

Building and evaluation of a PBPK model for cimetidine in healthy adults

Version	1.0-OSP9.1
based on <i>Model Snapshot</i> and <i>Evaluation Plan</i>	https://github.com/Open-Systems-Pharmacology/Cimetidine-Model/releases/tag/v1.0
OSP Version	9.1
Qualification Framework Version	2.2

This evaluation report and the corresponding PK-Sim project file are stored at:

<https://github.com/Open-Systems-Pharmacology/OSP-PBPK-Model-Library/>

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1 Introduction

Cimetidine is a histamine H₂ receptor antagonist that inhibits stomach acid production. It is mainly used as an antacid for the treatment of gastric and duodenal ulcers, Zollinger-Ellison syndrome and esophageal reflux.

The herein presented model was developed and published by Hanke et al. ([Hanke 2020](#)).

Cimetidine is mainly excreted unchanged via the kidneys (40–80% of the dose) with a high renal clearance of 400 ml/min. Metabolism is reported to account for 25– 40% of the total elimination of cimetidine, with less than 2% of the dose excreted unchanged with the bile. Cimetidine inhibits several transporters and CYP enzymes and it is recommended by the FDA as strong inhibitor of OCT2/MATE and as weak inhibitor of CYP3A4 and CYP2D6 for the use in clinical DDI studies and drug labeling.

The cimetidine model was established using 27 clinical studies, covering a dosing range from 100 to 800 mg. The final model applies active uptake of cimetidine into the liver by OCT1, uptake into the kidney by OAT3 and secretion from the kidney into the urine by MATE1, as well as an unspecific hepatic clearance and passive renal glomerular filtration.

The herein presented model building and evaluation report evaluates the performance of the PBPK model for cimetidine in (healthy) adults.

2 Methods

2.1 Modeling Strategy

The general concept of building a PBPK model has previously been described by e.g. Kuepfer et al. ([Kuepfer 2016](#)). The relevant anthropometric (height, weight) and physiological information (e.g. blood flows, organ volumes, binding protein concentrations, hematocrit, cardiac output) in adults was gathered from the literature and has been previously published ([Willmann 2007](#)). This information was incorporated into PK-Sim® and was used as default values for the simulations in adults.

Variability of plasma proteins and CYP enzymes are integrated into PK-Sim® and described in the publicly available PK-Sim® Ontogeny Database Version 7.3 ([PK-Sim Ontogeny Database Version 7.3](#)) or otherwise referenced for the specific process. The final model applies active uptake of cimetidine into the liver by OCT1, uptake into the kidney by OAT3 and secretion from the kidney into the urine by MATE1, as well as an unspecific hepatic clearance and passive renal glomerular filtration. The transporters were integrated into the PBPK model using the ([PK-Sim Ontogeny Database Version 7.3](#)) and is described in detail in [Hanke 2020](#).

First, a base PBPK model was built using clinical data including single and multiple dose studies with intravenous and oral applications of cimetidine to find an appropriate structure to describe the pharmacokinetics in plasma. This PBPK model was developed using a typical European individual adjusted to the demography of the respective study population.

Oral administration of cimetidine in the fasted state frequently produces two plasma concentrations peaks. These double peaks are probably caused by the phasic gastrointestinal motility that controls gastric emptying in the fasted state. To describe the very different shapes of the observed mean cimetidine plasma profiles, split dose administration protocols for all studies of cimetidine administered orally in the fasted state were optimized in a NONMEM analysis (see [Hanke 2020](#)). The resulting split dose administration protocols were then implemented and used for the PBPK modeling of the respective cimetidine studies.

Unknown parameters (see below) were identified using the Parameter Identification module provided in PK-Sim®. Structural model selection was mainly guided by visual inspection of the resulting description of data and biological plausibility.

Details about input data (physicochemical, *in vitro* and clinical) can be found in [Section 2.2](#).

Details about the structural model and its parameters can be found in [Section 2.3](#).

2.2 Data

2.2.1 In vitro / physico-chemical Data

A literature search was performed to collect available information on physicochemical properties of cimetidine. The obtained information from literature is summarized in the table below.

Parameter	Unit	Value	Source	Description
MW	g/mol	252.34	Wishart 2006	Molecular weight
pK _{a1}	6.93	(base)	Avdeef 2001	Acid dissociation constant
pK _{a2}	13.38	(acid)	Wishart 2006	Acid dissociation constant
Solubility (pH)	mg/L	24.00 (6.8)	Avdeef 2001	Water solubility
logP		0.48	Avdeef 2001	Partition coefficient between octanol and water
f _u	%	78.00	Taylor 1978	Fraction unbound in plasma
B/P ratio		0.98	Somogyi 1983	Blood to plasma ratio
OCT1 K _m	μmol/l	2600	Umehara 2007	Michaelis-Menten constant
OAT3 K _m	μmol/l	149	Tahara 2005	Michaelis-Menten constant
MATE1 K _m	μmol/l	8.0	Ohta 2005	Michaelis-Menten constant
OCT1 K _i	μmol/l	104	Ito 2012	Inhibition constant for competitive inhibition
OCT2 K _i	μmol/l	124	Ito 2012	Inhibition constant for competitive inhibition
MATE1 K _i	μmol/l	3.8	Ito 2012	Inhibition constant for competitive inhibition
CYP3A4 K _i	μmol/l	268	Wrighton 1994	Inhibition constant for competitive inhibition

2.2.2 Clinical Data

A literature search was performed to collect available clinical data on efavirenz in healthy adults.

2.2.2.1 Model Building

The following studies were used for model building:

Publication	Arm / Treatment / Information used for model building
Bodemar 1981	Peptic ulcer patients receiving a single intravenous dose of 200 mg and oral doses of 200, 400 and 800 mg
Morgan 1983	Peptic ulcer patients receiving a single intravenous dose of 200 mg (5 min infusion)
Bodemar 1979	Healthy subjects receiving single oral doses of 200 and 400mg (tablet)
Walkenstein 1978	Healthy subjects receiving a single oral dose of 300mg (solution)
D'Angio 1986	Healthy subjects receiving a single oral dose of 300mg (tablet)

2.2.2.2 Model verification

The following studies were used for model verification:

Publication	Arm / Treatment / Information used for model verification
Grahnen 1979	Healthy subjects receiving a single intravenous dose of 100 mg and a single oral dose of 400 mg (tablet)
Larsson 1982	Peptic ulcer patients receiving a single intravenous dose of 200 mg
Mihaly 1984	Peptic ulcer patients receiving a single intravenous and a single oral dose of 200 mg
Morgan 1983	Peptic ulcer patients receiving a single intravenous dose of 200 mg (30 min infusion)
Lebert 1981	Healthy subjects receiving a single intravenous dose of 300 mg (2 min infusion)
Walkenstein 1978	Healthy subjects receiving a single intravenous dose of 300 mg (2 min infusion) and a single oral dose of 300 mg (tablet)
Kanto 1981	Healthy subjects receiving a single oral dose of 200 mg
Burland 1975	Healthy subjects receiving single oral doses of 200 mg solution and capsule
Bodemar 1979	Peptic ulcer patients receiving a single oral dose of 200 mg (tablet)
Bodemar 1981	Peptic ulcer patients receiving single oral doses of 800 mg and multiple oral doses of 200 and 400 mg
Barbhaiya 1995	Healthy subjects receiving multiple oral doses of 300 mg (tablet)
Somogyi 1981	Healthy subjects receiving a single oral dose of 400 mg (tablet)
Tiseo 1998	Healthy subjects receiving multiple oral doses of 800 mg (tablet)

2.3 Model Parameters and Assumptions

2.3.1 Absorption

Absorption observed in clinical studies can be fully explained by passive absorption.

2.3.2 Distribution

Cimetidine is reported to be actively taken up into the liver by OCT1 ([Umehara 2007](#)), into the kidney by OAT3 ([Tahara 2005](#)) and secreted from the kidney into the urine by MATE1 ([Ohta 2010](#)).

After testing the available organ-plasma partition coefficient and cell permeability calculation methods built in PK-Sim, observed clinical data was best described by choosing the partition coefficient calculation method by [Rodgers and Rowland](#) and cellular permeability calculation by [PK-Sim Standard](#).

A [Lipophilicity](#) of 1.66 was back-calculated from the blood-to-plasma ratio of 0.98 ([Somogyi 1983](#), [Hanke 2020](#)).

2.3.3 Metabolism, Elimination and Inhibition

Cimetidine is mainly excreted unchanged via the kidneys. Additionally, 25 to 40 % is hepatically metabolized via an unknown pathway.

Cimetidine inhibits several enzymes such as CYP3A4 and CYP2D6 as well as transporters such as OCT2, OCT2 and MATE.

2.3.4 Automated Parameter Identification

The parameter identification tool in PK-Sim has been used to estimate selected model parameters by adjusting to PK data of the clinical studies that were used in the model building process (see [Section 2.2](#)).

The result of the final parameter identification is shown in the table below:

Model Parameter	Optimized Value	Unit
Specific intestinal permeability	8.72E-7	cm/min
CL _{hep}	0.16	1/min
k _{cat} OCT1	8.66E+4	1/min
k _{cat} OAT3	5.75E+07	1/min
k _{cat} MATE1	32.37	1/min

3 Results and Discussion

The PBPK model for efavirenz was developed and evaluated using publicly available clinical pharmacokinetic data from studies listed in [Section 2.2.2](#).

The next sections show:

1. the final model parameters for the building blocks: [Section 3.1](#).
2. the overall goodness of fit: [Section 3.2](#).
3. simulated vs. observed concentration-time profiles for the clinical studies used for model building and for model verification: [Section 3.3](#).

3.1 Final input parameters

The compound parameter values of the final PBPK model are illustrated below.

Compound: Cimetidine

Parameters

Name	Value	Value Origin	Alternative	Default
Solubility at reference pH	24 mg/ml	Publication-Avdeef 2001	Measurement	True
Reference pH	6.8	Publication-Avdeef 2001	Measurement	True
Lipophilicity	1.655 Log Units	Parameter Identification	Measurement	True
Fraction unbound (plasma, reference value)	0.78	Publication-Taylor 1978	Measurement	True
Specific intestinal permeability (transcellular)	8.7196061807E-07 cm/min	Parameter Identification	Fit	True
Is small molecule	Yes			
Molecular weight	252.34 g/mol	Database-Drugbank		
Plasma protein binding partner	Unknown			

Calculation methods

Name	Value
Partition coefficients	Rodgers and Rowland
Cellular permeabilities	PK-Sim Standard

Processes

Systemic Process: Total Hepatic Clearance-Somogyi 1983

Species: Human

Parameters

Name	Value	Value Origin
Fraction unbound (experiment)	0.78	
Lipophilicity (experiment)	1.655 Log Units	
Plasma clearance	0 ml/min/kg	
Specific clearance	0.1600808777 1/min	Parameter Identification

Transport Protein: MATE1-Paper

Molecule: MATE1

Parameters

Name	Value	Value Origin
Transporter concentration	1 $\mu\text{mol/l}$	
Vmax	0 $\mu\text{mol/l/min}$	
Km	8 $\mu\text{mol/l}$	Parameter Identification
kcat	32.371673382 1/min	Parameter Identification

Transport Protein: OAT3-Paper

Molecule: OAT3

Parameters

Name	Value	Value Origin
Transporter concentration	1 $\mu\text{mol/l}$	
Vmax	0 $\mu\text{mol/l/min}$	
Km	149 $\mu\text{mol/l}$	Publication-Tahara 2005
kcat	57506485.67 1/min	Parameter Identification

Transport Protein: OCT1-Paper

Molecule: OCT1

Parameters

Name	Value	Value Origin
Transporter concentration	1 $\mu\text{mol/l}$	
Vmax	0 $\mu\text{mol/l/min}$	
Km	2600 $\mu\text{mol/l}$	Publication-Umehara 2007
kcat	86599.4219923521 1/min	Parameter Identification

Systemic Process: Glomerular Filtration-GFR

Species: Human

Parameters

Name	Value	Value Origin
GFR fraction	1	

Inhibition: OCT1-Ito 2012

Molecule: OCT1

Parameters

Name	Value	Value Origin
Ki	104 $\mu\text{mol/l}$	Publication-Ito 2012

Inhibition: OCT2-Ito 2012

Molecule: OCT2

Parameters

Name	Value	Value Origin
Ki	124 $\mu\text{mol/l}$	Publication-Ito 2012

Inhibition: MATE1-Ito 2012

Molecule: MATE1

Parameters

Name	Value	Value Origin
Ki	3.8 $\mu\text{mol/l}$	Other-NBI measurement

Inhibition: CYP3A4-Wrighton 1994

Molecule: CYP3A4

Parameters

Name	Value	Value Origin
Ki	268 $\mu\text{mol/l}$	

Formulation: Tablet

Type: Weibull

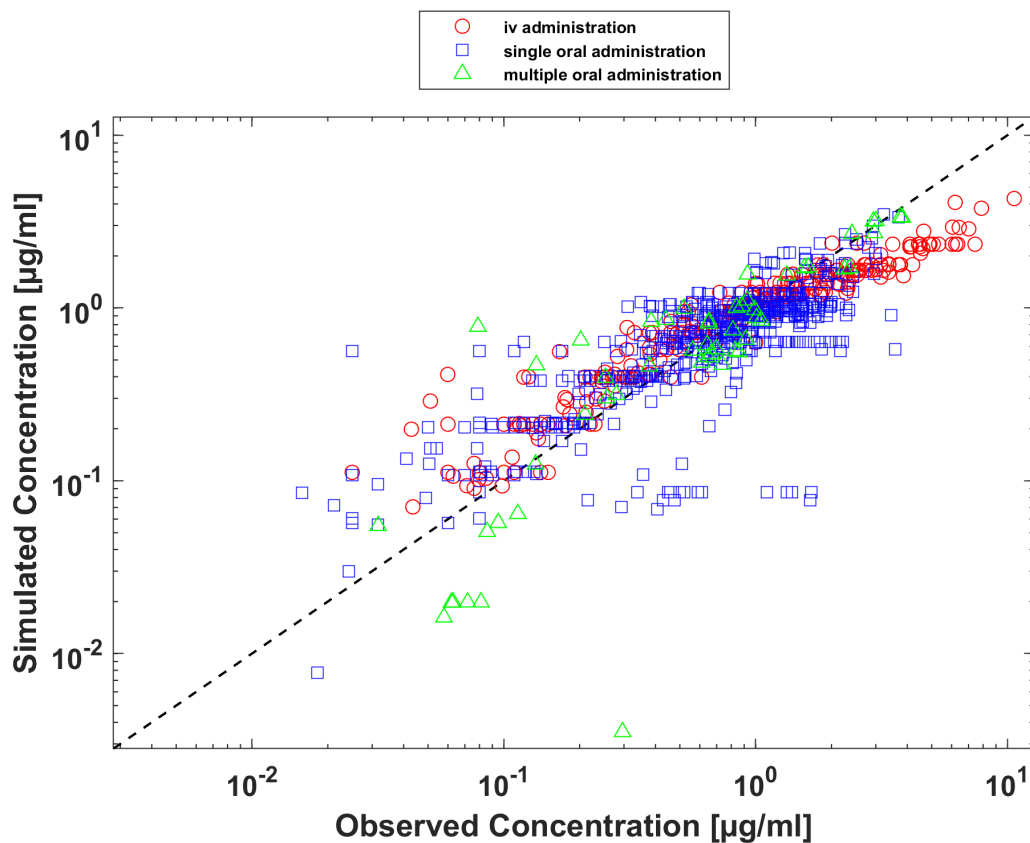
Parameters

Name	Value	Value Origin
Dissolution time (50% dissolved)	1 min	
Lag time	0 h	
Dissolution shape	10	
Use as suspension	Yes	

