## Protocol Flowchart

## **Research Nurse**

(RN)

- Patients referred from intermediate care, RN can check for potential eligible participants from clinic lists, theatre lists and referral letters.
- ◆ Highlight potential participants to the clinician.
- Patient added to screening log.

## Clinician in T&O

Clinic

- ⇒ Checks for eligibility. If **No**: Informs reason to RN who completes screening Log.
- ⇒ <u>If **Yes**</u>: Informs patient about study, gives patient information and refers patient to RN.
- ◆ Double checks eligibility.
- ♦ Allows patient enough time to read the information sheet and think about the study.
- ♦ Discuss treatment arms, study processes, answers questions, confirms willingness to participate.

## Research Nurse



If unwilling to participate Research Nurse reassures patient that it is OK, completes the <u>Screening log</u> including the reason why.



If patient is willing to participate:

 Collects signed Form. Informed Consent

- ◆ RN completes baseline pack and participant completes baseline questionnaire including the Constant Murley measurement.
- Participant is registered on to the START:REACTS database.
- Screening log is completed with the trial number.
- Intraoperative eligibility checks will be completed.
- If participant eligible continue to randomisation and receive an allocation.
- If patient ineligible the trial office needs to be notified. Log on to the randomisation database at the end of surgery and complete the randomisation form.
- A pre-prepared withdrawal letter will need to be sent to the participant by the RN informing the participant they are no longer taking part in the study.
- ♦ The participant will be followed up at 3,6 and 12 months to complete Constant Murley assessment and questionnaire (these time points are part of usual care).
- Participant will receive a START:REACTS rehabilitation physiotherapy package.
- Participants taking part in the **MRI sub-study** will receive a scan at 6 weeks and 6 months.