



Mpox Outbreak Affecting Many Fully Vaccinated Men: Chicago, May–June 2023

Emily A. G. Faherty, PhD
Epidemic Intelligence Service Officer
Chicago Department of Public Health

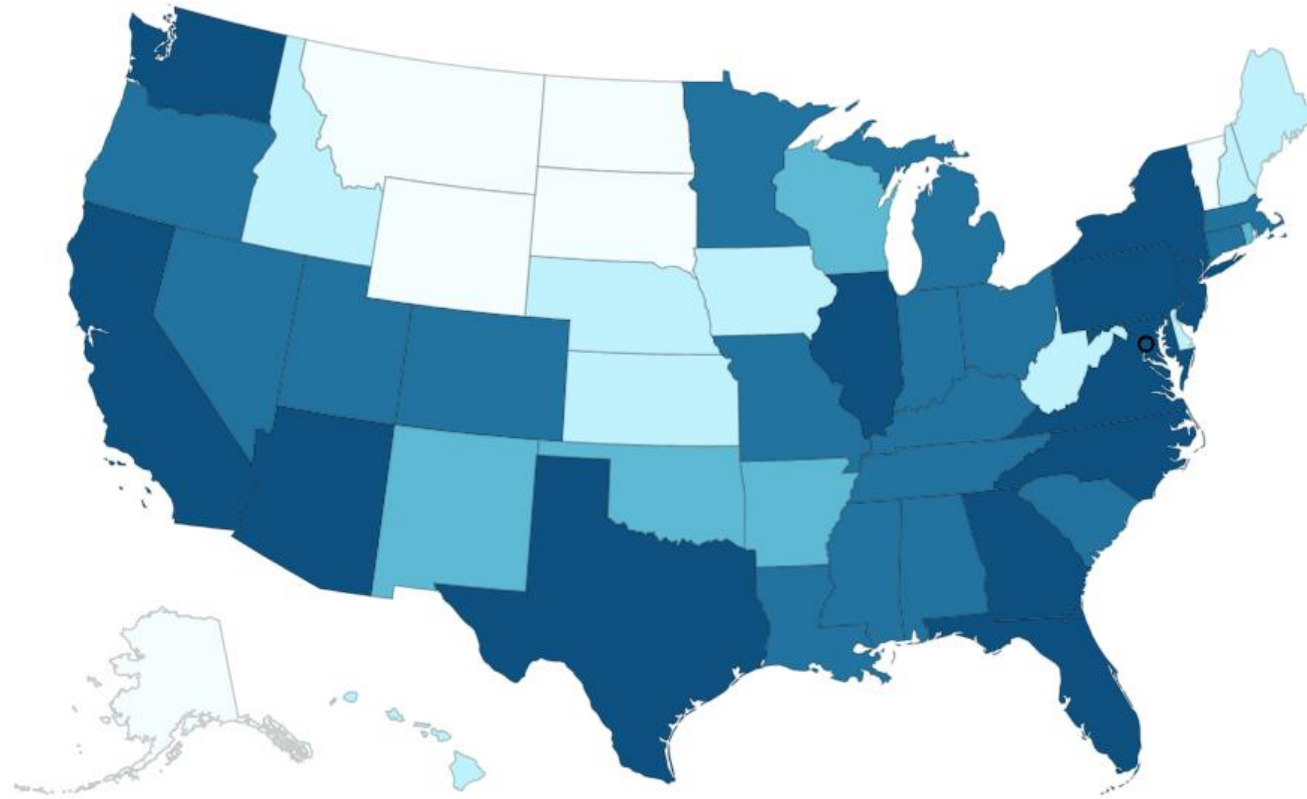
CDPH SID Bureau Provider Conference
June 27, 2024

Disclosure:

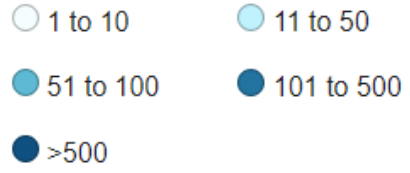
Presenter has no financial interest to disclose.

This continuing education activity is managed by The St. Louis STI/HIV Prevention Training Center and accredited by Missouri State Medical Association (MSMA) in cooperation with the Chicago Department of Public Health

There have been over 32,000 cases of mpox in the United States since 2022.



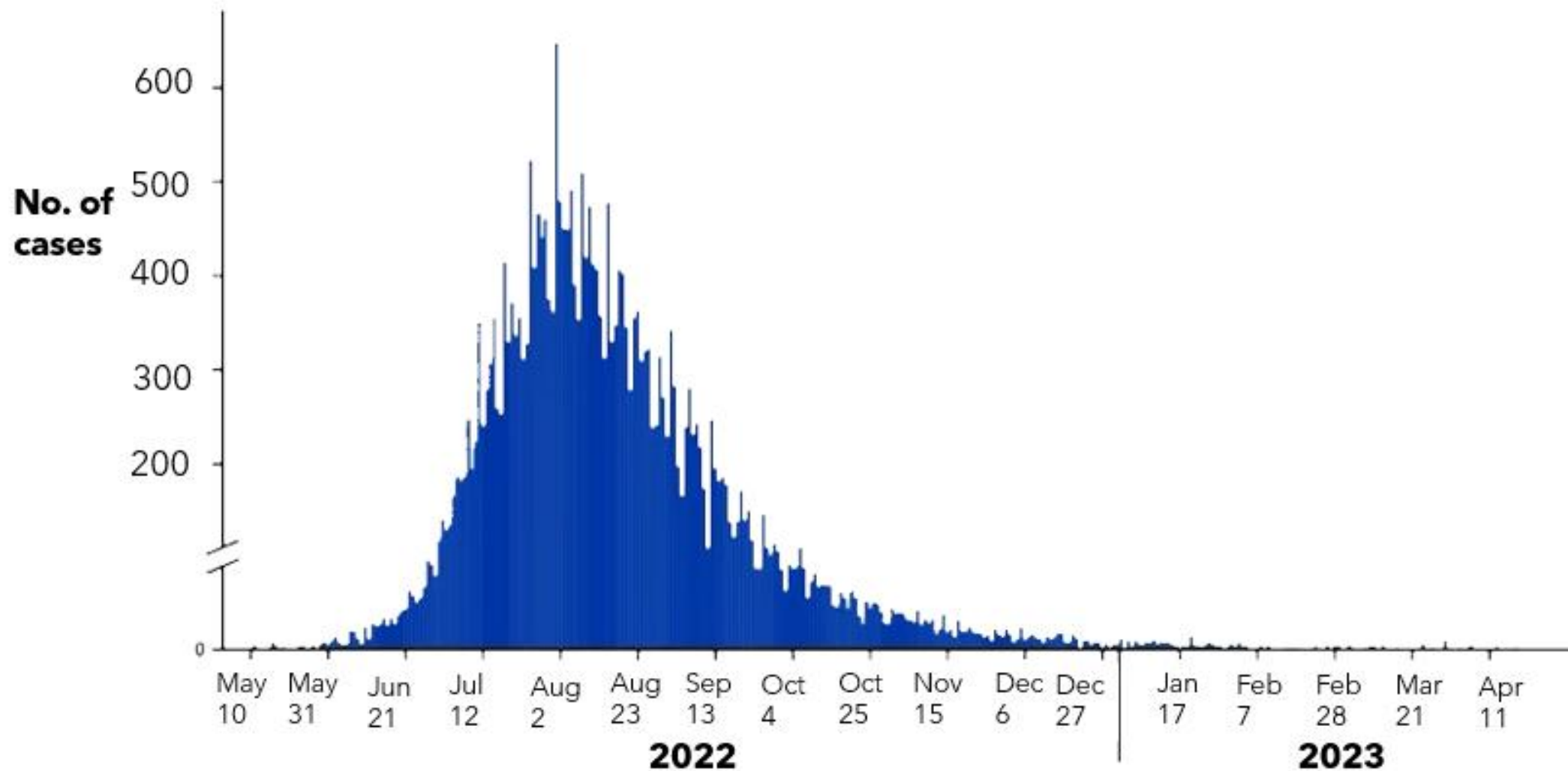
PR



Affected countries promoted vaccination and harm reduction to control mpox.



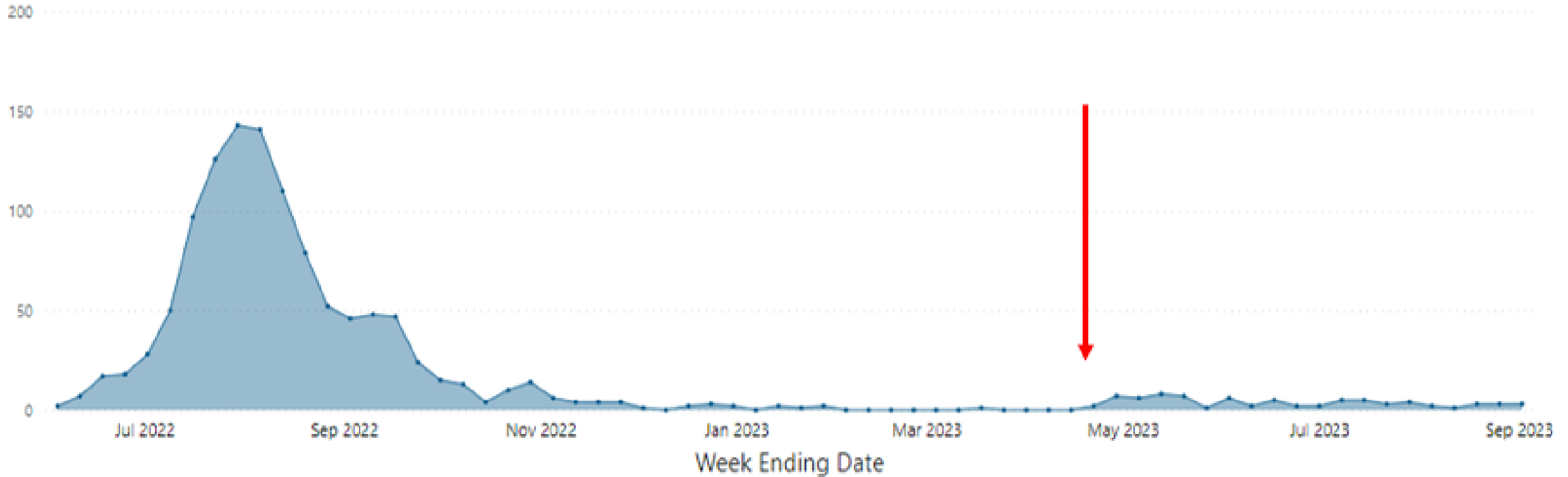
The mpox outbreak in 2023 had declined in most parts of the United States by early 2023.



