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CLINICAL PHARMACOLOGY AND THE GUT: PROCEEDINGS OF THE ...

Abstract. The complexity of integrating microbiota into clinical pharmacology, environmental toxicology, and opioid studies arises from bidirectional and multiscale interactions between humans and their many microbiota, notably those of the gut. Hosts and each microbiota are governed by distinct central dogmas, with genetics influencing transcriptomics, proteomics, and metabolomics.

The Microbiome and the Gut-Liver-Brain Axis for Central ...

The gut microbiota is of particular interest in pharmacology due to the manner in which gastrointestinal (GI) microbes are associated with a multitude of diseases 11, 12 and interact with xenobiotics. 13, 14 Drug designers may need to consider the high spatial heterogeneity and complexity of the gut microbiota: going from stomach to large intestine, there are gradients in pH, pressure, and the ...

The Microbiome and the Gut-Liver-Brain Axis for Central ...

Morphine and butorphanol also gave relief from visceral pain in the cecal distention model. Morphine may inhibit colonic, and butorphanol jejunal, motility. Whether xylazine or opiate mediated decreases in gut motility cause clinically important slowing of ingesta transit is controversial and requires further investigation.

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Selected Aspects of the Clinical Pharmacology of Visceral ...

The complexity of integrating microbiota into clinical pharmacology, environmental toxicology, and opioid studies arises from bidirectional and multiscale interactions between humans and their many microbiota, notably those of the gut. Hosts and each microbiota are governed by distinct central dogmas, with genetics influencing

The Microbiome and the Gut-Liver-Brain Axis for Central ...

Abstract This article reviews the pharmacology and physiology of opiate receptors and the current and potential uses of opioid agonists and antagonists in clinical gastroenterology. μ -receptors are involved in motor and sensory functions, and their modulation is established for treatment of diarrhea. κ -antagonists have potential to reverse endogenous (e.g., postoperative ileus) or ...

Opioids and the gut: pharmacology and current clinical ...

The complexity of integrating microbiota into clinical pharmacology, environmental toxicology, and opioid studies arises from bidirectional and multiscale interactions between humans and their many microbiota, notably those of the gut. Hosts and each microbiota are governed by distinct central dogma

The Microbiome and the Gut-Liver-Brain Axis for Central ...

The major effects of stress on gut physiology include: 1) alterations in gastrointestinal motility; 2) increase in visceral perception; 3) changes in gastrointestinal secretion; 4) increase in

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intestinal permeability; 5) negative effects on regenerative capacity of gastrointestinal mucosa and mucosal blood flow; and 6) negative effects on intestinal microbiota.

Stress and the gut: pathophysiology, clinical consequences ...

Clinical pharmacology encompasses all aspects of the relationship between drugs and humans. It is the only medical specialty in the NHS focusing on the safe, effective and economic use of medicines. It is a diverse discipline that both sustains and advances best healthcare.

What is clinical pharmacology | British Pharmacological ...

The field of clinical pharmacology is the study of all aspects of drugs as they relate to humans. St George's Clinical Pharmacology BSc is designed to provide you with a broad understanding of how drugs are developed, from discovery of molecules to treatment of patients.

Clinical Pharmacology - St George's, University of London

Gastrointestinal Pharmacology and Therapeutics consists of 59 chapters divided into 15 sections on all aspects of gastrointestinal pharmacology. In fact, there is considerable emphasis on gastrointestinal pathophysiology so the book could easily be called Gastrointestinal Pathophysiology, Pharmacology and Therapeutics.

Gastrointestinal pharmacology and therapeutics | Gut

In this article, we summarize recent findings, review the molecular mechanisms and the

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potential of polyphenols as dietary supplements for regulating gut microbiota and circadian rhythms, and discuss future research directions. Linked Articles. This article is part of a themed section on The Pharmacology of Nutraceuticals.

The roles of gut microbiota and circadian rhythm in the ...

The physicochemical property and permeability (typically obtained using Caco-2 system) data is the first necessity to predict the extent of absorption from the gut lumen to the intestinal epithelium (F a). Intrinsic clearance measured using the human microsome or hepatocytes is also needed to predict the gut (F g) and hepatic (F h) bioavailability. However, there are many issues with the correction of the inter-laboratory variability, hepatic cell membrane permeability, CYP3A4 dependency, etc.

:: TCP :: Translational and Clinical Pharmacology

Piet H. Graaf, The Role of the Microbiome in Central Nervous System Clinical Pharmacology: More Than a Gut Feeling, *Clinical Pharmacology & Therapeutics*, 10.1002/cpt.2036, 108, 5, (907-909), (2020). Wiley Online Library

The Promise and the Reality of ... - Wiley Online Library

The gut microbiome closely interacts with the host, and it has a major influence on drug response. Many studies have reported the possible microbial influences on drugs and the possible influences of drugs on the microbiome. This knowledge has led to a better understanding of intra- and inter-individual variabilities in clinical pharmacology.

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Abstract. The effect of intravenous and oral hyoscine butylbromide (HBB) on gut motility has been assessed by tape recording bowel sounds in 7 healthy volunteers, with the use of a double-blind procedure. In 6 of these intravenous administration of 8 mg. of HBB reduced the bowel sounds for 15 to 30 minutes; this was followed by a considerable increase in bowel sounds.

Effects of hyoscine butylbromide on gut motility ...

Gut bacterial communities directly metabolize certain drugs, reducing their bioavailability and influencing individual variation in drug response. In addition, some microbiome-produced compounds may affect drug pharmacokinetics and pharmacodynamics via altered expression of metabolizing enzymes and drug transporters or genes coding for drug target proteins, drug response phenotypes, and disease states.

ACCP Journals - American College of Clinical Pharmacology

Clinical Pharmacology and Quantitative Pharmacology, Clinical Pharmacology and Safety Sciences, R&D, at AstraZeneca – Much more research is needed, but we hope that our new findings may eventually provide a new way to repair neuronal deficits that are linked to many gut diseases.

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