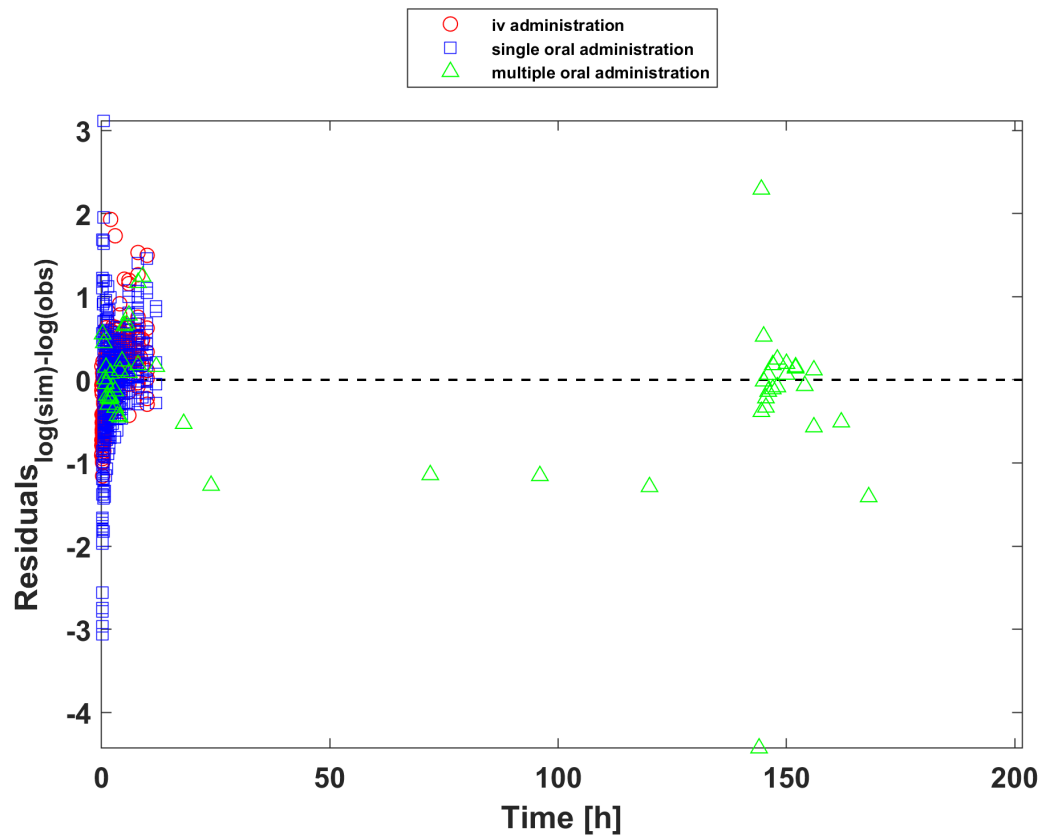
3.2 Diagnostics Plots

Below you find the goodness-of-fit visual diagnostic plots for the PBPK model performance of all data used presented in [Section 2.2.2](#).

The first plot shows simulated versus observed plasma concentration, the second weighted residuals versus time.



Goodness of fit plot for concentration in plasma



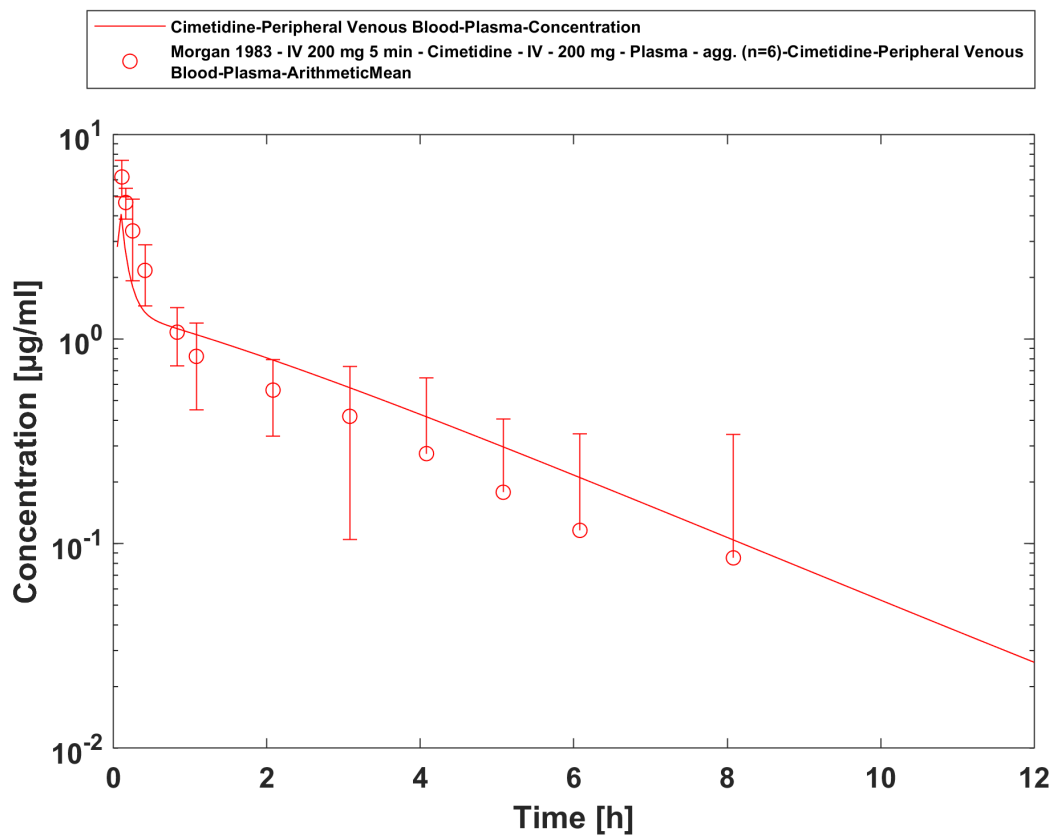
Goodness of fit plot for concentration in plasma

GMFE = 1.530217

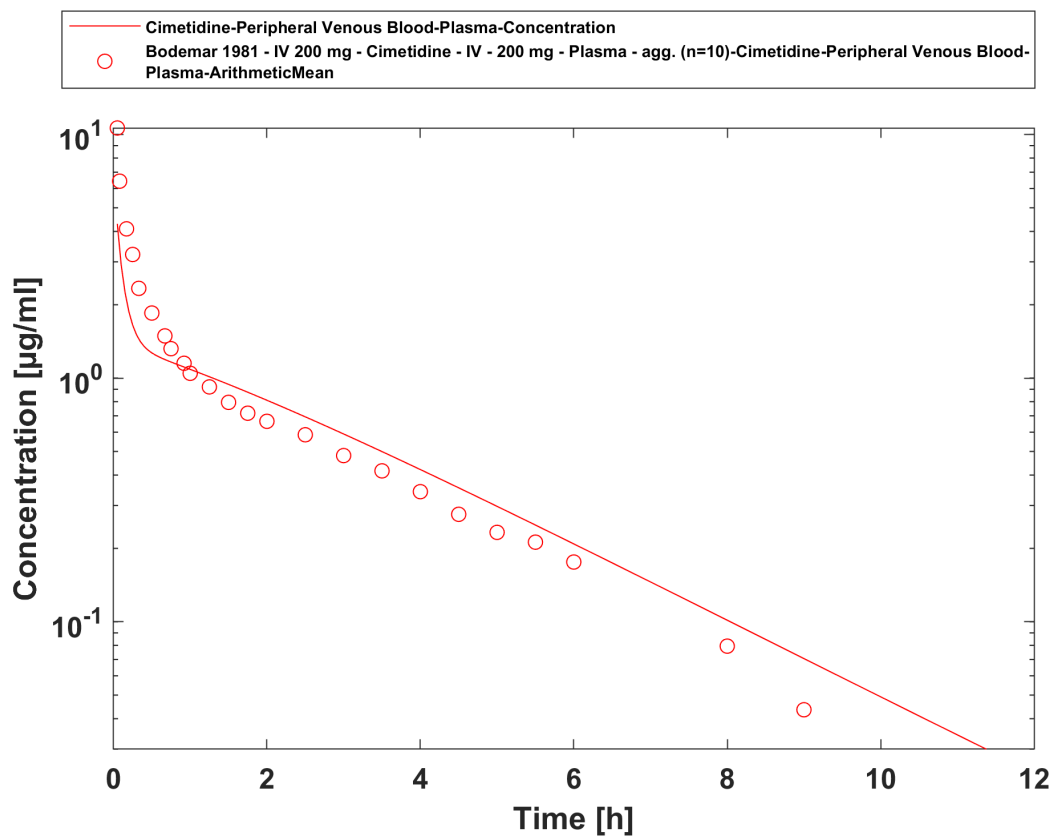
3.3: Concentration-Time Profiles

Simulated versus observed concentration-time profiles of all data listed in [Section 2.2.2](#) are presented below.

