From: **Paul** paulmozina@wi.rr.com
Subject: Fwd: CCFN 200043 Communication from the Milwaukee Health Department providing an update on its efforts responding to the COVID-19 pandemic.
Date: November 11, 2021 at 11:38 AM
To: ashanti Hamilton ahamil@milwaukee.gov, Murphy, Michael (Alderman) mmurph@milwaukee.gov, Bauman, Robert rjbauma@milwaukee.gov, Nikiya Dodd ald05@milwaukee.gov, scott.spiker@milwaukee.gov, Johnson, Cavalier Cavalier.Johnson2@milwaukee.gov, Perez, Jose JoseG.Perez@milwaukee.gov, JoCasta.Zamarripa@milwaukee.gov, Kovac, Nik nkovac@milwaukee.gov, Stamper II, Russell russell.stamperii@milwaukee.gov, Coggs, Milele mcoggs@milwaukee.gov, Dimitrijevic, Marina Marina@milwaukee.gov, Lewis, Chantia Chantia.Lewis@milwaukee.gov, Borkowski, Mark Mark.Borkowski@milwaukee.gov, Rainey, Khalif Khalif.Rainey@milwaukee.gov, mayor@milwaukee.gov

For the record, I notified the Public Safety and Health Committee that the MHD is violating the requirements of the Emergency Use Authorization for the Pfizer COVID-19 vaccines.

- The MHD is not providing informed consent at their vaccine clinics and thereby recklessly endangering naive potential vaccine recipients.
- The MHD is not informing potential vaccine recipients that, as the the “Vaccination Provider”, the MHD is responsible for reporting any adverse events that they may experience to the Vaccine Adverse Event Reporting System (VAERS), and that, therefore, they should notify the MHD if they experience any adverse events.
- The MHD is not alerting potential vaccine recipients of the Serious Adverse Events that have been reported to VAERS (and explicitly mentioned in the Pfizer Fact Sheet) and that they are at risk for the same.

You, and the MHD, are free to ignore the MANDATORY REQUIREMENTS imposed on those who administer vaccines under emergency use authorization (and you may get away with it), but you are not free to ignore the moral responsibility that accompanies those who choose to remain willfully ignorant, and with careless disregard, put the lives of innocent, naive, trusting people (including some as young as 5 years-old) at the risk of incurring lifelong damage from taking the experimental gene therapy injections.

I desperately hope that you will address these serious concerns with Health Commissioner Johnson and demand that the MHD fulfill its MANDATORY REQUIREMENTS as a HEALTHCARE PROVIDER ADMINISTERING the Pfizer covid-19 vaccine (similar requirements hold for the Moderna and Janssen injections but they are not explicitly declared to be MANDATORY).

Sincerely,

Paul Mozina

> Begin forwarded message:
>
> **From:** Paul <paulmozina@wi.rr.com>
> **Subject:** **CCFN 200043 Communication from the Milwaukee Health Department providing an update on its efforts responding to the COVID-19 pandemic.**
> **Date:** November 10, 2021 at 5:49:11 PM CST
> **To:** "Dimitrijevic, Marina" <Marina@milwaukee.gov>, "Rainey, Khalif" <Khalif.Rainey@milwaukee.gov>, "Borkowski, Mark" <Mark.Borkowski@milwaukee.gov>, scott.spiker@milwaukee.gov
>
> Dear Chairwoman Dimitrijevic and Public Safety & Health Committee members,
>
> I hope to make a public comment in person at tomorrow's meeting.
>
> I have visited 5 different MHD covid vaccine clinics and no one at any of the clinics had any idea they were ***required*** to report to the Vaccine Adverse Event Reporting System (VAERS). Only a couple of the nurses I spoke to were even vaguely aware of VAERS and none of them take it seriously enough to warn potential vaccine recipients of the adverse events, including deaths, reported there. This is a crime. The MHD is not providing the information that people need to give informed consent.
>
> FACT SHEET FOR HEALTHCARE PROVIDERS ADMINISTERING VACCINE (VACCINATION PROVIDERS) EMERGENCY USE AUTHORIZATION (EUA) OF THE PFIZER-BIONTECH COVID-19 VACCINE TO PREVENT CORONAVIRUS DISEASE 2019 (COVID-19) FOR 5 THROUGH 11 YEARS OF AGE DILUTE BEFORE USE
>
> https://www.fda.gov/media/153714/download

Page 1
Vaccination providers enrolled in the federal COVID-19 Vaccination Program must report all vaccine administration errors, all serious adverse events, cases of Multisystem Inflammatory Syndrome (MIS) in adults and children, and cases of COVID-19 that result in hospitalization or death following administration of Pfizer-BioNTech COVID-19 Vaccine. See "MANDATORY REQUIREMENTS FOR PFIZER-BIONTECH COVID-19 VACCINE ADMINISTRATION UNDER EMERGENCY USE AUTHORIZATION" for reporting requirements.

Pages 12-13
MANDATORY REQUIREMENTS FOR PFIZER-BIONTECH COVID-19 VACCINE ADMINISTRATION UNDER EMERGENCY USE AUTHORIZATION[2]
Vaccination providers administering COMIRNATY (COVID-19 Vaccine, mRNA) must adhere to the same reporting requirements.
In order to mitigate the risks of using this unapproved product under EUA and to optimize the potential benefit of Pfizer-BioNTech COVID-19 Vaccine, the following items are required. Use of unapproved Pfizer-BioNTech COVID-19 Vaccine for active immunization to prevent COVID-19 under this EUA is limited to the following (all requirements **must** be met):
1. Pfizer-BioNTech COVID-19 Vaccine is authorized for use in individuals 5 years of age and older.

