

Recommendations for designing inclusive consent processes in trials



Clinical trials are essential to improving the health and well-being of all populations.



People with communication and/or decision-making needs are often excluded from trials.



Inclusive consent processes may enable people to make informed decisions about taking part.



The OPTIMISE recommendations can help ensure consent processes are designed to be accessible.



The toolkit will help you to use the recommendations, with links to information and resources.

OPTIMISE Domains



Content



Language



Format



Design



Support



Timing



Environment

Guiding Principles



Collaboration



Ethics



Reflection

**Link to OPTIMISE
recommendations and toolkit:**



[www.capacityconsentresearch.com/
optimise](http://www.capacityconsentresearch.com/optimise)

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