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Ponce De Leon Center

# Clinical Presentation and Complications of Mpox in Atlanta During the 2022 Outbreak

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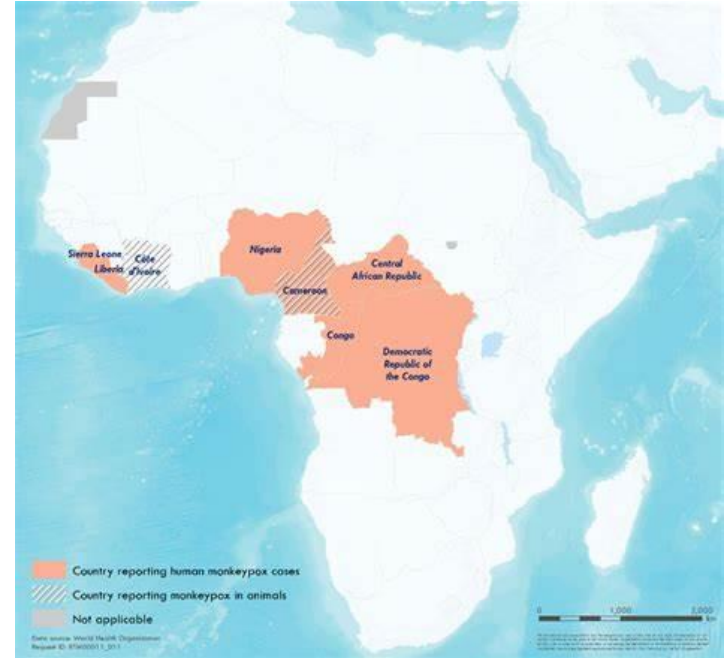


Serious  
Communicable  
Diseases Program



# Endemic Mpox

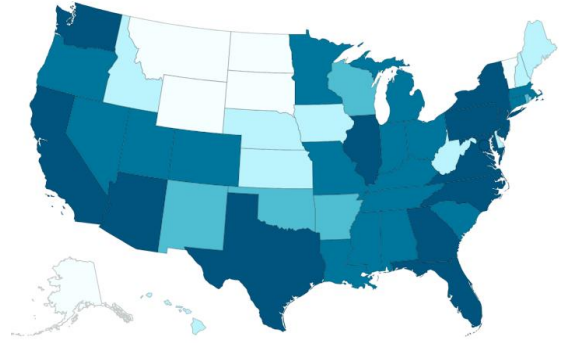
- Clade I (Congo Basin/ Central African)
  - Higher mortality (~ 10%)
- Clade II (West African)
  - Lower mortality (~ 3%)<sup>1</sup>



# 2022 Global Mpox Outbreak

- In spring 2022, clade II monkeypox virus (since renamed “mpox virus”) began spreading in numerous non-endemic regions.
- The global outbreak of mpox starting in 2022 is the largest in history to occur outside the continent of Africa<sup>2,3</sup>.
- There have been over 90,000 confirmed global mpox cases involving over 100 non-endemic countries<sup>4</sup>.
- Transmission predominantly occurs via sexual and intimate contact.
- Cases disproportionately affected certain groups<sup>5</sup>:
  - Men who have sex with men (MSM): ~ 95% of cases
  - People living with HIV (PLWH) ~ 40% of cases

# 2022 Mpox Outbreak in the U.S.



- 32,063 cases (as of 1/10/24)
- 58 deaths
- U.S. South was the region with the highest case counts<sup>6</sup>
- Notable population disparities<sup>7</sup>:
  - People living with HIV (PLWH):
    - 0.4% of population; 38% of mpox cases; 94% of deaths
  - Black (Non-Hispanic) individuals:
    - 12.1% of population; 33% of mpox cases; 86% of deaths

# Atlanta Patient Cohort

- Cohort of all individuals diagnosed with mpox disease at EUH, EUHM, Grady/Ponce, and the Atlanta VA from 6/1/2022- 10/7/2022.
- Medical charts were reviewed in detail to collect data on demographics, exposures, clinical course, management, and outcomes.
- Data collection was a collaborative effort of more than 20 people from the Emory ID Department, Emory Center For AIDS Research (CFAR), and Grady Ponce de Leon Center.

