

AEFI Reporting Form

Note: This form is for health care practitioner use only. If you are a member of the general public and need to report an adverse event following immunization, please call Health Link at 811 or contact your health provider.

1. Today's Date*

15/11/2021

2. Reporter(Your) LastName*

CHAN

3. Reporter(Your) FirstName*

GREGORY

4. Reporter (Your) SiteType:*

Physician

5. If Other SiteType, please specify:

6. Reporter(Your) Phone#*

(only numbers, no -,())

7. Immunization Facility:

Name	SiteType
Maskwacis Health Services	Public Health

If other SiteType, please specify

8. Immunization Facility Phone#

(only numbers, no -,())

9. Immunizing Facility Address

Building No/Street/PO Box	City/Town	Province	Postalcode(A1A 1A1)
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10. Select Zone: (Click [here](#) to determine zone.)

Central ▼

11. Patient LastName*

Minde

12. Patient FirstName*

Jenelle

13. If Patient is a minor, then Parent/Guardian

LastName	FirstName	RelationToPatient
-	-	-

14. Patient Sex at Birth:*

F-Female ▼

15. Patient/Guardian Phone#*

(only numbers, no -,())

780-839-838

16. PHN/ULI Info:

106220230

17. Date Of Birth(dd/mm/yyyy)*

24/07/1991

18. Patient Address:

Building No/Street/POBox	City/Town	Province	PostalCode(A1A 1A1)
Box 1228	MASKWACIS	AB	TOC 1N0

19. Date Of Immunization(dd/mm/yyyy):*

21/10/2021

20. Time of Immunization(If Known)(00:00:00)

00:00:00

21. List all the vaccines given on date of immunization.

Immunization Information:

Vaccine Code	Manufacturer	LotNo.
COVMODmF ▼	MODTH-Moc ▼	
Influenza (Aç ▼	Unknown/Otl ▼	

- Please Sel ▼
- Please Sel ▼
- Please Sel ▼
- Please Sel ▼
- Please Sel ▼

22. Adverse Event Info:

Symptoms	Started Date (mm/dd/yyyy)	Resolved?	Resolved Date (mm/dd/yyyy)
Other ▼	14/11/2021	No ▼	
None ▼		None ▼	
None ▼		None ▼	
None ▼		None ▼	

23. If other, describe including Started date & Resolved date:

Tinnitus

24. Additional Information:

new onset of RIGHT ear tinnitus this evening at ~2100h
 no recent trauma or injury or fall or loud noise/music
 normal vital signs
 RIGHT tympanic membrane is normal, LEFT TM normal, throat normal

25. Patient is aware that you are reporting the AEFI description and patient contact information to Alberta Health Services AEFI follow-up.*

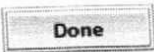
Yes ▼

26. Patient is aware Alberta Health services AEFI program may be contacting them.*

Yes ▼

Note: Hit Done button to Submit the Form.

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AEFI Reporting Form

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1. Today's Date*
18/11/2021
2. Reporter(Your) LastName*
CHAN
3. Reporter(Your) FirstName*
Gregory
4. Reporter (Your) SiteType:*
Physician
5. If Other SiteType, please specify:

6. Reporter(Your) Phone#*
(only numbers, no -,())

7. Immunization Facility:

Name	SiteType	If other SiteType, please specify
Ponoka Professional Pharma	-- Please Select --	

8. Immunization Facility Phone#
(only numbers, no -,())

9. Immunizing Facility Address

Building No/Street/PO Box	City/Town	Province	Postalcode(A1A 1A1)

5011-48 AVE

PONOKA

AB

T4J 1R5

10. Select Zone: (Click [here](#) to determine zone.)

Central

11. Patient LastName*

Dux

12. Patient FirstName*

Janelle

13. If Patient is a minor, then Parent/Guardian

LastName

FirstName

RelationToPatient

14. Patient Sex at Birth:*

F-Female

15. Patient/Guardian Phone#*

(only numbers, no -,())

403-704-3385

16. PHN/ULI Info:

515301420

17. Date Of Birth(dd/mm/yyyy)*

04/12/1984

18. Patient Address:

**Building
No/Street/POBox**

City/Town

Province

**PostalCode(A1A
1A1)**

RR1 SITE 17, B1

PONOKA

AB

T4J 1R1

19. Date Of Immunization(dd/mm/yyyy):*

30/09/2021

20. Time of Immunization(If Known)(00:00:00)

00:00:00

21. List all the vaccines given on date of immunization.

Immunization Information:

Vaccine Code

Manufacturer

LotNo.

COVMODrr

MODTH-Mc

-- Please S

-- Please S

- Please Select
- Please Select
- Please Select
- Please Select

22. Adverse Event Info:

Symptoms	Started Date (mm/dd/yyyy)	Resolved?	Resolved Date (mm/dd/yyyy)
Other <input type="checkbox"/>	10/09/2021	No <input type="checkbox"/>	
None <input type="checkbox"/>		None <input type="checkbox"/>	
None <input type="checkbox"/>		None <input type="checkbox"/>	
None <input type="checkbox"/>		None <input type="checkbox"/>	

23. If other, describe including Started date & Resolved date:

dyspnea

24. Additional Information:

possibly had COVID in September, then 1st injection of Moderna Sep 30/21
 persistent dyspnea, positive D-dimer, other tests negative
 awaiting cardiologist and lung function tests; stress testing done

25. Patient is aware that you are reporting the AEFI description and patient contact information to Alberta Health Services AEFI follow-up.*

Yes

26. Patient is aware Alberta Health services AEFI program may be contacting them.*

Yes

Note: Hit Done button to Submit the Form.

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PS

Jan 11, 2022

[redacted] from AEFI-587-779-9556

Looking to speak to CG regarding symptoms
? is chest pain related to accident or vaccination
?did the pain only appeared after vaccination
?is this a previous condition
?do you think she can have further doses

She has already had two doses so not time sensitive

She Will get notes from PGH health records regarding visit on NOV 9/2021 , saw Dr [redacted]

Would like to take a quick minute so that the file can be closed

Thurs or Fri between 8:45-3:30

Note to Dr CG to review

Panel Manager: [redacted], Jan 11, 2022, 9:50AM

Jan 19, 2022

PS

[redacted] from AEFI called

587-779-9556

wanting to speak to Dr CG about reaction after vaccine
Advised a fax with questions and concerns should be sent
she is working from home so can not do that at this time

Says bike accident in 2020 and presented to emerg on NOV 19/2021 and saw Dr [redacted]

had cardiac tests done and were normal
was diagnosed then with Pietze Syndrome

Wants to know:

- do you think her symptoms were related to the vaccine?
- do you think it is a Pre Existing condition?
- Is this muscle skeletal or cardiac?
- Had muscular Skelatal between ribs when she had the bike accident.
- Do you agree with the diagnosis of Tietze Syndrom from Dr [redacted] in emerg?
- Any plans to do more cardiac appt/testing in the near future?

Only the dr or a nurse can call her back and relay the answers to these questions.

Note given to Dr CG to review

Panel Manager: [redacted] Jan 19, 2022, 10:10AM

Feb 7, 2022

PS

[redacted] AEFI called

587-779-9556

Wanting to close the file on this pt

If a nurse can call her back with the questions as listed about from previous calls

Panel Manager: [redacted] Feb 7, 2022, 1:51PM

Feb 8, 2022

GC/PS

[redacted] AEFI

wanting to close file

would like an appt to discuss, she said has a few questions to finish up

[REDACTED] took the call
they decided to make her an appt to discuss
Mar 3/2022 for a call
Panel Manager: [REDACTED], Feb 8, 2022, 12:01PM

Feb 8, 2022

KS

[REDACTED] AEFI called regarding multiple attempts to contact Dr. [REDACTED] regarding patient history.
Asked Penny to make an appointment for Dr. [REDACTED] to call [REDACTED] the morning of March 3rd
2022.

Feb 8, 2022

GC/PS

[REDACTED] 587-779-9556
AEFI
booked an appt to speak to GC regarding patient as he did the AEFI form
same questions as she will be asking CG on her phone appt Mar 3/2022

- Wants to know:
- do you think her symptoms were related to the vaccine?
- do you think it is a Pre Existing condition?
- Is this muscle skeletal or cardiac?
- Had muscular Skelatal between ribs when she had the bike accident.
- Do you agree with the diagnosis of Tietze Syndrom from Dr [REDACTED] in emerg?
- Any plans to do more cardiac appt/testing in the near future?

Panel Manager: [REDACTED], Feb 8, 2022, 2:58PM

Feb 9, 2022

GC

Dr. Gregory Chan, February 9, 2022, 10:24AM
**** COVID-19 telephone conversation ****
discussed with AEFI team
they want to speak with Dr. Gilbert - I informed them that Dr. Gilbert does not want to speak
with them
asking if chest pain was pre-existing -> chart review, chest pain mentioned in Aug 2020 and
none since
wondering if it is a rib out
asking whether I recommend another dose - I cannot make this recommendation
advised that echo and MRI are not helpful now
Dr. Gregory Chan, February 9, 2022, 10:34AM

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1. Today's Date*
18/11/2021

2. Reporter(Your) LastName*
CHAN

3. Reporter(Your) FirstName*
GREGORY

4. Reporter (Your) SiteType:*
Physician

5. If Other SiteType, please specify:

6. Reporter(Your) Phone#*
(only numbers, no -,())
[REDACTED]

7. Immunization Facility

Name	SiteType	If other SiteType, please specify
Shopper's Drug Mart Ponoka	Pharmacy	<input checked="" type="checkbox"/>

8. Immunization Facility Phone#
(only numbers, no -,())
[REDACTED]

9. Immunizing Facility Address

Building No/Street/PO Box	City/Town	Province	Postalcode(A1A 1A1)
[REDACTED]	[REDACTED]	[REDACTED]	[REDACTED]

10. Select Zone: (Click [here](#) to determine zone.)

Central

11. Patient LastName*

Cunningham

12. Patient FirstName*

Kendall

13. If Patient is a minor, then Parent/Guardian

LastName	FirstName	RelationToPatient
-	-	-

14. Patient Sex at Birth:*

F-Female

15. Patient/Guardian Phone#*

(only numbers, no -,())

50-363-184

16. PHN/ULI Info:

[Redacted]

17. Date Of Birth(dd/mm/yyyy)*

[Redacted]

18. Patient Address:

Building No/Street/POBox	City/Town	Province	PostalCode(A1A 1A1)
5015 50th Ave	Ponoka	AB	T4J 0C1

19. Date Of Immunization(dd/mm/yyyy):*

21/10/2021

20. Time of Immunization(If Known)(00:00:00)

00:00:00

21. List all the vaccines given on date of immunization.

Immunization Information:

Vaccine Code	Manufacturer	LotNo.
COVPBmR <input type="checkbox"/>	BPF-BioNTi <input type="checkbox"/>	
-- Please Si <input type="checkbox"/>	-- Please Si <input type="checkbox"/>	

- Please Select
- Please Select
- Please Select
- Please Select
- Please Select

22. Adverse Event Info:

Symptoms	Started Date (mm/dd/yyyy)	Resolved?	Resolved Date (mm/dd/yyyy)
Pain and/or Sw	10/27/2021	No	
None		None	
None		None	
None		None	

23. If other, describe including Started date & Resolved date:

chest pain

24. Additional Information:

chest pain since SECOND dose of PFIZER Oct 21/21
 HR increased from baseline
 intermittent chest pain, paroxysmal
 slight increase in WBC

25. Patient is aware that you are reporting the AEFI description and patient contact information to Alberta Health Services AEFI follow-up.*

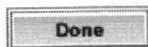
Yes

26. Patient is aware Alberta Health services AEFI program may be contacting them.*

Yes

Note: Hit Done button to Submit the Form.

