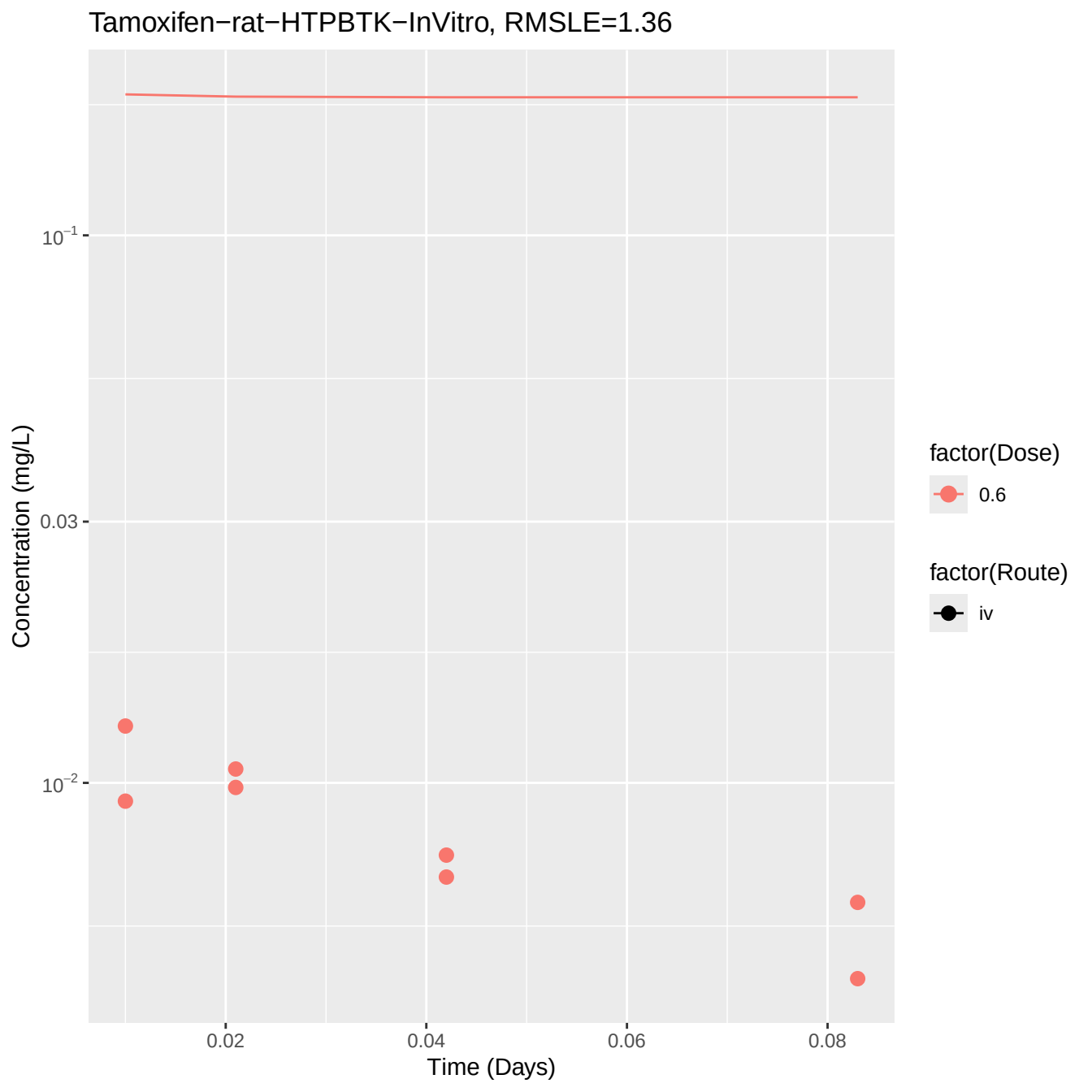
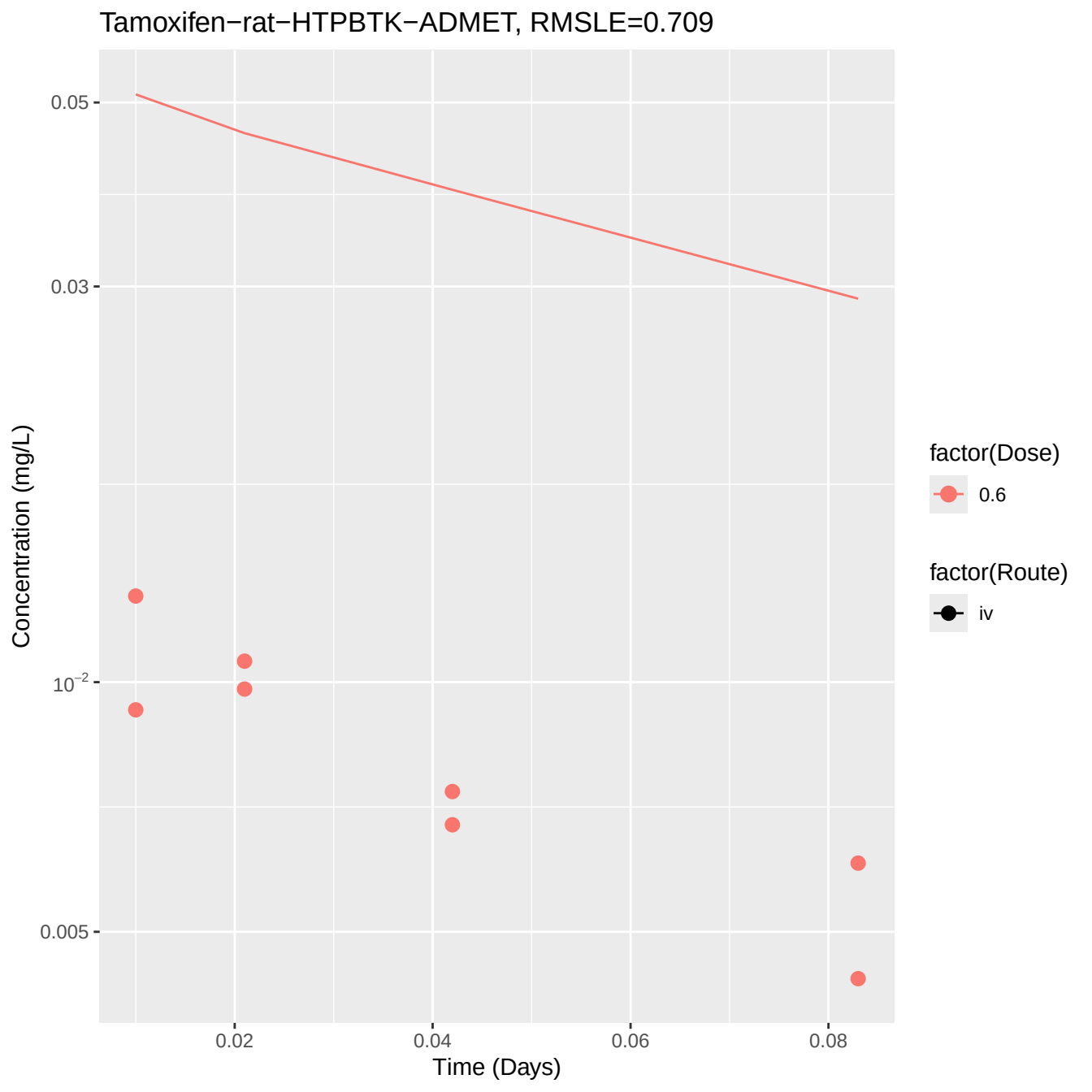
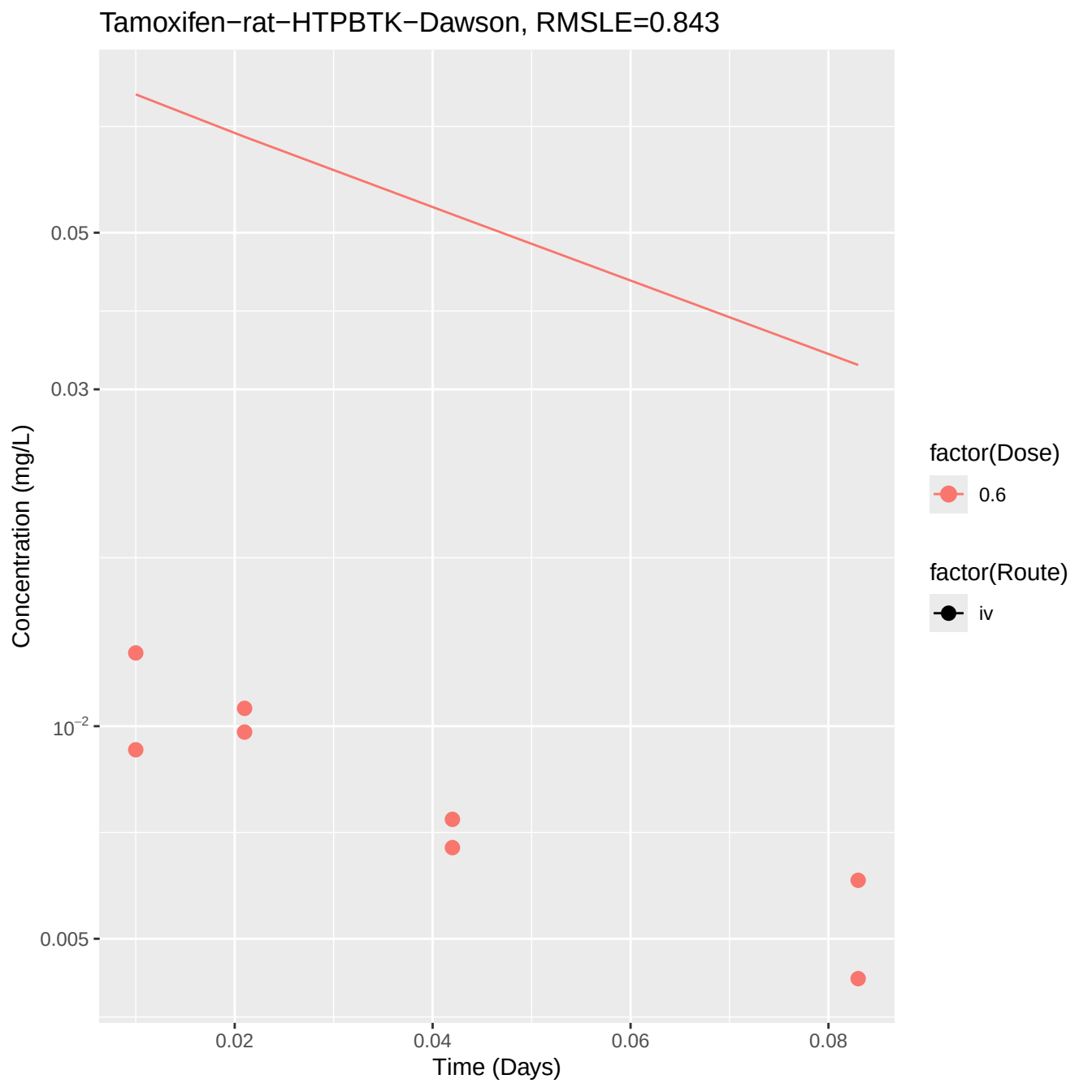
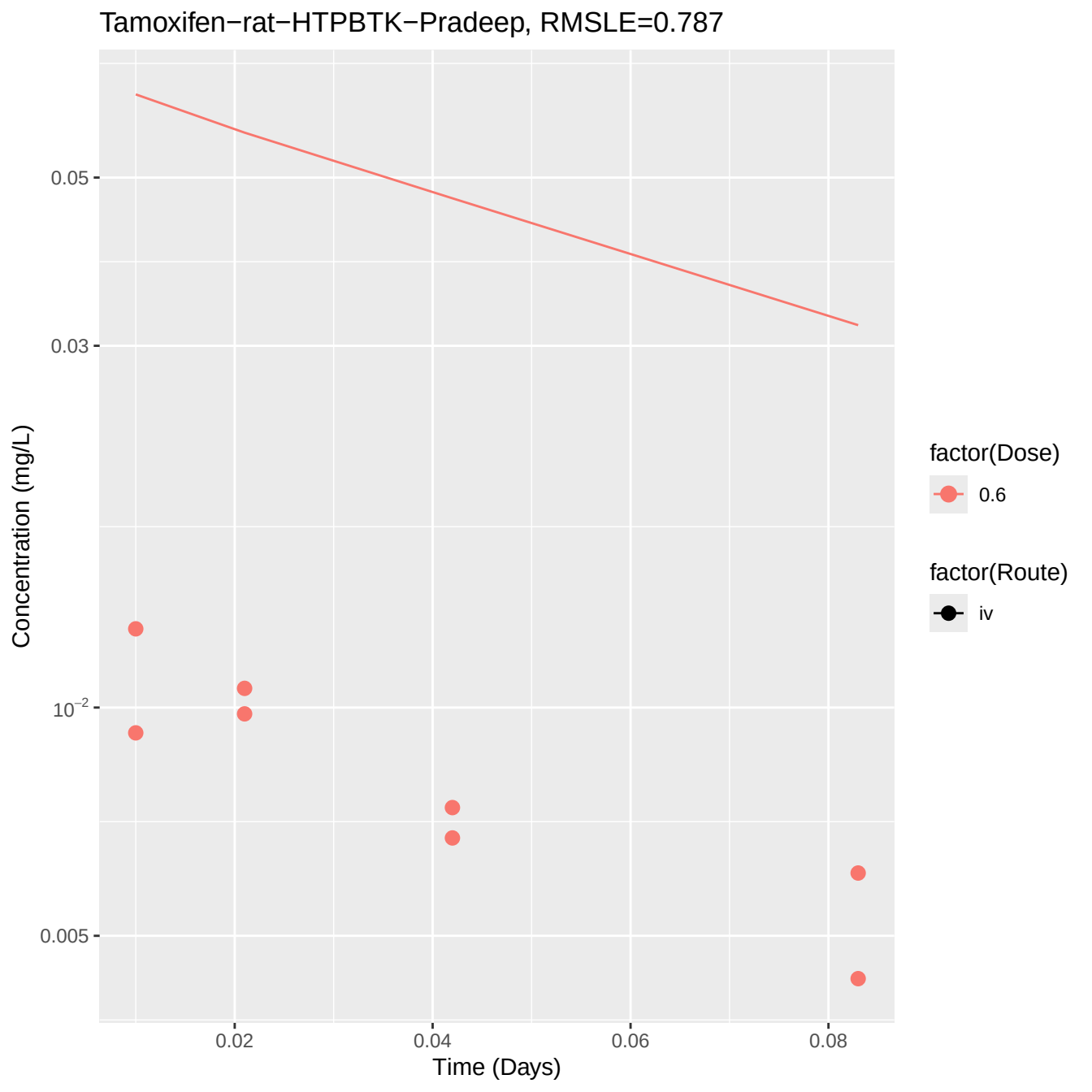
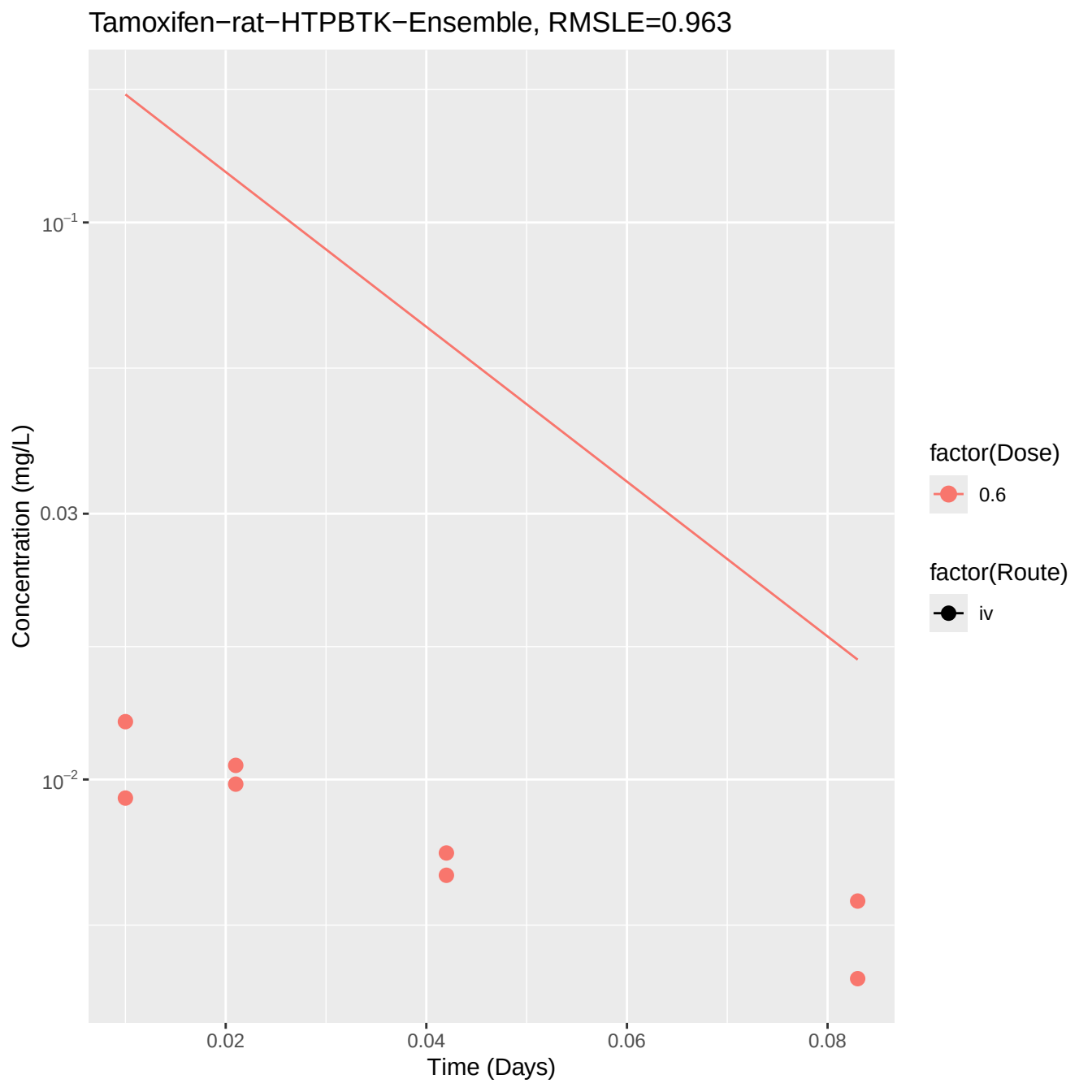
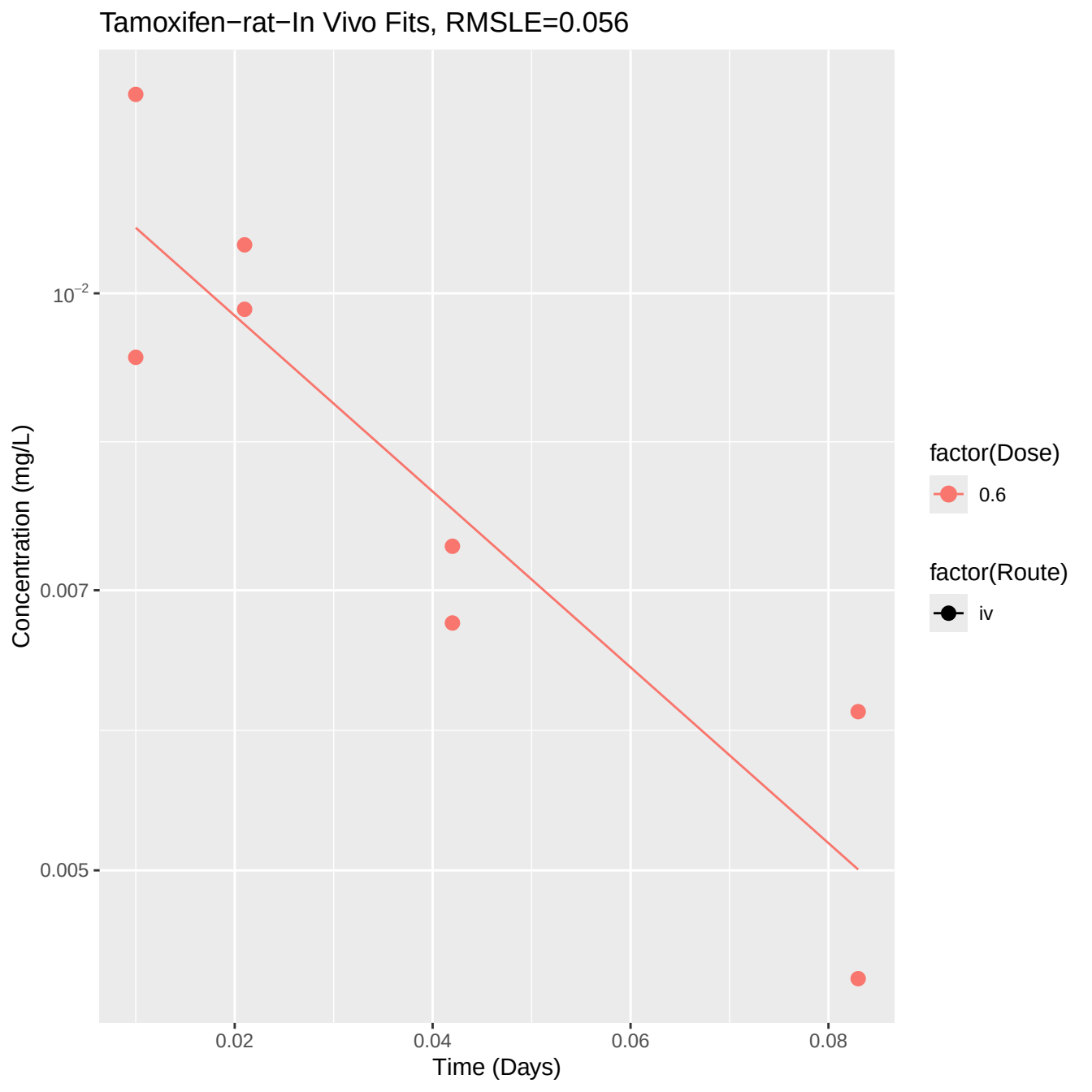

