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### BACKGROUND

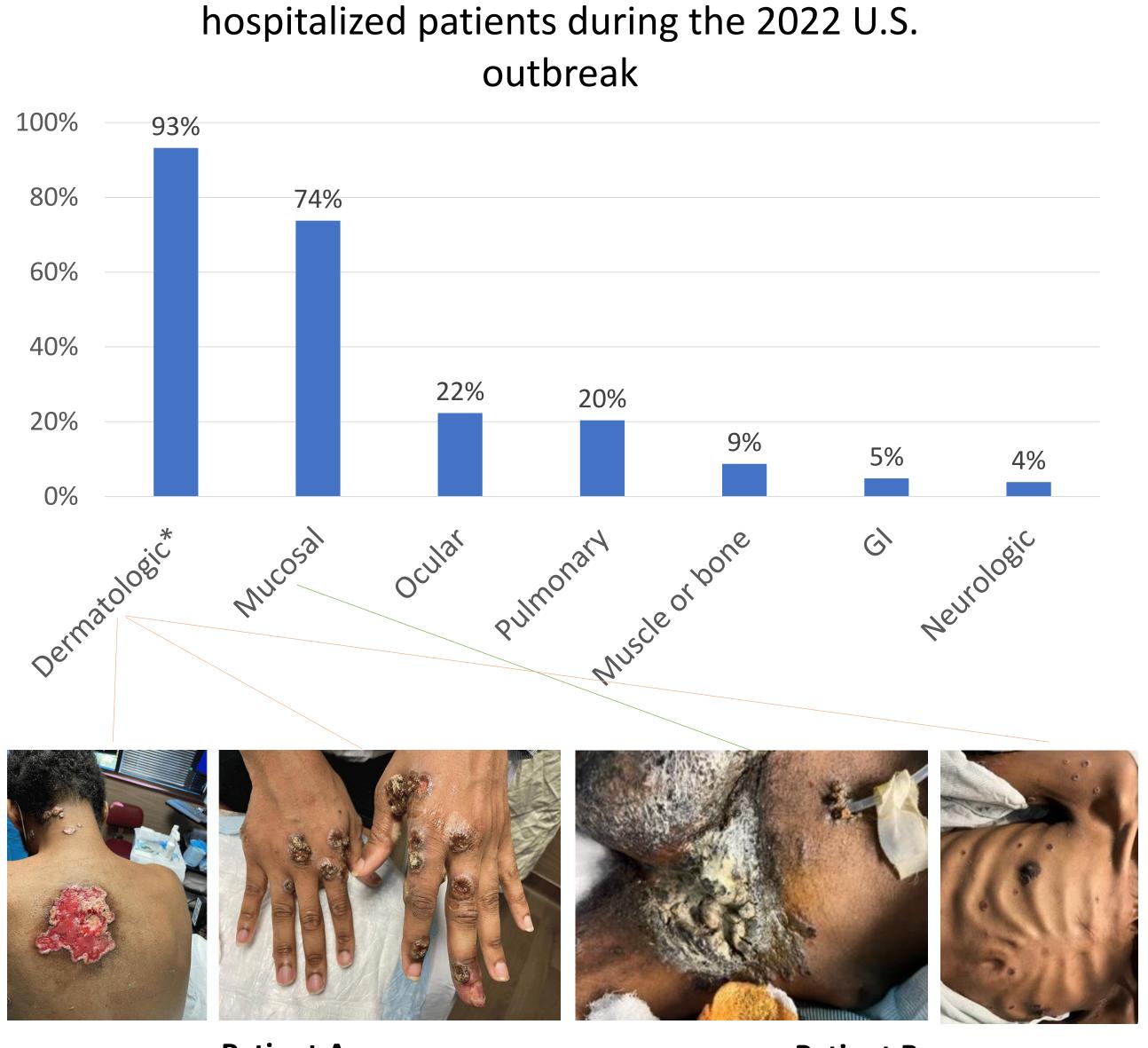
• What are the demographics, clinical trajectories, and HIVrelated care considerations for patients hospitalized with mpox during the 2022 mpox outbreak?