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Jan 7, 2022

[REDACTED] from AHS -Adverse reaction to vaccine
587-943-1987
would like to speak to Dr Chan regarding the plan for her and also another symptom that was
not written on form that she mentioned when AHS called her
note to Dr Chan to call her
Panel Manager: [REDACTED], Jan 7, 2022, 12:54PM

discussed with [REDACTED], agreed that itching is an adverse reaction, will be reported to MOH
Dr. Gregory Chan, January 7, 2022, 1:59PM

PS

Jan 10, 2022

[REDACTED] from AHS
discussed reaction and recommendations with Dr Horne
Pt is ok to have additional covid vaccine, recommended 30 min required after shot for viewing
note to Dr Chan to advise
Panel Manager: [REDACTED], Jan 10, 2022, 1:08PM

GC/BR

Jan 28, 2022

Jan 28, 2022, 9:46AM

Action - patient brought to room by BR

S: left ear infection

O: HEENT TMs left TM swollen, right normal throat normal

A: OM

P: inflammation of TM

AEFI panel has stated that benefits of booster outweigh risk of COVID

Dr. Gregory Chan, January 28, 2022, 10:11AM



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1. Today's Date*

2. Reporter(Your) LastName*

3. Reporter(Your) FirstName*

4. Reporter (Your) SiteType:*

5. If Other SiteType, please specify:

6. Reporter(Your) Phone#*

(only numbers, no -,())

7. Immunization Facility:

Name	SiteType	If other Site type, please specify
<input type="text" value="Rexall Ponoka"/>	<input type="text" value="Pharmacy"/>	<input type="text"/>

8. Immunization facility phone#

(only numbers, no -,())

9. Immunizing Facility Address

Building No/Street/PO Box	City/Town	Province	Postalcode(A1A 1A1)
BAY 3, 4502-50 ST	PONOKA	AB	T4J 1J5

10. Select Zone: (Click [here](#) to determine zone.)

Central

11. Patient LastName*

[Redacted]

12. Patient FirstName*

[Redacted]

13. If Patient is a minor, then Parent/Guardian

LastName	FirstName	RelationToPatient
-	-	-

14. Patient Sex at Birth:*

F-Female

15. Patient/Guardian Phone#*

(only numbers, no -,())

[Redacted]

16. PHN/ULI Info:

[Redacted]

17. Date Of Birth(dd/mm/yyyy)*

[Redacted]

18. Patient Address:

Building No/Street/POBox	City/Town	Province	PostalCode(A1A 1A1)
[Redacted]	[Redacted]	[Redacted]	[Redacted]

19. Date Of Immunization(dd/mm/yyyy):*

28/05/2021

20. Time of Immunization(If Known)(00:00:00)

00:00:00

21. List all the vaccines given on date of immunization.

Immunization Information:

Vaccine Code	Manufacturer	LotNo.
COVPBmRNA	BPF-BioNTec	
-- Please Sel	-- Please Sel	
-- Please Sel	-- Please Sel	
-- Please Sel	-- Please Sel	
-- Please Sel	-- Please Sel	

22. Adverse Event Info:

Symptoms	Started Date (mm/dd/yyyy)	Resolved?	Resolved Date (mm/dd/yyyy)
Other	05/28/2021	No	
None		None	
None		None	
None		None	

23. If other, describe including Started date & Resolved date:

Pruritis

24. Additional Information:

1st injection May 28/21 - had itching immediately afterward, rapid HR, severe, called 811 and then advised to call 911 treated on site by EMS
 told to take Benadryl before the 2nd dose (as per paramedics) - physician in Red Deer giving cortisone injections also suggested this
 2nd injection July 7/21 - had itching worsened and flushing immediately afterwards, was given an injection of Benadryl at the pharmacy
 itching persists today, worse in warm environments

25. Patient is aware that you are reporting the AEFI description and patient contact information to Alberta Health Services AEFI follow-up.*

Yes

26. Patient is aware Alberta Health services AEFI program may be contacting them.*

Yes

Note: Hit Done button to Submit the Form.

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Done

[REDACTED]
Jan 21, 2022

[REDACTED] from AHS-780-306-2178

AEFI

wanting an appt or call to discuss

?how do you want to proceed

? doesn't know if he qualifies for an exemption

?what do you want from them

?referred to a cardiologist

Advised to fax

she said they have done that and all info has been sent

wondering about holter monitor, not on netcare

wanted to speak to him directly

advised an appt would be best

she said that was fine , advised a week + away

she said no to far

note to Dr Chan to advise

Panel Manager: [REDACTED] Jan 21, 2022, 2:43PM

Jan 28, 2022

discussed with [REDACTED] regarding event

states that mom thought he was still limited between COVID and vaccine

I recall that he was able to participate in sports, but then after the vaccine he was severely limited

recently able to continue with physical activity

Dr. Gregory Chan, January 28, 2022, 5:13PM

Jan 31, 2022

Jan 31, 2022, 9:52AM

Action - patient brought to room by AE

lost appetite and some fatigue from COVID infection

lost about 20lb

had some issues with stamina

was able to return to practice and workouts and dry run training

was coming back to usual self

collateral history - mom states that

then had injection then had the incident and then amnesia of the event

fatigued, pale and took weeks to recover

GC

GC

AEFI Reporting Form

Note: This form is for health care practitioner use only. If you are a member of the general public and need to report an adverse event following immunization, please call Health Link at 811 or contact your health provider.

1. Today's Date*
25/11/2021

2. Reporter(Your) LastName*
CHAN

3. Reporter(Your) FirstName*
GREGORY

4. Reporter (Your) SiteType: *
Physician

5. If Other SiteType, please specify:

6. Reporter(Your) Phone#*
(only numbers, no -,())

7. Immunization Facility:

Name	SiteType	If other SiteType, please specify
Save-On-Foods Trinity Hills	Pharmacy	

8. Immunization Facility Phone#
(only numbers, no -,())

9. Immunizing Facility Address

Building No/Street/PO Box	City/Town	Province	Postalcode(A1A 1A1)
420 NA'A PLAZA	CALGARY	AB	T3H 6A4

10. Select Zone: (Click [here](#) to determine zone.)

Calgary

11. Patient LastName*

Bishop

12. Patient FirstName*

Liam

13. If Patient is a minor, then Parent/Guardian

LastName

FirstName

RelationToPatient

Bishop

Jennifer

Mother

14. Patient Sex at Birth:*

M-Male

15. Patient/Guardian Phone#*

(only numbers, no -,())

403-963-3330

16. PHN/ULI Info:

61076321

17. Date Of Birth(dd/mm/yyyy)*

01/03/2009

18. Patient Address:

**Building
No/Street/POBox**

City/Town

Province

**PostalCode(A1A
1A1)**

16 GREENBRI

CALGAR

AB

T2B 6J2

19. Date Of Immunization(dd/mm/yyyy):*

29/10/2021

20. Time of Immunization(If Known)(00:00:00)

14:00:00

21. List all the vaccines given on date of immunization.

Immunization Information:

Vaccine Code	Manufacturer	LotNo.
COVPBmR <input type="checkbox"/>	BPF-BioNTi <input type="checkbox"/>	
-- Please S <input type="checkbox"/>	-- Please S <input type="checkbox"/>	
-- Please S <input type="checkbox"/>	-- Please S <input type="checkbox"/>	
-- Please S <input type="checkbox"/>	-- Please S <input type="checkbox"/>	

22. Adverse Event Info:

Symptoms		Started Date (mm/dd/yyyy)	Resolved?	Resolved Date (mm/dd/yyyy)
Other	v	10/30/2021	Yes	10/30/2021
Other	v	10/30/2021	No	
None	v		None	
None	v		None	

23. If other, describe including Started date & Resolved date:

Syncope (resolved), dyspnea/palpitations (unresolv

24. Additional Information:

had COVID Sept 15/21, then recovered - NO SYMPTOMS after recovery
 had some dizziness then lost weight as well, bad cough as well
 after 1 month - was encouraged to have vaccine as per government and to play hockey
 had 1st dose of PFIZER Oct 29/21 1400 - Save on Foods Trinity Hills
 Oct 30/21 early morning found on the ground (0600h)
 mom could not find a pulse, unresponsive, 911 EMS arrived
 brought to hospital to be checked at ACH
 felt light in the chest
 no headache
 very fatigued
 investigations performed, then released
 told by ER physician that this was an adverse reaction to the vaccine
 no further syncope

Nov 2/21 "Dryland episode" - doing more physical activity then appeared pale, was feeling cold
 headache, then felt like he was going to pass out
 heart not feeling right
 could not participate in sports due to recurrent symptoms

has tried practices since, no weight training since (due to cold sensation, pale appearance,
 malaise when working out)

occasional headache and dizziness, consistently appears pale after skating
 some palpitations with minimal activity

25. Patient is aware that you are reporting the AEFI description and patient contact information to Alberta Health Services AEFI follow-up.*

Yes v

26. Patient is aware Alberta Health services AEFI program may be contacting them.*

Yes v

Note: Hit Done button to Submit the Form.

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conducting research, providing for health services, provider education, internal management purposes, planning and resource allocation, health system management, public health surveillance and health policy development.

Approved Date
(mm/dd/yyyy)

Reviewed

Done

Comments

10/05/17

✓

✓

10/05/17

✓

✓

10/05/17

✓

✓

✓

✓

✓

✓

✓

✓

✓

✓

✓

✓

22. I am not in the system and I have not been notified.

System involved in the notification process.

23. Additional information:

and their data is not in the system. We are currently in the process of updating the system and will be able to provide you with the data you need. We are currently in the process of updating the system and will be able to provide you with the data you need. We are currently in the process of updating the system and will be able to provide you with the data you need.

Program is intended to be used by all staff.

Left list in the list.

as a result.

very helpful.

Thank you for your assistance. We will be able to provide you with the data you need.

only if it appears that this was an error on the part of the system.

on the part of the system.

the list is not in the system. We are currently in the process of updating the system and will be able to provide you with the data you need.

please let us know if you have any questions.

Thank you for your assistance.

we will be able to provide you with the data you need.

Thank you for your assistance.

we will be able to provide you with the data you need.

Thank you for your assistance.

24. I am not in the system and I have not been notified.

System involved in the notification process.

Yes

✓

Yes

✓

25. I am not in the system and I have not been notified.

Additional information: We are currently in the process of updating the system and will be able to provide you with the data you need. We are currently in the process of updating the system and will be able to provide you with the data you need. We are currently in the process of updating the system and will be able to provide you with the data you need.

AEFI Reporting Form

Note: This form is for health care practitioner use only. If you are a member of the general public and need to report an adverse event following immunization, please call Health Link at 811 or contact your health provider.

