) (10), Netherlands (0.18%) (11) and Thailand (0.24%) (12), but lower than that of Iran (1.76%) (13), Turkey (7.2%) (14), India (1.39%) (15) and Bahrain (4%) (16).

The confirmed CH incidence is 1/1694 in this population. It is known that the incidence of CH could
vary significantly from one ethnic group to another (1/370 (17) to 1/10,000 (18)). Currently, the overall incidence of CH in Chinese population is not known, two previous studies based on populations in Zhejiang and Shanghai provided the incidences for Han Chinese as 1:1678 and 1:3009, respectively. Guangxi Zhuang Autonomous Region is dominated with Zhuang minority, a distinct ethnic group from Han. Our study revealed that the incidence of CH among newborns in Guangxi Zhuang Autonomous Region is similar to that of Han Chinese in Zhejiang (6) and slightly higher than the average worldwide incidence (1/2000–1/4000) (1, 2, 3).

We have used a relative low cutoff value of TSH in this newborn screening program. 97% of the screening test was done within 3–14 days of life. For the first time, we plotted the mean TSH levels in the newborn period based on a large cohort of population and noticed a rapid decrease of TSH level in the first several days (Fig. 1). Noticeably, the positive screening rates also showed a similar decline. More importantly, the confirmed positive rates or false-positive rate also correlated with the average level of TSH at different days after birth. This finding challenged the validity of using one cutoff value of TSH for newborn CH screening at any days after birth. Further study can help to determine the right cutoff value for samples collected at different days after birth.

Recent studies suggest that the TSH levels, weight, gender, FT$_4$ levels, gestational weeks and the thyroid morphology may correlate with the etiology of CH (19, 20, 21). In our study, there were statistical differences between HT and CH regarding TSH levels, gender composition, FT$_4$ levels and thyroid morphology composition. There were also statistical differences between PCH and TCH regarding the TSH levels, FT$_4$ levels and the thyroid morphology, but no significant differences in gestation weeks, gender and weight were found between the two groups. It demonstrated the correlation between the three
factors (TSH levels, FT\textsubscript{4} levels and thyroid morphology) and hypothryoidism.

Nearly all screening programs report a female preponderance among CH patients, often approaching a 2:1 female-to-male ratio (1, 5). In our study, this ratio was 1.03:1. The difference may be caused by the population gender bias among screened newborns (male:female 1.17) and etiological diversity of CH in different ethnic groups. The PCH-to-TCH rate in our CH patients is about 1:1. It is similar to that recently reported by other authors, who found rates ranging from 38 to 46% (19, 21, 22). Therefore, according to our and other authors’ results, more than one-third of the infants diagnosed with CH by neonatal screening will not require lifelong L-T\textsubscript{4} therapy, and reliable markers that would allow earlier detection of TCH might be useful.

Treatment plan and prognosis differ significantly for patients with PCH or TCH. Often, it is not known what CH subtype each newborn-screen-positive patient has until L-T\textsubscript{4} withdrawal at approximately 2–3 years of age. It is beneficial to have an earlier differentiation. In the past few years, several studies (19, 20, 21, 23, 24, 25, 26, 27, 28, 29) tried to identify the factors that could allow an early discrimination between the cases with either P or T CH. In a study conducted by Unuvar and coworkers (19), T4 dose required to maintain a euthyroid state was the parameter that distinguished between TCH and PCH. Rabbiosi and coworkers (21), in a study conducted in Italy, reported major risk factors for PCH were prematurity, thyroid hypoplasia at diagnosis, first-degree familial history of goiter/nodules and high L-T\textsubscript{4} requirements at follow-up. Hashemipour and coworkers, in a study conducted in Iran, reported PCH was associated with higher initial TSH and T4 levels than TCH. Skordis and coworkers (27) stated that L-T\textsubscript{4} requirements >4.9 μg/kg/day at 12 months or >4.27 μg/kg/day at 24 months are highly suggestive of PCH; L-T\textsubscript{4} requirements <1.7 μg/kg/day at 12 months or <1.45 μg/kg/day at 24 months are highly suggestive of TCH. Messina and coworkers (20) reported the initial measurements of the serum TSH level and the required doses of L-T\textsubscript{4} therapy for maintaining normal thyroid hormone levels, growth and development may have a predictive role for differentiating PCH from TCH.

So far, although a few studies have been conducted with the purpose of determining how best to discriminate between cases with either TCH or PCH before the age of 3 years, the evolution of CH remains difficult to predict, mainly due to the lack of large-scale, long-term and in-depth studies. In our study, we evaluated the effectiveness using L-T\textsubscript{4} dose, as well as TSH and FT\textsubscript{4} levels at diagnosis as predictors for an early differentiation using so far the largest cohort of patients. Using multiple statistical methods, we found that children with a higher initial L-T\textsubscript{4} dose and requiring increasing doses of L-T\textsubscript{4} during the first few years of treatment are likely to have PCH. Conversely, patients with a lower initial L-T\textsubscript{4} dose
and requiring decreasing doses are likely to have TCH. High TSH levels and low FT4 levels at diagnosis were found to be significant predictors for PCH. The L-T4 doses, however, are the best independent predictor. An L-T4 dose requirement at 90 days >30 μg/day demonstrated an accuracy of 93.9% for our prospective cohort.

In conclusion, the incidence of CH and hyperthyrotropinemia among newborns in Guangxi, Zhuang Autonomous Region, China was 1/1694 and 1/1359, respectively. The permanent CH and transient CH ratio is close to 1; thus, the estimated PCH incidence is 1/3708. There were statistical differences between hyperthyrotropinemia and CH regarding TSH levels, sex, FT4 levels and thyroid morphology; there were also statistical differences between PCH and TCH regarding the same four factors. Children with CH requiring increasing L-T4 doses during the first years of treatment are at risk for PCH; children with CH demonstrating decreased L-T4 dependency to maintain normal TSH values over time are likely to have TCH. An L-T4 dose requirement at 90 days >30 μg/day is highly suggestive of PCH.

Supplementary data
This is linked to the online version of the paper at http://dx.doi.org/10.1530/EC-17-0289.

Declaration of interest
The authors declare that there is no conflict of interest that could be perceived as prejudicing the impartiality of the research reported.

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