

| Research Ethics Committee Reference Number | Integrated Research Application System Number | Name of Trial | Target Number Of Patients Agreed? | Minimum Number Of Patients Agreed | Maximum Number Of Patients Agreed | Target Date To Recruit Patients Agreed? | Date Agreed to recruit target number of patients | Total Number Of Patients Recruited At The Agreed Target Date | Date That The Trial Closed To Recruitment | Total Number Of Study Participants Recruited | Reason For Closure Of Trial |
|--|---|--|-----------------------------------|-----------------------------------|-----------------------------------|---|--|--|---|--|-----------------------------|
| 14/YH/0086 | 147377 | RESPOND: Repositionable Lotus Valve System-Post Market Evaluation of Real World Clinical Outcomes | Range Agreed | 15 | 30 | Date Agreed | 31/12/2015 | 121 | 01/10/2016 | 128 | Recruitment Finished |
| 15/LO/1600 | 182262 | A Phase II, Open Label, Randomized Study of GDC-0810 Versus Fulvestrant in Postmenopausal Women with Advanced or Metastatic ER + / HER 2 - Breast Cancer Resistant to Aromatase Inhibitor Therapy | Number Agreed | 6 | 6 | Date Agreed | 02/03/2018 | 1 | 04/10/2016 | 1 | Withdrawn By Sponsor |
| 16/NI/0034 | 194752 | The Medtronic CoreValve™ Evolut R™ FORWARD Study | Number Agreed | 5 | 5 | Date Agreed | 01/08/2017 | 11 | 01/11/2016 | 11 | Recruitment Finished |
| 16/EE/0098 | 201052 | DIASOLVE: A randomised, crossover investigation to evaluate and compare the effectiveness, safety and feasibility of a novel dedicated Over-The-Wire FFR Infusion Microcatheter (HYPEREM™IC) for measuring fractional flow reserve (FFR) using intra-coron | Number Agreed | 5 | 5 | Date Agreed | 30/11/2016 | 7 | 10/11/2016 | 7 | Recruitment Finished |
| 13/NW/0002 | 114402 | Prospective, randomised, controlled investigation comparing the safety and performance of 032-11 Surgical Haemostat 2g Applicator with FLOSEAL? Haemostatic Matrix as an adjunctive haemostat in cardiac surgery and thoracic aortic surgery | Range Agreed | 10 | 32 | Date Agreed | 28/02/2014 | 5 | 01/12/2016 | 32 | Recruitment Finished |
| 13/NI/0138 | 120046 | Portico I Study International long-term follow-up study of patients | Number Agreed | 10 | 10 | Date Agreed | 30/06/2018 | 10 | 01/12/2016 | 10 | Recruitment Finished |
| 13/LO/0821 | 128690 | A Phase 3 Open Label Safety Study of Elvitegravir/Cobicistat/Emtricitabine/Tenofovir Alafenamide Single Tablet Regimen in HIV1 Positive Patients with Mild to Moderate Renal Impairment | Number Agreed | 5 | 5 | Not Available / Not Agreed | | | 30/11/2016 | 1 | Recruitment Finished |
| 14/SC/0225 | 147356 | A Phase 3, Randomized, Double-Blind, Switch Study to Evaluate F/TAF in HIV 1 Positive Subjects who are Virologically Suppressed on Regimens containing FTC/TDF | Number Agreed | 8 | 8 | Not Available / Not Agreed | | | 21/11/2016 | 6 | Recruitment Finished |
| 14/YH/1269 | 140294 | A Phase 3, OpenLabel, Randomised, Parallel, 2Arm, MultiCenter Study of BMN 673 versus Physician's Choice in Germline BRCA Mutation Subjects with Locally Advanced and/or Metastatic Breast Cancer, Who Have Received No More than 2 Prior Chemotherapy Reg | Range Agreed | 1 | 2 | Not Available / Not Agreed | | | 20/04/2017 | 2 | Recruitment Finished |
| 14/YH/0088 | 151005 | Phase I study of KHK2823 in Patients with Acute Myeloid Leukaemia or Myelodysplastic Syndrome | Range Agreed | 4 | 8 | Date Agreed | 01/02/2017 | 4 | 30/06/2017 | 4 | Recruitment Finished |
| 15/NW/0543 | 181870 | A Multicentre, Randomised, Doubleblind, Placebo controlled, Phase 3 Study Evaluating the Efficacy and Safety of Two Doses of Anifrolumab in Adult Subjects with Active Systemic Lupus Erythematosus | Range Agreed | 2 | 3 | Date Agreed | 30/04/2017 | 0 | 30/06/2017 | 0 | Recruitment Finished |
| 16/LO/0586 | 200579 | An Open-Label Study to Evaluate the Efficacy and Safety of Ocrelizumab in Patients with Relapsing Remitting Multiple Sclerosis who have a Suboptimal Response to an Adequate Course of Disease-Modifying Treatment | Number Agreed | 4 | 4 | Date Agreed | 31/07/2017 | 0 | 03/03/2017 | 0 | Recruitment Finished |

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| 15/YH/0478 | 186697 | A Phase 3, Double-Blind, Randomized, Long-Term, Placebo-Controlled, Multicenter Study Evaluating the Safety and Efficacy of Obeticholic Acid in Subjects with Nonalcoholic Steatohepatitis | Range Agreed | 5 | 10 | Date Agreed | 31/10/2016 | 0 | 18/08/2017 | 0 | Withdrawn By Host |
| 16/EM/0386 | 211113 | A 12-week double-blind, randomized, multicenter study comparing the efficacy and safety of once monthly subcutaneous 140 mg AMG 334 against placebo in adult episodic migraine patients who have failed 2-4 prophylactic treatments | Number Agreed | 4 | 4 | Date Agreed | 22/08/2016 | 2 | 03/05/2017 | 2 | Recruitment Finished |