NHS
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Health Research

| Id | Research Ethics Committee Reference Number | Name of Trial | Date of Receipt of Valid Research Application | Date of NHS Permission | First Patient Recruited? | Date of First Patient Recruited | Duration between VRA and NHS Permission | Duration between NHS Permission and First Patient | Duration between VRA and First Patient | Benchmark Met | A - Permissions delayed/denied | B - Suspended by sponsor | C - Closed by sponsor | D - Sponsor Delays |
|---|---|---|---|---|---|---|---|---|---|---|---|---|---|---|
| 69681 | 14/WM/0160 | "Pegasus - An Open Pilot TwoArm<br>Randomised Controlled Trial Comparing Pressure Garment Therapy with No Pressure<br>Garment Therapy for the Prevention of Abnormal Scarring after Burn Injury." | 13/01/2015 | 27/01/2015 | Yes | 30/01/2015 | 14 | 3 | 17 | Yes | | | | |
| 69682 | 13/EM/0118 | "ELFIN - A multicentre<br>randomised placebocontrolled<br>trial of prophylactic enteral lactoferrin supplementation to prevent lateonset invasive infection in very preterm infants." | 13/01/2015 | 13/01/2015 | Yes | 13/01/2015 | 0 | 0 | 0 | Yes | | | | |
| 69683 | 14/SC/1219 | AIRWAYS 2 - Airway Management in cardiac arrest patients | 14/04/2015 | 27/04/2015 | Yes | 13/07/2015 | 13 | 77 | 90 | No | | | | |
| 69684 | 13/LO/1365 | LeoPARDS-Levosimendan for the Prevention of Acute oRgan Dysfunction in Sepsis | 05/05/2015 | 18/05/2015 | Yes | 23/06/2015 | 13 | 36 | 49 | Yes | | | | Y |
| 69685 | 12/EE/0274 | CRASH 3 | 27/03/2015 | 02/04/2015 | Yes | 12/04/2015 | 6 | 10 | 16 | Yes | | | | |
| 69686 | 14/NW/1258 | ARAMIS - Phase 3 efficacy & safety study of ODM201 in high risk nonmet CRPC | 19/05/2015 | 03/06/2015 | No | | 15 | | | | | | | Y |
| 69687 | 14/NE/1176 | UK FROST (United Kingdom Frozen Shoulder Trial) - Multi-centre randomised controlled trial with economic evaluation and nested qualitative study comparing early structured physiotherapy versus manipulation under anaesthesia versus arthoscopic capsular release for patients referred to in secondary care with a frozen shoulder (Adhesive Capsulitis) | 22/07/2015 | 22/07/2015 | Yes | 11/08/2015 | 0 | 20 | 20 | Yes | | | | |
| 69688 | 14/WA/1075 | HeadPoST v2.1 - Head Position in Acute Stroke Trial | 07/09/2015 | 21/09/2015 | Yes | 29/09/2015 | 14 | 8 | 22 | Yes | | | | |
| 69689 | 14/SC/1223 | CALIBER - A phase II randomised feasibility study of chemresection and surgical management in low risk non muscle invasive bladder cancer | 16/09/2015 | 29/09/2015 | No | | 13 | | | | | | | |
| 69690 | 15/EE/0168 | A randomised controlled trial investigating the effectiveness of two splinting methods in the non-operative management of volar plate injuries at the proximal interphalangeal joint | 25/06/2015 | 26/06/2015 | Yes | 14/07/2015 | 1 | 18 | 19 | Yes | | | | |
| 69691 | 15/YH/0134 | R-3D-2 pilot study - Randomised Controlled Trial On the Impact of Surgical Rehearsal Strategies In Rectal Cancer Surgery Using 3D Models by Using 2 Methods | 22/07/2015 | 04/08/2015 | Yes | 21/09/2015 | 13 | 48 | 61 | Yes | | | | |
| 69692 | 14/LO/1539 | CANDLE - Nocturna 400 treatment for Diabetic Retinopathy.  Pilot Study to demonstrate and evaluate the care pathway for NHS adoption | 26/06/2015 | 06/07/2015 | Yes | 14/08/2015 | 10 | 39 | 49 | Yes | | | | |
| 69693 | 13/SW/0132 | MASTER  - Male synthetic sling versus Artificial urinary Sphincter Trial for men with urodynamic stress incontinence after prostate surgery: Evaluation by Randomised controlled trial | 12/10/2015 | 23/10/2015 | Yes | 20/11/2015 | 11 | 28 | 39 | Yes | | | | |
| 69695 | 15/WA/0241 | DEPICT2 - A Multicentre, Randomized, Double-Blind, Placebo-controlled, Parallel Group, Phase 3 Study to Evaluate the Efficacy and Safety of Dapagliflozin as an Add-on to Insulin Therapy in Subjects with Type 1 Diabetes Mellitus - Study 2 | 14/12/2015 | 24/12/2015 | Yes | 20/01/2016 | 10 | 27 | 37 | Yes | | | | |

| E - Staff availability issues | F - No patients seen | G - No patients consented | H - Contracting delays | I - Rare diseases | J - Other | Comments | Reasons for delay correspond to: | Errors |
|---|---|---|---|---|---|---|---|---|
| | | | | | | we are only doing the questionnaire part of the study | Please Select... | |
| | | | | | | We are a continuing care site therefore do not consent patients. | Neither | |
| | | | | Y | | No control over the number of patients admitted. We get recruits for those who live to consent at discharge. All others go to YAS. | Neither | |
| | | | Y | | | Delays in obtaining signed contract from sponsor also in them supplying drugs | Sponsor | |
| | | | | | | | Please Select... | |
| | | | | | | We received a letter from the Sponsor opening recruitment from 11.06.2015. The first patient was recruited within 30 days but due to courier issues at the sponsor site in Ireland the blood samples were damaged and useable. Therefore, the patient became ineligible. | Sponsor | |
| | | | | | | | Please Select... | |
| | | | | | | | Please Select... | |
| | | Y | | | | Patients screened but haven't had a recurrence, therefore ineligible for the study | Neither | |
| | | | | | | | Please Select... | |
| | | | | | | | Please Select... | |
| | | | | | | no patients meeting eligibility criteria were identified | Neither | |
| | | | | | | | Please Select... | |
| | | | | | | | Please Select... | |