

## Consent and consultation arrangements for research with adults lacking capacity

Type of research study	Who should be asked	What they should be asked	What is given	What documentation is needed*
Clinical trial (CTIMP) in England and Wales	Legal representative:  1. Person independent of trial and providing care (e.g relative)  OR  2. Doctor primarily responsible for adult's treatment, or another independent person nominated by healthcare provider	Presumed will of potential participant	Informed consent	Information Sheet for Legal Representative Consent Form for Legal Representative
Clinical trial (CTIMP) in Scotland	Legal representative:  1. Guardian or welfare attorney authorised to take decisions about research OR  2. Nearest relative OR  3. Doctor primarily responsible for adult's treatment, or another independent person nominated by healthcare provider	Presumed will of potential participant	Informed consent	Information Sheet for Legal Representative Consent Form for Legal Representative
Research that is not a clinical trial (non- CTIMP) in England and Wales	Consultee:  1. Unpaid person with an interest in the person's welfare  OR  2. Person in a professional capacity who is independent of project	Wishes and feelings of potential participant	Advice as to the person's wishes and feelings and whether they would decline to take part if they had capacity	Information Sheet for Consultee Declaration Form for Consultee
Research that is not a clinical trial (non- CTIMP) in Scotland	Guardian or welfare attorney authorised to take decisions about research  OR  Nearest relative	Informed consent	Informed consent	Information Sheet for Nearest Relative/Guardian or Welfare Attorney Consent Form for Nearest Relative/Guardian or Welfare Attorney

<sup>\*</sup> In addition to participant information such as accessible/pictorial versions, and information and consent documents for participants who regain or recover capacity