TOX/2021/14

COMMITTEE ON TOXICITY OF CHEMICALS IN FOOD, CONSUMER PRODUCTS AND THE ENVIRONMENT

Second draft statement on the potential effects that excess iodine intake may have during preconception, pregnancy and lactation.

Background

1. The Scientific Advisory Committee on Nutrition (SACN) last considered maternal diet and nutrition in relation to offspring health in its reports on 'The influence of maternal, fetal and child nutrition on the development of chronic disease in later life' (SACN, 2011) and on 'Feeding in the first year of life' (SACN, 2018). In the latter report, the impact of breastfeeding on maternal health was also considered. In 2019, SACN agreed to conduct a risk assessment on nutrition and maternal health focusing on maternal outcomes during pregnancy, childbirth and up to 24 months after delivery; this would include the effects of chemical contaminants and excess nutrients in the diet.

2. SACN agreed that, where appropriate, other expert Committees would be consulted and asked to complete risk assessments e.g. in the area of safety advice. A provisional list of chemicals was proposed by SACN and updated and amended following discussions by COT who will be guiding the toxicological risk assessment process.

3. Iodine is an essential micronutrient in the human diet, required for the production of thyroid hormones including thyroxine. These hormones are necessary for cell metabolism, growth and development at all stages of life. The most visible manifestation of iodine deficiency is goitre - an enlargement of the thyroid gland in the neck but there is concern, that in the fetus, infant, and young child, more modest changes may impair psychomotor development in the absence of overt thyroid enlargement in the mother (SACN, 2014). Excess iodine may lead to compensatory thyroid hypertrophy.

4. The National Diet and Nutrition Survey (NDNS) has collected spot urine samples for iodine analysis since 2013. The findings show that urinary iodine concentrations in children and adults in the standard NDNS age groups meet the World Health Organisation (WHO) criteria that a population with no iodine deficiency should have median urinary iodine concentrations of between 100 μ g/L and 199 μ g/L and fewer than 20% of the population below 50 μ g/L. The median urinary iodine for women of childbearing age (16 to 49 years) was 98 μ g/L with 21% of the population below 50 μ g/L in this group. They also fail to meet the stricter criteria for pregnant and lactating women which is a median urinary iodine concentration of 150 - 249 μ g/L (WHO, 2007; SACN, 2014). Time trend analysis on urinary iodine in the 9-11 report show that there is no statistically significant changes over time for any age/sex groups (SACN, 2014).

5. The COT was asked to consider whether exposure to excess intake of iodine would pose a risk to maternal health in discussion paper (TOX/2020/54)¹ and in a draft statement (TOX/2020/61). The Committee noted that further information on thyroid hormones in the body should be provided. It was also suggested that reasoning based on pathological or physiological considerations should be included to better explain why some individuals, other than those with underlying thyroid disorders, were particularly susceptible to the effects of excess iodine. Members suggested that it should also be noted that sensitive individuals were unable to escape the Wolff-Chaikoff effect, where excess iodine down-regulates thyroid hormone. Members noted that more information was needed on the fact that any iodisation fortification was voluntary in the UK. Members also agreed that further information on whether iodophor disinfectants were still authorised in the UK was needed.

5. The Committee are asked to consider the draft statement which includes further information on other iodine containing components in the body, mandatory iodisation schemes in the UK and whether iodophor disinfectants are still authorised in the UK², presented in Annex A. The revisions are given in tracked changes.

Questions for the Committee

Members are asked to consider the following questions:

- a) Do members have any comments on the structure and content of the statement?
- b) Does the Committee have any further comments?

¹ TOX/2020/54 is available on the <u>COT website</u>

² COT Final Minutes December 2020 is available on the <u>COT website</u>

This is a draft statement for discussion. It does not reflect the final views of the Committee and should not be cited.

Secretariat

March 2021

ANNEX A to TOX/2021/14

COMMITTEE ON TOXICITY OF CHEMICALS IN FOOD, CONSUMER PRODUCTS AND THE ENVIRONMENT

Second draft statement on the potential effects that excess iodine intake may have during preconception, pregnancy and lactation.

Background

1. In the environment, iodine is usually found in the form of iodate salts or organo-iodide compounds synthesised by algae and bacteria. Iodate is reduced in the gastrointestinal tract to iodide which is the biologically active from of iodine (SACN, 2014).

lodine function

lodine is essential in the human diet primarily because it is required 2. for the synthesis of the thyroid hormones tri-iodo- and tetra-iodothyronine (T3 and T4 or thyroxine). The thyroid hormones exert effects on a wide range of bodily functions such as basal metabolism, brain development and bone growth, especially in the fetus, which is exposed to iodine via the placenta, so maternal underexposure by iodine deficiency can have profound effects on both mother and offspring. Both chronic iodine deficiency and excess may lead to compensatory thyroid hypertrophy, known as goitre, in adults and children. lodine deficiency in pregnancy is also associated with an increased risk of miscarriage, stillbirth and congenital abnormality. lodine deficiency in the developing fetus can lead to congenital hypothyroidism (previously known as cretinism), which can have serious effects on both physical and mental development. Clinical features of acute iodine toxicity include vomiting and diarrhoea, seizure, delirium and collapsing. Sensitivity reactions such as iodide mumps following treatment with iodine-containing medication may also occur (EVM, 2003).

lodine function and status in pregnancy

3. Once iodine is consumed it is quickly absorbed in the body and enters the bloodstream. The thyroid captures the circulating iodine by oxidising it so it can be used to create the thyroid hormones triiodothyronine (T3) and thyroxine (T4). The thyroid hormones help regulate metabolism and ensure

that the heart, brain and other organs function in a healthy manner. The majority of iodine is found mainly in the thyroid glands; however, non-hormonal iodine can also be found in the mammary glands, the eyes, the gastric mucosa, the cervix and the salivary glands (Ahad et al, 2010).

4. Thyroid function and iodine levels are altered during pregnancy. In early gestation, maternal thyroid hormone production increases in response to the Thyroid Stimulating Hormone (TSH) (Glinoer, 2001). It is possible that the increase in glomerular filtration rate (GFR) results in a decrease to the circulating pool of plasma iodine (Glinoer, 2007; Gaberscek and Zaletel, 2011); it is unclear if this might be mitigated by increased renal retention of iodine. Additionally, any fall in plasma iodine concentrations would also be attributable, in part at least, to expansion of the plasma volume. A proportion of maternal thyroid hormone is transferred to the fetus, as in iodine via the placental NIS (Sodium Iodide symporter) (Glinoer, 2001; Zimmermann, 2009).