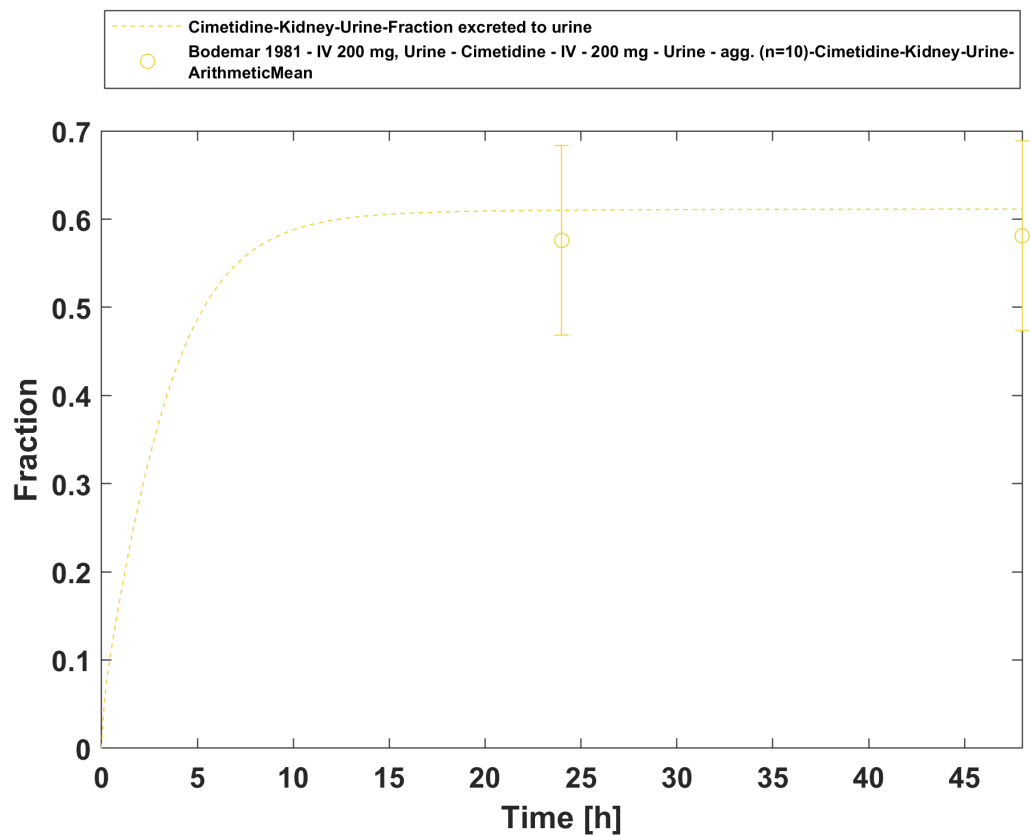
3.3.1 Model Building



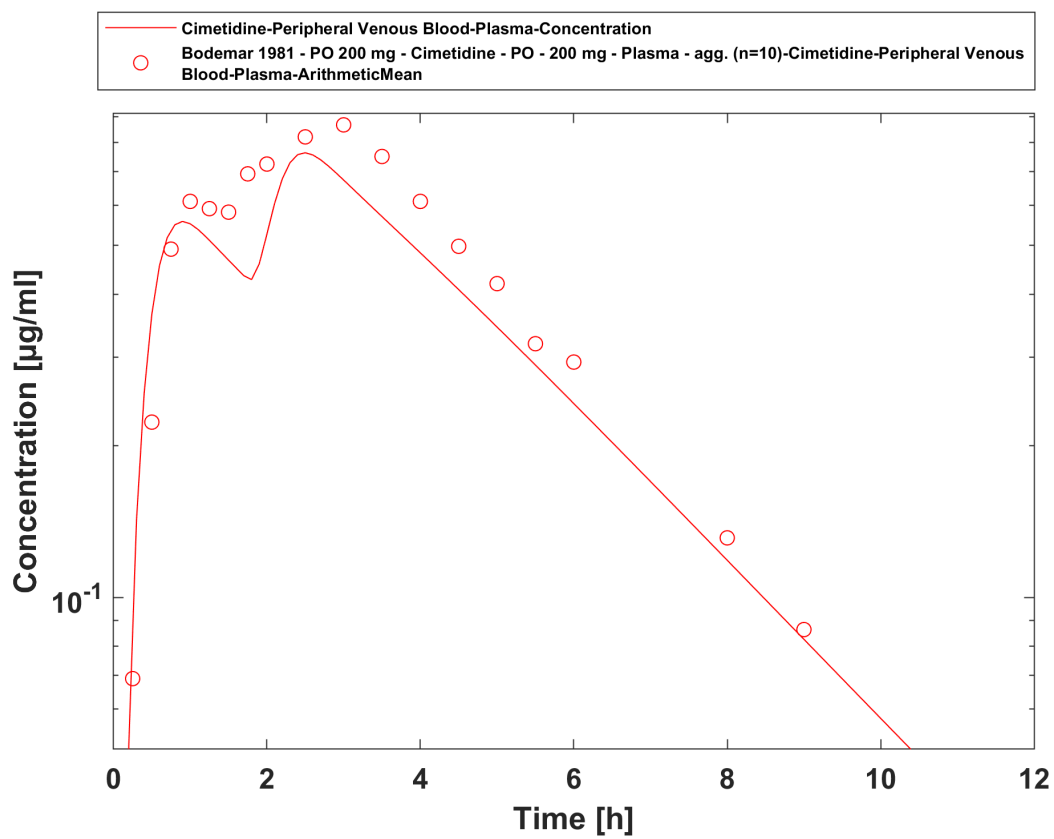
iv 200 mg (5 min),Morgan 1983, n=6



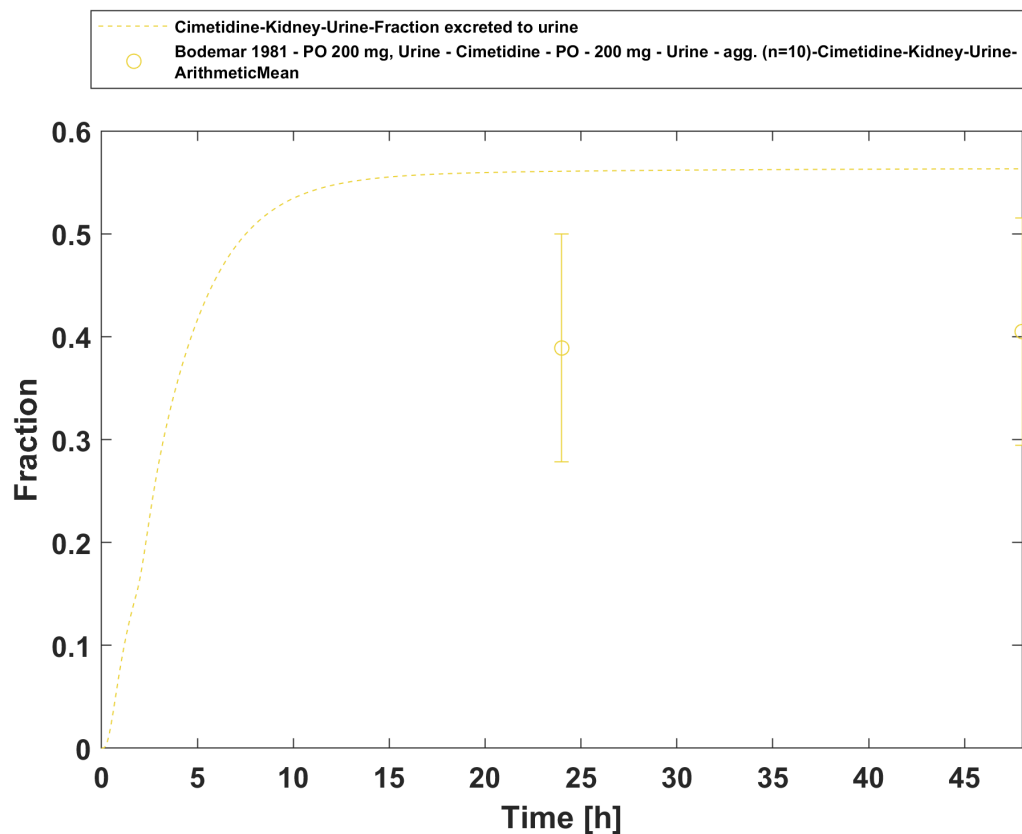
iv 200 mg, Bodemar 1981, n=10



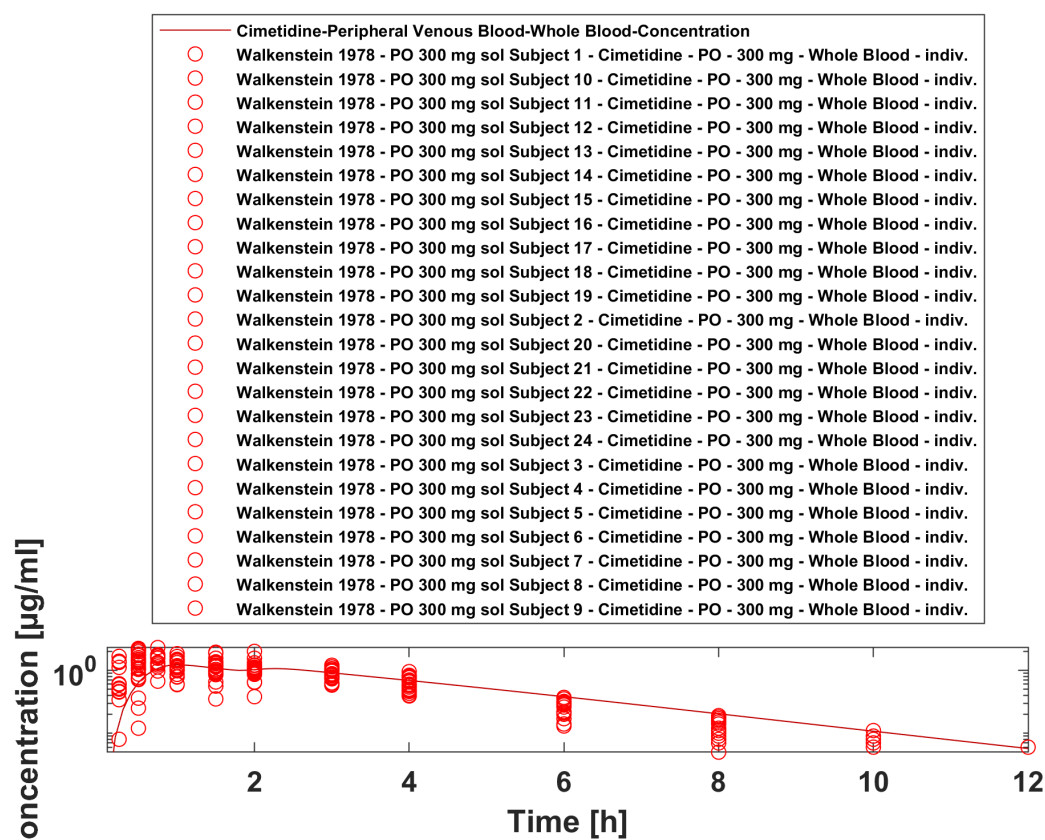
iv 200 mg, Bodemar 1981, n=10, urine



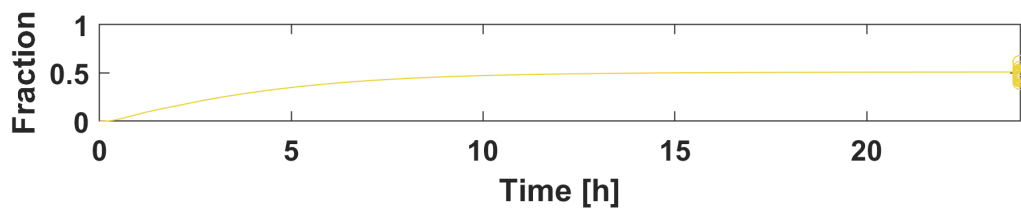
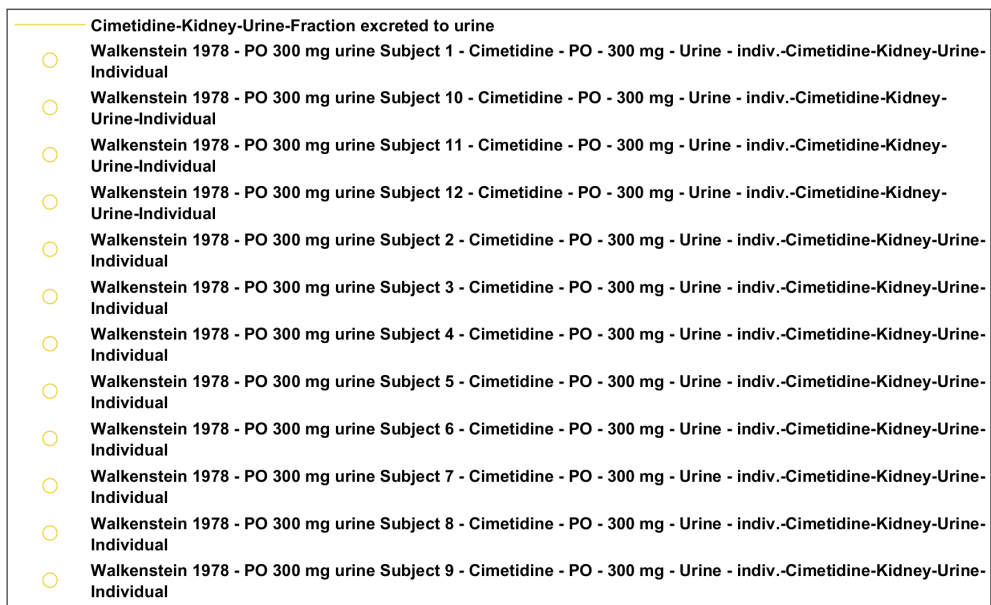
po 200 mg, Bodemar 1981, n=10



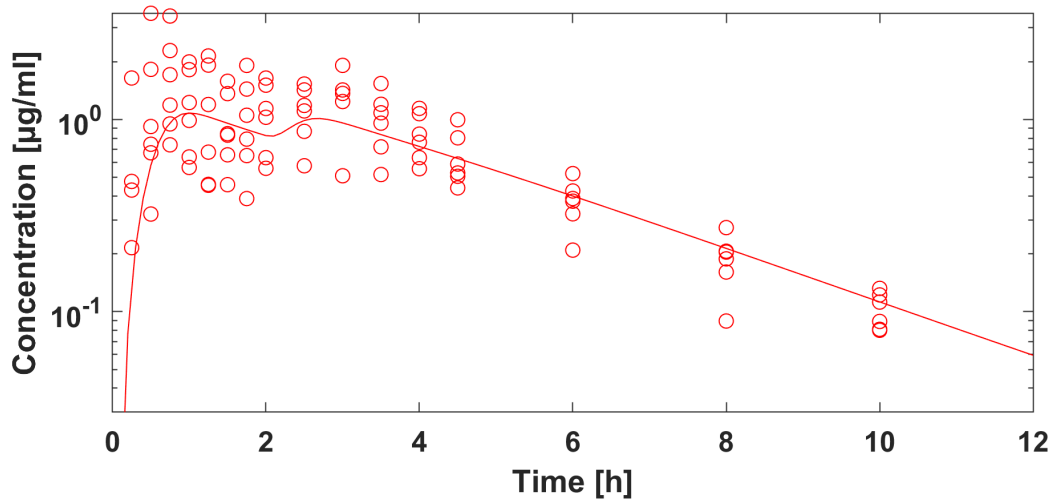
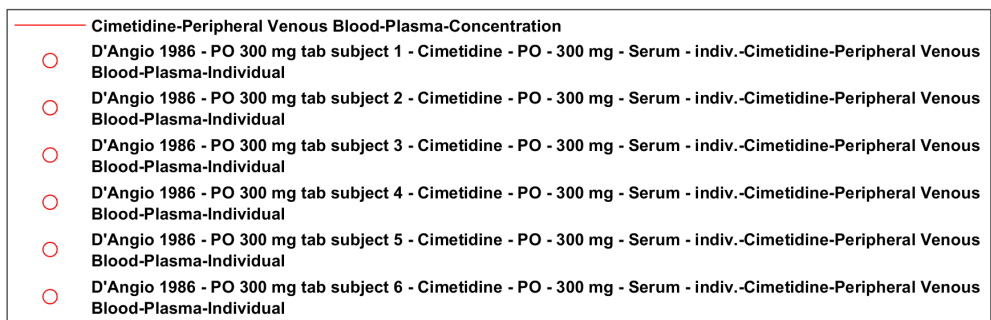
po 200 mg, Bodemar 1981, n=10, urine



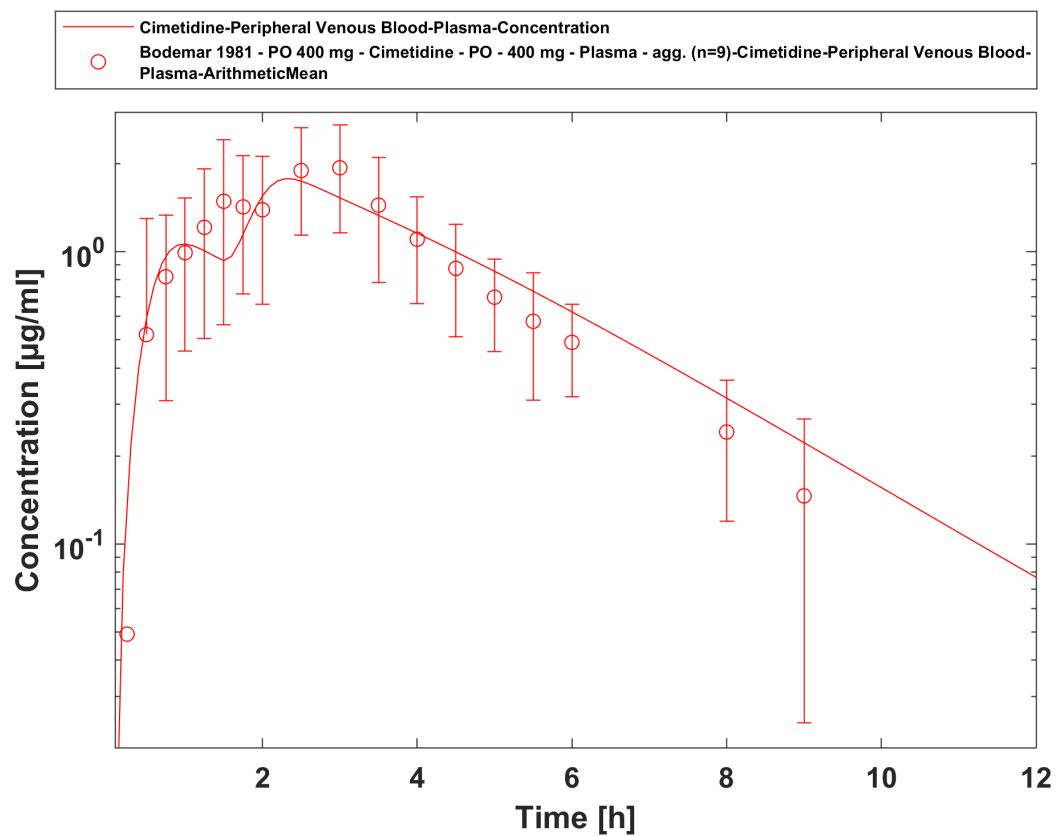
po 300 mg (sol), Walkenstein 1978, n=24



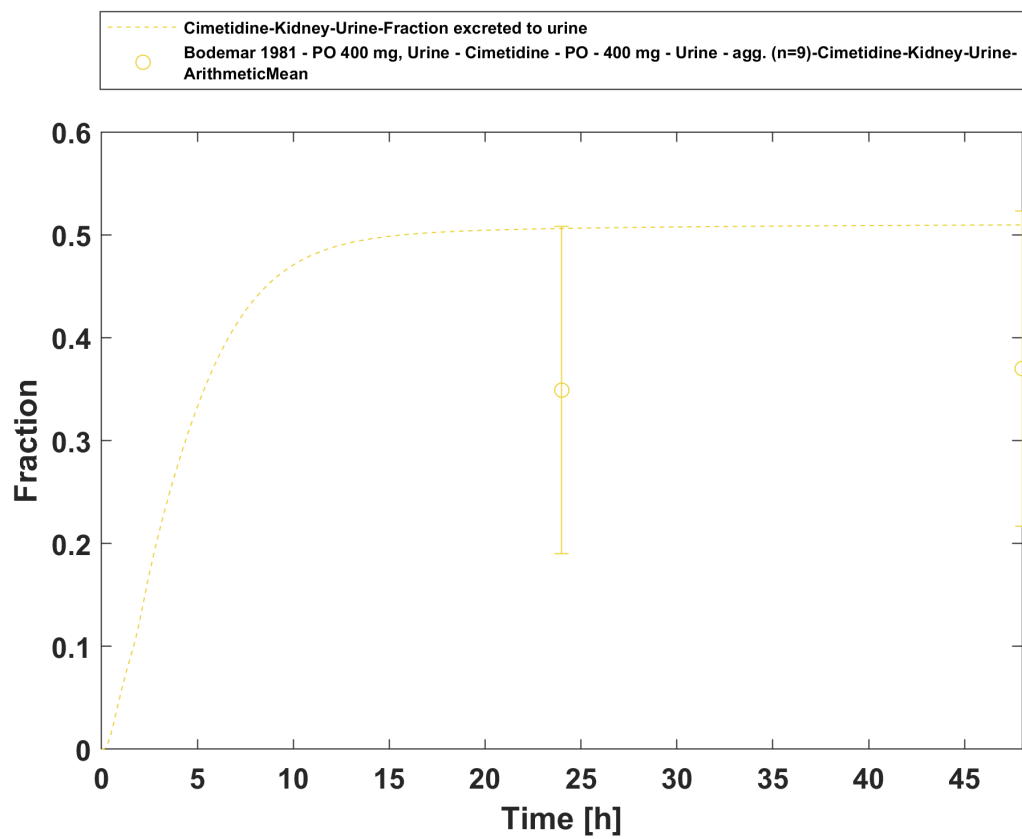
po 300 mg (sol), Walkenstein 1978, n=24, urine



