Procedure for:

Clinical trials with therapeutic products capable of emitting ionising radiation

Radiation source as a component of a medicinal product (radiopharmaceuticals)

Category A trials

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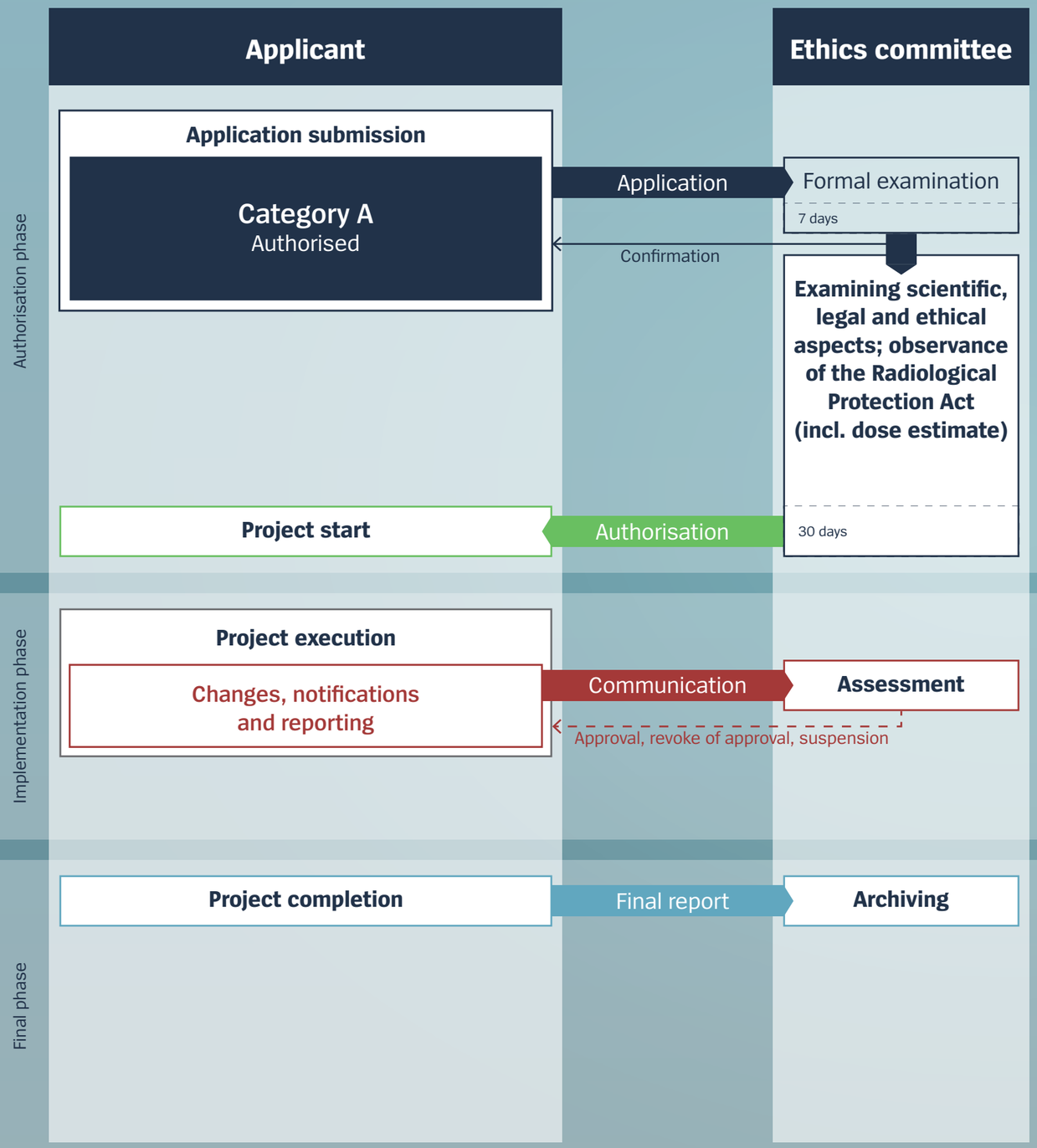

Relevant legal texts:

Authorisation phase
Art. 24–26 ClinO, Annex 3 nr 5 ClinO
Art. 28 RPO (Dose limit)

Implementation phase
Art. 29, 37, 40–43 ClinO

Final phase
Art. 38 ClinO