5. Urinary iodine concentration (UIC) is a good indicator of short-term iodine status as >90% of dietary iodine eventually appears in urine (Rohner, 2014). A median UIC of $\leq 100 \mu g/L$ specifies insufficient iodine intake, a median UIC of $\geq 100 \mu g/L$ indicates adequate iodine intake for women who are breastfeeding their infants and children under two years of age (Harding, 2017). UIC levels of >500 $\mu g/kg$ are considered excessive in pregnant women (WHO, UNICEF and ICCIDD, 2007).

6. As an alternative to UIC, Katko et al (2018) used plasma thyroglobulin (Tg) as a measure of iodine status in 164 pregnant Hungarian women in week 16 of pregnancy, the period corresponding to major brain development. They found that UIC corresponded with daily iodine intake whereas Tg corresponded with long-term storage levels of iodine.

7. Sixteen healthy lactating American women with no known history of thyroid disease were administered 600 μ g oral potassium iodide (KI) (456 μ g iodine) after an overnight fast. Iodide measurements were taken from breastmilk and urine at baseline and hourly for 8 hours following iodine intake. All dietary iodine ingested during the study period was also measured. Baseline breastmilk and urine iodine levels were 45.5 μ g/L and 67.5 μ g/L, respectively. Following 600 μ g KI administration, median increase in breastmilk iodine levels above baseline was 280.5 μ g/L and median peak breastmilk iodine concentration was 354 μ g/L. Median time to peak breastmilk iodine levels following KI administration was 6 hours (IQR 5– 7). Dietary iodine sources provided an additional 36–685 μ g iodine intake during the 8-hour study. It was concluded that breastmilk iodine concentrations can be interpreted in relation to recent iodine intake (Leung et al, 2012).

8. Although, iodine-deficiency is a known issue in maternal health, excess iodine intake may also occur and may potentially affect maternal health. Sources of excess iodine intake may occur via the consumption of naturally occurring iodine in water supplies, seaweed, iodine supplements, medication and milk which contains iodine resulting from feed supplements and iodophor disinfectants. Most healthy individuals are unaffected by excess iodine, however susceptible individuals with autoimmune thyroid disease such as Hashimoto's Disease³ and Grave' Disease⁴ and neonates are likely to respond adversely to excess iodine.

Excessive iodine dietary intake and/or iodine supplementation during 9. pregnancy may cause adverse effects on maternal and fetal thyroid function, birth outcomes, and offspring growth and development (Le et al, 2018). Excess iodine briefly inhibits thyroid hormone synthesis through decreasing organification⁵ by an autoregulatory mechanism known as the acute Wolff-Chaikoff effect (Wolff et al. 1949); The Wolff-Chaikov effects thus prevents the production of excess thyroid hormones in response to the intake of high levels of iodine. Healthy individuals can escape the Wolff-Chaikoff effect via downregulation of the sodium/iodide symporter in the thyrocytes, after which normal thyroid hormone synthesis restarts (Eng et al, 1999; Markou et al, 2001). Although the mechanism is not completely understood, individuals with underlying or autoimmune thyroid disease fail to escape the Wolff-Chaikoff effect. Under 36 weeks of gestational age, the immature fetal thyroid gland is also unable to escape from the acute Wolff-Chaikoff effect, making the fetus more susceptible to iodine-induced hypothyroidism (Markou et al, 2001).

Excess iodine – human health studies

10. In some countries such as Denmark, iodine deficiency in the population has been addressed through the mandatory iodisation of all salt, including table salt and salt in bread. However, monitoring instigated prior to mandatory fortification in Denmark found an increase in thyroid autoantibodies and higher levels of thyroid autoimmune disease in the 15 years following fortification (Laurberg et al, 2008; Rasmussen et al, 2008; Pedersen et al, 2011; Bliddal et al, 2015). Although iodised salt is readily available in UK supermarkets, the

³ Hashimoto disease: A condition where the immune system attacks the thyroid which leads to an underactive thyroid gland also known as hypothyroidism (Mayo clinic, 2021).

⁴ Graves disease: Is a disorder where there is an overproduction of thyroid hormones also known as hyperthyroidism (Mayo clinic, 2021).

⁵ Organification: A biochemical process that takes place in thyroid gland. lodine is incorporated into thyroglobulin for the production of thyroid hormone (Comprehensive Pharmacy Review)

UK does not have a mandatory fortification scheme for iodine (SACN, 2014). The UK government recommends that salt intakes should be reduced due to health reasons. (NHS, 2018; WHO, 2013).

11. The association of thyroid nodules (TNs)⁶ and iodine intake in pregnant women was investigated in a study by (Gao et al, 2019). Serum and spot urine samples from 2353 pregnant women were collected. Urine iodine concentration (UIC) and creatinine (Cr) level were determined in spot urine samples, serum thyroid hormones and thyroid autoantibodies. The UIC and UIC to creatinine ratio (I/Cr ratio) was found to be significantly higher in pregnant women with TNs. Thyroglobulin levels, age, pre-pregnancy body mass index and living in an iodine-excessive region were associated with TNs. The I/Cr ratio was not a significant risk factor for TNs in pregnant women in their second trimester.

12. The effects of high iodine intake on thyroid function in pregnant and lactating women was investigated in China. An epidemiological study was conducted among 130 pregnant women and 220 lactating women aged 19 - 40 years in areas that had a drinking water iodine content of >300 and 50-100 µg/L in Shanxi in 2014. Urinary iodine levels and blood thyroid stimulating hormone levels were examined. It was concluded that excess iodine intake might increase the risk of subclinical hypothyroidism in pregnant women and lactating women. It was suggested that iodine nutrition and thyroid function should be monitored in women, pregnant women and lactating women who live in areas with high environmental iodine. (Ren et al, 2018).

