
THE PATIENT VOICE

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Online Survey

Anonymous online survey that was sent globally to people living with CHB

Telephone Interviews

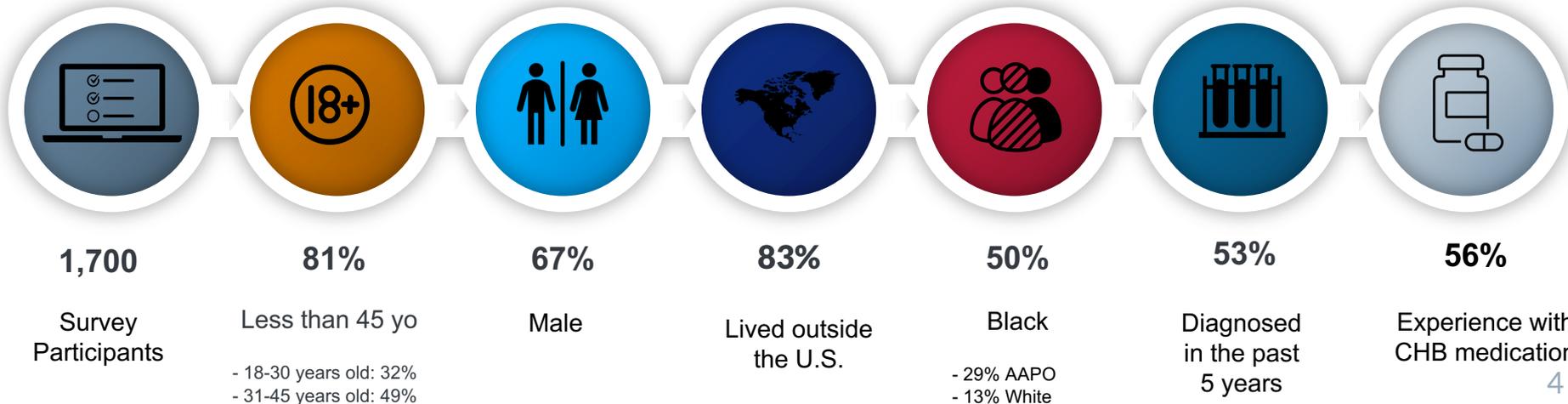
In-depth telephone interviews with people living with CHB in the U.S.

