A novel ESR2 frameshift mutation predisposes to medullary thyroid carcinoma and causes inappropriate RET expression

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Meeting abstracts from the 64th British Thyroid Association Annual Meeting

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S1 Clinical Features and Evidence-Based Management of Graves’ Orbitopathy
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Thyroid Research 2017, 10(Suppl 1):S1

Graves’ orbitopathy (GO) is the main extrathyroidal manifestation of Graves’ disease. When fully expressed, it is characterized by inflammatory soft tissue changes, exophthalmos, ocular dysmotility causing diplopia, and, rarely, sight-threatening dysthyroid optic neuropathy (DON). The prevalence of GO among Graves’ patients seems lately declining, probably due to early diagnosis, early intervention on risk factors associated with its occurrence or progression (smoking, uncontrolled thyroid dysfunction), early correction of hyper and hypothyroidism. Only about 25–30% of newly diagnosed Graves’ hyperthyroids are affected with GO, which is usually mild and rarely progressive. Assessment of activity and severity of GO according to standardized criteria is fundamental to plan management. The European Thyroid Association and the European Group on Graves’ Orbitopathy (EUGOGO) have recently published the first guideline on management of GO. Mild GO usually requires only a watchful strategy, in addition to local measures (eye drops, ointments) and removal of risk factors. Intravenous glucocorticoids (ivGCs) are the first-line treatment for moderate-to-severe and active GO, as demonstrated by randomized clinical trials. When ivGCs fail or GO recurs after treatment withdrawal, options include a second course of ivGCs, oral GCs combined with orbital radiotherapy or cyclosporine, rituximab. Evidence that the any of the above treatment be effective in the context of a poor response to a first course of ivGCs is limited and should be investigated in larger studies. In addition to rituximab, ongoing investigations are exploring the role of other biologics targeting, e.g., the IGFR-1 receptor or the IL-6 receptor, and results will probably available in 1–2 years. When GO has been treated medically and is inactive, rehabilitative surgery (orbital decompression, squint surgery, eyelid surgery) is often needed.

S2 Role of T3 and TRH in the hypothalamic regulation of energy metabolism
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Thyroid Research 2017, 10(Suppl 1):S2

The relation between thyrotoxicosis, the clinical syndrome resulting from exposure to excessive thyroid hormone, and the autonomic nervous system remains enigmatic. Recent experiments from our lab and others have shown that T3 may act within several hypothalamic nuclei to modulate hepatic glucose metabolism and brown adipose tissue (BAT) activity. These effects are mediated via pre-autonomic neurons connecting the paraventricular and ventromedial nuclei (PVN and VMH) with these peripheral organs via neural pathways. Intrahypothalamic effects of T3 on glucose metabolism in the liver could be modulated by selective hepatic sympathetic and parasympathetic denervation. Thyroid hormone appeared to stimulate hepatic glucose production via a sympathetic pathway, representing a novel central route for thyroid hormone action. As most of the experiments were performed in a rather acute setting we recently developed a model for chronic intrahypothalamic T3 administration in rats, allowing for the assessment of chronic metabolic effects of intrahypothalamic T3 administration. In addition to T3, we now investigate the role of intrahypothalamic TRH in energy metabolism. Further elucidation of intrahypothalamic effects of thyroid hormones on autonomic outflow to metabolic organs will add to our understanding of the metabolic effects of hyperthyroidism.

S3 Triiodothyronine, Deiodinase 3 and Stem Cell-derived Cardiomyocyte Maturation
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Thyroid Research 2017, 10(Suppl 1):S3

Functional cardiomyocytes can be derived from both human embryonic (hESCs) and induced pluripotent (hiPSCs) stem cells. These pluripotent stem cell-derived cardiomyocytes (hPSC-CM) hold great potential for regenerative medicine and in vitro screening. However, the phenotype is immature and more closely resembles that of foetal/neonatal cardiomyocytes. In the adult heart thyroid hormone has pleiotropic effects on contractility and energy metabolism. The active form of thyroid hormone, triiodothyronine (T3), is known to play a role during foetal cardiomyocyte maturation and has recently been shown to promote cardiac differentiation and enhance hPSC-CM maturation in vitro. During development the enzyme, deiodinase 3 (D3), protects the foetus against maternal thyroid hormone by metabolism of T3 into inactive products. Down regulation of D3 at appropriate times during development allows physiological growth and maturation of cardiomyocytes in response to thyroid hormone.
We show that stimulation of the thyroid hormone pathway can enhance the maturation of hPSC-CM and that further investigation into the contribution of individual components of this pathway is warranted.