<b>Demographics</b>	
<b>Median age (IQR, years)</b>	35 (30-41)
<b>Gender identity</b>	
• Cisgender man	384 (97.2%)
• Transgender woman	10 (2.5%)
• Cisgender woman	1 (0.3%)
<b>Race</b>	
• Black	335 (84.8%)
• White	18 (4.6%)
• Other/ unknown	42 (10.6%)
<b>Ethnicity</b>	
• Hispanic/Latinx	21 (5.3%)
<b>Sexual practices</b>	
• Sex with cisgender men	342 (86.6%)
<b>Unhoused</b>	30 (7.6%)
<b>Uninsured</b>	229 (58.0%)
<b>HIV-positive</b>	324 (82.0%)

# Clinical Presentation

Most Common Presenting Symptoms:	
• Rash	366 (92.7%)
• Fever/ chills	175 (44.3%)
• Lymphadenopathy	144 (36.5%)
• Fatigue/ malaise	120 (30.4%)
• Rectal pain	105 (26.6%)
• Sore throat	91 (23.0%)
• Myalgias	63 (15.9%)
• Headache	32 (8.1%)

## Differences in 2022 outbreak clinical presentation compared to endemic mpox:

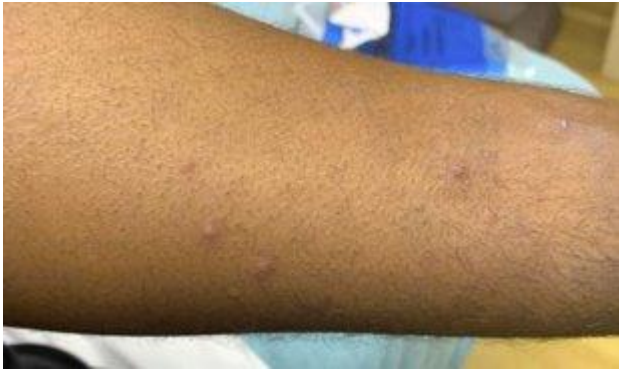
- Less prodromal symptoms
  - Less fevers, chills, lymphadenopathy, myalgias
- Rash generally more mild
  - Fewer lesions
  - More localized
    - Rash may be localized to only skin around the anorectum, genitals, or mouth/ lips (sites of sexual contact)
- More common mucosal involvement of rectum, urethra, and/or oropharynx
- Frequent concomitant chlamydia, syphilis, gonorrhea, and/or HIV diagnoses.



<b>Mucosal Involvement</b>	
• Anorectal	89 (22.5%)
• Oral/ pharyngeal	82 (20.8%)
• Urethral	26 (6.6%)
• Ocular	3 (0.8%)
• Nasal	2 (0.5%)
<b>Number of mucosal sites involved:</b>	
• None	228 (57.7%)
• One site	135 (34.2%)
• Two or more sites	32 (8.1%)

# Clinical Presentation: Timing

<b>Median days from symptom onset to mpox testing</b>	<b>5 (IQR: 3-7)</b>
<b>Prior healthcare visit(s) for mpox symptoms without mpox testing</b>	<b>91 (23.0%)</b>



(All clinical images used with patient consent)

# Mpox Rash: The 6 P's

Morphology: Papular → Pustular

Location: Peri-anal (often with Proctitis)

Penis

Pharynx

Periphery (limbs > torso)



(All clinical images used with patient consent)

# Rash Progression

(Papule → pustule → crust  
→ scab → healing)



# Diagnosis

- Mpox PCR swab
- Technique: Unroof and swab 1-2 skin lesions
  - If no obvious skin lesions, then swab the affected site (e.g. rectal swab, pharyngeal swab)
- Concomitant STI screening
  - GC/CT NAAT throat/ rectum/ urine
  - RPR
  - HIV screen
  - +/- HBV and HCV serologies



Complications	
• Any Complication	68 (17.2%)
• Bacterial superinfection (cellulitis/ abscess)	38 (9.6%)
• Anorectal abscess	8 (2.0%)
• Pharyngeal abscess	4 (1.0%)
• Bacteremia	4 (1.0%)
• Colitis/ GI bleeding requiring transfusion	13 (3.3%)
• Delayed rash healing > 4 weeks	10 (2.5%)
• Conjunctivitis	4 (1.0%)
• Phimosi	3 (0.8%)
• Myocarditis	2 (0.5%)

# Outcomes

Outcomes	
Hospital Admission	66 (16.7%)
ICU Admission	4 (1.0%)
Death	1 (0.3%)
Recovery	394 (99.7%)

The vast majority of patients recovered without sequelae.

