Ask first - shoot later!

Questions to ask the pharmacist when seeking to give informed consent before receiving a Covid booster…

1) Can you please point me in the direction of the research that supports this current booster - alone and in combination with a flu shot? I am looking for long term studies in animals and humans showing the necessity, efficacy and safety of these products.

2) Why are so many of the known adverse events of this injection NOT included on the consent form that needs to be signed prior to injection? Have you seen https://dailyclout.io/home/#pfizer-reports ?

3) Assuming human trials were done on this version of the booster, what before and after follow up was done on the trial subjects? Since MRIs, D-Dimer tests and antibody profiles have pointed to vaccine damage in recipients of earlier versions of this product, was this type of follow up done on trial subjects to ensure they had no effects of this newer version?

4) Given that we are no longer in an emergency situation, the terms of the interim orders no longer apply. Has Health Canada now reverted to its former vaccine approval process (involving three independent placebo controlled trials with a combined total of at least 9000 subjects)? If not, why not?

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