Tamoxifen−rat−HTPBTK−InVitro, RMSLE=1.36
Concentration (mg/L)
10^{-1}
0.03
10^{-2}
0.02
0.04
0.06
0.08
Time (Days)
factor(Dose)
0.6
factor(Route)
iv

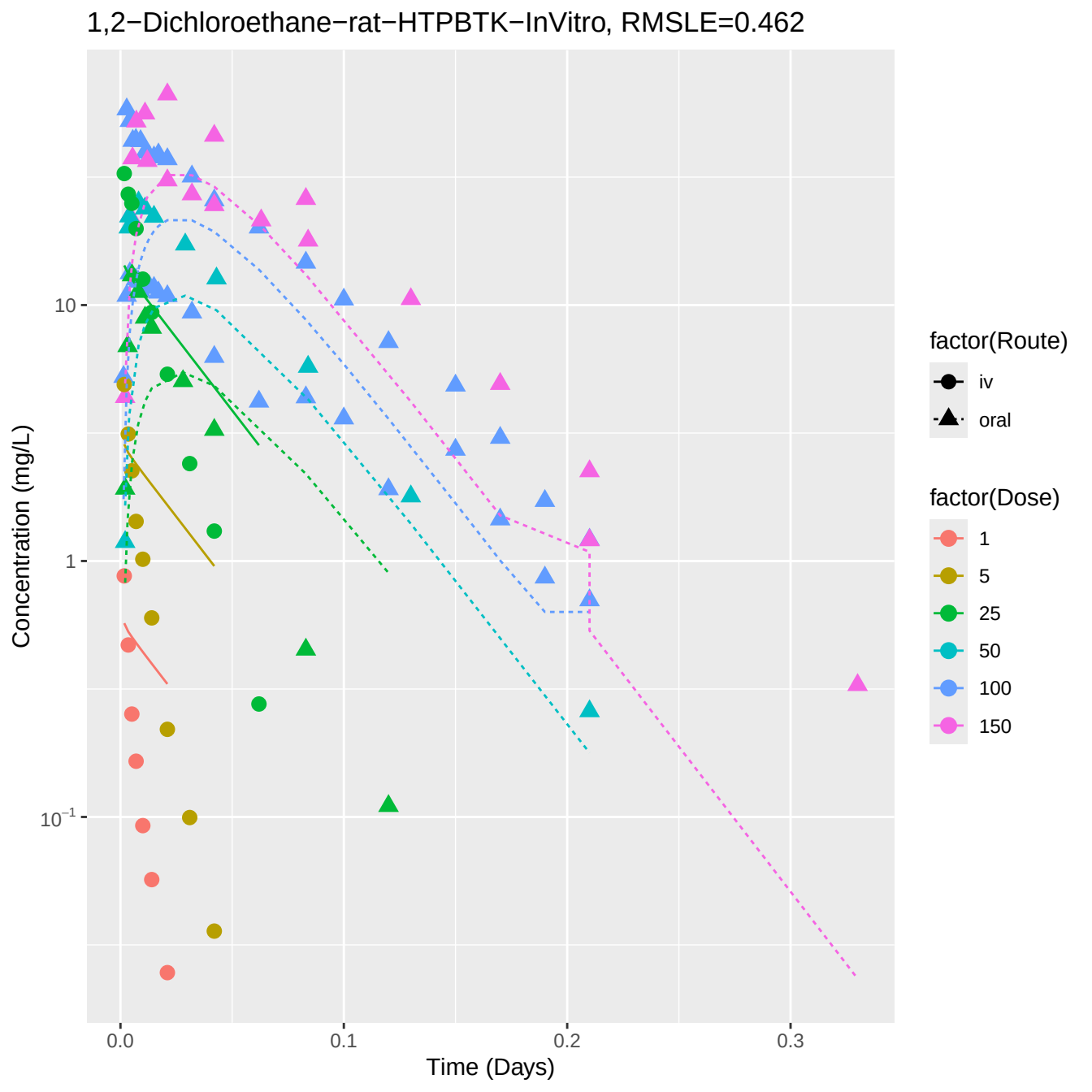

Tamoxifen−rat−HTPBTK−ADMET, RMSLE=0.709
Concentration (mg/L)
0.05
0.03
10^{-2}
0.005
0.02
0.04
0.06
0.08
Time (Days)
factor(Dose)
0.6
factor(Route)
iv

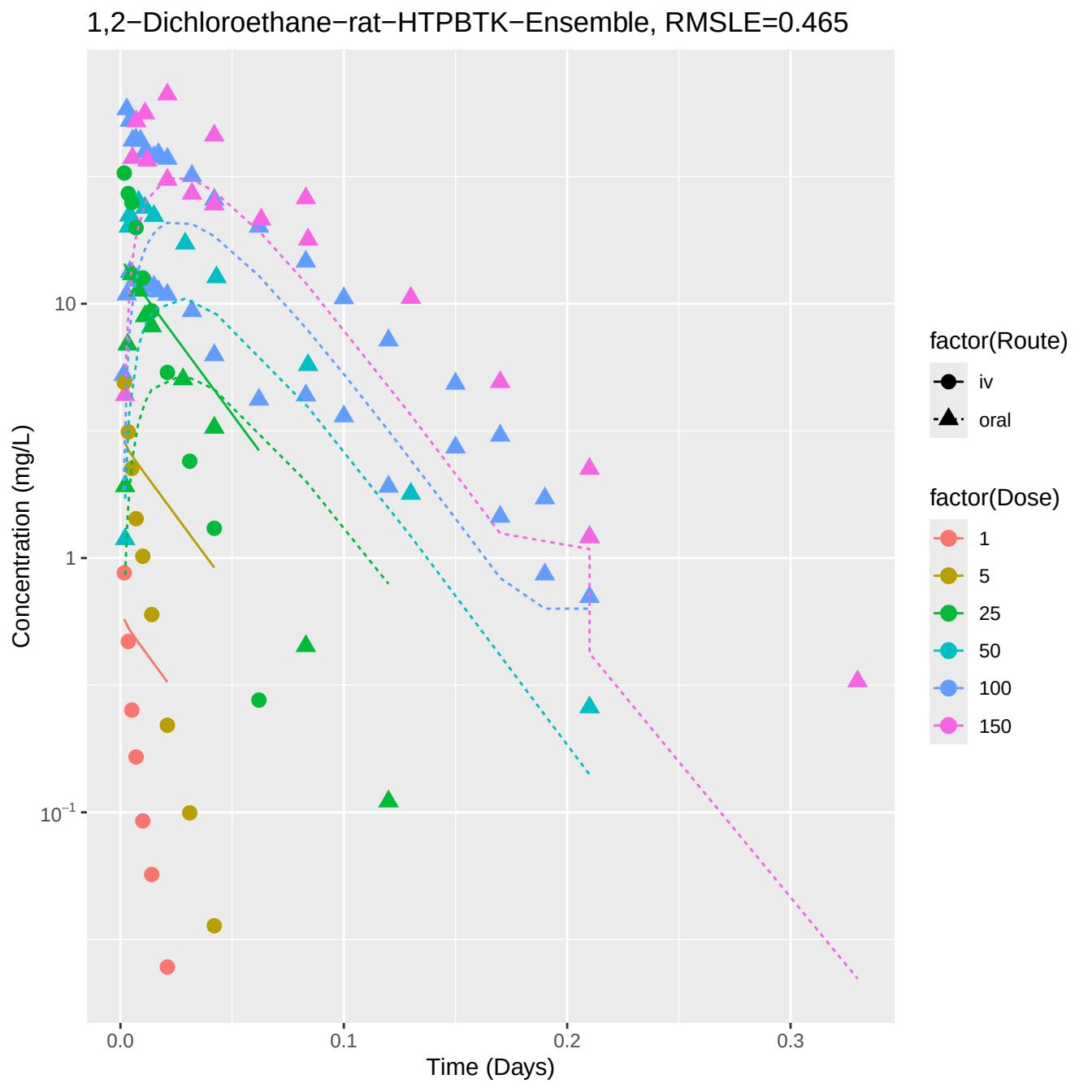

Tamoxifen−rat−HTPBTK−Dawson, RMSLE=0.843
Concentration (mg/L)
0.05
0.03
10^{-2}
0.005
0.02
0.04
0.06
0.08
Time (Days)
factor(Dose)
0.6
factor(Route)
iv

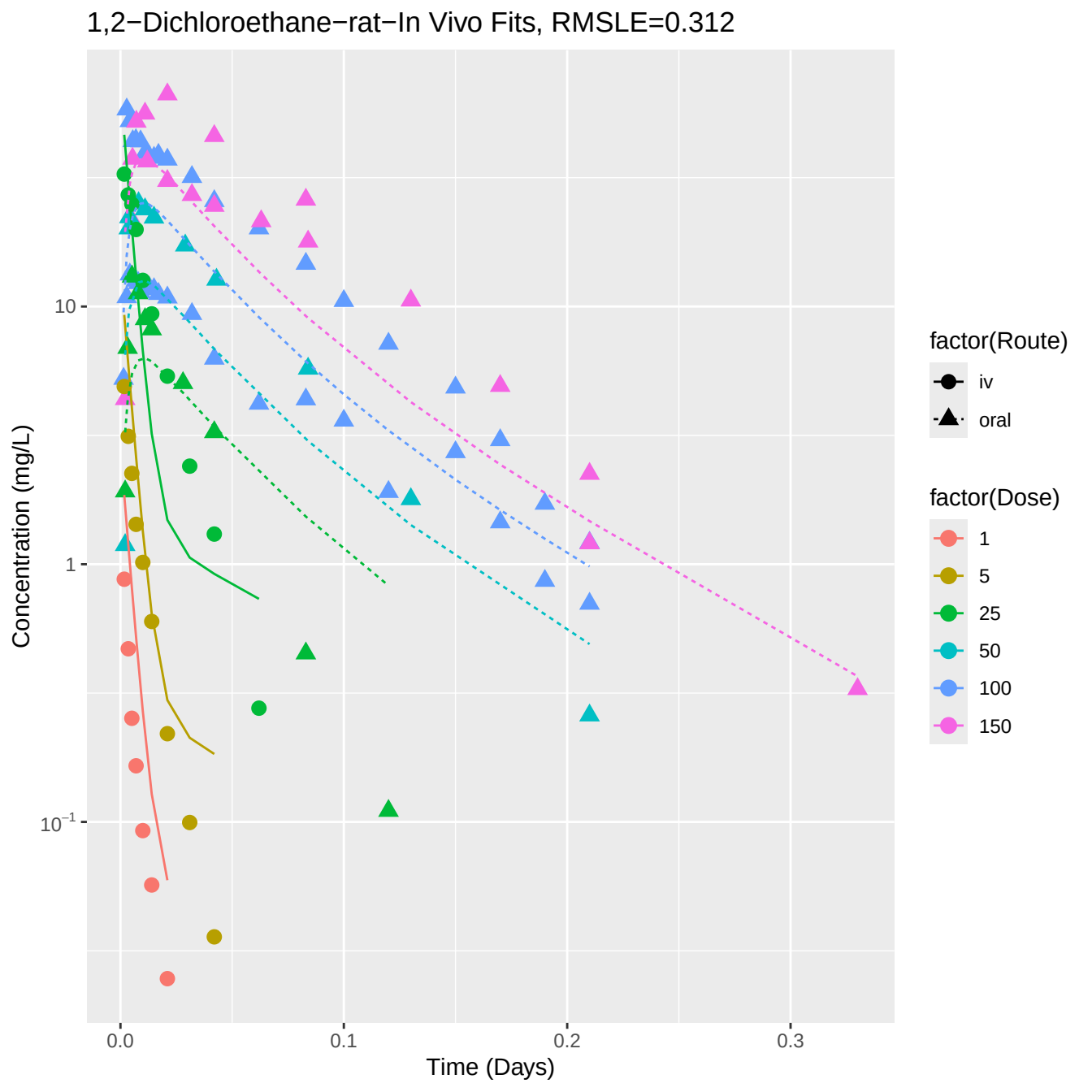

Tamoxifen−rat−HTPBTK−Pradeep, RMSLE=0.787
Concentration (mg/L)
0.05
0.03
10^{-2}
0.005
0.02
0.04
0.06
0.08
Time (Days)
factor(Dose)
0.6
factor(Route)
iv

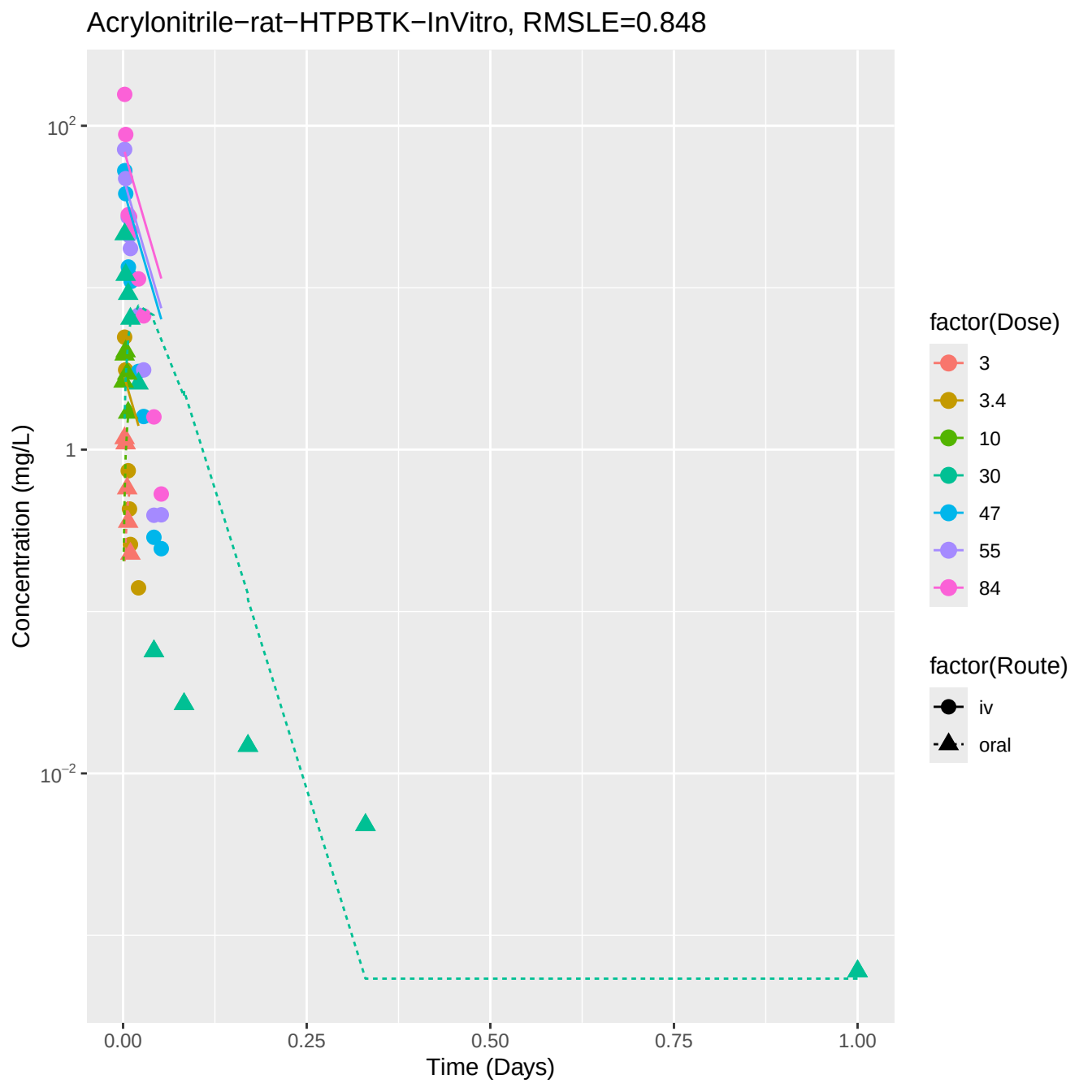

Tamoxifen−rat−HTPBTK−Ensemble, RMSLE=0.963
Concentration (mg/L)
10^{-1}
0.03
10^{-2}
0.02
0.04
0.06
0.08
Time (Days)
factor(Dose)
0.6
factor(Route)
iv

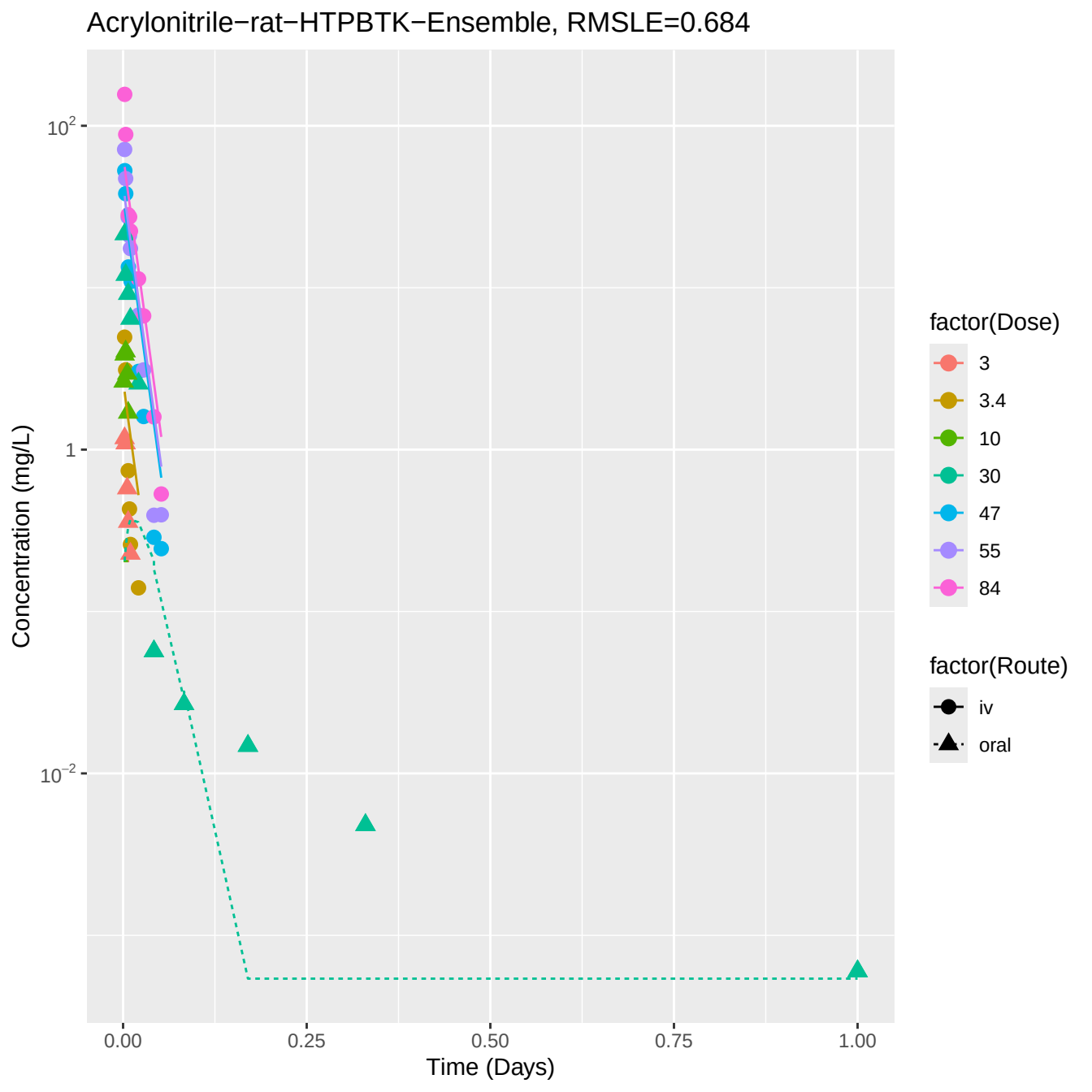

Tamoxifen−rat−In Vivo Fits, RMSLE=0.056
Concentration (mg/L)
10^{-2}
0.007
0.005
0.02
0.04
0.06
0.08
Time (Days)
factor(Dose)
0.6
factor(Route)
iv