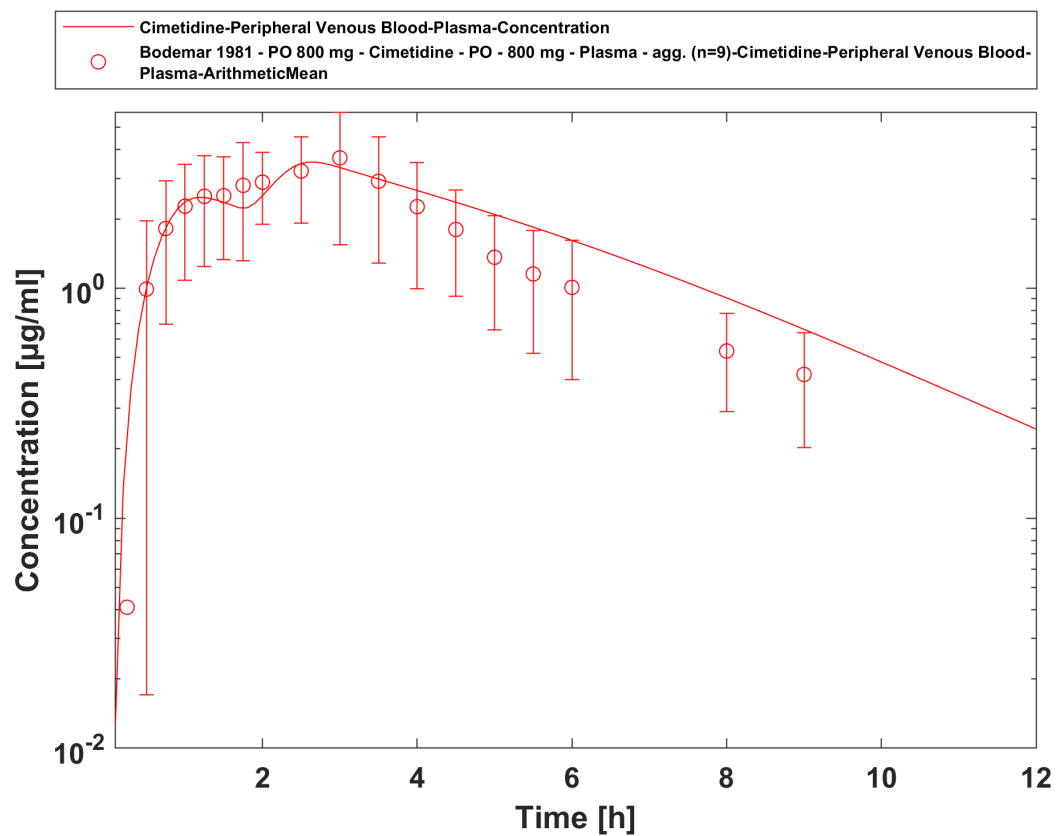
po 300 mg (tab), D'Angio 1986, n=6



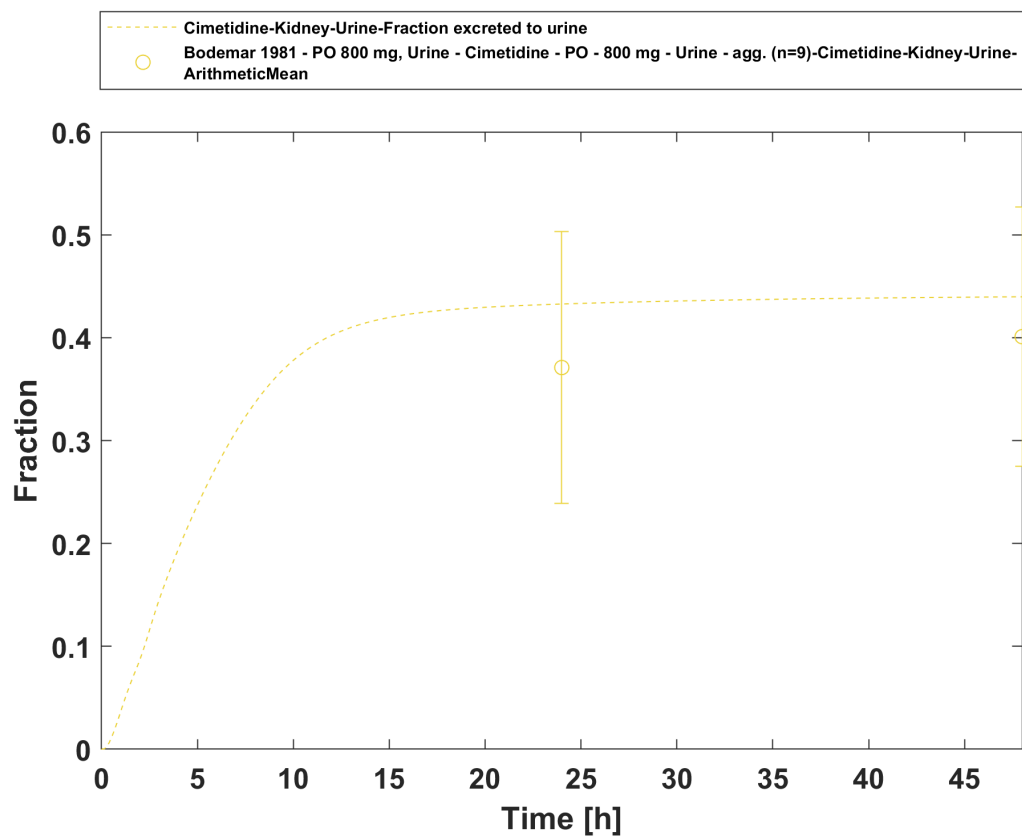
po 400 mg, Bodemar 1981, n=9



po 400 mg, Bodemar 1981, n=9, urine

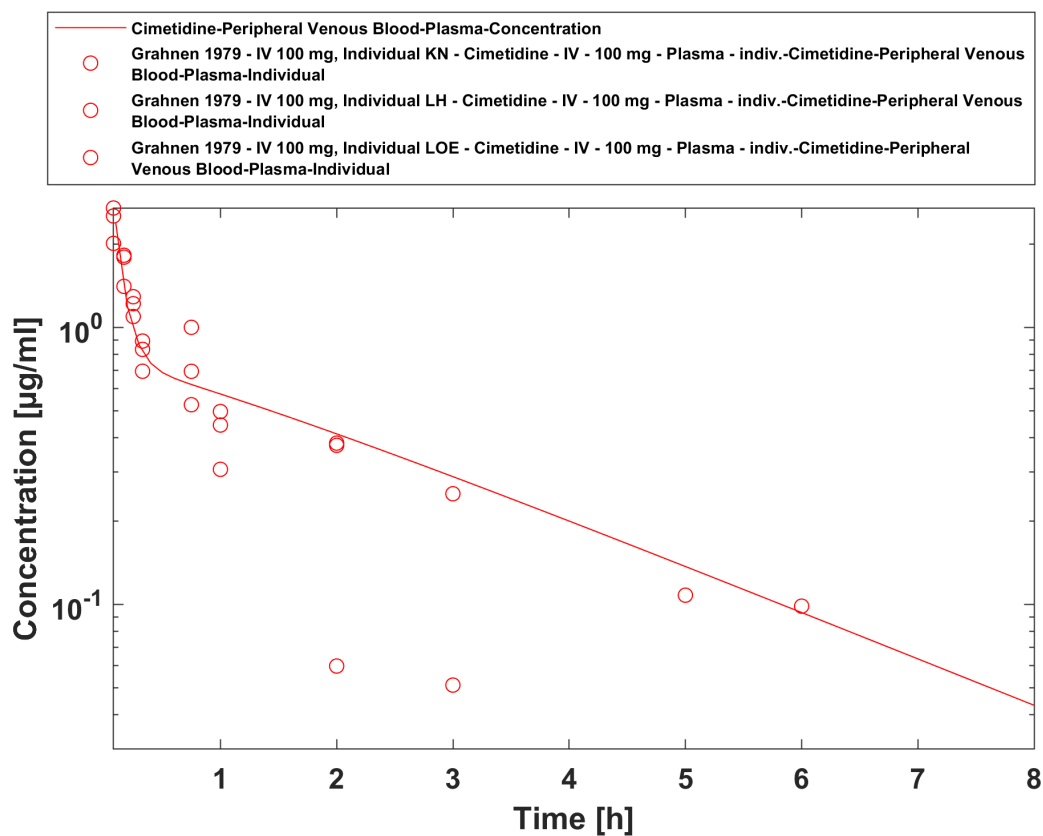


po 800 mg, Bodemar 1981, n=9

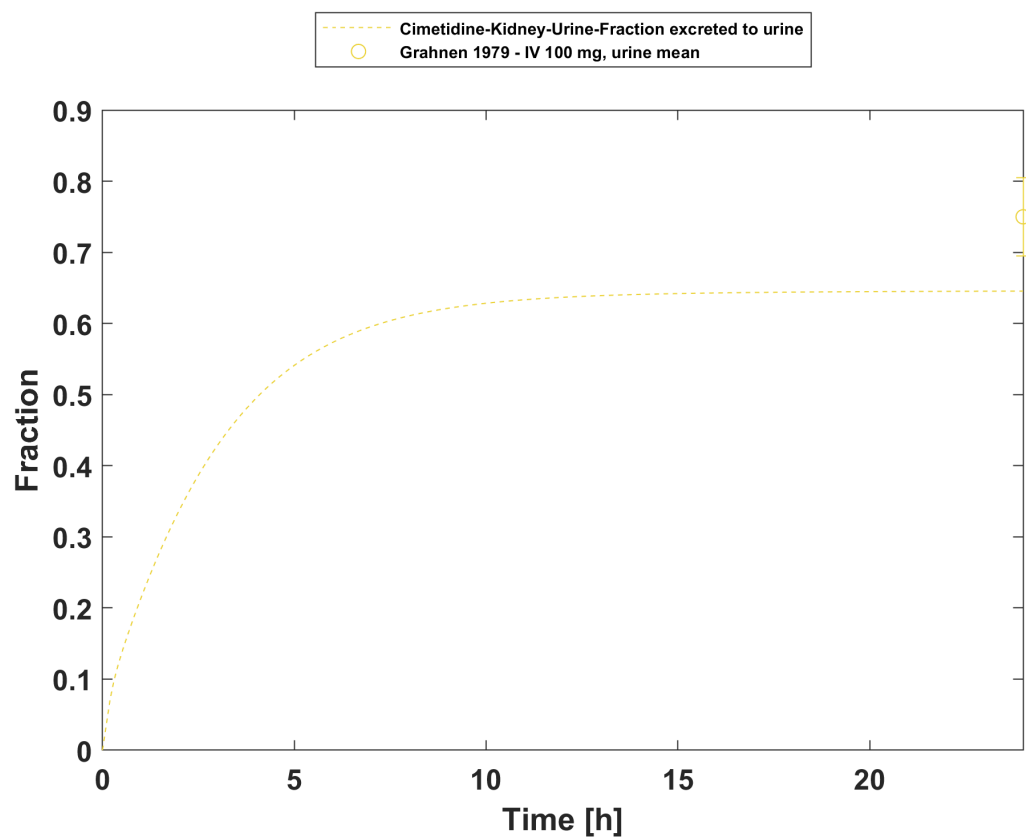


po 800 mg, Bodemar 1981, n=9, urine

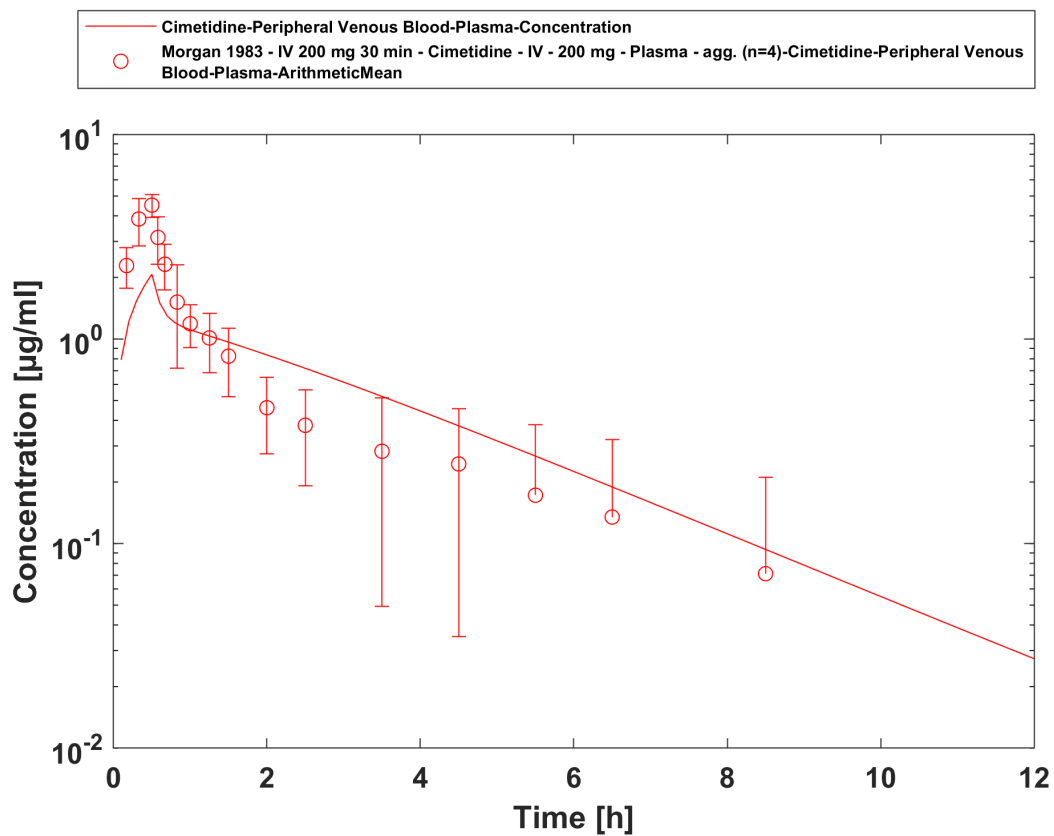
3.3.2 Model Validation



