Day 18: Healthcare Inequality

People with disabilities face disadvantages to accessing the healthcare system compared with people without disabilities due to social and environmental inequalities such as lower educational levels, lower income, and higher unemployment. An inability to navigate the medical labyrinth of services, whether physically or programmatically, creates significant barriers. This may result from any number of situations, such as an inability to connect with a healthcare professional who will see them, physical limitations to visiting a doctor’s office, and an inability to understand or adhere to a course of treatment.

Additionally, people with disabilities may not be prioritized by the healthcare system or are not taken seriously. Consider these common healthcare scenarios within the disability community:

- During the COVID-19 pandemic, many accounts arose of doctors and medical facilities prioritizing critical resources based on disability, with disabled people being pushed to the back of the line.
- People with psychiatric diagnoses often face disbelief from medical professionals who mistake physical health symptoms for mental health disabilities; as a result, they are disproportionately likely to die within 15 days of release from an ER visit.
- Individuals with chronic illnesses often wait years for a physician to listen to their concerns and seriously pursue diagnostic testing, and even longer to receive an accurate diagnosis.
- People with disabilities may receive over-medication and reproductions healthcare because medical professionals assume they are not sexually active or do not want to have children.
- Many people with fragrance-reactive disabilities lack access to healthcare facilities because the fragranced cleaning products used in building maintenance can trigger seizures, migraines, and anaphylaxis.

In many states, Medicaid provides Home and Community Based Services (HCBS) to disabled residents so they can live independently in the community rather than being locked into institutions, such as residential nursing facilities and psychiatric hospitals. But the level of care provided varies drastically from state to state. A person who receives 24/7 care from direct support staff in one state may move to another state with less funding for HCBS, they could end up institutionalized.

Additionally, Medicaid often pays extremely low wages to these direct support workers, leading to a staffing shortage.

Learn More

Read:
- Many doctors have negative perceptions of patients with disabilities — and that impacts quality of care, study finds
- My Experience with Diagnostic Overshadowing
- Health Disparities and People with Disabilities

Watch:
- How Health Care Makes Disability a Trap (13:50, includes captions)
  https://youtu.be/7Lfxle9UwCI
- Navigating Health Care with a Disability: Our Stories, a Focus on People with Disabilities (2:59, includes captions)
  https://www.youtube.com/watch?v=VgbPFV0i3vA

Listen:
- As Hospitals Fear Being Overwhelmed by COVID-19, Do the Disabled Get the Same Access? (12:00)
  (transcript here: https://www.npr.org/transcripts/945056176)

Discussion:
- Think about the last time you visited a healthcare professional. If you're non-disabled, how might your experience differ if you had a disability?
- How might the healthcare experiences of people with visible disabilities differ from those of people with non-apparent disabilities?
- How does the medical model of disability impact the quality of healthcare that people with disabilities receive?
- Why do you think many states are more willing to pay for people with disabilities to live in nursing homes than to pay for Home and Community Based Services (HCBS)?

[Image description: Illustration by Cap Pannell of a large red cross where the left arm is lowered to create a stairway and a person in a wheelchair faces it, unable to proceed.]

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