

## PHARMACOKINETIC ABBREVIATIONS

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<b>Variable</b>	<b>Abbreviation</b>
Absorption rate constant	Ka
AUC	mg*hr/L or mcg*hr/mL; AUC <sub>0-24 hr</sub> ; AUC <sub>0-tau</sub> or AUC <sub>0-τ</sub> ; AUC <sub>0-inf</sub> or AUC <sub>0-∞</sub>
Bioavailability	F
Clearance	CL; L/hr/kg or L/hr/m <sup>2</sup>
Creatinine clearance	CrCL
Elimination rate constant	ke
Estimate creatinine clearance	eCrCL
Estimated glomerular filtration rate	eGFR
Glomerular filtration rate	GFR
Half-life	t <sub>1/2</sub>
Intercompartmental rate constants	K <sub>12</sub> and K <sub>21</sub>
Maximum rate of metabolism	V <sub>max</sub>
Maximum serum concentration	C <sub>max</sub>
Mean residence time	MRT
Minimum serum concentration	C <sub>min</sub>
Model-informed drug development	MIDD
Modeling and simulation	M&S
Nonrenal clearance	CL <sub>NR</sub>
Pharmacokinetics	PK
Physiologically-based PK	PBPK
Plasma Concentration	C <sub>p</sub>
Population PK	PopPK
Renal clearance	CL <sub>R</sub>
Residence rate constant at steady state	k <sub>ss</sub>
Serum concentration	C <sub>p</sub>
Time to maximum concentration	T <sub>max</sub>
Volume of distribution	V <sub>d</sub>
Volume of distribution at steady state	V <sub>dss</sub>
Volume of the peripheral compartment	V <sub>p</sub>
Volume of the central compartment	V <sub>c</sub>