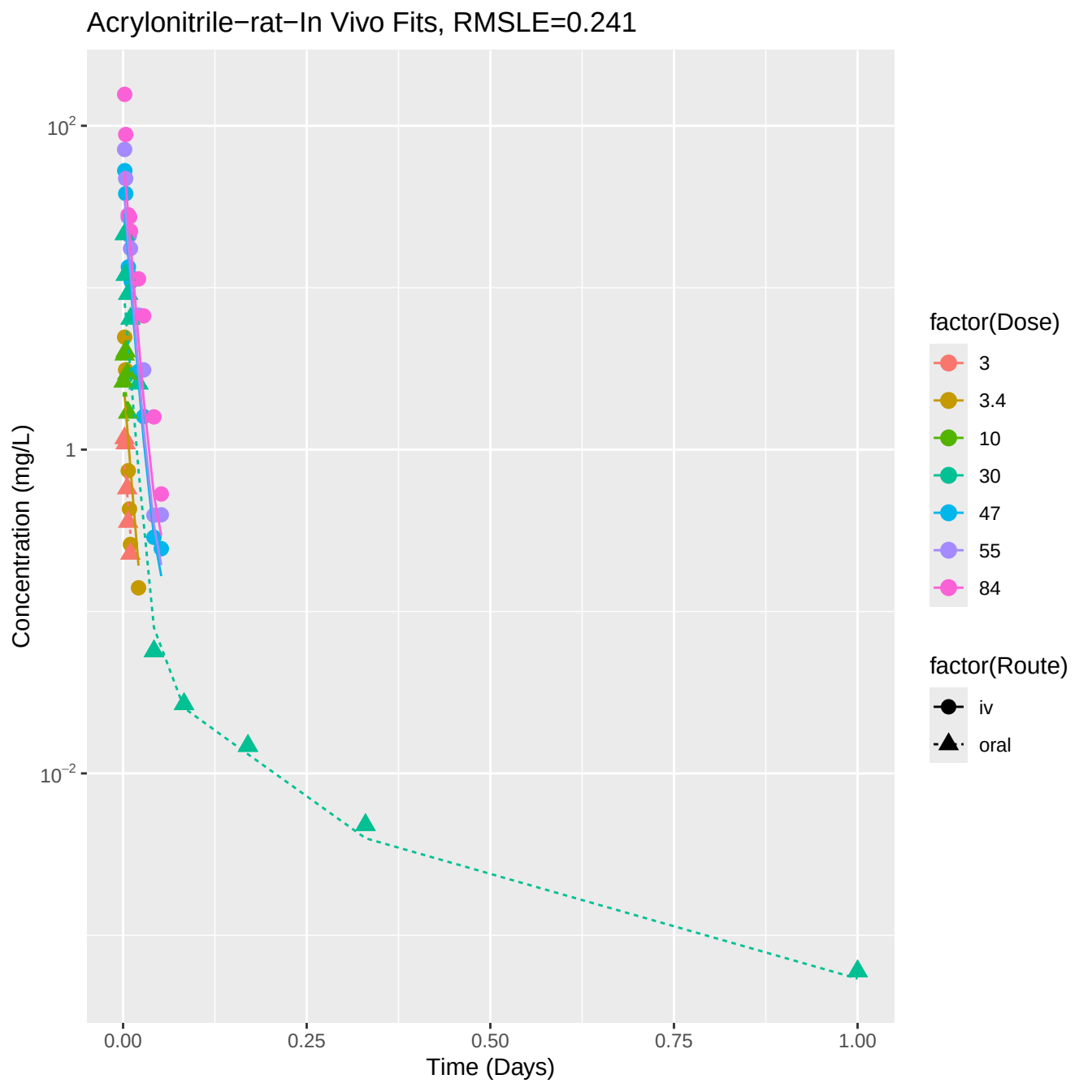

1,2−Dichloroethane−rat−HTPBTK−InVitro, RMSLE=0.462
Concentration (mg/L)
10
1
10^{-1}
0.0
0.1
0.2
0.3
Time (Days)
factor(Route)
iv
oral
factor(Dose)
1
5
25
50
100
150

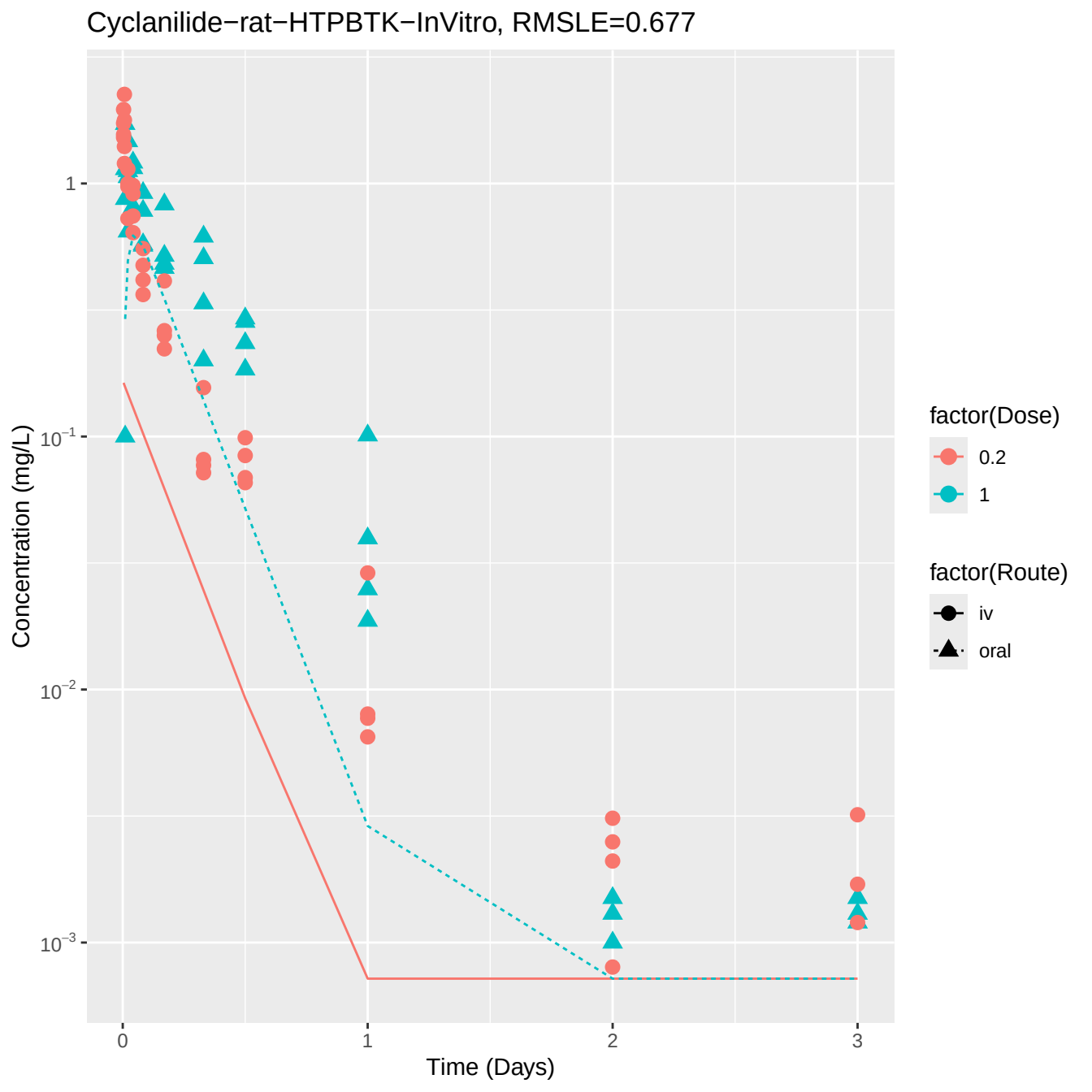

1,2−Dichloroethane−rat−HTPBTK−Ensemble, RMSLE=0.465
Concentration (mg/L)
10
1
10^{-1}
0.0
0.1
0.2
0.3
Time (Days)
factor(Route)
iv
oral
factor(Dose)
1
5
25
50
100
150

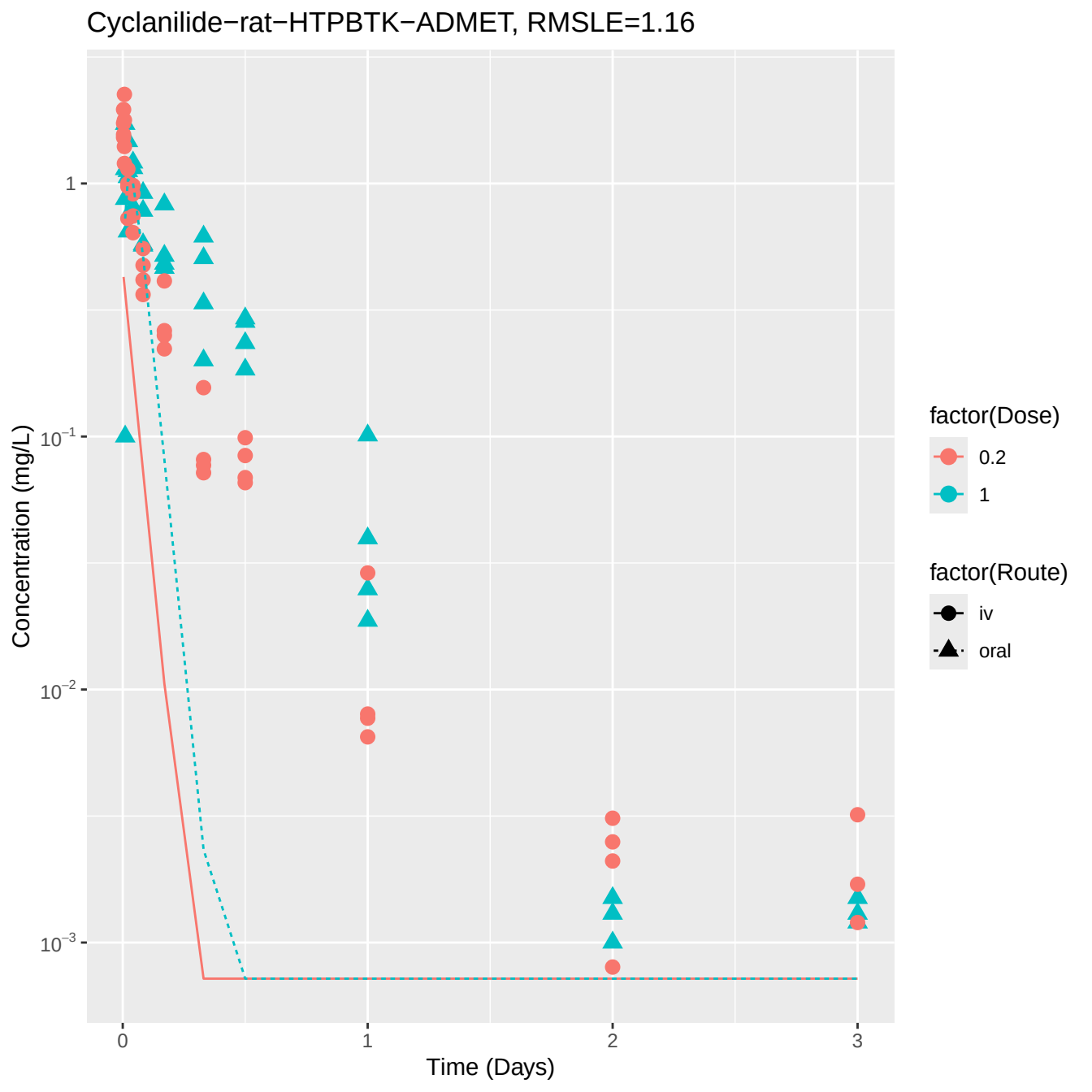

1,2−Dichloroethane−rat−In Vivo Fits, RMSLE=0.312
Concentration (mg/L)
10
1
10^{-1}
0.0
0.1
0.2
0.3
Time (Days)
factor(Route)
iv
oral
factor(Dose)
1
5
25
50
100
150

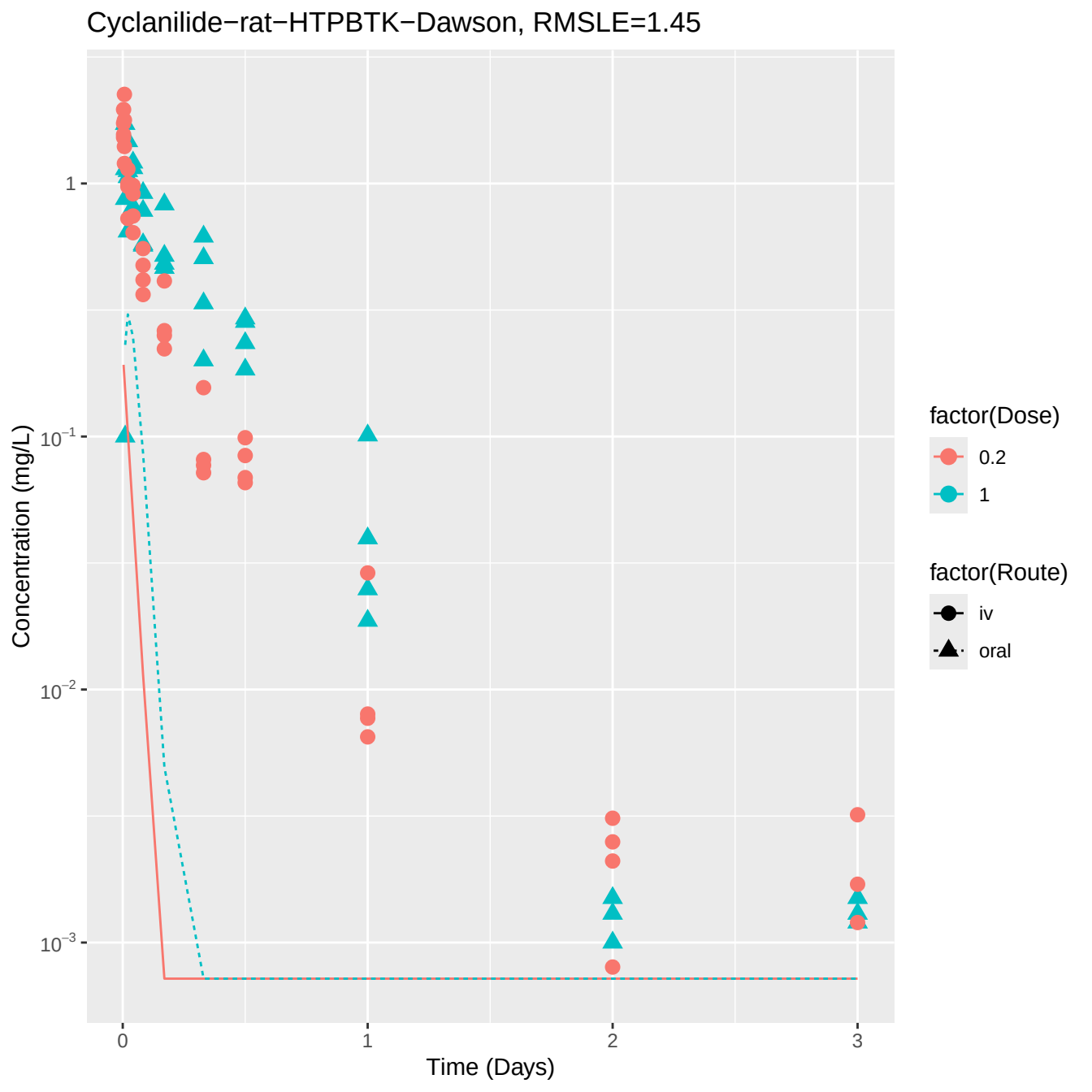

Acrylonitrile−rat−HTPBTK−InVitro, RMSLE=0.848
Concentration (mg/L)
10^2
1
10^{-2}
0.00
0.25
0.50
0.75
1.00
Time (Days)
factor(Dose)
3
3.4
10
30
47
55
84
factor(Route)
iv
oral

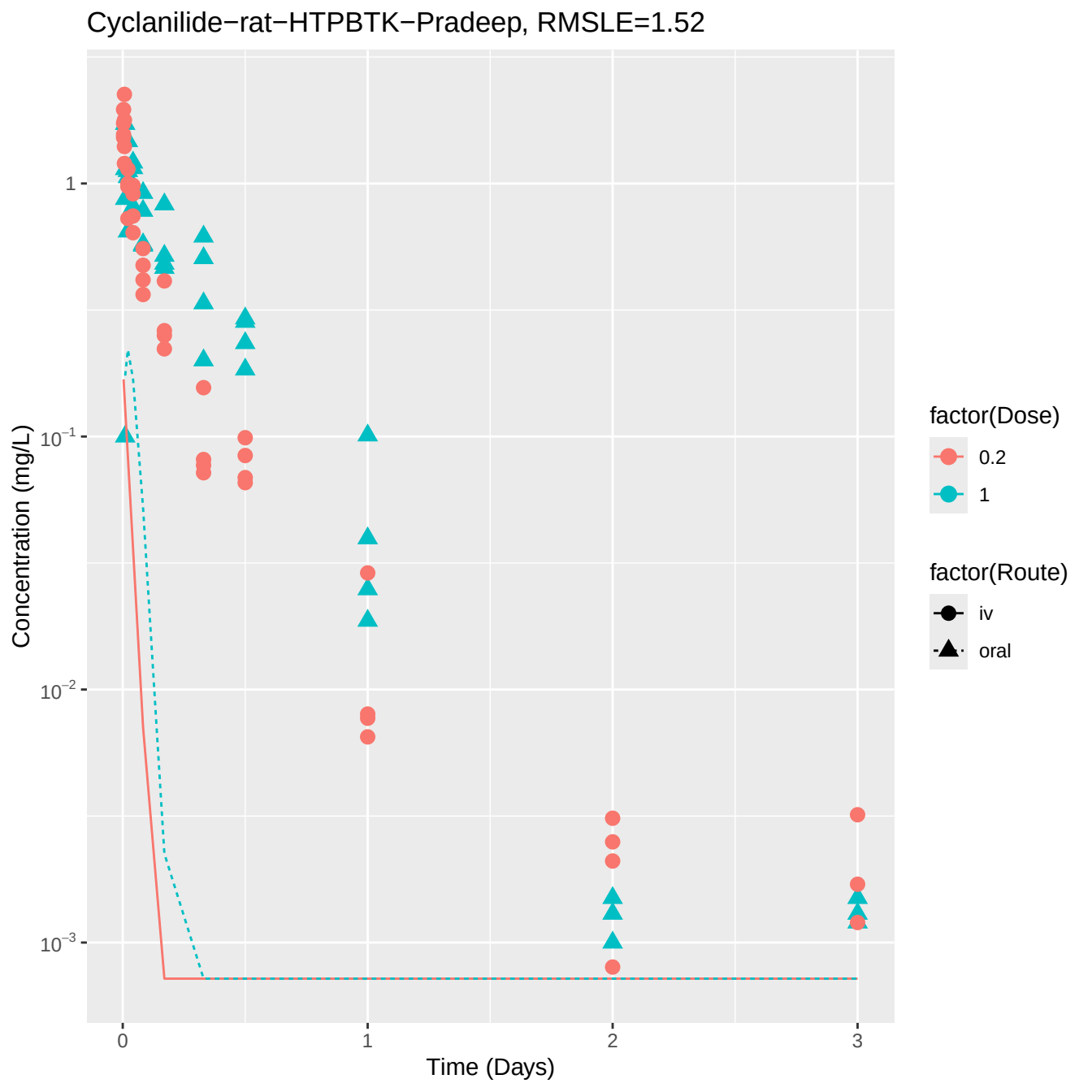

Acrylonitrile−rat−HTPBTK−Ensemble, RMSLE=0.684
Concentration (mg/L)
10^2
1
10^{-2}
0.00
0.25
0.50
0.75
1.00
Time (Days)
factor(Dose)
3
3.4
10
30
47
55
84
factor(Route)
iv
oral

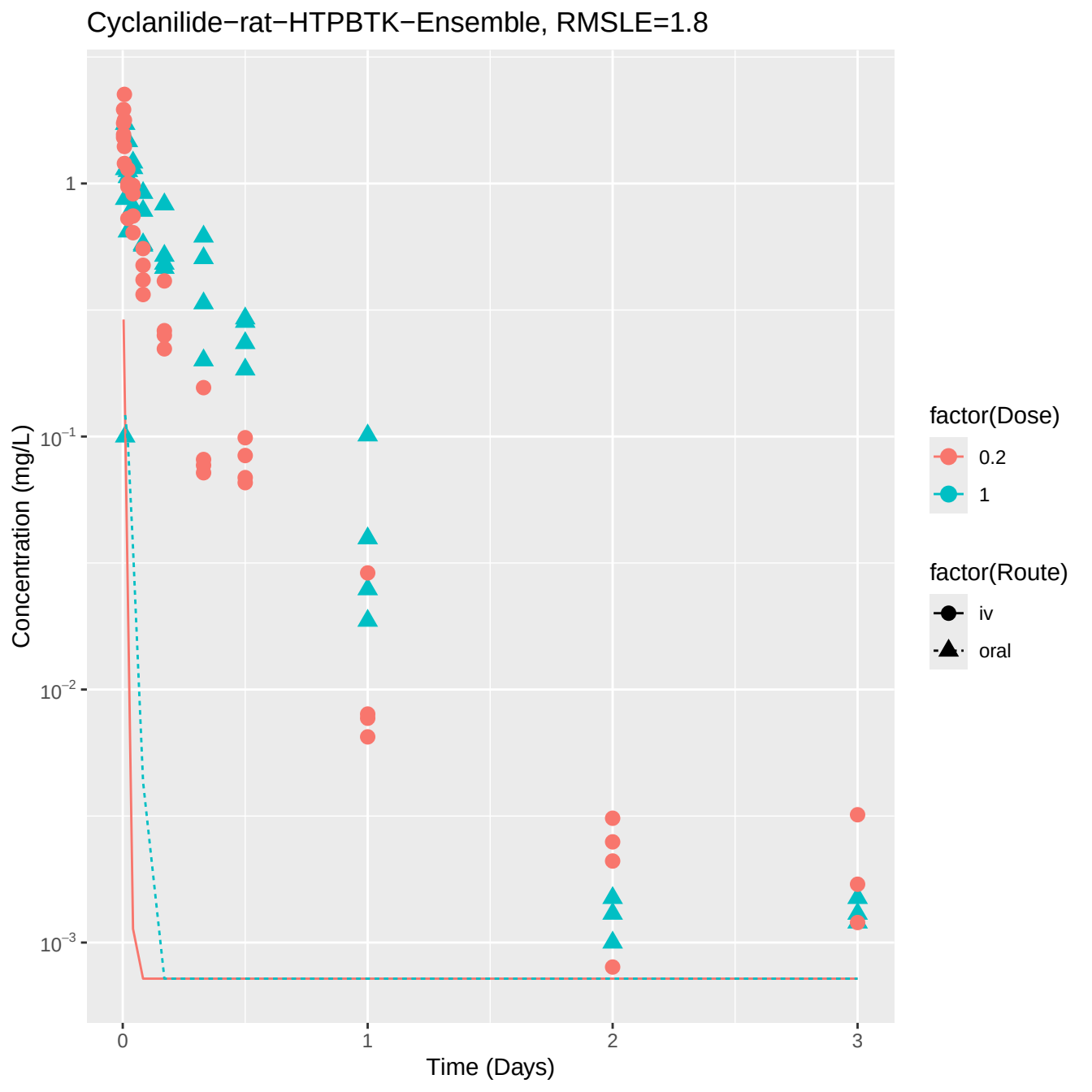

Acrylonitrile−rat−In Vivo Fits, RMSLE=0.241
Concentration (mg/L)
10^2
1
10^{-2}
0.00
0.25
0.50
0.75
1.00
Time (Days)
factor(Dose)
3
3.4
10
30
47
55
84
factor(Route)
iv
oral

