



Utrecht University

Differences in health technology assessment recommendations between European jurisdictions: the role of practice variations

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WHO Collaborating Centre for
Pharmaceutical Policy and Regulation



National Health Care Institute

Disclosures

- Partly funded by the Dutch National Health Care Institute

Study background

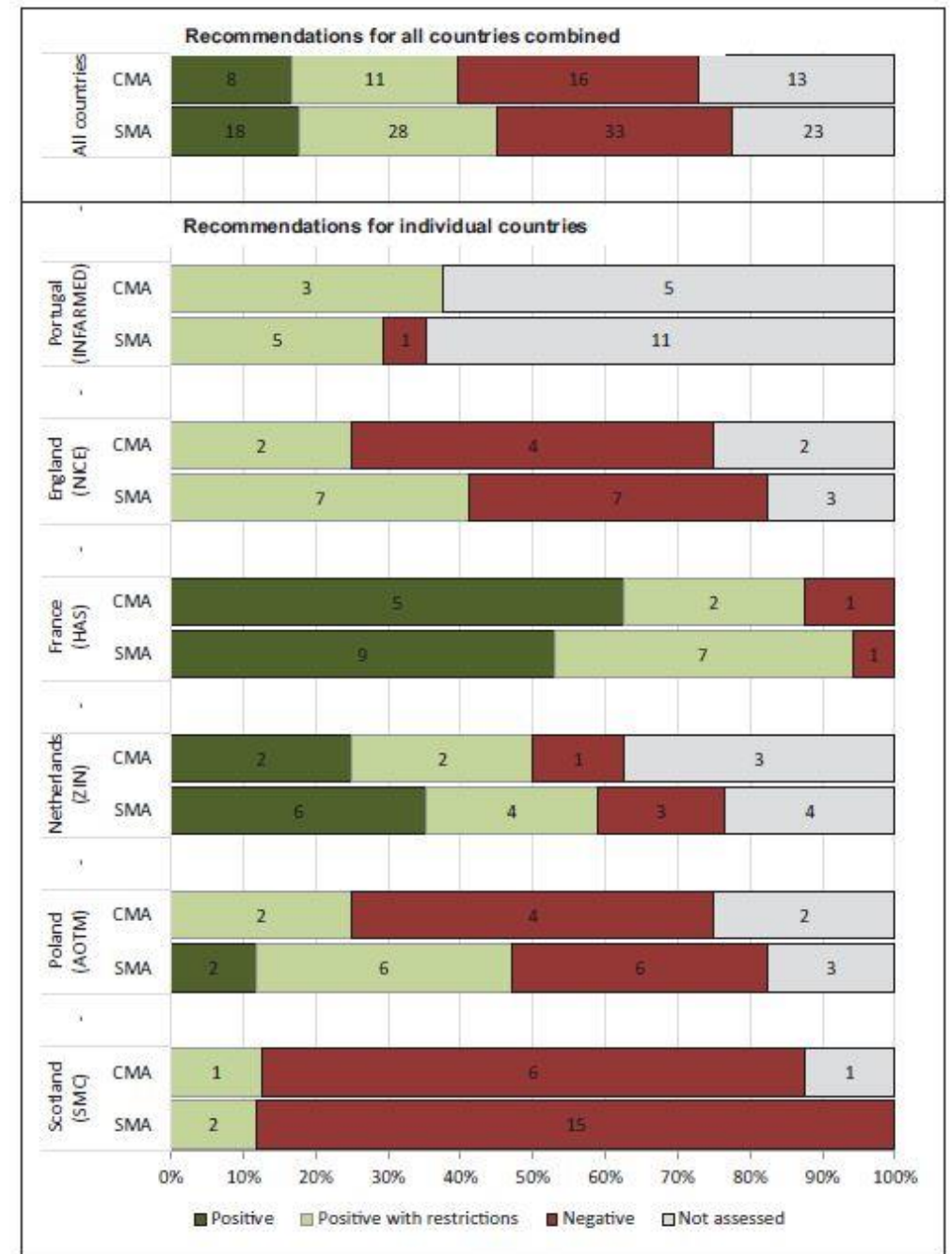
- HTA informs patient access
- Divergent opinions in Europe
- Divergent patient access

Because of evidence?

or

Because of practices?

