

Participant identifier

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Participant initials

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Eligibility CRF

Date eligibility completed: _____	
Participant NHS number: _____	
1.	Please select the participant's age range: - Less than 18 (Ineligible) 18—49 50—64 65+ 18—49—Eligible for UC or COLCH 50+ —Eligible for UC, BUDE or COLCH
2.	Does the participant have at least one of the comorbidities/conditions listed below? - Weakened immune system due to a serious illness or medication (e.g. chemotherapy) - Heart disease or high blood pressure - Asthma or lung disease - Known diabetes - Liver disease - Stroke or neurological problem If Yes and age range = 18-49—Eligible for COLCH If Yes and age range = 50-64—Eligible for COLCH and BUDE
2a)	If No, does the participant have a body mass index (BMI) at or above 35 kg/m²?
2b)	If No, is the participant experiencing shortness of breath as part of a Covid-19 illness? Shortness of breath can make it hard to breathe deeply and you may feel winded or as if you can't get enough air into your lungs. Unlike many other conditions that cause shortness of breath, this symptom can persist and quickly escalate in people with COVID-19. If Yes—Eligible for COLCH If No—Ineligible
3.	Is the participant able to use an inhaler?
4.	Is the participant currently admitted to hospital?
5.	Has the participant previously participated in the PRINCIPLE trial?

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6.	Does the participant have a known allergy to inhaled corticosteroids?	Yes ☐ No ☐
7.	Is the participant currently prescribed corticosteroids by inhaler, by mouth or had a corticosteroid injection in the past month?	Yes ☐ No ☐
8.	Does the participant have any of the following conditions: • Galactose intolerance • Lapp lactase deficiency • Glucose-galactose malabsorption	Yes ☐ No ☐
9.	Is the participant pregnant or planning on becoming pregnant within the next few weeks?	Yes ☐ No ☐
10.	Is the participant breastfeeding or planning on starting during the course of the trial?	Yes ☐ No ☐
11.	Is the participant a pre-menopausal female that is capable of becoming pregnant?	Yes ☐ No ☐
11 a)	If yes, is the participant using a highly effective method of contraception* and prepared to use this for the 28 days of the study? * The implant, the coil, and male or female sterilisation will be acceptable for participating in the trial. The injection and most forms of hormonal contraception will also be considered acceptable for the trial, if used in combination with condoms or other barrier methods. However, condoms alone won't be sufficient during the study.	Yes ☐ No ☐
12.	Is the participant hypersensitive/allergic to any of the following: • Colchicine • Lactose • Pregelatinised Maize Starch • Stearic Acid • Purified Water • Ethanol (96%)	Yes ☐ No ☐

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13.	Does the participant have a clinically significant blood dyscrasia?	Yes ☐ No ☐
14.	Does the participant have a known severe renal impairment or require dialysis?	Yes ☐ No ☐
15.	Does the participant have severe liver disease?	Yes ☐ No ☐
16.	Is the participant currently taking any of the following drugs: • Clarithromycin • Cobicistat • Colchicine • Cyclosporin • Diltiazem • Disulfiram • Erythromycin • HIV protease inhibitors (e.g. ritonavir, atazanavir) • Itraconazole • Ketoconazole • Quinidine • Verapamil • Voriconazole	Yes ☐ No ☐
17.	Does the participant have inflammatory bowel disease or chronic diarrhoea?	Yes ☐ No ☐

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18.	Is there any other reason you would exclude this participant? If yes, please provide your reason in the comments box below.	Yes ☐ No ☐
Comments: 		

KEY

Exclusion for all arms

Exclusion criteria from inhaled corticosteroids

Exclusion criteria from colchicine

NOTES

Q 1 – 5 = Inclusion criteria

Q5 onwards = Exclusion criteria

Completed by: Print name Sign Date __ / __ / __

Reviewed by: Print name Sign Date __ / __ / __