ONLINE SURVEY

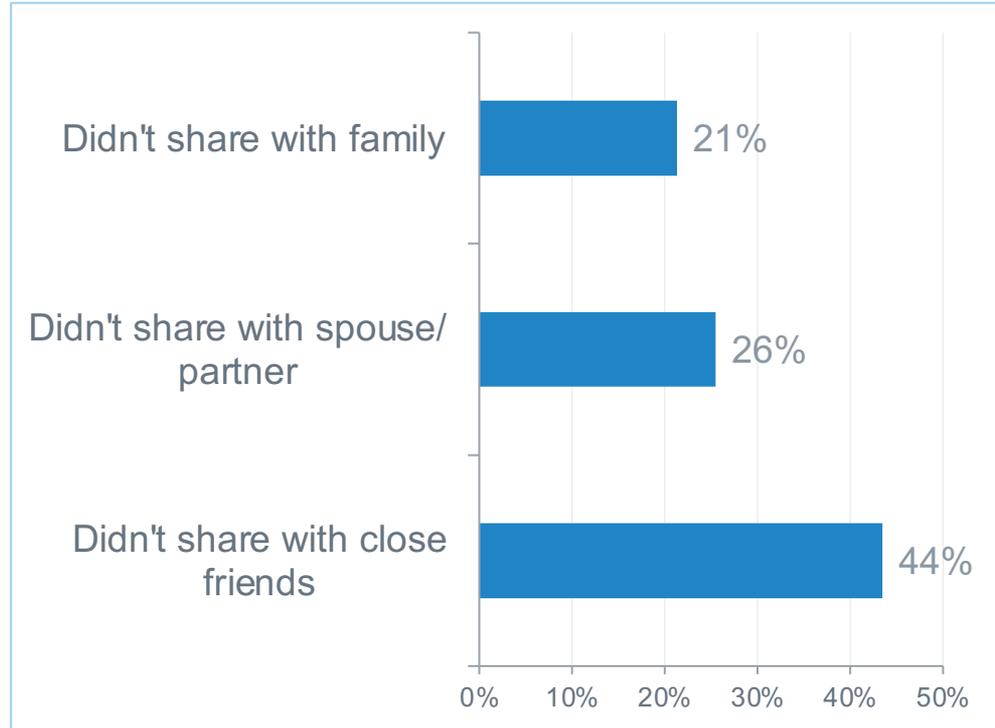
Patient Engagement Survey



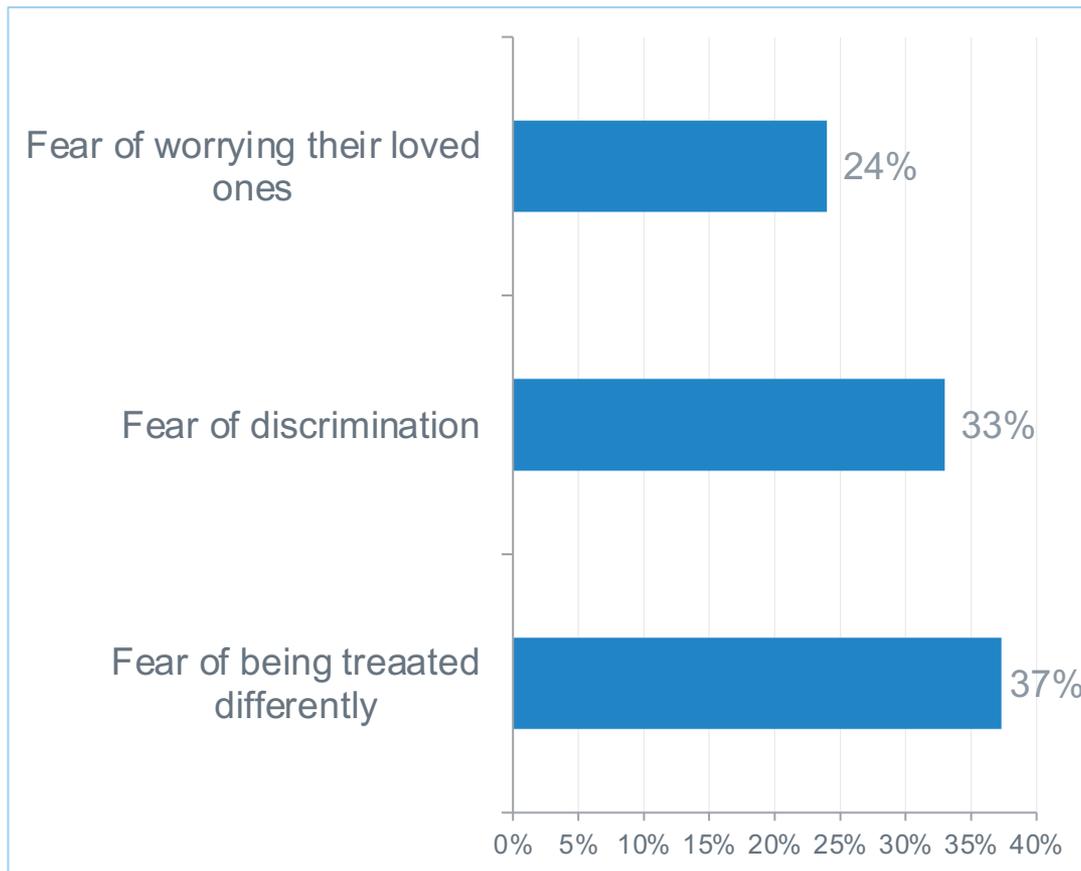
- ▷ Document the physical, emotional, and social impact of living with chronic hepatitis B (CHB)
- ▷ Treatment experiences, challenges and perceived value of current treatments
- ▷ Perspectives on future treatments and clinical trials