A cross-sectional study was performed amongst 111 lactating women 13. in a refugee camp in Algeria who lived in areas with high and very high iodine concentrations in drinking water. Breast milk iodine concentration (BMIC), UIC and the iodine concentration in the most commonly consumed foods/drinks were measured. A 24-h dietary recall was used to estimate iodine intake. Thyroid hormones and antibodies were measured in serum. Median UIC, BMIC and iodine intake across both areas were 350 µg/L, 479 µg/L and 407 µg/day, respectively. Thyroid dysfunction and/or positive thyroid antibodies were found in 33.3% of the women, of which 18.9% had hypothyroidism and 8.1% had hyperthyroidism and 6.3% had positive antibodies with normal thyroid function. Elevated thyroid antibodies were in total found in 17.1% of the women. There was no difference in the distribution of thyroid dysfunction or positive antibodies between high iodine and very high iodine drink water areas. It was found that BMIC could be considered as a good indication for iodine status among lactating women. There was an association found

⁶ Thyroid nodule (TNs): TNs are solid or fluid-filled lumps that form within the thyroid (<u>Mayoclinic, 2020</u>).

between BMIC and thyroid dysfunction and positive antibodies, which indicated that thyroid dysfunction may be caused by an excessive iodine intake. The high prevalence of thyroid dysfunction and the chronically high intake of iodine might have adverse health consequences for the women and their children. (Akare et al, 2015).

14. Three years after the baseline study described above (Akare et al, 2015) a follow-up study was conducted. This included seventy-six children from the original study with a new sample of randomly selected children being included to increase the sample size to 289. At follow-up, urinary iodine, and blood levels of thyroid hormones and serum thyroglobulin were measured. Excessive iodine intake (defined as urinary iodine levels of \geq 300 µg/L) was identified in 88% of the group at baseline (children aged 0 – 6 months) and 72% at follow-up (3 years later). At follow-up, 24% of the study group had thyroid hormone disturbance and 9% had subclinical hypothyroidism. Children with subclinical hypothyroidism showed poorer growth and were more likely to be underweight than their healthy counterparts (Aakre et al, 2016).

15. Case reports were collected following the recall of a prenatal supplement containing excessive iodine levels in Brazil. In all cases, the infant was born with goitre. In the first case, a pregnant woman at 22 weeks' gestation was diagnosed with fetal goitre via a prenatal ultrasound examination. It was reported that she had taken 2 prenatal vitamin pills/day bought from a local pharmacy. The maternal UIC on the 24th week of gestation was 902 μ g/24 h (normal 100 – 460 μ g/24 h). The prenatal vitamin was analysed and found to contain 40 mg of potassium iodide per pill, so the pregnant woman was exceeding the recommended dose of 200 μ g by 200-fold. The prenatal vitamins were discontinued. No iodine doses were reported in the remaining case reports (de Vasconcellos Thomas and Collett-Solberg, 2009).

16. There is also evidence that excessive iodine supplementation during pregnancy can increase serum TSH concentration. In one observational study, pregnant women who ingested supplements containing >200 μ g /d were found to be at an increased risk of hyperthyrotropinemia and a TSH above 3 muU/ml (Rebagliato et al, 2010).

17. Human epidemiological studies have shown variations in the incidence of thyroid cancer. In the past, Iceland has been known for its population's excess iodine intake from seafood and milk. Between 1944 to 1964, surgical specimens from Iceland and northern Scotland were compared. The PTC (Papillary thyroid cancer):FTC (Follicular thyroid cancer) ratio was 6.5 in Iceland and 3.6 in Scotland. The age-specific incidence rates for papillary carcinoma were approximately five times higher in Iceland than in Scotland in

adults older than 35 years of age. It was hypothesised that high iodine intakes contributed to the high incidence of thyroid cancer. However, it was argued the high rates are due to the volcanic nature of the island (Williams et al, 1977; Feldt-Rasmussen, 2001). Between 2003 and 2008 the German population had sufficient iodine intakes from voluntary iodization of salt. The incidence rate of thyroid cancer rose from 2.7 to 3.4 (men) and from 6.5 to 8.9 (women) per 100,000 per year and was mainly PTC (Radespiel-Troger et al, 2014).

Excess iodine - animal studies

18. Rats were fed a diet containing excess iodine (~120 mg of iodine per day) or a control diet for 9 months. There was a 40% increase in thyroid weight, and histological changes included enlarged follicles with increased colloid lined by flattened epithelia, but no thyroid tumours were found (Correa et al, 1960).

Health based guidance values

19. The Expert Group on Vitamins and Minerals (EVM) looked in detail at the metabolism of iodine and the effect of excess iodine in 2003. EVM examined iodine but were unable to set a safe upper limit (SUL) due to lack of data. It was noted that supplemental doses of 0.5 to 1.5 mg iodine/day showed a decrease in the thyroid hormone levels (Paul et al, 1988; Chow et al, 1991; Gardner et al, 1988). The Saxena et al, 1962; Freund et al, 1966 studies indicated that the supplemental doses of 2 mg/day in addition to the dietary iodine resulted in the blockage of further iodine uptake. For guidance purposes, the EVM indicated that a level of 0.5 mg/day of supplemental iodine in addition to the background intake of 0.43 mg/day (equivalent to 0.015 mg/kg bw/day in 60 kg adult) would be unlikely to cause adverse effects in adults based on slight decreases in serum thyroid hormone levels at supplemental doses of 0-2 mg/day in a range of human studies.

20. In 2002, the European Scientific Committee on Food published an opinion on the tolerable upper intake levels of vitamins and minerals. For iodine, they set a tolerable upper level (TUL) for total iodine intake of 600 μ g/day, based on the finding that biochemical changes in TSH levels and the TSH response to TRH administration were marginal and not associated with any clinical adverse effects at estimated intakes of 1700 and 1800 μ g/day. The studies on which these UL estimates were based were all only of short duration, with a small number of individuals but the results were supported by a 5- year study of exposure with an iodide intake of 30 μ g/kg bw/day

(equivalent to1800 μ g iodide/day for a 60 kg adult), where there was no clinical thyroid pathology. An uncertainty factor (UF) of 3 was considered adequate and provided the UL for adults The UL of 600 μ g was also considered to be acceptable for pregnant and lactating women based on of lack of adverse effects at significantly higher exposures (SCF, 2006).