McQuiston JH, Braden CR, Bowen MD, et al. The CDC Domestic Mpox Response – United States, 2022-2023. *MMWR Morb Mortal Wkly Rep* 2023;72:547-552. DOI: <http://dx.doi.org/10.15585/mmwr.mm7220a2>

In May 2023, an increased number of cases were reported in Chicago.

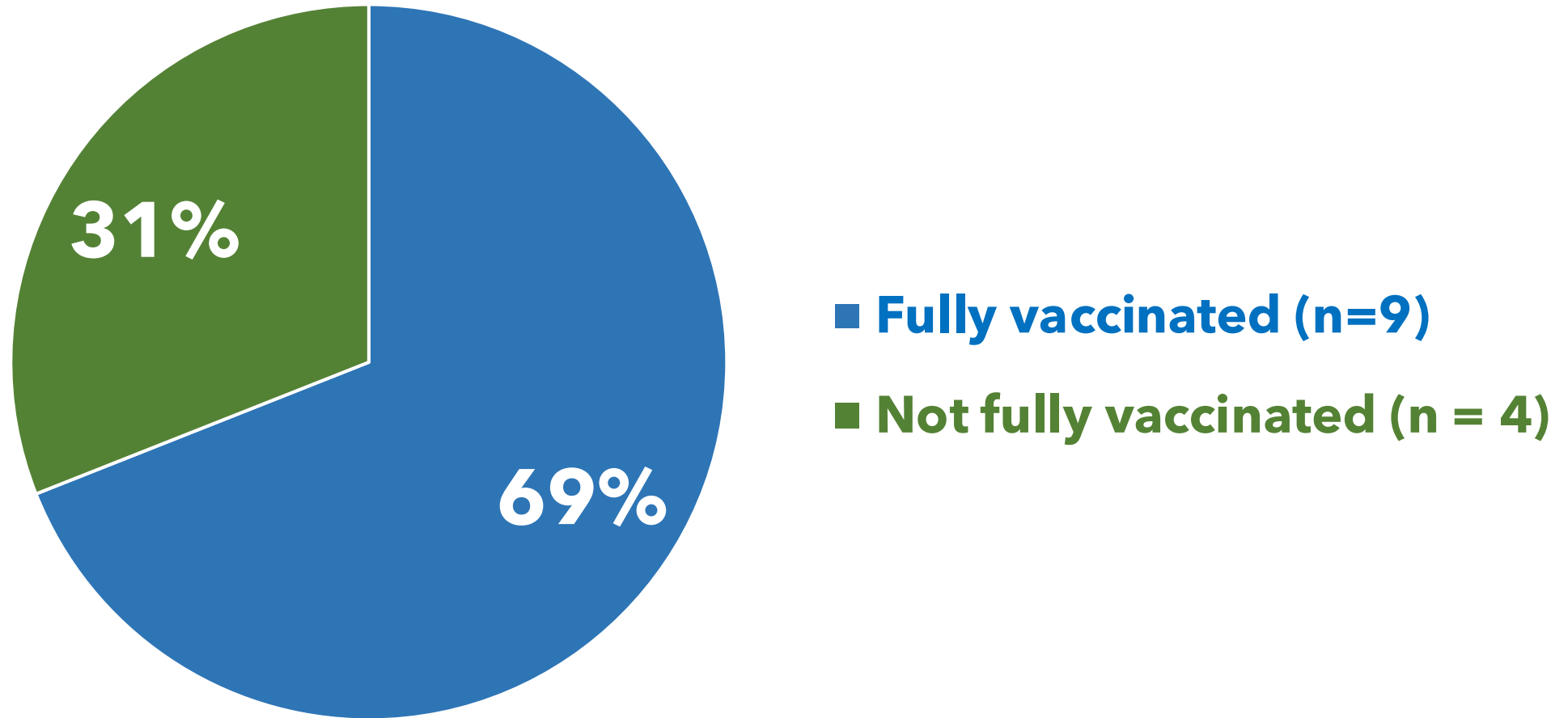
Cases of mpox diagnosed in Chicagoans during 6/4/2022-8/31/2023



CDPH Mpox Dashboard

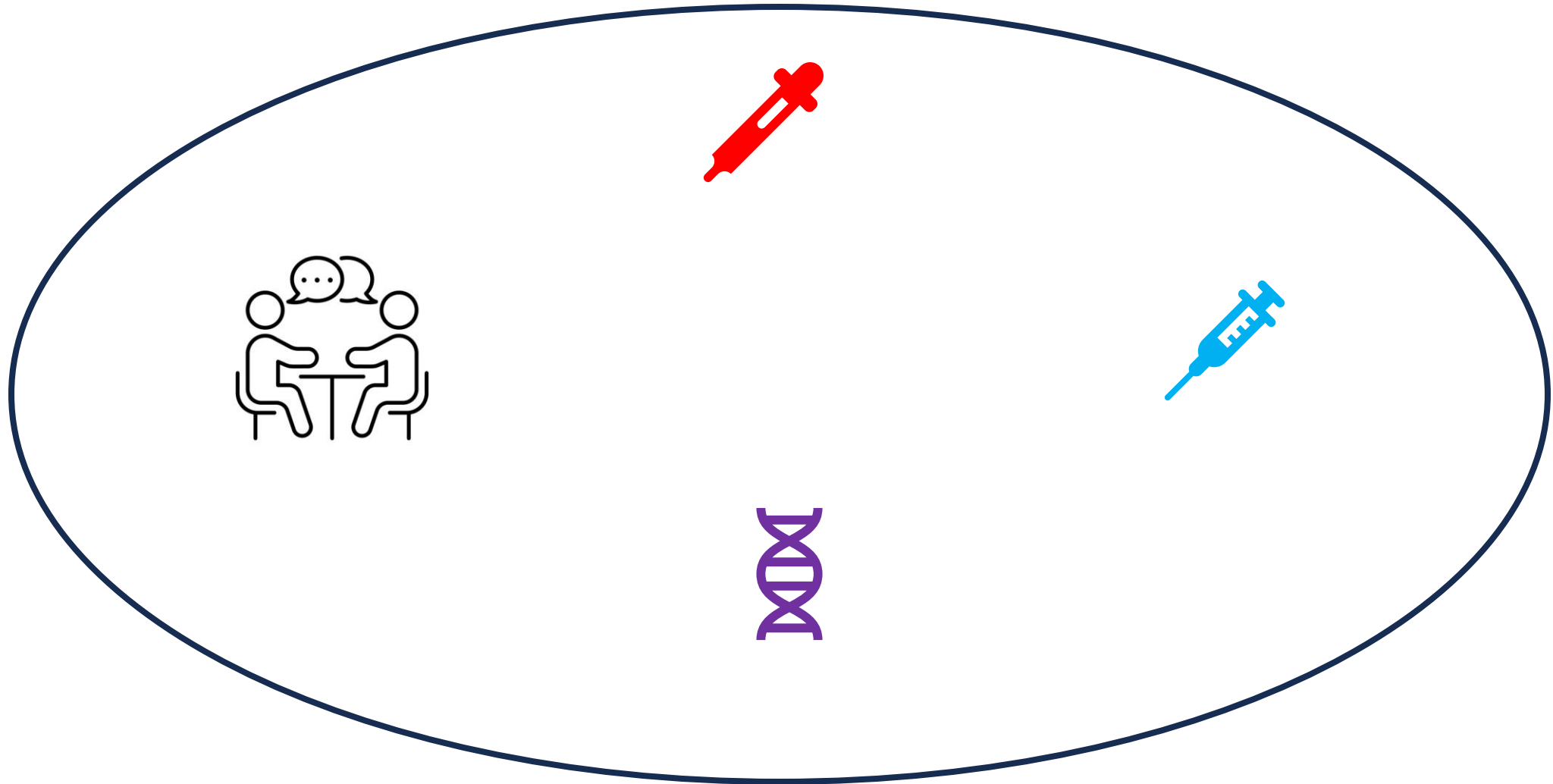
https://www.chicago.gov/city/en/depts/cdph/provdrs/infectious_disease/supp_info/mpox-home/mpox-dashboard.html

A concerning proportion of mpox patients initially identified were fully vaccinated.