Lipska et al. CP&T. 2015



Objectives

- Impact of practice differences

On

- Comparisons of HTA body recommendations

For

- Conditionally approved drugs

Methods

Inclusion

Conditionally approved drugs (N=27) up until June 2017

England/Wales, France, Germany, The Netherlands and Scotland

Methods

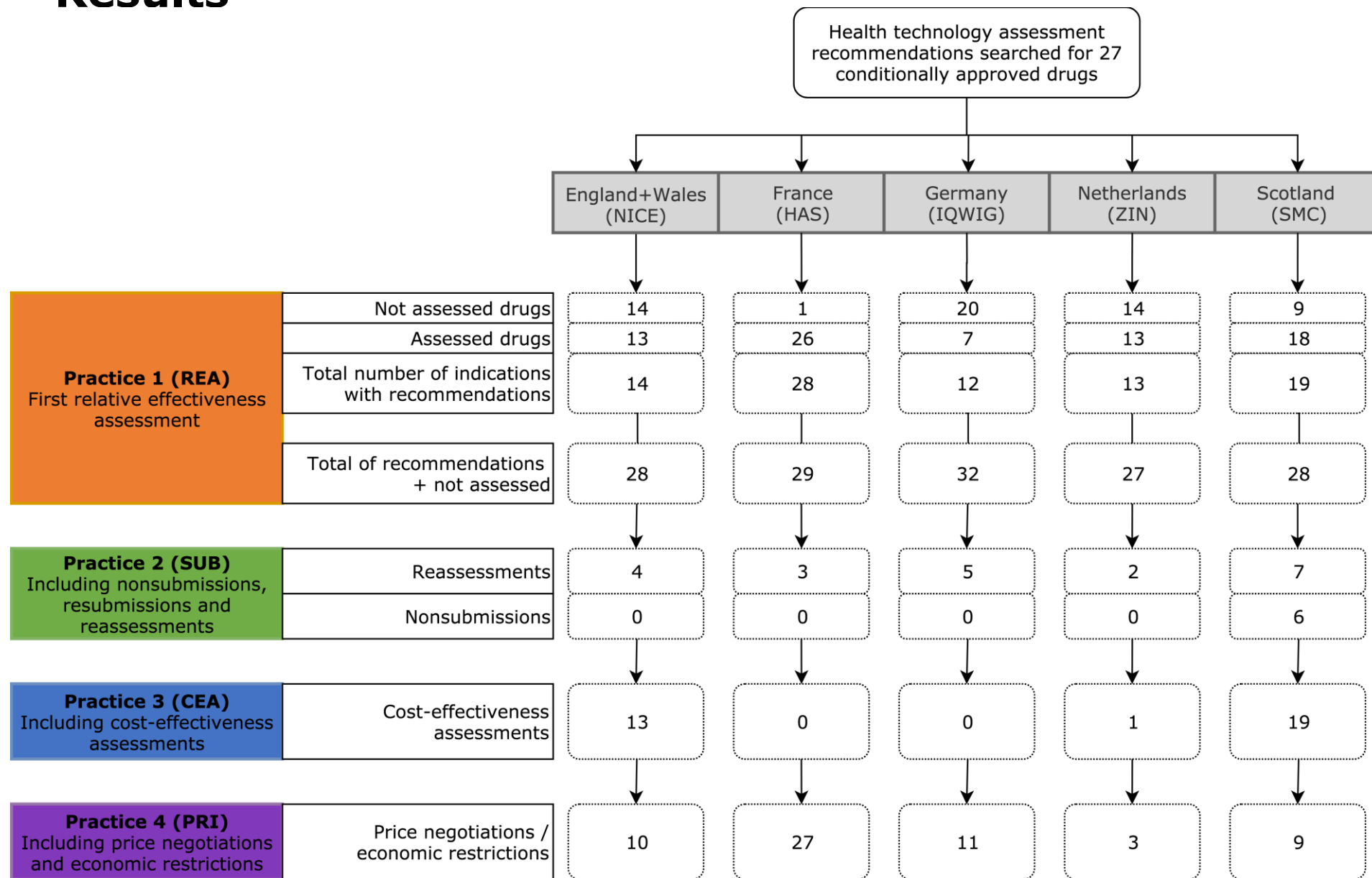
Practices assessed

	REA	SUB	CEA	PRI
<i>First relative effectiveness assessment</i>	Yes	Yes	Yes	Yes
<i>Reassessments / resubmissions / non-submissions</i>	No	Yes	Yes	Yes
<i>First and subsequent cost-effectiveness assessments</i>	No	No	Yes	Yes
<i>Economic restrictions / price negotiations</i>	No	No	No	Yes

