Freedom of Information Request	FOI 22-274	5th July 2022

1. **Firstly, I want them to provide a list of all staff who have disclosed links to the medical devices manufacturer Philips and what those links are, whether it be employment, if they have accepted benefits from them, if they have been involved in their events etc - going back as far as you can.**
   Please find attached the full register of Declarations of interest for the Health Board. Please note the Board Member list is also available on the Health Boards web pages.

2. **Secondly, can you confirm how many patients you have identified in your Health Board who have experienced adverse affects from the Philips CPAP devices, both prior to the Philips CPAP recall of June 2021 and after**
   The Health Board has not logged or reported any reportable Adverse Incidents involving Philips CPAP ventilators to date.

3. **as well as tell me if you have made or received any Yellow Card reports in relation to this matter.**
   The Health Board does not receive Yellow Card reports directly. Yellow Card reporting is the method by which both individuals and representatives from Health Institutions report specific concerns involving a Medical Device, or a Medicinal Product to the Medicines and Healthcare products Regulatory Agency (MHRA). The outcome of the MHRA's investigations, which can be prompted by it receiving Yellow Card reports, alongside other methods, are published as Safety Alerts and Notices. The Health Board has acted on all such notices relating to Philips CPAP ventilators to date. You may wish to request this information from the Medicines and Healthcare products Regulatory Agency (MHRA).