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pharmacokinetics.  
Pharmacokinetics can  
be simply described as  
the study of 'what the  
body does to the drug'  
and includes: • the rate  
and extent to which  
drugs are absorbed  
into the body and  
distributed to the body  
tissues. • the rate and  
pathways by which

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Pharmacokinetics and pharmacodynamics describe, respectively, the amount of drug in the body at a given time and the pharmacologic effects caused by the drug. 1 Pharmacokinetics describes the movement of a drug into, within, and out of the body over time, whereas pharmacodynamics explains the effects the

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drug has on the body  
that result in a clinical  
response.

**General Principles of  
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pharmacokinetics. 5th  
ed. Baltimore:  
Lippincott Williams and  
Wilkins; 2010. Winter  
MA, Guhr KN, Berg GM.  
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weights and serum  
creatinine  
concentrations on the

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bias and accuracy of  
the Cockcroft-Gault  
equation.

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learn about the  
principles of clinical  
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Understanding the dose-concentration-effect  
relationship is a

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fundamental component of clinical pharmacology. Interpreting data arising from observations of this relationship requires the use of mathematical models; i.e. pharmacokinetic (PK) models to describe the relationship between dose and concentration and pharmacodynamic (PD) models describing the relationship between



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concentration ...

**Understanding and  
applying  
pharmacometric  
modelling and ...**

The pharmacokinetics  
was best described by  
a two-compartment  
model with linear  
absorption and  
combined linear and  
Michaelis-Menten  
elimination. The  
subcutaneous  
bioavailability of  
Brodalumab was 55%,

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absorption rate was  $0.30 \text{ day}^{-1}$ , and body-weight was found to affect the volume of distribution and clearance.

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