

# A prospective randomised trial of the use of tourniquet in total knee replacements

<b>Submission date</b> 28/09/2007	<b>Recruitment status</b> No longer recruiting	<input type="checkbox"/> Prospectively registered <input type="checkbox"/> Protocol
<b>Registration date</b> 28/09/2007	<b>Overall study status</b> Completed	<input type="checkbox"/> Statistical analysis plan <input type="checkbox"/> Results
<b>Last Edited</b> 21/11/2019	<b>Condition category</b> Surgery	<input type="checkbox"/> Individual participant data <input type="checkbox"/> Record updated in last year

**Plain English summary of protocol**  
Not provided at time of registration

## Contact information

**Type(s)**  
Scientific

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## Additional identifiers

**Protocol serial number**  
N0244192302

## Study information

**Scientific Title**  
A prospective randomised trial of the use of tourniquet in total knee replacements

**Study objectives**

To investigate the short-term outcomes of inflating the tourniquet only during the cementation process.

**Ethics approval required**

Old ethics approval format

**Ethics approval(s)**

Not provided at time of registration

**Study design**

Randomised controlled trial

**Primary study design**

Interventional

**Study type(s)**

Treatment

**Health condition(s) or problem(s) studied**

Surgery: Total knee replacement (TKR)

**Interventions**

Intervention: Tourniquet only inflated during cementation of the replacement.

Control: Tourniquet inflated throughout procedure as in routine practice.

**Intervention Type**

Procedure/Surgery

**Phase**

Not Specified

**Primary outcome(s)**

Blood loss and change in haemoglobin level day 2 post-operatively

**Key secondary outcome(s)**

1. Range of movement six weeks post-op
2. Post-operative wound problems
3. Operative time
4. Blood transfusion
5. Complication rate

**Completion date**

01/11/2007

**Eligibility****Key inclusion criteria**

All patients undergoing total knee replacement at Stepping Hill Hospital under two named consultant surgeons

**Participant type(s)**

Patient

**Healthy volunteers allowed**

No

**Age group**

Adult

**Sex**

All

**Key exclusion criteria**

1. History of peripheral vascular disease with absent popliteal or distal pulses
2. Calcification of artery on x-ray of knee
3. Previous vascular surgery on side of operation

**Date of first enrolment**

01/11/2006

**Date of final enrolment**

01/11/2007

**Locations****Countries of recruitment**

United Kingdom

England

**Study participating centre**

Stepping Hill Hospital

Stockport

United Kingdom

SK2 7JE

**Sponsor information****Organisation**

Record Provided by the NHSTCT Register - 2007 Update - Department of Health

**Funder(s)**

**Funder type**

Government

**Funder Name**

Stockport NHS Foundation Trust

## Results and Publications

**Individual participant data (IPD) sharing plan****IPD sharing plan summary**

Not provided at time of registration

**Study outputs**

Output type	Details	Date created	Date added	Peer reviewed?	Patient-facing?
<a href="#">Participant information sheet</a>	Participant information sheet	11/11/2025	11/11/2025	No	Yes