21. JECFA established a provisional Maximum Tolerable Daily Intake (PMTDI) of 17 μ g/kg bw/day (equivalent to 1020 μ g/day for a 60 kg adult) for iodine from all sources, based on an epidemiological study in which 750 men and women were exposed to iodinated water in prison and consumed 1-2 mg of iodine per day for various time periods. Adverse effects were observed in four women who had previous thyroid issues before entering prison and became more symptomatic receiving the iodinated water supply. Out of 15 tested inmates, two had impaired organification of thyroidal iodine (JECFA, 1989).

lodine exposures in maternal health

Sources of iodine exposure

Food

22. Iodine levels have been measured in the composite food samples of The 2014 Total Diet Study (TDS) (FSA, 2016). The richest dietary sources of iodine are fish and seafood, seaweed, eggs and dairy products.

23. Seaweed has been used for centuries as a staple food in Asian countries. In some Asian countries, it is customary to serve seaweed soup to new mothers (Moon et al, 2009). Seaweed consumption is also popular in people who are on a plant-based diet (EFSA, 2019). A study found that iodine in seaweed contributed significantly to diets of vegans with excessive iodine intake (Eveleigh et al, 2020). EFSA conducted an analysis and risk assessment on seaweed. Estimated mean exposures to iodine ranged from 94.9 - 11, $512.3 \mu g/day$. Estimated 95th percentile exposures to iodine ranged from ranged from $86.0 - 18,677.2 \mu g/day$ (EFSA, 2019).

Cows' milk and milk products

24. Iodine based compounds have been used to clean the udders of cows during milk collection and these leach onto the milk, adding to the natural levels of iodine present. A survey of cows' milk carried out in 1998-

1999 found that the overall mean iodine concentration in cows' milk was 311 μ g/kg. Mean iodine concentrations were found to be lower in summer (200 μ g/kg) compared to winter (430 μ g/kg). The higher concentrations in winter may reflect greater use of supplemented compound feedstuffs during this period. At these levels, the COT concluded that the concentrations of iodine in cows' milk were unlikely to pose a risk to health, even in those children who are high level consumers (COT, 2000). However, in the Total Diet Study conducted in 2014, the overall mean iodine concentration in cows' milk was 0.26 μ g/kg (FSA, 2016). There are currently no approved iodophors in the UK on the Veterinary Medicine Directorate list.

Drinking water

25. Iodine was detected at low levels (8 μ g/L) in tap water in the 2014 TDS (FSA, 2016).

Supplements

A range of supplements that contain iodine are available to women 26. who are trying to conceive as well as pregnant women and breastfeeding mothers. These supplements contain multiple components and can contain up to 150 µg of iodine per dose. Women are usually advised to consult their doctor before taking these supplements. The NHS states that adults need to have a daily iodine intake of 140 µg However, most people should be able to obtain the required amount of iodine by eating a varied and balanced diet. People who follow a strict vegan diet and do not consume any fish are advised to take an iodine supplement (NHS, 2020). The UK Dietary Reference Values (DRVs) do not include an increment in iodine for pregnant or lactating women. COMA advised on the premise that women of reproductive age should have customary intakes that would enable them to manage pregnancies without any need for supplements. In its statement on iodine and health, SACN considered there was insufficient evidence to substantiate revisions to the UK DRVs for iodine for pregnant and lactating women (SACN, 2014). No data specifically focusing on the influence of regular use of iodine supplements on levels of iodine in breast milk of UK mothers were identified.

Environmental – Dust and soil

27. Iodine levels in soil are highly variable. A median value of 5.9 mg/kg and a 90th percentile value of 14.2 mg/kg have been reported for UK soil by the British Geological Survey (BGS 2016). No specific value for dust was identified from the literature.

Air

28. According to the expert panel on Air Quality Standards, concentrations of particle bound iodine in UK air between 1996 and 2001 ranged from 0.8 $x10-6 - 2.0 \times 10-6 \text{ mg/m}^3$ (DEFRA, 2006).

Medication

29. Iodine is used as a topical antiseptic, which can result in absorption through the skin. Excessive iodine intakes can occur following the ingestion of iodide-containing pharmaceuticals for the treatment of asthma, bronchitis, cystic fibrosis, chronic obstructive pulmonary disease, goitre. Iodine is also found in tropical antiseptics, mouthwashes, vaginal solutions and burn and wound treatments (SCF, 2006). Absorption in infants appears to be greater than in adults (Leung and Braverman, 2014). There are other medications that also contain iodine, which may be released metabolically into the systemic circulation.

Exposure assessment

Exposure estimates based on the TDS

30. A Total Diet Study (TDS) is described as a complementary approach to traditional monitoring and surveillance programs (EFSA, 2011). The TDS is used to calculate population dietary exposure to a range of chemicals in food and to assess the safety and/or nutritional quality of food. TDSs involve selecting, collecting and analysing commonly consumed food purchased at retail level, processing the food as for consumption, pooling the prepared food items into food groups that are representative, homogenising the pooled samples and analysing them for harmful and/or beneficial chemical substances (FSA, 2019).

31. Table 2 summarises total dietary exposures to iodine calculated using the iodine concentrations determined from food groups in 2014 Total Diet Study (TDS). The exposure assessment was carried out for women of childbearing age (16 - 49 years old) using food groups and consumption data from years 1 - 8 of the NDNS survey (Bates et al., 2014; 2016; 2018). The NDNS (Bates et al., 2014; 2016; 2018) does not provide data for pregnant or lactating women so while data is based on women of childbearing age, this data may not necessarily be representative of the maternal diet. Mean chronic iodine exposures from the total diet of women aged 16- 49 years old was 1.7

 $\mu g/kg$ bw/day and the 97.5th percentile chronic exposure was 3.7 $\mu g/kg$ bw/day.

32. This TDS comprises of 27 food groups. The food groups making the highest contribution to iodine exposure in the TDS were milk, followed by fish and seafood. It should be noted that it is advised that pregnant women should avoid particular types of cheese, dairy products, meat, eggs, fish and avoid drinking alcohol (NHS, 2020b). Therefore, pregnant or breastfeeding women may have a different diet compared to non-pregnant or breastfeeding women in the same age range as they may choose to increase or decrease consumption of certain foods or drinks due to this advice.