1. Today's Date*
26/11/2021

2. Reporter(Your) LastName*
CHAN

3. Reporter(Your) FirstName*
GREGORY

4. Reporter (Your) SiteType:*
Physician

5. If Other SiteType, please specify:

6. Reporter(Your) Phone#*
(only numbers, no -,())

7. Immunization Facility:

Name	SiteType	If other SiteType, please specify
Rexall Ponoka	Pharmacy	

8. Immunization Facility Phone#
(only numbers, no -,())

9. Immunizing Facility Address

Building No/Street/PO Box	City/Town	Province	Postalcode(A1A 1A1)

10. Select Zone: (Click [here](#) to determine zone.)

Central

11. Patient LastName*

Denis

12. Patient FirstName*

Jayden

13. If Patient is a minor, then Parent/Guardian

LastName	FirstName	RelationToPatient
-	-	-

14. Patient Sex at Birth:*

F-Female

15. Patient/Guardian Phone#*

(only numbers, no -,())

403-783-1960

16. PHN/ULI Info:

873549880

17. Date Of Birth(dd/mm/yyyy)*

14/11/1996

18. Patient Address:

Building No/Street/POBox	City/Town	Province	PostalCode(A1A 1A1)
5001-42A STRE	PONOKA	AB	T4J 1M3

19. Date Of Immunization(dd/mm/yyyy):*

15/10/2021

20. Time of Immunization(If Known)(00:00:00)

00:00:00

21. List all the vaccines given on date of immunization.

Immunization Information:

Vaccine Code	Manufacturer	LotNo.
COVPBmR <input type="checkbox"/>	BPF-BioNT <input type="checkbox"/>	
-- Please S <input type="checkbox"/>	-- Please S <input type="checkbox"/>	

- Please Select
- Please Select
- Please Select
- Please Select

22. Adverse Event Info:

Symptoms	Started Date (mm/dd/yyyy)	Resolved?	Resolved Date (mm/dd/yyyy)
Severe Diarrhea <input type="checkbox"/>	11/04/2021	No <input type="checkbox"/>	
None <input type="checkbox"/>		None <input type="checkbox"/>	
None <input type="checkbox"/>		None <input type="checkbox"/>	
None <input type="checkbox"/>		None <input type="checkbox"/>	

23. If other, describe including Started date & Resolved date:
diarrhea

24. Additional Information:

severe diarrhea and abdominal cramping 1st week of November (2 weeks after the 2nd injection)
no blood or mucous, but watery stools
cramping better with imodium and pro-biotics
still having 4-5 bowel movements per day

25. Patient is aware that you are reporting the AEFI description and patient contact information to Alberta Health Services AEFI follow-up.*

Yes

26. Patient is aware Alberta Health services AEFI program may be contacting them.*

Yes

Note: Hit Done button to Submit the Form.

Alberta Health Services (AHS) respects your confidentiality and privacy. Your information is collected, used, disclosed and protected according to the provisions of provincial and federal legislation. Your health information is collected by AHS in accordance with section 20 of the Health Information Act (HIA). The purpose of this collection is primarily for: providing health services, determining eligibility for health services, processing payments for health services, conducting research, providing for health services, provider education, internal management purposes, planning and resource allocation, health system management, public health surveillance and health policy development.

Done

GC/PS

Jan 12, 2022

request for medical records from AEFI
verbal consent has been given from pt to go ahead and send
Sent by fax to 780-342-0248
ok'd by Dr Chan
Panel Manager: [REDACTED], Jan 12, 2022, 10:17AM

GC/PS

Jan 12, 2022

call from [REDACTED] with AHS-587-943-1665
wanting information regarding vaccine after effects -AEFI
call with Dr Chan wanted
?requires further testing
?recommendations for further vaccines
spoke to Dr Chan
he said to tell them that he writes all the information as discussed with pt about symptoms and
follows all guidelines as per website on AEFI form before submitting
if more information is required they can send a fax
Advised to speak to the patient also
Panel Manager: [REDACTED], Jan 12, 2022, 10:50AM

AEFI Reporting Form

Note: This form is for health care practitioner use only. If you are a member of the general public and need to report an adverse event following immunization, please call Health Link at 811 or contact your health provider.

1. Today's Date*
26/11/2021

2. Reporter(Your) LastName*
CHAN

3. Reporter(Your) FirstName*
GREGORY

4. Reporter (Your) SiteType:*
Physician

5. If Other SiteType, please specify:

6. Reporter(Your) Phone#*
(only numbers, no -,())

7. Immunization Facility:

Name	SiteType	If other SiteType, please specify
Rexall Ponoka	Pharmacy <input checked="" type="checkbox"/>	

8. Immunization Facility Phone#
(only numbers, no -,())

9. Immunizing Facility Address

Building No/Street/PO Box	City/Town	Province	Postalcode(A1A 1A1)

10. Select Zone: (Click [here](#) to determine zone.)

Central

11. Patient LastName*

ANUSZEWSKI

12. Patient FirstName*

Leanne

13. If Patient is a minor, then Parent/Guardian

LastName	FirstName	RelationToPatient
-	-	-

14. Patient Sex at Birth:*

F-Female

15. Patient/Guardian Phone#*

(only numbers, no -,())

403-358-1785

16. PHN/ULI Info:

667236900

17. Date Of Birth(dd/mm/yyyy)*

25/09/1973

18. Patient Address:

Building No/Street/POBox	City/Town	Province	PostalCode(A1A 1A1)
1 STIG-48 AV	PONOKA	AB	T4J 1J5

19. Date Of Immunization(dd/mm/yyyy):*

15/10/2021

20. Time of Immunization(If Known)(00:00:00)

00:00:00

21. List all the vaccines given on date of immunization.

Immunization Information:

Vaccine Code	Manufacturer	LotNo.
COVPBmR <input checked="" type="checkbox"/>	BPF-BioNTi <input checked="" type="checkbox"/>	
-- Please Si <input checked="" type="checkbox"/>	-- Please Si <input checked="" type="checkbox"/>	

- Please Select
- Please Select
- Please Select
- Please Select
- Please Select

22. Adverse Event Info:

Symptoms	Started Date (mm/dd/yyyy)	Resolved?	Resolved Date (mm/dd/yyyy)
Other	11/23/2021	No	
None		None	
None		None	
None		None	

23. If other, describe including Started date & Resolved date:

Tinnitus

24. Additional Information:

new onset of tinnitus this week, recent immunization is the only new medication/procedure in the last 4 weeks. NO PREVIOUS TINNITUS.
 no symptoms when waking up, but low tone develops and increases/decreases through the day
 physical exam is normal aside from mild irritation in the nasal passage

25. Patient is aware that you are reporting the AEFI description and patient contact information to Alberta Health Services AEFI follow-up.*

Yes

26. Patient is aware Alberta Health services AEFI program may be contacting them.*

Yes

Note: Hit Done button to Submit the Form.

Alberta Health Services (AHS) respects your confidentiality and privacy. Your information is collected, used, disclosed and protected according to the provisions of provincial and federal legislation. Your health information is collected by AHS in accordance with section 20 of the Health Information Act (HIA). The purpose of this collection is primarily for: providing health services, determining eligibility for health services, processing payments for health services, conducting research, providing for health services, provider education, internal management purposes, planning and resource allocation, health system management, public health surveillance and health policy development.



AEFI Reporting Form

Note: This form is for health care practitioner use only. If you are a member of the general public and need to report an adverse event following immunization, please call Health Link at 811 or contact your health provider.

1. Today's Date*
29/11/2021

2. Reporter(Your) LastName*
CHAN

3. Reporter(Your) FirstName*
GREGORY

4. Reporter (Your) SiteType:*
Physician

5. If Other SiteType, please specify:
[Empty field]

6. Reporter(Your) Phone#*
(only numbers, no -,())
[Redacted]

7. Immunization Facility:

	Name	SiteType	If other SiteType, please specify
	Maskwacis Health Center	Public Health	

8. Immunization Facility Phone#
(only numbers, no -,())
[Redacted]

9. Immunizing Facility Address

Building No/Street/PO Box	City/Town	Province	Postalcode(A1A 1A1)
--	------------------	-----------------	--------------------------------

10. Select Zone: (Click [here](#) to determine zone.)

Central ▼

11. Patient LastName*

Currie

12. Patient FirstName*

Warner Jarvis

13. If Patient is a minor, then Parent/Guardian

LastName	FirstName	RelationToPatient
-	-	-

14. Patient Sex at Birth:*

M-Male ▼

15. Patient/Guardian Phone#*

(only numbers, no -,())

80-585-2280

16. PHN/ULI Info:

614772610

17. Date Of Birth(dd/mm/yyyy)*

18/04/1963

18. Patient Address:

Building No/Street/POBox	City/Town	Province	PostalCode(A1A 1A1)
BOX 2626	MASKWACIS	AB	T0C 1N0

19. Date Of Immunization(dd/mm/yyyy):*

18/11/2021

20. Time of Immunization(If Known)(00:00:00)

00:00:00

21. List all the vaccines given on date of immunization.

Immunization Information:

Vaccine Code	Manufacturer	LotNo.
COVMODmF ▼	MODTH-Moc ▼	
-- Please Sel ▼	-- Please Sel ▼	

-- Please Sel ▼ -- Please Sel ▼
 -- Please Sel ▼ -- Please Sel ▼
 -- Please Sel ▼ -- Please Sel ▼

22. Adverse Event Info:

Symptoms	Started Date (mm/dd/yyyy)	Resolved?	Resolved Date (mm/dd/yyyy)
Other ▼	11/28/2021	No ▼	
None ▼		None ▼	
None ▼		None ▼	
None ▼		None ▼	

23. If other, describe including Started date & Resolved date:

peripheral neuropathy

24. Additional Information:

3rd dose of Moderna Nov 18/21. 1st dose Jan 22/21, 2nd dose Feb 26/21.
 2nd visit to ER with migrating/progressive neurological symptoms starting in the last 12 hours. This is 10 days post 3rd dose. He noticed some weakness to hands last night then came to the ER. Sent home. The lower extremities seem to be involved today (not yesterday) BP is slightly low 98/72, HR normal.
 MED: Epilepsy; HTN; dyslipidemia; Brain injury remote Hx; Atrial fibrillation

25. Patient is aware that you are reporting the AEFI description and patient contact information to Alberta Health Services AEFI follow-up.*

Yes ▼

26. Patient is aware Alberta Health services AEFI program may be contacting them.*

Yes ▼

Note: Hit Done button to Submit the Form.

Alberta Health Services (AHS) respects your confidentiality and privacy. Your information is collected, used, disclosed and protected according to the provisions of provincial and federal legislation. Your health information is collected by AHS in accordance with section 20 of the Health Information Act (HIA). The purpose of this collection is primarily for: providing health services, determining eligibility for health services, processing payments for health services, conducting research, providing for health services, provider education, internal management purposes, planning and resource allocation, health system management, public health surveillance and health policy development.

Done

AEFI Reporting Form

Note: This form is for health care practitioner use only. If you are a member of the general public and need to report an adverse event following immunization, please call Health Link at 811 or contact your health provider.