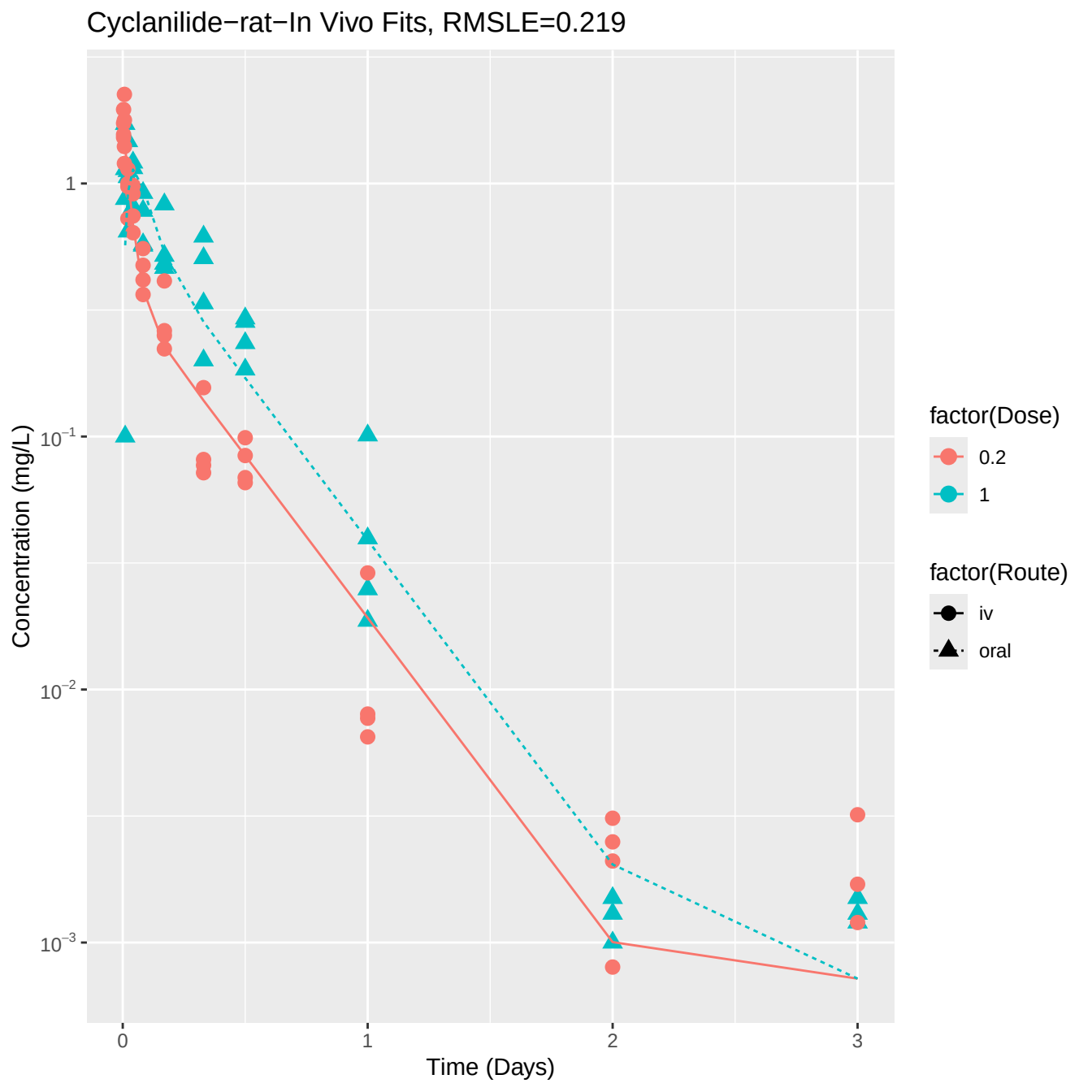

Cyclanilide−rat−HTPBTK−InVitro, RMSLE=0.677
Concentration (mg/L)
1
10^{-1}
10^{-2}
10^{-3}
0
1
2
3
Time (Days)
factor(Dose)
0.2
1
factor(Route)
iv
oral

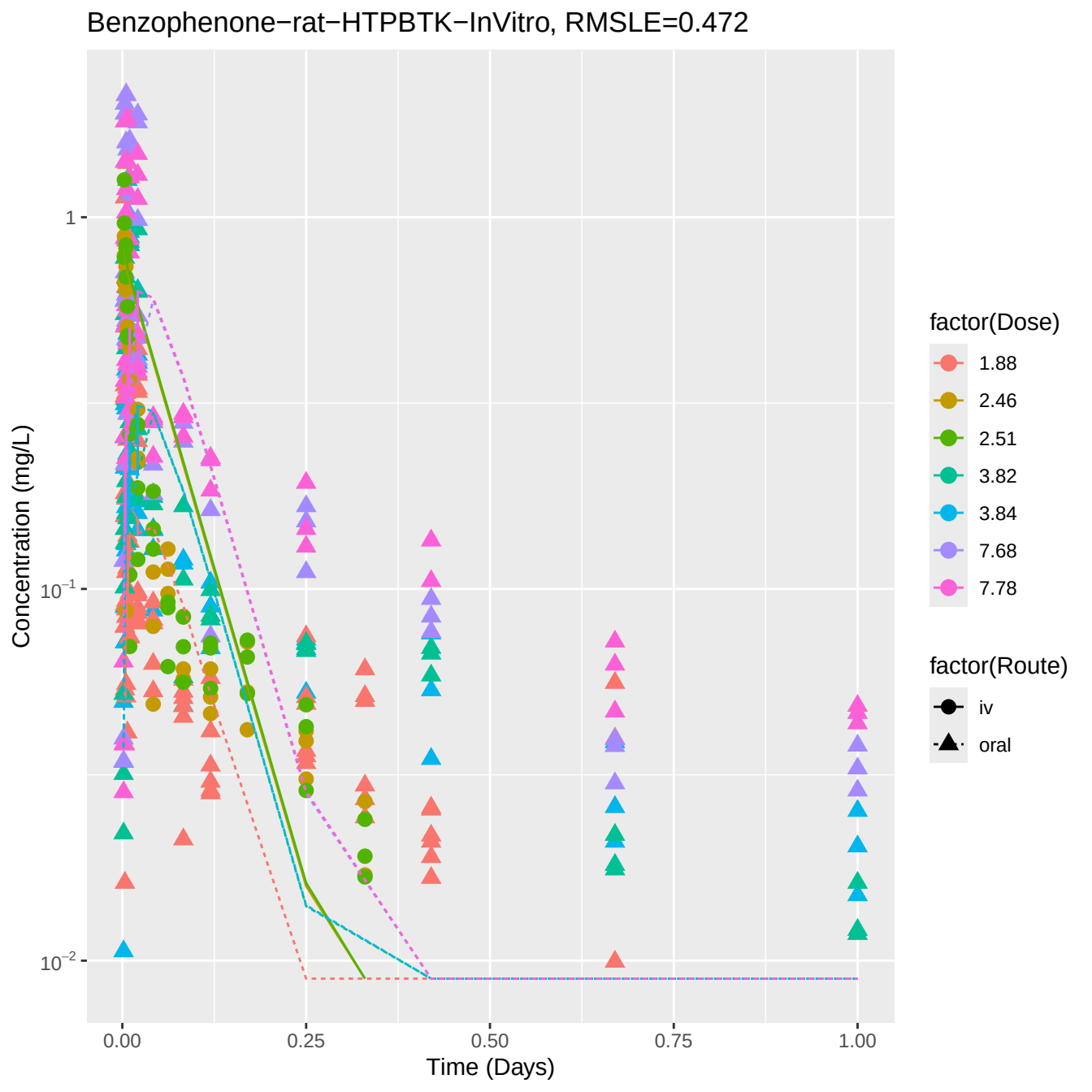

Cyclanilide−rat−HTPBTK−ADMET, RMSLE=1.16
Concentration (mg/L)
1
10^{-1}
10^{-2}
10^{-3}
0
1
2
3
Time (Days)
factor(Dose)
0.2
1
factor(Route)
iv
oral

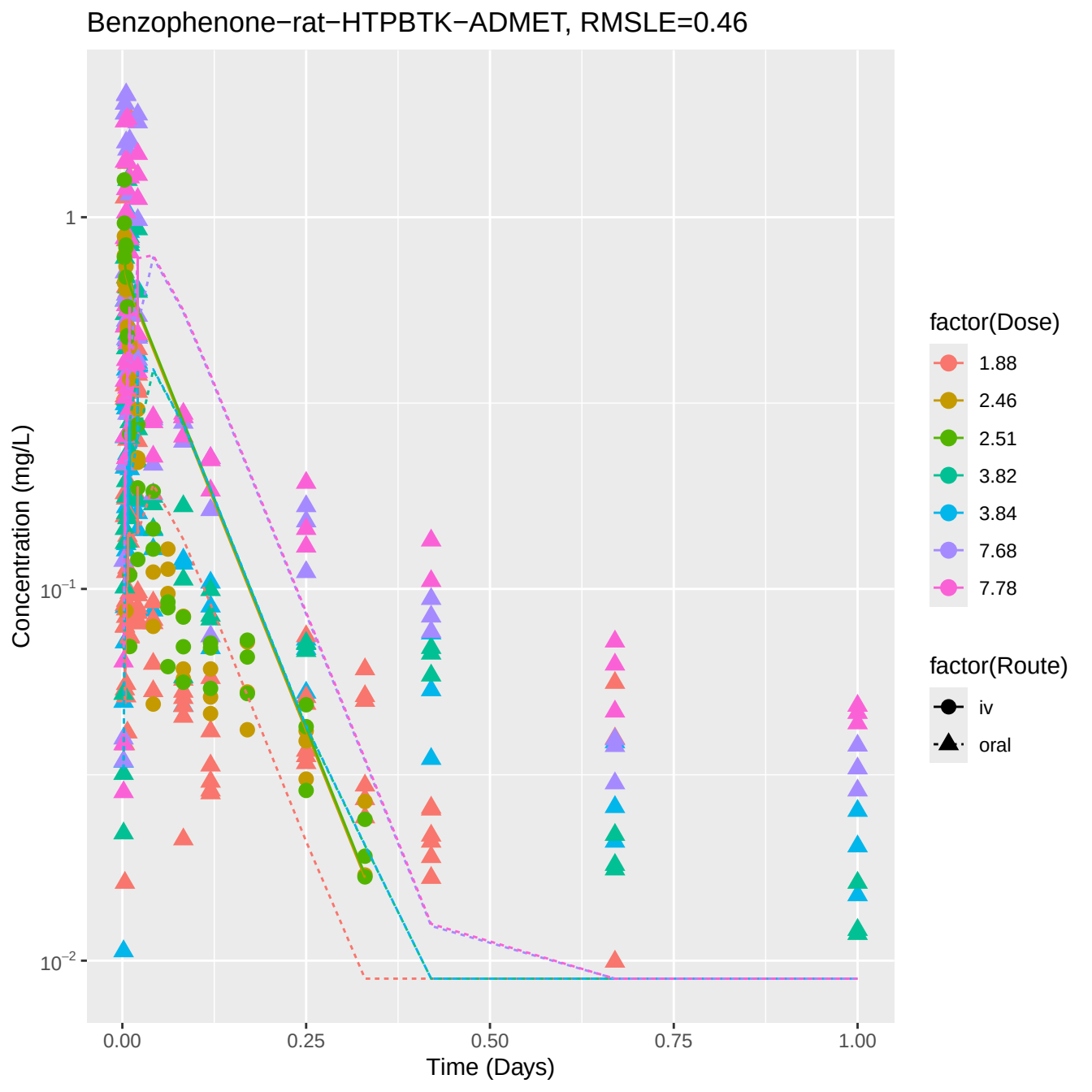

Cyclanilide−rat−HTPBTK−Dawson, RMSLE=1.45
Concentration (mg/L)
1
10^{-1}
10^{-2}
10^{-3}
0
1
2
3
Time (Days)
factor(Dose)
0.2
1
factor(Route)
iv
oral

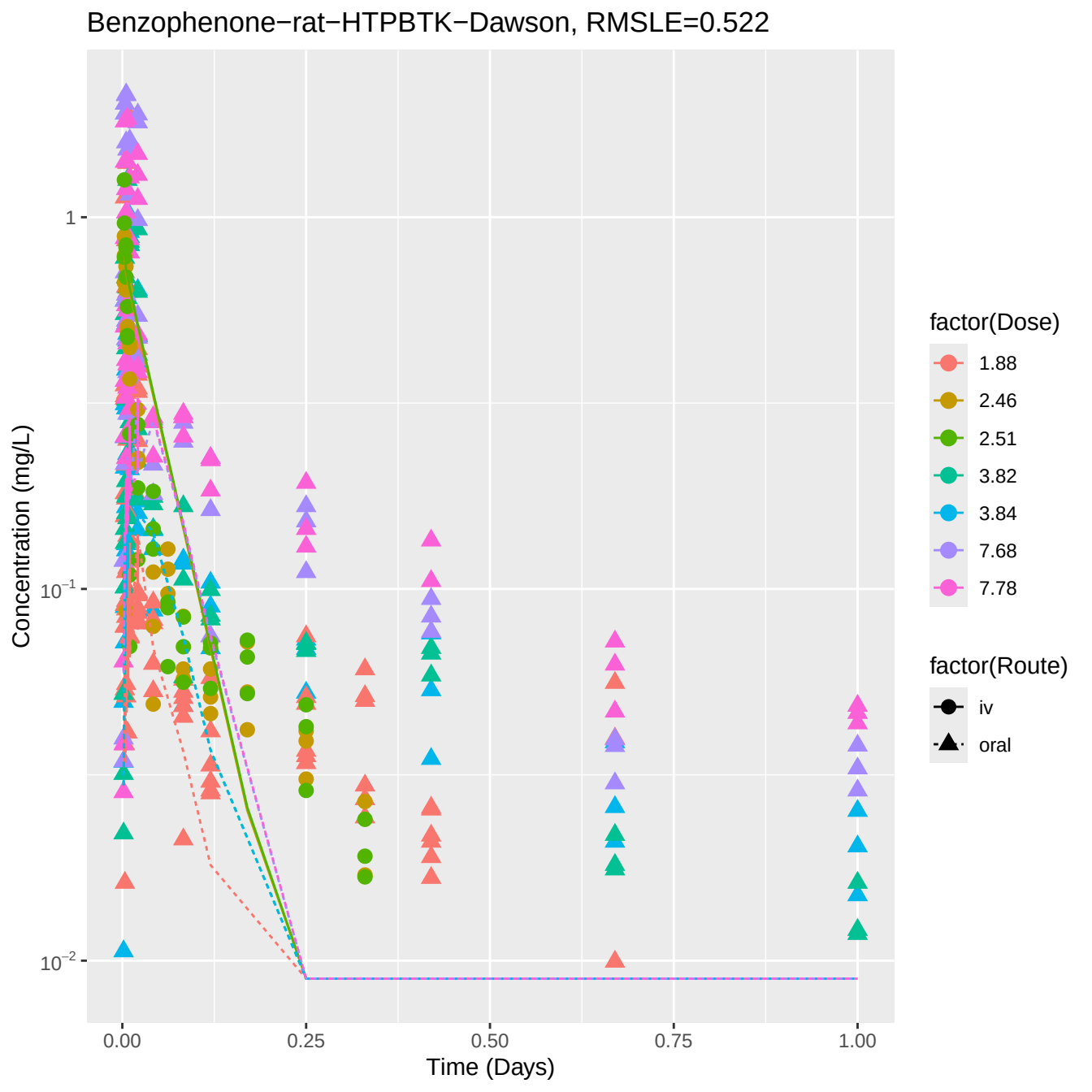

Cyclanilide−rat−HTPBTK−Pradeep, RMSLE=1.52
Concentration (mg/L)
1
10^{-1}
10^{-2}
10^{-3}
0
1
2
3
Time (Days)
factor(Dose)
0.2
1
factor(Route)
iv
oral

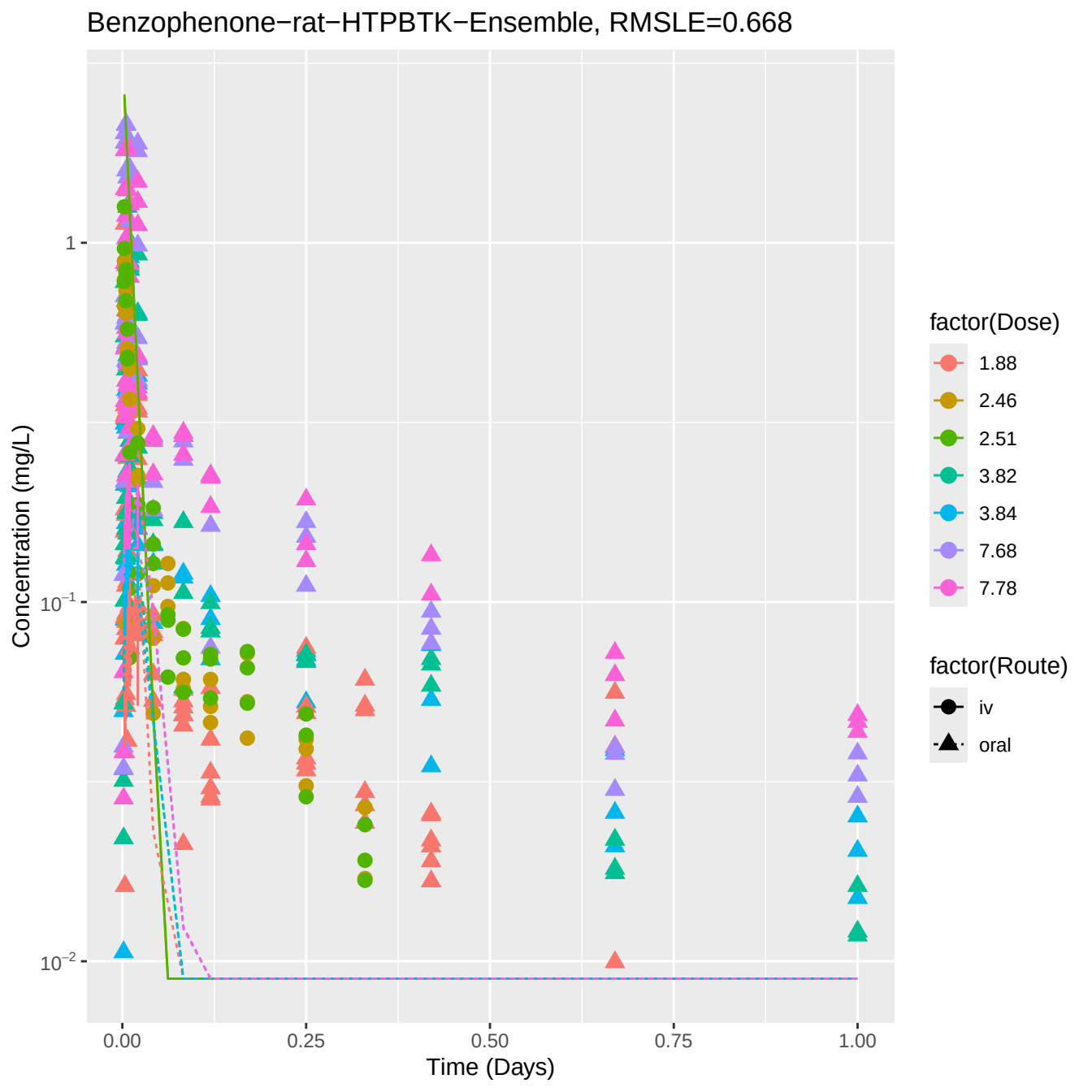

Cyclanilide−rat−HTPBTK−Ensemble, RMSLE=1.8
Concentration (mg/L)
1
10^{-1}
10^{-2}
10^{-3}
0
1
2
3
Time (Days)
factor(Dose)
0.2
1
factor(Route)
iv
oral

