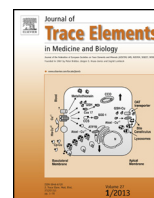




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Short communication

Iodine nutrition status amongst neonates in Kangra district, Himachal Pradesh

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ABSTRACT

Iodine deficiency (ID) is an endemic health problem in Kangra district, Himachal Pradesh (HP) state. ID leads to mental retardation, deaf mutism, squint, dwarfism, spastic diplegia, neurological defects and congenital anomalies. Iodine nutrition status amongst neonates can be assessed by estimating thyroid stimulating hormone (TSH). The present study was conducted with an objective to assess the iodine nutrition status amongst Neonates in Kangra district, HP. All of the hospitals in the district which provide obstetric services were enlisted, of which three were selected for this survey. A total of 613 umbilical cord blood samples of neonates were collected on filter paper and analyzed for TSH. WHO (2007) reported that that a <3% frequency of TSH concentrations above 5 mIU/L in samples collected 3–4 days after birth indicates iodine sufficiency in a population. In our study we found that 73.4% of the neonates had TSH levels of more than 5 mIU/L, thus indicating ID in the population studied. Iodine deficiency continues to be a public health problem in Kangra district, Himachal Pradesh.

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Introduction

Iodine deficiency disorders (IDD) is a serious global public health problem estimated to affect about one billion people across the world [1]. ID in neonates leads to cretinism including mental deficiency with a mixture of mutism, spastic diplegia, squint, hypothyroidism and short stature. Iodine deficiency amongst neonates lowers circulating thyroxine level and raises serum TSH. Hence, iodine deficient neonates have higher serum TSH concentration than do iodine sufficient neonates. In iodine-sufficient populations, about one in 4000 neonates may have congenital hypothyroidism, usually because of thyroid dysplasia. Thyroid stimulating hormone affects proper development of the central nervous system, particularly its myelination [2].

Kangra district, HP is a known endemic region for ID. The salt iodization programme in Kangra district was initiated in 1962, and thus has been running for over 50 years. Studies are available on the status of iodine nutrition in School Age Children (SAC) during the iodization programme which has documented the prevalence of ID as 12.1% (1999), 19.8% (2007) and 15.8% (2013), respectively

[3–5]; however, there is a lack of data on the status of iodine nutrition amongst neonates from this district. The present study was conducted with an objective to assess the iodine nutrition status amongst neonates in Kangra district, HP, with an aim to provide evidence to state health authorities to strengthen the IDD programme, if required.

Materials and methods

The study was carried out from May 2013–July 2013. In Kangra district a total of nine Hospitals/Community Health Centres which conducted more than 100 deliveries per annum were selected. Of these, three hospitals were randomly selected. The 613 births occurring consecutively in these three hospitals during the study period were included for estimation of iodine nutrition amongst neonates. The blood samples were collected after the written consent of the mothers (of the neonates). The umbilical cord blood sample was taken on a filter paper and was transported to the Central Laboratory in Delhi for estimation of TSH. Caesarean delivery, delivery in which iodine preparations were used and pregnant mothers who were on anti thyroid therapies were excluded from the study.

Umbilical cord blood collection

Cord blood was collected before placental delivery within five minutes after birth to avoid clotting. One drop of blood was applied

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to filter paper. The spots were dried at room temperature and the filter papers were sealed and kept in a freezer until assayed in the laboratory. The samples were stored at 4 °C before analysis.

The samples were estimated for TSH by using sandwich enzyme linked immuno-sorbent assay (ELISA) method. Dry blood spots were eluted in anti-TSH antibodies coated with micro wells and were incubated with peroxidase labelled anti-TSH monoclonal antibodies. After washing, the unbound antibodies were washed off and the bound conjugate remained in the micro well. These bound conjugates further react with substrate 3,3',5,5'-tetramethylbenzidine (TMB) and produce a colour product. The concentration of TSH is directly proportional to the colour produced. Absorbance was read at 450 nm and a value of TSH was expressed in the units' mIU/l of blood. In order to measure the concentration of TSH in the test sample, the calibration standards and controls were used. The calibration standards and controls were assayed for producing a standard curve of TSH by O.D versus TSH concentration (mIU/l). Therefore, by comparing the O.D of the test samples to this standard curve, the concentration of the TSH was determined [6–8].

