# Multi-centre evaluation of atlas-based and deep learning contouring using a modified Turing Test

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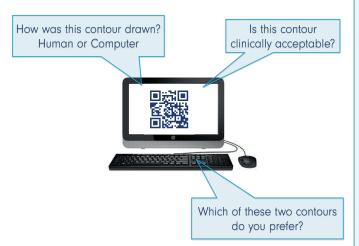
# Objective

While quantitative assessment of autocontouring quality is useful, frequently used measures do not necessarily indicate clinical acceptability or benefit. In contrast, clinical based assessment metrics, such as time saved with autocontouring or subjective evaluations, are both time consuming to perform and difficult to implement in a multi-centre evaluation.

Inspiration is taken from the Artificial Intelligence community to propose an assessment method based on the "Turing Test". The objective of this study was to perform a multi-centre evaluation of two autocontouring methods using this approach.

# Materials and Methods

A website was set up to facilitate multi-centre comparison, showing images and contours in a blinded fashion.



Use the QR-code to try it for yourself at http://www.autocontouring.com

Clinical test cases from a single institution (40 thoracic, 20 prostate)

Types of contour Existing clinical

- Multi-atlas contouring
  - (Mirada Workflow Box 1.4)
  - Deep Learning Contouring (DLC) (Mirada DLCExpert<sup>TM</sup> prototype)

Clinical participants (clinicians, dosimetrist or technicians)

Institutions participated



Randomly chosen questions answer by each participant

# Results

### Thoracic segmentation

Itour preference in blind side-by-side comparison

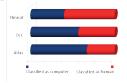
Classification of contours

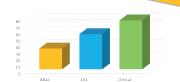
Acceptance of contours (%)

# Prostate segmentation

Classification of contours

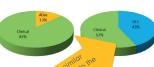
Acceptance of contours (%





Contour preference in blind side-by-side comparison





### Conclusions

The Turing Test style assessment method provided an easy way to perform web-based multi-centre validation of autocontouring.

This study found that autocontours may be confused with clinical ones, when reviewed blindly. DLC showed increased clinical acceptance for prostate OAR contouring compared to atlas contouring. For thoracic imaging, DLC contours were accepted at a similar rate to clinical ones.

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