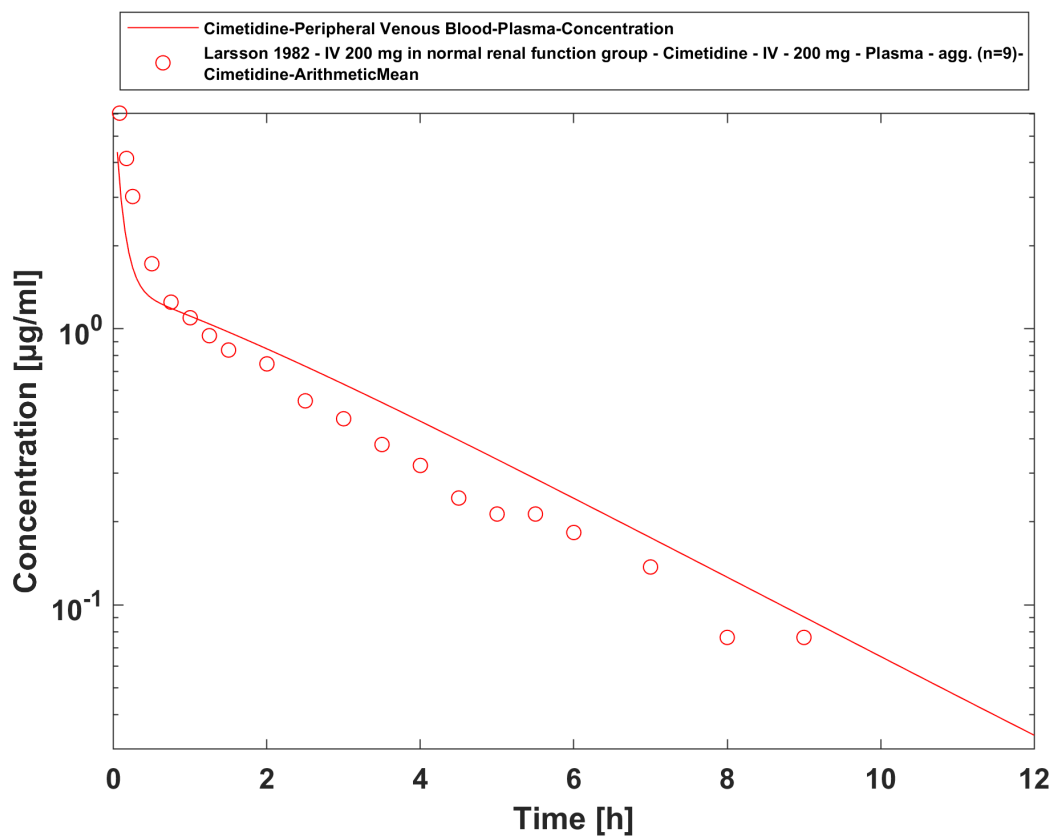
iv 100 mg (5 min), Grahnen 1979, n=3



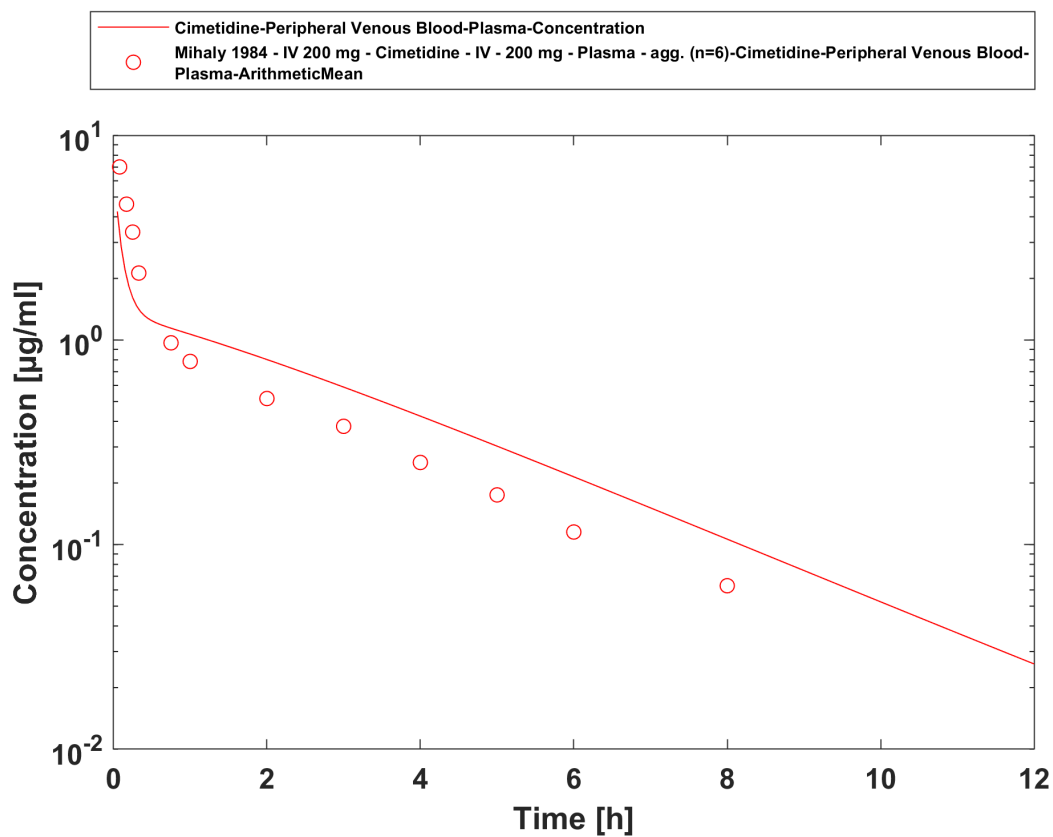
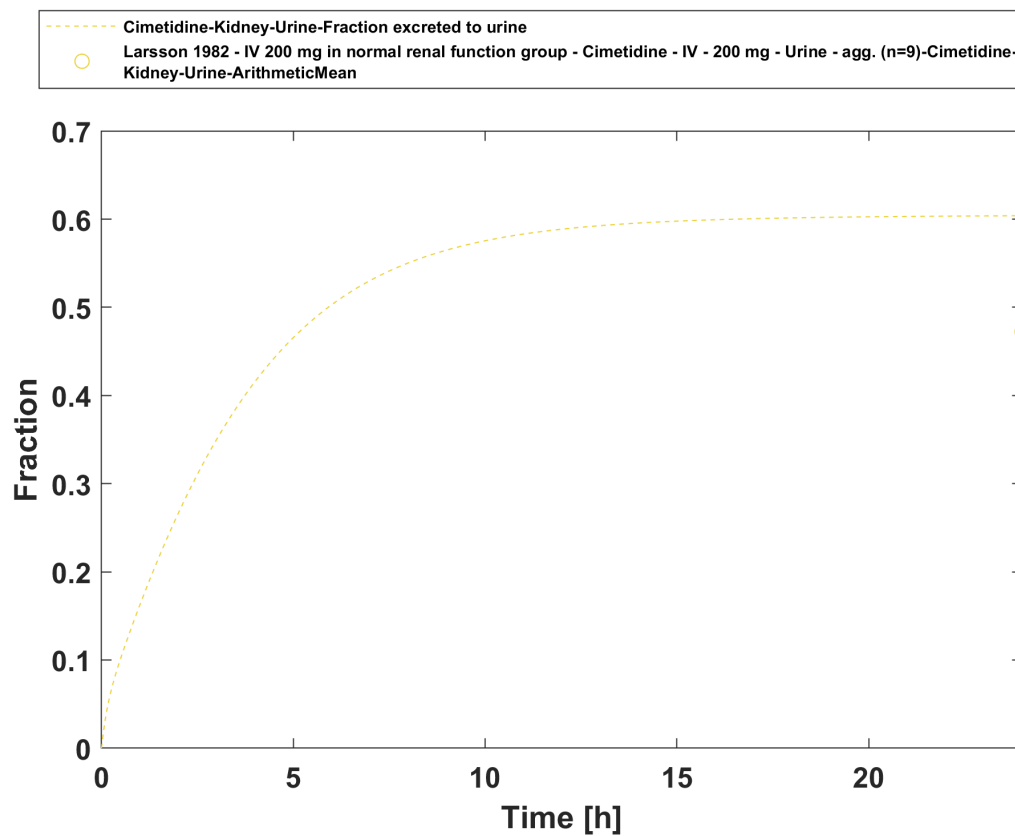
iv 100 mg (5 min), Grahnen 1979, n=3, urine

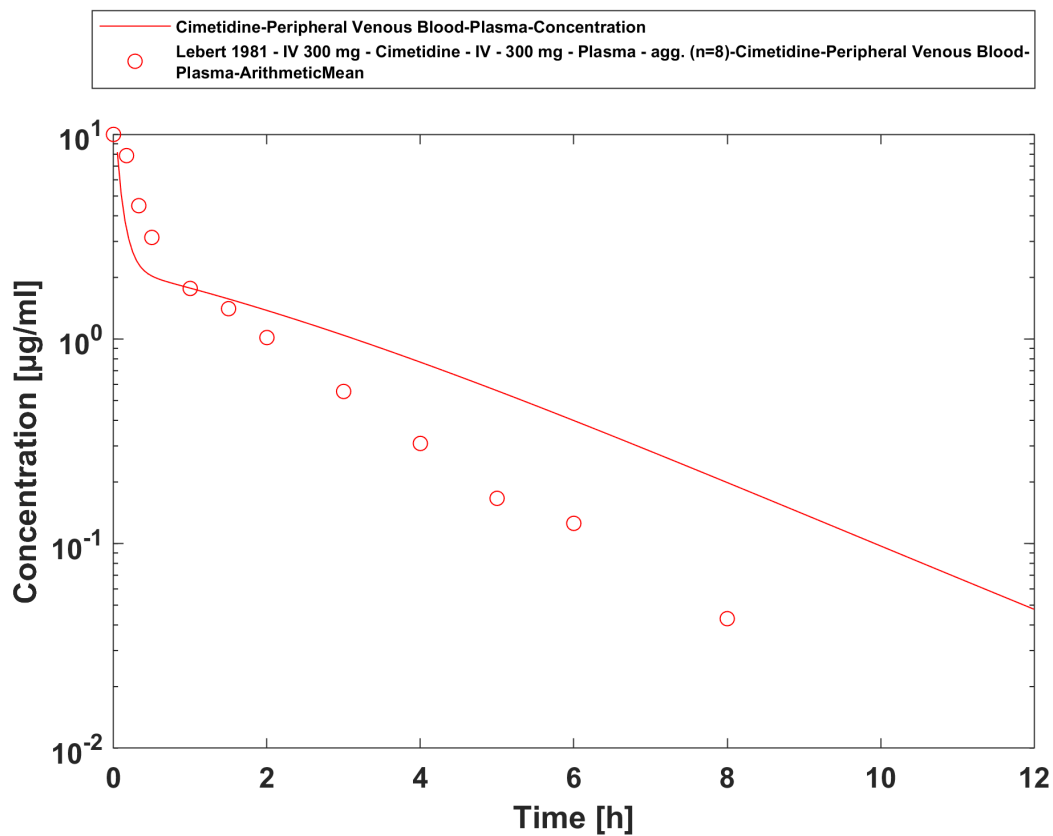


iv 200 mg (30 min),Morgan 1983, n=4

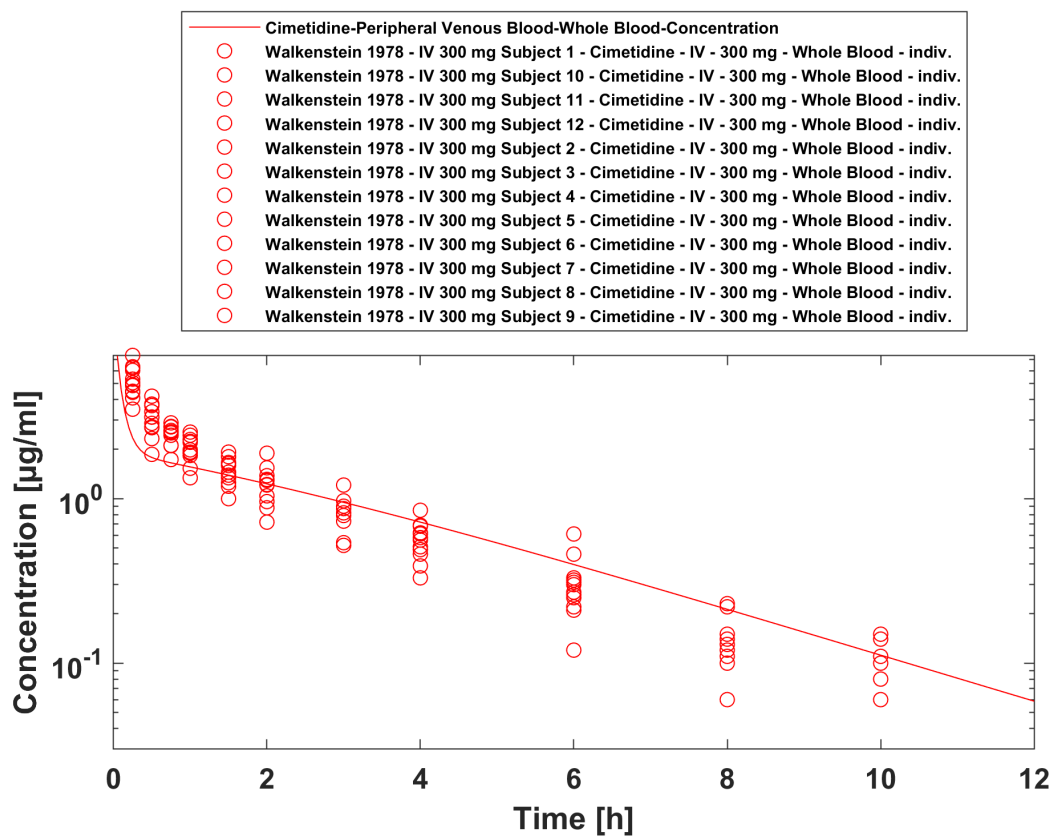


iv 200 mg, Larsson 1982, n=9

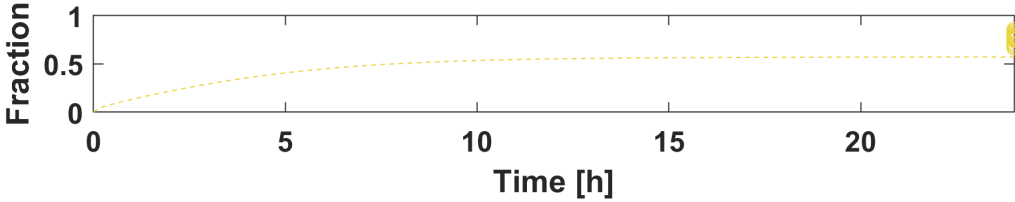
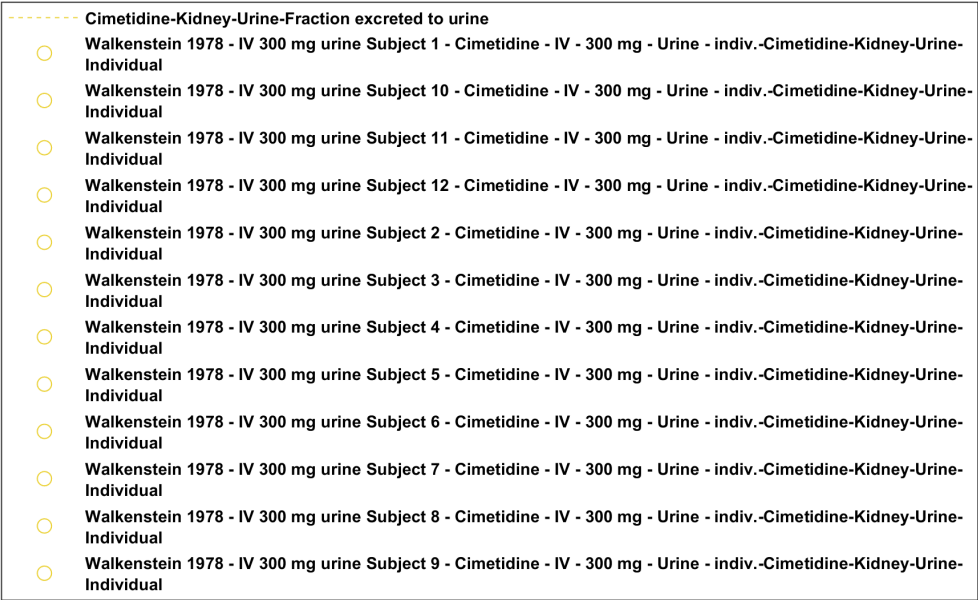




