ATTACK TRIAL OVERVIEW

DATABASE SEARCH AND INVITATION USING ATTACK TOOLKIT AND DOCMAIL

↓

RESEARCH NURSES PERFORM REMOTE CONSENT CONSULTATIONS

↓

ELIGIBLE PATIENTS PRESCRIBED ASPIRIN (75mg OD) OR USUAL CARE

↓

GP TO COMPLETE EVENT FORMS AND SAE FORMS AS INDICATED

REGULAR AUTOMATED SEARCH OF PRACTICE RECORDS FOR FOLLOW-UP INFORMATION

TRIAL INITIATION

Will be completed by training videos in conjunction with a Study Site File

POST-INITIATION:

- Automated database search for suitable patients:
  - Aged 18+
  - Evidence of CKD (based on blood & urine results)
- GP checks search list produced, removes unsuitable patients and signs and dates each page of the printed list
- Practice sends out invitation letters to suitable patients via Docmail from the ATTACK toolkit

CONSENT CONSULTATIONS

- Consent data entered on ATTACK toolkit / database by consenting nurse

RANDOMISATION

- Eligible participants will be randomised to the trial by the Regional Centre
- GP practices will be notified of the result and will prescribe aspirin (75mg od) to assigned participants
- Practice to add an alert to **all** trial participants' electronic records to notify Practice members of their participation in a clinical trial

PATIENT FOLLOW-UP

- SAE reporting for events not exempted by Protocol (see Fact Sheet 2 'SAE and endpoint event reporting')
- Complete Event Form for any potential endpoint events
- Practice records will be interrogated regularly for follow-up data via the ATTACK toolkit. Please remember to code any bleeding events!
- Copies of the Informed Consent Form and Data Capture Records will be sent to you for your records, and we recommend that these are scanned into the patients' notes.

MONITORING VISITS

- If these occur, the monitor will need access to Practice patient data.