Consumer- based and population-based exposures estimates based on foods in the Total Diet Study (TDS)

Table 2. Estimated chronic exposure for iodine from the total diet in women aged 16 – 49 years old (TDS 27 groups) (Bates et al., 2014; 2016; 2018)

Consumers (n)	Chronic						
	exposure	exposure					
	Mean (µg/kg bw/day)	P97.5 th (µg/kg bw/day)	Respondents in population group (n)				
1874	1.7	3.7	1874				

Exposure estimates for seaweed

33. As noted in paragraph 22, seaweed has been noted as a rich source of iodine, particularly in people on a plant-based diet or from a culture where dietary seaweed consumption is high. For this reason, iodine levels in seaweed have been used to give exposure estimates.

34. A search within the recipes database of the NDNS (Bates et al., 2014; 2016; 2018) was conducted to retrieve seaweed and seaweed products which had been recorded in the survey. These can be seen in column one of table 3. Column three is the assumption made for the type of seaweed in each food, where it has not been specified. These assumptions were based on common uses of seaweed.

Table 3.Seaweed or seaweed-containing foods recorded in the NDNS(Bates et al., 2014; 2016; 2018)

Food as recorded in NDNSDescription of foodAssumed type of seaweed
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Higher nature energy breakfast shake dry powder	Meal replacement shake	Kelp/ kombu
Laverbread	Welsh seaweed dish	Laver seaweed aka Nori
Sushi, tuna based	NA	Nori
Sushi, vegetarian	NA	Nori
Sushi, salmon based	NA	Nori
Soup with tofu and seaweed	NA	Wakame
Vecon	Vegetable stock	Kelp/ kombu
Seaweed wakame dried raw	NA	Wakame

35. It should be noted that levels of iodine vary between types of seaweed; this can be seen in the EFSA 'Analysis and Risk Assessment of Seaweed' where kelp (Saccharina latissima) had a higher iodine concentration in comparison to other seaweed species (EFSA, 2019). In this case, iodine concentration in kelp ranged from $333.0 - 4,782.2 \mu g/g$ freeze dried weight. Three other species had lower ranges of $137.8-451.2 \mu g/g$, $105.2-961.4 \mu g/g$ and $17.2-20.8 \mu g/g$ of freeze-dried weight (fdw).

36. The levels of iodine vary in different types seaweeds (Yeh et al, 2014) where kelp/ kombu can be compared to wakame and nori, which is more specific to the data available from the NDNS. In this case, the range for nori was $29.3 - 45.8 \mu g/g$, for wakame $93.9 - 185.1 \mu g/g$ and for kombu 241- 4921 $\mu g/g$ in dried seaweed weight.

37. Seaweed harvested from different countries may also have differing levels of iodine, this is also illustrated by Yeh et al, 2014. 10 samples each of nori, wakame and kombu harvested from China, Japan, Korea and Taiwan were analysed with varied results (Yeh et al, 2014).

38. Another uncertainty was that seaweed consumption based on the foods in table 3 and NDNS survey data for women of childbearing age (16 – 49 years) yielded consumption data based on very few consumers (Bates et al., 2014; 2016; 2018). Exposure or consumption estimates based on very few consumers should be treated with caution, particularly 97.5th percentile estimates based on less than 60 consumers. Furthermore, as mentioned previously, the NDNS does not consider pregnant or lactating women so these estimates may not be representative of the target population. The chronic consumption estimates can be seen in table 4.

Table 4. Chronic consumption estimates of seaweed in women aged 16 - 49 years old (Bates et al., 2014; 2016; 2018)

Consumers (n)	g/ person/day*		g/kg bw/day*		
	Mean	P97.5 th	Mean	P97.5 th	Respondents in population group (n)
36	1.1	3.8	0.017	0.060	1874

*rounded to 2.s.f

39. Further to the data considered from literature in paragraphs 33 - 35, real time data was consulted. The Rapid Alert for Food and Feed (RASFF Portal) is a tool that provides information on public health warnings issued by food safety authorities and food companies. It also provides the latest information on food recall notices. Between March 2019 to August 2020, there were 37 reported incidences whereby levels of iodine were detected in seaweed samples at above the alert, recall or 'information for follow-up' level according to the RASFF Portal. The concentrations of iodine in these seaweed samples ranged from 25 µg/g to 20,620 µg/g. Countries of origin in order of contribution were: South Korea, China, Germany, Spain, Japan, Belgium and Austria. These are similar to countries from which samples were derived by Yeh et al, 2014. This range is much greater than that seen by the EFSA analysis of kelp (EFSA, 2019) and is considered to be a worst-case scenario.

40. After reviewing all of the information in paragraphs 35– 36 and tables 3 and 4, exposure estimates were calculated using NDNS seaweed consumption data and the iodine concentration range measured in 16 kelp samples by EFSA, $333.0 - 4,782.2 \mu g/g$ fdw (EFSA, 2019). Due to the various uncertainties discussed such as the varying levels of iodine in different types of seaweed and the limited consumption data available in the NDNS this was considered the most realistic scenario.

Table 5: Chronic exposure estimates using chronic consumption data from table 4 and the iodine concentration range measured in kelp by EFSA, 2019 (minimum: 333.0 μ g/g fdw, maximum: 4,782.2 μ g/g fdw).

Age	Numbe	Chronic	Chronic	Chronic	Chronic
group	r of	exposur e	exposure μg/	exposure μg/kg	exposur e
	consu mers	μg/ person/ day*	person/ day*	bw/day*	μg/kg bw/day*

		Mean		97.5th Percentile		Mean		97.5th Percentil e	
woman 16-49 vears	36	Min	Max	Min	Max	Min	Max	Min	Max
		360	5200	1300	18000	5.5	79	20	290

*rounded 2.s.f

41. The exposure data in table 5 should be treated with caution because it is based on consumption data from a limited number of consumers, particularly the 97.5th percentile estimates. Another important caveat is that the use of unprocessed seaweed biomass is a conservative approach for exposure assessments which is likely to lead to an overestimation. The minimum and maximum seaweed iodine values are only based on 16 samples. The effects of cooking and processing as well as bioavailability are not taken into account for this assessment as the data from EFSA 2019 is based on freeze dried samples of seaweed (EFSA, 2019).