O1 Controlled Antenatal Thyroid Study: Obstetric Outcomes
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Thyroid Research 2017, 10(Suppl 1):O1

Background
Suboptimal thyroid function in pregnancy is associated with adverse obstetric outcomes but it is unclear whether levothyroxine treatment, initiated during pregnancy is beneficial. We investigated whether correction of abnormal thyroid function during pregnancy is associated with improved obstetric outcomes.

Methods
Retrospective analysis of the Controlled Antenatal Thyroid Screening (CATS) study with obstetric outcomes obtained through data-linkage in the Secure Anonymised Information Linkage (SAIL) databank. Setting: Welsh participants from CATS. Participants: 13,506 pregnant women; 12,874 women had normal thyroid function, 320 had subclinical hypothyroidism (SCH), 281 had isolated hypothyroxinemia (IH) and 31 had overt hypothyroidism. Main Outcome Measures: Odds of stillbirths (fetal demise after 24 weeks gestation), Caesarean section (SD 639) vs. 3558 g (SD 532) p = 0.004. Untreated women with SCH and IH had an increased risk of stillbirth compared to women with normal thyroid function OR = 5.73 (95%CI 1.74, 18.9) p = 0.003. Caesarean sections, prematurity and abnormal birth-weight by thyroid and treatment status.

Results
Untreated women with SCH had increased odds of stillbirth compared to women with normal thyroid function OR = 5.73 (95%CI 1.74, 18.9) p = 0.003. No stillbirths occurred in women receiving levothyroxine. In analysis of women with IH, untreated women had an increased risk of early (<37 weeks) caesarean section than those who received levothyroxine (6% vs 0%) p = 0.006. Untreated women with IH also had earlier mean gestational age 38.8 (SD 2.34) weeks vs. 39.7 (SD 1.94) weeks p = 0.002 and lower mean birth-weight 3353 g (SD 639) vs. 3558 g (SD 532) p = 0.004.

Conclusion
Both SCH and IH are associated with adverse obstetric outcomes. In IH levothyroxine treatment is associated with favourable effects on gestational age at delivery and early Caesarean section rates. Levothyroxine may also have some protective impact on stillbirths in SCH.

O2 Pharmacological enhancement of radioiodine uptake using Src kinase inhibitors
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Thyroid Research 2017, 10(Suppl 1):O2

Background
Radioiodine imaging, cell ablation and treatment of metastases in thyroid cancer rely on the presence of a functional sodium iodide symporter (NIS) in the basolateral plasma membrane (PM) of thyrocytes. Augmenting NIS PM localisation represents an important therapeutic strategy for increasing radioiodine delivery. We previously described a mechanism by which NIS is internalised by pituitary tumor-transforming gene-binding factor (PBF) in thyroid cells, significantly reducing radioiodine uptake. Importantly, we demonstrated that PBF phosphorylation at Y174 by Src kinase mediated NIS repression, which could be rescued by the Src family kinase (SKF) inhibitor PP1.

Methods
We have now replicated these findings in breast cancer cells, further elucidated the mechanism of repression and identified a more potent inhibitor of PBF-pY174.

Results
In MCF-7 and MDA-MB-231 breast cancer cells with either exogenous or all-trans retinoic acid/dexamethasone-induced NIS expression, PBF significantly repressed radioiodine uptake and this was reversible with PP1 treatment. PBF-Y174 is also a critical part of an endocytosis motif and mutation results in PM accumulation. Mutation of a predicted Src consensus sequence (EEN170-172AAA) abrogated pY174 and radioiodine uptake repression, confirming Src-dependent Y174 phosphorylation and demonstrating that NIS repression is mediated by phospho-PBF and not PM-bound PBF. Treatment with the SKF inhibitor dasatinib potently inhibited PBF-pY174 and restored radioiodine uptake. In the presence of dasatinib-resistant Src (T341I), dasatinib no longer rescued PBF repression of NIS, indicating that Src and no other SKF mediates PBF phosphorylation.