1. Today's Date*

21/12/2021

2. Reporter(Your) LastName*

Chan

3. Reporter(Your) FirstName*

Gregory

4. Reporter (Your) SiteType:*

Physician

5. If Other SiteType, please specify:

6. Reporter(Your) Phone#*

(only numbers, no -,())

7. Immunization Facility

Name	SiteType	If other SiteType, please specify
Parson's Pharmacy	Pharmacy	

8. Immunization Facility Phone#

(only numbers, no -,())

9. Immunizing Facility Address

Building No/Street/PO Box	City/Town	Province	Postalcode(A1A 1A1)

1C - 4755 49TH

RED DEER

AB

T4N 1T6

10. Select Zone: (Click [here](#) to determine zone.)

Central ▼

11. Patient LastName*

Fleming

12. Patient FirstName*

Margo

13. If Patient is a minor, then Parent/Guardian

LastName

FirstName

RelationToPatient

-

-

-

14. Patient Sex at Birth:*

F-Female ▼

15. Patient/Guardian Phone#*

(only numbers, no -,())

403-505-6703

16. PHN/ULI Info:

533766410

17. Date Of Birth(dd/mm/yyyy)*

28/02/1981

18. Patient Address:

Building
No/Street/POBox

City/Town

Province

PostalCode(A1A
1A1)

103

RIMBEY

AB

T0C 2J0

19. Date Of Immunization(dd/mm/yyyy):*

21/06/2021

20. Time of Immunization(If Known)(00:00:00)

00:00:00

21. List all the vaccines given on date of immunization.
Immunization Information:

Vaccine Code

Manufacturer

LotNo.

COVPBmR ▼

BPF-BioNTx ▼

-- Please Sx ▼

-- Please Sx ▼

-- Please Sr ▼	-- Please Sr ▼	
-- Please Sr ▼	-- Please Sr ▼	
-- Please Sr ▼	-- Please Sr ▼	

22. Adverse Event Info:

Symptoms	Started Date (mm/dd/yyyy)	Resolved?	Resolved Date (mm/dd/yyyy)
Rash ▼	07/19/2021	No ▼	
None ▼		None ▼	
None ▼		None ▼	
None ▼		None ▼	

23. If other, describe including Started date & Resolved date:

rash

24. Additional Information:

initial rash on right arm, clustered raised erythematous lesions
 then lesions on plantar surface 3rd digit right foot - blister with erythematous background,
 additional lesions on 1st digit with bruising
 now lesions on left arm raised with hypopigmentation at the periphery, raised, pruritic -
 targetoid lesions clustered circular lesions 2-3 grouped

25. Patient is aware that you are reporting the AEFI description and patient contact information to Alberta Health Services AEFI follow-up.*

Yes ▼

26. Patient is aware Alberta Health services AEFI program may be contacting them.*

Yes ▼

Note: Hit Done button to Submit the Form.

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Done

Jan 31, 2022

SG

note [redacted] AHS AEFI said no contraindications for next vaccine

AEFI Reporting Form

Note: This form is for health care practitioner use only. If you are a member of the general public and need to report an adverse event following immunization, please call Health Link at 811 or contact your health provider.

1. Today's Date*

6/1/2022

2. Reporter(Your) LastName*

CHAN

3. Reporter(Your) FirstName*

GREGORY

4. Reporter (Your) SiteType:*

Physician

5. If Other SiteType, please specify:

6. Reporter(Your) Phone#*

(only numbers, no -,())

7. Immunization Facility:

Name

Ponoka Professional Pharma

SiteType

Pharmacy

If other SiteType, please specify

8. Immunization Facility Phone#

(only numbers, no -,())

9. Immunizing Facility Address

Building No/Street/PO Box

City/Town

Province

Postalcode(A1A 1A1)

5011-48 AVE

PONOKA

AB

T4J 1J3

10. Select Zone: (Click [here](#) to determine zone.)

Central ▼

11. Patient LastName*

Surbey

12. Patient FirstName*

Jack

13. If Patient is a minor, then Parent/Guardian

LastName

FirstName

RelationToPatient

-

-

-

14. Patient Sex at Birth:*

M-Male ▼

15. Patient/Guardian Phone#*

(only numbers, no -,())

403-588-0597

16. PHN/ULI Info:

684162010

17. Date Of Birth(dd/mm/yyyy)*

21-10-1951

18. Patient Address:

Building
No/Street/POBox

City/Town

Province

PostalCode(A1A
1A1)

5822-61 STREET

PONOKA

AB

T4J 1L3

19. Date Of Immunization(dd/mm/yyyy):*

23/12/2021

20. Time of Immunization(If Known)(00:00:00)

00:00:00

21. List all the vaccines given on date of immunization.

Immunization Information:

Vaccine Code

Manufacturer

LotNo.

COVMODmF ▼

MODTH-Moc ▼

-- Please Sel ▼

-- Please Sel ▼

-- Please Sel ▼ -- Please Sel ▼
 -- Please Sel ▼ -- Please Sel ▼
 -- Please Sel ▼ -- Please Sel ▼

22. Adverse Event Info:

Symptoms	Started Date (mm/dd/yyyy)	Resolved?	Resolved Date (mm/dd/yyyy)
Pain and/or Swe ▼	01/05/2022	No ▼	
Other ▼	12/23/2021	Yes ▼	12/25/2021
None ▼		None ▼	
None ▼		None ▼	

23. If other, describe including Started date & Resolved date:

fatigue, malaise, arm pain

24. Additional Information:

OTHER - fatigue malaise arm pain on same day as the injection (3rd dose)
 doses 1 and 2 had no arm pain or fatigue or malaise
 PAIN - chest pain onset 2230 Jan 5/22, left sided. Pleuritic. Some dyspnea occurs when deep breathing. Chest pain continued today. Had episode Dec 21/21 but it resolved.
 ECG normal, CK troponin normal. D-dimer 0.13. Lipase >800.
 CT chest/abdo/pelvis pending
 exercise stress test pending
 advised to take aspirin daily until stress test result available

25. Patient is aware that you are reporting the AEFI description and patient contact information to Alberta Health Services AEFI follow-up.*

Yes ▼

26. Patient is aware Alberta Health services AEFI program may be contacting them.*

Yes ▼

Note:Hit Done button to Submit the Form.

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Done

Jan 31, 2022

PS

with AHS-587-782-4345

recommendations from Medical officer of Health

Recommends further dose of Covid 19 vaccines can not be made at this time until further research is available. AHS has reported suspected myocarditis to Alberta Health

Note to Dr Chan to review

Panel Manager: Jan 31, 2022, 3:44PM

Feb 8, 2022

PS

[REDACTED] AEFI 587-786-6448

wondering if shortness of breath is related to the infection of Covid or from the vaccine
any upcoming testing

booked in for a call as per DR CHan to discuss

Panel Manager: [REDACTED] Feb 8, 2022, 11:12AM

Feb 10, 2022

GC

Dr. Gregory Chan, February 10, 2022, 10:22AM

**** COVID-19 telephone conversation ****

discussion with AEFI

chronology fits with vaccine

advised

Dr. Gregory Chan, February 10, 2022, 10:23AM

AEFI Reporting Form

Note: This form is for health care practitioner use only. If you are a member of the general public and need to report an adverse event following immunization, please call Health Link at 811 or contact your health provider.

1. Today's Date*
17/1/2022

2. Reporter(Your) LastName*
Chan

3. Reporter(Your) FirstName*
Gregory

4. Reporter (Your) SiteType:*
Physician

5. If Other SiteType, please specify:

6. Reporter(Your) Phone#*
(only numbers, no -,())

7. Immunization Facility:

Name	SiteType	If other SiteType, please specify
Ponoka Shoppers Drug Mart	Pharmacy	▼

8. Immunization Facility Phone#
(only numbers, no -,())

9. Immunizing Facility Address

Building No/Street/PO Box	City/Town	Province	Postalcode(A1A 1A1)

10. Select Zone: (Click [here](#) to determine zone.)

Central

11. Patient LastName*

[Redacted]

12. Patient FirstName*

Darien

13. If Patient is a minor, then Parent/Guardian

LastName	FirstName	RelationToPatient
-	-	-

14. Patient Sex at Birth:*

M-Male

15. Patient/Guardian Phone#*

(only numbers, no -,())

[Redacted]

16. PHN/ULI Info:

[Redacted]

17. Date Of Birth(dd/mm/yyyy)*

[Redacted]

18. Patient Address:

Building No/Street/POBox	City/Town	Province	PostalCode(A1A 1A1)
[Redacted]	PONOKA	AB	T4J 1R3

19. Date Of Immunization(dd/mm/yyyy):*

17/11/2021

20. Time of Immunization(If Known)(00:00:00)

00:00:00

21. List all the vaccines given on date of immunization.

Immunization Information:

Vaccine Code	Manufacturer	LotNo.
COVPBmR <input type="checkbox"/>	BPF-BioNTi <input type="checkbox"/>	
-- Please Si <input type="checkbox"/>	-- Please Si <input type="checkbox"/>	

- Please Select
- Please Select
- Please Select
- Please Select

22. Adverse Event Info:

Symptoms	Started Date (mm/dd/yyyy)	Resolved?	Resolved Date (mm/dd/yyyy)
Other <input type="checkbox"/>	20/12/2021	No <input type="checkbox"/>	
None <input type="checkbox"/>		None <input type="checkbox"/>	
None <input type="checkbox"/>		None <input type="checkbox"/>	
None <input type="checkbox"/>		None <input type="checkbox"/>	

23. If other, describe including Started date & Resolved date:

Dyspnea with minimal exertion

24. Additional Information:

2nd COVID injection Nov 17/21
 mid December has noticed an increase in dyspnea with MINIMAL exertion progressively worse
 had COVID Dec 31/21
 dyspnea is persistent, can get dyspnea with 2 steps or talking
 previously active, weight lifting, sports (competitive)
 arranging ECHO and PFT and labwork

25. Patient is aware that you are reporting the AEFI description and patient contact information to Alberta Health Services AEFI follow-up.*

Yes

26. Patient is aware Alberta Health services AEFI program may be contacting them.*

Yes

Note: Hit Done button to Submit the Form.

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AEFI Reporting Form

Note: This form is for health care practitioner use only. If you are a member of the general public and need to report an adverse event following immunization, please call Health Link at 811 or contact your health provider.

1. Today's Date*

17/1/2022

2. Reporter(Your) LastName*

CHAN

3. Reporter(Your) FirstName*

GREGORY

4. Reporter (Your) SiteType:*

Physician

5. If Other SiteType, please specify:

6. Reporter(Your) Phone#*

(only numbers, no -,())

[Redacted]

7. Immunization Facility:

Name
IDA Ponoka

SiteType
Pharmacy

If other SiteType, please specify

8. Immunization Facility Phone#

(only numbers, no -,())

[Redacted]

9. Immunizing Facility Address

Building
No/Street/PO
Box

City/Town

Province

Postalcode(A1A
1A1)

5020 50 ST

PONOKA

AB

T4J 1S3

10. Select Zone: (Click [here](#) to determine zone.)

Central ▼

11. Patient LastName*

Jensen

12. Patient FirstName*

Peter

13. If Patient is a minor, then Parent/Guardian

LastName

FirstName

RelationToPatient

-

-

-

14. Patient Sex at Birth:*

M-Male ▼

15. Patient/Guardian Phone#*

(only numbers, no -,())

403-704-5351

16. PHN/ULI Info:

172254900

17. Date Of Birth(dd/mm/yyyy)*

01/08/1946

18. Patient Address:

Building
No/Street/POBox

City/Town

Province

PostalCode(A1A
1A1)

5512-57 AVE UN

PONOKA

AB

T4J 1V7

19. Date Of Immunization(dd/mm/yyyy):*

28/09/2021

20. Time of Immunization(If Known)(00:00:00)

00:00:00

21. List all the vaccines given on date of immunization.

Immunization Information:

Vaccine Code

Manufacturer

LotNo.

