

From the Commissioner Anthony Barrowclough QC

Our ref: C.65/88

The Rt Hon Dr David Owen MP House of Commons London SWIA OAA Parliamentary Commissioner for Administration Church House Great Smith Street London SW1P 3BW

01-212 0190/ GRO-C

/4 March 1988

Thank you for your letter of 2 March giving me further information about the

case of your constituent, Mr **Andrew Security** whose complaint against the Departmen of Health and Social Security (DHSS) you have asked me to investigate.

I am grateful for the additional details you have provided, and I note that Mr **Mr** - for whom I have the very greatest sympathy - was aware of your intention to refer the matter to me. But I am afraid I remain very doubtful Common whether the facts presented provide me with a sufficient basis for embarking on an investigation. For that I would need not only <u>prima facie</u> evidence of maladministration on DHSS's part but also evidence that any maladministration which there may have been was relevant in terms of causation and foreseeability to the injustice suffered by Mr **Mr** (or other haemophiliacs who need to **M** (or other haemophiliacs who need to **M** (or **C**).

I am at something of a disadvantage in assessing the prima facie merits of the transformer of the decision you announced (in your Written Answer to Mr George Cunningham decision you announced (in your Written Answer to Mr George Cunningham decision of 22 January 1975) regarding intended self-sufficiency as soon as practicable with the domestic production of Factor VIII. But on the face of things, the purpose of the domestic production envisaged in 1975 was to secure the best available treatment for haemophiliacs at a lower cost than was involved in using imported products. If discern a similar emphasis in the Answers you gave (on 22 April 1975) to the best decision was made to relative contamination risks in any of the Answers concerned (or, so far the domestic decision decision to Parliament by the government of your decision decision to achieve early self-sufficiency.

In rather the same vein (and we are concerned here, of course, with the period that the before the HIV virus was known about) both imported and UK-produced blood products would presumably have been tested for all known blood infections before being he put on the market. Does it not follow - given the state of medical knowledge that the time - that the UK product would have been seen as carrying neither a GRO-A lesser nor greater risk than that of imported Factor VIII? After discovery were not will be the HIV virus, both would surely be tested to ensure that they were not will be the time were not will be the time at the time are also to ask myself, in relation to your whether that it was imported Factor VIII which infected Mr **Wereme** in the first place.

I do not doubt that you will find these further comments disappointing. I hope, which however, that they will help to explain to your constituent why - as things stand - I do not see his case as one which I can take up. If there is additional

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evidence which you or he wish to submit and which you feel might lead me to a different conclusion, I shall of course be happy to consider it. (If you do send me any further material, perhaps you would be kind enough to include a copy of the letter which you told me you had sent to the Secretary of State last November, and of the reply he sent you in February?.)

your in an GRO-C