Your ref: PO-1286639
Guy Opperman MP ref: GO4464

By signed for Post

NOTICE. 21 August 2021

To: Nadhim Zahawi MP

Minister for Business and Industry
Minister for COVID Vaccine Deployment
Department for Business, Energy & Industrial Strategy
Department of Health & Social Care
39 Victoria Street
London SW1H 0EU

From: Tim Coulter Architect

52A Painshawfield Road
Stocksfield, Northumberland
NE43 7QY

Strictly for the personal attention of Nadhim Zahawi concerning:

My letter dated 9 December 2020 forwarded by Guy Opperman resulting in Nadhim Zahawi's reply dated 22 April 2021

Claims:

Thank you for your letter dated 22 April 2021 outlining the following two claims.

1. Your paragraph 3: "We have now accepted the recommendation from the independent Medicine and Healthcare products Regulatory Agency (MRHA) to authorise three COVID-19 vaccines for use, including Pfizer/BioNTech, Oxford/AstraZeneca, and Moderna. This follows months of rigorous clinical trials and a thorough analysis of the data by experts at the MHRA, who have concluded that all three vaccines met the regulator's strict standards of safety, quality, and effectiveness".
2. Your paragraph 4: "The UK currently operates a system of informed consent for vaccinations. Our objective is to vaccinate as many people as possible, in line with the advice of the Joint

Committee on Vaccination and Immunisation. We will continue to provide clear information to the public, encouraging people to seek NHS's advice so that they have the right information to make an informed choice".

Please provide the documentation by the experts at the MHRA concluding that all three vaccines met the regulator's strict standards of safety, quality and effectiveness. Please provide these these standards and the test results showing that these products meet these standards.

Please provide the documentation that prior to vaccination that each vaccine recipient receives the information to make an informed consent. These include the following:

- The vaccine insert.
- The contents of the vaccine to be used.
- How it works.
- That there is no liability clause in the vaccination agreement.
- That the vaccine manufacturers do not claim their products stop infection or transmission and the public are now informed of the resulting breakthrough cases.
- That the manufacturer or the person carrying out the injection (Doctors, Nurses, and others) do not carry any liability whatsoever for damage to individuals.
- That they are informed that before commencement of the vaccination programme that no studies were carried out on: animals, pregnant women, children, people with co-morbities and the elderly.
- That they are informed of all the possible side effects so far identified, including death, micro and macro blood clotting in many parts of the body causing heart attacks, strokes, lung problems, myocarditis, Guillaim Barre's syndrome, Bell's Palsy and any other side effects identified to date.
- That they are informed that these products use experimental technology where there have been no long term safety studies that might identify all possible side effects. These include studies of toxicity, pharmacokinetics (to what part of the body does the injected material spread) and pharmacodynamics (what does the injected material do when it gets there).
- That they are now informed that these products (injected gene therapy treatments) have produced more adverse reactions worldwide than any other recent vaccination programme.
- That due to the accelerated release of the experimental products there are no long term studies of suspected side effects such as genetic damage, autoimmune diseases, cancer, sterility in men and women, antibody dependant enhancement and other as yet unknown damage.
- That the risk of death for anyone under the age of 60 from Covid 19 is minuscule.

- That there are several published early home treatment protocols which became known as early as April 2020. It is reported they can reduce hospitalisation and death by up to 85% which for people under and over 60 who become infected, can produce robust natural immunity.

Please provide a sworn affadavit supporting your above claims in 1 & 2 numbered above, backed by your commercial liability of a minimum of £2,000,000.

Please supply your reply to this **Notice** within 14 days of the above date.

Failure to reply will mean that I will conclude that you agree with me :

A. That you have no documentation supporting your above claims.

B. That all three vaccines fail to meet the regulator's strict standards of safety, quality and effectiveness.

C. That the public has not been informed concerning the risks involved and are unable to make an informed consent.

D. That so called vaccination is generally by coercion.

E. That you are spreading false information.

F. That you should as Minister responsible in Parliament forthwith shut down the Government's so called vaccination programme.

I enclose a copy of your letter of 22 April 2021 for your records. This contains your claims.

Signed

Tim Coulter Architect