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| 1. **Does this application make sense?**
 |
| **Question** | **Reference documents** | **Y/N** |
| 1. Has an Independent Scientific Review (ISR) been conducted and provided?
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| 1. Is the application GDPR compliant?
 | Ethics Application, Participant information sheet, consent form |  |
| 1. Has a project code been provided (if funded or healthcare research – involves NHS)?
 | Ethics Application |  |
| 1. Has a Data Management Plan been provided?
 | Ethics Application |  |
| 1. Is all information provided in the ethics application? (Also has too much information been provided?)
 | Ethics application: only need to state what they intend to do, protocol should not be integral to this. |  |
| 1. Is it clear what is involved in the study?
 | Ethics application, Participant information sheet |  |
| 1. If there is more than one part to the study, has this been made consistently clear throughout the application?
 | Ethics application, Participant information sheet (if appropriate)/consent form (if participants need to consent to more than 1 study part) – consider more than one consent form. |  |
| 1. If there are different groups of participants, is it clear how they differ?
 | Ethics application, Participant information sheet |  |
| 1. Is it clear if different participants are required to do different things
 | Ethics application, Participant information sheet (perhaps more than one version) |  |
| 1. Is more than one consent form appropriate?
 | Ethics application, Consent form(s) |  |
| 1. Is it clear how potential participants are identified?
 | Ethics application |  |
| 1. Is it clear how potential participants are initially approached?
 | Ethics application |  |
| 1. Is it clear that participants aren’t being harassed/pestered during recruitment?
 | Ethics application |  |
| 1. Is it clear how participants obtain the information sheet?
 | Ethics application |  |
| 1. Is it clear participants have a minimum of 24 hours to review the information sheet before agreeing to take part?
 | Ethics application, Participant information sheet |  |
| 1. Is it clear how participants can decide they want to take part?
 | Ethics application, Participant information sheet |  |
| 1. Is it clear why certain people have been invited to take part in the study?
 | Ethics application, Participant information sheet |  |
| 1. Are sufficient contact details made available so potential participants/participants can contact a member of the study team?
 | Ethics application, Participant information sheet |  |
| 1. Is it clear to participants what the study is about?
 | Participant information sheet |  |
| 1. Is it clear if all sections on the consent form are compulsory to take part (e.g. sometimes there is a non-compulsory option to sign and agree to be contacted RE future related research).
 | Consent form |  |
| 1. Is a clear complaints procedure identified in the information sheet?
 | Participant information sheet |  |
| 1. Is it clear how participants agree to take part?
 | Ethics application, Participant information sheet |  |
| 1. Is it clear what participants are agreeing to when they sign up to the study?
 | Ethics application, Participant information sheet/Consent form |  |
| 1. If participants are vulnerable, is there a process in place to ensure these individuals are safe and looked after?
 | Ethics application, Participant information sheet |  |
| 1. If there are capacity issues, has this been sufficiently taken into account for the consent process?
 | Ethics application/ You decide/seek advice from ethics panel if unsure, Participant information sheet/Consent form |  |
| 1. Are there any rewards/travel expenses for participants taking part?
 | Ethics application, Participant information sheet |  |
| 1. If there are rewards for participation, is this a token gesture or coercion?
 | You decide/seek advice from ethics panel if unsure |  |
| 1. If there is a reward for participation, how do participants receive it?
 | Ethics application, Participant information sheet |  |
| 1. If there is a reward for participation, what do participants have to do to receive it? (i.e. complete the study/show up?)
 | Ethics application, Participant information sheet |  |
| 1. Has it been made clear that participants have the right to withdraw at any time?
 | Ethics application, Participant information sheet/Consent form |  |
| 1. Is it clear how the data will be stored?
 | Ethics application, Participant information sheet |  |
| 1. Is it clear that data will be anonymised?
 | Ethics application, Participant information sheet/Consent form |  |
| 1. Is it clear how data will be anonymised?
 | Ethics application |  |
| 1. Is it clear that all participant information and data will be kept confidential at all times?
 | Ethics application, Participant information sheet/Consent form |  |
| 1. Where applicable, is it clear when confidentiality could be broken and why?
 | Ethics application, Participant information sheet |  |
| 1. Have staff lone working policies been put in place where applicable?
 | Ethics application |  |
| 1. Have measures been put in place to protect participants in the study raises areas of sensitivity?
 | Ethics application, Participant information sheet |  |
| 1. Have participants been informed of any potential risks posed by the study/procedures?
 | Ethics application, Participant information sheet |  |
| 1. If a risk to participants and/or researcher(s) is posed, is it justified?
 | Ethics application, Participant information sheet |  |
| 1. Are audio recorders used during the research?
 | Ethics application, Participant information sheet/Consent form |  |
| 1. Are proper processes in place in regards to what happens to the audio recording, when it should be deleted, and what happens to any transcriptions (i.e. anonymised and how)?
 | Ethics application, Participant information sheet |  |
| 1. Is all information on consent forms, participant information sheets and ethics application consistent?
 | Ethics application/ Participant information sheet/ Consent form |  |
| 1. Is the way the Participant information sheet and consent forms are written/presented appropriate for the population?
 | Participant information sheet/Consent form |  |
| 1. Is there a clear inclusion/exclusion criteria?
 | Ethics application, Participant information sheet |  |
| 1. Is it clear what data will be collected? (e.g questionnaires, audio/video recording etc).
 | Ethics application, Participant information sheet |  |
| 1. Is it clear what will happen to the data throughout the study and at the end of the study?
 | Ethics application, Participant information sheet |  |
| 1. Has a protocol been provided?
 | Protocol (note this is an ADDITIONAL document, not there to inform the ethics application) |  |
| 1. Has it been made clear who the data custodian is?
 | Ethics application |  |
| 1. Is it clear who is involved in the study? i.e. research team, other organisations.
 | Ethics application, Participant information sheet (if other organisations are involved). |  |
| 1. Is it clear who is funding the research?
 | Ethics application, Participant information sheet |  |
| 1. Are refreshments available?
 | Ethics application, Participant information sheet |  |

*Please note the above are generic examples of questions reviewers will ask when reviewing applications. There may be other separate issues that need addressing outside the remit of this document.*