

Clinical Trials Potentially Impacted by COVID-19

| Company | Drug | Indication | Clinical trial | Anticipated timings (pre COVID-19) | Status/guidance | Modality | COVID-19 risk factors |
|-----------|------------|-------------------------------|------------------------------|--|---|-----------|--|
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| Galapagos | Filgotinib | Ulcerative colitis | Phase III (SELECTION) | Headline data 2Q20E | Fully enrolled, read-out unaffected | Oral (QD) | |
| | | Crohn's disease | Phase III (DIVERSITY) | Headline data 2Q21E | Enrolment paused due to COVID-19 | Oral (QD) | Co-primary endpoint requires endoscopy |
| | | Psoriatic arthritis | Phase III (PENGUIN) | Headline data 1H22E | Enrolment paused due to COVID-19 | Oral (QD) | |
| | | Ankylosing spondylitis | Phase III | Expected to start 1Q20E (JEFe) | Initiation now expected 2H20E due to COVID-19 | Oral (QD) | |
| | | Testicular safety | Phase II (MANTA & MANTA-RAY) | Safety data 2Q20E (JEFe) | Enrolment paused due to COVID-19 | Oral (QD) | |
| | | Uveitis | Phase II (Humboldt) | Headline data 1H21E | Enrolment paused due to COVID-19 | Oral (QD) | |
| | GLPG1690 | Idiopathic pulmonary fibrosis | Phase III (ISABELA) | Interim futility analysis 1Q21E, headline data 1H22E | >50% patient enrolled sufficient for interim analysis; full enrolment was expected YE20 | Oral (QD) | High risk patient population, endpoints require monthly follow ups |
| | | Systemic sclerosis | Phase IIa (NOVESA) | Headline data 3Q20E | Fully enrolled Dec 2019 | Oral (QD) | |
| | GLPG1972 | Osteoarthritis | Phase IIa (ROCCELLA) | Headline data 4Q20E | Fully enrolled Jun 2019 | Oral (QD) | Primary endpoint requires an MRI |
| | GLPG1205 | Idiopathic pulmonary fibrosis | Phase IIa (PINTA) | Headline data 3Q20E | Fully enrolled early-2020 | Oral (QD) | High risk patient population |

Source: Company data; Jefferies research. Notes: QD, once a day; BID, twice a day; QW, once a week; LPLV, last patient last visit