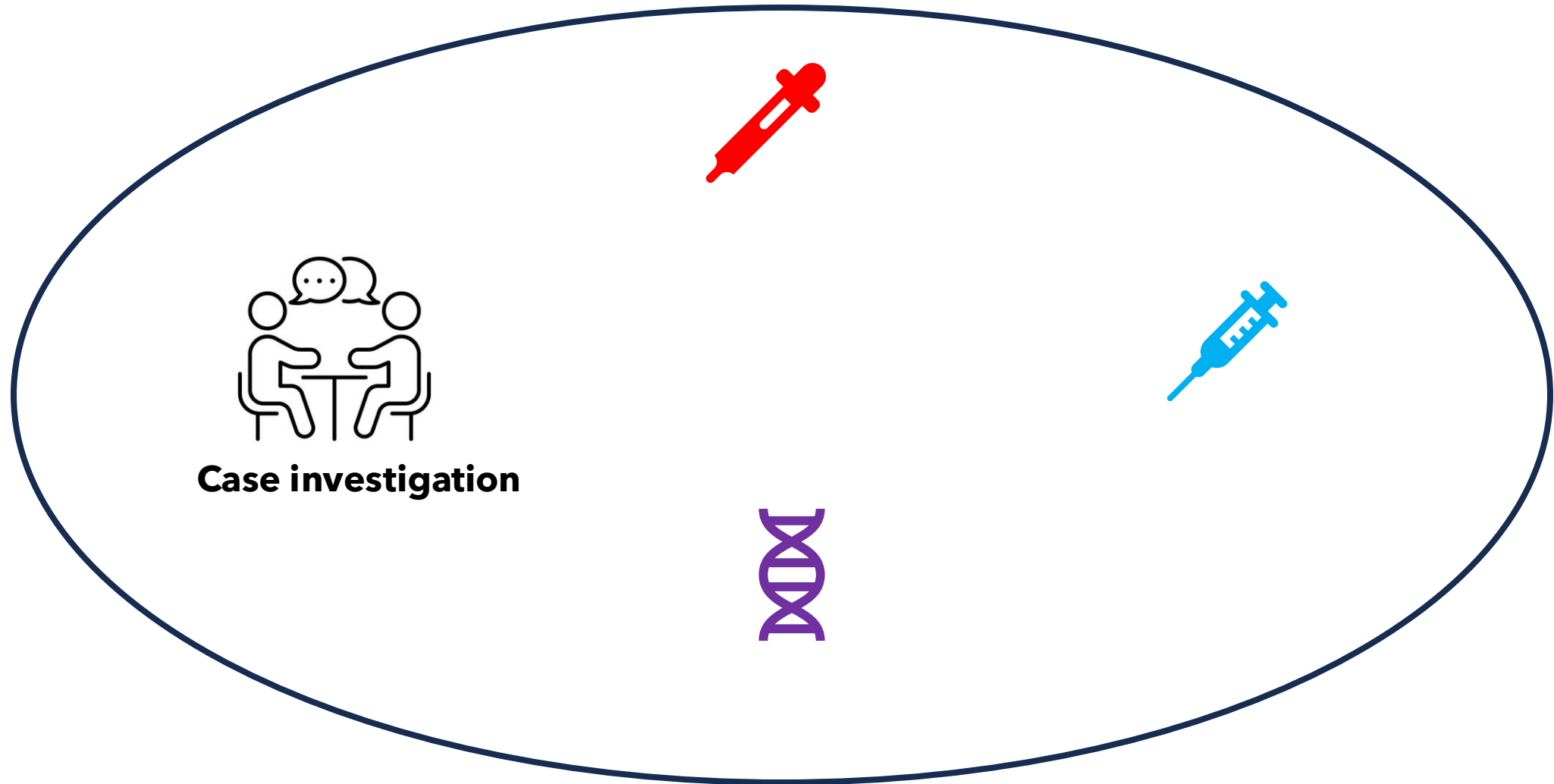


Cases reported between April 17 and May 5, 2023

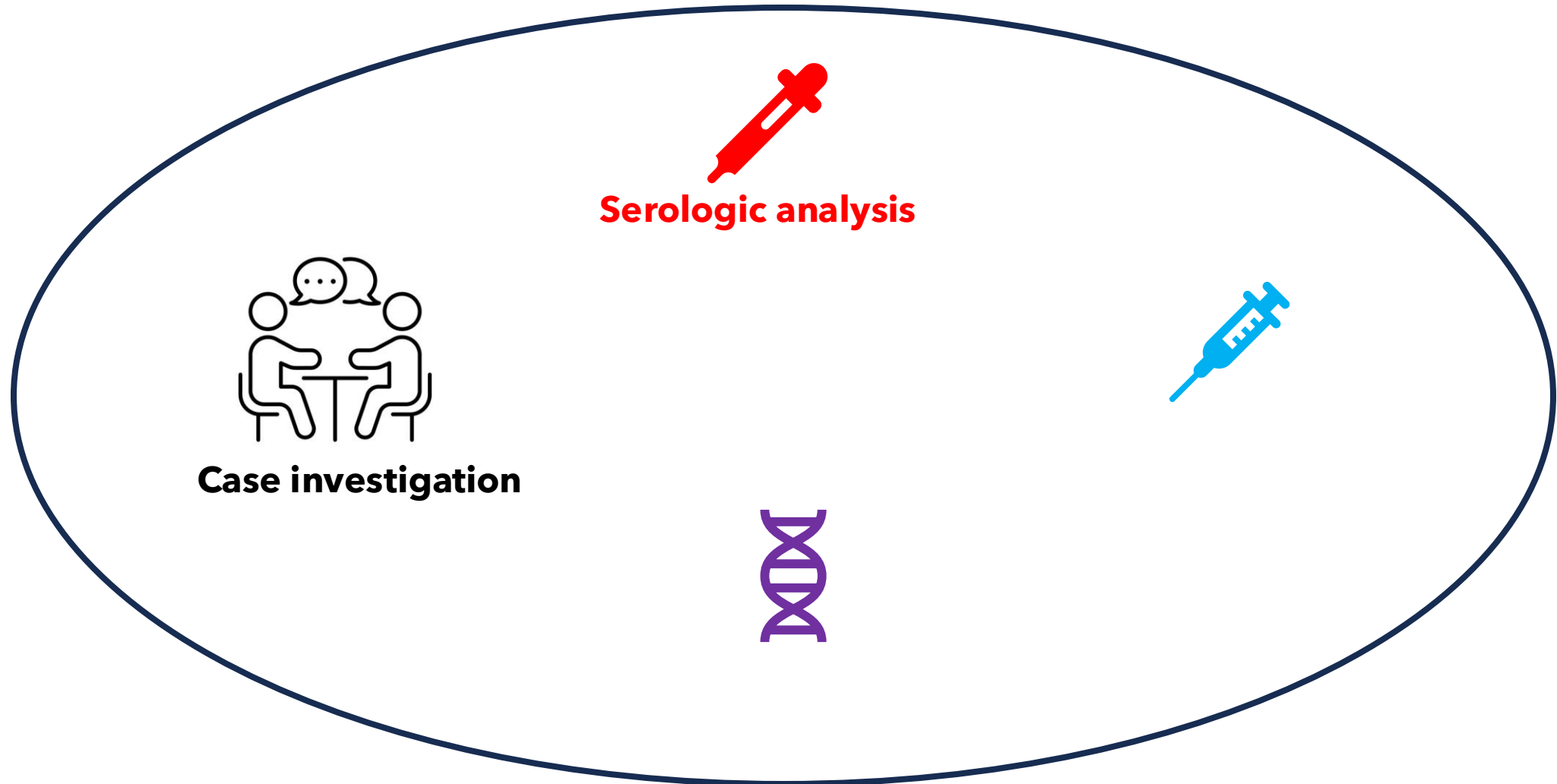
CDPH and CDC launched a multidisciplinary investigation to explain the cluster.



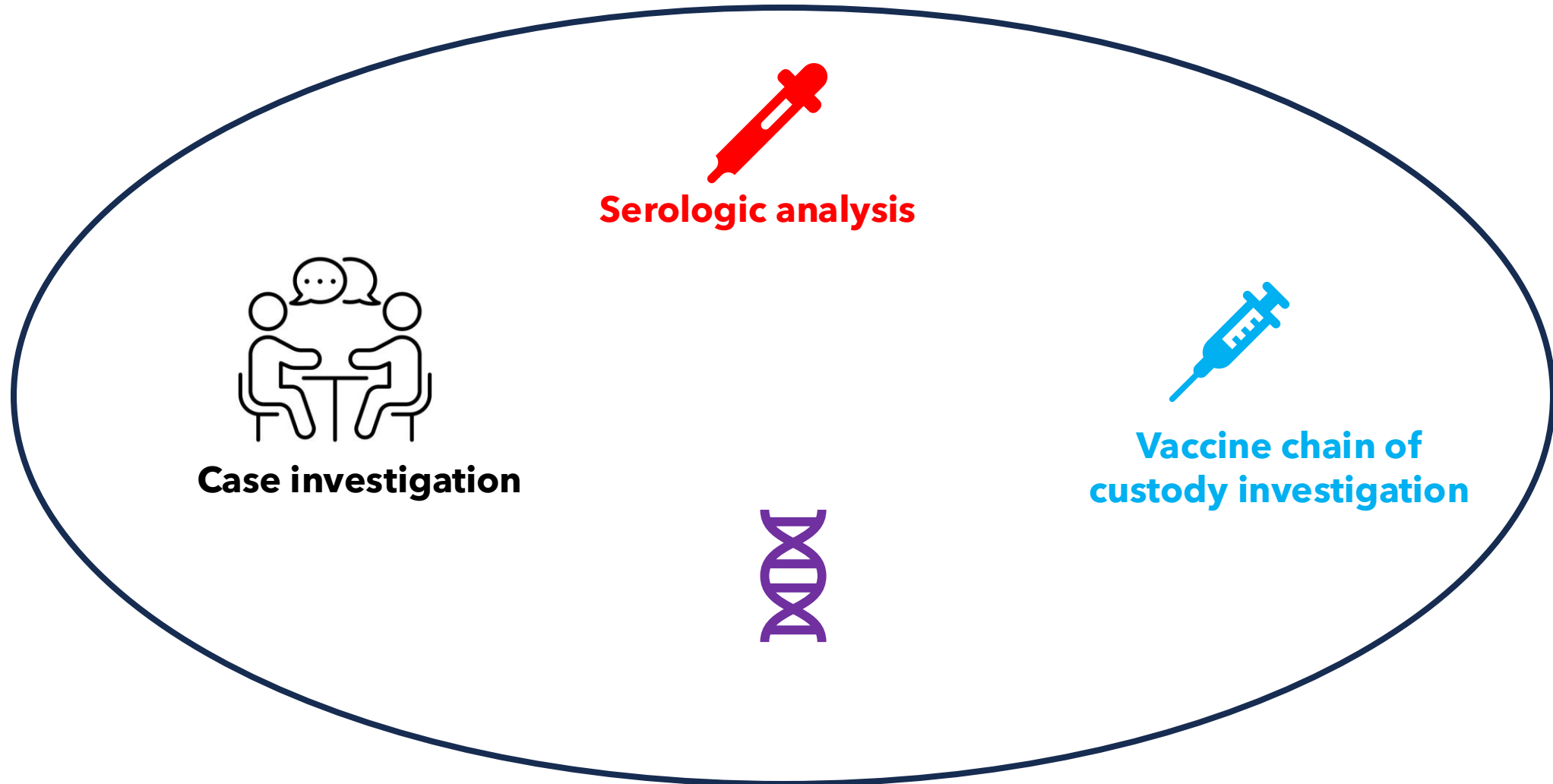
CDPH and CDC launched a multidisciplinary investigation to explain the cluster.



