ORD MANUSCRIPT COVER SHEET

**Title:** Great Title about *invivoPKfit* and the CvTdb

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Labs/Centers: CCTE

**External collaborators:**

* Showa Pharmaceutical University

**Internal Peer Reviewers:** TBD

**Target Journal:** Possibly: Computational Toxicology, Journal of Pharmacokinetics and Pharmacodynamics, Journal of Statistical Software

**Four bullet point summary:**

* words
* words
* words
* words

**One sentence description:** This collaborative trial demonstrates that …

**Chemicals Involved:** ~100 CvTdb chemicals with oral and/or intravenous administration routes

**Great Title about *invivoPKfit* and the CvTdb**

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# Abstract

The Concentration versus Time Database (CvTdb) is a public repository of toxicokinetic (TK) data developed by the Center for Computational Toxicology and Exposure at the U.S. Environmental Protection Agency (EPA). The CvTdb contains manually curated experimental time-course data of chemical compound concentrations in body fluids and tissues, along with descriptive metadata, from hundreds of publications. The CvTdb is a “living” repository, to which new data can be added as they become available. Recent efforts have been made to 1) Expand the scope of in-house data curation and 2) Integrate sets of CvT data shared by other institutions. To standardize the format of data extracted from numerous publications, by several curators, we developed an R Shiny application that checks completed template curation files for both format and data correctness. New data sets include those from collaborators of Showa Pharmaceutical University with more than 200 compounds. Standardized data reporting allows for analysis of TK trends across studies. For example, we observe that among replicate observations (those with common chemical/reference/dose/route/timepoint data), 85.9% are within two-fold of the mean concentration. When the data are subset by route, we observe that 84.0% of oral administration observations, and 89.7% of intravenous administration observations, are within two-fold of the mean concentration. Data are systematically analyzed using invivoPKfit, an R package that fits standardized 1- and 2- compartmental TK models to all data associated with a particular compound, including data that spans multiple references. We used invivoPKfit to fit models and estimate TK parameters, such as volume of distribution and elimination half-life. Overall, the CvTdb serves as a platform for assessing TK trends across a large and standardized set of data, as well as for calibrating and validating TK models.

# Introduction

## General

* Toxicokinetics (TK) describes the absorption, distribution, metabolism, and excretion (ADME) of a chemical compound in the body as a function of time
* TK allows for the prediction of internal tissue concentrations as a function of chemical exposure and provides critical information for the assessment of a risk posed by a chemical to public health
* This data is referred to as “concentration *versus* time” (CvT) data
* TK involves mathematically describing CvT data such that extrapolation is possible

## CvTdb and *invivoPKfit* can Help

* There is a need for a standardized and reproducible approach to fitting pharmacokinetic models
* A standardized approach would make one data set for a chemical intercomparable with other data sets
* We address this problem with two tools – the Concentration *versus* Time Database (CvTdb) and the R package, *invivoPKfit*
* The CvTdb is a public repository of standardized TK data developed by the Center for Computational Toxicology and Exposure (CCTE) at the U.S. Environmental Protection Agency (EPA)
* *invivoPKfit* is an R package developed by CCTE to fit generic 1- and 2-compartment models to the diverse set of data within the CvTdb
* We estimate model parameters for the CvTdb using *invivoPKfit* and compare them to those reported in Lombardo, which compiles parameters from models tailored to specific sets of data

## Review on other packages

* There are other software packages to fit models to CvT data, notably *PK*, *pmxTools*, and *PKconverter*
* *invivoPKfit* differentiates itself from these other tools by allowing for meta-analyses between data of the same chemical but from different studies…
* Maybe a table describing them?

# Methods

## *invivoPKfit* Models

* *invivoPKfit* fits generic 1- and 2- compartment models to all data
* These were copied from the Boomer page, but can be cited from

## How *invivoPKfit* Finds Solutions to Models

* *invivoPKfit* estimates the most likely parameter values for generic 1- and 2- compartment models describing the TK of a compound given CvT observations
* We use the R package *optimx* to optimize the log-likelihood equation and estimates the parameters and sigma
* Confidence intervals on the estimated parameters were calculated using the Hessian of the likelihood function to estimate standard deviations of the log parameter (Bartlett, 1953a,b).

## *In vivo* Data

EPA has developed a public database of concentration vs. time data for building, calibrating, and evaluating TK models [27]. Curation and development of the database are ongoing, but when this study began there were 101 chemicals with either rat or human *in vivo* blood or plasma concentration vs. time data. The *in vivo* measured concentration vs. time values are available as Supplemental Table 2.