Ethical approval was obtained from the ethics committee of the All India Institute of Medical Sciences, New Delhi.

Sample size

Keeping in view the anticipated prevalence of 2.9% [9], a confidence level of 95%, absolute precision of 2.0 and a design effect of 2, a total sample size of 541 was calculated. However we studied a total of 613 subjects.

Statistical analysis

The TSH normality and abnormality was compared in gender by using chi-square test.

Results

A total of 613 umbilical cord blood samples of neonates (324 males and 289 females) on filter papers were collected and analyzed for TSH. Three hundred and thirty six (54.8%) samples had TSH level of <10 mIU/l, 217 (35.4%) had 10 to <20 mIU/l and 60 (9.8%) had \geq 20 mIU/l. It was also found that 73.4% of the neonates had TSH levels of more than 5 mIU/l, thus indicating prevalence of ID in a population studied. No statistically significant difference of gender on the TSH levels was observed.

Discussion

Neonates are the most vulnerable group for ID. Raised TSH in neonates is an indicator for ID. WHO (2007) reported that a <3% frequency of TSH concentrations above 5 mIU/l in samples collected 3–4 days after birth indicates iodine sufficiency in a population [2]. In the present study 73.4% of the neonates have TSH levels of more than 5 mIU/l, thus indicating widespread presence of ID. This could be possibly due to higher percentage of families consuming salt with iodine content of less than 15 ppm [4,5]. Moreover, ID in soil leads to crops deficient in iodine and consumption of food deficient in iodine leads to poor iodine status in the population. Low birth weight could also be a reason for high TSH level amongst 8.4% of the neonates.

Many studies have attempted to apply the frequency of neonatal TSH values of more than 5 mIU/l in determining population iodine status and monitoring intervention programmes, and although some have proven to be successful, most have provided conflicting or uncertain data [10–17]. This is due to many technical issues that

remain unresolved on the use of neonatal TSH screening for monitoring iodine status, making it doubtful as a sensitive and reliable quantitative tool [18].

Neonatal TSH should not be used as the sole indicator for monitoring ID control programmes, especially when samples are collected less than 48 hours after birth, as there are no established reference intervals. And the cord blood sample taken immediately after delivery could have false positive high values of TSH due to physiological neonatal TSH surge that elevates TSH level [19].

Studies have documented that ID during the initial phase of life affects the developing brain. Thyroid stimulating hormone affects proper development of the central nervous system [2]. Earlier studies conducted in different countries of the world reported the high TSH level of more than 5 mIU/l in; Western Uganda (20–32%), Estonia (18.1%), Italy (14.4%), Spain (9.08%), Thailand (8.9%), Australia (6.8%), Poland (3–5%) and Iran (3.6%), respectively [10–17].

A study conducted in West Bengal reported TSH levels of more than 5 mIU/l in 2.9% of the neonates [9]. An Indian study conducted amongst 1200 neonates reported the high TSH level of more than 20 mIU/l in 1.83% of the neonates [20]. The prevalence of IDD found in neonates in the current study was much higher than that found in urban metropolitan hospitals.

Our earlier studies conducted in Kangra district amongst SAC [3–5] have also documented that ID was a public health problem. Findings of our present study substantiated that there is a high prevalence of ID amongst neonates in Kangra district. There is a need to review the implementation of the iodized salt programme in Kangra district. Furthermore, there is an urgent need for a neonatal screening programme for early detection of children with ID.

Conflict of interest

The authors declare no conflict of interest.