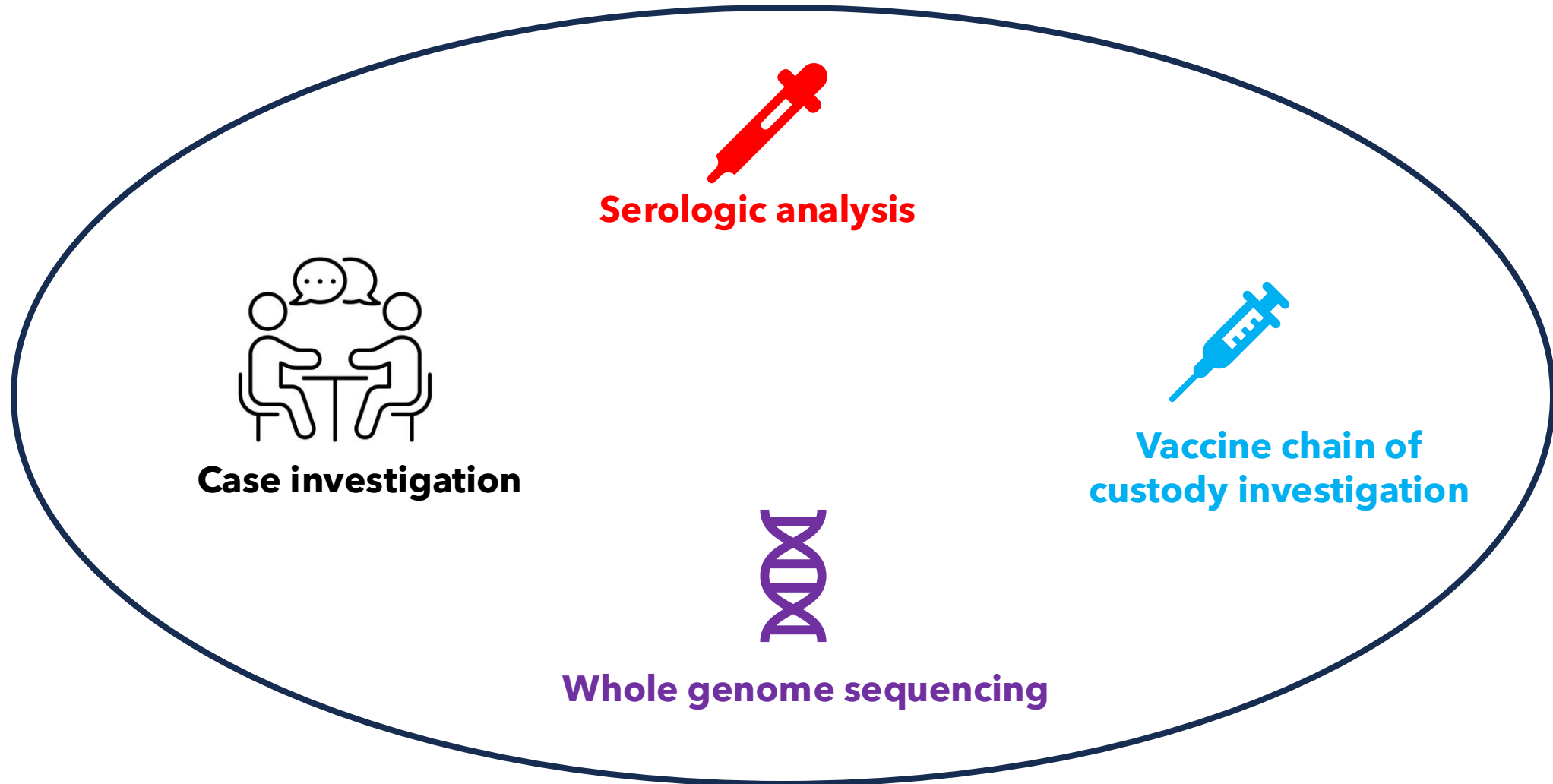
CDPH and CDC launched a multidisciplinary investigation to explain the cluster.



CDPH and CDC launched a multidisciplinary investigation to explain the cluster.



CDPH and CDC launched a multidisciplinary investigation to explain the cluster.



Case definition:

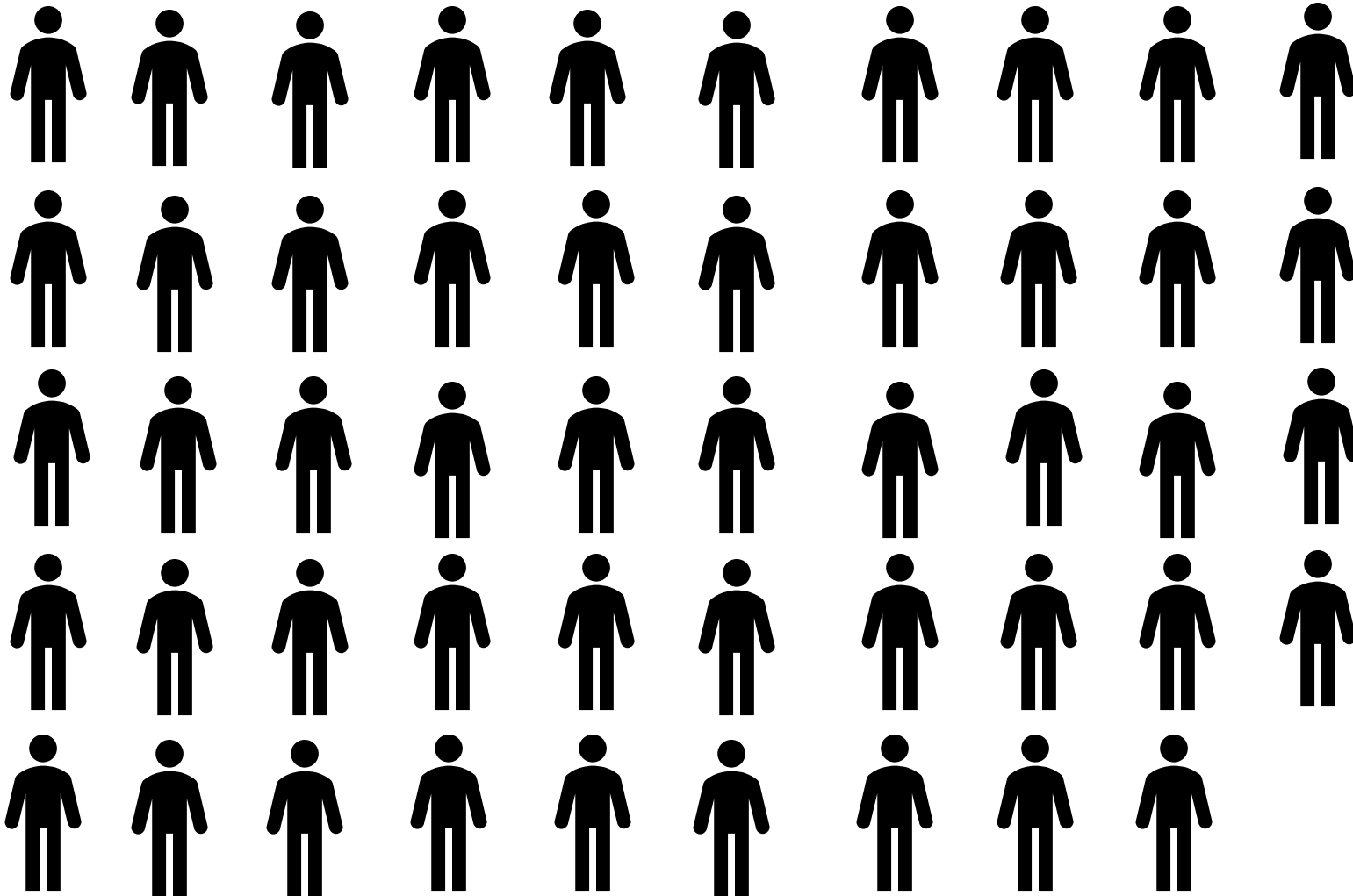
- Signs or symptoms of mpox in person with epidemiologic risk factors for mpox and laboratory confirmation identified March 18–June 27, 2023.
 - **Confirmed:** monkeypox virus positive
 - **Probable:** orthopoxvirus positive

Vaccination status:

