National Health Technology Assessment in Turkey After a Decade:

Are Key Principles Followed?

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Introduction

The Health Technology Assessment (HTA) agency of the Turkish Ministry of Health (MoH) was first established in 2012 [1]. The country does not have detailed national guidelines, but the National HTA Directive sets out the process of undertaking HTA [2]. The strategy document prepared by the national HTA agency for 2019-2023 lists compliance with the key HTA principles developed by Drummond as a strategy for the institutionalisation of HTA in the country [3]. The current study aimed to provide an overview of the country's HTA progress since establishing the agency a decade ago, critically reviewing the National HTA Directive and the national HTA reports published by the Turkish MoH's HTA agency.

Methods

National HTA Directive and the HTA reports were accessed on the national HTA agency's website. First, the National HTA Directive was assessed to understand whether the key principles of HTA were included. Second, the compliance of the HTA reports with the principles, and the Directive were evaluated. The principles published by Drummond et al. were chosen as an assessment tool because the agency's strategy document included them as guiding principles. Assessments were completed by two researchers independently, and any discrepancies were dissolved through discussion.

Results

The study included all publicly available national HTA reports (n=8). Most reports (n=5) were published before the establishment of the National HTA Directive in 2019. The reports focused on various topics, including foetal anomalies, cancer, and obesity. The decision-maker was not stated in any of the reports, and none was peer-reviewed. A summary of the assessment of the Directive and HTA reports against the key principles is provided in Table 2. The overall compliance with the key principles was poor.

Table 1. Summary of the included HTA reports

| First author & year | Disease area | Intervention | Comparator | Economic evaluation conducted |
|------------------------------------|-----------------------------------|---|--|-------------------------------|
| Tecirli 2020 | Fetal chromosomal anomalies | Combined test (CT) only | CT+ triple test, CT+ quadruple test, CT+ cfDNA | Cost-effectiveness analysis |
| Arslan 2019 | Sepsis | Molecular rapid tests + Blood culture test | Blood culture test | Cost-utility |
| Mahagaonkar 2019 | Rheumatoid arthritis | bDMARDs | csDMARDs | Cost analysis |
| Kockaya 2018 | Cancer | Cytoreductive surgery + hyperthermic intraperitoneal chemotherapy | Cytoreductive surgery alone | None |
| Ozturk 2018 | Male circumcision | Single-use devices that require minimal surgical implementation | Traditional surgery | Cost analysis |
| Gunal 2017 | Renal insufficiency | Peritoneal dialysis | Haemodialysis | Cost-utility |
| Sener* 2014 | Obesity | Surgery for obesity | No surgery | Cost analysis |
| Karadayı 2013 (short report) | Erectile dysfunction | Electroshock wave treatment (ESW) | Usual care (not defined) | None |

Table 2. Compliance with Drummond's key principles of HTA

| | STRUCTURE OF HTA | Included in Directive? | Compliance in HTA reports |
|----|--|------------------------|---------------------------|
| 1 | Was the goal and scope of the HTA explicit and relevant to its use? | Partly | 50% |
| 2 | Was the HTA report an unbiased and transparent exercise? | No | 6% |
| 3 | Did the HTA report include all relevant technologies? | Partly | 63% |
| 4 | Was there a clear system for setting priorities for HTA? | Partly | 0% |
| | METHODS OF HTA | | |
| 5 | Did the HTA report incorporate appropriate methods for assessing costs and benefits? | Partly | 69% |
| 6 | Did the HTA consider a wide range of evidence and outcomes? | Yes | 25% |
| 7 | Was a full societal perspective considered? Was not having a societal perspective justified? | No | 13% |
| 8 | Was uncertainty surrounding estimates explicitly characterised? | No | 13% |
| 9 | Were issues of generalizability and transferability considered and addressed? | No | 0% |
| | PROCESSES FOR CONDUCTING HTA | | |
| 10 | Were all key stakeholder groups actively engaged? | Yes | 50% |
| 11 | Have all available data been seeked actively by those undertaking HTA? | No | 13% |
| 12 | Have the implementation of HTA findings been monitored? | Yes | Can't tell |
| | USE OF HTA IN DECISION MAKING | | |
| 13 | Was the HTA published before the implementation of the technology? | No | 13% |
| 14 | Were the HTA findings communicated appropriately to different decision makers? | Partly | Can't tell |
| 15 | Was the link between HTA findings and decision-making processes transparent and clearly defined? | No | 0% |
| | • | | |
| | Overall compliance with the principles | Poor 20% | Poor 25% |
| | | 30% | 25% |

Conclusion

The role of HTA in healthcare decision-making in Turkey remains unclear. The study findings provide important learnings for the HTA agencies, researchers, and policymakers in countries where HTA is developing. The link between HTA findings and decisionmaking processes needs to be explicitly defined and formalised. The HTA reports published by the national HTA agency of the Ministry of Health lack consistency, and significant variations exist in how the evidence is gathered and assessed. There is an urgent need to establish and adopt national HTA guidelines that are at the international standards.

References

- 1. Research Development and Health Technology Evaluation Department, National Health Technology Assessment Strategy (2019-2023). 2019, Ministry of Health.
- 2. Turkish Ministry of Health (MoH), National Health Technology Assessment Directive. 2019.
- 3. Drummond, M.F., et al., Key principles for the improved conduct of health technology assessments for resource allocation decisions. International journal of technology assessment in health care, 2008. 24(3): p. 244-258.