## Compartmental Model Fits

For each chemical with CvT data, parameters were estimated for empirical one- and two-compartment toxicokinetic models using R package “invivoPKfit” (<https://github.com/USEPA/CompTox-ExpoCast-invivoPKfit>). Between the one- and two-compartment models, the one with the lower Akaike Information Criterion (AIC) value – indicating model parsimony – was selected [37]. The empirical model fit was then used as a “best case” prediction scenario for comparison with PBTK parameterize by *in vitro* or QSPR predictions.

For both models a half-life was calculated from the terminal elimination rate as thalf = ln(2)/kelim. For the two-compartment model the volume of distribution at steady-state was used as Vd. For both models, clearance was calculated as Cltot = Vd \* kelim. The estimated TK parameters for both models are provided as Supplemental Table 3.

# Results

## Evaluation Chemicals and Predictions

There were 102 chemicals present in the CvTdb (Sayre, 2020) as of September 2019 that had plasma concentration data following either oral or intravenous doses given to rats or humans. These chemicals included: 57 from the Toxic Substances Control Act (TSCA) active inventory [38], 20 pharmaceuticals, 24 pesticides, 99 that are found in consumer products, 7 per- and poly-fluorinated substances (PFAS) [39], and 64 that are part of the ToxCast screening program.

Of the 102 chemicals, there were 10 chemicals that could only be predicted by OPERA and were omitted from the rest of the analysis (Supplemental Table 5). Two chemicals, Oxoacetic acid--water (1/1) and Nitrite have CvT data but do not have either measured or predicted values for both Clint and fup from any model to date.

The data for each chemical was fit using maximum likelihood estimation to one and two compartment empirical pharmacokinetic models, with separate fits for each combination of compound and species for which there were data. Maximum likelihood estimates could not be obtained for either model for 8 chemicals (listed in Supplemental Table 4). These chemicals were withheld from subsequent analysis. For each remaining chemical the better of the one or two compartment models was used on the basis of model parsimony.

Eliminating chemicals where the CvT data could not be described by an empirical model or for which there were only one QSPR that could make predictions left 83 chemicals with *in vivo* CvT data and 68 chemicals with *in vitro* measure fup and Clint.

For each QSPR we removed predictions where the predicted values for a given chemical were within 1% for both fup and Clint assuming these values reflected the chemical data present in the training set and the model method allowing for recall of the measurements. This only affected 21 chemicals as predicted by OPERA.

We summarize the chemical-specific properties and predictions in Figure 1. In Figure 1 similar chemicals (rows) and properties/predictions (columns) are clustered together based on Euclidean distance. All properties/predictions were centered (mean changed to zero) and scaled (divided by standard deviation) such that the value reflects the number of standard deviations from the mean. Interestingly, the first division between clusters in Figure 1 places all the Clint measurements and predictions on the one side and all the fup measurements and predictions on the other. The physico-chemical properties are divided between those two clusters, with Octanol:Water (partition coefficient, PC), Octanol:Air PC, Molecular Weight, Boiling Point, and Melting Point all clustering with Clint. Water solubility, vapor pressure, and the Henry’s law constant all clustered with fup.

## Level 3 Analysis

We then proceed on to the third level of evaluation, in which we use the QSPR predictions to predict toxicokinetic summary parameters – volume of distribution (Vd), half-life for elimination from the body (thalf), and whole-body clearance (Cltot) – and compare the predictions to the values estimated from the empirical fits to the CvT data. The values predicted for each method are provided in Supplemental Table 8.

In Figure 9 we examine predicted vs. observed thalf. As summarized in Table 4, none of the models are very successful – the highest coefficient of variation is 0.15 for IFS-QSAR, while QSARINS-Chem and HTTK with ADMET both had an R2 of 0.11. The RMSLE for all models, including HTTK with y-randomized data, was just above 1 (a factor of 10x).

The models were distinctly better than the y-randomization for predicting Vd. As shown in Figure 10 and summarized in Table 4, the HTTK algorithm for predicting Vd [34] when used with y-randomized data had no skill. The models performed similarly to the measured *in vitro* data when used with the Vd algorithm – R2 ranged from 0.10 to 0.16 with the measured data being the worst. For all models the RMSLE again indicated a factor of 10x.

In Figure 11 we examine predictions for CLtot, which depends on both elimination rate (inverse of thalf) and Vd. As summarized in Table 4 the y-randomized predictions reassuringly have no skill at predicting *in vivo* clearance, while the combination of HTTK and ADMET predicted values had the most, with a R2 of 0.32 and a RMSLE indicating a factor of 17x. Both IFS-QSAR and QSARINS-Chem had comparable R2 of 0.25 and 0.2 (respectively) and an RMSLE indicating a factor of ~20x. The other QSPRs performed about as well as using the *in vitro* measured data.