Risk characterisation

42. In 2003, the EVM set a guidance level for iodine of 15 μ g/kg bw/day that would not be expected to cause adverse effects in the majority of the population. It should be noted that this is a guidance level only as there was insufficient data from human or animal studies to establish a Safe Upper Level for iodine. Therefore, its applicability to maternal health is uncertain. The exposure estimates in Table 2 are below the guidance value. As seen in table 5, the estimated mean exposure for seaweed based on the minimum value of 333.0 μ g/g freeze dry weight (fdw) of iodine was 5.5 μ g/kg and is below the guidance value. The estimated mean exposure for seaweed based on the maximum value of 4782.2 μ g/g fdw of iodine was 79 μ g/kg bw/day and is above the set guidance value. The estimated 97.5th percentile exposures for seaweed based on the minimum and maximum values of 333.0 μ g/g and 4782.2 μ g/g fdw iodine was 20 and 288 μ g/kg bw/day, respectively. The estimated exposures are above the guidance value.

43. JECFA established a provisional Maximum Tolerable Daily Intake (PMTDI) of 17 μ g/kg bw/day for iodine from all sources, based on the same longer- term studies in adults used by SCF in 2002 to support the TUL. The exposure estimates in Table 2 are below the PMTDI. As seen in table 5, the

estimated mean exposure for seaweed based on the minimum value of 333.0 μ g/g fdw of iodine is below the PMTDI. The estimated mean exposure for seaweed based on the maximum value of 4782.2 μ g/g fdw of iodine is above the PMTDI. The 97.5th percentile exposure estimates for seaweed based on the minimum (333.0 μ g/g fdw) and maximum iodine (4782.2 μ g/g fdw) values are above the PMTDI.

44. The SCF established a UL of 600 μ g/day for pregnant and lactating women based on of lack of adverse effects at significantly higher exposures (SCF, 2000). The exposure estimates in table 2 are below the UL. The estimated mean exposure for seaweed based on the minimum value of 333.0 μ g/g fdw of iodine was below the UL. The estimated mean exposure based on the maximum value of 4782.2 μ g/g fdw iodine was above the UL. The estimated 97.5th percentile exposures for seaweed based on the minimum (333.0 μ g/g fdw) and maximum iodine (4782.2 μ g/g fdw) values are above the UL.

Conclusions of the Committee

45. Iodine is essential in the diet to produce thyroid hormones which are involved in cell metabolism, growth and development. However, both exposure to excess iodine and iodine deficiency in pregnancy may cause adverse effects on maternal and fetal thyroid function, birth outcomes, offspring growth and development.

46. Iodine status in the individual is difficult to assess since healthy people excrete excess iodine in urine so it is not a good marker for individual exposure. However, iodine deficiency and excess can be identified in a population by looking at the distribution of urinary iodine.

47. Iodine intake from the diet is within the HBGVs set by bodies such as EFSA, JECFA and the EVM for adults and would not be a risk to maternal health.

48. There are no concerns with regard to excess iodine in the general population, however, exposure to excess iodine in high seaweed consumers could pose a potential risk to maternal health. Currently available data was not sufficient to enable a risk benefit assessment to be performed.

Secretariat March 2021

References

Aakre I, Strand TA, Bjøro T, Norheim I, Barikmo I, Ares S, Alcorta MD, Henjum S. Thyroid Function among Breastfed Children with Chronically Excessive Iodine Intakes. Nutrients 2016, 8, 398; doi:10.3390/nu8070398

Aakre I, Bjøro T, Norheim I, Strand TA, Barikmo I, Henjum S (2015) Excessive iodine intake and thyroid dysfunction among lactating Saharawi women, J. Trace Elem, Med. Biol. 31 (637-641)

Ahad F, Ganie SA (2010). Iodine, Iodine metabolism and Iodine deficiency disorders revisited. Indian J Endocrinol Metab. 2010;14(1):13-17.

Bates, B.; Lennox, A.; Prentice, A.; Bates, C.; Page, P.; Nicholson, S.; Swan, G. (2014) National Diet and Nutrition Survey Results from Years 1, 2, 3 and 4 (combined) of the Rolling Programme (2008/2009 – 2011/2012) Available at: https://www.gov.uk/government/uploads/system/uploads/attachment_data/file/ 31099 5/NDNS_Y1_to_4_UK_report.pdf

Bates, B.; Cox, L.; Nicholson, S.; Page, P.; Prentice, A.; Steer, T.; Swan, G. (2016) National Diet and Nutrition Survey Results from Years 5 and 6 (combined) of the Rolling Programme (2012/2013 – 2013/2014) Available at: https://assets.publishing.service.gov.uk/government/uploads/system/uploads/ attach ment_data/file/551352/NDNS_Y5_6_UK_Main_Text.pdf

Bates, B.; Cox, L.; Nicholson, S.; Page, P.; Prentice, A.; Steer, T.; Swan, G. (2018) National Diet and Nutrition Survey Results from Years 7 and 8 (combined) of the Rolling Programme 2014/2015 – 2015/2016. Available at: <u>https://assets.publishing.service.gov.uk/government/uploads/system/uploads/attachment_data/file/699241/NDNS_results_years_7_and_8.pdf</u>

Bath SC, Rayman MP. A review of the iodine status of UK pregnant women and its implications for the offspring. Environ Geochem Health. 2015;37(4):619-629. doi:10.1007/s10653-015-9682-3

Bliddal S, Boas M, Hilsted L, Friis-Hansen L, Tabor A and Feldt-Rasmussen U. (2015) Thyroid function and autoimmunity in Danish pregnant women after an iodine fortification program and associations with obstetric outcomes. European Journal of Endocrinology. 173. 709-718

British Geological Survey (BGS) (2016) Map of soil iodine in the UK available at: <u>http://www.ukso.org/nsi/lodine.html</u>

Carswell F, Kerr MM, Hutchinson JH (1970). Congenital goitre and hypothyroidism after topical iodine in pregnancy and lactation. Lancet i:1241-1243.

Chow, C.C., Phillips, D.I.W., Lazarus, J.H., Parkes, A.B. (1991) Effect of low dose iodide supplementation on thyroid function in potentially susceptible subjects: are dietary iodide levels in Britain acceptable? Clinical Endocrinology 34, 423-416.