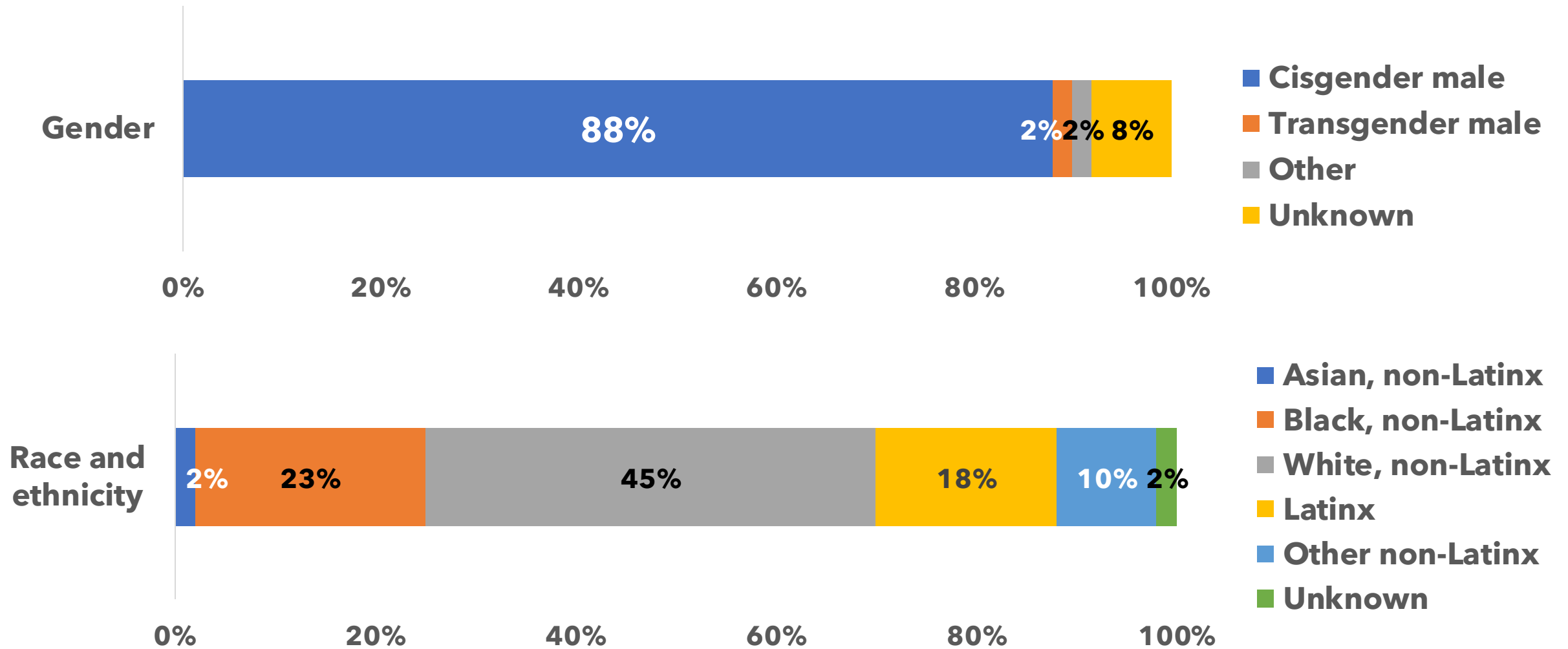
Fully vaccinated (FV): 2 doses of JYNNEOS or 1 dose of ACAM2000 before infection

Not fully vaccinated (NFV): ≤ 1 dose of JYNNEOS and no dose of ACAM2000

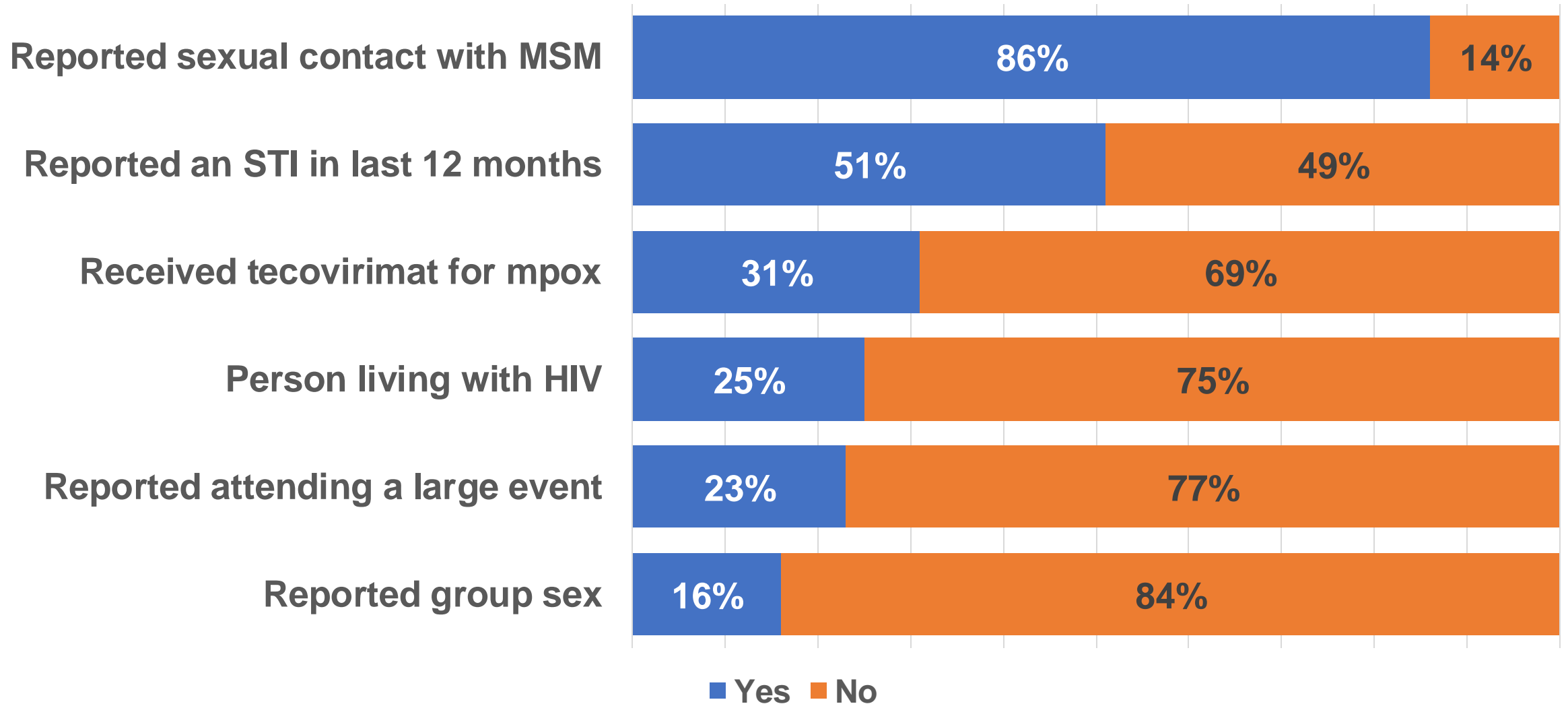
We identified 49 cluster patients,



Most cluster patients were cisgender males, and White patients were the largest group.



We examined clinical factors and behaviors associated with mpox transmission among patients.



Characteristics by vaccination status

28 (57%) patients were fully vaccinated (FV)

21 (43%) were not fully vaccinated (NFV)

Characteristics by vaccination status

28 (57%) patients were fully vaccinated (FV)

21 (43%) were not fully vaccinated (NFV)



Median sex partners reported by FV patients = 3

Median sex partners reported by NFV patients = 1

Characteristics by vaccination status

28 (57%) patients were fully vaccinated (FV)

21 (43%) were not fully vaccinated (NFV)



Median sex partners reported by FV patients = 3

Median sex partners reported by NFV patients = 1

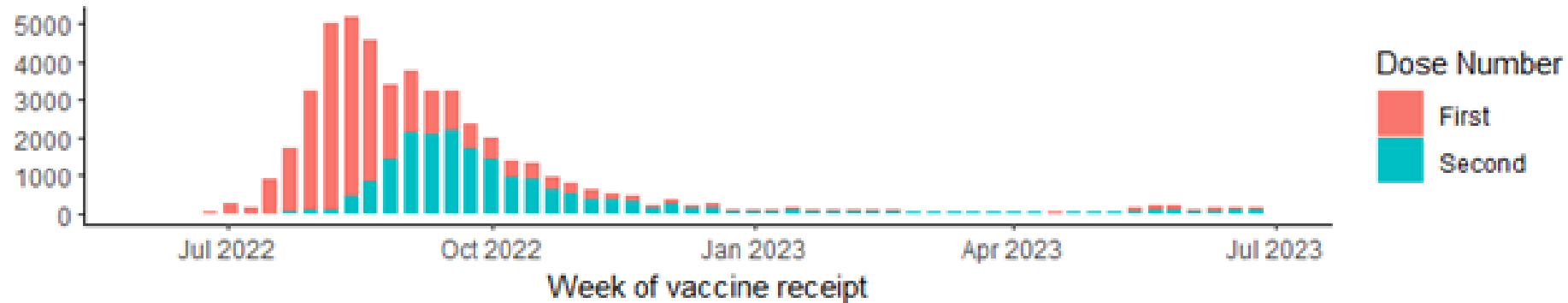


2 patients were hospitalized, both unvaccinated

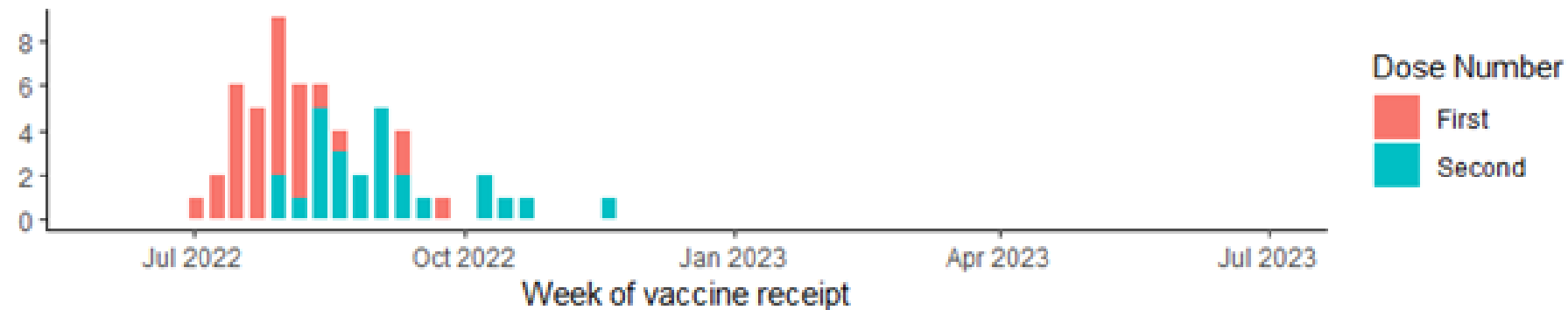
One with advanced HIV (CD4 count <200 cells/mm³)

Timing of vaccination for patients in this cluster was similar to all cases.