Conclusions
Taken together, these data suggest that Src inhibition can effectively enhance radioiodine uptake in multiple tumour types, with implications for improving outcomes in thyroid cancer and utilising NIS for the treatment of other tumours.

O3 Comparative analysis of human and mouse expression data identifies proto-oncogene PTTG and PBF-associated genes in thyroid cancer
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Thyroid Research 2017, 10(Suppl 1):O3

Background
Whilst the proto-oncogene PTTG and its binding partner PBF have been shown to be up-regulated in differentiated thyroid cancer, there is a paucity of information regarding their co-expression and specific roles in tumour progression. In particular, PTTG and PBF have both been reported to modulate the tumour suppressor p53, whose activity is impaired in most human cancers. Therefore, the role of PTTG and PBF in thyroid tumorigenesis may involve disruption of p53 pathways that are central to DNA-damage repair (DDR), cell growth and apoptosis.

Methods
In the present study we investigated the association of PTTG and PBF with p53-related genes in the TCGA thyroid cancer dataset, as well as in a bi-transgenic murine model (Bi-Tg) overexpressing PTTG and PBF specifically in the thyroid gland.

Results
Characterisation of primary murine Bi-Tg thyrocytes revealed that co-expression of PTTG and PBF caused extensive repression of DDR genes (39/82 genes; P < 0.05). Of these, 31 genes were down-regulated >1.5-fold, including genes with key roles in maintaining genomic integrity such as Brca1. Irradiation exposure to increase intracellular p53 further showed significant differences in overall DDR gene expression (n = 82 genes) between irradiated Bi-Tg and wild-
type thyrocytes ($P = 2.4 \times 10^{-5}$) that was greater than either PBF-Tg ($P = 1.5 \times 10^{-3}$) or PTGT-Tg thyrocytes ($P = NS$). By comparison in the TCGA dataset, there were striking correlations with PTGT and PBF in well-characterised p33-related gene panels ($P < 0.05$; 82–96 genes per panel; $n = 322$ unmatched TCGA tumour samples). Importantly, nearly half of the significant DDR gene alterations in Bi-Tg thyrocytes were also present in TCGA comparing samples with either low or high PTGT/PBF mRNA levels. Furthermore, the overall survival ($P = 0.0002$) and disease-free survival ($P = 0.02$) was poorer for TCGA individuals with high tumoural PTGT/PBF expression ($n = 20$) than for all other patients ($n = 235$).

**Conclusions**

Altogether our findings provide important insights into the association of p33-related genes with PTGT and PBF in thyroid tumourigenesis.

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**O4**

**Graves’ disease with fluctuating thyroid status and hypothyroidism with positive anti-TSH receptor antibody levels - distinctive autoimmune side-effects following alemtuzumab therapy for multiple sclerosis**

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*Thyroid Research 2017, 10(Suppl 1):O4*

**Background**

Alemtuzumab, a highly effective, newly-licensed, treatment for multiple sclerosis (MS), is associated with autoimmune side-effects - notably Graves’ disease (GD) with a reportedly indolent course. Aim To determine the type, frequency and course of thyroid dysfunction (TD) in a cohort of alemtuzumab-treated MS patients in Cambridge & Cardiff.

**Methods**

Case records of alemtuzumab-treated patients who developed TD were reviewed.

**Results**

Overall, 40% (102/249; 81 F, 21 M) of alemtuzumab-treated patients were reviewed. To determine if TD were reviewed.

**Conclusions**

Altogether our findings provide important insights into the association of p33-related genes with PTGT and PBF in thyroid tumourigenesis.
large cohort of inpatients evaluating their hospitalisation frequency, comorbidities and all-cause mortality.

Methods
A nested case-control study was conducted using a cohort admitted to a large tertiary centre (2007–2012). 671 hyperthyroid subjects were identified, matched (1:4) by age, gender and year of admission with hypo- and euthyroid inpatients and followed until 31/12/2015.

