Articles

Costs and benefits of iodine supplementation for pregnant women in a mildly to moderately iodine-deficient population: a modelling analysis



Summary

Background Results from previous studies show that the cognitive ability of offspring might be irreversibly damaged as a result of their mother's mild iodine deficiency during pregnancy. A reduced intelligence quotient (IQ) score has broad economic and societal cost implications because intelligence affects wellbeing, income, and education outcomes. Although pregnancy and lactation lead to increased iodine needs, no UK recommendations for iodine supplementation have been issued to pregnant women. We aimed to investigate the cost-effectiveness of iodine supplementation versus no supplementation for pregnant women in a mildly to moderately iodine-deficient population for which a population-based iodine supplementation programme—for example, universal salt iodisation—did not exist.

Methods We systematically searched MEDLINE, Embase, EconLit, and NHS EED for economic studies that linked IQ and income published in all languages until Aug 21, 2014. We took clinical data relating to iodine deficiency in pregnant women and the effect on IQ in their children aged 8–9 years from primary research. A decision tree was developed to compare the treatment strategies of iodine supplementation in tablet form with no iodine supplementation for pregnant women in the UK. Analyses were done from a health service perspective (analysis 1; taking direct health service costs into account) and societal perspective (analysis 2; taking education costs and the value of an IQ point itself into account), and presented in terms of cost (in sterling, relevant to 2013) per IQ point gained in the offspring. We made data-supported assumptions to complete these analyses, but used a conservative approach that limited the benefits of iodine supplementation and overestimated its potential harms.

Findings Our systematic search identified 1361 published articles, of which eight were assessed to calculate the monetary value of an IQ point. A discounted lifetime value of an additional IQ point based on earnings was estimated to be £3297 (study estimates range from £1319 to £11967) for the offspring cohort. Iodine supplementation was cost saving from both a health service perspective (saving £199 per pregnant woman [sensitivity analysis range -£42 to £229]) and societal perspective (saving £4476 per pregnant woman [sensitivity analysis range £540 to £4495]), with a net gain of 1.22 IQ points in each analysis. Base case results were robust to sensitivity analyses.

Interpretation Iodine supplementation for pregnant women in the UK is potentially cost saving. This finding also has implications for the 1.88 billion people in the 32 countries with iodine deficiency worldwide. Valuation of IQ points should consider non-earnings benefits—eg, health benefits associated with a higher IQ not germane to earnings.

Funding None.

Introduction

Severe iodine deficiency in pregnancy is linked to impaired neurodevelopment of the unborn child, manifesting in a permanent reduction in intelligence quotient (IQ) and cretinism in some children.^{1,2} A systematic review of iodisation programmes and trials in China reported an 8.7 (95% CI 6.3-11.1) IQ point difference in children born to women in severely deficient regions with and without iodine supplementation during pregnancy and after birth.² However, strong evidence exists^{1,2} that this cognitive impairment is prevented by iodine supplementation in pregnancy. Although the cognitive effects of severe iodine deficiency in pregnant women are established, the effect of mild iodine deficiency is less clear.3 In two cohort studies in the UK and Australia, 9-year-old children of women who had a urinary iodine concentration suggestive of mild iodine deficiency during their pregnancy had reduced education outcomes⁴ and decreased IQ scores⁵ compared with children of iodine-replete women. By contrast, a large Spanish cohort study, ⁶ which undertook cognitive assessment of infants at a median age of 16 months, did not report a significant association between iodine supplementation and cognitive outcomes.

The UK is one of a decreasing number of countries that does not have any iodine fortification of food or salt and some of the UK population is now believed to have become mildly iodine deficient.⁷ At present, no national guidance for iodine supplementation has been issued to pregnant women, even though pregnancy and lactation lead to increased iodine requirements.⁸⁹

A reduced IQ in infancy has broad future economic societal costs because cognitive development has effects on health outcomes, educational attainment, and lifetime earnings. A reduced IQ is associated with an increased

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Research in context

Evidence before this study

We searched PubMed, with no restrictions on date or language, with the keywords "iodine", "dietary supplements", "pregnancy", "intelligence", "cognition", "child development", and "costs and cost analysis". We identified no studies reporting the cost-effectiveness of iodine supplementation in pregnancy in a mildly iodine-deficient population. We identified a 2013 systematic review, which focused on mildly to moderately iodine-deficient populations, which concluded that "the impact of maternal iodine supplementation on newborn neurodevelopment remains uncertain due to lack of appropriate controlled intervention trials". One other systematic review of research in mildly to moderately iodine-deficient populations from 2009 reported benefits of maternal iodine supplementation on maternal thyroid indices, but underlined the need for further data for infant neurodevelopment. In two cohort studies in the UK and Australia, 9-year-old children of women who had a urinary iodine concentration suggestive of

rate of mortality,¹⁰ an increased risk of suicide,¹¹ psychiatric illness,^{12,13} and an increased incidence of heart disease.^{14,15} An increased IQ is postulated to have a positive effect on an individual's health-improvement behaviour,¹⁶ and those with increased childhood IQ scores are significantly more likely to have higher educational attainment and earnings by the age of 25 years.¹⁷

Here, we report the results of a model-based economic evaluation using the best available data from the existing published scientific literature, a systematic literature search, and expert clinical input. Because of the need for data-supported assumptions to complete the analysis, we used an approach that limited the benefits of iodine supplementation and overestimated its potential harms as far as possible. We aimed to use economic evaluation to compare the costs and benefits of a strategy of iodine supplementation tablets with a strategy of no iodine supplementation for pregnant women in a mildly to moderately iodine-deficient population.

