HB248 Witness Testimony for Cristy Berg

My name is Cristy Berg and I'm a VP of Communications for a medical device company. I strongly support House Bill 248. To those paying attention, COVID-19 has been an entirely created pandemic narrative built on false-positive tests and inflated case counts. A huge portion of the population are not even at risk, most especially, our children. There is a 99.9x survival rate AND proven treatments which are being withheld from doctors and therefore the public. I ask you WHY? The CDC continues to manipulate data that props-up "vaccine" effectiveness, hiding breakthrough cases and blaming the unvaccinated for outbreaks. Where is the study that proves this? Where is the data? FACT: There were no trials – other than animal trials which were stopped because they all died. There is no human data.

Recently, a study analyzed VAERS data (a resource most American's don't even know exists) and showed the COVID shots are likely cause of deaths, spontaneous abortions, cardiovascular, neurological and immunological adverse events. This is causing great and unnecessary harm to innocent people! The shot is being inaccurately marketed as a vaccine, but does not fit the definition of a vaccine. It is a shot – more like a bioweapon – that is proving to do nothing to protect those that have taken it! We are now using celebrity, illegal bribes and peer pressure to force people to comply which is only serving to feed fear and divide our country. When did it become ok to ask someone about their personal health choices? When have the vaccinated ever been so fearful of others post-vaccination? Why don't they feel protected? Why are they still so afraid? Why do we need to point out who is and who is not? What is going on here is an obvious and deliberate act of deception.

We went from randomized placebo-controlled studies right to what we have now: the population <u>almost</u> being *forced* inoculated – that is not a randomized placebo-controlled trial, the gold-standard for FDA approval. Pfizer has applied with the FDA for a full-on approval. How can they do that if they have not completed the trials for efficacy and safety? They stopped those parts of the trial and they are just giving the shot to everyone! How and why would the FDA accept this data? Where is the priority to safety?

Thank you for the opportunity to provide you with feedback on the need for and urgency of House Bill 248.