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*Absorption and
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Metabolism And

The colon is believed to be a suitable absorption site for peptides and protein drugs for the following reasons; (i) less diversity, and intensity of digestive Colon targeted drug delivery system increases the absorption of poorly absorbable drug due to high retention time of the

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colon.enzymes, (ii)
comparative proteolytic
activity of colon mucosa
is much less than that
observed in the small
intestine, thus CDDS
protects peptide drugs
from hydrolysis, and
enzymatic degradation
in duodenum ...

*COLON TARGETED
DRUG DELIVERY
SYSTEM: A REVEIW*

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ON PRIMARY...

1293 drug absorption from colon the colonic absorption of drugs can differ significantly from absorption in the small intestine as a

consequence of several physiological physicochemical and biopharmaceutical factors the permeability through passive transport is thought to

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be lower in colonic than
in other intestinal tissue
because of the smaller
surface area and tighter
junctions in the

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The synergistic action of
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intestinal drug metabolism and transport is also discussed. Despite the complicated regulatory factors, the biopharmaceutics drug disposition classification system (BDDCS) put forward by Wu and Benet may help us better predict the effect of transporters on drug absorption.

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Absorption And
*Intestinal Transporter-
Associated Drug
Absorption and ...*

Similar to gastric
absorption, passive
intestinal absorption can
be modulated by
luminal pH, intestinal
motility, mucin
production, as well as
surface area or
absorptive capacity. A
unique consideration in

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the neonate (term or preterm) is that the colon can be a site of significant absorption of nutrients and pharmaceuticals, whereas in older children and adults, the colon exhibits less absorptive capacity.

*Physiology of the
Neonatal
Gastrointestinal System*
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The small intestine plays an important role in the absorption and metabolism of oral drugs. In the current evaluation system, it is difficult to predict the precise absorption and metabolism of oral drugs. In this study, we generated small intestinal epithelial-like cells from human

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Functional Comparison
of Human Colonic
Carcinoma Cell Lines
and Primary Small
Intestinal Epithelial
Cells for Investigations
of Intestinal Drug
Permeability and First-
Pass Metabolism. To

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further the development of a model for simultaneously assessing intestinal absorption and first-pass metabolism in vitro, Caco-2, LS180, T84, and fetal human small intestinal epithelial cells (fSIECs) were cultured on permeable inserts, and the integrity of cell monolayers, CYP3A4 activity, and t

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1. Drug Metab Dispos.
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transporters and
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