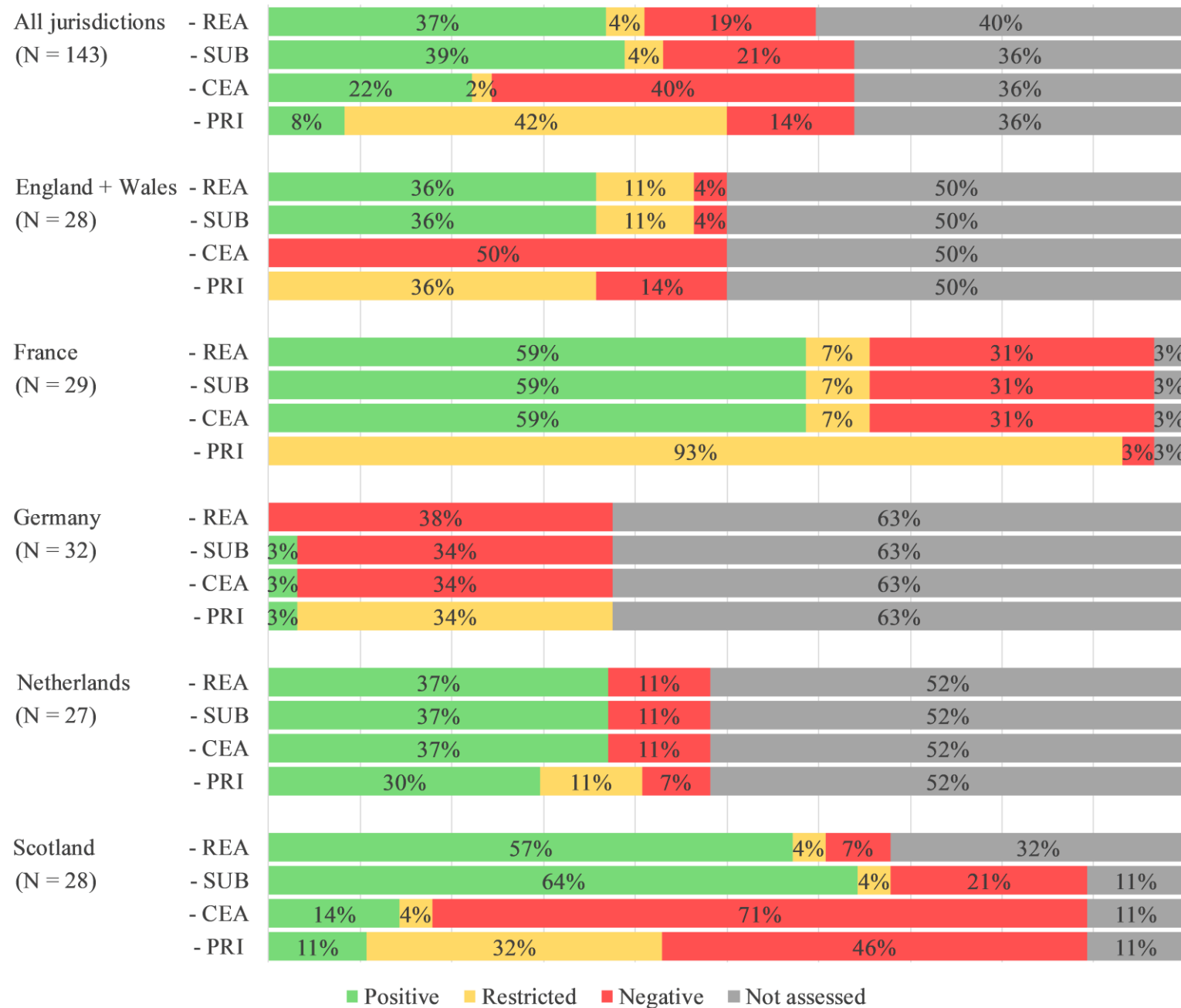
Outcomes of interest

Recommendation outcome (positive, restricted, negative)

Results



Results



Conclusions

Variations in HTA practices:

- Have substantial and significant impact on comparisons
- Should always be considered when comparing HTA bodies

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