All vaccine doses given for mpox in Chicago



Vaccine doses given for mpox to cluster patients



Timing from vaccination to mpox disease



**Median:
9 months**

**Range:
5-12 months**



Serologic analysis revealed expected immune responses (n=13).



10 FV patients and **3 NFV patients (1 dose)** had serologic analysis

IgG **All positive** (median IgG = 1.28 (IQR = 1.25-1.38))

IgM **12 positive** (median IgM = 0.26 (IQR = 0.19-0.49))

Serologic analysis revealed expected immune responses.



10 FV patients and **3 NFV patients (1 dose)** had serologic analysis

IgG **All positive** (median IgG = 1.28 (IQR = 1.25-1.38))

IgM **12 positive** (median IgM = 0.26 (IQR = 0.19-0.49))

IgM values did not vary by number of doses received.

Vaccine Product Investigation



Vaccines given to cluster patients all came from 9 shipments from 6 lots at the Strategic National Stockpile (SNS) distributed throughout the United States.

Vaccine Product Investigation



Vaccines given to cluster patients all came from **9** shipments from **6** lots at the Strategic National Stockpile (SNS) distributed throughout the United States.



In chain of custody, temperature deviations were monitored and reported, but **none likely to affect product stability were detected.**

Vaccine Product Investigation



Vaccines given to cluster patients all came from **9** shipments from **6** lots at the Strategic National Stockpile (SNS) distributed throughout the United States.



In chain of custody, temperature deviations were monitored and reported, but **none likely to affect product stability were detected.**

We interviewed staff about **34 doses** given to cluster patients:

68% in a clinical setting,

32% at vaccine events, including

6% administered outdoors.



Vaccine Product Investigation



Vaccines given to cluster patients all came from **9** shipments from **6** lots at the Strategic National Stockpile (SNS) distributed throughout the United States.



In chain of custody, temperature deviations were monitored and reported, but **none likely to affect product stability were detected.**

We interviewed staff about **34 doses** given to cluster patients:

68% in a clinical setting,

32% at vaccine events, including

6% administered outdoors.



Facilities reported no temperature excursions.

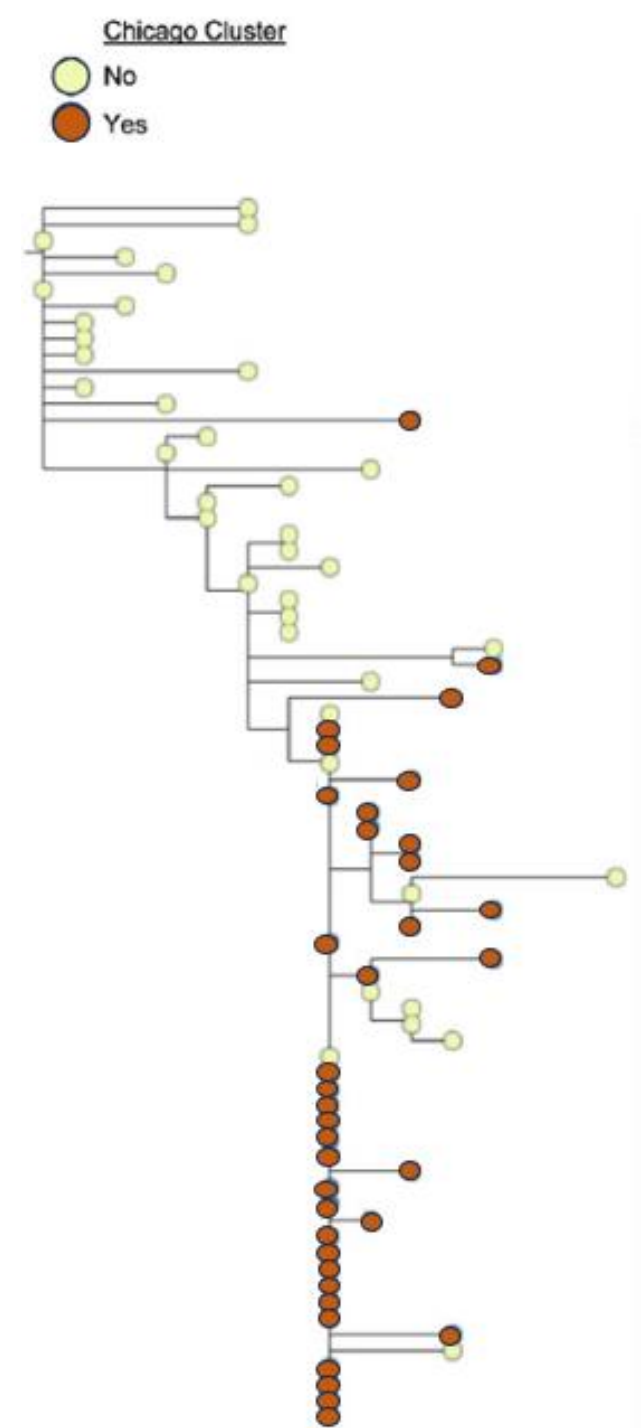
Only minor vaccine administration concerns were detected, affecting 2-3 doses.

Whole Genome Sequencing

37 specimens: 14 (38%) FV and **23 (62%) NFV patients**

Whole Genome Sequencing

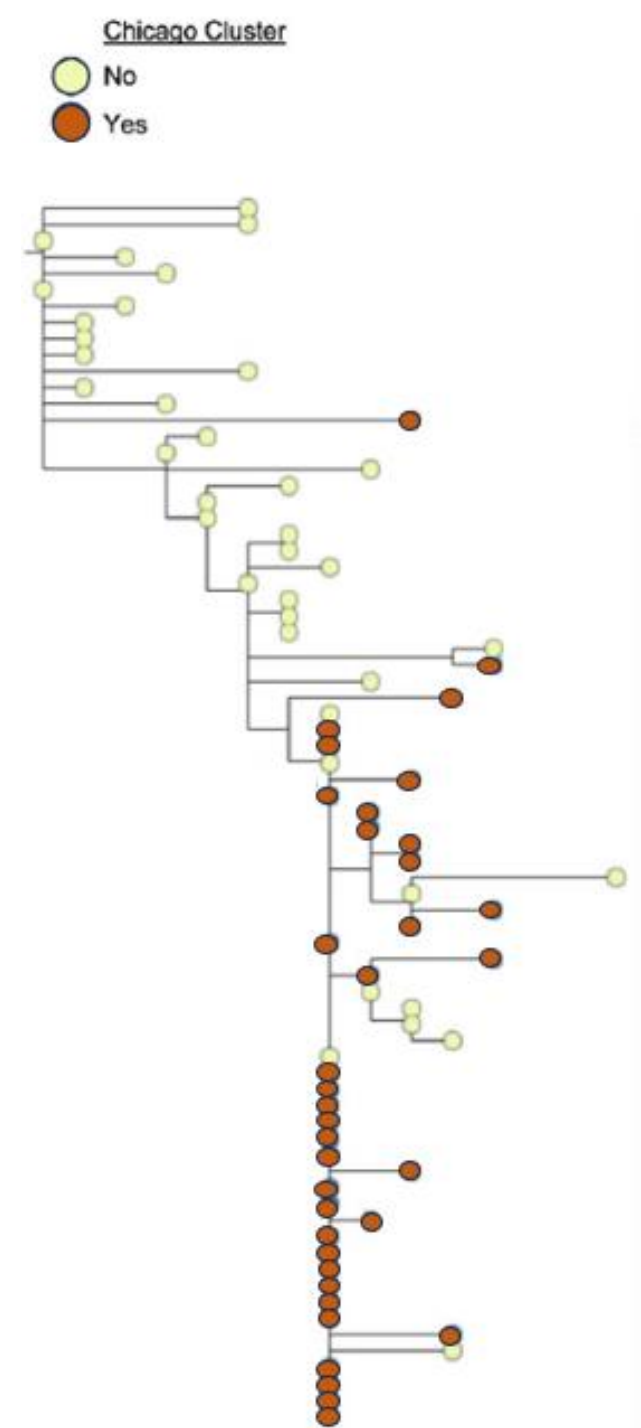
37 specimens: 14 (38%) FV and **23 (62%) NFV patients**