In prior mpox outbreaks, severe manifestations of disease and poor outcomes have been reported among people with HIV, particularly those with AIDS. During the 2022 international mpox outbreak, CDC staff provided clinical consultations to providers caring for patients with mpox regarding clinical management and therapeutics.

### METHODS

- Descriptive analysis of hospitalized U.S. patients aged  $\geq$ 18 years with confirmed mpox for whom CDC was consulted (between 8-10-22 to 11-22-22)
- Data obtained from provider reports during consultation with health departments or clinicians

Figure 1: Categories of mpox manifestations in



Patient A

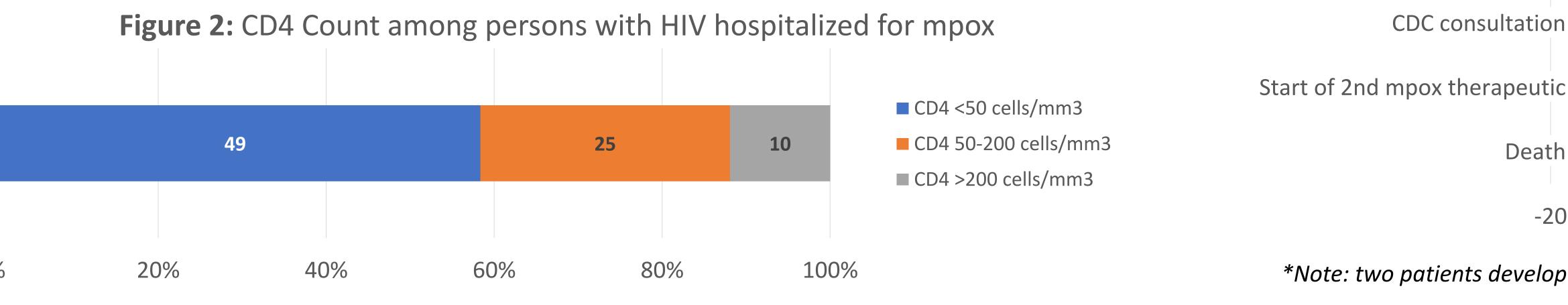
\*Note: the remaining 7% of patient consult data lacked information on presence or absence of dermatologic manifestations

# **Characteristics of Patients Hospitalized with Mpox during the** 2022 U.S. Outbreak

Inadequately treated HIV i most commonly associated this study population. More hospitalized with mpox diedpatients with symptoms su tested for HIV. Engaging a ensure ART initiation and adh health priority.

### RESULTS

- Of the patients for whom CDC was consulted over 60% (62) were Black, and 21% (22) were experier
- A range of systemic manifestations was reported for patients hospitalized with mpox (Figure 1).
- Nearly all patients (95%; 98) had immunocompromising conditions:
- HIV: 87% (90) (84 with available CD4 counts; 77 with ART data)
- CD4 < 200 cells/mm<sup>3</sup>: 88% (74/84) (Figure 2)
- ART at time of mpox diagnosis: 18% (14/77)
- Other: history of solid organ or stem cell transplant, cancer on chemotherapy, autoimmune conditions requiring steroids, and pregnancy (all 3 women)
- All patients with HIV who died had CD4 < 200 cells/mm<sup>3</sup>
- No patients with HIV who died (and had ART data) had been on ART at the time of mpox diagnosis
- Median times from symptom onset to initiation of tecovirimat, ART, and any second mpox therapeutic (vaccinia immunoglobulin, cidofovir, and/or brincidofovir) (Figure 3) were 10, 34, and 45 days.



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infection was the risk factor	•
d with severe to fatal mpox in	
e than one fifth of patients	•
-of whom 91% had AIDS. All	•
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all people with HIV in care to	•
nerence remains a critical public	

er the study period (n=103), 97% (100) were male,	ļ
encing homelessness.	

• ICU-level care was required by 22% (23) of patients, and 21% (22) died. All but two who died had HIV.

## **ADDITIONAL KEY INFORMATION**

### CONCLUSIONS

• In immunocompromised persons, mpox can be a severe and systemic illness with a high mortality.

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• Black and unstably housed populations were especially affected—echoing inequities in the HIV care continuum.

• There were notable delays from symptom onset to initiation of ART and mpox therapies beyond tecovirimat.

• All patients with suspected mpox should be tested for HIV and promptly initiated on ART if indicated.

