S40 Oral Presentations

OP136 A Comparison Of Health Technology Assessment Recommendations In Australia, Canada, And England: Is There Opportunity For Further Alignment?

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Introduction: Over the past decade there have been increasing interactions between health technology assessment (HTA) agencies through international networks at the policy level and European joint actions at the product level. A pilot project is underway to explore collaboration beyond Europe between HTA agencies in Australia, Canada, and the UK. This study the compared HTA recommendations of new active substances (NAS) appraised by Australia's Pharmaceutical Benefits Advisory Committee (PBAC), Canada's CADTH, and England's National Institute for Health and Care Excellence (NICE).

Methods: Using publicly available data and established benchmarking methodology, we examined 45 NAS appraised by PBAC, CADTH, and NICE between 2017 and 2021. Analysis was performed to assess rollout time from regulatory to HTA recommendation, and to the first HTA recommendation.

Results: Most products were submitted to the Europe Medicine Agency first (89%). However, 71 percent of NAS in Australia and 69 percent in Canada were submitted to HTA in parallel with regulatory review, which shortened overall rollout time. The median HTA submission gap among the three agencies was 140 days in Australia, 102 days in Canada, and 8 days in England. PBAC had the highest number of negative recommendations (51%), followed by CADTH (18%), and NICE (5%). The congruence of HTA decisions was highest between CADTH and NICE (56%), compared with PBAC and CADTH (11%), and PBAC and NICE (20%)

Conclusions: To achieve a more collaborative HTA process it is necessary to understand the rollout time in these jurisdictions. This study identified submission gaps among the regulatory agencies, but there may be more synergy in the future as regulators in the three jurisdictions work collaboratively through the Access Consortium. Currently, the submission gap for the three HTA agencies was mostly within six months, making collaboration on joint assessment possible. We observed divergences in HTA recommendations due to the methodology and decision criteria applied by each agency. Therefore, collaboration on assessment should build on the clinical aspects, although HTA decisions should be grounded in the local context.

OP137 A Qualitative Exploration Of The National Institute For Health And Care Excellence's Impact On International Health Technology Assessment

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Introduction: The National Institute For Health And Care Excellence (NICE) is widely acknowledged as a seminal health technology assessment (HTA) body, known for its transparent and accountable approach to decision-making. This research aimed to investigate the impact of NICE methodology and decisions on international HTA bodies. We sought to identify direct and indirect factors that may influence an international HTA body's methods or outcomes. To the best of our knowledge, this is the first research to use a qualitative approach to understand the influence of NICE on other HTA bodies. Methods: We conducted 13 semi-structured qualitative interviews with HTA and market access experts from industry and academia from nine countries (Brazil, Israel, Italy, Japan, Poland, Saudi Arabia, South Korea, Sweden, and the United Arab Emirates). The interview script was organized into three main sections: comparing NICE methods and processes with other HTA bodies; the impact of specific NICE decisions; and Likert scale questions (to allow for comparability of opinions).

Results: Most interviewees believed their local HTA body would consider NICE's decision when evaluating a medicine. However, the way and extent to which NICE influences HTA varied across countries. The most common means of considering a NICE decision was as background information or context for an HTA evaluation. Generally, interviewees suggested that negative NICE decisions had more impact on local decision-making than positive decisions. Nine of the 13 interviewees agreed or strongly agreed that their country's HTA body considers the decisions of other HTA bodies in their decision-making process. Eleven of the 13 interviewees agreed or strongly agreed that the development of their country's HTA body methods and processes was influenced by NICE.

Conclusions: NICE is perceived to be a seminal HTA body, with continued influence on HTA agencies in other countries. However, the mechanisms and extent of this influence varies considerably between countries. We suggest that implicit factors are likely to contribute more to NICE's influence than individual decisions. Nevertheless, further research is needed to reveal these factors and increase efficiency in international HTA decision-making processes.