*Questions continued…*

5) Have you personally made an effort to locate the data on the bio distribution, toxicology, reproductive toxicity, etc. of this (or any previous) Covid-19 products you may have administered? If not, who in your professional organization is known to have accessed this data? Is this data age-stratified? Does the data differentiate between those with zero, one, two, three or more previous injections to see if potential adverse events are dose dependent? Or is it known that this data does not exist for these products?

6) Are you aware of patients who have had long term disabilities, increased cancer rates or other adverse effects with an onset weeks or months after previous Covid-19 injections? What treatment protocols have you been able to provide for them? How well have these protocols helped mitigate the symptoms? If I were to also experience adverse events, do you have the expertise to take on my care?

*If your physician or your pharmacist cannot provide answers to these basic questions, it is not possible for you to provide the necessary informed consent for these injections.*

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