Methods

Model structure

We developed a decision tree model in TreeAgePro 2014 (TreeAge Software, Williamstown, MA, USA) to represent two alternative strategies—iodine supplementation versus no iodine supplementation. The model pathways (figure) represent the alternative clinical pathways undertaken by pregnant women.

We are interested in the incremental costs and effects of giving universal iodine supplementation to pregnant women. The model pathways represent the alternative (simplified) clinical pathways undertaken by pregnant women. Some of the pregnancies will be unsuccessful and we will not realise any IQ gain related benefits of iodine supplementation. Successful pregnancies without mild iodine deficiency during their pregnancy had reduced educational outcomes and decreased IQ scores compared with children of iodine-replete mothers.

Added value of this study

To our knowledge, this study is the first model of the costeffectiveness of iodine supplementation in pregnant women in a mildly to moderately iodine-deficient population. We used an economic model of best available evidence and assumptions that do not favour iodine supplementation and identified that universal iodine supplementation before pregnancy and during pregnancy and lactation increased the child's IQ by 1.22 points, saved the UK NHS £199 and society £4476 per pregnant woman.

Implications of all the available evidence

Available evidence suggests that a policy of iodine supplementation during pregnancy would be benefical. Ideally, a randomised controlled trial would be undertaken to confirm our findings.

complications due to iodine supplementation intervention whose mothers were iodine deficient presupplementation receive IQ gain related benefits for their newborn baby. Pregnant women who were iodine sufficient without supplementation with normal thyroid function receive no IQ gains for their offspring. We have stacked the cards against iodine supplementation being cost effective with very conservative assumptions. We assumed that some women suffer adverse thyroid dysfunction as a result of iodine supplementation. An increased incidence of pregnancy losses and complications for each type of thyroid dysfunction is caused by iodine supplementation. Pregnancy complications push up the costs from iodine supplementation. Hypothyroidism and isolated hypothyroxinaemia induced by iodine supplementation are assumed to lose IQ points for the offspring. IQ points were subtracted for the children of mothers with adverse thyroid dysfunction from iodine supplementation who were iodine deficient presupplementation.

Variables and their respective sources are listed in table 1. Data for the probability of a pregnant woman in the UK being iodine deficient is based on the only UK data available and refers to a selective cohort of women.⁵ The proportion of pregnant women in each iodine status category—mild to moderate (median urinary iodine concentration [UIC] of 50–149 μ g/L) and severe (UIC <50 μ g/L)—and the subsequent effect on a child's IQ were based on a cohort study⁵ that examined the association between iodine status during the first trimester of pregnancy and the IQ of their children at age 9 years.

The cost of iodine tablets is based on the cost of local supermarket multivitamin tablets for pregnant women (\pounds 3.50 for 30 tablets typically containing 140 µg or 150 µg of iodine).³⁷ A daily dose of 150 µg of iodine is recommended by the American Thyroid Association³⁸ and

the European Thyroid Association³⁹ for euthyroid pregnant and lactating women. Annual incremental health and services costs and public sector costs, including education by a child's IQ category, are taken from a study⁴⁰ that looked at the costs associated with neurological impairment when children were aged 11 years old. The incremental childhood cost of preterm birth and the health-care cost associated with stillbirth are taken from published sources.^{41,42} We discounted costs at the standard annual rate of $3.5\%^{24}$ and updated them to 2013 prices using a subset of the Consumer Price Index, covering price inflation in education, health, and social protection.⁴³ Public sector costs included in the model consist of health and social services costs, and education costs.

We did an additional systematic search of the scientific literature to establish a monetary value for an IQ point to use in the economic evaluation (appendix p 2). To complete the analysis, some pragmatic assumptions were needed and were informed by the scientific literature and expert opinion. As far as possible, assumptions were conservative. Our model assumptions are listed in panels 1 and 2.

Analyses

We did two separate analyses. In the first analysis (analysis 1), we used a health service perspective in which direct health service costs are taken into account. In the second analysis (analysis 2), we used a societal perspective that additionally takes into account education costs and the value of an IQ point itself.

For both analyses, we assumed that IQ follows the conventional normal distribution with a mean of 100 and

a standard deviation of 15. We used Z tables to calculate the reduction in the proportion of children in the lower IQ categories as a result of iodine supplementation (appendix p 9). The present value health and social services costs for the first 16 years of life are calculated for children with a mild neurodevelopmental impairment (IQ scores of 82–92); this is also calculated for children with a moderate or severe neurodevelopmental disability (IQ scores of ≤81), but with an additional cost of special education. These costs savings are then adjusted for survivors by use of UK life tables.⁴⁴

The monetary value of an IQ point (analysis 2 only) was identified by the systematic search done as part of this study (appendix p 2). Analyses 1 and 2 are both presented in terms of their disaggregated cost (in sterling) and outcomes in the form of a cost-consequence analysis, and IQ points gained are reported as natural units.