Acknowledgment

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References

- [1] Hetzel BS, Potter BJ, Dulberg EN. The iodine deficiency disorders; nature, pathogenesis and epidemiology. In: Bourne GH, editor. Aspects of some vitamins, mineral and enzymes in health and disease. World review of nutrition and dietetics, vol. 6. 1990. p. 59–119.
- [2] Assessment of iodine deficiency disorders and monitoring their elimination. A guide for programme Managers. WHO/UNICEF/ICCIDD. Geneva: World Health Organization; 2007.
- [3] Kapil U, Sohal KS, Sharma TD, Tandon M, Pathak P. Assessment of iodine deficiency disorders using the 30 clusters approach in District Kangra, Himachal Pradesh, India. J Tropical Pediatr 2000;46:264–6.
- [4] Kapil U, Sharma TD, Singh P. Iodine status and goiter prevalence after 40 years of salt iodization in the Kangra district, India. Indian J Pediatr 2007;74:135–7.
- [5] Kapil U, Pandey RM, Kabra M, Jain V, Sareen N, Bhadoria AS, et al. Status of iodine deficiency in district Kangra, Himachal Pradesh after 60 years of salt iodization. Eur J Clin Nutr 2013;67:827–8.
- [6] Slazyk WE, Hannon WH. In: Therrell BL, editor. Laboratory methods for Neonatal Screening. Washington, DC: American Public Health Association; 1993. p. 23.
- [7] Westgard JO, Klee GG. In: Burtis CA, Ashwood R, editors. Tietz textbook of clinical chemistry. 3rd ed. Philadelphia, PA: W.B. Saunders Co.; 1999. p. 384–8.
- [8] Spencer CA, Takeuchi M, Kazarosyan M, MacKenzie F, Beckett GJ, Wilkinson E. Inter laboratory/intermethod differences in functional sensitivity of immuno-metric assays of thyrotropin (TSH) and impact on reliability of measurement of subnormal concentrations of TSH. Clin Chem 1995;41(3):367–74.
- [9] Chakraborty I, Chatterjee S, Bhadra D, Mukhopadhyaya BB, Dasgupta A, Purkait B. Iodine deficiency disorders among the pregnant women in a rural hospital of West Bengal. Indian J Med Res 2006;123:825–9.

- [10] Ehrenkranz J, Fualal J, Ndizihiwe A, Clarke I, Alder S. Neonatal age and point of care TSH testing in the monitoring of iodine deficiency disorders: findings from western Uganda. *Thyroid* 2011;21:183–8.
- [11] Mikelsaar RV, Viikmaa M. Neonatal thyroid stimulating hormone screening as an indirect method for the assessment of iodine deficiency in Estonia. *Horm Res* 1999;52:284–6.
- [12] Costante G, Grasso L, Ludovico O, Marasco MF, Nocera M, Schifino E, et al. The statistical analysis of neonatal TSH results from congenital hypothyroidism screening programs provides a useful tool for the characterization of moderate iodine deficiency regions. *J Endocrinol Invest* 1997;20:251–6.
- [13] Peris Roig B, Calvo Rigual F, Tenias Burillo JM, Merchante Alfaro A, Presencia Rubio G, Miralles Dolz F. Iodine deficiency and pregnancy. Current situation. *Endocrinol Nutr* 2009;56:9–12.
- [14] Jaruratanasirikul S, Sangsupawanich P, Koranantakul O, Chanvitan P, Ruaengrairatanaroj P, Sriplung H, et al. Maternal iodine status and neonatal thyroid-stimulating hormone concentration: a community survey in Songkhla, southern Thailand. *Public Health Nutr* 2009;12:2279–84.
- [15] Rahman A, Savige GS, Deacon NJ, Francis I, Chesters JE. Increased iodine deficiency in Victoria, Australia: analysis of neonatal thyroid-stimulating hormone data, 2001 to 2006. *Med J Aust* 2010;193:503–5.
- [16] Oltarzewski M, Szyborski J. Neonatal hypothyroid screening in monitoring of iodine deficiency and iodine supplementation in Poland. *J Endocrinol Invest* 2003;26:27–31.
- [17] Najafi M, Khodaei GH, Bahari M, Sabahi M, Farsi MM, Kiani F. Neonatal thyroid screening in a mild iodine deficiency endemic area in Iran. *Indian J Med Sci* 2008;62:113–6.
- [18] Li M, Eastman CJ. Neonatal TSH screening: is it a sensitive and reliable tool for monitoring iodine status in populations? *Best Prac Res Clin Endocrinol Metab* 2010;24(1):63–75.
- [19] Büyükgöbüz A. Newborn screening for congenital hypothyroidism. *J Clin Res Pediatr Endocrinol* 2013;5:8–12.
- [20] Manglik AK, Chatterjee N, Ghosh G. Umbilical cord blood TSH levels in term neonates: a screening tool for congenital hypothyroidism. *Indian Pediatr* 2005;42:1029–32.