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soluble to achieve

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the majority of drug

modification and

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strategies focus on

improving solubility.

The higher the

concentration of

drug in solution, and

the longer the drug

stays in solution as

it travels through

the GIT, the higher

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Solubility and bioavailability are the overwhelming challenges in drug development. We

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have a range of practical, proven approaches for overcoming these barriers.

Approximately 40% of all drug s currently on the market, and 90% of the compounds at present in development for future use, are reported to show

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covers all aspects of

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**focus is placed on
methods for**

determining the

**parameters relevant
to bioavailability.**

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First, several BCS

II/IV drugs, such as

naproxen,

phenytoin, and

diazepam, have an

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absolute bioavailability (F) > 90%. 6, 7 Second, although poorly water?soluble, lipophilic compounds are generally expected to show a better solubilization in gastrointestinal fluids in the presence of food and, thus, a better

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oral absorption in
the fed state, 9
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phenomenon of dissolution of solute in solvent to give a homogenous system, is one of the important parameters to achieve desired concentration of drug in systemic circulation for desired (anticipated) pharmacological response. Low

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aqueous solubility is the major problem encountered with formulation development of new chemical entities as well as for the generic development.

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Below are some common techniques

for overcoming

limited drug

solubility and

producing a finished

product that has

high bioavailability

and therapeutic

effect: Amorphous

solids – Amorphous

solid dispersions

utilize techniques

such as hot melt

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extrusion or spray
drying to
incorporate an API
into a polymer
matrix and maintain
it in an amorphous
form.

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...

**PEC is a new
generation of**

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phosphamide ester
anti-hepatitis B virus

drug. It is a prodrug

of tenofovir and can

be rapidly

metabolized to

tenofovir. However,

its poor solubility in

water (0.219 mg

mL⁻¹ at 25 °C) has

limited its oral

bioavailability. In

this study, we aimed

to improve the

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**mechanisms. The
much revised and
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surveys current in
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properties needed to
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**bioavailability of any
new drug candidate.**

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Drug developers
from industry and

academia present all

the factors

governing drug

bioavailability,

complete with

practical examples

and real-life data.

Part I focuses on

solubility and

gastrointestinal

absorption, while

the second

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discusses in vitro and in vivo measurements of physicochemical properties, such as membrane permeability and solubility.

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