# Discussion

# Acknowledgements

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# Conflict of Interest

Please declare any COI here

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The views expressed in this publication are those of the authors and do not necessarily represent the views or policies of the U.S. EPA. Reference to commercial products or services does not constituteendorsement.

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# Figures

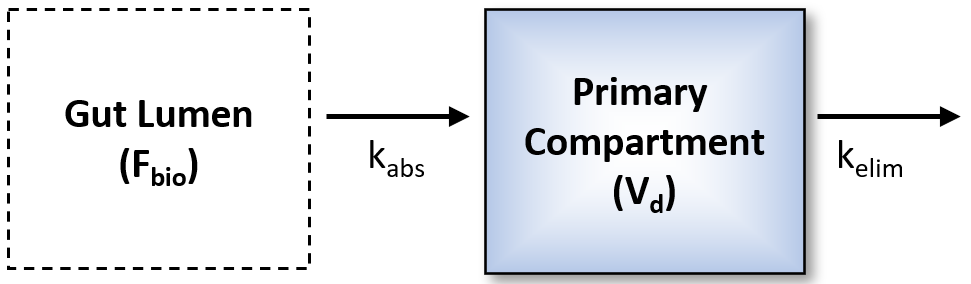


Figure 1: test

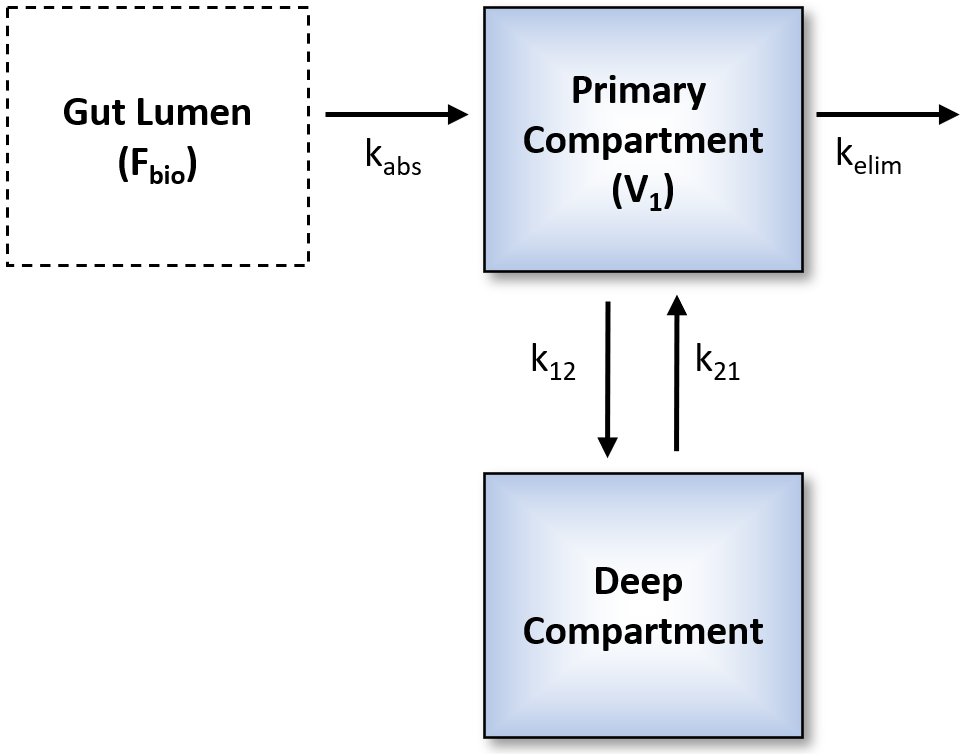


Figure 2: test

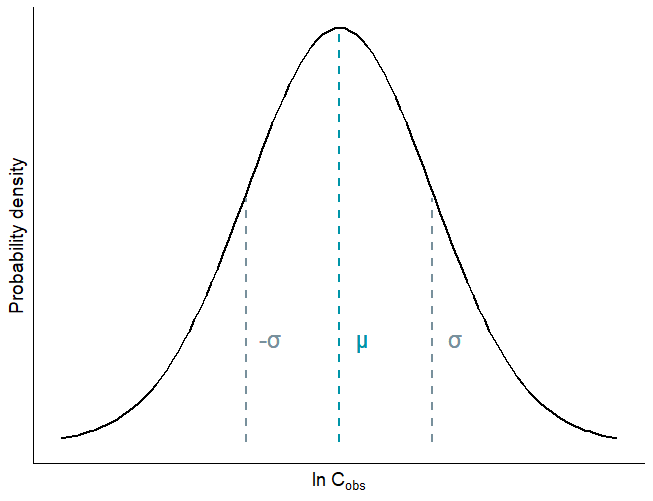


Figure 3: Log-normal distribution. Here are more details.