COVPBmRN ▼

BPF-BioNTec ▼

-- Please Sel ▼

-- Please Sel ▼

-- Please Sel ▼	-- Please Sel ▼	<input type="text"/>
-- Please Sel ▼	-- Please Sel ▼	<input type="text"/>
-- Please Sel ▼	-- Please Sel ▼	<input type="text"/>

22. Adverse Event Info:

Symptoms	Started Date (mm/dd/yyyy)	Resolved?	Resolved Date (mm/dd/yyyy)
Other ▼	10/14/2021	No ▼	
None ▼	<input type="text"/>	None ▼	<input type="text"/>
None ▼	<input type="text"/>	None ▼	<input type="text"/>
None ▼	<input type="text"/>	None ▼	<input type="text"/>

23. If other, describe including Started date & Resolved date:

urinary retention

24. Additional Information:

patient started to have urinary retention about 2 weeks after the 2nd COVID vaccine progressively worse each week November, AFTER THE VACCINE and AFTER THE ONSET OF SYMPTOMS, had treatment for prostate cancer (radiation). Operative report from December cystoscopy - "The bladder was examined, and there were no gross abnormalities seen. No tumor, masses, or any stones."

Prior to 2nd dose (September 2021) - no urinary retention symptoms

25. Patient is aware that you are reporting the AEFI description and patient contact information to Alberta Health Services AEFI follow-up.*

Yes ▼

26. Patient is aware Alberta Health services AEFI program may be contacting them.*

Yes ▼

Note:Hit Done button to Submit the Form.

Alberta Health Services (AHS) respects your confidentiality and privacy. Your information is collected, used, disclosed and protected according to the provisions of provincial and federal legislation. Your health information is collected by AHS in accordance with section 20 of the Health Information Act (HIA). The purpose of this collection is primarily for: providing health services, determining eligibility for health services, processing payments for health services, conducting research, providing for health services, provider education, internal management purposes, planning and resource allocation, health system management, public health surveillance and health policy development.

Done

Feb 7, 2022

PS

[redacted] from AEFI team 587-773-2035
wants to discuss her symptoms and if any treatments and when D Chan spoke to her
as per dr chan I have made her a telephone appt under pts name
Panel Manager: [redacted] Feb 7, 2022, 9:43AM

Feb 8, 2022

GC

Dr. Gregory Chan, February 8, 2022, 11:52AM
**** COVID-19 telephone conversation ****
discussed AEFI with [redacted]
wanting to clarify, but I simply REITERATED the previous information on the submitted form
Dr. Gregory Chan, February 8, 2022, 11:52AM

AEFI Reporting Form

Note: This form is for health care practitioner use only. If you are a member of the general public and need to report an adverse event following immunization, please call Health Link at 811 or contact your health provider.

1. Today's Date*
19/1/2022

2. Reporter(Your) LastName*
Chan

3. Reporter(Your) FirstName*
Gregory

4. Reporter (Your) SiteType:*
Physician

5. If Other SiteType, please specify:

6. Reporter(Your) Phone#*
(only numbers, no -,())
403-783-3399

7. Immunization Facility

Name	SiteType	If other SiteType, please specify
Pharmasave Clive	Pharmacy	

8. Immunization Facility Phone#
(only numbers, no -,())
403-784-0421

9. Immunizing Facility Address

Building No/Street/PO Box	City/Town	Province	Postalcode(A1A 1A1)

CLIVE

AB

10. Select Zone: (Click [here](#) to determine zone.)

Central

11. Patient LastName*

van Eaton

12. Patient FirstName*

Stephanie

13. If Patient is a minor, then Parent/Guardian

LastName	FirstName	RelationToPatient
----------	-----------	-------------------

14. Patient Sex at Birth:*

F-Female

15. Patient/Guardian Phone#*

(only numbers, no -,())

403-550-4085

16. PHN/ULI Info:

579816051

17. Date Of Birth(dd/mm/yyyy)*

18/01/1989

18. Patient Address:

Building No/Street/POBox	City/Town	Province	PostalCode(A1A 1A1)
BOX 449	CLIVE	AB	T0C 0Y0

19. Date Of Immunization(dd/mm/yyyy):*

25/10/2021

20. Time of Immunization(If Known)(00:00:00)

00:00:00

21. List all the vaccines given on date of immunization.

Immunization Information:

Vaccine Code	Manufacturer	LotNo.
COVPBmR <input type="checkbox"/>	BPF-BioNTi <input type="checkbox"/>	FF2595
-- Please S <input type="checkbox"/>	-- Please S <input type="checkbox"/>	

- Please Select
- Please Select
- Please Select
- Please Select
- Please Select
- Please Select

22. Adverse Event Info:

Symptoms	Started Date (mm/dd/yyyy)	Resolved?	Resolved Date (mm/dd/yyyy)
Other	10/28/2021	Yes	12/01/2021
None		None	
None		None	
None		None	

23. If other, describe including Started date & Resolved date:
numbness around mouth

24. Additional Information:

patient described numbness around the mouth developing days after SECOND DOSE of Pfizer NEVER has symptoms like this previously took several weeks to resolve did not seek medical attention, no treatment

25. Patient is aware that you are reporting the AEFI description and patient contact information to Alberta Health Services AEFI follow-up.*

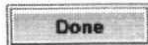
Yes

26. Patient is aware Alberta Health services AEFI program may be contacting them.*

Yes

Note: Hit Done button to Submit the Form.

Alberta Health Services (AHS) respects your confidentiality and privacy. Your information is collected, used, disclosed and protected according to the provisions of provincial and federal legislation. Your health information is collected by AHS in accordance with section 20 of the Health Information Act (HIA). The purpose of this collection is primarily for: providing health services, determining eligibility for health services, processing payments for health services, conducting research, providing for health services, provider education, internal management purposes, planning and resource allocation, health system management, public health surveillance and health policy development.



Jan 31, 2022

GC/PS

Verbal consent given to send AHS requested notes for AEFI
faxed to 1-780-342-0248
Panel Manager: [redacted], Jan 31, 2022, 12:05PM

Jan 31, 2022

PS

[redacted] from AHS-780-702-2653
all info was sent and now asking for
1. Current Diagnosis
2. Treatment plan
3. Assessment plan
I have printed off the AEFI form as I believe all information is there, will confirm with Dr Chan
and if he wants to add anything
Will contact [redacted]
Panel Manager: [redacted], Jan 31, 2022, 3:06PM

Feb 1, 2022

GC

discussion with [redacted] from AEFI
no physical exam done
I explained that the chronology seems to fit with a vaccine injury
It doesn't meet criteria according to AEFI
Dr. Gregory Chan, February 1, 2022, 3:40PM

Feb 2, 2022

PS

AHS asking for reports for pt
advised they were already sent to [redacted] on Jan 31/2022
will send again and will provide AEFI form as well
Panel Manager: [redacted], Feb 2, 2022, 11:38AM

Feb 3, 2022

GC/PS

as per [redacted] at AEFI 780-702-2653
After review no contradiction of COVID 19 Vaccine .
Panel Manager: [redacted], Feb 3, 2022, 4:04PM

AEFI Reporting Form

Note: This form is for health care practitioner use only. If you are a member of the general public and need to report an adverse event following immunization, please call Health Link at 811 or contact your health provider.

1. Today's Date*
19/1/2022

2. Reporter(Your) LastName*
Chan

3. Reporter(Your) FirstName*
Gregory

4. Reporter (Your) SiteType:*
Physician

5. If Other SiteType, please specify:

6. Reporter(Your) Phone#*
(only numbers, no -,())

7. Immunization Facility:

Name	SiteType	If other SiteType, please specify
Rexall Ponoka	Pharmacy	

8. Immunization Facility Phone#
(only numbers, no -,())

9. Immunizing Facility Address

Building No/Street/PO Box	City/Town	Province	Postalcode(A1A 1A1)

10. Select Zone: (Click [here](#) to determine zone.)

Central ▼

11. Patient LastName*

Obers

12. Patient FirstName*

Sandra

13. If Patient is a minor, then Parent/Guardian

LastName	FirstName	RelationToPatient
-	-	-

14. Patient Sex at Birth:*

F-Female ▼

15. Patient/Guardian Phone#*

(only numbers, no -,())

93-783-048

16. PHN/ULI Info:

642641030

17. Date Of Birth(dd/mm/yyyy)*

09/09/1961

18. Patient Address:

Building No/Street/POBox	City/Town	Province	PostalCode(A1A 1A1)
6412-51 STREET	PONOKA	AB	T4J 1E2

19. Date Of Immunization(dd/mm/yyyy):*

03/12/2021

20. Time of Immunization(If Known)(00:00:00)

00:00:00

21. List all the vaccines given on date of immunization.

Immunization Information:

Vaccine Code	Manufacturer	LotNo.
COVPBmR ▼	BPF-BioNTi ▼	
-- Please S ▼	-- Please S ▼	

- Please Sign
- Please Sign
- Please Sign
- Please Sign
- Please Sign

22. Adverse Event Info:

Symptoms	Started Date (mm/dd/yyyy)	Resolved?	Resolved Date (mm/dd/yyyy)
Other <input type="checkbox"/>	12/04/2021	No <input type="checkbox"/>	
None <input type="checkbox"/>		None <input type="checkbox"/>	
None <input type="checkbox"/>		None <input type="checkbox"/>	
None <input type="checkbox"/>		None <input type="checkbox"/>	

23. If other, describe including Started date & Resolved date:

Severe symptoms - fatigue, malaise, myalgias

24. Additional Information:

patient HAD COVID Sept 15, 2021
 was advised to have vaccination anyways, and had 1st dose Dec 3, 2021
 symptoms include myalgia, arthralgias, chest pain, back pain
 progressive each day and week since vaccine
 severe enough to have multiple ER visits Dec 5, Dec 19
 clinic visit Dec 15 - also noted worsening dyspnea; referral to respirology and
 echocardiogram - I was concerned about myocarditis
 echocardiogram normal
 Dec 5 mild elevation of D-dimer 0.54, Hb 115, CRP 56; VQ scan negative for PE
 Dec 15 - seen and advised to try short course of high dose aspirin 325mg daily
 Dec 19 hemoglobin 106
 Jan 13 labwork showed decline in hemoglobin WITHOUT EVIDENCE of hemorrhage (Hb ~94),
 normocytic
 massive elevation of ferritin > 2192 and CRP > 226
 being treated for polymyalgia rheumatica (suspected) with prednisone
 iron for anemia
 awaiting consult for endoscopy to investigate other causes of anemia

25. Patient is aware that you are reporting the AEFI description and patient contact information to Alberta Health Services AEFI follow-up.*

Yes

26. Patient is aware Alberta Health services AEFI program may be contacting them.*

Yes

Note: Hit Done button to Submit the Form.

