

from Mr Nick Hunt
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on behalf of all the signatories,

30 August 2026

Dear <MP or Peer (see distribution list at foot of letter)>

Re: Mandatory Fortification of Flour with Folic Acid

We are a group of medical and safety management professionals with grave concerns about the newly mandated addition of folic acid to flour. We believe that as a Parliamentarian who qualified with a medical degree, you are best placed to understand the issues involved. The claims of benefit are not as clear cut as many of us were taught in medical school and the evidence that risks outweigh benefits has grown to be substantial.

The addition of synthetic folic acid to flour from 13 December 2026 became law in the UK via SI2024/1162, an amendment to the Bread and Flour Regulations 1998. As a Statutory Instrument, there was no parliamentary debate. There was also no full, scientific, Impact Assessment; just a *de minimis* financial assessment by DEFRA with limited DHSC input. The stated aim of the mandate is to increase blood folate levels in pregnant women in order to reduce the risk of neural tube defects (NTDs). The Government estimates the policy will result in a reduction of about 200 NTD-affected pregnancies annually, approximately 20% of UK cases.

However, behind this headline are some serious risks that were not presented to Parliamentarians when SI2024/1162 was laid. We explain these below and would be grateful to know your views. If you share our concerns about the seriousness of this issue, we ask for your support in ensuring a proper parliamentary discussion and debate.

The Hungarian Folic Acid Trial

The strongest evidence for a reduction in NTDs comes from animal work and randomised controlled trials using high doses of folic acid in high-risk women. However, there was also one randomised controlled trial using moderate doses in healthy women: the Hungarian Periconceptional Supplementation Trial (Czeizel & Dundas, 1992)¹. In this trial *“Women planning a pregnancy (in most cases their first) were randomly assigned to receive a single tablet of a **vitamin supplement (containing 12 vitamins, including 0.8 mg of folic acid; 4 minerals; and 3 trace elements)** or a **trace-element supplement (containing copper, manganese, zinc, and a very low dose of vitamin C)** daily for at least one month before conception and until the date of the second missed menstrual period or later.”*

The trial found 0 NTDs in the multivitamin (folic acid) arm versus 6 NTDs in the control arm. These results led to public health authorities recommending folic acid for pregnant women and declaring that further research would be unethical.

¹ <https://www.nejm.org/doi/10.1056/NEJM199212243272602>

However, it is less well known that the same researchers published a safety analysis² two years later, in 1994, which revealed **70 excess fetal deaths (mainly miscarriages) in the supplemented arm**. This paper is behind a paywall, and the abstract does not mention the excess deaths, but Table 2 of the full report shows this data clearly, as shown below:

Table 2. Intention-to-treat analysis of pregnancy outcomes

Evaluated pregnancies ended in	Multivitamin		Trace element	
	No.	%	No.	%
Termination				
First trimester	6	0.2	6	0.2
Second trimester after diagnosis of fetal defect	3	0.1	13	0.5
Fetal death				
Chemical pregnancy	55	2.0	40	1.5
Ectopic pregnancy	7	0.2	4	0.2
Miscarriage	301	10.8	251	9.4
Stillbirth	11 ^a	0.4	9	0.3
Subtotal	374	13.4	304	11.4
Livebirth	2410	86.3	2337	87.9
Total	2793	100.0	2660	100.0

^a Three stillbirths occurred in twin pregnancies in which the other twin was liveborn

Table 2 | from Czeizel, Dudás & Métneki (1994), intention-to-treat analysis.⁽⁸⁾

Therefore, for a pregnant woman, despite the 1 in 450 possibility of NTD benefit, there was also a 1 in 50 risk of pregnancy loss. **This translates as 9 pregnancies lost for every 1 NTD prevented.** Moreover, the authors noted that **all the claimed NTD benefit might have resulted from prenatal death of affected fetuses.**

These losses have been explained away as "good deaths" i.e. pregnancies that folic acid sustained just long enough for a loss that would have passed unnoticed to become a recognised miscarriage. This ignores the effect of a pregnancy loss on the parents and does not fit with the data. If folic acid were carrying early conceptions further, the earliest losses (the chemical pregnancies) should have also fallen. However, they rose. This safety analysis has consistently been ignored, and pregnant women have not been informed of the risks of folic acid supplementation, only its claimed benefits.

Risks to Human Health from Synthetic Folic Acid

Synthetic folic acid is not the same as natural folate (found in foods like leafy greens and citrus fruits). It is a manufactured molecule that is converted in the body by an inefficient enzyme into usable folate. This takes many hours and even days at higher doses. The risk to human health arises from circulating, unmetabolised, folic acid, which is associated with a range of harms. For example, it acts as a partial agonist at the folate receptor, blocking actual folate from binding and being absorbed into the cells (including the brain), thereby making any folate deficiency worse.

² <https://pubmed.ncbi.nlm.nih.gov/7979565/>

The NHS has long warned³ of the risks of folic acid to certain groups; including **anyone with cancer, a coronary stent, B12 deficiency (particularly if undiagnosed), or a previous allergic reaction to it**, who are all advised by the NHS to completely avoid folic acid supplementation. This fact alone should preclude it from being imposed on the whole public as a mass, uncontrolled medication through fortification of flour. Flour is a staple food which is also hidden in many processed foods, so it will be almost impossible to avoid exposure if you have any of these conditions.

There is also **strong evidence that folic acid increases the risk of cancer⁴, with an overall increase of 7%** (RR 1.07 [95% CI 1.00 to 1.14]) reported in a meta-analysis of 10 randomised trials. Prostate cancer showed an increase of 24% (RR1.24 [1.03-1.49]).

Folic Acid Fortification is Unethical and Unscientific

Prior to SI2024/1162, the UK had rejected folic acid fortification of flour for nearly 30 years, and it continues to be rejected by nearly every other Western European country. We are gravely concerned that the UK is now speeding, without proper parliamentary scrutiny of serious health risks, towards mandatory folic acid supplementation of flour which will, by implication, be in all food products containing flour (including pasta, cakes, pizza, burgers, battered fish, biscuits, soup, pies, ready meals, baby food, breakfast cereals).

This is unethical mass medication, without consent, of the entire population (including men, postmenopausal women and children who have no possibility of any benefit) for a small potential benefit to reduce NTDs in a very small and specific group. **It is not scientific or wise**, as it ignores the disproportionately higher risk of fetal death than NTD benefit in pregnant women taking folic acid, as well as the health risks to the wider population from circulating unmetabolised folic acid.

How are those whom the NHS warns about folic acid supposed to avoid it? Or anyone else who does not want to ingest it due to health concerns e.g. its potential for cancer promotion?

Please let us know your views and whether you will press for parliamentary debate about this. If you are willing, we would welcome a meeting or further discussion.

Yours sincerely

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³ <http://www.nhs.uk/medicines/folic-acid/>

⁴ <https://bmjopen.bmj.com/content/2/1/e000653>

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