To investigate the robustness of the base case results, we did a comprehensive sensitivity analysis in which the main motivation was to further disadvantage the effect of iodine supplementation in the model and assess the effect on the results (appendix p 8). Most of the changes explored, for both analyses 1 and 2, were arbitrary where, for example, any gains as a result of iodine supplementation were halved and any detrimental effects as a result of the supplementation were doubled. A probabilistic sensitivity analysis was not appropriate in this case because we were already using estimates for the worst case scenario rather than using central mean estimates. We identified evidence on the benefits of iodine supplementation using systematic reviews and from experts on iodine status and supplementation in pregnant women. We identified three

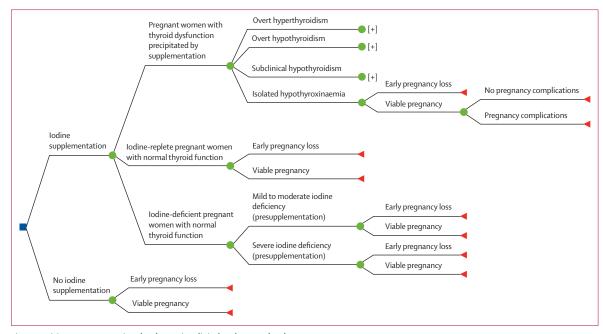


Figure: Decision tree representing the alternative clinical pathways taken by pregnant women The model is identical at every node ending in [+].

See Online for appendix

cohort studies of iodine status in pregnancy and IQ or cognitive development in infants and children⁴⁻⁶ and two trials of iodine supplementation in pregnant women.^{45,46}

Role of the funding source

There was no funding source for this study. All authors made the collective decision to submit for publication.

Results

Our systematic search identified 1361 published articles, of which eight studies⁴⁷⁻⁵⁴ passed quality criteria and were assessed to calculate the monetary value of an IQ point (appendix p 4). The quality criteria were as follows: an individual's IQ is used and is not a proxy; variables are

clearly specified; IQ measure follows a conventional

	Data	Sources	
Probability of a pregnant woman being iodine deficient (%)	67%	Bath and colleagues ⁵ ; Vanderpump and colleagues ¹ showed a similar proportion of iodine deficiency in 14–15 year old girls in the UK	
Iodine deficiency			
Proportion of iodine-deficient women who are mildly to moderately iodine deficient (%; urinary iodine to creatinine ratio [UIC] of 50–149 μg/L)	89%	Bath and colleagues ⁵	
Proportion of iodine-deficient women who are severely iodine deficient (UIC <50 µg/L)	11%	Bath and colleagues ⁵	
IQ gain			
IQ gain from supplementation in previously mildly iodine-deficient women	2.20	Bath and colleagues⁵	
IQ gain from supplementation in previously severely iodine-deficient women	3.00	Bath and colleagues ⁵	
lodine supplementation			
Duration of iodine supplementation with successful pregnancy and lactation (weeks)	78	Model assumption	
Duration of iodine supplementation with early pregnancy loss (weeks)	23	Model assumption	
Duration of iodine supplementation with late pregnancy loss (weeks)	47	Model assumption	
Pregnancy complications			
Baseline pregnancy risk of early pregnancy loss (%)	20%	Royal College of Obstetricians and Gynaecologists ¹	
Baseline pregnancy risk of stillbirth (%)	<1% of total births	UK stillbirth rate ²⁰	
Baseline pregnancy risk of preterm birth (%)	7% of livebirths	UK preterm birth rate ²¹	
Baseline pregnancy risk of pre-eclampsia (%)	8%	Duley ²²	
Pre-eclampsia cost per woman	£11370·00	Meads and colleagues ²³	
Discount rate for costs (%)	3.5%	NICE guide to the methods of technology appraisal ²⁴	
For a few women who might develop thyroid dysfunction as a result of iodine supp	lementation*		
Incremental incidence of thyroid dysfunction from iodine supplementation (%)	<1%	European Commission ²⁵	
IQ loss from overt and subclinical hypothyroidism	7.00	Haddow and colleagues ²⁶	
IQ loss from isolated hypothyroxinaemia	7.00	Model assumption based on equivalent neurodevelopmental test scores in subclinical hypothyroidism and isolated hypothyroxinaemia groups (Li and colleagues ²⁷)	
Incidence of early pregnancy loss from overt hyperthyroidism (%)	26%	Momotani and Ito ²⁸	
OR of stillbirth from overt hyperthyroidism†	8.42 (2.01-35.20)	Aggarawal and colleagues ²⁹	
OR of preterm birth from overt hyperthyroidism	16.50 (2.09–130.02)	Millar and colleagues ³⁰	
OR of pre-eclampsia from overt hyperthyroidism†	3.94 (2.47-6.29)	Aggarawal and colleagues ²⁹	
Incidence of early pregnancy loss from overt hypothyroidism (%)	30%	Glinoer ³¹	
OR for stillbirth from overt hypothyroidism	9.69 (2.92–32.16)	Allan and colleagues ³²	
OR for preterm birth from overt hypothyroidism	15.55 (3.62–66.81)	Ajmani and colleagues ³³	
Incidence of pre-eclampsia from overt hypothyroidism (%)	44%	Davis and colleagues ³⁴	
OR for early pregnancy loss from subclinical hypothyroidism	1.88 (1.13-3.15)	Wang and colleagues ³⁵	
OR of stillbirth from subclinical hypothyroidism	3.29 (1.32-8.21)	Allan and colleagues ³²	
OR for preterm birth from subclinical hypothyroidism	5.60 (2.30-13.58)	Ajmani and colleagues ³³	
OR for pre-eclampsia from subclinical hypothyroidism	3.39 (1.40-8.15)	Ajmani and colleagues ³³	
OR for preterm birth from isolated hypothyroxinaemia†	2.54 (1.42-4.54)	Korevaar and colleagues ³⁶	

Data are n (%), mean (SD, where available), or OR (95% CI). OR=odds ratio. *Assumption based on non-pregnant population iodine supplementation programmes that include elderly people. †Adjusted OR.

