

| Research Ethics Committee Reference Number | Integrated Research Application System Number | Name of Trial  | Target Number Of Patients Agreed? | Minimum Number Of Patients Agreed (Enter Same In Both If Only One Number) | Maximum Number Of Patients Agreed (Enter Same In Both If Only One Number) | 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 |
|--|---|--|-----------------------------------|---|---|---|--|--|---|--|-----------------------------|
| 17/LO/0825                                 | 223457  | Randomised, Phase 2 , double blind, placebo controlled study to assess the safety and efficacy of Filgotinib, GS-9876 and GS-4059 in adult subjects with Active Sjogren's syndrome | Number Agreed                     | 2   | 2   | Date Agreed                             | 22/04/2018                                       | 0  | 05/09/2018                                | 0  | Recruitment Finished        |
| 14/SC/1161                                 | 155743  | E-Ultimaster registry  | Number Agreed                     | 30  | 30  | Date Agreed                             | 01/12/2015                                       | 250  | 02/05/2018                                | 391  | Recruitment Finished        |
| 15/LO/0769                                 | 155035  | Left Atrial Appendage Occlusion Study II (LAAOS III)   | Range Agreed                      | 2   | 50  | Date Agreed                             | 01/06/2018                                       | 13   | 08/06/2018                                | 13   | Recruitment Finished        |
| 17/LO/0736                                 | 225746  | CN2002012  | Number Agreed                     | 5   | 5   | Date Agreed                             | 27/07/2018                                       | 1  | 27/07/2018                                | 1  | Recruitment Finished        |
| 17/EM/0361                                 | 234065  | 215MS202 Efficacy & Safety of BIIB033 as an Add-on Therapy in RMS  | Number Agreed                     | 2   | 2   | Not Available / Not Agreed              |  |  | 17/08/2018                                | 0  | Recruitment Finished        |
| 18/WM/0017                                 | 236521  | Post-Market Clinical Investigation of the Clareon® IOL   | Number Agreed                     | 10  | 10  | Not Available / Not Agreed              |  |  | 28/09/2018                                | 13   | Recruitment Finished        |
| 17/EM/0281                                 | 229054  | 204862 Phase 3 switch study -TAF regimen to DTG + 3TC in HIV-1 adults  | Range Agreed                      | 5   | 10  | Not Available / Not Agreed              |  |  | 10/05/2018                                | 1  | Recruitment Finished        |
| 16/SC/0416                                 | 210405  | D5290C00003 - MEDI8897 Against RSV in Healthy Preterm Infants  | Range Agreed                      | 2   | 5   | Date Agreed                             | 30/11/2017                                       | 5  | 30/11/2018                                | 5  | Recruitment Finished        |

|            |        |   |               |   |   |             |            |   |            |   |                      |
|------------|--------|---|---------------|---|---|-------------|------------|---|------------|---|----------------------|
| 17/EE/0431 | 234555 | A Phase 2B, Randomized, Double-Blind, Active-Comparator-Controlled, Dose- Ranging Clinical Trial to Evaluate the Safety, Tolerability, Antiretroviral Activity, and Pharmacokinetics of MK-8591 Given in Combination with Doravirine (DOR) and Lamivudine | Number Agreed | 4 | 4 | Date Agreed | 31/03/2018 | 1 | 20/04/2018 | 1 | Recruitment Finished |
| 18/NE/0132 | 242937 | "A PHASE III, RANDOMIZED, MULTICENTER, OPEN-LABEL, TWO-ARM STUDY TO EVALUATE THE PHARMACOKINETICS, EFFICACY, AND SAFETY OF SUBCUTANEOUS ADMINISTRATION OF THE FIXED-DOSE COMBINATION OF PERTUZUMAB AND TRASTUZUMAB & CHEMOTHERAPY IN PATIENTS WITH HER2 P | Range Agreed  | 4 | 8 | Date Agreed | 01/07/2019 | 5 | 24/12/2018 | 5 | Recruitment Finished |
| 17/YH/0228 | 222492 | CALM- 2 – CONTROLLING AND LOWERING BLOOD PRESSURE WITH THE MOBIUSHD®  | Number Agreed | 2 | 2 | Date Agreed | 31/05/2019 | 0 | 19/02/2019 | 0 | Withdrawn By Host    |