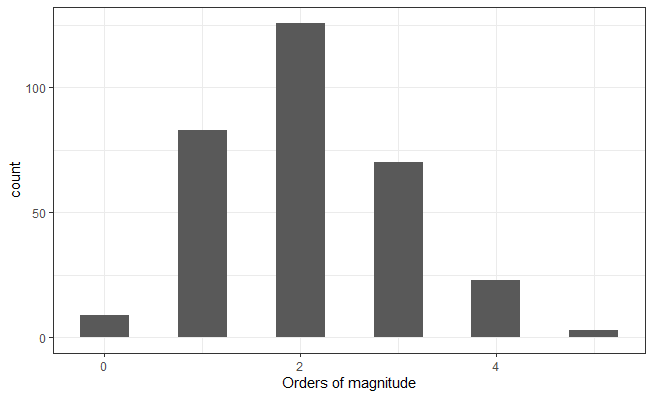


Figure : Orders of magnitude difference for individual series. The "2" average is a case for the taking the log-normal distribution, as otherwise there would be a bias toward higher numbers.

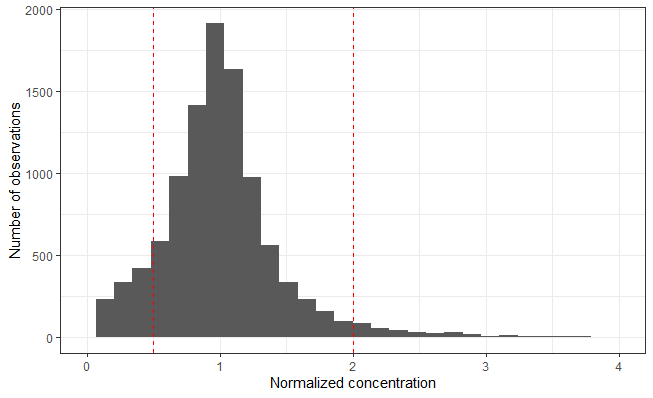


Figure 5: The CvTdb data was grouped into unique combinations of reference/route/dose/timepoint. Groups without replicate timepoints were removed. If all replicate timepoints corresponded to NA (non-detect) concentration values, those data were removed. If some, but not all replicate timepoints corresponded to NA concentration values, those NA concentration values were set equal to the limit of quantitation (LOQ). If the LOQ was unknown, it was assumed to equal 0.45 of minimum concentration value present. Each concentration value was normalized by the mean concentration value of each group. The red dashed lines mark the upper and lower bounds of data within two-fold of the mean (0.5 and 2).

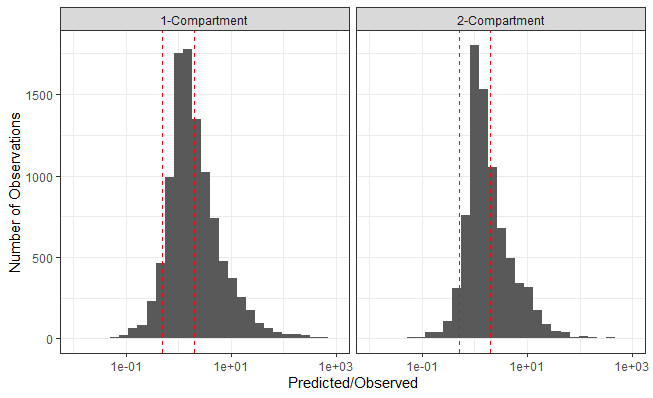


Figure 6: Concentrations were predicted by applying estimated model parameters to the timepoints present in the data. Predicted concentrations were normalized by observed concentrations at their corresponding timepoints. The red dashed lines mark the lower and upper bounds of predicted data within two-fold of observed data.