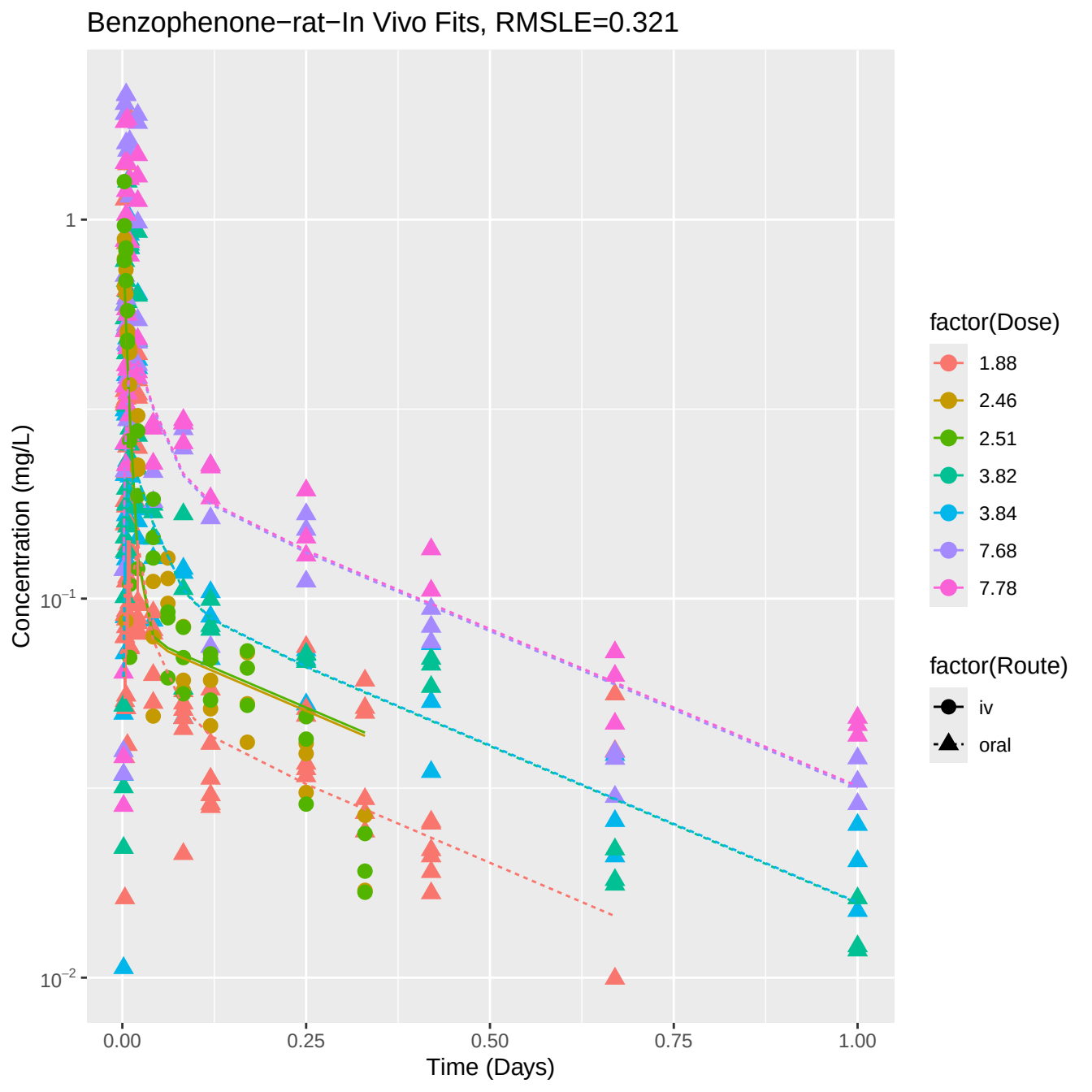

Cyclanilide−rat−In Vivo Fits, RMSLE=0.219
Concentration (mg/L)
1
10^{-1}
10^{-2}
10^{-3}
0
1
2
3
Time (Days)
factor(Dose)
0.2
1
factor(Route)
iv
oral

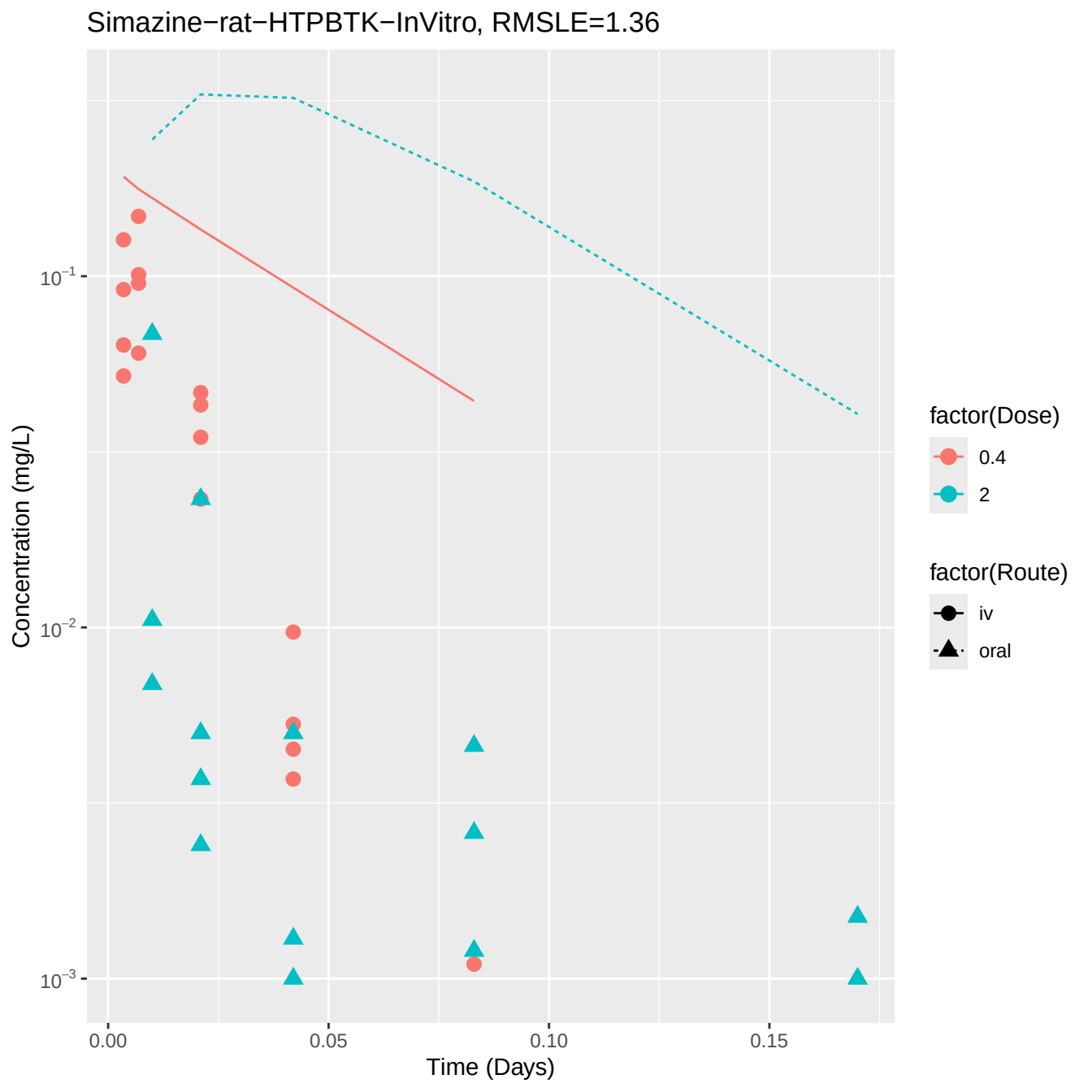

Benzophenone−rat−HTPBTK−InVitro, RMSLE=0.472
Concentration (mg/L)
1
10^{-1}
10^{-2}
0.00
0.25
0.50
0.75
1.00
Time (Days)
factor(Dose)
1.88
2.46
2.51
3.82
3.84
7.68
7.78
factor(Route)
iv
oral

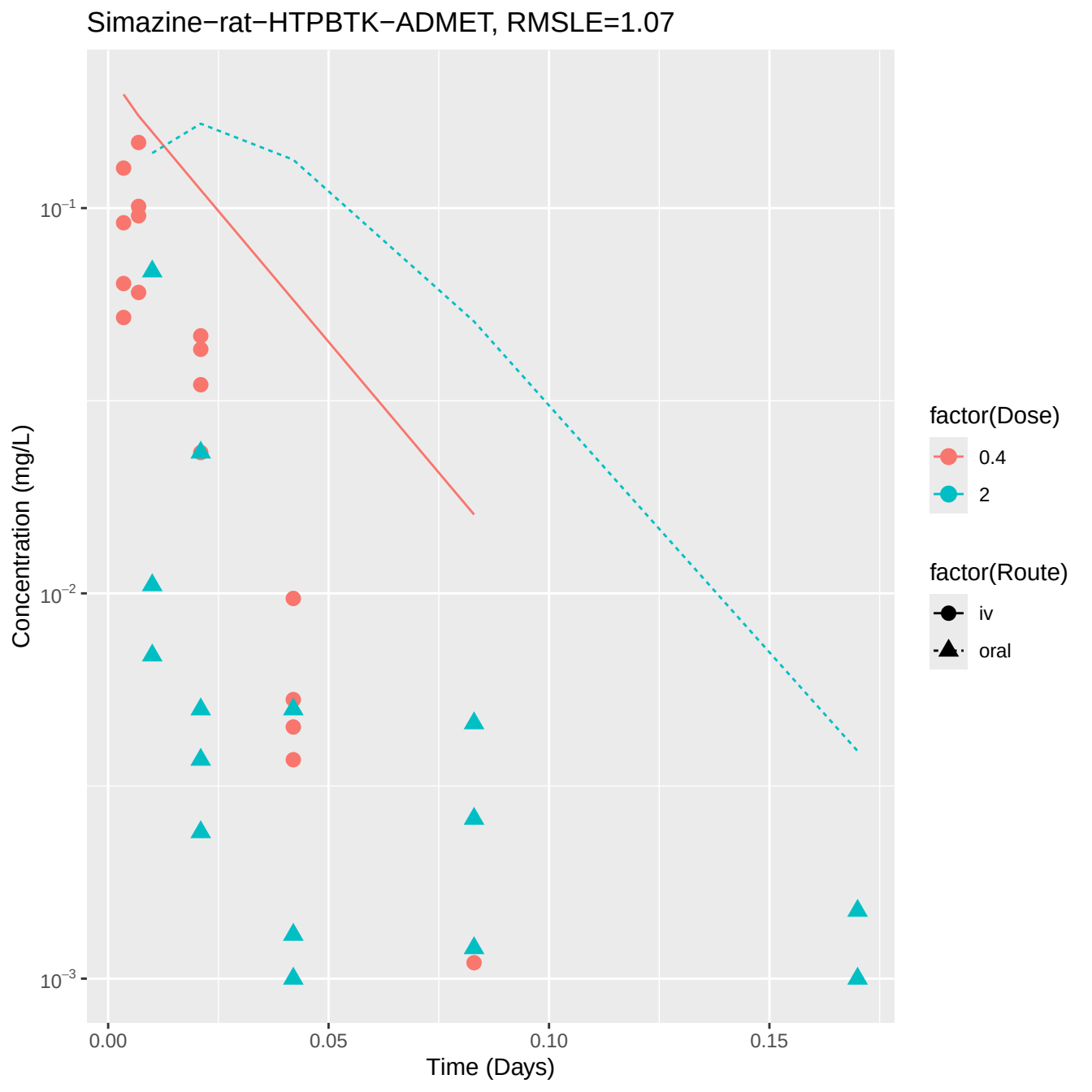

Benzophenone−rat−HTPBTK−ADMET, RMSLE=0.46
Concentration (mg/L)
1
10^{-1}
10^{-2}
0.00
0.25
0.50
0.75
1.00
Time (Days)
factor(Dose)
1.88
2.46
2.51
3.82
3.84
7.68
7.78
factor(Route)
iv
oral

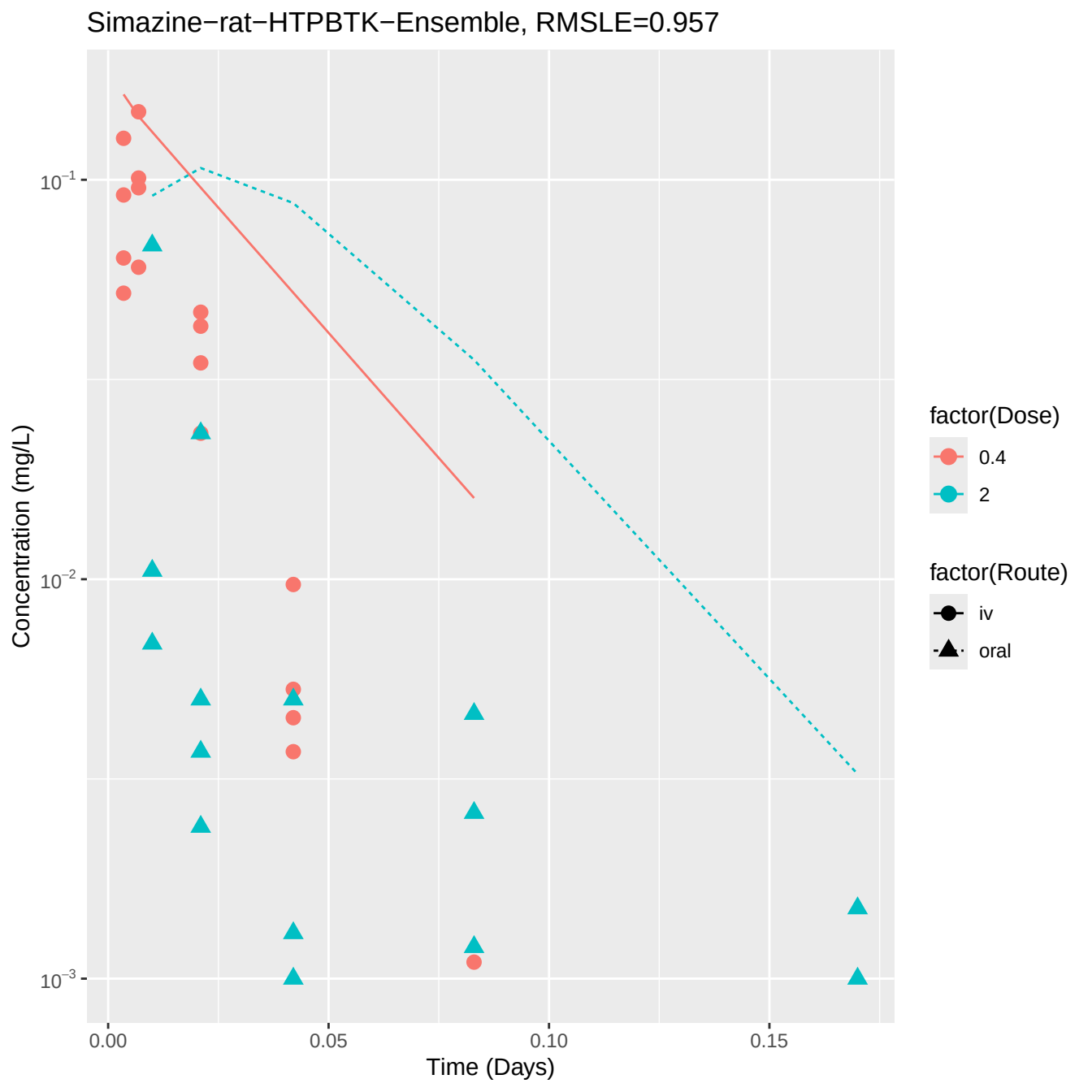

Benzophenone−rat−HTPBTK−Dawson, RMSLE=0.522
Concentration (mg/L)
1
10^{-1}
10^{-2}
0.00
0.25
0.50
0.75
1.00
Time (Days)
factor(Dose)
1.88
2.46
2.51
3.82
3.84
7.68
7.78
factor(Route)
iv
oral

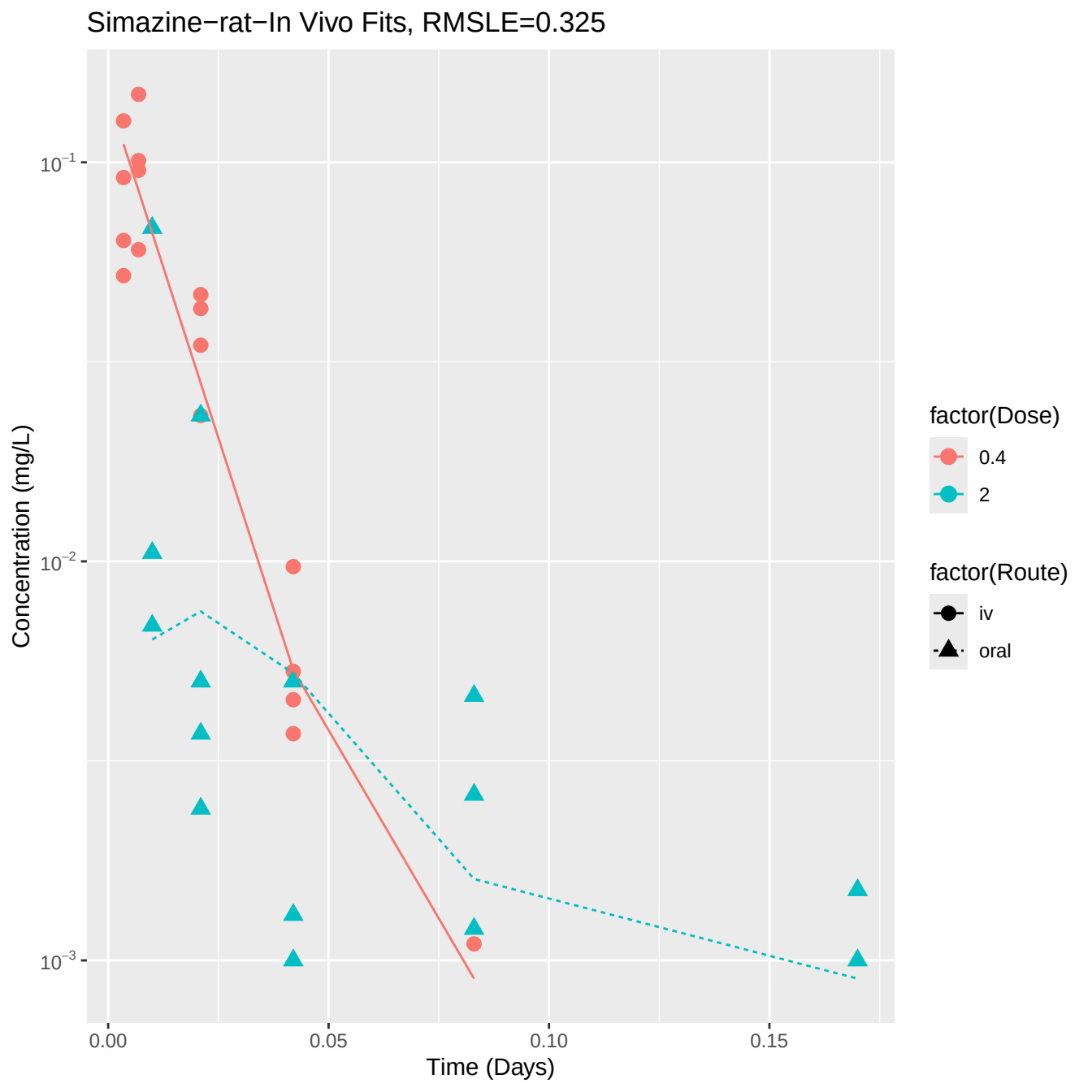

Benzophenone−rat−HTPBTK−Ensemble, RMSLE=0.668
Concentration (mg/L)
1
10^{-1}
10^{-2}
0.00
0.25
0.50
0.75
1.00
Time (Days)
factor(Dose)
1.88
2.46
2.51
3.82
3.84
7.68
7.78
factor(Route)
iv
oral

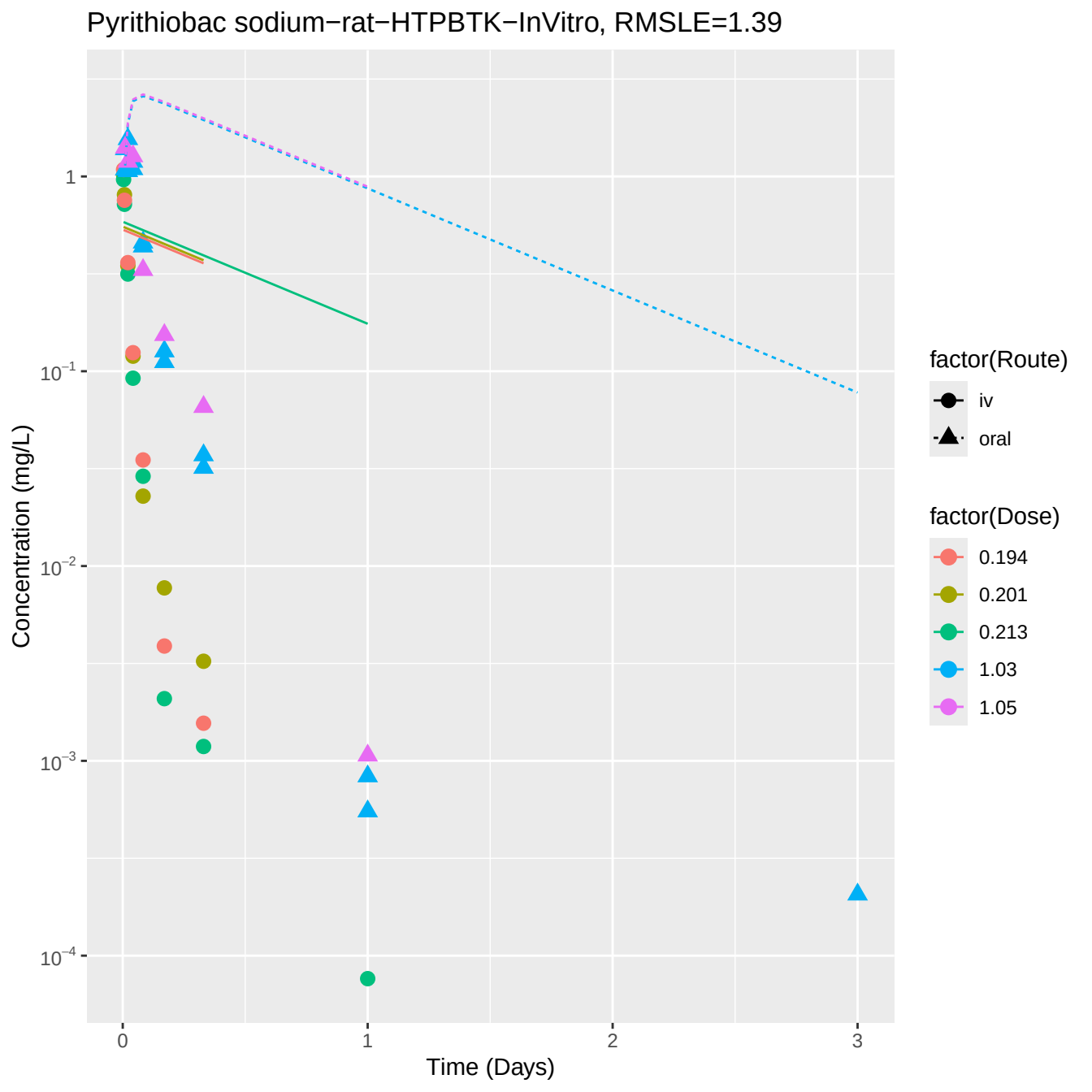

Benzophenone−rat−In Vivo Fits, RMSLE=0.321
Concentration (mg/L)
1
10^{-1}
10^{-2}
0.00
0.25
0.50
0.75
1.00
Time (Days)
factor(Dose)
1.88
2.46
2.51
3.82
3.84
7.68
7.78
factor(Route)
iv
oral

