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### **OBJECTIVES**

The purpose of health technology assessment (HTA) agencies globally is to make evidence-based evaluations of the value of new health interventions. Increasingly, HTA agencies ask manufacturers to include reference to health equity in their submissions. Following an earlier pilot study, this study aims to understand whether and how three cost effectiveness-based HTA agencies (NICE [United Kingdom], CDA [Canada] and PBS [Australia]) ask manufacturers to provide evidence relating to health equity, and how frequently they refer to that evidence in their assessment reports

### **METHODS**

A comparison of the most recent HTA agency guidance (including submission templates) was undertaken to assess the nature of agency requests for health equity data or insight. Analysis was performed to assess the presence and role of health equity-related drivers and detractors of value in HTA assessments by NICE, CDA and PBS in 2024. All 68 pharmaceuticals assessed in non-terminated single technology appraisals (STAs), multi-technology appraisals (MTAs), or highly specialized technology (HST) appraisals by NICE in 2024 served as the index list for review of assessments by all three agencies. We considered elements of value in two broad categories; therapy area-related value (e.g. investment in indications that disproportionately impact vulnerable or underserved populations) and intervention-related value (features of the intervention that

| may impact equita   | able access)             |                      |                   |   |  |  |   |   |
|---|--------------------------|----------------------|-------------------|---|--|--|---|---|
| RESULTS<br>Figure 1   Health Equity Mentions in HTA Outcomes  |                          |                      |                   | <b>Drug + Indication</b>                            | HTA<br>Agency  | Health Equity<br>Considered  | Impact on Outcome   | Quotes from HTA Decisions Regarding Impact of<br>Health Equity on Outcome   |
|   |                          |                      |                   | Veklury®<br>remdesivir 100 MG FOR<br>COVID-19       |  | Discrimination relating  | To prevent potential inequality of access <b>due to age</b> , a positive                  | "The committee noted that <b>the issues raised could affect</b><br>[children] disproportionately The committee was willing to<br>accept an ICER slightly more than what is usually acceptable<br>because there are no licensed treatments available for children." <sup>4</sup>   |
|   | <b>NICE</b> <sup>1</sup> | CDA-2<br>AMC         | PBAC <sup>3</sup> | Evkeeza®<br>(evinacumab-dgnb)                       | Evkeeza<br>(evinacumab-dgnb)<br>Injection<br>Homozygous familial<br>hypercholsterolemia<br>Casgeys<br>(exagamglogene autotemcel) | to a protected<br>characteristic (age)                                       | recommendation was awarded<br>despite being not cost-effective<br>in that population      |   |
| Assessments<br>Examined   | 68                       | 45                   | 30                |   |  |  |   |   |
| Health Equity<br>Mentioned  | 42                       | 14                   | 4                 | (exagamglogene autotemcel)<br>Transfusion dependent |  | Disproportionate<br>prevalence and impact<br>among specific ethnic<br>groups | A stimate was acconted due to   | "The committee concluded that it was <b>willing to take health</b><br><b>inequality into account</b> in its decision making by <b>accepting a higher</b><br><b>cost-effectiveness estimate</b> than it otherwise would have done." <sup>6</sup>   |
| No Impact   | 36                       | 3                    | 3                 | FABHALTA®<br>(iptacopan) 200 mg<br>capsules         |  | Increased accessibility  | The ability of the product to<br>meet identified unmet needs<br>regarding more convenient | "There is an unmet need for effective therapies that provide a<br>more convenient route of administration [important] for patien<br>living in remote communities that may lack access to an infusion<br>center. [The committee] concluded that [FABHALTA] met some of<br>the needs and provide[s] an oral treatment option that can be<br>administered in a patient's home." <sup>7</sup> |
| Unclear Impact  | 3                        | 9                    | 1                 | Paroxysmal nocturnal<br>haemoglobinuria             |  |  |   |   |
| Positive Impact   | 3                        | 2                    | 0                 | ORGOVYX   | ORGOLYK.   |  | routes of administration (for<br>patients in remote<br>communities) was noted in the      |   |
| Non-terminated NICE list of products search   |                          |                      |                   | (relugolix) 120 mg<br>tablets                       |  |  | <b>rationale</b> for recommending reimbursement   | [ORGOVYX] met some of the needs identified by patients, such as<br>being convenient to take." <sup>8</sup>  |
| <ul> <li>NICE and the CDA are the most consistent with regards to<br/>examining health equity, often including sections dedicated to</li> </ul> |                          |                      |                   | Hormone-sensitive<br>prostate cancer                |  |  |   | Clinician Input: "[F]or <b>patients in remote areas</b> of Canada it <b>may</b><br><b>be particularly beneficial</b> ." <sup>8</sup>  |
| equality / ethics   | where health eq          | juity issues are dis | cussed and        | NICE was willing                                    | to accept h  | nigher cost-effective  | ness estimates than usually   | outcomes: in three positive examples identified,<br>y considered acceptable. Of note, for the CASGEVY   |