Whole Genome Sequencing

37 specimens: 14 (38%) FV and **23 (62%) NFV patients**

New sublineage: **mpox B.1 sublineage B.1.20.**



Whole Genome Sequencing

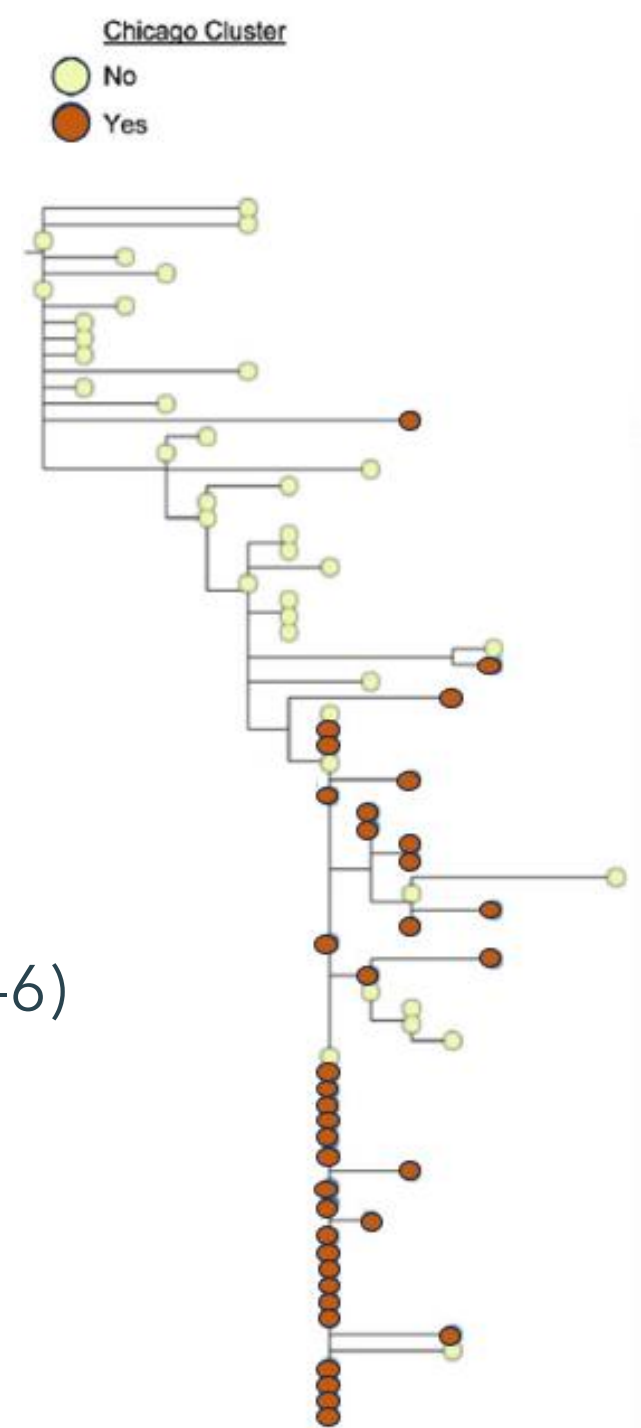
37 specimens: 14 (38%) FV and **23 (62%) NFV patients**

New sublineage: **mpox B.1 sublineage B.1.20.**

No genetic differences between **FV** and **NFV patients**

18 identical sequences and little genetic diversification

16 closely related (1-4 mutations); 2 less closely related (4-6)



Whole Genome Sequencing

37 specimens: 14 (38%) FV and **23 (62%) NFV patients**

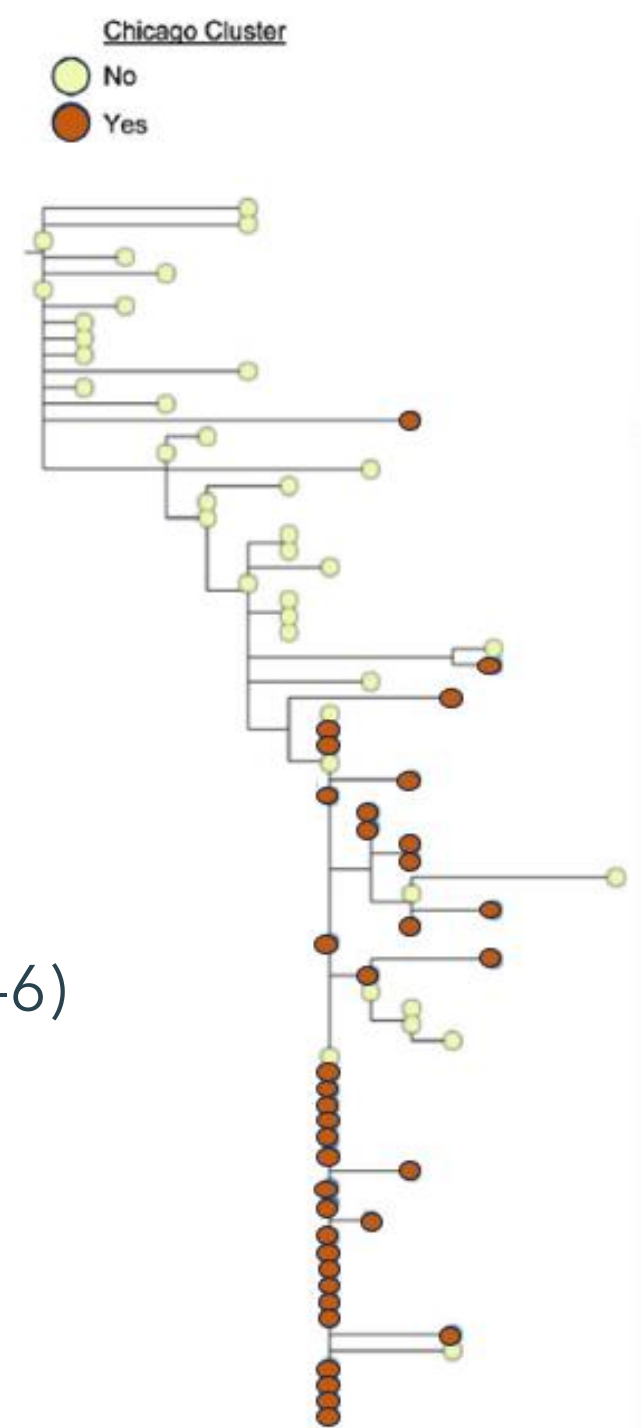
New sublineage: **mpox B.1 sublineage B.1.20.**

No genetic differences between **FV** and **NFV patients**

18 identical sequences and little genetic diversification

16 closely related (1-4 mutations); 2 less closely related (4-6)

Few amino acid changes in B.1.20 Chicago cluster



This investigation ruled out most hypotheses for the cluster emergence.



FV patients were less likely to have been hospitalized

More sexual partners 3 weeks before symptom onset.

This investigation ruled out most hypotheses for the cluster emergence.



FV patients were less likely to have been hospitalized
More sexual partners 3 weeks before symptom onset.



Patients sampled mounted **robust immune responses**.
Could not evaluate immune response after initial vaccination.

This investigation ruled out most hypotheses for the cluster emergence.



FV patients were less likely to have been hospitalized
More sexual partners 3 weeks before symptom onset.



Patients sampled mounted **robust immune responses**.
Could not evaluate immune response after initial vaccination.



Vaccine temperature excursions and vaccine administration breaches were **minor and unlikely to explain cluster**

This investigation ruled out most hypotheses for the cluster emergence.



FV patients were less likely to have been hospitalized
More sexual partners 3 weeks before symptom onset.



Patients sampled mounted **robust immune responses**.
Could not evaluate immune response after initial vaccination.



Vaccine temperature excursions and vaccine administration breaches were **minor and unlikely to explain cluster**



No pathogenic differences in genome sequences.

Conclusions and Recommendations



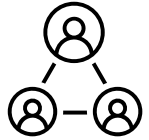
Monkeypox virus infections after vaccination can occur

Include mpox in differential diagnosis, even if previously vaccinated

Conclusions and Recommendations



Monkeypox virus infections after vaccination can occur
Include mpox in differential diagnosis, even if previously vaccinated

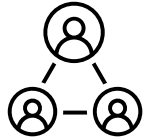


Syndemic approach critical for patients with mpox, HIV, other STIs, homelessness, and other mental health issues

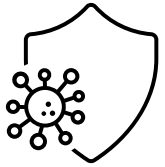
Conclusions and Recommendations



Monkeypox virus infections after vaccination can occur
Include mpox in differential diagnosis, even if previously vaccinated

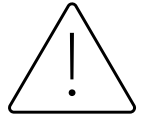


Syndemic approach critical for patients with mpox, HIV, other STIs, homelessness, and other mental health issues

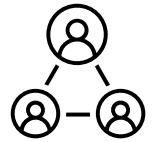


JYNNEOS vaccine effectiveness: 66%-88%
Clinical manifestations after vaccination may be less severe

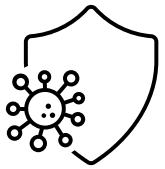
Conclusions and Recommendations



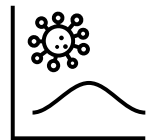
Monkeypox virus infections after vaccination can occur
Include mpox in differential diagnosis, even if previously vaccinated.



Syndemic approach critical for patients with mpox, HIV, other STIs, homelessness, and other mental health issues



JYNNEOS vaccine effectiveness: 66%-88%
Clinical manifestations after vaccination may be less severe

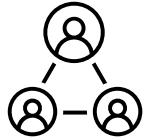


Future analyses of vaccine effectiveness over time are recommended

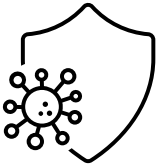
Conclusions and Recommendations



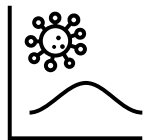
Monkeypox virus infections after vaccination can occur
Include mpox in differential diagnosis, even if previously vaccinated.



Syndemic approach critical for patients with mpox, HIV, other STIs, homelessness, and other mental health issues



JYNNEOS vaccine effectiveness: 66%-88%
Clinical manifestations after vaccination may be less severe.



Future analyses of vaccine effectiveness over time are recommended



Strong recommendation for JYNNEOS vaccine

Acknowledgments

Patients and Health Provider Participants

Chicago Department of Public Health

Taylor Holly	Massimo Pacilli
Hillary Spencer	David Kern
Ashley M. Becht	Janna Kerins
Gordon Crisler	Irina Tabidze
Michael Wasz	Christopher Shields
Patrick Stonehouse	Rebecca Pavlatos
Christy Zelinski	Donna Peace
Alyse Kittner	Van Quach
Dorothy Foulkes	Usha Samala
Kendall W. Anderson	Shamika Smith
Tiffany Evans	Saadeh Ewaidah
Brian Borah	Spencer Gorelick
Stephanie R. Black	Stephanie Gretsich

Centers for Disease Control and Prevention

Yasmin P. Ogale	Laura A. S. Quilter
Lavinia Nicolae	Andrea M. McCollum
Amber Staton	Agam K. Rao
Carla Hardnett	Whitni Davidson
Michael B. Townsend	Nhien Wyn
William C. Carson	Kimberly Wilkins
Panayampalli S. Satheshkumar	Faisal Minhaj
Christina L. Hutson	Dawn Broussard
Crystal M. Gigante	Ian Kracalik
Laura A. S. Quilter	

Regional Innovative Public Health Laboratory

Hannah J. Barbian

US Strategic National Stockpile

Susan Gorman

Thank you!