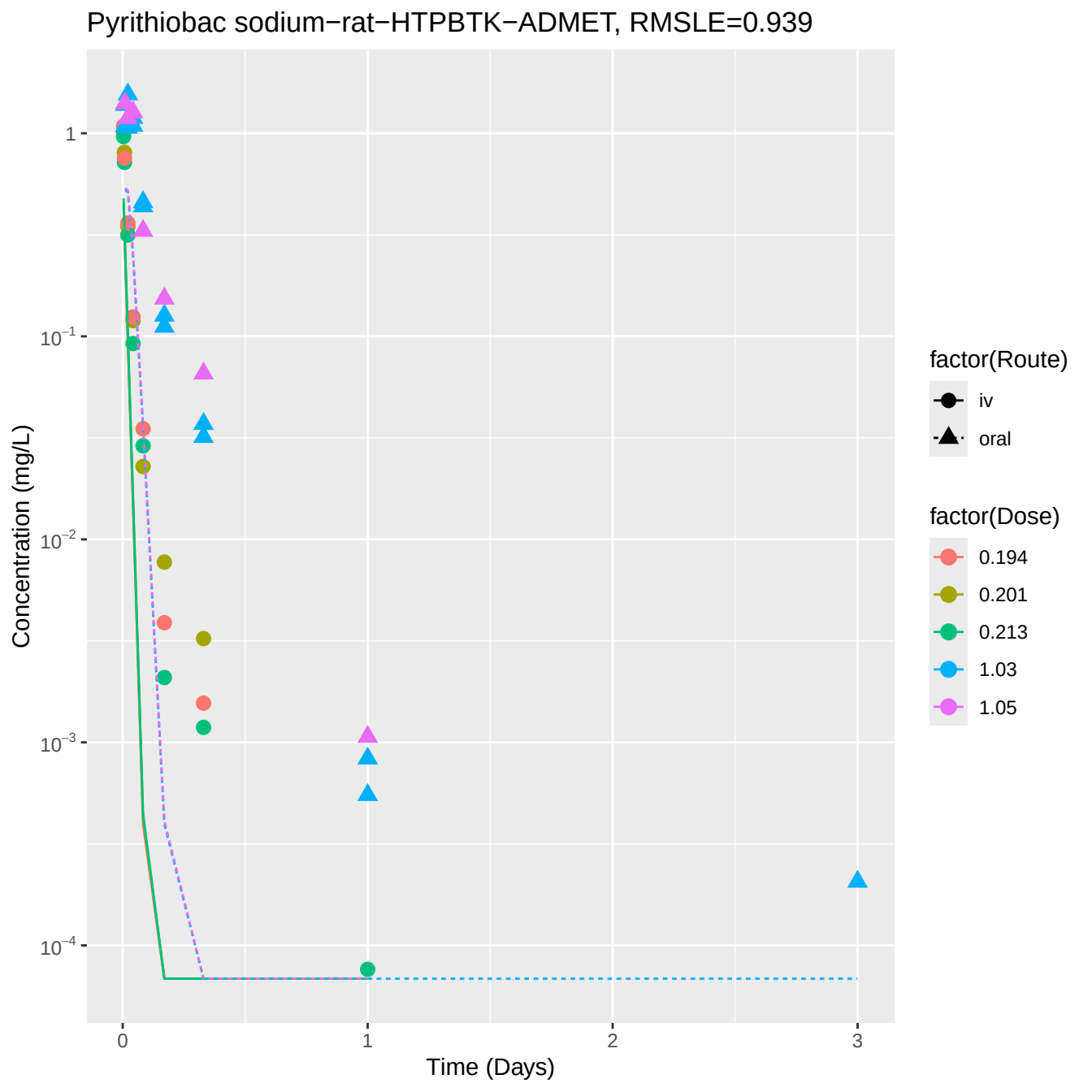

Simazine−rat−HTPBTK−InVitro, RMSLE=1.36
Concentration (mg/L)
10^{-1}
10^{-2}
10^{-3}
0.00
0.05
0.10
0.15
Time (Days)
factor(Dose)
0.4
2
factor(Route)
iv
oral

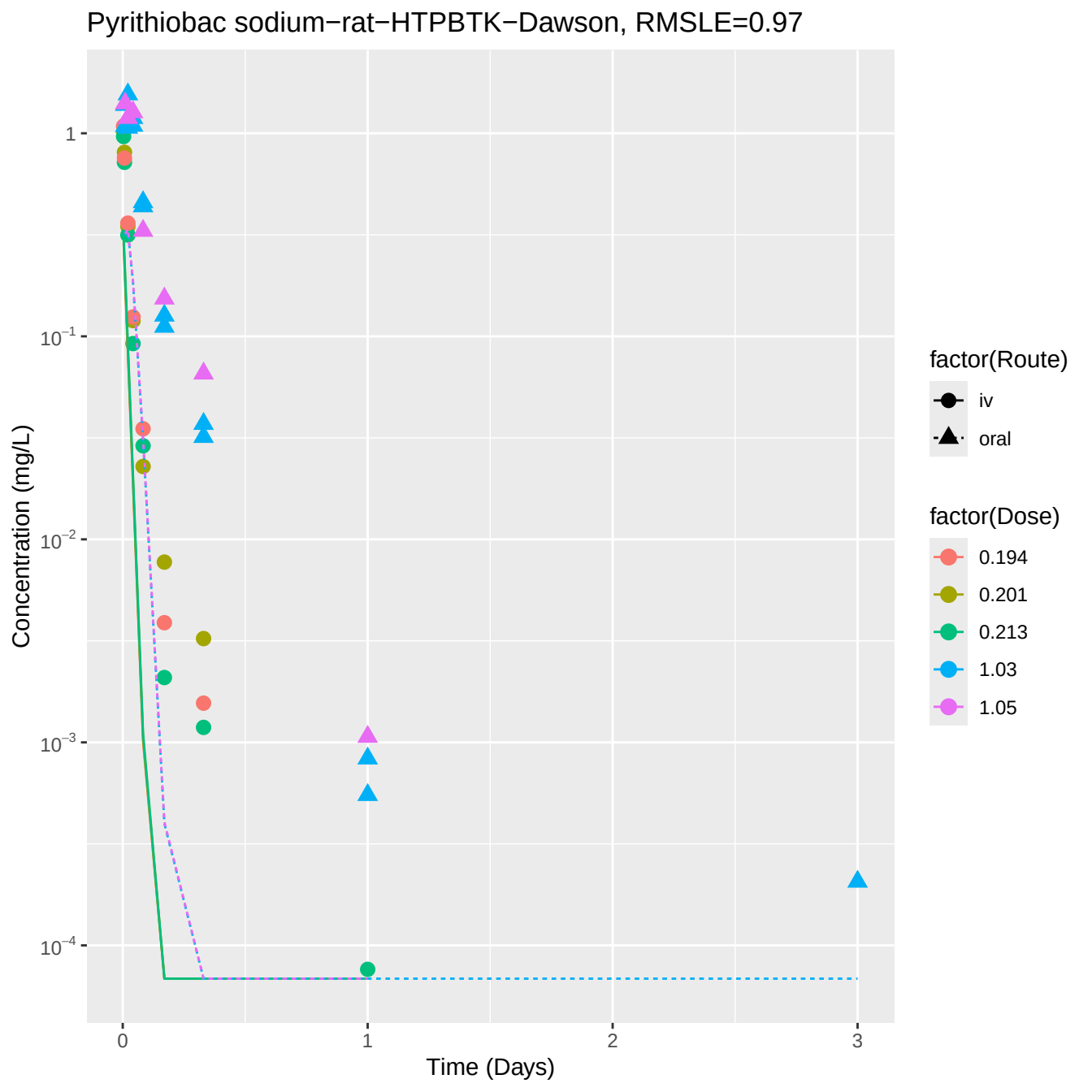

Simazine−rat−HTPBTK−ADMET, RMSLE=1.07
Concentration (mg/L)
10^{-1}
10^{-2}
10^{-3}
0.00
0.05
0.10
0.15
Time (Days)
factor(Dose)
0.4
2
factor(Route)
iv
oral

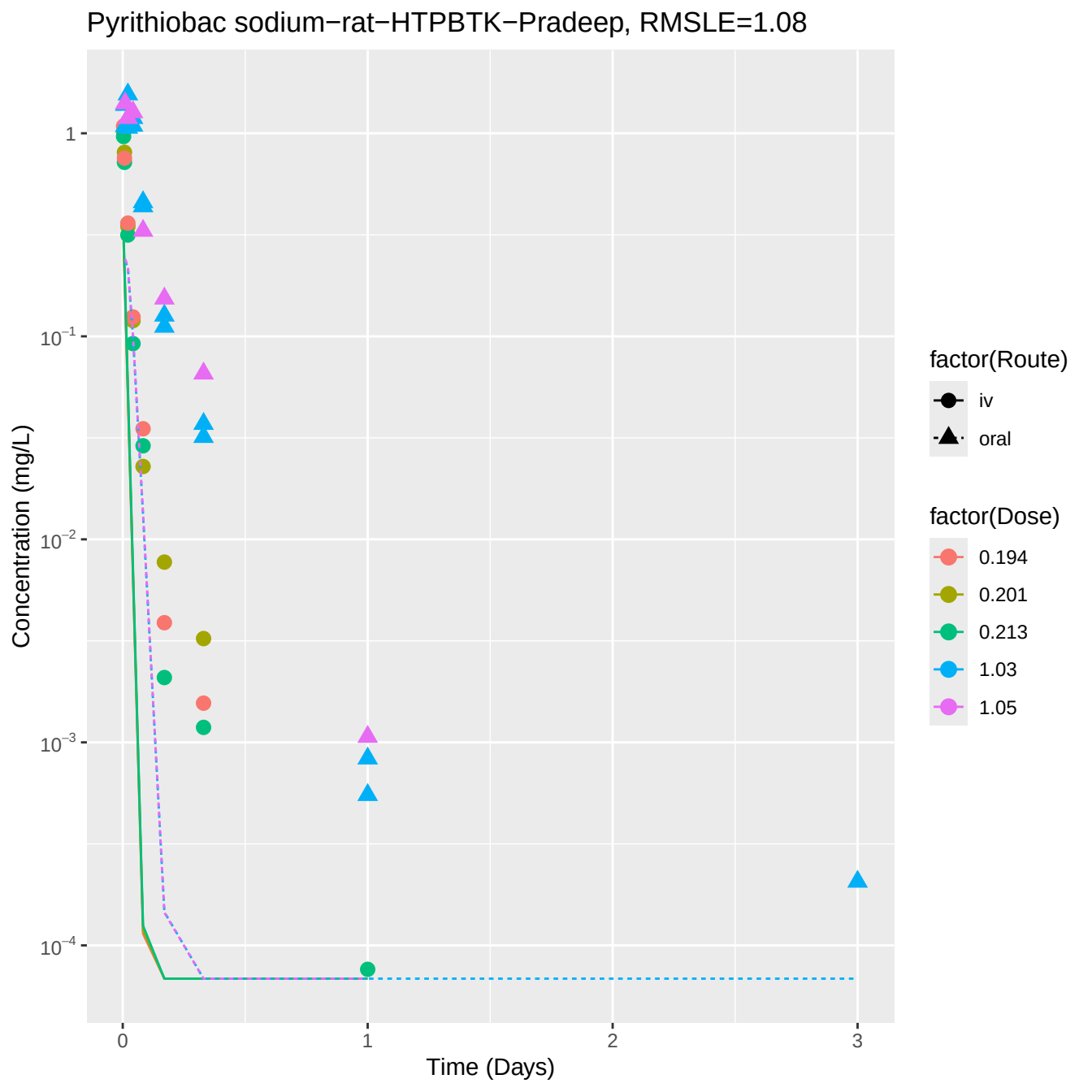

Simazine−rat−HTPBTK−Ensemble, RMSLE=0.957
Concentration (mg/L)
10^{-1}
10^{-2}
10^{-3}
0.00
0.05
0.10
0.15
Time (Days)
factor(Dose)
0.4
2
factor(Route)
iv
oral

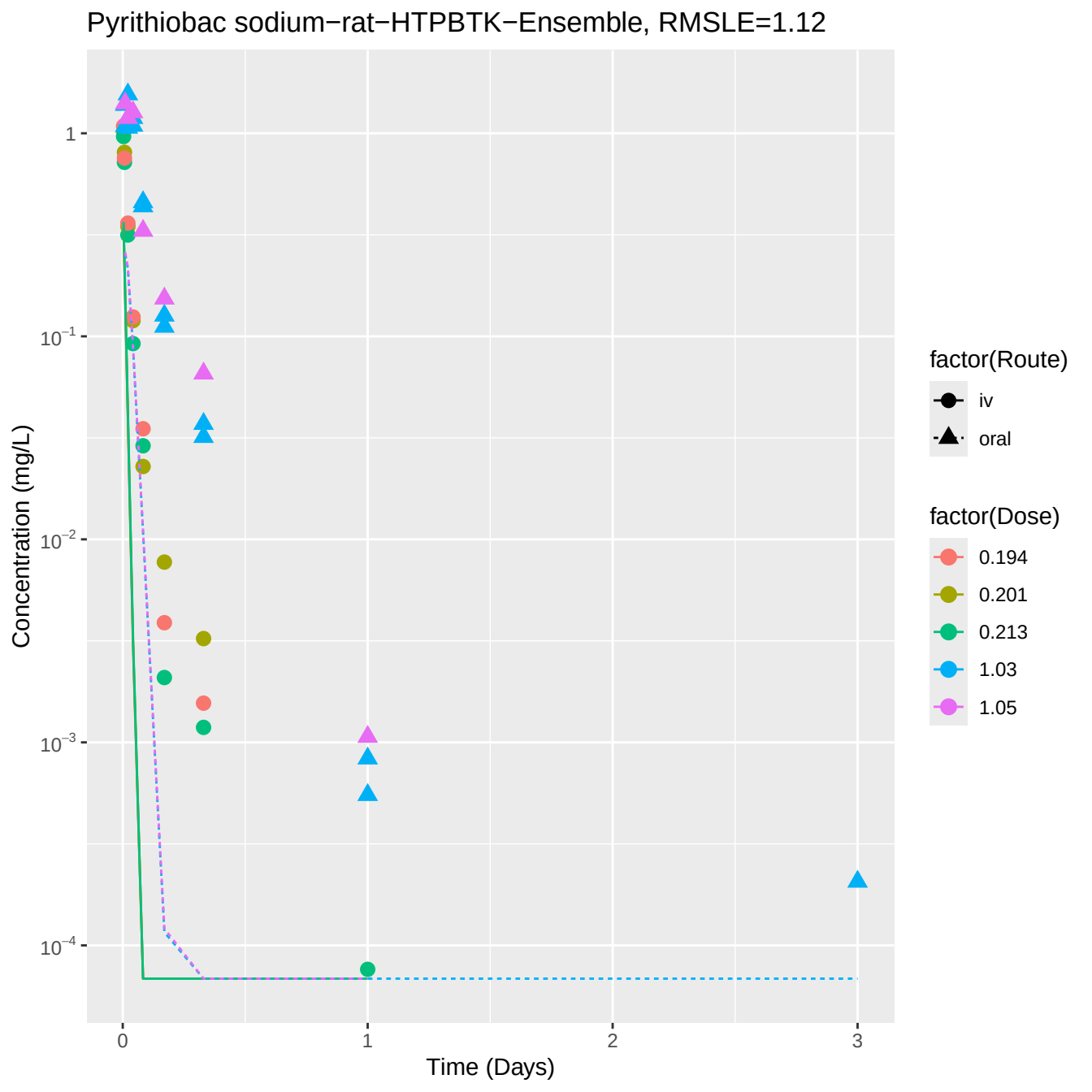

Simazine−rat−In Vivo Fits, RMSLE=0.325
Concentration (mg/L)
10^{-1}
10^{-2}
10^{-3}
0.00
0.05
0.10
0.15
Time (Days)
factor(Dose)
0.4
2
factor(Route)
iv
oral

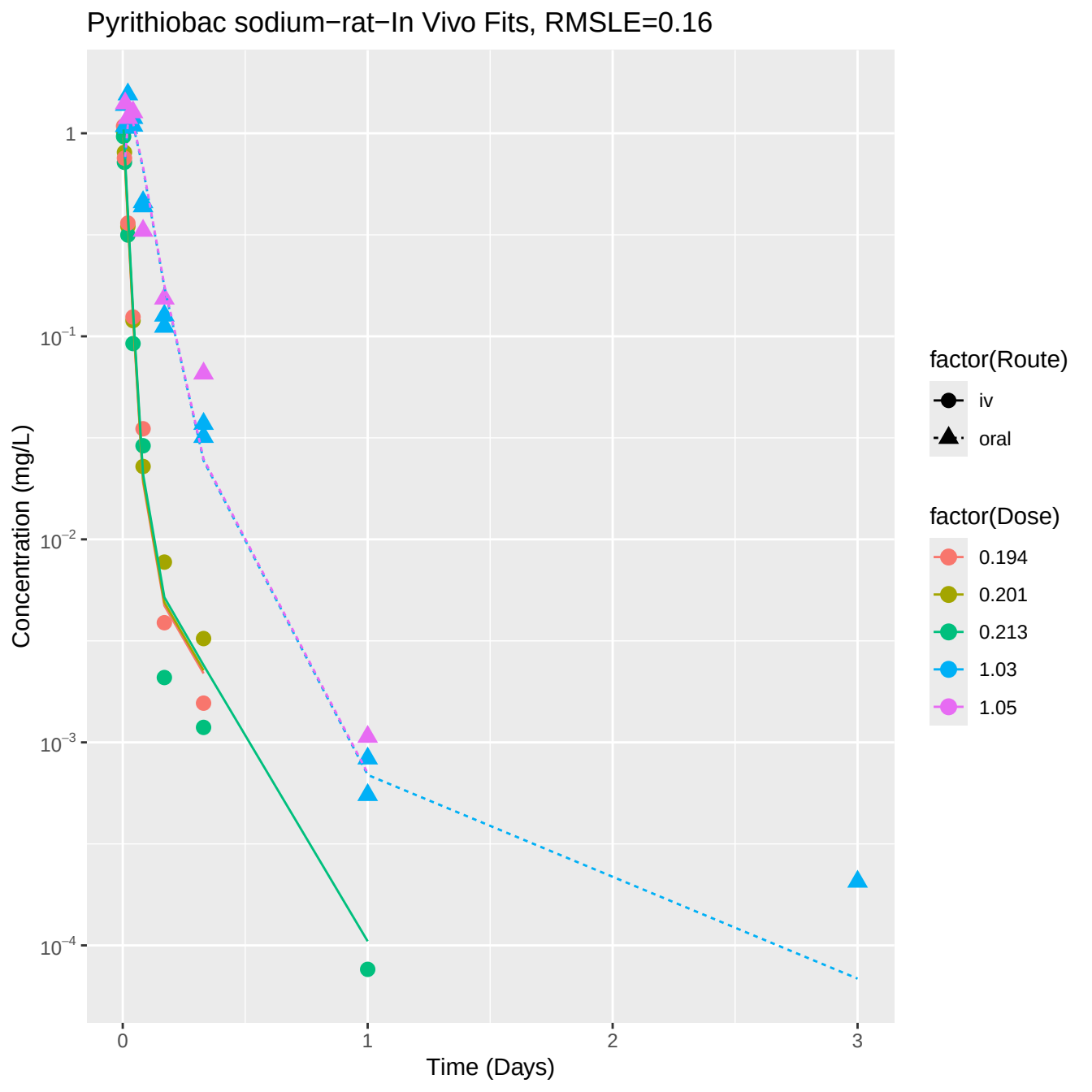

Pyrithiobac sodium−rat−HTPBTK−InVitro, RMSLE=1.39
Concentration (mg/L)
1
10^{-1}
10^{-2}
10^{-3}
10^{-4}
0
1
2
3
Time (Days)
factor(Route)
iv
oral
factor(Dose)
0.194
0.201
0.213
1.03
1.05