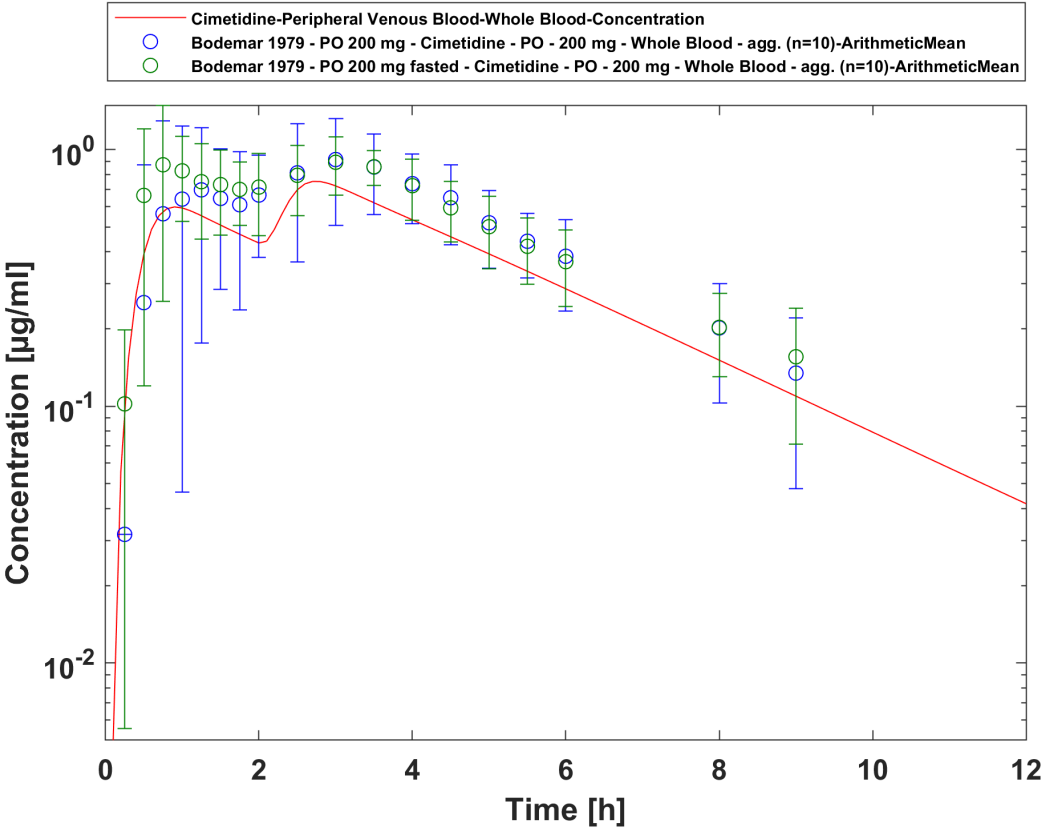
iv 300 mg (2 min), Lebert 1981, n=1



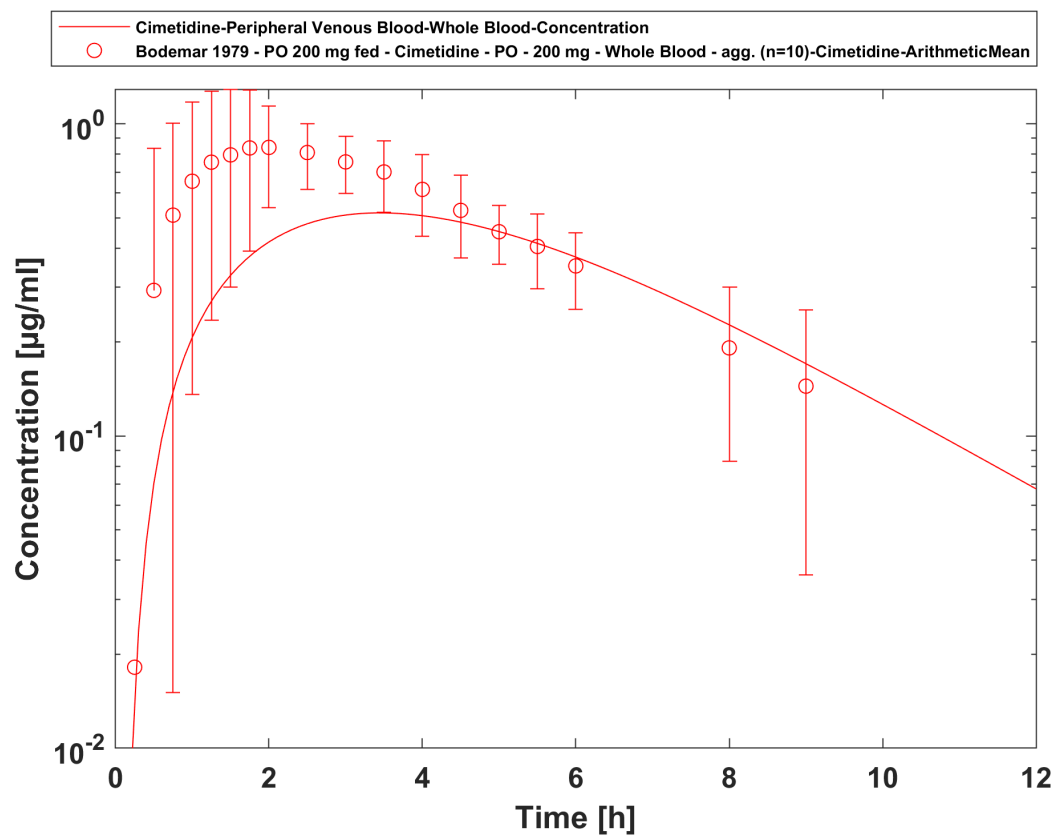
iv 300 mg (2 min), Walkenstein 1978, n=12



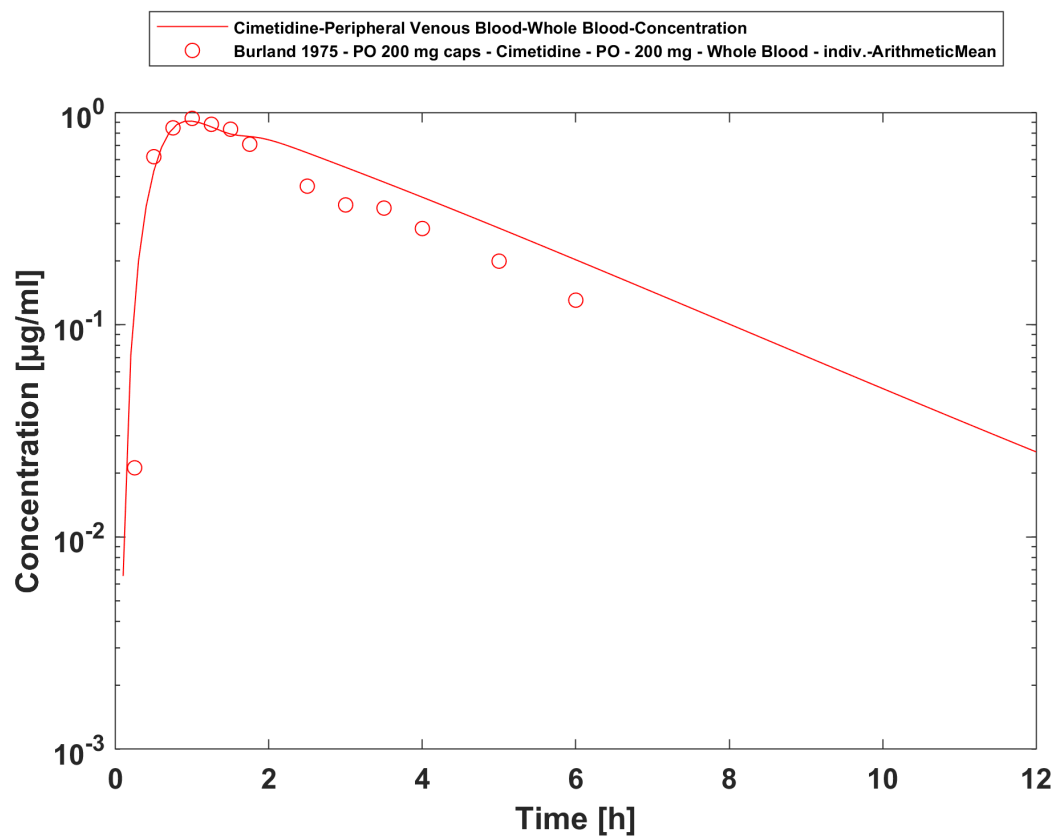
iv 300 mg (2 min), Walkenstein 1978, n=12



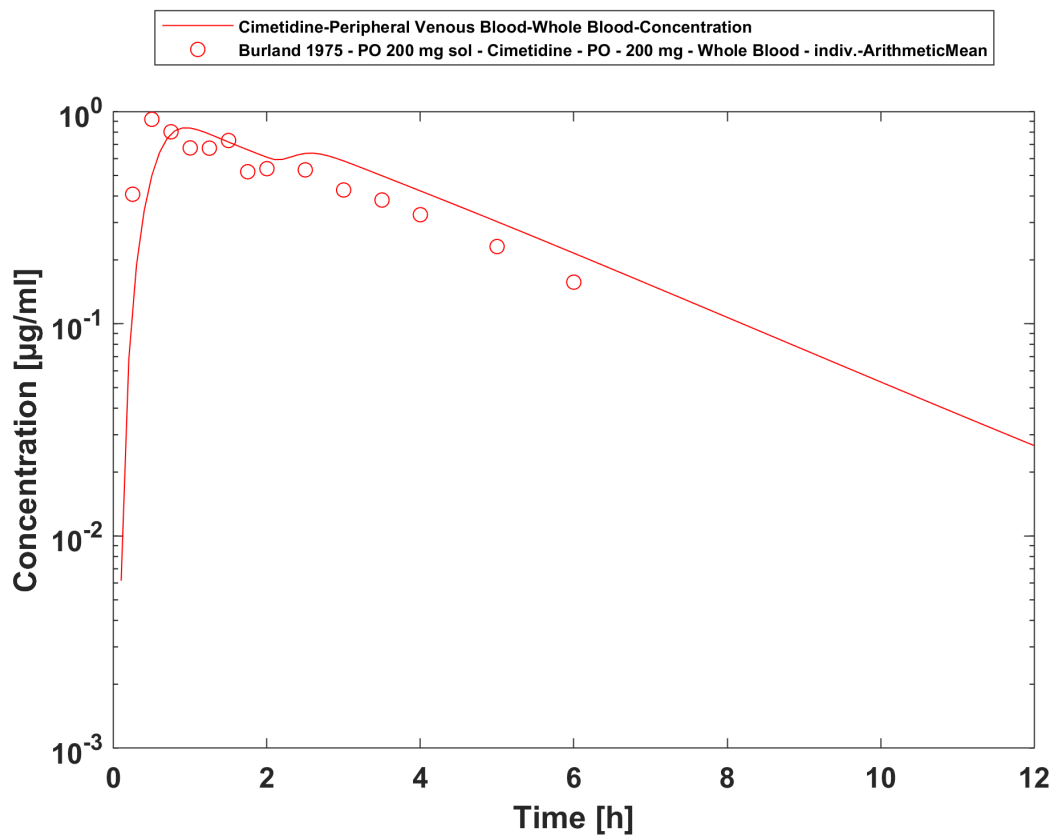
po 200 mg (tab), Bodemar 1979 (fasted)



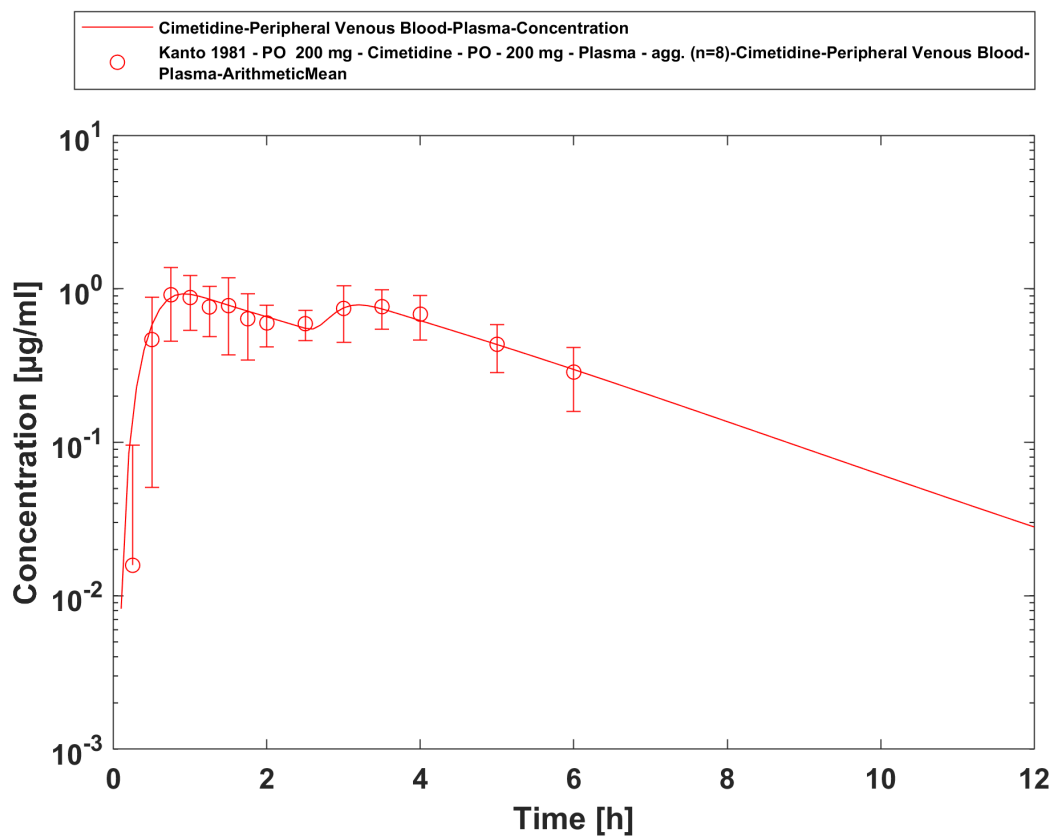
po 200 mg (tab), Bodemar 1979 (fed)