Table 1: List of variables assuming the worst case scenario (least favourable to iodine supplementation)

normal distribution with a mean of 100 and standard deviation of 15 or sufficient information is included in the study to allow the IQ measure's distribution to be converted into one (for cross study comparability); and the results reported in currency form have the applicable year stated.

Most of the studies valued an IQ point on the basis of its effect on an individual's income (appendix p 3). The issue of differences in scaling of IQ tests hindered the comparability across studies. The value of an IQ point, derived from the systematic search and applied to the unborn cohort, comes from the lifetime earnings premium of an additional IQ point. This is calculated to be \pounds 3297 (study estimates range from \pounds 1319 to \pounds 11967; after adjustment with life tables).

With the use of base case assumptions, the results of both analyses show that the iodine supplementation strategy was less costly and more effective compared with no supplementation (table 2). From the UK NHS perspective, where only health-related costs were taken into account (analysis 1), iodine supplementation was cost saving, with an expected positive net present value of \pounds 199 per pregnant woman (sensitivity analysis range $-\pounds$ 42 to \pounds 229) and an increase of an average 1·22 IQ points for the unborn infants. From a societal perspective (analysis 2), iodine supplementation was cost saving with an expected positive net present value of \pounds 4476 per pregnant woman (sensitivity analysis range \pounds 540 to \pounds 4495), and an average increase of 1·22 IQ points for the unborn infants.

For the pregnant women for whom thyroid dysfunction was precipitated by iodine supplementation, an average cost of more than \pounds 91000 each would have to be incurred to negate the overall benefits arising from the iodine-deficient pregnant women without thyroid dysfunction taking iodine supplementation in the model looking at the NHS perspective alone.

The sensitivity analysis supported the cost saving indication of base case results (table 2; appendix p 8). Iodine supplementation remained cost saving in all the sensitivity scenarios undertaken with one exception: in analysis 1 (health service perspective) where we assumed zero IQ gain for children of the previously mild to moderately iodine-deficient women, the results suggested an additional cost of \pounds 42 per pregnant woman for a gain of 0.17 IQ points for their offspring (table 2).

Discussion

To our knowledge, this study is the first to estimate the cost-effectiveness of a policy of iodine supplementation during pregnancy and lactation in a population with mild to moderate iodine deficiency. The analyses showed that iodine supplementation saved money and improved IQ. The results were supported by all the sensitivity analysis scenarios apart from the most extreme scenario, in which supplementation of mildly iodine-deficient pregnant women did not improve IQ. A key strength of the analysis was the use of very conservative assumptions to limit the benefits of iodine supplementation and potentially

Panel 1: Model assumptions relating to the women

- Women take daily iodine tablets 13 weeks before pregnancy, throughout pregnancy, and for an additional 26 weeks while breastfeeding. Cessation of iodine tablets occurs at the end of the lactation period or if there is a pregnancy loss event.
- Iodine supplementation will only benefit women who were iodine deficient before supplementation.
- All iodine-deficient women will be iodine replete with supplementation and they adhere to taking the daily supplementation.
- IQ gains will be different depending on the severity of iodine deficiency before supplementation; the presupplementation iodine-deficient women were subclassified into severe and mild or moderate iodine-deficient categories (table 1).
- All pregnancies are singleton.

Panel 2: Model assumptions relating to the pregnancy losses and complications

- Women with early pregnancy losses take daily tablets for 23 weeks on average and women with late pregnancy loss take tablets for 47 weeks on average (these both include the 13 weeks of taking the iodine tablets before pregnancy).
- 0.25%²⁵ of women of reproductive age have adverse thyroid dysfunction as a result of the iodine supplementation. Without existing evidence, the types of thyroid dysfunction precipitated by the iodine supplementation in the population are split evenly into the four following groups:
 - Overt hyperthyroidism (undetectable thyroid-stimulating hormone [TSH] with raised free T_4 [fT₄])
 - Overt hypothyroidism (increased TSH with low fT₄)
 - Subclinical hypothyroidism (increased TSH with normal fT₄)
 - Isolated hypothyroxinaemia (normal TSH with low fT₄).
- None of the thyroid dysfunctions were diagnosed and treated.
- An increased incidence of pregnancy losses and complications for each type of thyroid dysfunction is caused by iodine supplementation. Only pregnancy complications incurring significant costs (pre-eclampsia, preterm birth before 37 completed weeks of gestation, and stillbirth loss after 24 completed weeks of gestation) are used in the model.
- Although pre-eclampsia and preterm births often occur together, they are separate events in the model.
- The infants whose mothers had overt and subclinical hypothyroidism during
 pregnancy have losses of cognition of 7 IQ points.²⁶ On the basis of equivalent
 neurodevelopmental test scores,²⁷ this loss is also assumed for infants whose mothers
 had isolated hypothyroxinaemia during pregnancy. This assumption relates to the
 hypothyroidism that is induced by iodine supplementation in a few women and in this
 case is not related to iodine deficiency.
- Children of women with adverse thyroid dysfunction who had iodine supplementation and were iodine deficient before supplementation did not have an increase in IQ after supplementation.

overestimate adverse outcomes arising from supplementation. In view of the preliminary work for this study showing that iodine in pregnancy was unequivocally cost saving, the aim of our analysis was to explore the robustness of this indication by solely focusing on the worst case possible.