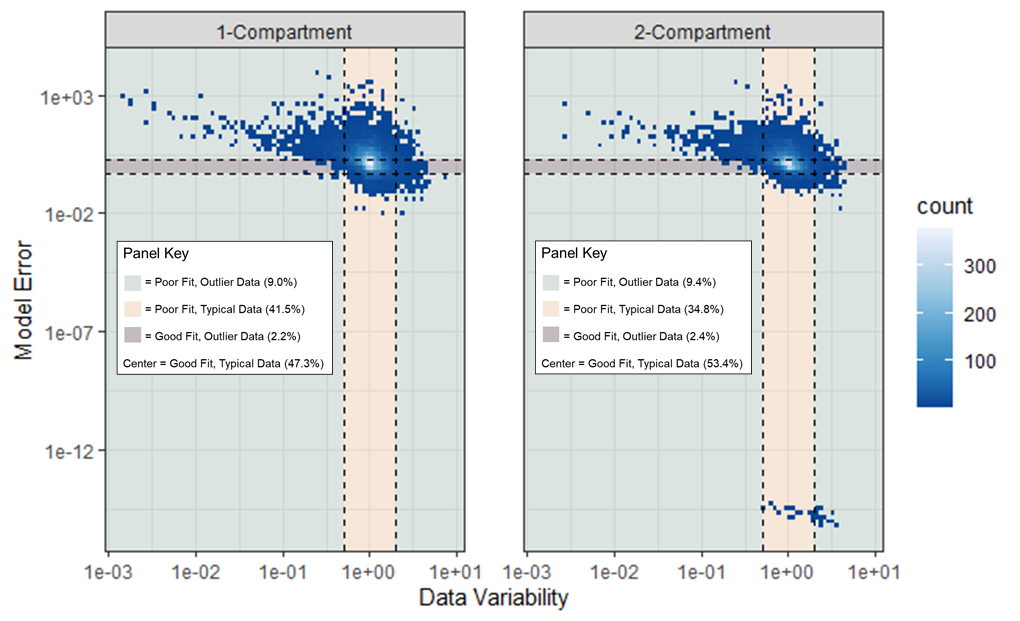
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Figure 7: Density plot of Model Error (predicted concentration / observed concentration) versus Data Variability (concentration normalized by mean concentration of replicate timepoints; see Figure 3). Colored panels define the ‘goodness’ of both the data and the model, based on whether those points are within two-fold of the mean (x-axis) and two-fold of the observed concentration (y-axis). The “Center” panel (the densest area) represents data where points correspond to both a Data Variability within two-fold of the mean and a Model Error within two-fold of the observed concentrations. Under the 1-Compartment Model, 47.3% of points lie in the “Center” area. Under the 2-Compartment Model, 53.4% of points lie in the “Center” area.

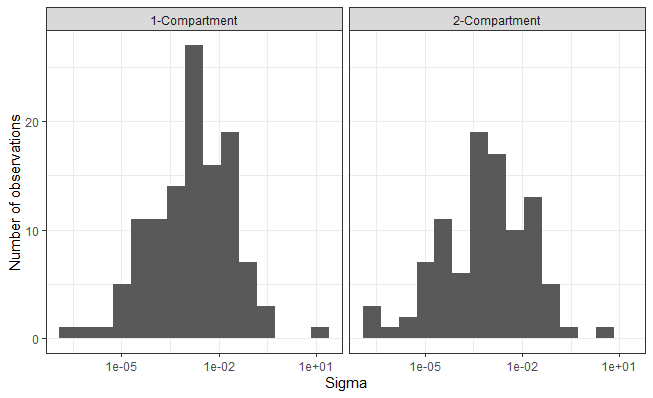


Figure :

# Tables

# Supplemental Tables

Supplemental Table 1: Empirical (one and two compartment model) toxicokinetic parameter estimates

Supplemental Table 3: Empirical (one and two compartment model) toxicokinetic parameter estimates