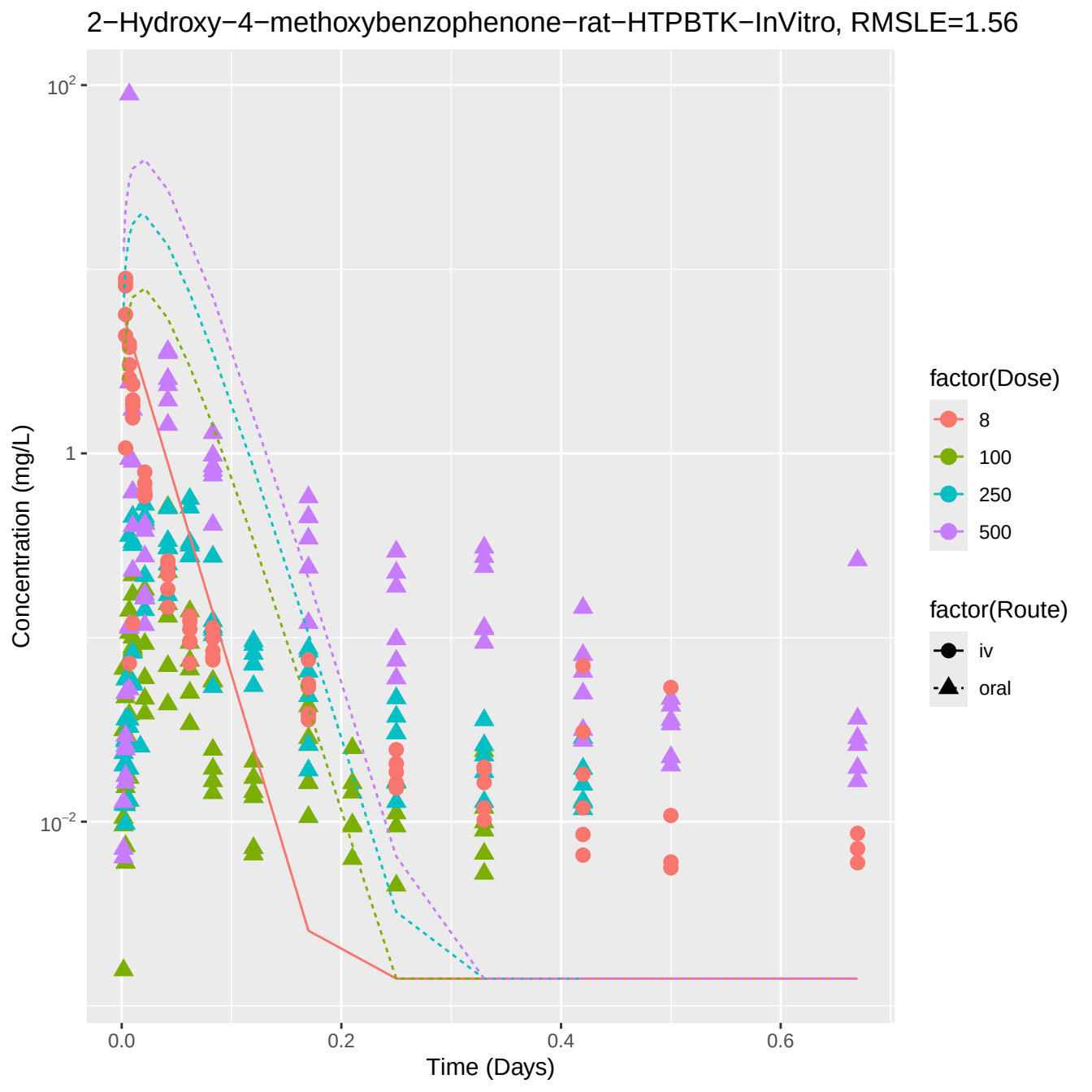

Pyrithiobac sodium−rat−HTPBTK−ADMET, RMSLE=0.939
Concentration (mg/L)
1
10^{-1}
10^{-2}
10^{-3}
10^{-4}
0
1
2
3
Time (Days)
factor(Route)
iv
oral
factor(Dose)
0.194
0.201
0.213
1.03
1.05

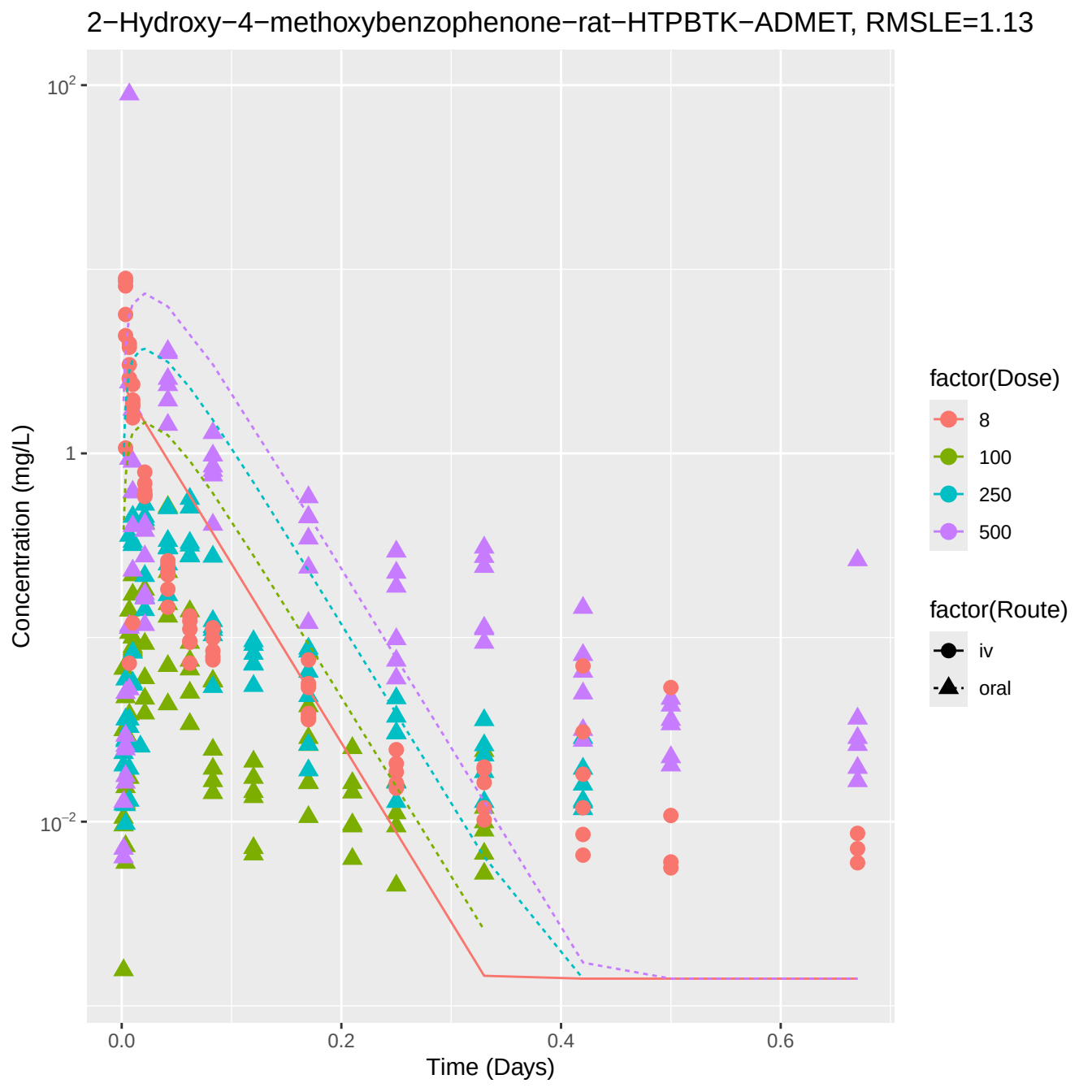

Pyrithiobac sodium−rat−HTPBTK−Dawson, RMSLE=0.97
Concentration (mg/L)
1
10^{-1}
10^{-2}
10^{-3}
10^{-4}
0
1
2
3
Time (Days)
factor(Route)
iv
oral
factor(Dose)
0.194
0.201
0.213
1.03
1.05

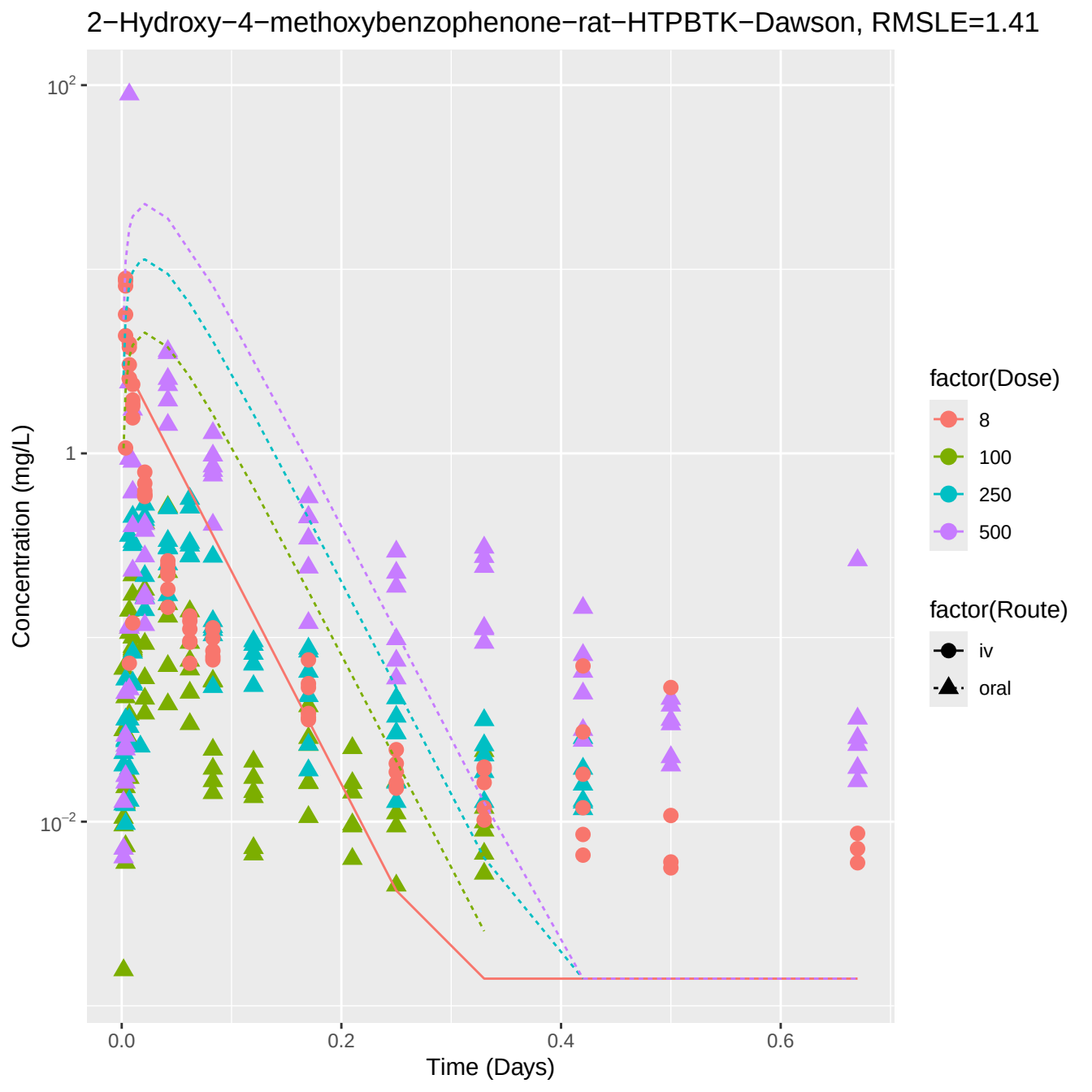

Pyrithiobac sodium−rat−HTPBTK−Pradeep, RMSLE=1.08
Concentration (mg/L)
1
10^{-1}
10^{-2}
10^{-3}
10^{-4}
0
1
2
3
Time (Days)
factor(Route)
iv
oral
factor(Dose)
0.194
0.201
0.213
1.03
1.05

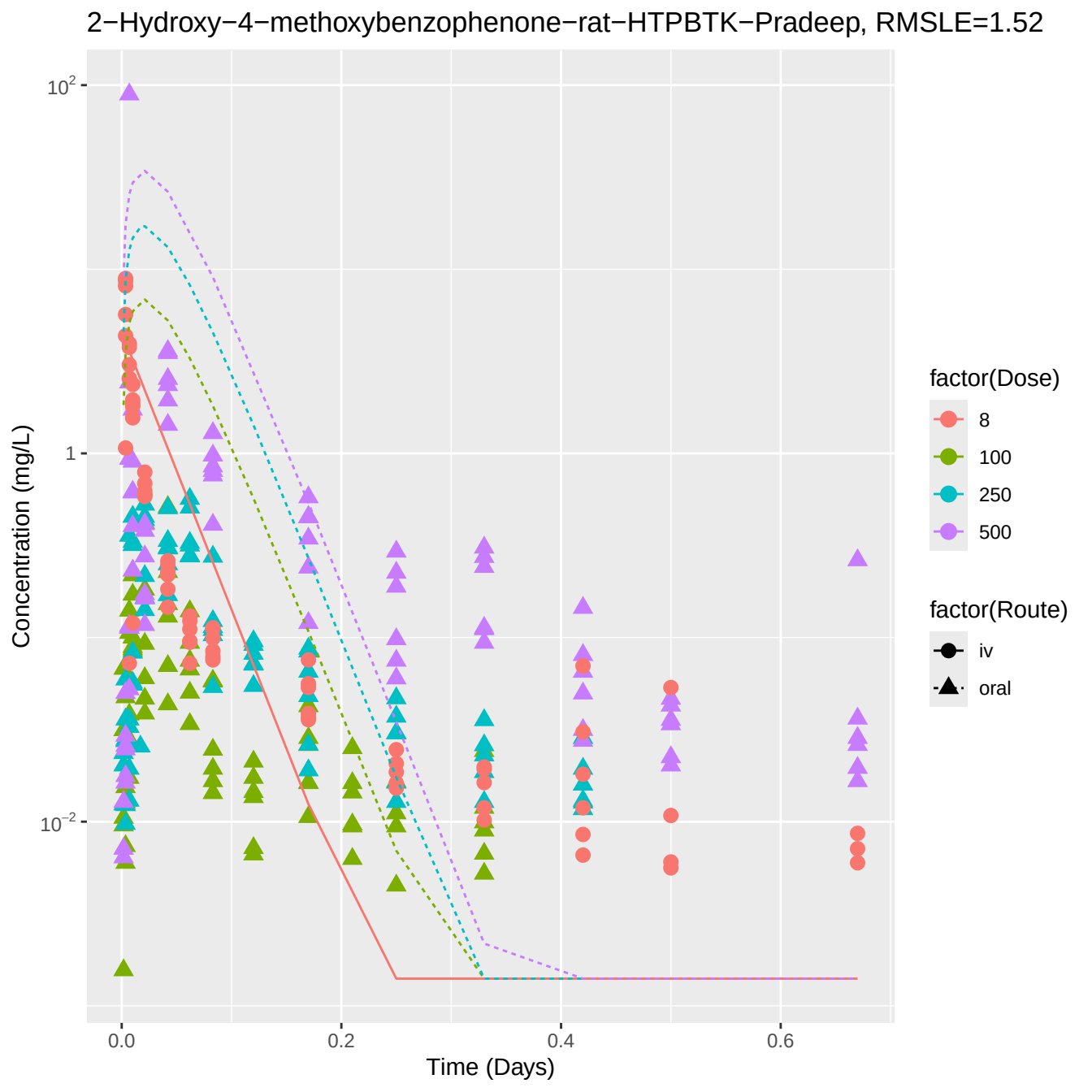

Pyrithiobac sodium−rat−HTPBTK−Ensemble, RMSLE=1.12
Concentration (mg/L)
1
10^{-1}
10^{-2}
10^{-3}
10^{-4}
0
1
2
3
Time (Days)
factor(Route)
iv
oral
factor(Dose)
0.194
0.201
0.213
1.03
1.05

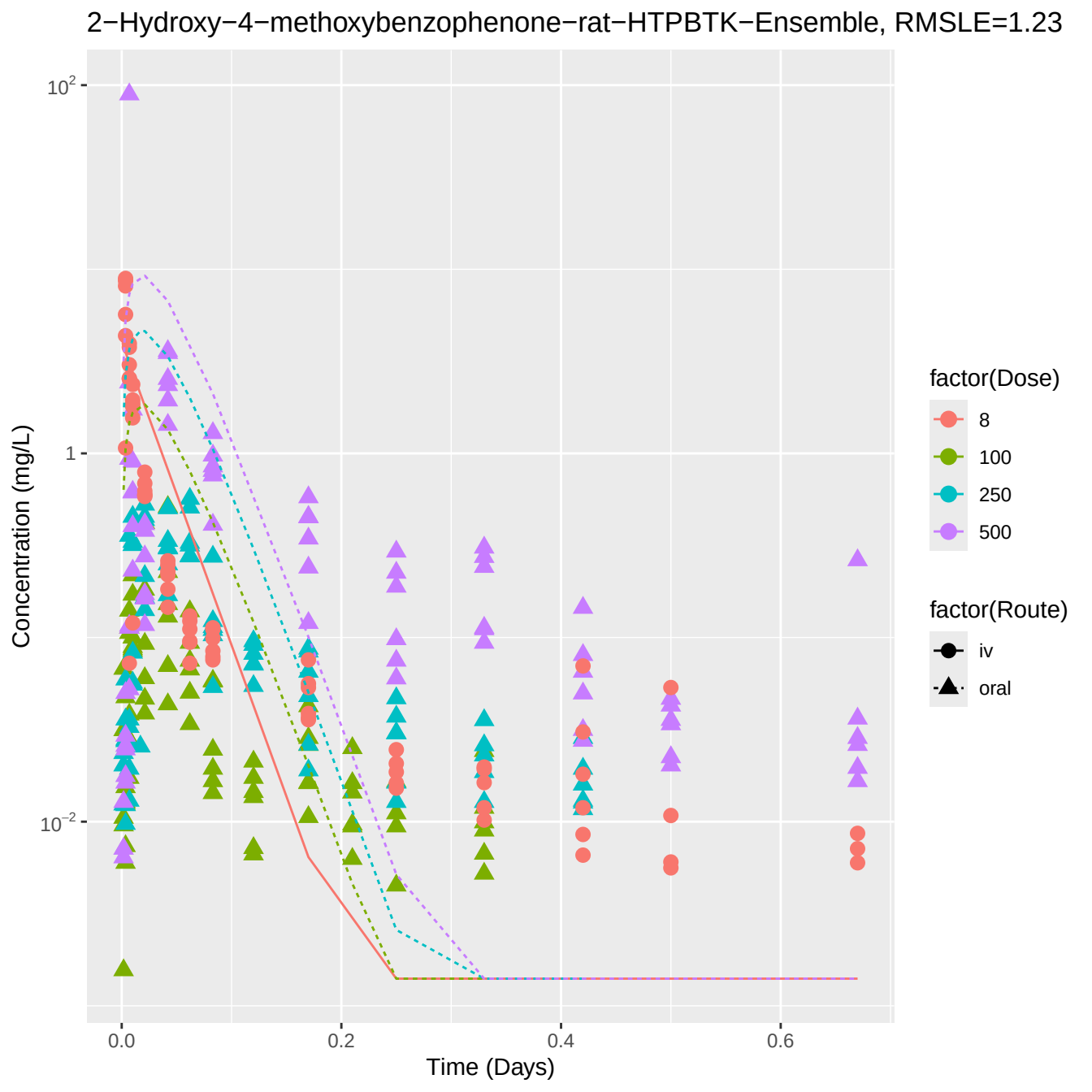

Pyrithiobac sodium−rat−In Vivo Fits, RMSLE=0.16
Concentration (mg/L)
1
10^{-1}
10^{-2}
10^{-3}
10^{-4}
0
1
2
3
Time (Days)
factor(Route)
iv
oral
factor(Dose)
0.194
0.201
0.213
1.03
1.05

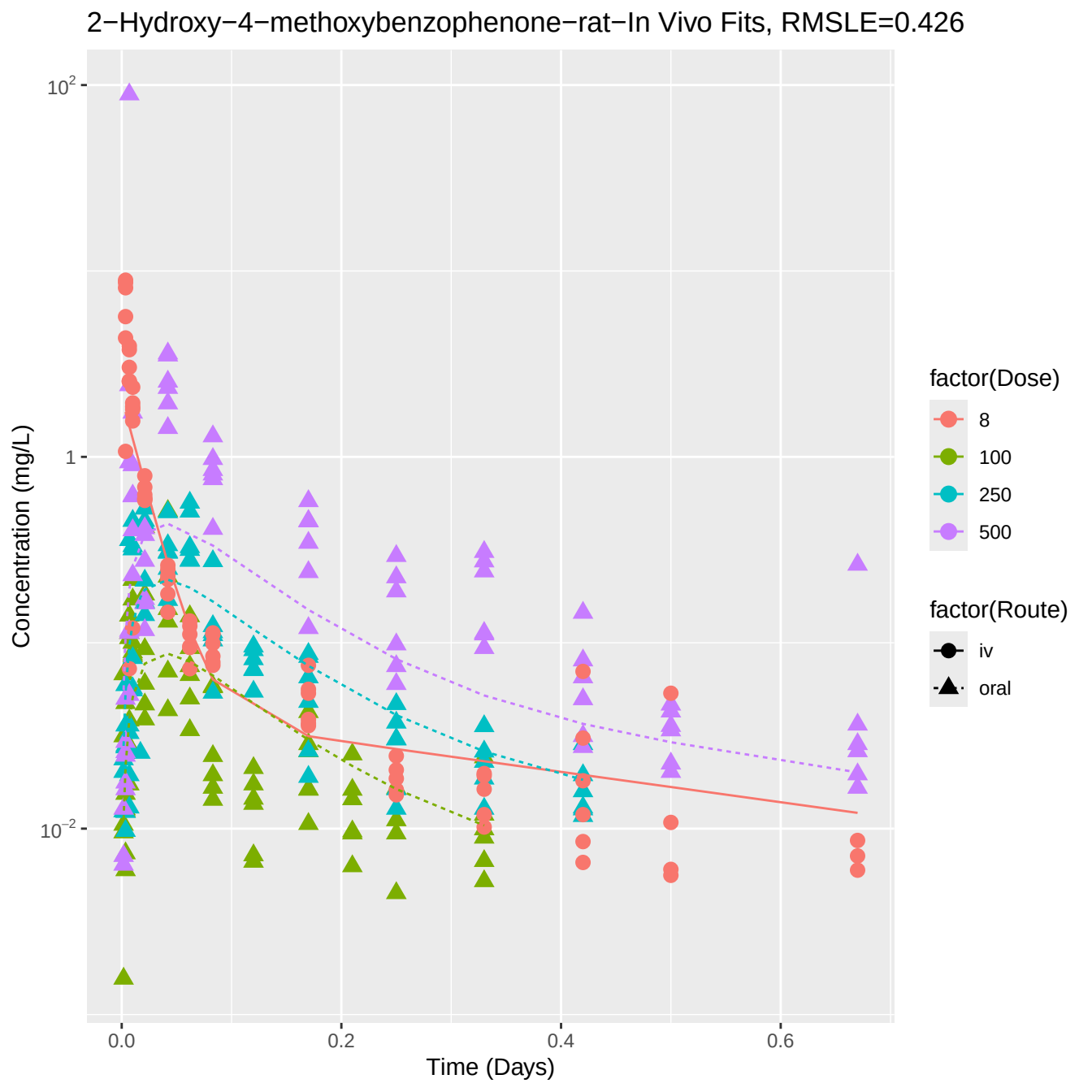

2−Hydroxy−4−methoxybenzophenone−rat−HTPBTK−InVitro, RMSLE=1.56
Concentration (mg/L)
10^2
1
10^{-2}
0.0
0.2
0.4
0.6
Time (Days)
factor(Dose)
8
100
250
500
factor(Route)
iv
oral