The monetary value of an IQ point used was also intentionally conservative and excluded voluntary work and any earnings that happen after the UK retirement age. These factors potentially act to underestimate the true monetary value associated with an additional IQ

	Cost saving analysis 1 from NHS perspective (£)	Cost saving analysis 2 from societal perspective (£)	IQ points gained	
Base case results	199	4476	1.22	
Sensitivity analysis scenarios				
IQ gain for severe iodine deficiency equal to mild to moderate deficiency	189	4302	1.18	
1 IQ point gain from iodine supplementation	46	1900	0.53	
No IQ gain for mild to moderate iodine deficiency	-42	540	0.17	
Prevalence of iodine deficiency halved	59	2178	0.61	
Doubled early pregnancy loss	145	3352	0.92	
Doubled cost of iodine tablets	148	4452	1.22	
Doubled cost discount rate (from 3.5% to 7%)	144	1608	1.22	
No thyroid dysfunction	229	4495	1.23	
Health costs savings halved (analysis 1 only)	60		1.22	
Value of an IQ point halved (analysis 2 only)		2409	1.22	
No real wage growth (analysis 2 only)		3239	1.22	
Willingness to pay figure* for an additional IQ point used (analysis 2 only)		1832	1.22	
Exclusion of public sector costs (analysis 2 only)		3953	1.22	
*Willingness to pay was used from a US based study ^{so} that used preference to elicit a figure from parents deciding on chelation therapy for their children.				

Table 2: Summary of base case results and and sensitivity analysis scenarios

point. The IQ earnings premium used in the model is based on an estimate from a US study⁴⁸ in which earnings came from the years 1974 and 1990. In today's technologically driven high skill economy, the earnings benefit from an additional IQ point might be more valuable for a worker than in previous decades.

Health and public sector costs relating to childhood neurological impairment were taken from a study⁴¹ that recorded incremental costs during a 1 year period in midchildhood (aged 11 years). The model assumed that these annual costs are the same for each year of childhood when in reality they are very likely to vary. Exclusion of these costs in the societal perspective sensitivity analysis did not change the direction of the results. No account has been made of public sector savings resulting from IQ improvement at the upper end of the IQ scale.

For the present value of lifetime earnings, a real wage growth of 1% per year in the future was assumed.⁴⁷ However, the sensitivity analysis also took into account a zero real wage growth scenario. This scenario did not prevent the iodine supplementation intervention from remaining cost saving.

This study has several limitations. First, one argument is that if most of the IQ gains will provide an absolute shift in the population IQ distribution, the relative IQ differences remain largely unchanged, negating most of the earnings advantages stemming from the gains in IQ points for workers. A possible response to this is that, generally, economies compete at a worldwide level and the addition of a more intelligent workforce in the future should help with productivity-linked earning gains. Second, some women might already take supplementation, which would mean that the overall modelled benefits might be overstated, but the analysis sought to identify the benefits of iodine supplementation to an individual compared with no supplementation.

We also used the sensitivity analysis to measure a person's willingness to pay for an additional IQ point instead of the monetary value of an IQ point derived from earnings. This analysis was done to allay any issues about using earnings as a basis for the value of an IQ point. However, when the monetary value of an IQ point is excluded (NHS perspective), the result showed that iodine supplementation was still cost saving.

A further possible weakness of our analysis is that iodine status is not identified at the individual level for the pregnant woman and, therefore, some wastage when iodine-sufficient women receive occurs supplementation. unnecessary However, iodine supplementation for iodine-replete pregnant women with normal thyroid function is not likely to cause any harm in most pregnancies. Despite our assumptions of harm in our model, no evidence thus far suggests that iodine supplementation induces thyroid dysfunction in pregnancy. Moreover, iodine is included in some proprietary pregnancy supplements. However, some evidence²⁵ of induction of thyroid dysfunction in the nonpregnant population does exist. Severe iodine deficiency has been associated with increases in pregnancy loss and complications; rectification of the iodine status of mildly to moderately iodine-deficient women is likely to decrease rather than increase pregnancy loss, although no studies have investigated this thus far. At present, no acceptable test for the assessment of individual iodine status exists; tests for assessment of population iodine status are available, but they require collection of urine from a large sample size, which is both cumbersome and costly. Furthermore, the testing needed for a targeted programme also causes delay, whereas the evidence suggests⁴⁵ that the benefits of iodine supplementation are increased with earlier treatment.

The limitations in our analysis relate to the limitations in the evidence. Although the evidence for the benefit of iodine supplementation in populations who are severely iodine deficient is clear.¹² the evidence of benefit in mildly iodine-deficient populations has not been established.³ Two of the three prospective studies^{45,46} of iodine supplementation in women from mildly iodine-deficient areas have shown improvements in child cognition, but these are limited by not being randomised studies, risk of bias, and small sample sizes. A large cohort study, with a short follow-up, provides evidence of potential harm.6 Maternal consumption of 150 µg per day or more of iodine from supplements was related to a 1.7 (95% CI 0.9-3.0) times higher risk of a child's mental scale score being less than 85 (derived from the Bayley Scales of Infant Development test), but this was not statistically significant. This study was not included in the model because the association was not statistically significant, the assessments were done at the age of 16 months, which is less robust than the later ages used in Bath and Hynes' study, and a third of the population used iodised salt. We based the IQ gains on an observational study, which was a cohort study comprising highly educated older women,⁵ which was a major limitation, but is the most robust information available in the absence of high quality experimental evidence. Although systematic reviews exist, ^{2,355,56} none of them provides an IQ change in a mildly iodine-deficient non-supplemented population.