# Risk Factors Associated with Severe Disease

- Older age
- Immunocompromised states
  - Non-HIV (e.g. hematologic malignancy, immunosuppressive meds)
  - HIV infection
    - Lower CD4+ count (< 200 cells/  $\mu$ L)
    - Non-suppressed HIV viral load (> 200 copies/ mL)
- Unhoused status
- Mucosal site involvement at presentation



# Management

- Supportive Care
  - Pain management:
    - NSAIDs
    - Topical dibucaine for anorectal irritation
    - Short-course opiates for severe pain (e.g. necrotic skin lesions, mucosal involvement)
  - Antihistamines (e.g. hydroxyzine) for itching
  - Acetaminophen for fever
  - Stool softeners if proctitis
  - Antibiotics if signs of skin lesion bacterial superinfection
  - IV fluids if difficulty with PO fluid intake



# Management

- Antivirals
  - **Tecovirimat**
    - Efficacy shown in animal models (if started within 5-7 days of mpox inoculation)
    - Favorable safety profile in humans
    - Efficacy in humans has not been conclusively demonstrated
      - A small observational matched-cohort of people with HIV with mpox suggests tecovirimat may be beneficial if started early after symptom onset<sup>8</sup>.
    - Access: ACTG STOMP trial ([www.stomptpox.org](http://www.stomptpox.org)) - RCT of tecovirimat vs placebo
    - Conclusion: Offer STOMP enrollment to any patient with suspected or confirmed mpox.
  - **Cidofovir/ Brincidofovir and Vaccinia Immune Globulin (VIGIV)**
    - IV-only; case reports of use as salvage therapy in severe cases.

# Management: Refractory Disease

- Some people with very advanced HIV (CD4+ < 100 cells/ $\mu$ L) develop non-healing necrotic skin lesions that may take months to recover.
  - Requires aggressive supportive care: wound care, pain management, treatment of bacterial superinfections, etc.
  - In our Atlanta cohort, 3 patients (0.8%) had hospital LOS  $\geq$  30 days.
- Restarting antiretroviral therapy (ART)
  - In general, prompt reinitiation of ART is recommended.
  - A large global case series including individuals with mpox and advanced HIV raised c/f an immune reconstitution inflammatory syndrome (IRIS) phenomenon which can paradoxically exacerbate disease severity<sup>9</sup>.
    - It may be appropriate to delay ART initiation in mpox patients who are critically ill.
    - In our Atlanta cohort, of the 9 patients with CD4+ < 100 cells/ $\mu$ L who were non-adherent to ART prior to presentation and were promptly restarted on ART, no paradoxical worsening of mpox disease or IRIS-like phenomenon was observed.

# Prevention

What: JYNNEOS vaccine

Who: Any cisgender man or transgender female who is sexually active with > 1 male partners within the past 6 months. Other high-risk individuals can be vaccinated on a case-by-case basis (see [www.cdc.gov/poxvirus/mpox/vaccines](https://www.cdc.gov/poxvirus/mpox/vaccines)).

When: 2 doses, given 4 weeks apart.