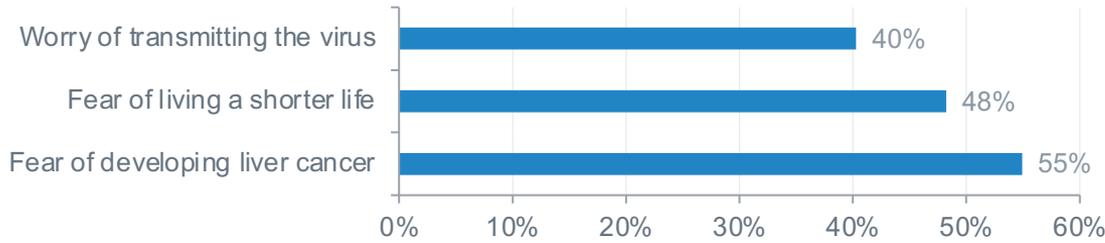
Sharing CHB Status with Close Circle



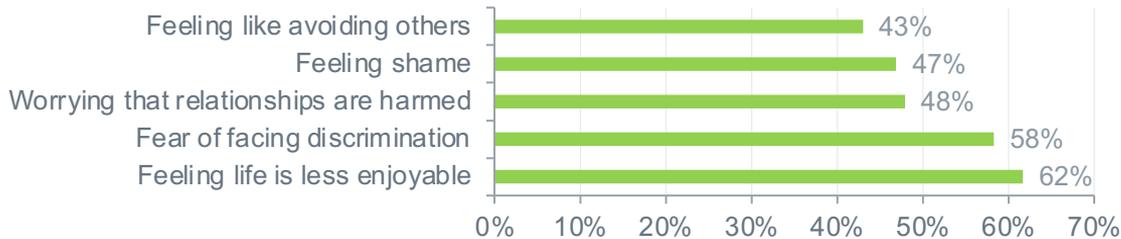
Reasons for not sharing CHB Status



Fears related to disease prognosis

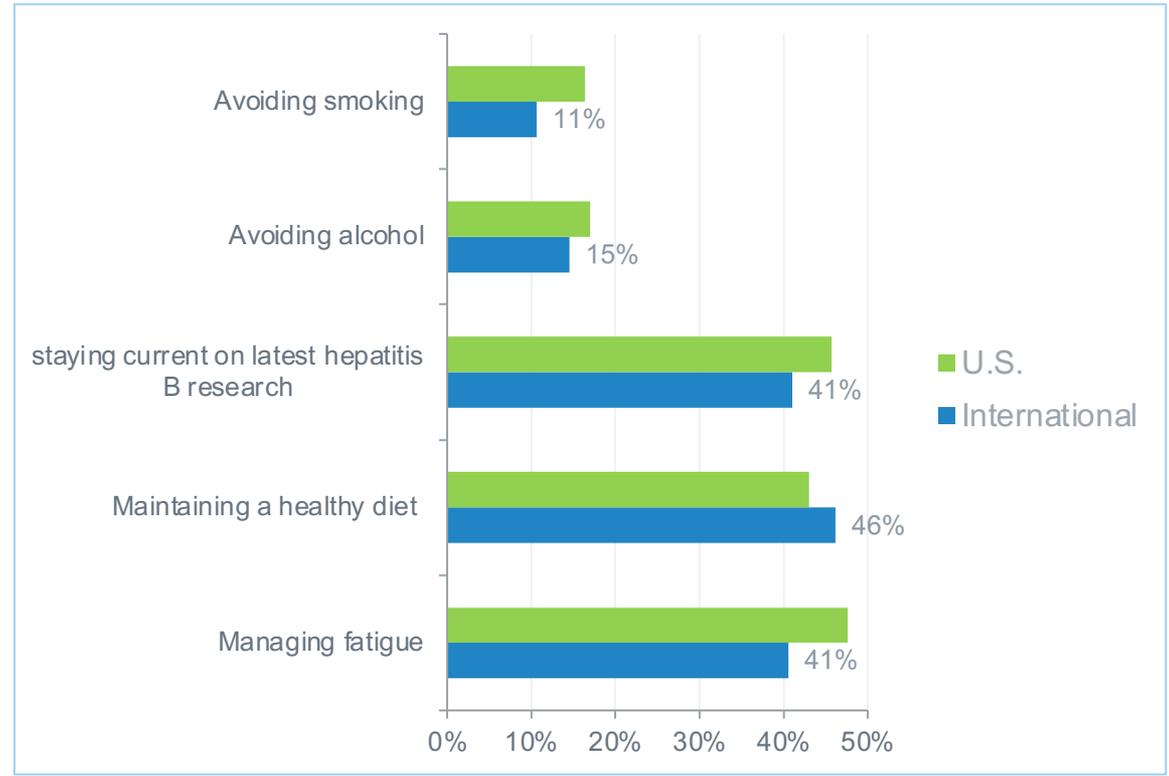


Emotional Impact on Social Aspects of Life

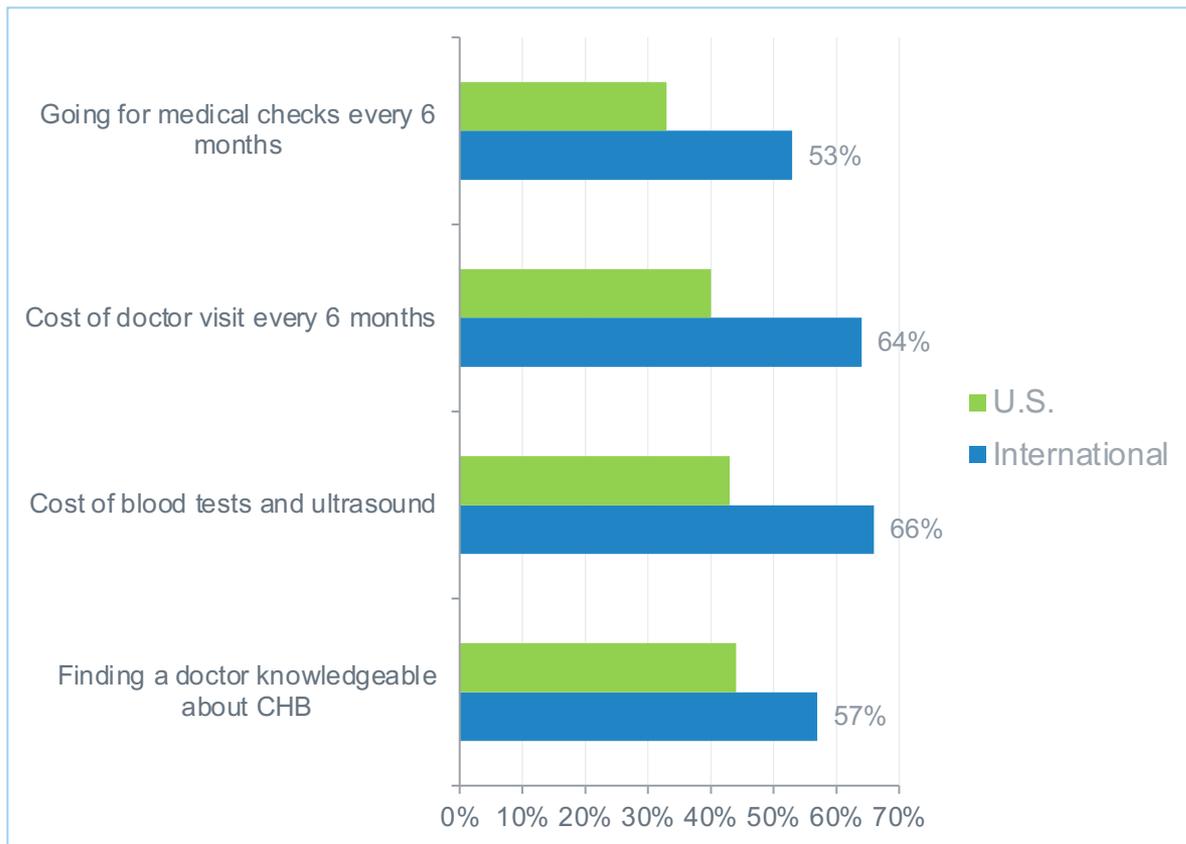


