A Double Blind Randomised Study Comparing The Skin

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A Double Blind Randomised Study
Methods The study was a randomized, double blind, placebo controlled study conducted on 2 parallel arms. The experimental products were dry cat food with 1% scFOS (Profeed®, TEREOS) or without scFOS.

A Randomised Double-Blind, Placebo Controlled Study Evaluating the Effects of Short-Chain Fructo-Oligosaccharides (scFOS) on Cat Stools Odors
Zotatifin acts on a human protein vital for viral replication. Credit: Viktor Forgacs / Unsplash. eFFECTOR Therapeutics has dosed the first participant in the Phase Ib clinical trial of zotatifin ...

eFFECTOR begins dosing in Phase Ib trial of zotatifin in Covid-19
Hufford, Maja Hojnik, Philip J. Mease, Arthur Kavanaugh, the SPIRIT-P3 Study Group SPIRIT-P3 was a multicenter, randomized, double-blind withdrawal study that enrolled biologic-naive adult ...

Withdrawing Ixekizumab in Patients With Psoriatic Arthritis Who Achieved Minimal Disease Activity: A Randomized, Double-Blind Withdrawal Study
pylori. The aim of this study was to compare the eradication efficacy and tolerability profile of a 7-day course of RBC plus clarithromycin and amoxicillin, RBC plus clarithromycin, and omeprazole ...

Efficacy and Tolerability of Three Regimens for Helicobacter pylori Eradication A Multicentre, Double-Blind, Randomised Clinical Trial
Acticor Biotech, a clinical stage biotechnology company developing an innovative drug for the acute phase of ischemic stroke and thrombotic diseases, ...

Acticor Biotech Announces the Completion of Enrollment in its GARDEN Clinical Trial, a COVID-19-induced Acute Respiratory Distress Syndrome Efficacy Study
Methods: Multicentre, randomised, double-blind, parallel-group clinical trial in 64 outpatients with GAD (DSM-IV criteria). Patients were assigned to mexazolam 1mg three times daily (n = 32 ...

Mexazolam and Alprazolam in the Treatment of Generalised Anxiety Disorder: A Double-Blind, Randomised Clinical Trial
Methods 230 adults starting Achilles rupture non-surgical management within 12 days of injury were randomised to PRP injection or dry needle insertion to the rupture gap, under local anaesthetic.

3 Platelet rich plasma for acute achilles tendon rupture: a double-blind, multicentre, randomised, placebo-controlled trial
Dicerna announced interim results from its Phase 1 trial of belcesiran for treatment of alpha-1 antitrypsin deficiency-associated liver disease.

Dicerna Announces Interim Results From Phase 1 Trial of Belcesiran for Treatment of Alpha-1 Antitrypsin Deficiency-Associated Liver Disease
Merck (MSD) has reported data from a Phase IIa trial that showed an encouraging safety and tolerability profile of oral medication islatravir for pre-exposure prophylaxis (PrEP) of human ...

Merck’s islatravir shows favourable safety profile in Phase IIa HIV trial
LONGJUMEAU, France, June 8, 2021 /PRNewswire/ -- The French biotech TargEDys specialized in breakthrough microbiome-based solutions announces the publication of its study conducted in 230 ...

TargEDys announces the publication of the first double-blind, randomized, placebo-controlled clinical trial showing Hafnia al...
InDex Pharmaceuticals Holding AB (publ) today announced that the Swedish Medical Products Agency (MPA) has given approval to start the phase III clinical study CONCLUDE in Sweden. The study will ...

InDex Pharmaceuticals receives first regulatory approval to start the phase III study CONCLUDE with cobitolimod
Humanigen, Inc. (Nasdaq: HGEN) ("Humanigen"), today announced its development and commercialization partners for South Korea and the Philippines, Telcon RF Pharmaceutical, Inc. ("Telcon") and KPM Tech ...

Humanigen’s Partner in South Korea Receives Ministry of Food and Drug Safety (MFDS) Approval to Conduct Phase 1 Study of Lenzilumab
Bio-Thera Solutions, Ltd. (688177.SH), a commercial-stage pharmaceutical company, today announced that the first patient has been dosed in a Phase III ...

Bio-Thera Solutions Announces the First Patient Dosed in Phase III Clinical Trial for BAT2206, a Proposed Biosimilar of Stelara® (Ustekinumab)
HANGZHOU, China and SHAOXING, China, July 22, 2021 /PRNewswire/ -- Ascletis Pharma Inc. (HKEX code: 1672) today announces that China National ...

China NMPA Approves Phase III Clinical Trial of ASC40 Combined with Bevacizumab for Treatment of Patients with Recurrent Glioblastoma
CET; regulated information - Galapagos NV (Euronext & Nasdaq: GLPG) reports positive topline results with tyrosine kinase 2 (TYK2) inhibitor GLPG3667 in a Phase 1b study in psoriasis patients.

Galapagos reports positive topline results with selective TYK2 inhibitor GLPG3667 in Phase 1b psoriasis study
Shares of Cytokinetics Inc. soared 33.0% in premarket trading on Monday after the company said a Phase 2 clinical trial for its experimental hypertrophic cardiomyopathy treatment yielded positive ...

Cytokinetics shares jump based on results from Phase 2 clinical trial for experimental heart-disease drug
Madrigal Pharmaceuticals, Inc. (NASDAQ:MDGL) announced today its first patient dosed in a planned 52-week open label ...

Madrigal Pharmaceuticals Offers Patients Resmetirom in a Planned Open Label Active Treatment Extension of the Phase 3 MAESTRO-NAFLD-1 Clinical Study
SPIRIT-P3 was a multicenter, randomized, double-blind withdrawal study that enrolled biologic-naive adult patients with PsA to open-label ixekizumab (160 mg at week 0, 80 mg every two weeks [IXE ...

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