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The colon is believed to be
a suitable absorption site
for peptides and protein

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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 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 ON PRIMARY*

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

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in other intestinal tissue
because of the smaller
surface area and tighter
junctions in the

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The synergistic action of
intestinal drug metabolism
and transport is also
discussed. Despite the
complicated regulatory
factors, the
biopharmaceutics drug

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disposition classification
system (BDDCS) put forward
by Wu and Benet may help us
better predict the effect of
transporters on drug
absorption.

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 the neonate
(term or preterm) is that
the colon can be a site of
significant absorption of

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pharmaceuticals, whereas in
older children and adults,
the colon exhibits less
absorptive capacity.

*Physiology of the Neonatal
Gastrointestinal System ...*
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
induced pluripot ...

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Functional Comparison of
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Lines and Primary Small
Intestinal Epithelial Cells
for Investigations of
Intestinal Drug Permeability
and First-Pass Metabolism.
To further the development
of a model for

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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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enzymes (for the Gillette
Review Series). Pang KS(1).

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