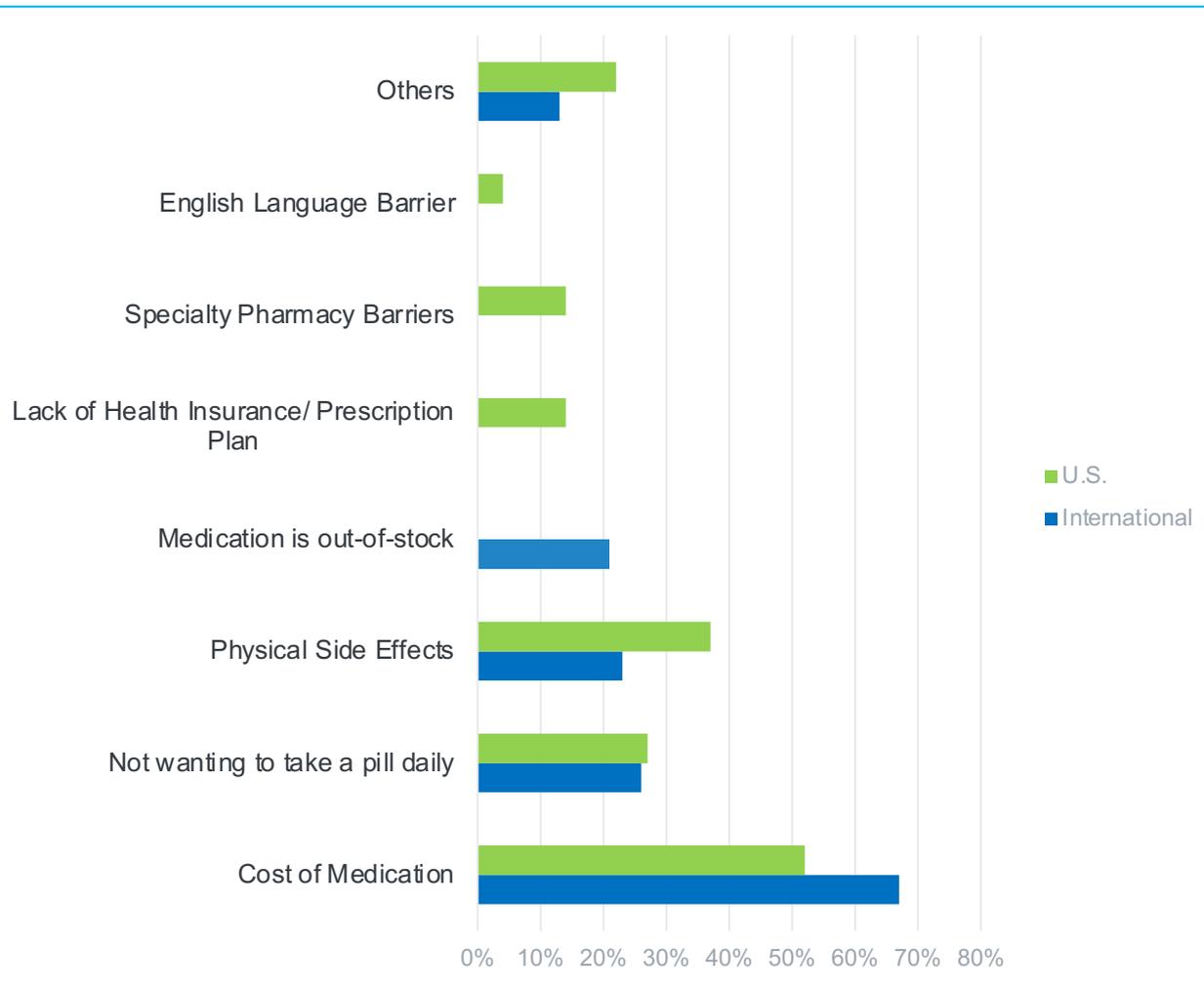
Emotional Burden of Living with CHB

Challenges Living with CHB



Challenges Managing CHB





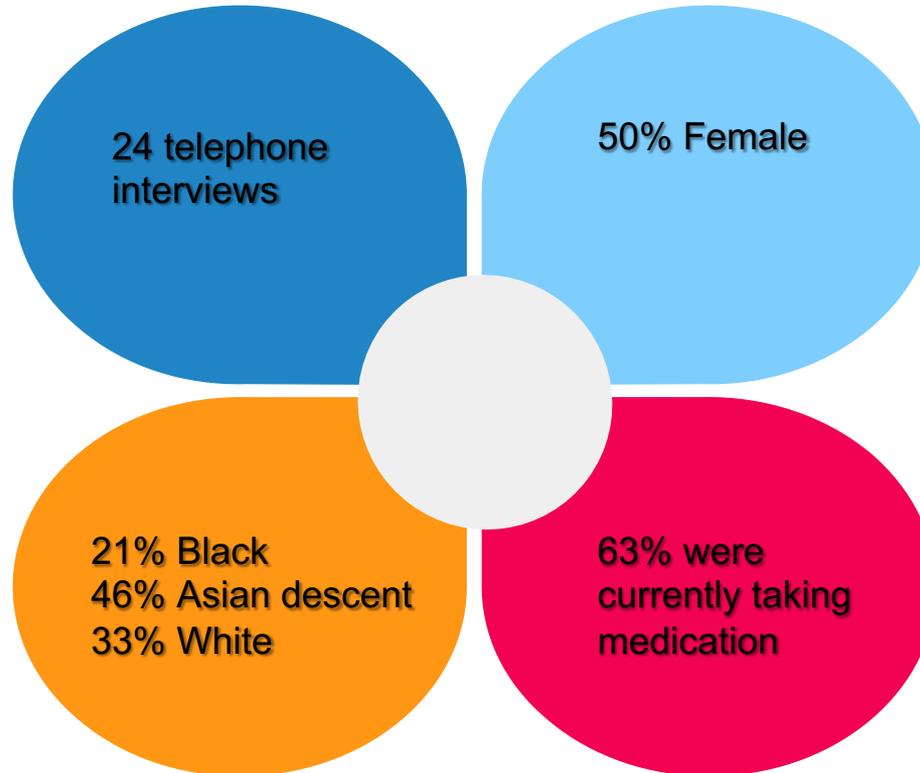
Barriers to Access to CHB Medication

Telephone Interviews



“... I haven't been in a romantic relationship for years and years and years, and you know that certainly