Committee of Toxicity (2000) Statement on iodine in cows' milk. Available at: <u>https://cot.food.gov.uk/sites/default/files/cot/iodin2.pdf</u>

Committee of Toxicity (2017) Statement on the potential risks from excess iodine in the diets of infants aged 0 -12 months and children aged 1 to 5 years. Available

at:https://webarchive.nationalarchives.gov.uk/20200808005653/https://cot.foo d.gov.uk/cotstatements/cotstatementsyrs/cot-statements-2017/statement-onthe-potential-risks-from-excess-iodine

Comprehensive Pharmacy Review, Leon Shargel, 6th edition, p1181

Correa P, Welsh RA (1960) The effect of excessive iodine intake on the thyroid gland of the rat. Arch Pathol. 1960:70:247 - 51

de Vasconcellos Thomas J and Collett-Solberg PF. (2009) Perinatal goitre with increased iodine uptake and hypothyroidism due to excess maternal iodine ingestion. Horm Res 72 (344-347).

DEFRA (Department for Environment, Food and Rural Affairs) (2006). Guidelines for Halogens and Hydrogen Halides in Ambient Air for Protecting Human Health Against Acute Irritancy Effects. Available at: http://webarchive.nationalarchives.gov.uk/20060715141954/http://www.defra.g ov.uk/ environment/airquality/aqs/halogens/fullreport.pdf

Department of Health and Social Security (DHSS) (1977) The composition of mature human milk. Available at:

https://assets.publishing.service.gov.uk/government/uploads/system/uploads/ attach ment_data/file/743819/The_Composition_of_Mature_Human_Milk 1977_.pdf

EFSA (2019) Analysis and Risk Assessment of Seaweed. Available at: https://efsa.onlinelibrary.wiley.com/doi/epdf/10.2903/j.efsa.2019.e170915

Eng PH, Cardona GR, Fang SL, Previti M, Alex S, Carrasco N, et al. Escape from the acute Wolff-Chaikoff effect is associated with a decrease in thyroid sodium/iodide symporter messenger ribonucleic acid and protein. Endocrinology. 1999;140:3404– 3410.

Expert Group on Vitamins and Minerals (EVM, 2003) Safe Upper Levels for Vitamins and Minerals.

https://cot.food.gov.uk/sites/default/files/vitmin2003.pdf

Eveleigh ER, Coneyworth LJ, Avery A, Welham SJM. Vegans, Vegetarians, and Omnivores: How Does Dietary Choice Influence Iodine Intake? A Systematic Review. Nutrients. 2020;12(6):1606. Published 2020 May 29. doi:10.3390/nu12061606 Feldt-Rasmussen U. Iodine and cancer. Thyroid. 2001;11:483-6

FSA (2019) Total diet study: metals and other elements. Available at: https://www.food.gov.uk/research/research-projects/total-diet-study-metalsand- other-elements

FSA (2016) Metals and other elements in the 2014 Total Diet Study. Available at: https://www.food.gov.uk/sites/default/files/media/document/measurement-of-the- concentrations-of-metals-and-other-elements-from-the-2014-uk-total-diet-study.pdf

Gardner, D.F., Centor, R.M., Utiger, R.D. (1988) Effects of low dose oral iodide supplementation on thyroid function in normal men. Clinical Endocrinology 28, 283-288.

Gao M, Chen W, Sun H, Fan L, Wang W, Du C, Chen Y, Lin L, Pearce EN, Shen J, Cheng Y, Wang C, Zhang W (2019) Excessive iodine intake is associated with formation of thyroid nodules in pregnant Chinese women.

Gaberšček S, Zaletel K. Thyroid physiology and autoimmunity in pregnancy and after delivery. Expert Review of Clinical Immunology 2011;7(5):697-707.

Glinoer D, de Nayer P, Delange F, Lemone M, Toppet V, Spehl M, Grün JP, Kinthaert J, Lejeune B (1995). A randomized trial for the treatment of mild iodine deficiency during pregnancy: Maternal and neonatal effects. J Clin Endocrin Metab 80: 258-269.

Glinoer D. Pregnancy and iodine. Thyroid 2001;11 (5): 471-81.

Glinoer D. The importance of iodine nutrition during pregnancy. Public Health Nutrition 2007;10(12A):1542-1546.

Freund, G., Thomas, W.C. Jr., Bird, E.D., Kinman, R.N., Black, A.P. (1966) Effect of iodinated water supplies on thyroid function. Journal of Clinical Endocrinology 26, 619-624.

Harding KB, Peña-Rosas JP, Webster AC, Yap CMY, Payne BA, Ota E, De-Regil LM (2017). Iodine supplementation for women during the preconception, pregnancy and postpartum period. Cochrane Database of Systematic Reviews 2017, Issue 3. Art. No.: CD011761. DOI: 10.1002/14651858.CD011761.pub2.

JECFA (1989) Monograph on iodine available at: http://www.inchem.org/documents/jecfa/jecmono/v024je11.htm

Katko M, Gazso AA, Hircsu I, Bhattoa HP, Molnar Z, Kovacs B, Andrasi, D, Aranyos J, Maka Ri, Veress L, Torok O, Bodor M, Samson L, Nagy LM (2018). Thyroglobulin level at week 16 of pregnancy is superior to urinary iodine concentration in revealing preconceptual and first trimester iodine supply Maternal and Child Nutr. 2018 14(1): e12470.

Laurberg P, Jorgensen T, Perrild H, Ovesen L, Knudsen N, Bulow Pedersen I, Rasmussen L B, Carle A and Vejbjerg P. (2006) The Danish investigation on iodine intake and thyroid disease, DANThyr: status and perspectives. European Journal of Endocrinology 155 219-228

Lee YA, Cho SW, Sung HK, et al. Effects of Maternal Iodine Status during Pregnancy and Lactation on Maternal Thyroid Function and Offspring Growth and Development: A Prospective Study Protocol for the Ideal Breast Milk Cohort. Endocrinol Metab (Seoul). 2018;33(3):395-402. doi:10.3803/EnM.2018.33.3.395

Leung AM, Pearce EN, Braverman LE (2011). Iodine nutrition in pregnancy and lactation. Endocrinology and Metabolism Clinics of North America 2011;40(4):765-77

Leung AM, Braverman LE, He X, Heeren T, Pearce EN. Breastmilk iodine concentrations following acute dietary iodine intake. Thyroid. 2012;22(11):1176- 1180. doi:10.1089/thy.2012.0294

Markou K, Georgopoulos N, Kyriazopoulou V, Vagenakis AG. lodine-induced hypothyroidism. Thyroid. 2001;11:501–510.

This is a draft statement for discussion. It does not reflect the final views of the Committee and should not be cited.

