

Participant name: Hospital ID: CHI/Date of Birth:

| Birth: | |
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| SOPHIST - SON Sponsor Chief Investigat IRAS number | tagliflozin in Patients with Heart failure Symptoms and Type 1 Diabetes University of Dundee-NHS Tayside tor Dr Ify Mordi 1007807 | | | | |
|--|--|--|--|--|--|
| Principal Investi | igator | | | | |
| Contact number | r | | | | |
| Contact email | | | | | |
| Visit 6 | | | | | |
| Date of visit: | Participant trial ID | | | | |
| Please tick to indicate the following have been completed: | | | | | |
| Confirmed participant's identity | | | | | |
| Participant has verbally given their consent to continue in the trial | | | | | |
| Concomitant medications have been reviewed | | | | | |
| Adverse events have been reviewed | | | | | |
| Education/review of glucose and ketone management undertaken including | | | | | |
| hypoglycaemic and DKA events, glucose/ketone readings, and insulin doses | | | | | |
| Particip | pant compliance with treatment | | | | |
| | | | | | |

The following must be filed in the participant's medical notes:

- Changes to concomitant medications since last visit
- Any adverse events since last visit
- Details of review of diabetes management including any changes to insulin or other diabetes medication
- Any hypoglycaemic and DKA events since last visit
- Any other notable findings and actions taken
- If the participant was withdrawn from the trial at this visit, document reason

Any further information of note:



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Visit has been carried out as per protocol