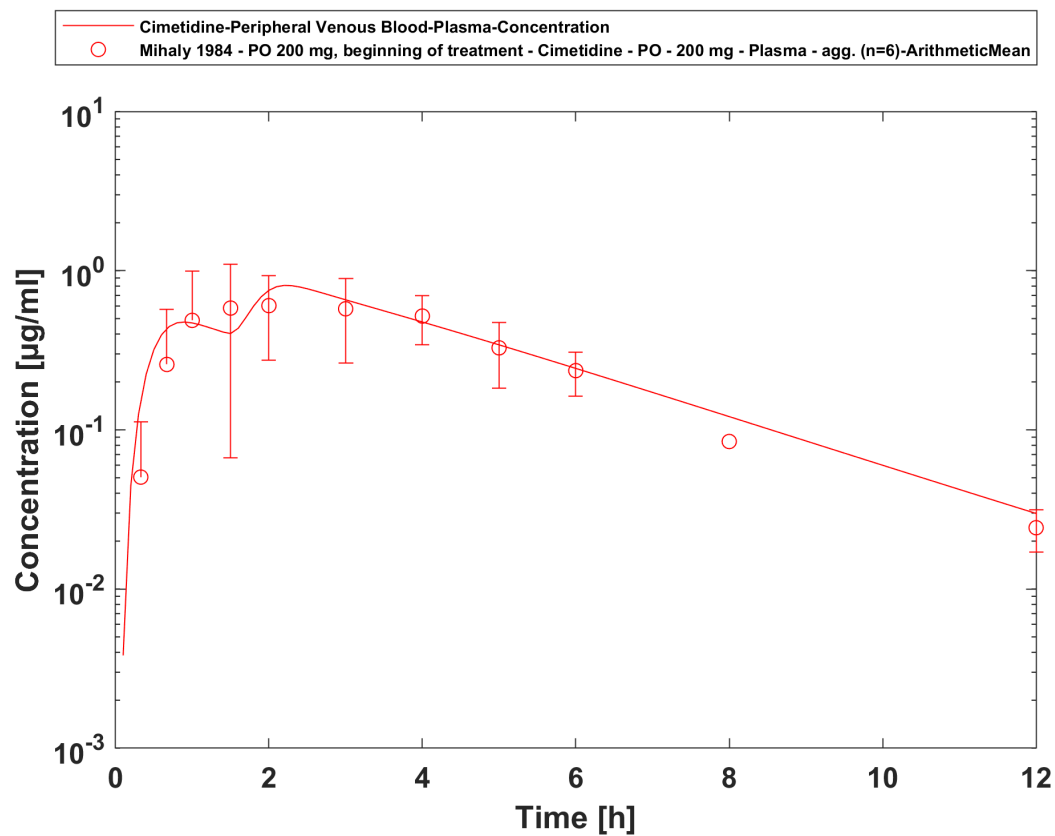
po 200 mg, Burland 1975, caps



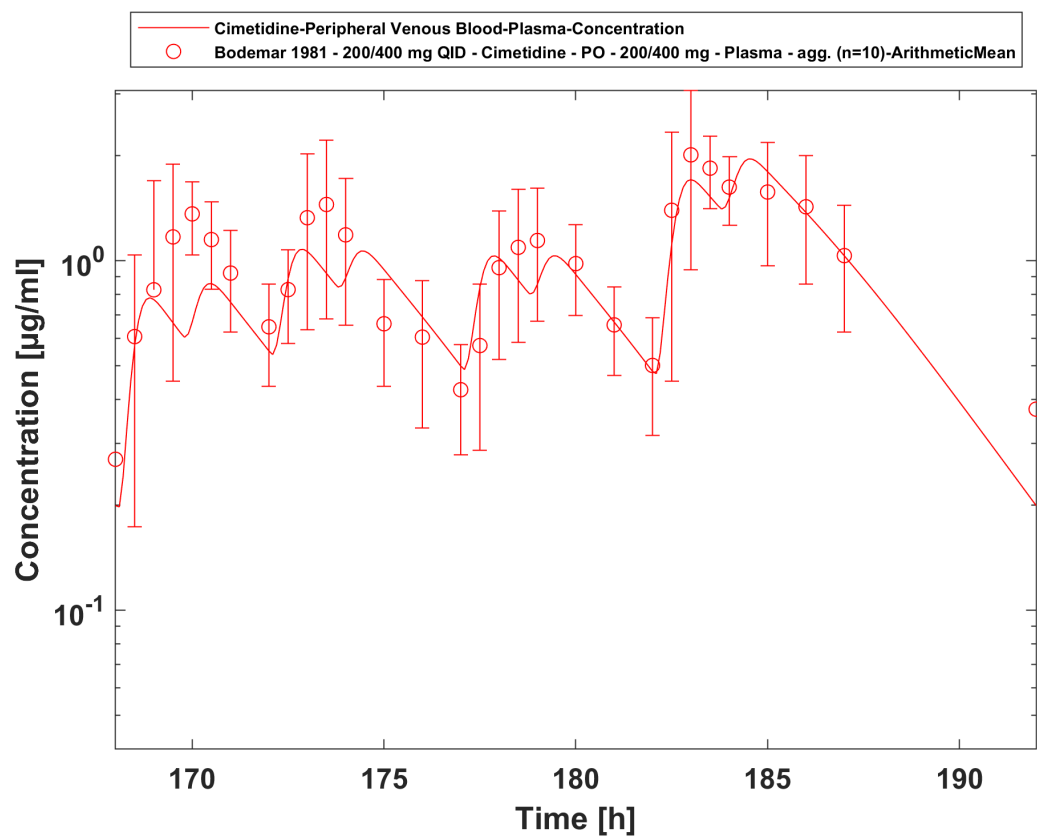
po 200 mg, Burland 1975, sol



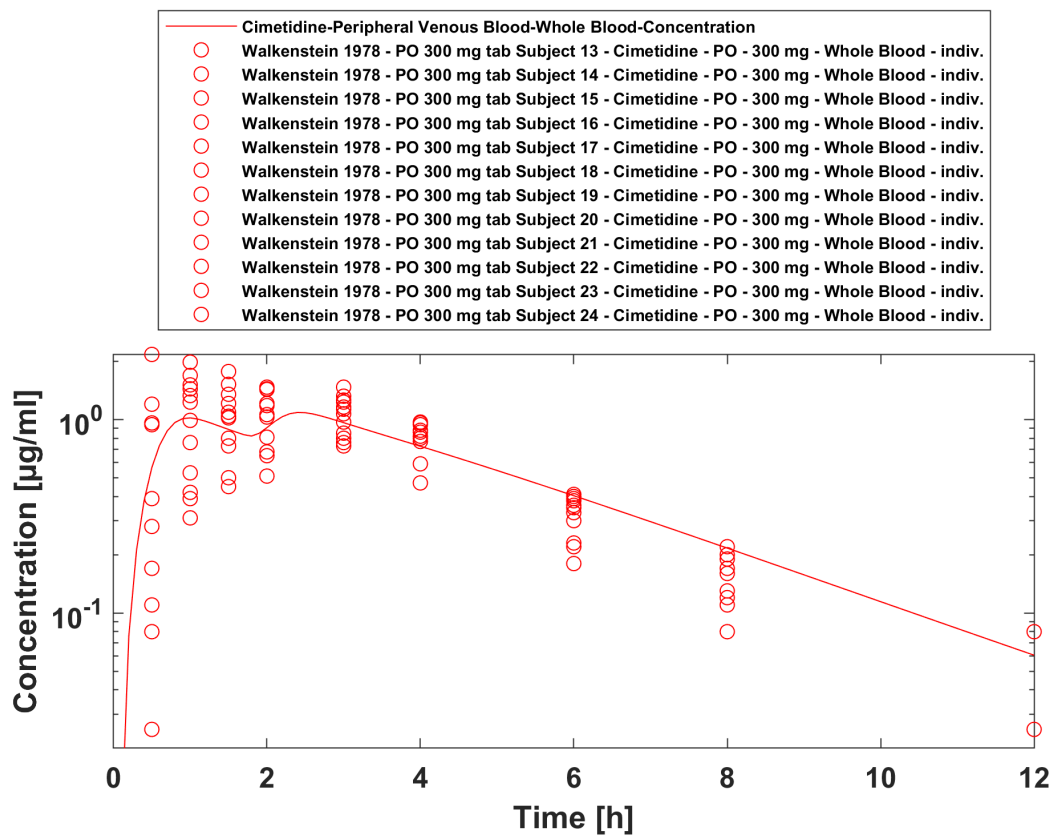
po 200 mg, Kanto 1981, n=8



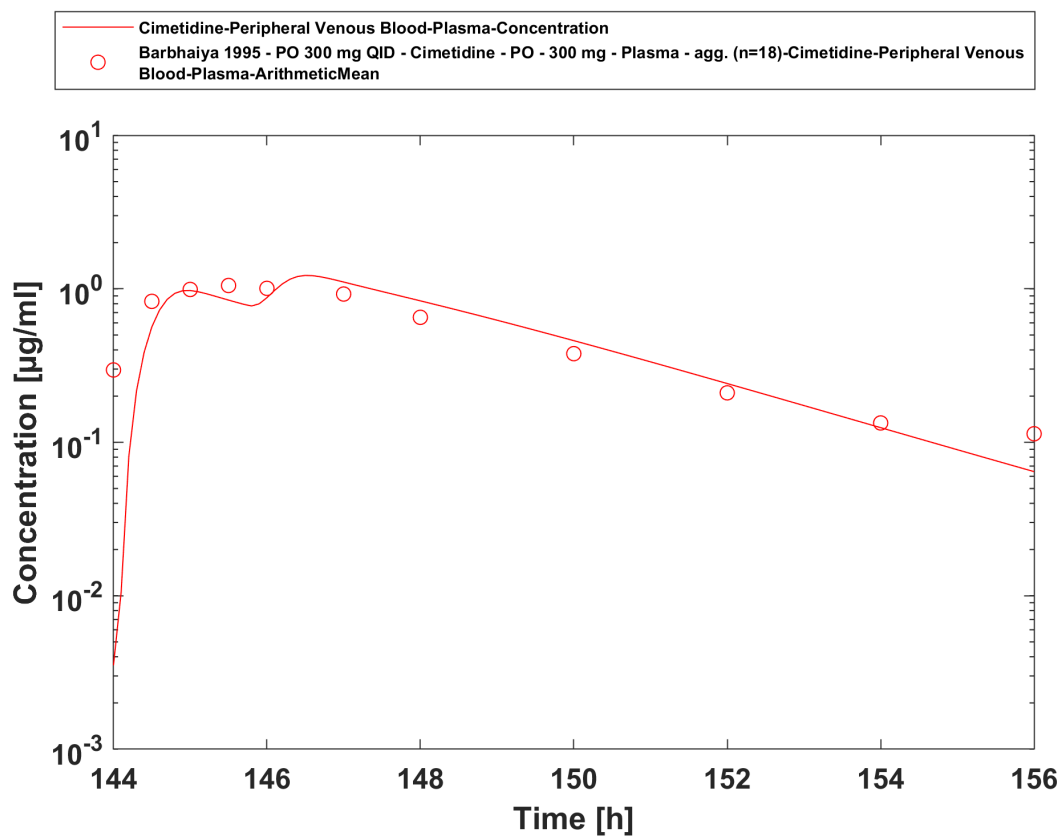
po 200 mg, Mihaly 1984, n=8



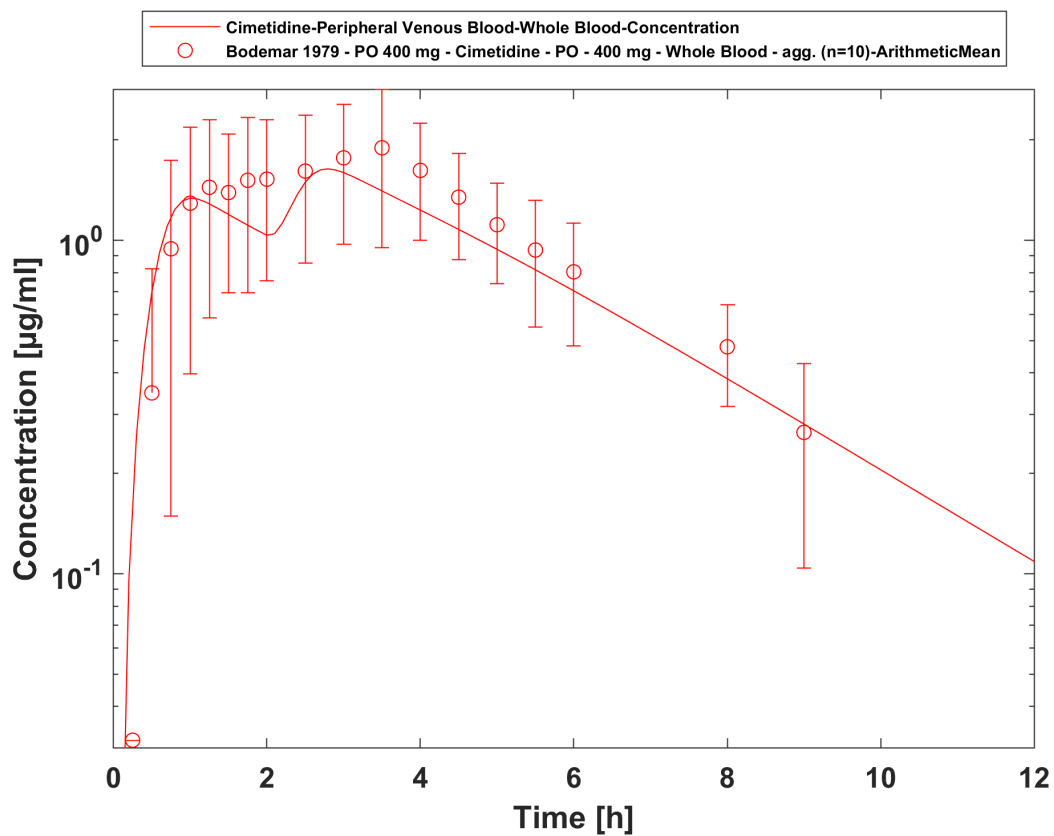
po 200/400 mg QID, Bodemar 1981 (fasted)



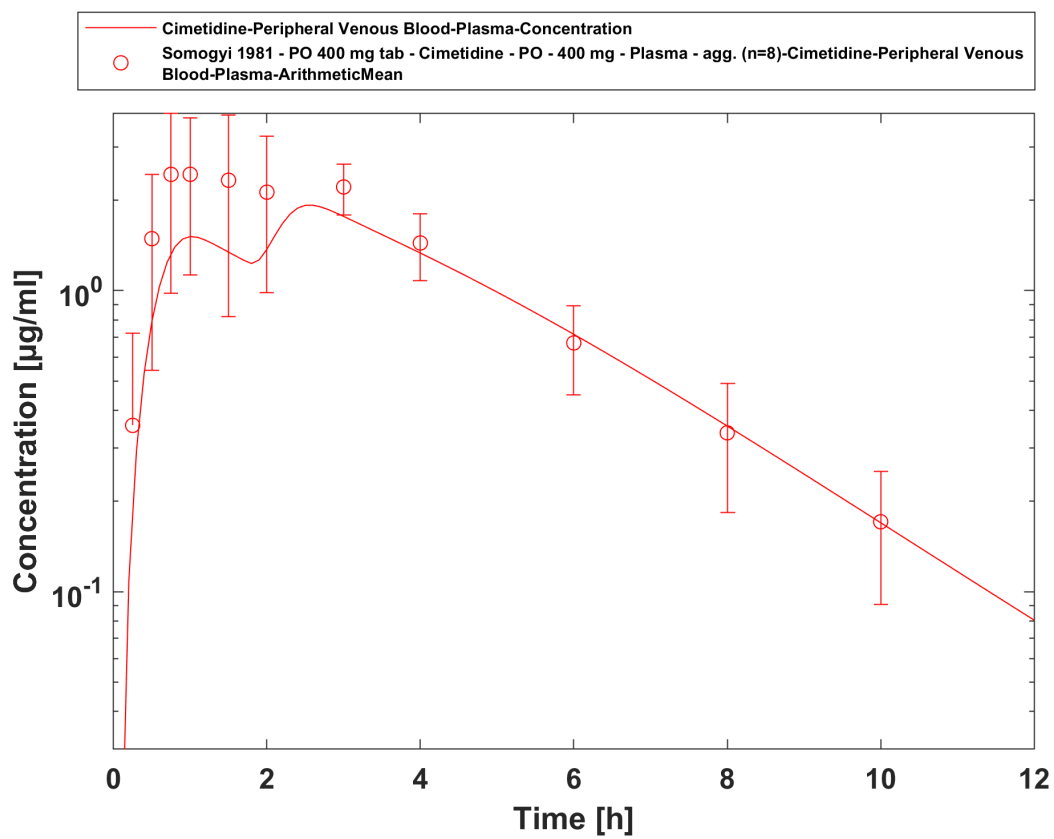
po 300 mg (tabl), Walkenstein 1978, n=12



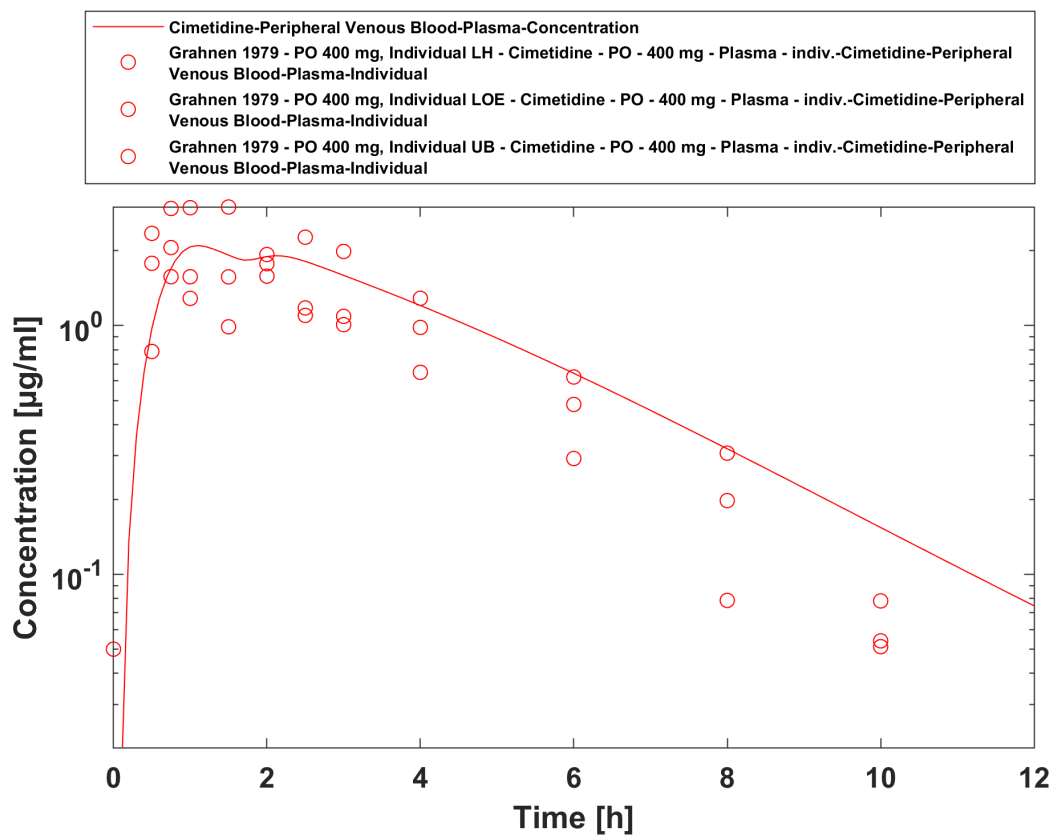
po 300 mg QID (sol), Barbhaiya 1995, n=18



