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Evaluating The Treatment Of Drug

Apremilast, an oral drug which inhibits the activity of PDE4 (Phosphodiesterase 4),

was selected for evaluation because it was hypothesized ... to incorporate additional potential treatment regimens ...

Global Coalition for Adaptive Research Announces the Discontinued Evaluation of Apremilast in REMAP-COVID

The US Food and Drug Administration (FDA) has approved drug Actemra (tocilizumab), used for arthritis, for the treatment of hospitalized ... FDA's Center for Drug Evaluation and Research.

Arthritis Drug for Treatment of COVID-19

In clinical trials of hospitalized patients with COVID-19, Actemra in addition to the routine care patients receive for treatment of ... the FDA's Center for Drug Evaluation and Research.

Coronavirus (COVID-19) Update: FDA Authorizes Drug for Treatment of COVID-19

She said they're working to put in place systems to collect data on participant success that will allow the court to evaluate its programs and treatment providers. And, she said, the drug ...

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The studies will use Praedicare's capability and expertise with the hollow fiber

system model of TB in combination with its clinical mapping algorithms to evaluate new drug combinations for ...

Praedicare Inc. Receives Grant to Evaluate Potential New Drug Regimens for Tuberculosis

Milland Dixon Dikio, retd, has said the scheme is planning to begin free medical evaluation of ex-agiators to determine the impact of drug abuse on their lives. Dikio, who spoke at the Bayelsa ...

Drug abuse: PAP to begin free medical evaluation of ex-agiators

for the drug Actemra (tocilizumab) -- used for arthritis -- for the treatment of hospitalised patients with Covid-19. Under the EUA, the drug can be administered only ot hospitalised adults and ...

US FDA okays arthritis drug for treatment of Covid-19

Concept Medical Inc., focused on vascular intervention drug delivery devices, has updated the progress of IMPRESSION (sIroliMus coated balloon angioPlasty versus plain balloon angioplasty in the ...

IMPRESSION – A randomised trial to evaluate the efficacy of MagicTouch Sirolimus Coated Balloon in Dysfunctional Fistula progresses rapidly.

June 15, 2021 /PRNewswire/ -- Today, the U.S. Food and Drug Administration

approved StrataGraft for the treatment of adult ... of FDA's Center for Biologics Evaluation and Research. " ...

FDA Approves StrataGraft for the Treatment of Adults with Thermal Burns

JB Chemicals & Pharmaceuticals (JBCPL) backed by US private equity giant KKR is evaluating assets that include ... in the Mumbai-based family-owned drug maker JBCPL a year ago for Rs 3,100 crore.

Exclusive: KKR-controlled drug maker JBCPL evaluating acquistion of select brands, mid-sized firm

Zydus Cadila on Monday said it has received tentative approval from the US health regulator to market epilepsy treatment drug Brivaracetam ... clinical trial for evaluation of COVID-19 therapy ...

Zydus Cadila gets USFDA tentative nod for epilepsy treatment drug

Singapore, June 23 (ANI/PRNewswire): Concept Medical Inc., focused on vascular intervention drug delivery devices ... plain balloon angioplasty in the tREatment of dialySis acceSs dysfunctION ...

IMPRESSION - A randomised trial to evaluate the efficacy of MagicTouch Sirolimus Coated Balloon in Dysfunctional Fistula progresses rapidly

The US Food and Drug Administration (FDA) has issued an emergency use

authorization (EUA) for the drug Actemra (tocilizumab) -- used for arthritis -- for the treatment of hospitalised patients ...

US FDA okays arthritis drug for treatment of Covid-19

The MagicTouch AVF Sirolimus Coated Balloon has already been granted the Breakthrough Device Designation by the U.S. Food and Drug Administration ... current standard treatment for patients ...