For more information, contact CDC
1-800-CDC-INFO (232-4636)
TTY: 1-888-232-6348 [cdc.gov](https://www.cdc.gov)

The findings and conclusions in this report are those of the authors and do not necessarily represent the official position of the U.S. Centers for Disease Control and Prevention.



Supplementary slides

Mpox is caused by a viral infection spread through contact with infected persons or animals.



Vaccine Product Investigation

Some vaccine administration concerns were identified:

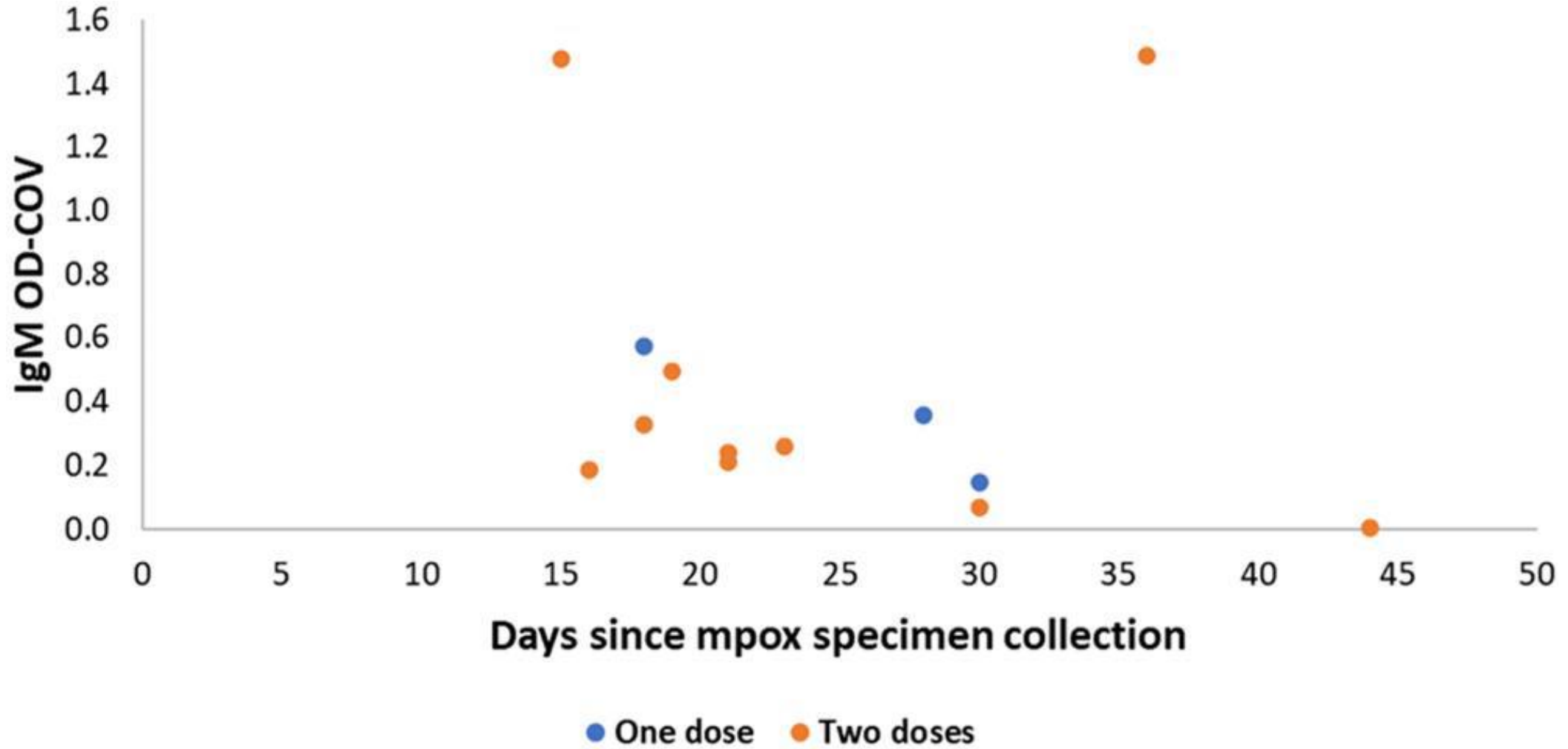
2 doses Administration in a room where ambient temperature $>25^{\circ}\text{C}$

1 dose Use of punctured vials

2 doses Use of pre-filled syringes before vaccine administration >2 minutes

These deviations did not solely affect cluster patients and could not explain vaccine compromise for the cluster.

Time from specimen collection to serology



Mucosal Areas Affected

Ocular



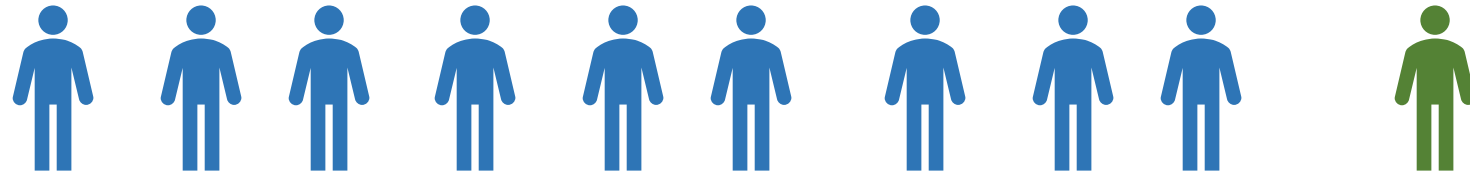
Oral



Genital



Rectal



Hospitalized patients



Advanced HIV (CD4 count <200 cells/mm³)
Fungating Lesions
Bacterial superinfection



Virally suppressed (viral load <200 cells/mm³)
Oropharyngeal Lesions
Pain swallowing

Tecovirimat Administration

15 Received Tecovirimat:

4 STOMP

11 EA IND Protocol / Open Label STOMP

Documented Indications:

1 - Ocular

2 - Oral

4 - Rectal lesions with painful/bloody defecation

1 - Fungating lesions

4 - PWH:

3 **FV** well controlled HIV (>500 cells/mm³, viral load <200 copies/ml)

1 **unvaccinated** with advanced HIV

Primary and secondary antibody responses

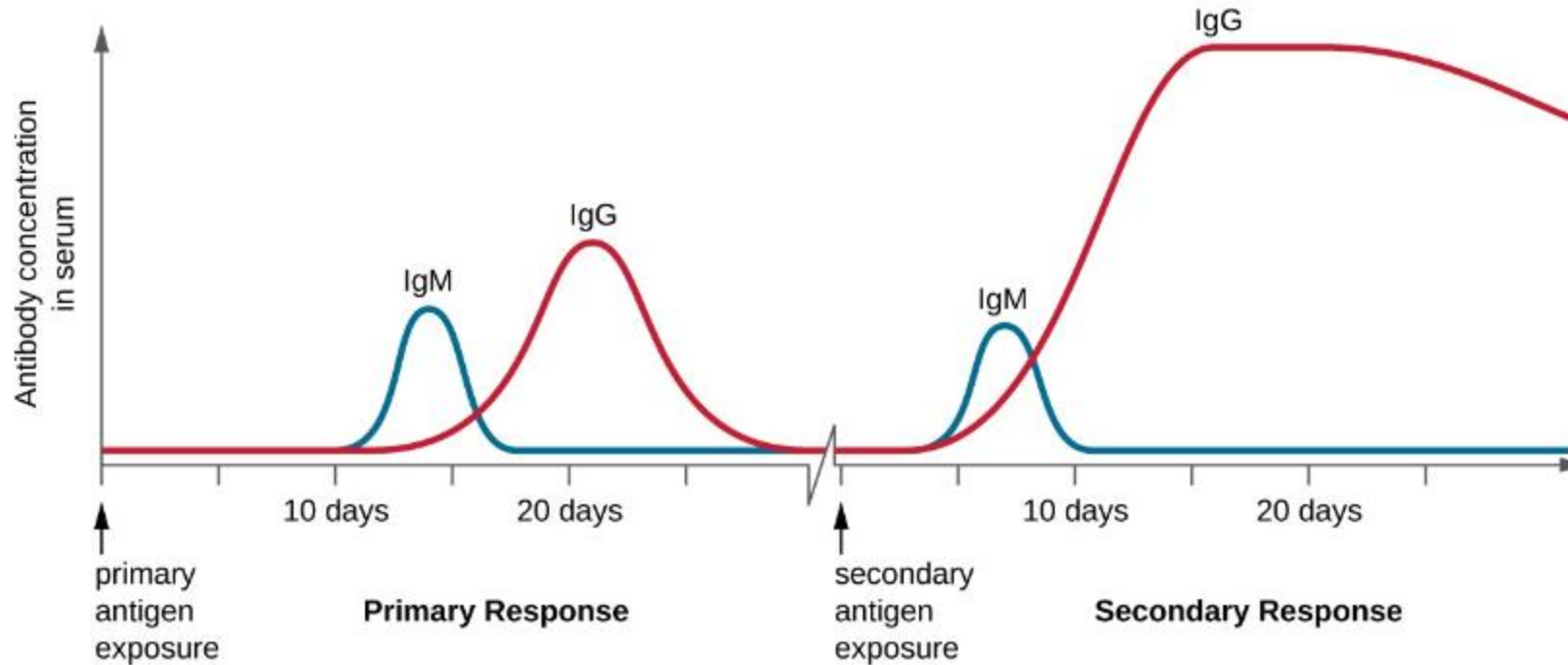


Figure 4. Compared to the primary response, the secondary antibody response occurs more quickly and produces antibody levels that are higher and more sustained. The secondary response mostly involves IgG.