The use of different IQ tests across studies raises the question of the comparability of findings. Bath and colleagues⁵ used the abbreviated form of the Wechsler Intelligence Scale for Children. Our base case estimate, taken from the study by Zax and Rees,⁴⁸ used the Henmon-Nelson Test of Mental Ability. Because these two intelligence tests are not perfectly correlated, an individual might get a slightly different IQ score from each test. We explored this possibility in the sensitivity analysis by modification of IQ gains and the results remained cost saving with one exception, which was based on an extreme scenario.

In the absence of randomised controlled trial evidence, our model results strengthen the case for universal iodine supplementation pre-conception and during pregnancy and lactation in mildly to moderately iodinedeficient populations. In our study, we only took into account iodine supplementation in tablet form. Fortification of food with iodine is another way of attaining iodine sufficiency; however, food fortification alone might not be enough to achieve iodine sufficiency for pregnant women.⁵⁷

Our findings have important implications worldwide. 32 countries have mild or moderate iodine deficiency identified from surveys of the iodine status of school-age children.⁵⁸ These countries have a population of 1.88 billion people and 241 million school-age children, so the potential effects of introduction of iodine supplementation for pregnant women could be substantial. The use of urinary iodine of school-age children to estimate the iodine status of pregnant women is likely to underestimate the prevalence of iodine deficiency during pregnancy.^{59,60} Reduced urinary iodine concentrations have been identified in pregnant women compared with school aged children, possibly because of an increased consumption of milk in children.⁵⁹ Additionally, pregnant women have an increased iodine requirement.⁶¹

A randomised controlled trial would provide the most robust evidence on which to base policy; however, such a study would need costly child developmental assessments. A randomised controlled trial with iodinedeficient pregnant women taking placebo iodine tablets has been described as unethical⁹ because iodine supplementation in pregnancy is already recommended by many national and international bodies, including WHO.⁸ On the basis of the best available evidence, this study emphasises the cost-effectiveness of an iodine supplementation strategy for pregnant women in the UK. Our findings are applicable to any country which is mildly to moderately iodine deficient, but has no universal salt or other iodisation programme.

Contributors

TER and KJ conceived and designed the study. KJ was responsible for the groundwork that led to the study being planned. KJ, KB, and SC provided clinical data used in the model-based analysis and clinical interpretation. MM did the scientific literature review, the model-based analysis, and interpreted the results under the supervision of TER. PB provided advice on use of data and modelling techniques. MM and TER wrote the first draft of the report. All authors provided input on the manuscript draft. TER is the guarantor for the study.

Declaration of interests

KJ is a member of the UK Iodine Group. All other authors declare no competing interests.

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References

- Pharoah POD, Buttfield IH, Hetzel BS. Neurological damage to the fetus resulting from severe iodine deficiency during pregnancy. *Lancet* 1971; 297: 308–10.
- P. Qian M, Wang D, Watkins WE, et al. The effects of iodine on intelligence in children: a meta-analysis of studies conducted in China. Asia Pac J Clin Nutr 2005; 14: 32–42.
- 3 Taylor PN, Okosieme OE, Dayan CM, Lazarus JH. Impact of iodine supplementation in mild-to-moderate iodine deficiency: systematic review and meta-analysis. *Eur J Endocrinol* 2013; **170**: R1–R15.
- 4 Hynes KL, Otahal P, Hay I, Burgess JR. Mild iodine deficiency during pregnancy is associated with reduced educational outcomes in the offspring: 9-year follow-up of the gestational iodine cohort. J Clin Endocrinol Metab 2013; 98: 1954–62.
- 5 Bath SC, Steer CD, Golding J, Emmett P, Rayman MP. Effect of inadequate iodine status in UK pregnant women on cognitive outcomes in their children: results from the Avon Longitudinal Study of Parents and Children (ALSPAC). *Lancet* 2013; 382: 331–17.
- 6 Rebagliato M, Murcia M, Álvarez-Pedrerol M, et al. Iodine supplementation during pregnancy and infant neuropsychological development INMA mother and child cohort study. *Am J Epidemiol* 2013; 177: 944–53.
- 7 Vanderpump MP, Lazarus JH, Smyth PP, et al. Iodine status of UK schoolgirls: a cross-sectional survey. *Lancet* 2011; 377: 2007–12.
- 8 WHO. Assessment of iodine deficiency disorders and monitoring their elimination: a guide for programme managers. 2007. http:// apps.who.int/iris/handle/10665/43781 (accessed May 1, 2014).
- 9 Pearce EN. Effects of iodine deficiency in pregnancy. J Trace Elem Med Biol 2012; 26: 131–33.
- 10 Whalley LJ, Deary IJ. Longitudinal cohort study of childhood IQ and survival up to age 76. *BMJ* 2001; **322**: 819.
- 11 Gunnell D, Magnusson PK, Rasmussen F. Low intelligence test scores in 18 year old men and risk of suicide: cohort study. BMJ 2005; 330: 167.
- 12 Batty GD, Mortensen EL, Osler M. Childhood IQ in relation to later psychiatric disorder Evidence from a Danish birth cohort study. *Br J Psychiatry*. 2005; 187: 180–1.
- 13 Koenen K, Moffitt T, Roberts A, et al. Childhood IQ and adult mental disorders: a test of the cognitive reserve hypothesis. *Am J Psychiatry* 2009; 166: 50–57
- 14 Batty GD, Mortensen EL, Nybo Andersen AM, Osler M. Childhood intelligence in relation to adult coronary heart disease and stroke risk: evidence from a Danish birth cohort study. *Paediatr Perinat Epidemiol* 2005; 19: 452–59.
- 15 Lawlor DA, Batty GD, Clark H, McIntyre S, Leon DA. Association of childhood intelligence with risk of coronary heart disease and stroke: findings from the aberdeen children of the 1950s cohort study. *Eur J Epidemiol* 2008; 23: 695–706.