|  |  |  |  |  |  |  |  |  |  |
| --- | --- | --- | --- | --- | --- | --- | --- | --- | --- |
| **Compound** | **CAS** | **Species** | **Reference** | **AIC\_1comp** | **AIC\_2comp** | **Model** | **Vdist** | **kelim** | **halflife** |
| 1,2-dichloroethane | 107-06-2 | rat | 18 | -279.8 | -275.9 | 1Comp | 3650 | 0.3431 | 2.02 |
| 1,4-dioxane | 123-91-1 | rat | 24, 6 | -367.4 | -368.1 | 2Comp | 658.8 | 0.02368 | 29.27 |
| 1-chloro-2-propanol | 127-00-4 | rat | 176 | -1391 | -1392 | 2Comp | 2382 | 1.993 | 0.3479 |
| 2,3,7,8-tetrachlorodibenzo-p-dioxin | 1746-01-6 | rat | 177 | -108.1 | -107 | 1Comp | 1452 | 0.001888 | 367.2 |
| 2,4-dichlorophenoxyacetic acid | 94-75-7 | rat | 164, 192 | -1670 | -1733 | 2Comp | 339.4 | 1.738 | 0.3989 |
| 2-hydroxy-4-methoxybenzophenone | 131-57-7 | rat | 129 | -4667 | -4683 | 2Comp | 14350 | 0.3336 | 2.078 |
| 2-methylimidazole | 693-98-1 | rat | 150 | -565.2 | -593 | 2Comp | 2440 | 0.02029 | 34.17 |
| 2-methyltetrahydrofuran | 96-47-9 | rat | 175 | -3240 | -3240 | 1Comp | 812.7 | 0.3836 | 1.807 |
| 4-methylimidazole | 822-36-6 | rat | 151 | -716.5 | -755.5 | 2Comp | 1274 | 0.7412 | 0.9351 |
| acrylonitrile | 107-13-1 | rat | 43 | -260.5 | -285.6 | 2Comp | 747.6 | 9.896 | 0.07004 |
| alachlor | 15972-60-8 | rat | 192 | -965.6 | -962.4 | 1Comp | 173200 | 0.07371 | 9.404 |
| alpha-thujone | 546-80-5 | rat | 130 | -2351 | -2390 | 2Comp | 10510 | 0.7298 | 0.9498 |
| alprazolam | 28981-97-7 | rat | 193 | -529.1 | NA | 1Comp | 2671 | 2.273 | 0.305 |
| anthraquinone | 84-65-1 | rat | 166 | -2502 | NA | 1Comp | 3786 | 0.133 | 5.21 |
| antipyrine | 60-80-0 | rat | 197 | -101.1 | -98.84 | 1Comp | 1133 | 0.5825 | 1.19 |
| benzo(a)pyrene | 50-32-8 | rat | 36, 38, 46, 59 | -777.5 | -884.6 | 2Comp | 3258 | 0.4873 | 1.422 |
| benzophenone | 119-61-9 | rat | 171 | -5427 | -5565 | 2Comp | 12870 | 0.2211 | 3.135 |
| bis 2-chloroethoxy methane | 111-91-1 | rat | 155 | -1090 | -1172 | 2Comp | 2323 | 3.444 | 0.2013 |
| bisphenol a | 80-05-7 | rat | 192 | -201.9 | -204.4 | 2Comp | 1929 | 4.372 | 0.1585 |
| boscalid | 188425-85-6 | rat | 192 | -798.1 | -855 | 2Comp | 11250 | 1.098 | 0.631 |
| bosentan | 147536-97-8 | rat | 198 | -225.5 | NA | 1Comp | 3156 | 0.2439 | 2.841 |
| bromochloroacetic acid | 5589-96-8 | rat | 132 | NA | -2067 | 2Comp | 3939 | 0.3587 | 1.932 |
| bromodichloromethane | 75-27-4 | rat | 154 | -923.6 | -977.7 | 2Comp | 19190 | 2.019 | 0.3433 |
| carbaryl | 63-25-2 | rat | 192 | -465.9 | -260.3 | 1Comp | 41640 | 4.485 | 0.1545 |
| carbendazim | 10605-21-7 | rat | 201 | -68.57 | -64.57 | 1Comp | 33150 | 0.245 | 2.829 |
| chloridazon | 1698-60-8 | rat | 192 | -893.7 | NA | 1Comp | 6923 | 0.08218 | 8.434 |
| chlorpyrifos | 2921-88-2 | rat | 202 | -364.5 | -366.8 | 2Comp | 43220 | 0.5533 | 1.253 |
| cyclanilide | 113136-77-9 | rat | 192 | -1223 | -1277 | 2Comp | 361.7 | 0.2194 | 3.159 |
| cyclosporin a | 59865-13-3 | rat | 203, 204 | -271.8 | -308.3 | 2Comp | 1605 | 0.3603 | 1.924 |
| diazinon-o-analog | 962-58-3 | rat | 192 | -884.7 | -904.1 | 2Comp | 271100 | 8.04E-05 | 8619 |
| dichloroacetic acid | 79-43-6 | rat | 134 | -699.9 | -699.6 | 1Comp | 528.2 | 1.767 | 0.3923 |
