# University of Glasgow

# **Transient TSH elevation in infants referred** on newborn screening – features, prevalence and trends

<sup>1</sup>Yasmine Ouarezki, <sup>2</sup>Jeremy Jones, <sup>3</sup>Moira Fitch, <sup>2</sup>M Guftar Shaikh, <sup>4</sup>Malcolm Donaldson <sup>1</sup>Etablissement Public Hospitalier Hassen Badi, El-Harrach, Algiers, Algeria; <sup>2</sup>Royal Hospital for Children, Govan Road, Glasgow, United Kingdom<sup>3</sup> Newborn Screening Laboratory, West of Scotland Genetic Services, Queen Elizabeth University Hospital, Glasgow, United Kingdom, <sup>4</sup> Section of Child Health, Glasgow University School of Medicine, Glasgow, United Kingdom

# Background

#### Up to 20% of infants referred on newborn congenital hypothyroidism (CH) screening are subsequently shown to have transient TSH elevation

#### FACULTE DE MEDECINE D'ALGER



rather than permanent CH. Accurate identification of such cases is important to avoid prolonged treatment with thyroxine and unnecessary clinic attendance.

Objective	Method
To determine the prevalence, trends and profile of infants with transient TSH elevation referred between August 1979 and December 2015 by the Scottish Newborn Screening Programme.	Analysis of infants referred during the study period with initial/repeat capillary TSH ≥50/≥25 mU/L (1979-82); ≥40/≥15 (1982-89); ≥40/≥10 (1989-2002); ≥25/≥8 (2002-15) in whom venous thyroid function tests subsequently became normal off thyroxine. Details of gestation, birthweight (BW), "sickness" and extra- thyroidal congenital malformations (CM) were recorded.



Of 2,202,191 newborns screened, 936 were referred by the screening laboratory during the study period including 630 (68.9%) with definite CH; 208 (22.8%) with transient TSH elevation and 58(6.35%) patients with status uncertain.

Groups	Definite CH N=630	Transient TSH elevation N=208
Birth weigh(kg) Mean	3.3 0.7-5.5	2.7 0.7-4.6
Gestational age (weeks) Mean Range	39.6 27-43	36.5 24-42
GA< 30 weeks(%) Sickness(%) Congenital malformations(%)	0.5 7.1 5.7	12.9 35.6 20.7
Capillary TSH(mU/l) median range Venous TSH(mU/l) median	167.5 102	37
FT4(pmol/l) median	6.6	15.15
range Second TSH sample(%)	12	50

<b>Etiologies of Transient TSH elevation</b>	N=208
Blocking maternal antibodies	3
Maternal carbimazole	1
Pendrin mutation	1
TSH receptor heterozygoty	1
Down syndrome	12

Table 2: Specific aetiologies of transient TSH elevation.

The incidence of transient TSH elevation was 6.6 and 5.1/yr between 1982-2004 and 2005-15. Of 43 transient cases with CM, 19 involved the digestive system/abdominal wall of which 15 were born  $\leq$  2004 when iodine antisepsis was largely discontinued in Scottish newborn units.

#### Second ISH sample(%)

Table 1: characteristics of CH and transient TSH elevation groups.

## Conclusion

Infants with low BW, extreme prematurity, sickness, additional malformations, Down syndrome and modest capillary/venous TSH elevation are particularly likely to have transient thyroid dysfunction especially if diagnostic imaging shows a eutopic thyroid gland. Such infants require careful re-evaluation at ≥ 3 years of age. Trends in transient TSH elevation will be influenced by capillary TSH cut-offs being altered, reduced iodine antisepsis usage, and the currently unknown dietary iodine status in Scottish mothers.

### REFERENCES

- A Foo, H Leslie, D J Carson. Confirming congenital hypothyroidism identified from neonatal screening. The Ulster Med J 2002, 71:38-41.
- V. V. Thaker et al. Iodine-Induced Hypothyroidism in Full-term Infants With Congenital Heart Disease: More Common Than Currently Appreciated? JCEM 2014, 99: 3521–26
- Rosanna Rovelli et al. Newborn of mothers affected by autoimmune thyroiditis: the importance of thyroid function monitoring in the first months of life . Italian J Ped 2010, 36:24