Alberta Health Services (AHS) respects your confidentiality and privacy. Your information is collected, used, disclosed and protected according to the provisions of provincial and federal legislation. Your health information is collected by AHS in accordance with section 20 of the Health Information Act (HIA). The purpose of this collection is primarily for: providing health services, determining eligibility for health services, processing payments for health services, conducting research, providing for health services, provider education, internal management

purposes, planning and resource allocation, health system management, public health surveillance and health policy development.

Done

Received Date (mm/dd/yyyy)	Received	STANDARD TIME (mm/dd/yyyy)	TYPE OF CASE
>	1st	12/04/2012	Case
>	2nd	>	Case
>	3rd	>	Case
>	4th	>	Case

12. From the following information, identify the following:
Government - Large private hospital

13. A patient with a fever and cough was admitted to the hospital on 12/04/2012. The patient was born in 1975 and has no known medical history. The patient is a male and is currently residing in the United States. The patient was admitted to the hospital with a fever of 101.5 degrees Fahrenheit and a cough that has been present for the past 3 days. The patient also reports a sore throat and a headache. The patient is currently being treated with antibiotics and is expected to be discharged on 12/06/2012.

14. From the following information, identify the following:
Government - Large private hospital

15. From the following information, identify the following:
Government - Large private hospital

Feb 17, 2022

GC/PS

██████████ AEFI

appt has been made to speak with Dr Chan

Panel Manager: ██████████, Feb 17, 2022, 3:51PM

Feb 18, 2022

GC

Dr. Gregory Chan, February 18, 2022, 9:52AM

discussion with AEFI ██████████

discussed Historical/Forensic Science vs Observation/Experimental Science

Dr. Gregory Chan, February 18, 2022, 9:56AM

AEFI Reporting Form

Note: This form is for health care practitioner use only. If you are a member of the general public and need to report an adverse event following immunization, please call Health Link at 811 or contact your health provider.

1. Today's Date*
11/2/2022

2. Reporter(Your) LastName*
CHAN

3. Reporter(Your) FirstName*
GREGORY

4. Reporter (Your) SiteType:*
Physician

5. If Other SiteType, please specify:

6. Reporter(Your) Phone#*
(only numbers, no -,())

7. Immunization Facility

Name	SiteType	If other SiteType, please specify
Ponoka Professional Pharma	Pharmacy	<input checked="" type="checkbox"/>

8. Immunization Facility Phone#
(only numbers, no -,())

9. Immunizing Facility Address

Building No/Street/PO Box	City/Town	Province	Postalcode(A1A 1A1)

10. Select Zone: (Click [here](#) to determine zone.)

Central

11. Patient LastName*

Jones

12. Patient FirstName*

Theresa

13. If Patient is a minor, then Parent/Guardian

LastName	FirstName	RelationToPatient
-	-	-

14. Patient Sex at Birth:*

F-Female

15. Patient/Guardian Phone#*

(only numbers, no -,())

403-783-6495

16. PHN/ULI Info:

787559120

17. Date Of Birth(dd/mm/yyyy)*

04/06/1964

18. Patient Address:

Building No/Street/POBox	City/Town	Province	PostalCode(A1A 1A1)
RT BOX 8, SIT	PONOKA	AB	T4J 1R1

19. Date Of Immunization(dd/mm/yyyy):*

07/01/2022

20. Time of Immunization(If Known)(00:00:00)

00:00:00

21. List all the vaccines given on date of immunization.

Immunization Information:

Vaccine Code	Manufacturer	LotNo.
COVMO <input checked="" type="checkbox"/>	MODTH- <input checked="" type="checkbox"/>	
-- Please <input checked="" type="checkbox"/>	-- Please <input checked="" type="checkbox"/>	

- Please
- Please
- Please
- Please
- Please

22. Adverse Event Info:

Symptoms	Started Date (mm/dd/yyyy)	Resolved?	Resolved Date (mm/dd/yyyy)
Pain and/or: <input checked="" type="checkbox"/>	01/09/2022	No <input checked="" type="checkbox"/>	
None <input checked="" type="checkbox"/>		None <input checked="" type="checkbox"/>	
None <input checked="" type="checkbox"/>		None <input checked="" type="checkbox"/>	
None <input checked="" type="checkbox"/>		None <input checked="" type="checkbox"/>	

23. If other, describe including Started date & Resolved date:
 shoulder pain

24. Additional Information:

patient had her booster injection Jan 7/22, and within 2-3 days had shoulder pain that was significantly worse, subjectively different from previous shoulder pain lasted 3-4 weeks
 now has pain that migrates from one area of the body to another, mild in severity similar episode in July/Aug 2021 after 2nd dose in June 2021 but it was MILD had pain in the shoulder/back after dry needling, stopped once she stopped dry needling never had muscle/arm/back pain like this with previous immunizations

25. Patient is aware that you are reporting the AEFI description and patient contact information to Alberta Health Services AEFI follow-up.*

Yes

26. Patient is aware Alberta Health services AEFI program may be contacting them.*

Yes

Note: Hit Done button to Submit the Form.

Alberta Health Services (AHS) respects your confidentiality and privacy. Your information is collected, used, disclosed and protected according to the provisions of provincial and federal legislation. Your health information is collected by AHS in accordance with section 20 of the Health Information Act (HIA). The purpose of this collection is primarily for: providing health services, determining eligibility for health services, processing payments for health services, conducting research, providing for health services, provider education, internal management purposes, planning and resource allocation, health system management, public health surveillance and health policy development.



[REDACTED]

Feb 22, 2022

GC/PS

[REDACTED] - AEFI 587-782-4345

"pt can go ahead with future vaccines for MRNA/Covid"
noet to Dr Chan

Panel Manager: [REDACTED] Feb 22, 2022, 11:33AM

AEFI Reporting Form

Note: This form is for health care practitioner use only. If you are a member of the general public and need to report an adverse event following immunization, please call Health Link at 811 or contact your health provider.

1. Today's Date*
11/2/2022

2. Reporter(Your) LastName*
CHAN

3. Reporter(Your) FirstName*
GREGORY

4. Reporter (Your) SiteType:*
Physician

5. If Other SiteType, please specify:

6. Reporter(Your) Phone#*
(only numbers, no -,())

7. Immunization Facility

Name	SiteType	If other SiteType, please specify
No Frills	Pharmacy <input checked="" type="checkbox"/>	

8. Immunization Facility Phone#
(only numbers, no -,())

9. Immunizing Facility Address

Building No/Street/PO Box	City/Town	Province	Postalcode(A1A 1A1)

5561 HWY 53

PONOKA

AB

T4J 1J8

10. Select Zone: (Click [here](#) to determine zone.)

Central

11. Patient LastName*

[REDACTED]

12. Patient FirstName*

SHELLY

13. If Patient is a minor, then Parent/Guardian

LastName	FirstName	RelationToPatient
-	-	-

14. Patient Sex at Birth:*

F-Female

15. Patient/Guardian Phone#*

(only numbers, no -,())

403-598-0198

16. PHN/ULI Info:

63300720

17. Date Of Birth(dd/mm/yyyy)*

10/10/1963

18. Patient Address:

Building No/Street/POBox	City/Town	Province	PostalCode(A1A 1A1)
1R4	PONOKA	AB	T4J 1R4

19. Date Of Immunization(dd/mm/yyyy):*

09/01/2022

20. Time of Immunization(If Known)(00:00:00)

00:00:00

21. List all the vaccines given on date of immunization.

Immunization Information:

Vaccine Code	Manufacturer	LotNo.
COVPBr <input checked="" type="checkbox"/>	BPF-Biol <input checked="" type="checkbox"/>	
-- Please <input checked="" type="checkbox"/>	-- Please <input checked="" type="checkbox"/>	

- Please
- Please
- Please

22. Adverse Event Info:

Symptoms	Started Date (mm/dd/yyyy)	Resolved?	Resolved Date (mm/dd/yyyy)
Other <input type="checkbox"/>	01/11/2022	No <input type="checkbox"/>	
None <input type="checkbox"/>		None <input type="checkbox"/>	
None <input type="checkbox"/>		None <input type="checkbox"/>	
None <input type="checkbox"/>		None <input type="checkbox"/>	

23. If other, describe including Started date & Resolved date:

hot flashes

24. Additional Information:

has had menopause and hot flashes which RESOLVED about 1.5 years ago had vaccine, 1st dose Jan 9/22 - then has had hot flashes again for the last 4 weeks, not resolving

25. Patient is aware that you are reporting the AEFI description and patient contact information to Alberta Health Services AEFI follow-up.*

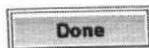
Yes

26. Patient is aware Alberta Health services AEFI program may be contacting them.*

Yes

Note: Hit Done button to Submit the Form.

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AEFI Reporting Form

Note: This form is for health care practitioner use only. If you are a member of the general public and need to report an adverse event following immunization, please call Health Link at 811 or contact your health provider.

1. Today's Date*

17/4/2022

2. Reporter(Your) LastName*

Chan

3. Reporter(Your) FirstName*

Gregory

4. Reporter (Your) SiteType:*

AHS Acute Care

5. If Other SiteType, please specify:

6. Reporter(Your) Phone#*

(only numbers, no -,())

7. Immunization Facility:

Name	SiteType	If other SiteType, please specify
Centennial Center for Mental	AHS Acute Care	

8. Immunization Facility Phone#

(only numbers, no -,())

9. Immunizing Facility Address

Building No/Street/PO Box	City/Town	Province	Postalcode(A1A 1A1)
---------------------------	-----------	----------	---------------------

46 STREET S

PONOKA

AB

T4J 1R8

10. Select Zone: (Click [here](#) to determine zone.)

Central ▼

11. Patient LastName*

Smith

12. Patient FirstName*

Greg

13. If Patient is a minor, then Parent/Guardian

LastName

FirstName

RelationToPatient

-

-

-

14. Patient Sex at Birth:*

M-Male ▼

15. Patient/Guardian Phone#*

(only numbers, no -,())

403-396-3702

16. PHN/ULI Info:

482304720

17. Date Of Birth(dd/mm/yyyy)*

05/01/1984

18. Patient Address:

Building
No/Street/POBox

City/Town

Province

PostalCode(A1A
1A1)

5102-51 STREET

PONOKA

AB

T4J 1E7

19. Date Of Immunization(dd/mm/yyyy):*

09/09/2021

20. Time of Immunization(If Known)(00:00:00)

00:00:00

21. List all the vaccines given on date of immunization.

Immunization Information:

Vaccine Code

Manufacturer

LotNo.