| diltiazem | 34933-06-7 | rat | 207 | -236.3 | -258.5 | 2Comp | 2745 | 3.392 | 0.2044 |
| dimethenamid | 87674-68-8 | rat | 192 | -911.5 | -964 | 2Comp | 159600 | 0.8059 | 0.8601 |
| di-n-butyl phthalate | 84-74-2 | rat | 135 | -49.46 | -45.46 | 1Comp | 1916 | 5.349 | 0.1296 |
| dl-camphor | 76-22-2 | rat | 165 | -851.9 | -895.5 | 2Comp | 6283 | 3.609 | 0.192 |
| emodin | 518-82-1 | rat | 160 | -713.7 | -668.6 | 1Comp | 715.3 | 2.152 | 0.3221 |
| etoxazole | 153233-91-1 | rat | 192 | -1539 | -1626 | 2Comp | 31930 | 0.711 | 0.9749 |
| fenarimol | 60168-88-9 | rat | 192 | -806 | -815 | 2Comp | 8003 | 0.2633 | 2.632 |
| flufenacet | 142459-58-3 | rat | 192 | -1414 | NA | 1Comp | 62440 | 0.03117 | 22.24 |
| fluorotelomer alcohol 8+2 | 678-39-7 | rat | 131 | -3425 | NA | 1Comp | 11420 | 0.1106 | 6.266 |
| formamide | 75-12-7 | rat | 172 | -1936 | -1934 | 1Comp | 460.5 | 0.03309 | 20.95 |
| formetanate hydrochloride | 23422-53-9 | rat | 192 | -101 | -97.04 | 1Comp | 25240 | 8.31E-08 | 8341000 |
| free carbon disulfide | 75-15-0 | rat | 167 | -268.3 | -280.3 | 2Comp | 3352 | 7.072 | 0.09802 |
| gemfibrozil | 25812-30-0 | rat | 163 | -1367 | -1393 | 2Comp | 1266 | 0.2384 | 2.907 |
| glyoxylic acid monohydrate | 563-96-2 | rat | 156 | -325.9 | -368.1 | 2Comp | 898.4 | 13.42 | 0.05164 |
| hexachlorobenzene | 118-74-1 | rat | 180 | -8917 | -3018 | 1Comp | 4790 | 0.000306 | 2265 |
| hexobarbital | 15307-86-5 | rat | 206, 209 | NA | -189.5 | 2Comp | 2152 | 0.3759 | 1.844 |
| ibuprofen | 15687-27-1 | rat | 210 | -185.5 | -211.9 | 2Comp | 687 | 0.8539 | 0.8118 |
| imazalil | 35554-44-0 | rat | 192 | -405.6 | -436.5 | 2Comp | 16370 | 3.427 | 0.2023 |
| imipramine | 50-49-7 | rat | 211, 212 | -215 | -228.6 | 2Comp | 39970 | 0.3045 | 2.276 |
| isoeugenol | 97-54-1 | rat | 157 | -4214 | -4493 | 2Comp | 25090 | 1.949 | 0.3556 |
| l-ephedrine | 299-42-3 | rat | 136 | -1884 | -1891 | 2Comp | 16250 | 0.4206 | 1.648 |
| methanol | 67-56-1 | rat | 17 | -64.13 | -60.13 | 1Comp | 992 | 0.3046 | 2.275 |
| methyl tert-butyl ether | 1634-04-4 | human | 51 | -355.9 | -363 | 2Comp | 893.1 | 1.343 | 0.5161 |
| methylene chloride | 75-09-2 | rat | 18 | -198.5 | -197.5 | 1Comp | 1883 | 0.4908 | 1.412 |
| methyleugenol | 93-15-2 | rat | 158, 170 | -5782 | -5836 | 2Comp | 3780 | 0.3344 | 2.073 |
| midazolam | 59467-70-8 | rat | 217 | -285.3 | NA | 1Comp | 3287 | 1.18 | 0.5876 |
| naphthalene | 91-20-3 | rat | 169 | -1721 | -1820 | 2Comp | 4317 | 1.104 | 0.6277 |
| nilvadipine | 75530-68-6 | rat | 219 | -570.8 | -572.6 | 2Comp | 5910 | 0.3737 | 1.855 |
| nitrite | 14797-65-0 | rat | 137 | -916.3 | -916.5 | 2Comp | 2766 | 0.4414 | 1.57 |
| novaluron | 116714-46-6 | rat | 192 | -1258 | -1282 | 2Comp | 7105 | 0.05488 | 12.63 |
| octylphenol | 140-66-9 | rat | 60 | -1465 | -1557 | 2Comp | 49920 | 0.3106 | 2.231 |
| ondansetron | 99614-02-5 | rat | 220 | -850.4 | -894.2 | 2Comp | 617.2 | 30.69 | 0.02259 |
| oxazepam | 604-75-1 | rat | 159 | -1789 | -1941 | 2Comp | 13120 | 0.1889 | 3.67 |
| oxymetholone | 434-07-1 | rat | 168 | -688.5 | NA | 1Comp | 5639 | 0.1631 | 4.251 |
| pentachlorophenol, purified | 87-86-5 | rat | 161 | -1585 | NA | 1Comp | 82.06 | 0.07683 | 9.021 |