would be scary if I met somebody, to tell them”

Telephone Interviews





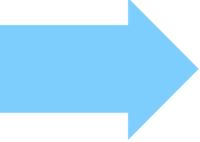
Physical and emotional impact of living with CHB:

- Pain and fatigue were particularly among the most frequently cited physical impact
- Stigma, isolation, anger and fear of developing liver cancer



Impact of CHB on career choices

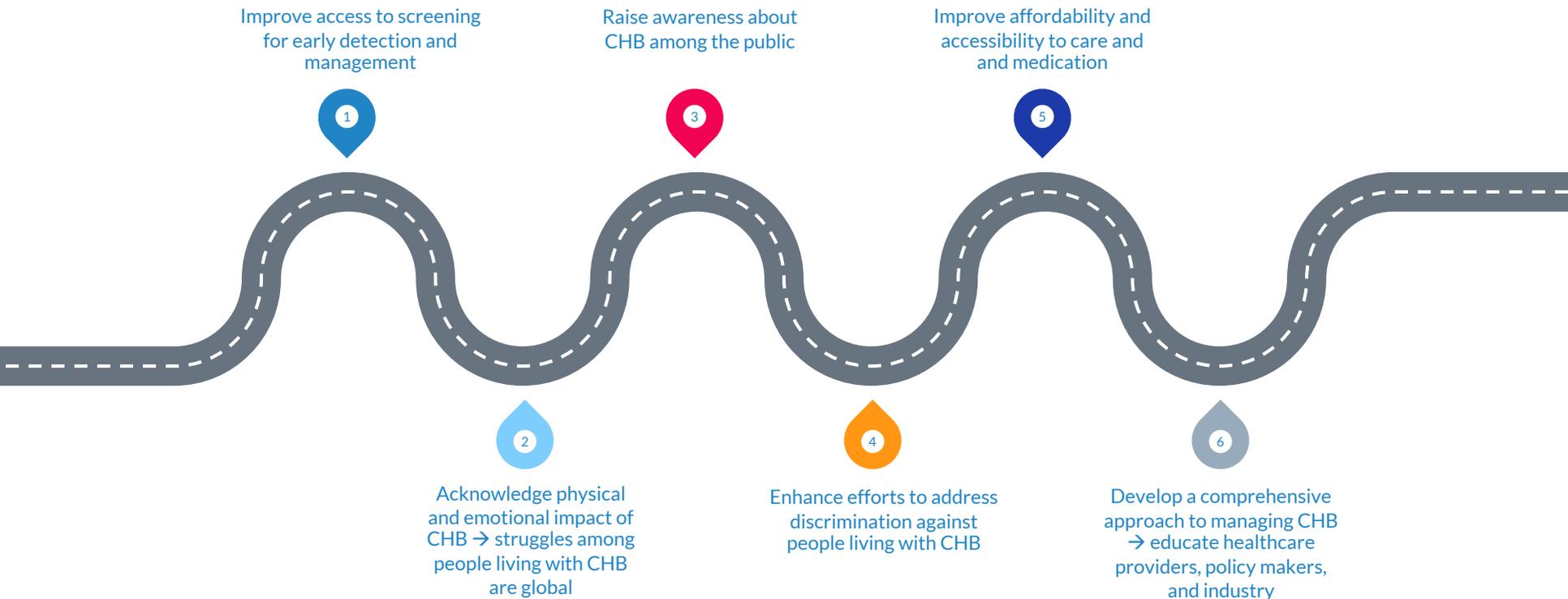
- Avoiding jobs that require disclosure
- Avoiding jobs that are physically taxing



Challenges managing CHB

- Finding a knowledgeable doctor
- Cost of care, even among those who have health insurance
- Taking a daily pill for life

Considerations for a Roadmap



Thanks!