COVPBmRN ▼

BPF-BioNTec ▼

-- Please Sel ▼

-- Please Sel ▼

-- Please Sel ▼	-- Please Sel ▼	
-- Please Sel ▼	-- Please Sel ▼	
-- Please Sel ▼	-- Please Sel ▼	

22. Adverse Event Info:

Symptoms	Started Date (mm/dd/yyyy)	Resolved?	Resolved Date (mm/dd/yyyy)
Other ▼	09/10/2022	No ▼	
None ▼		None ▼	
None ▼		None ▼	
None ▼		None ▼	

23. If other, describe including Started date & Resolved date:

change in bowel consistency

24. Additional Information:

prior to 1st COVID injection - normal bowel movement size and consistency was admitted to hospital (Centennial Center for Mental Health and Brain Injury), was given 1st injection as an inpatient within 24 hours of 1st injection - had smaller size flat bowel movements with episodes of diarrhea and small volume stool chronic abdominal pain, Left lower quadrant and left flank worse in the last 1-2 months presented to ER with abdominal pain and reported change in stool Slight increase in WBC/neutrophil count Slight elevation of ALT Plan - needs colonoscopy and abdominal US

Summary - since 1st injection, bowel movements have significantly changed in consistency and frequency, and have not returned to pre-injection bowel movement consistency/frequency

25. Patient is aware that you are reporting the AEFI description and patient contact information to Alberta Health Services AEFI follow-up.*

Yes ▼

26. Patient is aware Alberta Health services AEFI program may be contacting them.*

Yes ▼

Note: Hit Done button to Submit the Form.

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AEFI Reporting Form

Note: This form is for health care practitioner use only. If you are a member of the general public and need to report an adverse event following immunization, please call Health Link at 811 or contact your health provider.

1. Today's Date*

24/5/2022

2. Reporter(Your) LastName*

CHAN

3. Reporter(Your) FirstName*

GREGORY

4. Reporter (Your) SiteType:*

Physician

5. If Other SiteType, please specify:

6. Reporter(Your) Phone#*

(only numbers, no -,())

7. Immunization Facility:

Name

SiteType

**If other SiteType, please
specify**

Maskwacis Health Services

First Nations and Inuit He: ▼

8. Immunization Facility Phone#

(only numbers, no -,())

780-585-3830

9. Immunizing Facility Address

**Building
No/Street/PO
Box**

City/Town

Province

**Postalcode(A1A
1A1)**

MINDE AVE & W

MASKWACIS

AB

TOC 1N0

10. Select Zone: (Click [here](#) to determine zone.)

Central ▼

11. Patient LastName*

WOLFE

12. Patient FirstName*

CHRISTINA

13. If Patient is a minor, then Parent/Guardian

LastName

FirstName

RelationToPatient

-

14. Patient Sex at Birth:*

F-Female ▼

15. Patient/Guardian Phone#*

(only numbers, no -,())

587-729-0145

16. PHN/ULI Info:

927614310

17. Date Of Birth(dd/mm/yyyy)*

28/03/1956

18. Patient Address:

Building
No/Street/POBox

City/Town

Province

PostalCode(A1A
1A1)

5210-49 AVE

PONOKA

AB

T4J 1J1

19. Date Of Immunization(dd/mm/yyyy):*

11/05/2021

20. Time of Immunization(If Known)(00:00:00)

00:00:00

21. List all the vaccines given on date of immunization.

Immunization Information:

Vaccine Code

Manufacturer

LotNo.

COVMODmF ▼

MODTH-Moc ▼

-- Please Sel ▼

-- Please Sel ▼

-- Please Sel ▼ -- Please Sel ▼

-- Please Sel ▼ -- Please Sel ▼

-- Please Sel ▼ -- Please Sel ▼

22. Adverse Event Info:

Symptoms	Started Date (mm/dd/yyyy)	Resolved?	Resolved Date (mm/dd/yyyy)
Rash ▼	05/18/2021	No ▼	
None ▼		None ▼	
None ▼		None ▼	
None ▼		None ▼	

23. If other, describe including Started date & Resolved date:

Rash

24. Additional Information:

rash developed in LEFT antecubital fossa 1 week after 2nd MODERNA COVID vaccine
patient has never had eczema
rash has slowly increased in size, about 5x6cm

25. Patient is aware that you are reporting the AEFI description and patient contact information to Alberta Health Services AEFI follow-up.*

Yes ▼

26. Patient is aware Alberta Health services AEFI program may be contacting them.*

Yes ▼

Note:Hit Done button to Submit the Form.

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Done

Sep 19, 2022

GC

discussed with [REDACTED]
does not meet criteria because no bloodwork drawn immediately after 3rd dose
documented
recommendation to continue with the vaccines
Dr. Gregory Chan, September 19, 2022, 1:59PM

AEFI Reporting Form

Note: This form is for health care practitioners use only. If you are a member of the general public and need to report an adverse event following immunization, please call Health Link at 811 or contact your health provider.

Page 1 of 1

2022-09-15, 10:31 AM

1. Today's Date
15/9/2022
2. Reporter(Your) LastName*
CHAN
3. Reporter(Your) FirstName*
GREGORY
4. Reporter (Your) SiteType:*
Physician
5. If Other SiteType, please specify:
6. Reporter(Your) Phone#*
(only numbers, no -. (0))
7. Immunization Facility:

Name	SiteType	If other SiteType, please specify
Shoppers Drug Mart	Pharmacy	
8. Immunization Facility Phone#
(only numbers, no -. (0))

2022-09-15, 10:31 AM

9. Immunizing Facility Address

Building	No./Street/ PO Box	City/Town	Province	PostalCode(A1A 1A1)
	5015-50 STREET	PONOKA	AB	T4L 0C1
10. Select Zone: (Click here to determine zone.)
Calgary
11. Patient LastName*
[Redacted]
12. Patient FirstName*
[Redacted]
13. If Patient is a minor, then Parent/Guardian

Last Name	First Name	Relation to Patient
[Redacted]	[Redacted]	[Redacted]
14. Patient Sex at Birth:*
F-Female
15. Patient/Guardian Phone#*
(only numbers, no -. (0))
16. PHN/ULL Info:
[Redacted]
17. Date Of Birth(dd/mm/yyyy)*
[Redacted]
18. Patient Address:

Building	No./Street/POBox	City/Town	Province	PostalCode(A1A 1A1)
[Redacted]	[Redacted]	[Redacted]	[Redacted]	[Redacted]

19. Date Of Immunization(dd/mm/yyyy):-

23/12/2021

20. Time of Immunization(If Known)(00:00:00)

14:25:00

21. List all the vaccines given on date of immunization.

Immunization Information:

Vaccine Code	Manufacturer	LotNo.
COVBRNA	BPF-BioNTec	28-227-DK
-- Please Sel	-- Please Sel	
-- Please Sel	-- Please Sel	
-- Please Sel	-- Please Sel	

22. Adverse Event Info:

Symptoms	Started Date (mm/dd/yyyy)	Resolved? No	Resolved Date (mm/dd/yyyy)
Rash	01/10/2022	<input type="radio"/>	
None		<input type="radio"/>	
None		<input type="radio"/>	
None		<input type="radio"/>	

23. If other, describe including Started date & Resolved date:

bruising, ecchymosis

24. Additional Information:

about 2 weeks after booster injection - bruising and ecchymosis with minimal trauma not present prior to booster injection

25. Patient is aware that you are reporting the AEFI description and patient contact information to Alberta Health Services AEFI follow-up.

Yes

26. Patient is aware Alberta Health services AEFI program may be contacting them.

Yes

Note: Hit Done button to Submit the Form.

Alberta Health Services (AHS) collects your information and privacy information collected, used, disclosed and protected according to the provisions of provincial and federal legislation. Your health information is collected by AHS in accordance with section 20 of the Health Information Act (HIA). The purpose of this collection is primarily for: providing health services, determining eligibility for health services, processing payments for health services, conducting research, providing for health services, provider education, internal management purposes, planning and resource allocation, health system management, public health surveillance and health policy development.

Done

Feb 11/23
@ 2:17.

→ away till
Monday
can call then?

[REDACTED] RH 69460
Ponoka, AB [REDACTED] M [REDACTED]
[REDACTED]

[REDACTED] @ AHS.
[REDACTED]

AFI outcome
Does not meet
criteria for
myocarditis.

All cardio tests were
Normal

she asks that you
review the "criteria"
on the AHS website

A

Ps
I did advise
her that I was
sure you were
fully aware of
the "criteria"
A

AEFI Reporting Form

Note: This form is for health care practitioner use only. If you are a member of the general public and need to report an adverse event following immunization, please call Health Link at 811 or contact your health provider.

1. Today's Date*

30/12/2022

2. Reporter(Your) LastName*

CHAN

3. Reporter(Your) FirstName*

Gregory

4. Reporter (Your) SiteType:*

AHS Acute Care

5. If Other SiteType, please specify:

6. Reporter(Your) Phone#*

(only numbers, no -,())

7. Immunization Facility:

Name	SiteType	If other SiteType, please specify
Rexall Ponoka	Pharmacy	

8. Immunization Facility Phone#

(only numbers, no -,())

9. Immunizing Facility Address

Building No/Street/PO Box	City/Town	Province	Postalcode(A1A 1A1)
---------------------------	-----------	----------	---------------------

4502 50 ST

PONOKA

AB

T4J 1J5

10. Select Zone: (Click [here](#) to determine zone.)

Central ▼

11. Patient LastName*

MCCOMBS

12. Patient FirstName*

DUSTIN

13. If Patient is a minor, then Parent/Guardian

LastName

FirstName

RelationToPatient

--	--	--

14. Patient Sex at Birth:*

M-Male ▼

15. Patient/Guardian Phone#*

(only numbers, no -,())

519-718-4059

16. PHN/ULI Info:

679624071

17. Date Of Birth(dd/mm/yyyy)*

7/07/1985

18. Patient Address:

Building
No/Street/POBox

City/Town

Province

PostalCode(A1A
1A1)

4403-43 STREET

PONOKA

AB

T4J 1B8

19. Date Of Immunization(dd/mm/yyyy):*

22/10/21

20. Time of Immunization(If Known)(00:00:00)

00:00:00

21. List all the vaccines given on date of immunization.

Immunization Information:

Vaccine Code

Manufacturer

LotNo.

COVPBmRN ▼

BPF-BioNTec ▼

-- Please Sel ▼

-- Please Sel ▼

-- Please Sel ▼ -- Please Sel ▼

-- Please Sel ▼ -- Please Sel ▼

-- Please Sel ▼ -- Please Sel ▼

22. Adverse Event Info:

Symptoms	Started Date (mm/dd/yyyy)	Resolved?	Resolved Date (mm/dd/yyyy)
Pain and/or Swe ▼	10/24/2021	No ▼	
Other ▼	10/24/2021	No ▼	
None ▼		None ▼	
None ▼		None ▼	

23. If other, describe including Started date & Resolved date:

dyspnea, palpitations, dizziness

24. Additional Information:

Patient had 1st dose of PFIZER and 2 days later. Intermittent symptoms. Multiple ER and GP visits. Cardiology consultation Dec 5/21. Echocardiogram, Dec 2/21 and cardiac MRI Nov 25/22 unremarkable, though the tests occurred much after the vaccination event.

25. Patient is aware that you are reporting the AEFI description and patient contact information to Alberta Health Services AEFI follow-up.*

Yes ▼

26. Patient is aware Alberta Health services AEFI program may be contacting them.*

Yes ▼

Note:Hit Done button to Submit the Form.