Mayo Clinic (2020a) Hashimito's disease. Available at: <u>https://www.mayoclinic.org/diseases-conditions/hashimotos-</u> <u>disease/symptoms-causes/syc-20351855</u>

May Clinic (2020b) Graves' disease. Available at: <u>https://www.mayoclinic.org/diseases-conditions/graves-disease/symptoms-causes/syc-20356240</u>

Moleti M, Di Bella B, Giorgianni G, Mancuso A, De Vivo A, Alibrandi A, Trimarchi F, Vermiglio F. Maternal thyroid function in different conditions of iodine nutrition in pregnant women exposed to mild-moderate iodine deficiency: an observational study. Clin Endocrinol (Oxf) 2011;74:762–8.

Moon S, Jungyeon K (1999) lodine content of human milk and dietary iodine intake of Korean lactating mothers, International Journal of Food Sciences and Nutrition, 50:3, 165-171, DOI: 10.1080/096374899101201

NHS (2020) Iodine: Vitamins and minerals. Available at: https://www.nhs.uk/conditions/vitamins-and-minerals/iodine/

NHS (2018) Tips for a lower salt diet. Available at: https://www.nhs.uk/livewell/eat-well/tips-for-a-lower-salt-diet/

Paul, T., Meyers, B., Witorsch, R.J., Pino, S., Chipkin, S., Ingbar, S.H., Braverman, L.E. (1988) The effect of small increases in dietary iodine on thyroid function in euthyroid subjects. Metabolism 37, 121-124.

Pedersen I B, Knudsen N, Carle A, Vejbjerg P, Jorgensen T, Perrild H, Ovesen L, Rasmussen L B and Laurberg P. (2011) A cautious iodization programme bringing iodine intake to a low recommended level is associated with an increase in the prevalence of thyroid autoantibodies in the population. Clinical Endocrinology 75 120-126

Prado EL, Dewey KG. Nutrition and brain development in early life. Nutrition Reviews 2014;72(4):267-84. [DOI: doi:10.1111/nure.12102] Public Health England (2016) Government Dietary Recommendations: Government recommendations for energy and nutrients for males and females aged 1 – 18 years and 19+ years. Available at: https://assets.publishing.service.gov.uk/government/uploads/system/uploads/ attach ment_data/file/618167/government_dietary_recommendations.pdf

Radespiel-Troger M, Batzler WU, Holleczek B, Luttmann S, Pritzkuleit R, Stabenow R, et al. Im Namen der Gesellschaft der epidemologischen Krebsregister in Deutschland e V [Rising incidence of papillary thyroid carcinoma in Germany]. Bundesgesundheitsbl Gesundheitsforsch Gesundheitsschutz. 2014;57:84-92

Rasmussen L B, Carle A, Jorgensen T, Knudsen N, Laurberg P, Pedersen I B, Perrild H, Vejbjerg P and Ovesen L. (2008) Iodine intake before and after mandatory iodization in Denmatk: results from the Danish Investigation of Iodine Intake and Thyroid Diseases (DanThyr) study. British Journal of Nutrition 100 166-173

Rebagliato M, Murcia M, Espada M, Alvarez-Pedrerol M, Bolúmar F, Vioque J, Basterrechea M, Blarduni E, Ramón R, Guxens M, et al. . lodine intake and maternal thyroid function during pregnancy. Epidemiology 2010;21:62–9.

Ren YT, Jia QZ, Zhang XD, Guo BS, Zhang FF, Cheng XT, Wang YP (2018) Prevalence of thyroid function in pregnant and lactating women in areas with different iodine levels of Shanxi province. Chinese Journal of Epidemiology, vol 39, no. 5, pp. 609-613

Rohner F, Zimmermann M, Jooste P, Pandav C, Caldwell K, Raghavan R, et al (2014). Biomarkers of nutrition for development - iodine review. Journal of Nutrition 2014;144(8):1322S-42S.

SACN (2011) The influence of maternal, fetal and child nutrition on the development of chronic disease later in life: <u>https://assets.publishing.service.gov.uk/government/uploads/system/uploads/attach ment_data/file/339325/SACN_Early_Life_Nutrition_Report.pdf</u>

Scientific Advisory Committee on Nutrition (2014) SACN Statement on Iodine and Health. Available

at:https://assets.publishing.service.gov.uk/government/uploads/system/upload s/attac hment_data/file/339439/SACN_lodine_and_Health_2014.pdf

SACN (2018) Feeding in the first year of life:

https://www.gov.uk/government/publications/feeding-in-the-first-year-of-lifesacnreport

Scientific Committee on Food (SCF) / Scientific Panel on Dietetic Products, Nutrition and Allergies (NDA). (2006). Tolerable upper intake levels for vitamins and minerals.

http://www.efsa.europa.eu/sites/default/files/efsa_rep/blobserver_assets/ndat olerabl_euil.pdf Saxena, K. M., Chapman, E.M., Pryles, C.V. (1962) Minimal dosage of iodide required to suppress uptake of iodine-131 by normal thyroid. Science 138, 430-431.

The Association of UK Dieticians (BDA) (2021) Iodine: Food Fact Sheet. Available at: <u>https://www.bda.uk.com/resource/iodine.html</u>

World Health Organisation (WHO) (2013) Salt Reduction and Iodine Fortification Strategies in Public Health. Available online: http://apps.who.int/iris/bitstream/10665/101509/1/9789241506694_eng.pdf

WHO, UNICEF and ICCIDD. Assessment of the iodine deficiency disorders and monitoring their elimination. 2007 WHO/NHD/01.1 [online], http://whqlibdoc.who.int/publications/2007/9789241595827eng.pdf

Williams ED, Doniach L, Bjarnason O, Michie W. Thyroid cancer in an Iodide Rich Area – Histopathological Study. Cancer. 1977;39:215-22

Wolff J, Chaikoff IL, Goldberg RC, Meier JR. The temporary nature of the inhibitory action of excess iodine on organic iodine synthesis in the normal thyroid. Endocrinology. 1949;45:504–513.

Yeh TS, Hung NH, Lin TC (2014) Analysis of iodine content in seaweed by GC-ECD and estimation of iodine intake. Journal of Food and Drug Analysis 2014:22:189 – 196

Zimmerman MB (2009) Iodine deficiency. Endocr Rev. 2009 Jun; 30(4):376-408.