Results
A total of 31,400 person-years were analysed in 5,979 inpatients. There were 2,175 (36.4%) deaths. Hyperthyroidism was associated with a significant increase in all-cause mortality compared to hypo- and euthyroid inpatients (HR = 1.2 [95% CI:1.02–1.33], P = 0.03 and 1.3 [1.10–1.44], P = 0.001). Additionally, hyperthyroid inpatients were more frequently re-hospitalised (4.0 [3.5–4.5]) compared with controls in both groups (AORHyp = 2.1, P = 0.003; AOREu = 4.8, P < 0.001). They were more frequently admitted for circulatory conditions (CVD) (Nhyp = 230, 34.3%; Neu = 711, 27.1%, AORh = 1.5 (1.2–1.8), P < 0.001; Nh = 601, 22.4%, AORh = 1.9 (1.6–2.3), P < 0.001) while respiratory admissions were more common in hyperthyroid (N = 119, 17.7%) when compared with euthyroid (N = 357, 13.3%, AOR = 1.4 (1.1–1.8), P = 0.003) but not different from hypothyroid subjects; proportions of patients admitted for nervous and digestive causes were not significantly different. When considering recorded comorbidities, CVD was more frequent in hyperthyroid inpatients (Nhyp = 457, 68.1%) than matched controls (Nh = 1,679, 64.0%, AORh = 1.3 (1.1–1.6), P = 0.007; Nh = 1,442, 53.7%, AORh = 2.2 (1.8–2.8), P < 0.001) and presented more frequently with atrial fibrillation (Nhyp = 207, 30.8%; Nh = 441, 16.8%, AORh = 2.6 (2.1–3.2), P < 0.001; Nh = 339, 12.6%, AORh = 3.6 (2.9–4.5), P < 0.001) or heart failure (Nhyp = 105, 15.6%; AORh = 1.5 (1.2–1.9), P = 0.002; AORh = 2.6 (2.0–3.4), P < 0.001).

Conclusion
We conclude that hyperthyroidism in hospitalised patients is associated with additional health and economic burdens, significantly increasing likelihood of re-hospitalisation, cardiovascular morbidity and all-cause mortality.

P2 Controlled Antenatal Thyroid Screening (CATS) Study II: (ii) Effect of treatment of suboptimal gestation thyroid function (SGTF) on children’s cognition at age 9
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Background
The Controlled Antenatal Thyroid Screening (CATS) study was the first randomized controlled trial to investigate the effect of antenatal thyroid screening and maternal levothyroxine treatment on childhood cognition, evaluated at 3 years of age. Although CATS did not show treatment benefits, the childhood assessment may have been undertaken too early for differences to be apparent.

Methods
In the present study cognition was re-assessed in offspring of the Welsh CATS cohort aged 9 years using the WISC-IV (IQ). Groups comprised children of mothers with normal gestational thyroid function (GTf, n = 233) and those of mothers with treated (n = 118) and untreated (n = 101) SGTF, i.e. TSH in the highest 2.5% and/or fT4 in the lowest 2.5%. Analysis included a regression to explore the odds of IQ < 85 (1SD below the mean). A secondary analysis explored children’s Thr92Ala polymorphism and this was compared to maternal thyroid status.

Results
There were no difference in proportion of children with IQ < 85 between children of women with SGTF (10%) and normal GTf (6.4%) (p = 0.59) or between treated (8.5%) and untreated (11.9%) SGTF results (p = 0.29). Children homozygous for Thr92Ala and born to mothers with low maternal FT4 levels, indicated a trend for IQ < 85 compared to the normal GTf group: OR = 2.90 (95%CI 0.49,17.30) p = 0.24. Removing the treated group increased the odds of IQ < 85: OR = 13.7 (95%CI 1.63,115) p = 0.01.

Conclusion
Our results do not support a beneficial effect of antenatal thyroid screening on child cognition. One in seven of the population have the Ala92Ala genotype and are more at risk of adverse effects of SGTF. Further research is needed to clarify the impact of the Thr92Ala polymorphism and its interaction with maternal thyroid function on child cognition.