Alberta Health Services (AHS) respects your confidentiality and privacy. Your information is collected, used, disclosed and protected according to the provisions of provincial and federal legislation. Your health information is collected by AHS in accordance with section 20 of the Health Information Act (HIA). The purpose of this collection is primarily for: providing health services, determining eligibility for health services, processing payments for health services, conducting research, providing for health services, provider education, internal management purposes, planning and resource allocation, health system management, public health surveillance and health policy development.

Done

AEFI Reporting Form

Note: This form is for health care practitioner use only. If you are a member of the general public and need to report an adverse event following immunization, please call Health Link at 811 or contact your health provider.

1. Today's Date*

17/1/2023

2. Reporter(Your) LastName*

CHAN

3. Reporter(Your) FirstName*

GREGORY

4. Reporter (Your) SiteType:*

Physician

5. If Other SiteType, please specify:

6. Reporter(Your) Phone# *

(only numbers, no -,())

7. Immunization Facility

Name

SiteType

If other SiteType, please specify

Pharmacy

Pharmacy

unknown

8. Immunization Facility Phone#

(only numbers, no -,())

9. Immunizing Facility Address

Building
No/Street/PO
Box

City/Town

Province

Postalcode(A1A
1A1)

-

PONOKA

AB

10. Select Zone: (Click [here](#) to determine zone.)

Central

11. Patient LastName*

WALSH

12. Patient FirstName*

PHIM

13. If Patient is a minor, then Parent/Guardian

LastName

FirstName

RelationToPatient

-

-

-

14. Patient Sex at Birth:*

F-Female

15. Patient/Guardian Phone#*

(only numbers, no -,())

403-318-1389

16. PHN/ULI Info:

588505510

17. Date Of Birth(dd/mm/yyyy)*

23/02/1951

18. Patient Address:

Building No/Street/POBox

City/Town

Province

PostalCode(A1A 1A1)

1000

PONOKA

AB

T4J 1C9

19. Date Of Immunization(dd/mm/yyyy):*

16/03/2021

20. Time of Immunization(If Known)(00:00:00)

00:00:00

21. List all the vaccines given on date of immunization.

Immunization Information:

Vaccine Code

Manufacturer

LotNo.

COVMODm

MODTH-Mc

-- Please Se

-- Please Se

-- Please Se ▼	-- Please Se ▼	
-- Please Se ▼	-- Please Se ▼	
-- Please Se ▼	-- Please Se ▼	

22. Adverse Event Info:

Symptoms	Started Date (mm/dd/yyyy)	Resolved?	Resolved Date (mm/dd/yyyy)
Pain and/or Swε ▼	03/16/2021	Yes ▼	03/20/2021
Other ▼	03/16/2021	Yes ▼	03/20/2021
None ▼		None ▼	
None ▼		None ▼	

23. If other, describe including Started date & Resolved date:

diarrhea, weakness

24. Additional Information:

left sided face scalp face pain, then trigger migraine
 crawling on floor
 diarrhea
 recovered after about 1 week
 no reaction with Pfizer, went to Calgary 'in case she had an adverse reaction'

25. Patient is aware that you are reporting the AEFI description and patient contact information to Alberta Health Services AEFI follow-up.*

Yes ▼

26. Patient is aware Alberta Health services AEFI program may be contacting them.*

Yes ▼

Note:Hit Done button to Submit the Form.

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Done



AEFI Reporting Form

Note: This form is for health care practitioner use only. If you are a member of the general public and need to report an adverse event following immunization, please call Health Link at 811 or contact your health provider.

1. Today's Date*

10/2/2023

2. Reporter(Your) LastName*

CHAN

3. Reporter(Your) FirstName*

GREGORY

4. Reporter (Your) SiteType:*

Physician

5. If Other SiteType, please specify:

6. Reporter(Your) Phone#*

(only numbers, no -,())

[REDACTED]

7. Immunization Facility:

	Name	SiteType	If other SiteType, please specify
	UNKNOWN	-- Please Select --	

8. Immunization Facility Phone#

(only numbers, no -,())

9. Immunizing Facility Address

Building No/Street/PO Box	City/Town	Province	Postalcode(A1A 1A1)
--	------------------	-----------------	--------------------------------

- - -



AEFI Reporting Form

10. Select Zone: (Click [here](#) to determine zone.)

Central

11. Patient LastName*

12. Patient FirstName*

13. If Patient is a minor, then Parent/Guardian

LastName	FirstName	RelationToPatient
<input type="text" value="-"/>	<input type="text" value="-"/>	<input type="text" value="-"/>

14. Patient Sex at Birth:*

M-Male

15. Patient/Guardian Phone#*

(only numbers, no -,())

16. PHN/ULI Info:

17. Date Of Birth(dd/mm/yyyy)*

18. Patient Address:

Building No/Street /POBox	City/Town	Province	PostalCode(A1A 1A1)
<input type="text" value="CX 6/ SITE 10"/>	<input type="text" value="PONOMA"/>	<input type="text" value="AB"/>	<input type="text" value="T4J 1R3"/>

19. Date Of Immunization(dd/mm/yyyy):*

20. Time of Immunization(If Known)(00:00:00)

21. List all the vaccines given on date of immunization.

Immunization Information:

Vaccine Code	Manufacturer	LotNo.
<input type="button" value="COVPBmRN/ v"/>	<input type="button" value="BPF-BioNTec v"/>	<input type="text"/>

-- Please Sel ▾	-- Please Sel ▾	
-- Please Sel ▾	-- Please Sel ▾	
-- Please Sel ▾	-- Please Sel ▾	
-- Please Sel ▾	-- Please Sel ▾	

22. Adverse Event Info:

Symptoms	Started Date (mm/dd /yyyy)	Resolved?	Resolved Date (mm/dd /yyyy)
Other ▾	12/01/2021	Yes ▾	04/01/2022
Other ▾	03/16/2022	Yes ▾	18/07/2022
None ▾		None ▾	
None ▾		None ▾	

23. If other, describe including Started date & Resolved date:

1. CHEST PAIN, 2. elevated ferritin

24. Additional Information:

1. Chest pain - within 2 weeks of 2nd Pfizer COVID mRNA injection - developed chest pain when working under a car. Never had chest pain like this previously. Had seen his GP and she ordered a cardiology consultation.
 As per notes from Netcare Cardiology consultation:
 "As you know, this gentleman experienced some chest pain in early December of 2021. He was under a car changing oil. The pain lasted for about 20 minutes. There was a sensation of cramping over the area of his heart. There was no associated shortness of breath, nausea, or diaphoresis. He had further pain in early March of this year, which again lasted for about 20 minutes. He had been lifting some heavy boxes. There is no associated shortness of breath or nausea."
 Chest pain has since resolved, no exertional chest pain or pain at rest

2. Elevated ferritin.
 Known to have slightly elevated ferritin. Investigated prior to vaccination and max was mid 600s. After vaccination his March 2022 Ferritin was > 1100. After phlebotomy the number has reduced to normal range. Still followed by gastroenterologist in Red Deer

25. Patient is aware that you are reporting the AEFI description and patient contact information to Alberta Health Services AEFI follow-up.*

-- Please Select -- ▾

26. Patient is aware Alberta Health services AEFI program may be contacting them.*

-- Please Select -- ▾

Note: Hit Done button to Submit the Form.

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Resolved Date (mm/dd/yyyy)

Assigned

Assigned Date (mm/dd/yyyy)

System

Done

Additional Information

1. I first pain - within 2 weeks of the first COVID-19 infection - developed chest pain when working under a car. However, the chest pain like this previously had not hit me and was not related to a coronary artery disease.

As you know, this guideline requires an acute chest pain in early diagnosis of MI. He was under a car working with the pain lasted for about 30 minutes. There was a reduction of tearing over the area of his heart. There was no associated symptoms of cardiac arrest or dyspnea. He had further pain in early part of this year, when again lasted for about 30 minutes. It was not lifting some heavy boxes, there is no associated symptoms of breath or tearing. The pain has since resolved, no recurrent chest pain or risk at rest.

1 - Elevated troponin

When I had slightly elevated troponin. Troponin level after the infection and was also 0.009. After vaccination his troponin level was 0.109. After laboratory the number was reduced to normal range. Still followed by cardiac troponin to see how

25. Patient is aware that you are reporting the AHS description and patient contact information to Alberta Health Services AHS follow up.

26. I speak to Alberta Health Services AHS program that is contacting them.

AEFI Reporting Form

Note: This form is for health care practitioner use only. If you are a member of the general public and need to report an adverse event following immunization, please call Health Link at 811 or contact your health provider.

1. Today's Date*
14/11/2021

2. Reporter(Your) LastName*
CHAN

3. Reporter(Your) FirstName*
GREGORY

4. Reporter (Your) SiteType:*
Physician

5. If Other SiteType, please specify:

6. Reporter(Your) Phone#*
(only numbers, no -,())
[REDACTED]

7. Immunization Facility:

Name	SiteType	If other SiteType, please specify
Roots & Berries Pharmacy	Pharmacy	

8. Immunization Facility Phone#
(only numbers, no -,())
[REDACTED]

9. Immunizing Facility Address

Building No/Street/PO Box City/Town Province Postalcode(A1A 1A1)

10. Select Zone: (Click [here](#) to determine zone.)

Calgary ▼

11. Patient LastName*

BULL

12. Patient FirstName*

GREGORY

13. If Patient is a minor, then Parent/Guardian

LastName

FirstName

RelationToPatient

- - -

14. Patient Sex at Birth:*

M-Male ▼

15. Patient/Guardian Phone#*

(only numbers, no -,())

780-335-8681

16. PHN/ULI Info:

739795090

17. Date Of Birth(dd/mm/yyyy)*

27/04/1997

18. Patient Address:

Building
No/Street/POBox

City/Town

Province

PostalCode(A1A
1A1)

BOX 488 MASKWACIS AB TOC 1N0

19. Date Of Immunization(dd/mm/yyyy):*

18/10/2021

20. Time of Immunization(If Known)(00:00:00)

00:00:00

21. List all the vaccines given on date of immunization.

Immunization Information:

Vaccine Code

Manufacturer

LotNo.

COVPBmRN ▼

BPF-BioNTex ▼

-- Please Sel ▼

-- Please Sel ▼

-- Please Sel ▼ -- Please Sel ▼
 -- Please Sel ▼ -- Please Sel ▼
 -- Please Sel ▼ -- Please Sel ▼

22. Adverse Event Info:

Symptoms	Started Date (mm/dd/yyyy)	Resolved?	Resolved Date (mm/dd/yyyy)
Severe Diarrhea ▼	11/06/2021	No ▼	
None ▼		None ▼	
None ▼		None ▼	
None ▼		None ▼	

23. If other, describe including Started date & Resolved date:

diarrhea, cough, congestion

24. Additional Information:

2nd dose of PFIZER
 onset Nov 6/21 - diarrhea (non bloody, about 3 per day, no mucous), abdominal pain, cough (non-productive), sore throat (this has resolved on Nov 11/21)
 diarrhea has not resolved

25. Patient is aware that you are reporting the AEFI description and patient contact information to Alberta Health Services AEFI follow-up.*

Yes ▼

26. Patient is aware Alberta Health services AEFI program may be contacting them.*

Yes ▼

Note: Hit Done button to Submit the Form.

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Done