

With us today

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During COVID-19, we all witnessed

- Governments purchasing quantities of vaccines or medicines even before they were authorised, even before any health technology assessment
- HTA typically advises on
 - Relative efficacy of different vaccines or treatments
 - Which populations to benefit?
- Can it be made real-time? Can it inform decision makers and how?

First questions to the panel

- 1. How can HTA experts scan the horizon to be prepared to provide rapid reports for decision making? Is there a role for assessments at the EU level to achieve consistency and harmonise messages communicated to the public?
- 2. Can real-world data analysis help when clinical trial results contradict each other, or not provide enough information for all target populations?
- 3. How can HTA agencies respond to the rapidly changing evidence base where information can rapidly become obsolete and frequent updates are needed?
- 4. Misinformation and fake news are everywhere, how can HTA outcomes be communicated to the public against this backdrop?



And of course, your own questions and experiences!