

Performance in Delivering Qtr 2 2021-22

| Id     | 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 | Comments                              |
|--------|--|---|--|-----------------------------------|---|---|---|--|--|---|--|-----------------------------|---------------------------------------|
| 55180  | 20/EM/0049                                 | 273949  | M19-944: A Phase 3 Randomized, Placebo-Controlled, Double-Blind Program to Evaluate Efficacy and Safety of Upadacitinib in Adult Subjects with Axial Spondyloarthritis                                       | Number Agreed                     | 2   | 2   | Date Agreed                             | 20/05/2021                                       | 2  | 20/05/2021                                | 0  | Recruitment Finished        | No eligible patients                  |
| < Back |  |   |  |                                   |   |   |   |  |  |   |  |                             |                                       |
| 55181  | 19/ES/0100                                 | 261027  | A Phase 3, Double-Blind, Multicenter Study to Evaluate the Long-Term Safety and Efficacy of Baricitinib in Patients with Systemic Lupus Erythematosus (SLE)  | Number Agreed                     | 0   | 2   | Date Agreed                             | 31/01/2021                                       | 2  | 05/01/2021                                | 2  | Recruitment Finished        |                                       |
| 55182  | 18/NE/0023                                 | 230930  | Randomised, double-blind, placebo controlled multi-centre study to assess the efficacy, tolerability and safety of Enterosgel® in the treatment of Irritable Bowel Syndrome with Diarrhoea (IBS-D) in adults | Number Agreed                     | 5   | 5   | Date Agreed                             | 31/03/2021                                       | 4  | 21/04/2021                                | 4  | Recruitment Finished        | A number of patients failed screening |