- 16 Kilgour AH, Starr JM, Whalley LJ. Associations between childhood intelligence (IQ), adult morbidity and mortality. *Maturitas* 2010; 65: 98–105.
- 17 Fergusson DM, John Horwood L, Ridder EM. Show me the child at seven II: childhood intelligence and later outcomes in adolescence and young adulthood. J Child Psychol Psychiatry 2005; 46: 850–58.
- 18 Vanderpump MP, Lazarus JH, Smyth PP, et al. Iodine status of UK schoolgirls: a cross-sectional survey. *Lancet* 2011; 377: 2007–12.
- 19 Royal College of Obstetricians and Gynaecologists. Recurrent and late miscarriage: tests and treatment of couples 2012. Nov 19, 2014. https://www.rcog.org.uk/globalassets/documents/patients/patientinformation-leaflets/pregnancy/recurrent-and-late-miscarriage.pdf (accessed Nov 19, 2014).
- 20 Office for National Statistics. Birth summary tables, England and Wales, 2013. 2014. http://www.ons.gov.uk/ons/rel/vsob1/birthsummary-tables-england-and-wales/2013/index.html (accessed July 16, 2014).
- 21 Office for National Statistics. Gestation-specific infant mortality in England and Wales, 2011. 2013. http://www.ons.gov.uk/ons/rel/childhealth/gestation-specific-infant-mortality-in-england-and-wales/2011/ gest-spec-bulletin-2011.html (accessed July 15, 2014).
- 22 Duley L. The global impact of preeclampsia and eclampsia. Semin Perinatol 2009; **33**: 130–37.
- 23 Meads CA, Cnossen J, Meher S, et al. Methods of prediction and prevention of pre-eclampsia: systematic reviews of accuracy and effectiveness literature with economic modelling. *Health Technol Assess* 2008; 12: iii–iv.
- 24 NICE. Guide to the methods of technology appraisal. London: National Institute for Health and Care Excellence, 2013.
- 25 European Commission. Opinion of the Scientific Committee on Food on the tolerable upper intake level of iodine. Brussels: European Commission, 2002.
- 26 Haddow JE, Palomaki GE, Allan WC, et al. Maternal thyroid deficiency during pregnancy and subsequent neuropsychological development of the child. N Engl J Med 1999; 341: 549–55.
- 27 Li Y, Shan Z, Teng W, et al. Abnormalities of maternal thyroid function during pregnancy affect neuropsychological development of their children at 25–30 months. *Clin Endocrinol (Oxf)* 2010; 72: 825–29.
- 28 Momotani N, Ito K. Treatment of pregnant patients with Basedow's disease. Exp Clin Endocrinol 1991; 97: 268–74.
- 29 Aggarawal N, Suri V, Singla R, et al. Pregnancy outcome in hyperthyroidism: a case control study. *Gynecol Obstet Invest* 2014; 77: 94–99.
- 30 Millar LK, Wing DA, Leung AS, Koonings PP, Montoro MN, Mestman JH. Low birth weight and preeclampsia in pregnancies complicated by hyperthyroidism. *Obstet Gynecol* 1994; 84: 946–99.
- 31 Glinoer D. The thyroid in pregnancy: a European perspective. *Thyroid Today* 1995; 18: 1–11.
- 32 Allan WC, Haddow JE, Palomaki GE, et al. Maternal thyroid deficiency and pregnancy complications: implications for population screening. J Med Screen 2000; 7: 127–30.
- 33 Ajmani SN, Aggarwal D, Bhatia P, Sharma M, Sarabhai V, Paul M. Prevalence of overt and subclinical thyroid dysfunction among pregnant women and its effect on maternal and fetal outcome. J Obstet Gynaecol India 2014; 64: 105–10.
- 34 Davis LE, Leveno KJ, Cunningham FG. Hypothyroidism complicating pregnancy. *Obstet Gynecol* 1988; **72**: 108–12.
- 35 Wang S, Teng W, Li J, Wang W, Shan Z. Effects of maternal subclinical hypothyroidism on obstetrical outcomes during early pregnancy. J Endocrinol Invest 2012; 35: 322–25.
- 36 Korevaar TI, Schalekamp-Timmermans S, de Rijke YB, et al. Hypothyroxinemia and TPO-antibody positivity are risk factors for premature delivery: the generation R study. J Clin Endocrinol Metab 2013; 98: 4382–90.
- 37 MoneySuperMarket. Supermarket price comparison of vitabiotics pregnacare original vitamins 2014. May 14, 2014]. http://www. mysupermarket.co.uk/grocery-categories/vitamins_and_ supplements_for_women_in_asda.html (accessed May 14, 2014).
- 38 Stagnaro-Green A, Abalovich M, Alexander E, et al. Guidelines of the American Thyroid Association for the diagnosis and management of thyroid disease during pregnancy and postpartum. *Thyroid* 2011; 21: 1081–125.