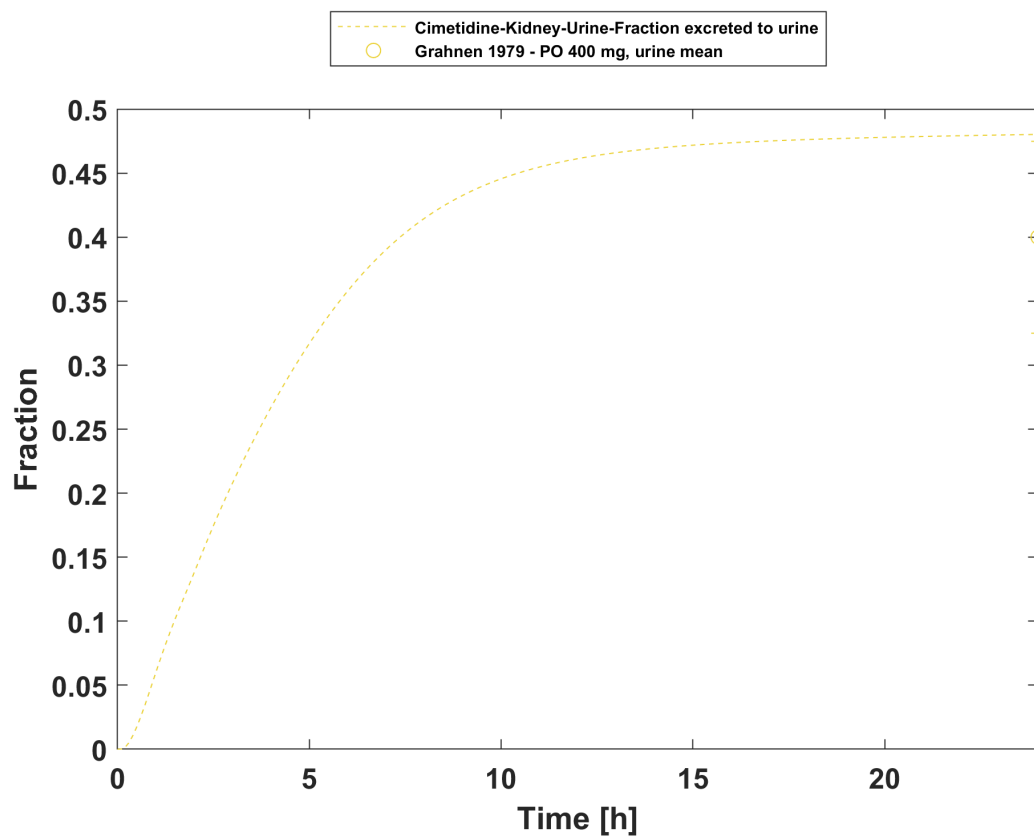
po 400 mg (tab), Bodemar 1979, n=10



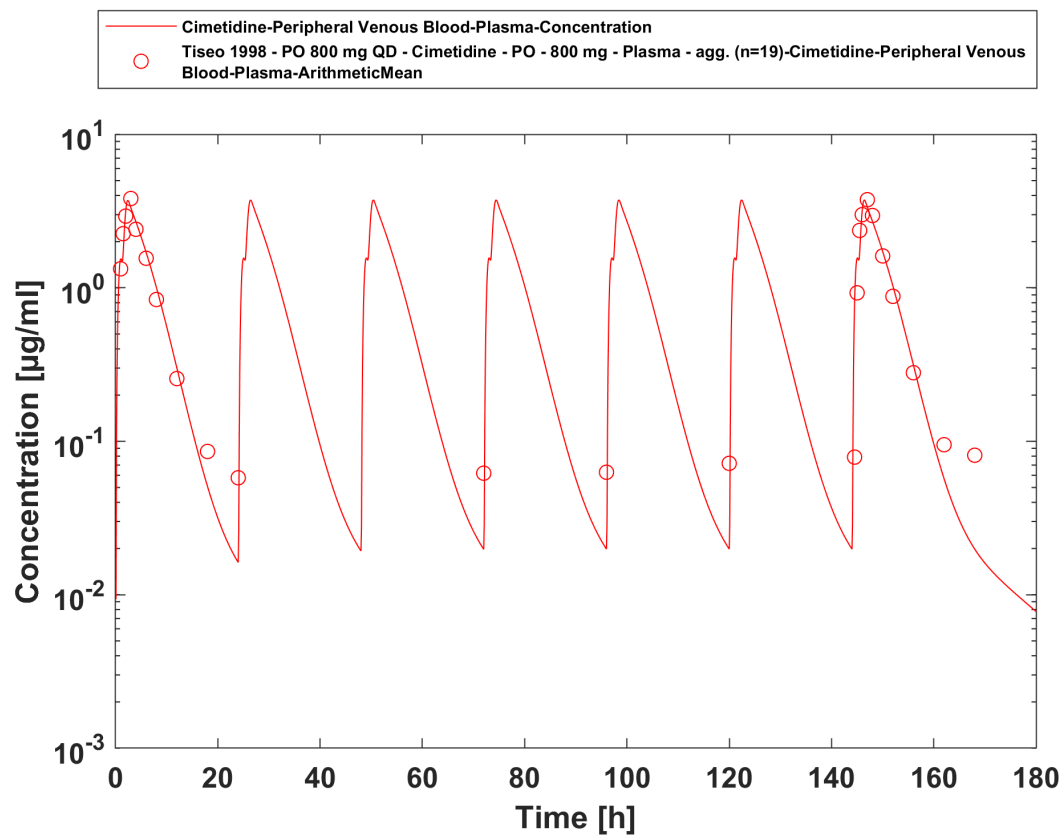
po 400 mg (tab), Somogyi 1981, n=8



po 400 mg (tab),Grahnen 1979, n=3



po 400 mg (tab),Grahnen 1979, n=3, urine



po 800 mg (tab) qd, Tiseo 1998, n=18

4 Conclusion

The herein presented PBPK model adequately describes the pharmacokinetics of cimetidine after intravenous and oral administration of single and multiple doses to healthy adults and peptic ulcer patients covering a broad dosing range from 100 to 800 mg. The established cimetidine PBPK model is verified for the use as a mild inhibitor of CYP3A4 drug in drug-drug interaction simulations.

5 References

Avdeef 2001 Avdeef A, Berger CM. pH-metric solubility. 3. Dissolution titration template method for solubility determination. *Eur J Pharm Sci.* 2001 Dec;14(4):281-91. doi: 10.1016/s0928-0987(01)00190-7. PMID: 11684402.

Barbhaiya 1995 Barbhaiya RH, Shukla UA, Greene DS. Lack of interaction between nefazodone and cimetidine: a steady state pharmacokinetic study in humans. *Br J Clin Pharmacol.* 1995 Aug;40(2):161-5. doi: 10.1111/j.1365-2125.1995.tb05771.x. PMID: 8562300; PMCID: PMC1365177.

Bodemar 1979 Bodemar G, Norlander B, Fransson L, Walan A. The absorption of cimetidine before and during maintenance treatment with cimetidine and the influence of a meal on the absorption of cimetidine--studies in patients with peptic ulcer disease. *Br J Clin Pharmacol.* 1979 Jan;7(1):23-31. doi: 10.1111/j.1365-2125.1979.tb00892.x. PMID: 760739; PMCID: PMC1429608.

Bodemar 1981 Bodemar G, Norlander B, Walan A. Pharmacokinetics of cimetidine after single doses and during continuous treatment. *Clin Pharmacokinet.* 1981 Jul-Aug;6(4):306-15. doi: 10.2165/00003088-198106040-00005. PMID: 7249489.

Burland 1975 Burland WL, Duncan WA, Hesselbo T, Mills JG, Sharpe PC, Haggie SJ, Wyllie JH. Pharmacological evaluation of cimetidine, a new histamine H₂-receptor antagonist, in healthy man. *Br J Clin Pharmacol.* 1975 Dec;2(6):481-6. doi: 10.1111/j.1365-2125.1975.tb00564.x. PMID: 9952; PMCID: PMC1402643.

D'Angio 1986 D'Angio R, Mayersohn M, Conrad KA, Bliss M. Cimetidine absorption in humans during sucralfate coadministration. *Br J Clin Pharmacol.* 1986 May;21(5):515-20. doi: 10.1111/j.1365-2125.1986.tb02834.x. PMID: 3755052; PMCID: PMC1401033.

Fee 1987 Fee JP, Collier PS, Howard PJ, Dundee JW. Cimetidine and ranitidine increase midazolam bioavailability. *Clin Pharmacol Ther.* 1987 Jan;41(1):80-4. doi: 10.1038/clpt.1987.13. PMID: 3802710.

Grahnén 1979 Grahnén A, von Bahr C, Lindström B, Rosén A. Bioavailability and pharmacokinetics of cimetidine. *Eur J Clin Pharmacol.* 1979 Nov;16(5):335-40. doi: 10.1007/BF00605632. PMID: 520401.

Greenblatt 1986 Greenblatt DJ, Locniskar A, Scavone JM, Blyden GT, Ochs HR, Harmatz JS, Shader RI. Absence of interaction of cimetidine and ranitidine with intravenous and oral midazolam. *Anesth Analg.* 1986 Feb;65(2):176-80. PMID: 2935051.

Ito 2012 Ito S, Kusuhara H, Yokochi M, Toyoshima J, Inoue K, Yuasa H, Sugiyama Y. Competitive inhibition of the luminal efflux by multidrug and toxin extrusions, but not basolateral uptake by organic cation transporter 2, is the likely mechanism underlying the pharmacokinetic drug-drug interactions caused by cimetidine in the kidney. *J Pharmacol Exp Ther.* 2012 Feb;340(2):393-403. doi: 10.1124/jpet.111.184986. Epub 2011 Nov 9. PMID: 22072731.

Kanto 1981 Kanto J, Allonen H, Jalonen H, Mäntylä R. The effect of metoclopramide and propantheline on the gastrointestinal absorption of cimetidine. *Br J Clin Pharmacol.* 1981 Jun;11(6):629-31. doi: 10.1111/j.1365-2125.1981.tb01184.x. PMID: 7272182; PMCID: PMC1402204.

Kuepfer 2016 Kuepfer L, Niederalte C, Wendl T, Schlender JF, Willmann S, Lippert J, Block M, Eissing T, Teutonico D. Applied Concepts in PBPK Modeling: How to Build a PBPK/PD Model. *CPT Pharmacometrics Syst Pharmacol.* 2016 Oct;5(10):516-531.

Larsson 1982 Larsson R, Erlanson P, Bodemar G, Walan A, Bertler A, Fransson L, Norlander B. The pharmacokinetics of cimetidine and its sulfoxide metabolite in patients with normal and impaired renal function. *Br J Clin Pharmacol*. 1982 Feb;13(2):163-70. doi: 10.1111/j.1365-2125.1982.tb01351.x. PMID: 7059413; PMCID: PMC1402003.

Lebert 1981 Lebert PA, Mahon WA, MacLeod SM, Soldin SJ, Fenje P, Vandenberghe HM. Ranitidine kinetics and dynamics. II. Intravenous dose studies and comparison with cimetidine. *Clin Pharmacol Ther*. 1981 Oct;30(4):545-50. doi: 10.1038/clpt.1981.201. PMID: 6269789.

Martinez 1999 Martínez C, Albet C, Agúndez JA, Herrero E, Carrillo JA, Márquez M, Benítez J, Ortiz JA. Comparative in vitro and in vivo inhibition of cytochrome P450 CYP1A2, CYP2D6, and CYP3A by H₂-receptor antagonists. *Clin Pharmacol Ther*. 1999 Apr;65(4):369-76. doi: 10.1016/S0009-9236(99)70129-3. PMID: 10223772.

Meyer 2012 Meyer M, Schneckener S, Ludewig B, Kuepfer L, Lippert J. Using expression data for quantification of active processes in physiologically based pharmacokinetic modeling. *Drug Metab Dispos*. 2012 May;40(5):892-901.

Mihaly 1984 Mihaly GW, Jones DB, Anderson JA, Smallwood RA, Louis WJ. Pharmacokinetic studies of cimetidine and ranitidine before and after treatment in peptic ulcer patients. *Br J Clin Pharmacol*. 1984 Jan;17(1):109-11. doi: 10.1111/j.1365-2125.1984.tb05010.x. PMID: 6318788; PMCID: PMC1463299.

Morgan 1983 Morgan DJ, Uccellini DA, Raymond K, Mihaly GW, Jones DB, Smallwood RA. The influence of duration of intravenous infusion of an acute dose on plasma concentrations of cimetidine. *Eur J Clin Pharmacol*. 1983;25(1):29-34. doi: 10.1007/BF00544010. PMID: 6617722.

Nishimura 2013 Nishimura M, Yaguti H, Yoshitsugu H, Naito S, Satoh T. Tissue distribution of mRNA expression of human cytochrome P450 isoforms assessed by high-sensitivity real-time reverse transcription PCR. *Yakugaku Zasshi*. 2003 May;123(5):369-75.

Ohta 2005 Ohta KY, Inoue K, Yasujima T, Ishimaru M, Yuasa H. Functional characteristics of two human MATE transporters: kinetics of cimetidine transport and profiles of inhibition by various compounds. *J Pharm Pharm Sci*. 2009;12(3):388-96. doi: 10.18433/j3r59x. PMID: 20067714.

PK-Sim Ontogeny Database Version 7.3 (<https://github.com/Open-Systems-Pharmacology/OSPSuite.Documentation/blob/38cf71b384cfc25cfa0ce4d2f3adddfd32757e13b/PK-Sim%20Ontogeny%20Database%20Version%207.3.pdf>)

Salonen 1986 Salonen M, Aantaa E, Aaltonen L, Kanto J. Importance of the interaction of midazolam and cimetidine. *Acta Pharmacol Toxicol (Copenh)*. 1986 Feb;58(2):91-5. doi: 10.1111/j.1600-0773.1986.tb00076.x. PMID: 2939688.

Smith 1984 Smith MS, Benyunes MC, Bjornsson TD, Shand DG, Pritchett EL. Influence of cimetidine on verapamil kinetics and dynamics. *Clin Pharmacol Ther*. 1984 Oct;36(4):551-4. doi: 10.1038/clpt.1984.218. PMID: 6478741.

Somogyi 1981 Somogyi A, Thielscher S, Gugler R. Influence of phenobarbital treatment on cimetidine kinetics. *Eur J Clin Pharmacol*. 1981;19(5):343-7. doi: 10.1007/BF00544584. PMID: 7238562.

Somogyi 1983 Somogyi A, Gugler R. Clinical pharmacokinetics of cimetidine. *Clin Pharmacokinet*. 1983 Nov-Dec;8(6):463-95. doi: 10.2165/00003088-198308060-00001. PMID: 6418428.

Tahara 2005 Tahara H, Kusuhara H, Endou H, Koepsell H, Imaoka T, Fuse E, Sugiyama Y. A species difference in the transport activities of H₂ receptor antagonists by rat and human renal organic anion and cation transporters. *J Pharmacol Exp Ther*. 2005 Oct;315(1):337-45. doi: 10.1124/jpet.105.088104. Epub 2005 Jul 8. PMID: 16006492.

Taylor 1978 Taylor DC, Cresswell PR, Bartlett DC. The metabolism and elimination of cimetidine, a histamine H₂-receptor antagonist, in the rat, dog, and man. *Drug Metab Dispos.* 1978 Jan-Feb;6(1):21-30. PMID: 23270.

Tiseo 1998 Tiseo PJ, Perdomo CA, Friedhoff LT. Concurrent administration of donepezil HCl and cimetidine: assessment of pharmacokinetic changes following single and multiple doses. *Br J Clin Pharmacol.* 1998 Nov;46 Suppl 1(Suppl 1):25-9. doi: 10.1046/j.1365-2125.1998.0460s1025.x. PMID: 9839762; PMCID: PMC1873814.

Umehara 2007 Umehara KI, Iwatsubo T, Noguchi K, Kamimura H. Functional involvement of organic cation transporter1 (OCT1/Oct1) in the hepatic uptake of organic cations in humans and rats. *Xenobiotica.* 2007 Aug;37(8):818-31. doi: 10.1080/00498250701546012. PMID: 17701831.

Walkenstein 1978 Walkenstein SS, Dubb JW, Randolph WC, Westlake WJ, Stote RM, Intoccia AP. Bioavailability of cimetidine in man. *Gastroenterology.* 1978 Feb;74(2 Pt 2):360-5. PMID: 620910.

Willmann 2007 Willmann S, Höhn K, Edginton A, Sevestre M, Solodenko J, Weiss W, Lippert J, Schmitt W. Development of a physiology-based whole-body population model for assessing the influence of individual variability on the pharmacokinetics of drugs. *J Pharmacokinet Pharmacodyn.* 2007, 34(3): 401-431.

Wishart 2006 Wishart DS, Knox C, Guo AC, Shrivastava S, Hassanali M, Stothard P, Chang Z, Woolsey J. DrugBank: a comprehensive resource for in silico drug discovery and exploration. *Nucleic Acids Res.* 2006 Jan 1;34(Database issue):D668-72. doi: 10.1093/nar/gkj067. PMID: 16381955; PMCID: PMC1347430.

Wrighton 1994 Wrighton SA, Ring BJ. Inhibition of human CYP3A catalyzed 1'-hydroxy midazolam formation by ketoconazole, nifedipine, erythromycin, cimetidine, and nizatidine. *Pharm Res.* 1994 Jun;11(6):921-4. doi: 10.1023/a:1018906614320. PMID: 7937537.