P3 Genetic variants associated with hypothyroidism and serum thyroid stimulating hormone levels (TSH) in Tayside (Scotland): a GoDARTS study
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Thyroid Research 2017, 10(Suppl 1):P3

Background
Hypothyroidism is the most common thyroid disorder. Genome-wide association studies (GWAS) have identified variants in FOXE1 that are associated with hypothyroidism, and PDE8B and CAPZB that are associated with TSH levels. However, the number of studies is limited and replication studies are desirable.

Methods
Patients were identified from a study of the Genetics of Diabetes Audit and Research Tayside (GoDARTS) recruited in Tayside, Scotland, between January 1996 and March 2014. Electronic Medical records (EMR)-derived phenotypes were ascertained and single-locus tests of association were performed via logistic regression under the assumption of an additive genetic model. Serum TSH levels were identified in 16,769 patients of white ethnicity, 2,664 with hypothyroidism, and PDE8B and CAPZB that are associated with hypothyroidism, and PDE8B and CAPZB that are associated with serum TSH levels, respectively.

Results
A FOXE1 variant (rs925489) was associated with hypothyroidism (OR = 0.77 [95%CI 0.71–0.83], P = 9.9 × 10-10). We found evidence that other genetic variants previously associated with hypothyroidism were also associated in this study. We found association with PTPN22 (rs4915077, P = 3.0 × 10-03), and HLA (rs2517532, P = 7.5 × 10-04), SH2B3 (rs3184504, P = 9.4 × 10-05), VAV3 (rs6679677, P = 3.0 × 10-04), SH2B3 (rs3184504, P = 9.4 × 10-05), and HLA (rs2517532, P = 7.5 × 10-04). The analysis found the association of PDE8B (rs4704397, P = 1.0 × 10-22), rs6885099, P = 1.1 × 10-32, and CAPZB (rs10798924; P = 8.2 × 10-21) with serum TSH levels in euthyroid individuals. These genetic variants explained 1.67, 1.57, and 0.82% of the variation of mean TSH levels respectively.

Conclusions
Our results replicated the previously reported association of FOXE1 with hypothyroidism, and PDE8B and CAPZB with serum TSH levels,
and emphasize the need for additional genetic studies in more diverse population.

P4
A novel ESR2 frameshift mutation predisposes to medullary thyroid carcinoma and causes inappropriate RET expression

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P5
Use of TSH-Receptor Antibodies (TRAb) in the assessment of new onset thyrotoxicosis within the North East of England

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Background
Causes of hyperthyroidism include Graves' disease (GD), toxic multinodular goitre and thyroiditis. British Thyroid Association guidelines recommend specialist referral once hyperthyroidism is diagnosed. TSH-receptor antibodies (TRAb) are more specific to GD than thyroid peroxidase antibodies (TPOAb). Regional guidelines recommend measuring TRAB in thyrotoxic patients. We aimed to assess TRAB use regionally in newly diagnosed thyrotoxicosis.

Method
We conducted a retrospective case review of patients with newly detected thyrotoxicosis from 1st March-31st August 2015 from 4 endocrinology centres. TSH, fT4, fT3, TPOAb and TRAb values; the requester and date requested were recorded. Any thyroid uptake scans done; and final diagnoses were noted.

Results
We analysed 209 records – 79% females (n = 166), 21% males (n = 43); the average age was 50.5 (17–96), 88.6% had TRAB requested. 76.2% requests were from endocrinologists, 15.7% from GPs, 5.9% from Biochemistry, and 2.2% from other physicians. 56.2% had positive TRABs. The commonest diagnosis was GD (55.0%), followed by multinodular goitre (10.5%) and thyroiditis (8.1%). GD was diagnosed using TRAb in 81.7%; and using thyroid uptake scans in 6% (n = 7). 2 patients had clinical features; in 12 patients, the reason for favouring a GD diagnosis was unclear. 19 patients (9%) were not referred to endocrinology. Average time between detection of thyrotoxicosis and TRAB request was 40 days (32–57 days). Time to TRAB request was shorter (average 9 days) if requested prior to referral to an endocrinologist. This lag was because most TRAB requests were only requested after referrals were received and reviewed by endocrinologists.