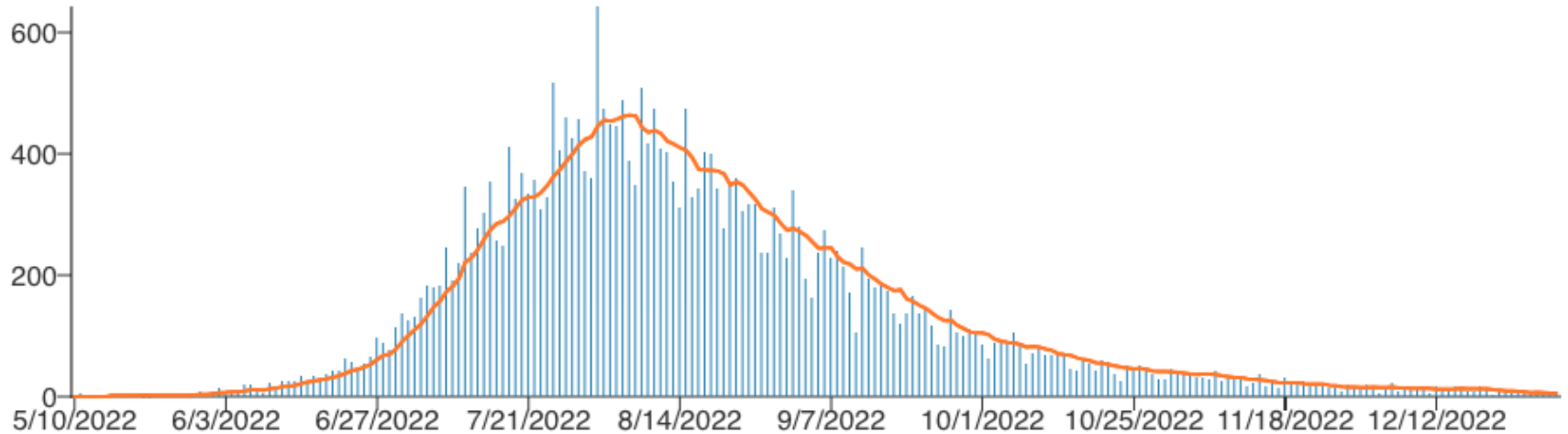
Where: Intradermal administration

Why: > 65% effective at preventing mpox<sup>10</sup>



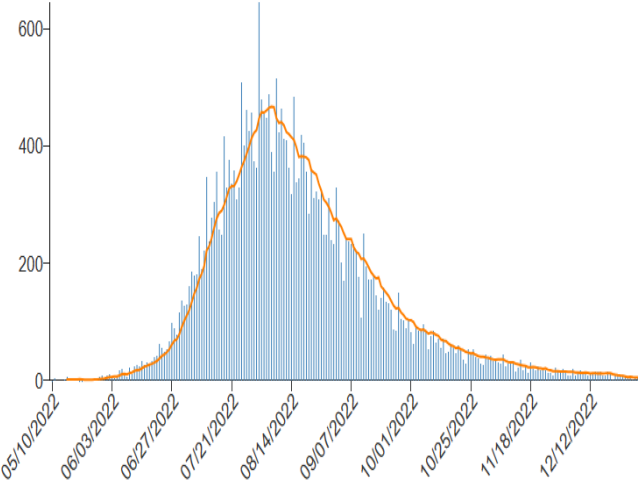
# Ending the Outbreak

- Likely predominantly due to high vaccine uptake in high-risk populations<sup>11</sup>.

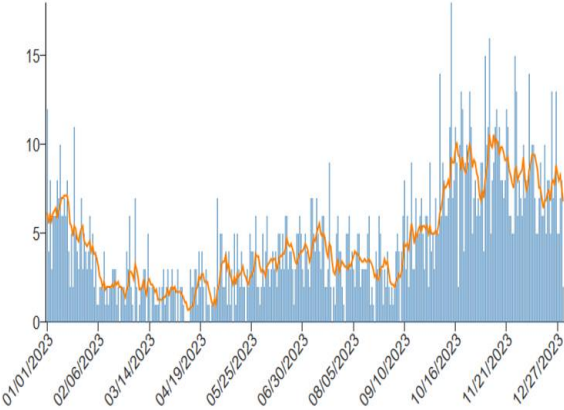


# Mpox: USA 2024

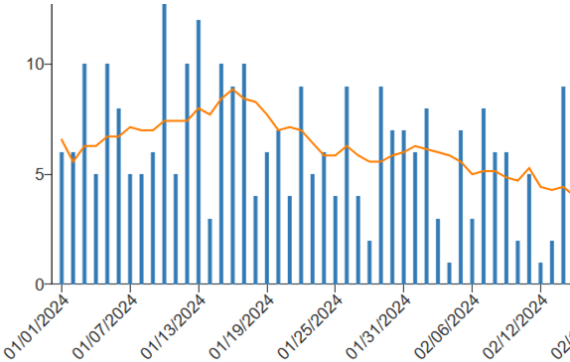
- While case numbers are quite low, they have not gone to zero<sup>6</sup>.



2022



2024



2023

# Endemic Mpox: 2024

- The largest clade I mpox outbreak in the history of the DRC has been ongoing since 2023, with 12,569 suspect mpox cases and 581 deaths<sup>12</sup>.
  - Travel-associated cases in the U.S. are possible and clinicians must remain vigilant.



Image:

[https://th.bing.com/th/id/R.5e8e866f9892ec505fe7fca70f00c702?rik=V%2ffvO%2bhwyQSog&riu=http%3a%2f%2fd2z7bwflv7old.cloudfront.net%2fcdn\\_image%2fexW\\_1200%2fimages%2fmaps%2fen%2fcg%2fcg-area.gif&ehk=pwpHurGjghnQO3uwkM%2beyvHty4Hc4NFEWtwR3wBDEvQ%3d&risl=&pid=ImgRaw&r=0](https://th.bing.com/th/id/R.5e8e866f9892ec505fe7fca70f00c702?rik=V%2ffvO%2bhwyQSog&riu=http%3a%2f%2fd2z7bwflv7old.cloudfront.net%2fcdn_image%2fexW_1200%2fimages%2fmaps%2fen%2fcg%2fcg-area.gif&ehk=pwpHurGjghnQO3uwkM%2beyvHty4Hc4NFEWtwR3wBDEvQ%3d&risl=&pid=ImgRaw&r=0)

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# Key Takeaways

- Mpox spreads through sexual and intimate contact and the 2022 outbreak disproportionately affected men who have sex with men (MSM).
  - Mpox rash “6 P’s”: Papular or Pustular rash located most commonly on the Perianus, Penis, Pharynx, and/or Periphery
  - HIV and other immunocompromised states are associated with more severe mpox, with risk correlating to the degree of immunosuppression.
  - Mucosal involvement (rectum, oropharynx, urethra, and other sites) can be extremely painful and warrants aggressive supportive care.
  - While case numbers are quite low compared to summer 2022, low-level community transmission continues to persist.
  - Refer any suspected mpox patient to the ACTG-STOMP trial.
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