| perfluorodecanoic acid | 335-76-2 | rat | 147 | -2278 | -2361 | 2Comp | 306.2 | 0.000804 | 862.6 |
| perfluorohexane-1-sulphonic acid â€“ potassium salt | 3871-99-6 | rat | 143 | -1153 | NA | 1Comp | 350.9 | 0.001978 | 350.5 |
| perfluorooctane sulfonate | 45298-90-6 | rat | 148 | -1869 | -1907 | 2Comp | 278.6 | 0.000965 | 718 |
| permethrin | 52645-53-1 | rat | 192 | -723.3 | NA | 1Comp | 64800 | 0.1253 | 5.53 |
| phenacetin | 62-44-2 | human | 14 | -715.9 | NA | 1Comp | 1502 | 0.4745 | 1.461 |
| phenacetin | 62-44-2 | rat | 1, 221, 62, 63, 67 | -355.9 | -385 | 2Comp | 1865 | 0.6633 | 1.045 |
| phenolphthalein | 77-09-8 | rat | 139 | -834.6 | -886.5 | 2Comp | 1408 | 1.153 | 0.601 |
| phenytoin | 57-41-0 | rat | 222, 223 | -256.1 | -262.5 | 2Comp | 3687 | 0.1736 | 3.993 |
| potassium perfluorobutane sulfonate | 29420-49-3 | rat | 144 | -1413 | -1452 | 2Comp | 204.1 | 0.2538 | 2.731 |
| primidone | 125-33-7 | rat | 140 | -463 | -459.9 | 1Comp | 1109 | 0.2159 | 3.211 |
| propamocarb hydrochloride | 25606-41-1 | rat | 192 | -595.7 | -603.1 | 2Comp | 8878 | 2.974 | 0.2331 |
| propyzamide | 23950-58-5 | rat | 192 | -1635 | NA | 1Comp | 9560 | 0.1086 | 6.383 |
| pyridine | 110-86-1 | rat | 149, 162 | -2362 | -2369 | 2Comp | 316.2 | 0.02765 | 25.07 |
| pyrithiobac sodium | 123343-16-8 | rat | 192 | -604 | NA | 1Comp | 678.3 | 0.13 | 5.331 |
| resmethrin | 10453-86-8 | rat | 192 | -794.5 | -823.1 | 2Comp | 48620 | 1.48 | 0.4683 |
| s-bioallethrin | 28434-00-6 | rat | 192 | -553.5 | -555.1 | 2Comp | 33980 | 0.7573 | 0.9152 |
| simazine | 122-34-9 | rat | 192 | -651.3 | -650 | 1Comp | 3592 | 1.939 | 0.3575 |
| solvent red1 | 1229-55-6 | rat | 66 | -275.9 | -281 | 2Comp | 2154 | 1.058 | 0.6552 |
| tamoxifen | 10540-29-1 | rat | 141 | -357.8 | 282 | 1Comp | 605100 | 14.33 | 0.04837 |
| tert-amyl methyl ether | 994-05-8 | human | 51 | -342.8 | -365.5 | 2Comp | 5393 | 1.04 | 0.6662 |
| tetrachloroethylene | 127-18-4 | rat | 4 | -81.9 | -79.47 | 1Comp | 12730 | 0.08628 | 8.034 |
| tetralin | 119-64-2 | rat | 152 | -1825 | -2004 | 2Comp | 6321 | 1.725 | 0.4019 |
| thiodiglycolic acid | 123-93-3 | rat | 155 | -442.9 | -489 | 2Comp | 976.6 | 4.789 | 0.1447 |
| tolbutamide | 64-77-7 | rat | 225, 226 | -139.7 | NA | 1Comp | 182.9 | 0.1618 | 4.284 |
| trichloroethylene | 79-01-6 | rat | 18, 29 | -250.9 | -313 | 2Comp | 318.8 | 11.58 | 0.05988 |
| triclosan | 3380-34-5 | rat | 192 | -334.7 | -326.2 | 1Comp | 617.9 | 0.06498 | 10.67 |
| valproic acid | 99-66-1 | rat | 227, 228 | -201.8 | NA | 1Comp | 220.9 | 1.101 | 0.6294 |
| wyeth-14643 | 50892-23-4 | rat | 173 | -1050 | NA | 1Comp | 448.8 | 0.3273 | 2.118 |

Supplemental Table 4: Chemicals that could not be fit by either a one- or two-compartment model using R package invivoPKfit

|  |  |  |
| --- | --- | --- |
| **DTXSID** | **PREFERRED\_NAME** | **CASRN** |
| DTXSID0021125 | Phenolphthalein | 77-09-8 |
| DTXSID2020139 | Benzo(a)pyrene | 50-32-8 |
| DTXSID2021103 | Pentachloroanisole | 1825-21-4 |
| DTXSID5032442 | Imidacloprid | 138261-41-3 |
| DTXSID8021359 | Tolbutamide | 64-77-7 |
| DTXSID8022292 | Permethrin | 52645-53-1 |
| DTXSID8023393 | Ondansetron | 99614-02-5 |
| DTXSID9032329 | Bensulide | 741-58-2 |