• Identifying and engaging all people with HIV in care remains a critical public health priority to reduce mpoxassociated morbidity and mortality.

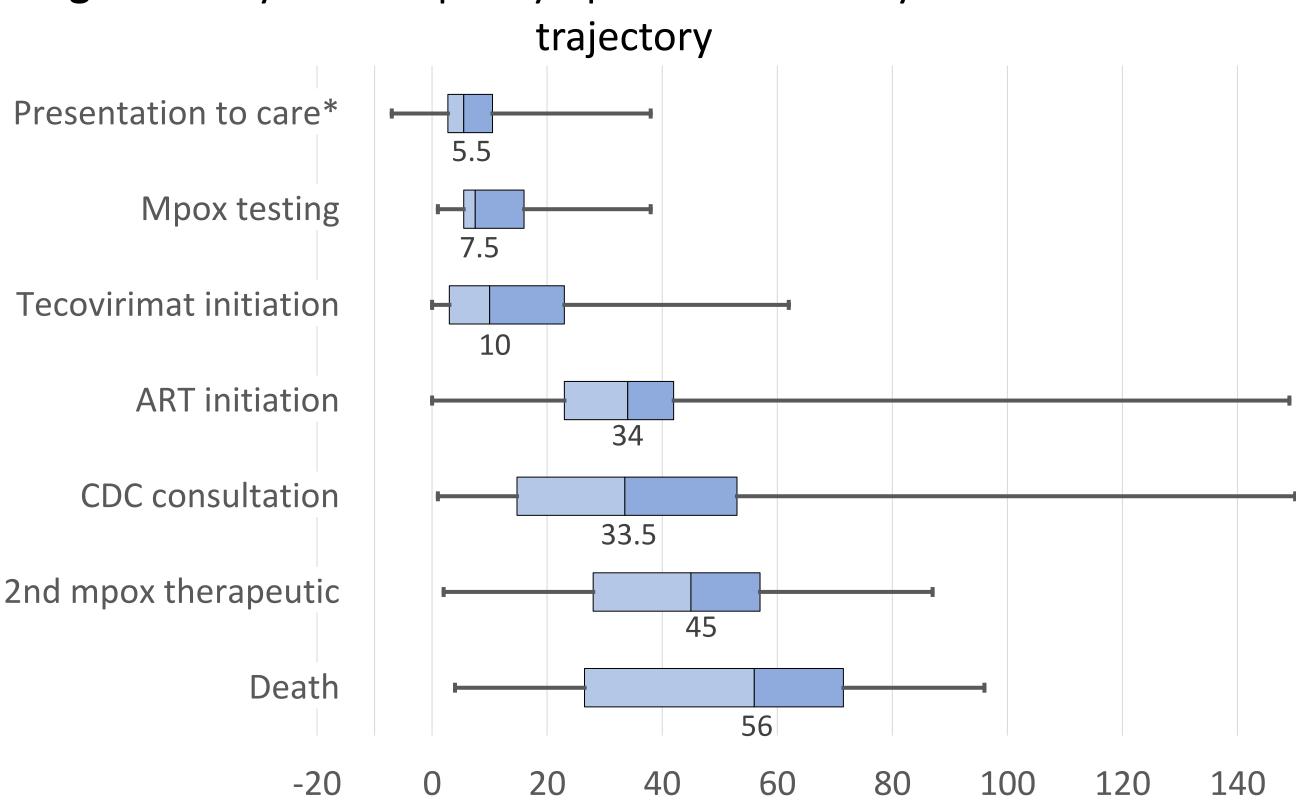
• Limitations: all data was based on provider reporting. No confirmatory medical record review was performed.

• Acronyms: ART (antiretroviral therapy)

• Photos used with permission from patients or next of kin. • Photo reference (Patient A): Miller MJ et al. MMWR Morb Mortal Wkly Rep 2022;71:1412–1417.

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• We thank our health department and clinical colleagues, and the patients and families who consented to sharing photographs to help others affected by this disease.



**Figure 3**: Days from mpox symptom onset to key events in illness

\*Note: two patients developed mpox rash while hospitalized, after presenting for other symptoms (pneumonia, small bowel obstruction)

Patient B