- 39 Lazarus J, Brown RS, Daumerie C, Hubalewska-Dydejczyk A, Negro R, Vaidya B. 2014 European Thyroid Association Guidelines for the Management of Subclinical Hypothyroidism in Pregnancy and in Children. *Eur Thyroid J* 2014; **3**: 76–94.
- 40 Petrou S, Johnson S, Wolke D, Marlow N. The association between neurodevelopmental disability and economic outcomes during mid-childhood. *Child Care Health Dev* 2013; 39: 345–57.
- Mangham LJ, Petrou S, Doyle LW, Draper ES, Marlow N. The cost of preterm birth throughout childhood in England and Wales. *Pediatrics* 2009; **123**: e312–27.
- 42 Mistry H, Heazell AE, Vincent O, Roberts T. A structured review and exploration of the healthcare costs associated with stillbirth and a subsequent pregnancy in England and Wales. *BMC Pregnancy Childbirth* 2013; **13**: 236.
- 43 Office for National Statistics. Consumer Price Inflation, December 2013. Feb 15, 2014. http://www.ons.gov.uk/ons/rel/cpi/consumerprice-indices/december-2013/stb---consumer-price-indices--december-2013.html (accessed Feb 15, 2014).
- 44 Office for National Statistics. United Kingdom national life tables: 1980–1982 to 2011–2013. 2014. http://www.ons.gov.uk/ons/rel/ lifetables/national-life-tables/2011-2013/stb-uk-2011-2013.html (accessed Oct 10, 2014).
- 45 Berbel P, Mestre JL, Santamaria A et al. Delayed neurobehavioral development in children born to pregnant women with mild hypothyroxinemia during the first month of gestation: the importance of early iodine supplementation. *Thyroid* 2009; **19**: 511–19.
- 46 Velasco I, Carreira M, Santiago P, et al. Effect of iodine prophylaxis during pregnancy on neurocognitive development of children during the first two years of life. J Clin Endocrinol Metab 2009; 94: 3234–41.
- 47 Schwartz J. Societal benefits of reducing lead exposure. *Environ Res* 1994; 66: 105–24.
- 48 Zax JS, Rees DI. IQ, Academic performance, environment, and earnings. *Rev Econ Stat* 2002; 84: 600–16.
- 49 Fletcher J. Friends or Family? Revisiting the effects of high school popularity on adult earnings. 2013. National Bureau of Economic Research Working Papers: 19232. http://www.nber.org/papers/ w19232 (Sept 4, 2014).
- 50 Lutter RW. Valuing children's health: a reassessment of the benefits of lower lead levels. AEI—Brookings Joint Center Working Paper No. 00-02. 2000. http://papers.ssrn.com/sol3/papers.cfm?abstract_ id=243537 (Feb 11, 2014).
- 51 Mueller G, Plug E. Estimating the effect of personality on male and female earnings. *Ind Lab Relat Rev* 2006; **60**: 3–22.
- 52 Salkever DS. Updated estimates of earnings benefits from reduced exposure of children to environmental lead. *Environ Res* 1995; **70**: 1–6.
- 53 de Wolff P, van Slijpe ARD. The relation between income, intelligence, education and social background. *Europ Econ Rev* 1973; 4: 235–64.
- 54 Zagorsky JL. Do you have to be smart to be rich? The impact of IQ on wealth, income and financial distress. *Intelligence* 2007; 35: 489–501.
- 55 Bougma K, Aboud FE, Harding KB, Marquis GS. Iodine and mental development of children 5 years old and under: a systematic review and meta-analysis. *Nutrients* 2013; 5: 1384–416.
- 56 Zhou SJ, Anderson AJ, Gibson RA, Makrides M. Effect of iodine supplementation in pregnancy on child development and other clinical outcomes: a systematic review of randomized controlled trials. *Am J Clin Nutr* 2013; **98**: 1241–54.
- 57 Clifton V, Hodyl N, Fogarty P, et al. The impact of iodine supplementation and bread fortification on urinary iodine concentrations in a mildly iodine deficient population of pregnant women in South Australia. *Nutr J* 2013; 12: 32.
- 58 Andersson M, Karumbunathan V, Zimmermann MB. Global iodine status in 2011 and trends over the past decade. J Nutr 2012; 142: 744–50.
- 59 Gowachirapant S, Winichagoon P, Wyss L, et al. Urinary iodine concentrations indicate iodine deficiency in pregnant Thai women but iodine sufficiency in their school-aged children. J Nutr 2009; 139: 1169–72.
- 60 Wong EM, Sullivan KM, Perrine CG, Rogers LM, Peña-Rosas JP. Comparison of median urinary iodine concentration as an indicator of iodine status among pregnant women, school-age children, and nonpregnant women. *Food Nutr Bull* 2011; 32: 206–12.
- 61 Zimmermann MB, Jooste PL, Pandav CS. Iodine-deficiency disorders. *Lancet* 2008; **372**: 1251–62.