Conclusions
TRAB is commonly used regionally with good sensitivity and specificity. We propose that it should be added on by biochemistry labs in all newly thyrotoxic patients, as this could expedite diagnosis, minimizing use of inappropriate antithyroid medications. Availability of results before endocrinology consultations would also facilitate prompt treatment and better communication with patients.
Results
Fourteen patients (two male, twelve female) with a mean age of 41 years (16–74) and weight 89 kg (53–124) underwent the test. Cause of hypothyroidism was autoimmune in 8, total thyroidectomy in 5 and congenital hypothyroidism in 1. Before testing, mean daily levothyroxine dose was 209 μg and 14 (100–300). In 9 there was adequate absorption of levothyroxine and TSH normalised in 4 weeks confirming poor compliance. In one patient TSH normalised after receiving the higher dose while two patients required higher amount in divided dose to normalise TSH. One patient on continuous ambulatory peritoneal dialysis had inadequate FT4 response and failed to normalise TSH at 8 weeks. TSH however normalised following renal transplantation. Malabsorption due to total pancreatectomy was the cause of TSH non-suppression in one. TFT results available on 10 patients, 6 months or more after the test showed that 70% of the patients were maintaining TSH within normal range.

Conclusion
Thyroxine absorption test was useful in differentiating poor compliance from other causes of persistently raised TSH. Some patients who fail to suppress TSH after 4 weeks may require a higher dose of levothyroxine administered once or twice weekly. Splitting dose as twice weekly may be useful as in some individuals as t1/2 may be shorter. Offering continuation as once or twice weekly dose may help improve compliance over long term.

P7 Diagnosis and workup of thyrotoxicosis – a secondary care audit
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Thyroid Research 2017, 10(Suppl 1):P7

Background
Identification of the cause of thyrotoxicosis is important as this translates to different treatments and prognosis. Using the current NICE clinical knowledge summary (CKS), BTA guidelines for the use of thyroid function tests, and a recent clinical review by B Vaidya et al (2014), audit standards for appropriate diagnosis of thyrotoxicosis were established.

Methods
All new patients at the hospital thyroid clinic in January 2015 were screened. Data was gathered from referral letters, clinic notes, and hospital information system. 37 patients were eligible.

Results
Treatment was initiated by the GP at the time of diagnosis in 13 of the 37 (35.1%). 19 patients had thyroid antibodies checked (51.35%) however only 1 of the 19 had TRAbs included (5.55%). At initial endocrine OPA 32 of 36 had their TRAbs checked (88.9%) and 3 of 35 (8.5%) had an RIUS or USS ordered.

Conclusions
51% of GPs ordered thyroid antibodies, but only 1 patient had TRAbs requested, suggesting that GPs were not aware of the benefit of measuring TRAbs (as the diagnostic hallmark of Graves) or did not have access to this test. The NICE CKS does not ask the primary care physician to measure TRAbs, although most clinical experts suggest its measurement on diagnosis of thyrotoxicosis. As TRAbs was usually only ordered at the first outpatient appointment, only 8% of patients had RIUS/USS ordered at their first hospital appointment. 35% of patients had carbimazole initiated by GPs. This has implications for diagnosis and long-term management, especially in the absence of appropriate antibody testing and mean time from referral to first appointment of 35 days. A proposed change in our pathway is the automatic addition of TRAbs testing in all newly detected cases of thyrotoxicosis when noted by our laboratory.
**Results**
Participants had a mean (SD) age 31(5) years, median (IQR) BMI 24.4(22.0, 28.3) kg/m², 41% were primiparous, 10% smoked during pregnancy, 34% took iodine (>140 μg/pill) containing vitamins. 96% were Caucasian. Median (IQR) UIC was 88.0 (54.1, 157.5) μg/l, which is consistent with iodine deficiency by WHO criteria. A total of 224/308 (73%) of women had UIC values <150 μg/l. Increasing milk intake was associated with higher UIC (p = 0.003). There was no difference in median (IQR) UIC between those women who took iodine containing vitamins (n = 106) and those who did not (n = 202): 88.5(54.5, 170.5) vs 88.0 (53.8, 150.0) μg/l, p = 0.6. There was no correlation between median UIC and TSH (p = 0.6) or FT4 (p = 0.1).

**Conclusion**
Iodine deficiency in pregnancy is common in South-west England. Further research is needed to develop optimum prevention and treatment strategies.