2. The vaccination provider must communicate to the individual receiving the Pfizer-BioNTech COVID-19 Vaccine or their caregiver, information consistent with the "Vaccine Information Fact Sheet for Recipients and Caregivers" prior to the individual receiving Pfizer-BioNTech COVID-19 Vaccine.

3. The vaccination provider must include vaccination information in the state/local jurisdiction's Immunization Information System (IIS) or other designated system.

4. The vaccination provider is responsible for mandatory reporting of the following to the Vaccine Adverse Event Reporting System (VAERS):
• vaccine administration errors whether or not associated with an adverse event,
• serious adverse events* (irrespective of attribution to vaccination),
• cases of Multisystem Inflammatory Syndrome (MIS) in adults and children, and
• cases of COVID-19 that result in hospitalization or death.

Complete and submit reports to VAERS online at https://vaers.hhs.gov/reportevent.html. For further assistance with reporting to VAERS call 1-800-822-7967. The reports should include the words "Pfizer-BioNTech COVID-19 Vaccine EUA" in the description section of the report.
5. The vaccination provider is responsible for responding to FDA requests for information about vaccine administration errors, adverse events, cases of MIS in adults and children, and cases of COVID-19 that result in hospitalization or death following administration of Pfizer-BioNTech COVID-19 Vaccine to recipients.

* Serious adverse events are defined as:
• Death;
• A life-threatening adverse event;
• Inpatient hospitalization or prolongation of existing hospitalization;
• A persistent or significant incapacity or substantial disruption of the ability to conduct normal life functions;
• A congenital anomaly/birth defect;
• An important medical event that based on appropriate medical judgement may jeopardize the

individual and may require medical or surgical intervention to prevent one of the outcomes listed above.

OTHER ADVERSE EVENT REPORTING TO VAERS AND PFIZER INC.

Vaccination providers may report to VAERS other adverse events that are not required to be reported using the contact information above.

To the extent feasible, report adverse events to Pfizer Inc. using the contact information below or by providing a copy of the VAERS form to Pfizer Inc. **Website**	**Fax number**	**Telephone number**
www.pfizersafetyreporting.com	1-866-635-8337	1-800-438-1985

2 Vaccination providers administering COMIRNATY (COVID-19 Vaccine, mRNA) must adhere to the same reporting requirements Revised: 29 October 2021

This is what the Milwaukee Health Department is NOT DOING and it is deeply troubling. The main objective of the MHD is to "get needles in arms", not provide informed consent.

The VAERS Report attached below is just Wisconsin and it was extracted on 11/9/21. It is well documented that there is a serious Under Reporting Factor that needs to be considered when looking at VAERS data with estimates ranging from only 1% reported to 30-40% reported. I suspect that many of these adverse events were associated with people in the City of Milwaukee.

The Vaccine Adverse Event Reporting System (VAERS) Results

Request Form | Results | Map | Chart | Report | About

Dataset Documentation | Other Data Access | Help for Results | Printing Tips | Help with Exports

Save | Export | Reset

Quick Options | More Options

Top | Notes | Citation | Query Criteria

Messages:

- VAERS data in CDC WONDER are updated every Friday. Hence, results for the same query can change from week to week.
- These results are for 10,434 total events.
- Rows with zero Events Reported are hidden. Use Quick Options above to show zero rows.

Vaccine	Event Category	Events Reported	Percent (of 10,434)
COVID19 (COVID19 (JANSSEN)) (1203)	Death	9	0.09%
	Life Threatening	16	0.15%
	Permanent Disability	14	0.13%
	Congenital Anomaly / Birth Defect *	2	0.02%
	Hospitalized	70	0.67%
	Emergency Room *	177	1.70%
	Office Visit *	209	2.00%
	None of the above	624	5.98%
	Total	**1,121**	**10.74%**
COVID19 (COVID19 (MODERNA)) (1201)	Death	92	0.88%
	Life Threatening	88	0.84%
	Permanent Disability	77	0.74%
	Congenital Anomaly / Birth Defect *	1	0.01%
	Hospitalized	462	4.43%
	Emergency Room / Office Visit **	2	0.02%
	Emergency Room *	734	7.03%
	Office Visit *	922	8.84%
	None of the above	3,261	31.25%
	Total	**5,639**	**54.04%**
COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	Death	78	0.75%
	Life Threatening	100	0.96%
	Permanent Disability	74	0.71%
	Congenital Anomaly / Birth Defect *	5	0.05%
	Hospitalized	489	4.69%
	Existing Hospitalization Prolonged	5	0.05%
	Emergency Room *	924	8.86%
	Office Visit *	1,172	11.23%
	None of the above	2,871	27.52%
	Total	**5,718**	**54.80%**
Total		**12,478**	**119.59%**

Note: Submitting a report to VAERS does not mean that healthcare personnel or the vaccine caused or contributed to the adverse event (possible side effect).
*** These values are only available from VAERS 2.0 Report Form, active 06/30/2017 to present.**

** These value are only available from VAERS-1 Report Form, active 07/01/1990 to 06/29/2017.

> Attached is the FDA document linked above from which the information provided above was excerpted.

Annotated 28Oct...1_.pdf

> Sincerely,
>
> Paul Mozina