- committee
- In contrast, most health equity mentions in PBAC reports originate from third-party input, with no indication of how the equity concern was considered by the HTA committee

Ask A Question:

# HEALTH EQUITY AND HTA: IS EQUITY DATA SHAPING RECOMMENDATIONS

| Table 1       Distribution of Health Equity Factors and Impact on Assessment |  |  |      |   |     |      |  |  |  |  |  |
|--|--|--|------|---|-----|------|--|--|--|--|--|
|  | Type of Health Equity-Related Factors Addressed or Mentioned |  |      |   |     |      |  |  |  |  |  |
| Impact on<br>assessment  |  | - <b>Related</b> (e.g., dis<br>pact of disease ba<br>age, gender, etc) |      | Intervention related (e.g., geographical barriers in relation to treatment accessibility) |     |      |  |  |  |  |  |
|  | NICE   | CDA  | PBAC | NICE  | CDA | PBAC |  |  |  |  |  |
| No impact  | 30   | _  | 2    | 9   | 3   | 1    |  |  |  |  |  |
| Unclear Impact   | 2  | 4  | _    | 1   | 6   | 1    |  |  |  |  |  |
| Positive Impact  | 3  | _  | _    | 1   | 2   | _    |  |  |  |  |  |
| Total  | 35   | 4  | 2    | 11  | 11  | 2    |  |  |  |  |  |

Note: Assessments that mentioned more than one type of health equity factor are listed in this table separately for each health equity type (i.e., an assessment mentioning both indication-related factors and intervention related accessibility is counted in both).

- The majority of NICE assessments that mentioned health equity referred to therapy area-related health equity. Most of these assessments acknowledged how the indication may be more prevalent in certain populations (e.g., minority ethnic groups) but note that because the final recommendation does not restrict access to treatment for some people over others, the committee agreed that this was not a potential equalities issue. Similar results were noted in the case of intervention-related equity factors (e.g., geographical barriers to access)
- In cases where CDA assessment reports mentioned health equity, the most frequent factor referenced was the impact of geographic barriers to treatment on equitable access
- While PBAC reports referred to health equity considerations raised by physician or patient advocacy groups, these considerations were rarely mentioned in the discussion points for the committee's decision making

**Table 2** | Instances of Positive impact of Health Equity Factors on Assessment

- assessment, the company accounted for health inequalities in its submission using a distributional cost-effectiveness analysis (DCEA) to create an equity-weighted ICER (the only 2024 assessment identified to do so)
- In Canada, while the **CDA assessments do not clearly indicate** how health equity may have impacted the outcomes, equity considerations are noted in the rationale for the final recommendation

### **DISCUSSION & CONCLUSION**

- considering health equity in their decision making, with varying frequency; other agencies are likely to follow
- Suggest: NICE, CDA and PBAC are already referencing or • Currently there is no widely accepted quantitative
- methodology for measuring health equity (outside of ICER) • NICE and the CDA most consistently referenced health equity, with clear evidence of health equity considerations impacting NICE assessment outcomes, while health equity considerations are cited in CDA assessments without clear
- decision impact • While health equity considerations are referenced in PBAC reports, they are primarily acknowledged as input from clinician or patient advocacy groups; the health equity considerations were acknowledged in the decision-making for only one out of four assessments that mentioned health equity considerations
- Direct evidence of health equity considerations impacting the outcome of assessments is still relatively unusual and often unclear
- At the time of writing, medical benefit-focused markets (such as DEU and FRA) do not mention health equity in their evaluations

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## **ABBREVIATIONS**

**CDA:** Canada Drug Agency; **DCEA:** Distributional Cost-Effectiveness Analysis; HTA: Health Technology Assessment; ICER: Incremental Cost-Effectiveness Ratio; **NICE:** National Institute for Health and Care Excellence; **PBAC**: Pharmaceutical Benefits Advisory Committee; **PBS**: Pharmaceutical Benefits Scheme