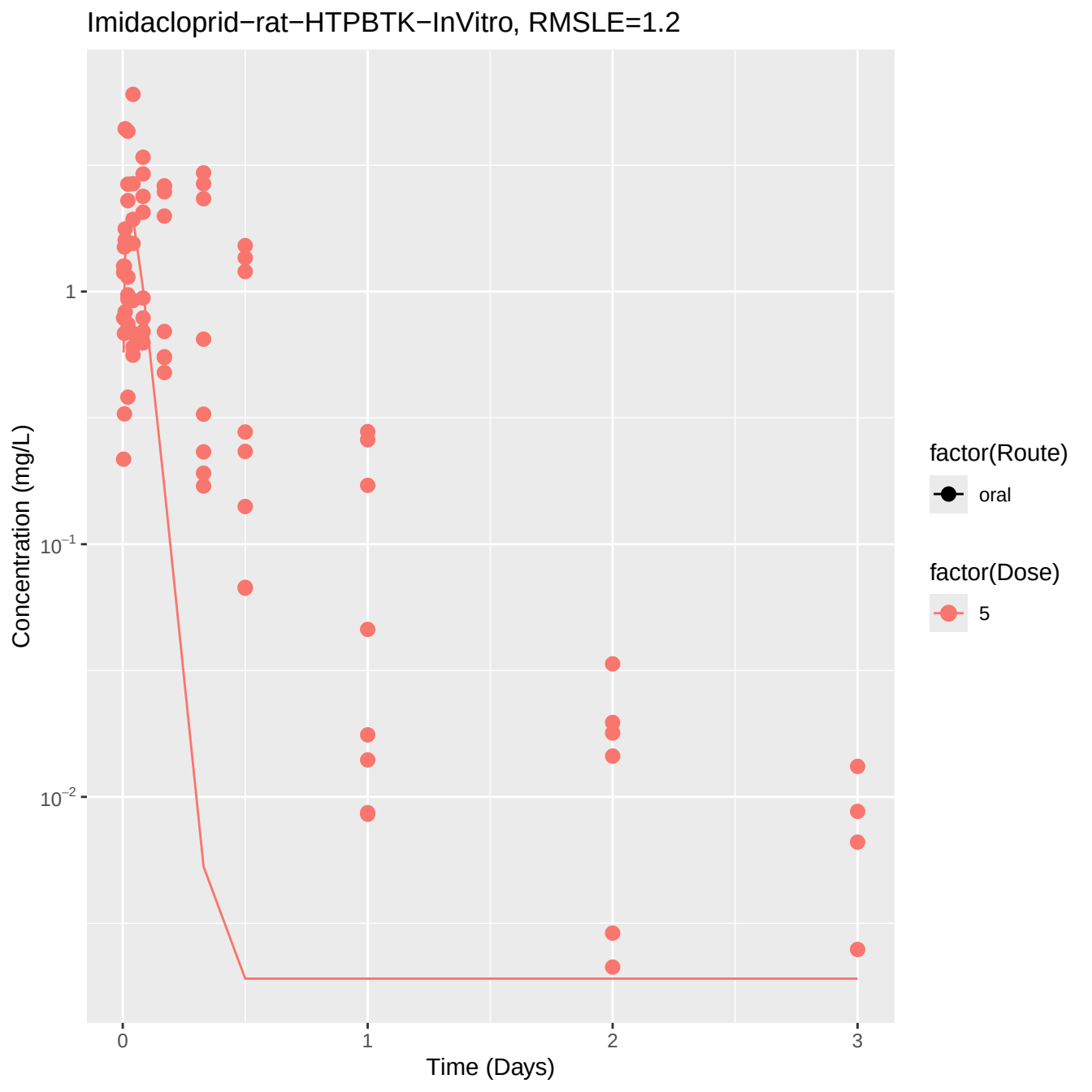

2−Hydroxy−4−methoxybenzophenone−rat−HTPBTK−ADMET, RMSLE=1.13
Concentration (mg/L)
10^2
1
10^{-2}
0.0
0.2
0.4
0.6
Time (Days)
factor(Dose)
8
100
250
500
factor(Route)
iv
oral

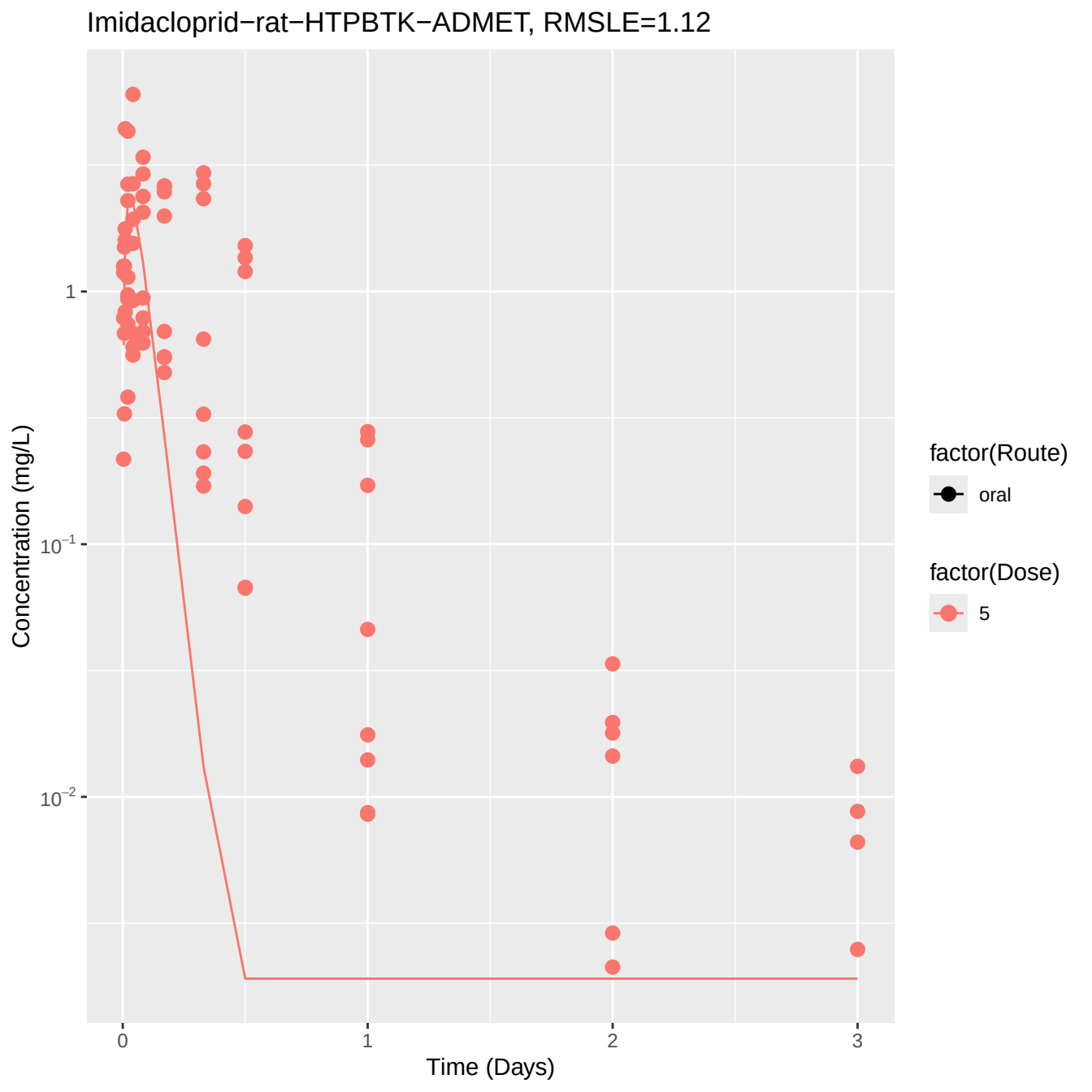

2−Hydroxy−4−methoxybenzophenone−rat−HTPBTK−Dawson, RMSLE=1.41
Concentration (mg/L)
10^2
1
10^{-2}
0.0
0.2
0.4
0.6
Time (Days)
factor(Dose)
8
100
250
500
factor(Route)
iv
oral

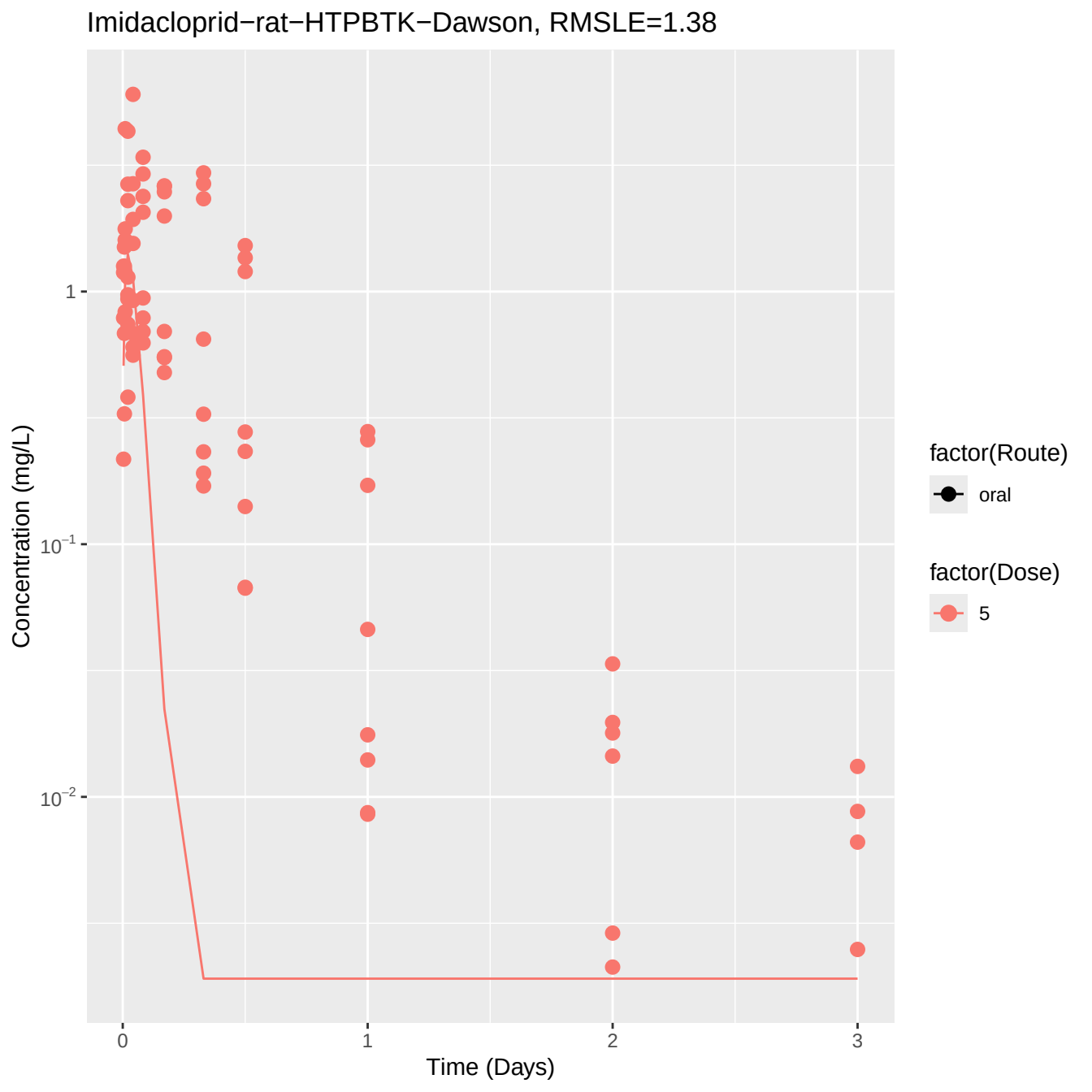

2−Hydroxy−4−methoxybenzophenone−rat−HTPBTK−Pradeep, RMSLE=1.52
Concentration (mg/L)
10^2
1
10^{-2}
0.0
0.2
0.4
0.6
Time (Days)
factor(Dose)
8
100
250
500
factor(Route)
iv
oral

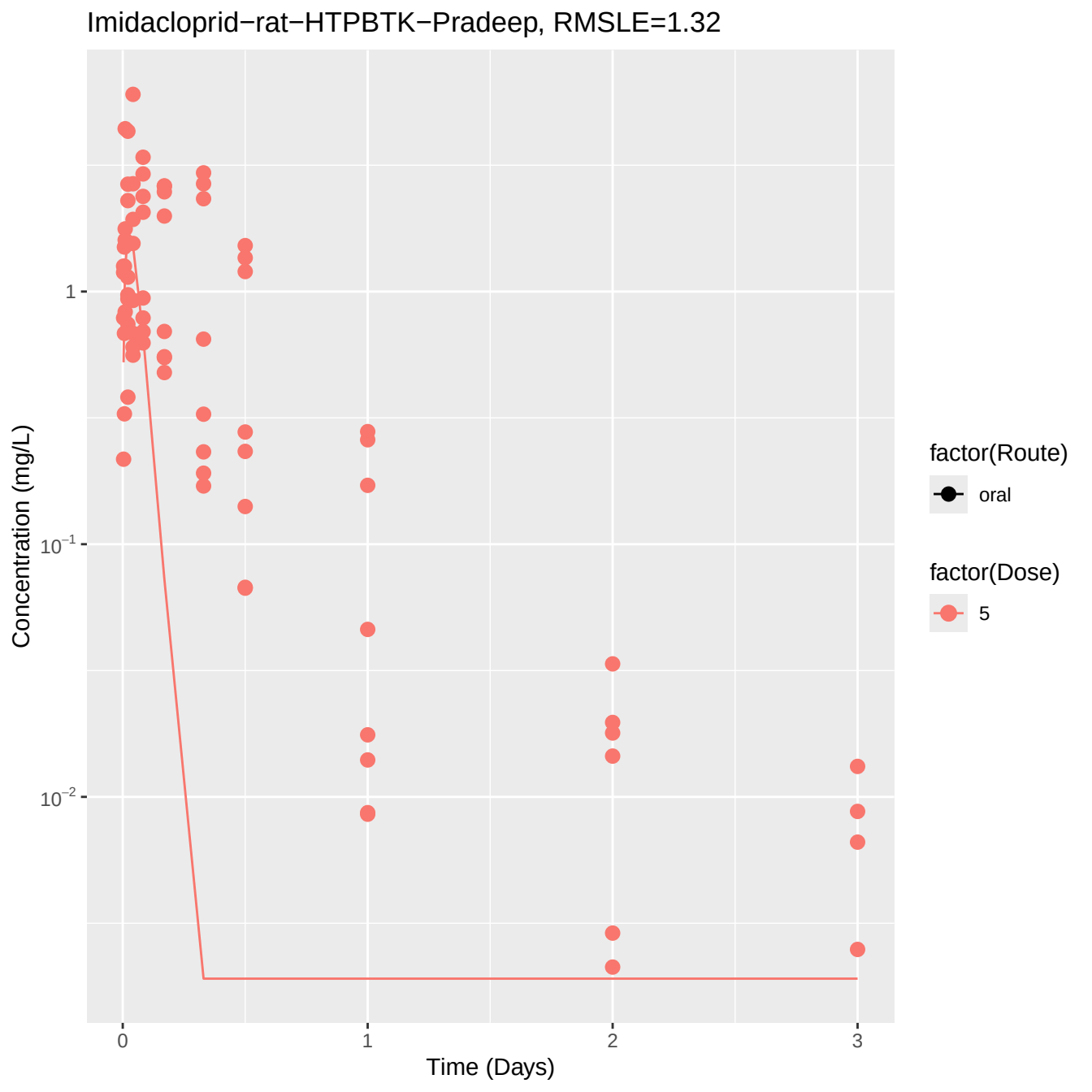

2−Hydroxy−4−methoxybenzophenone−rat−HTPBTK−Ensemble, RMSLE=1.23
Concentration (mg/L)
10^2
1
10^{-2}
0.0
0.2
0.4
0.6
Time (Days)
factor(Dose)
8
100
250
500
factor(Route)
iv
oral

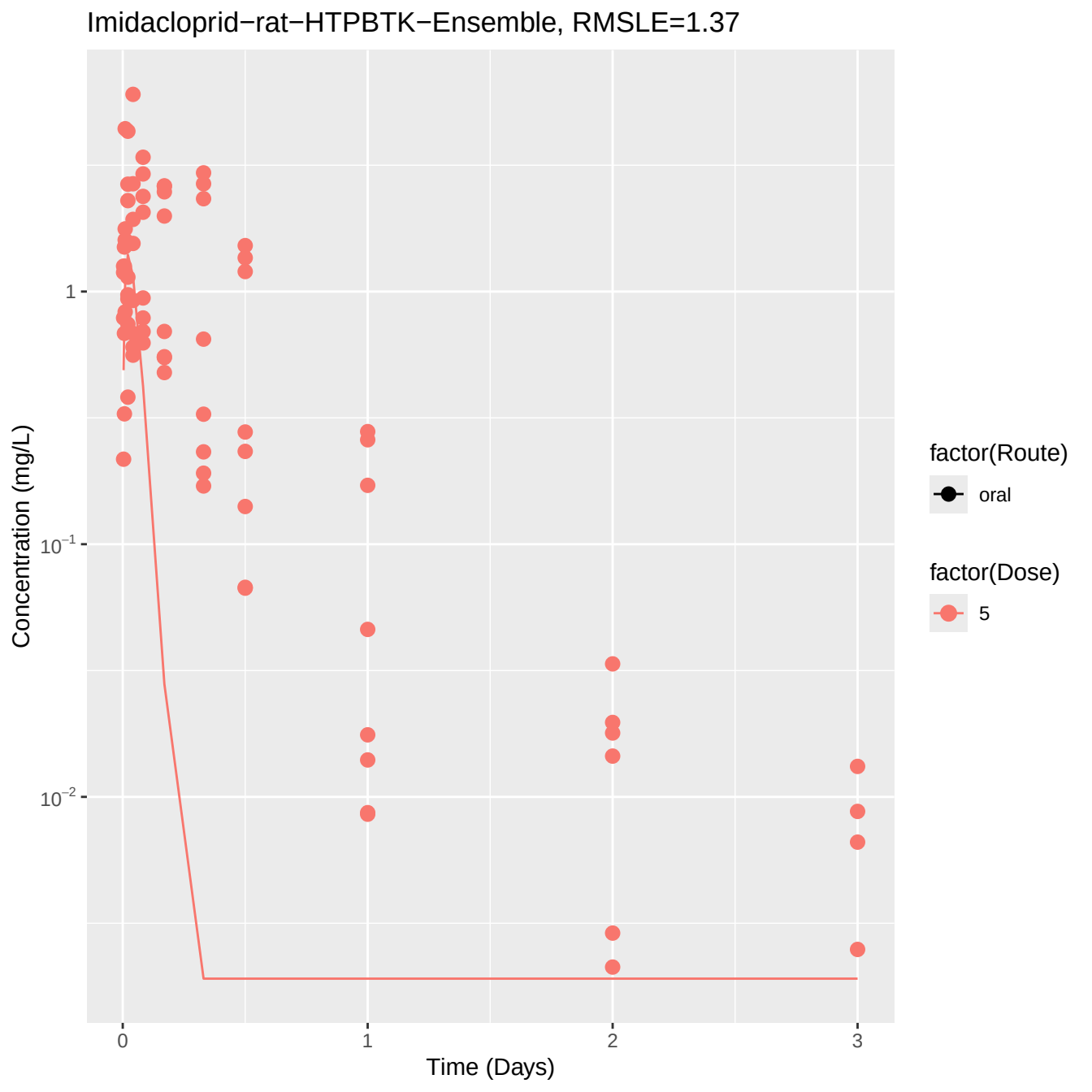

2−Hydroxy−4−methoxybenzophenone−rat−In Vivo Fits, RMSLE=0.426
Concentration (mg/L)
10^2
1
10^{-2}
0.0
0.2
0.4
0.6
Time (Days)
factor(Dose)
8
100
250
500
factor(Route)
iv
oral

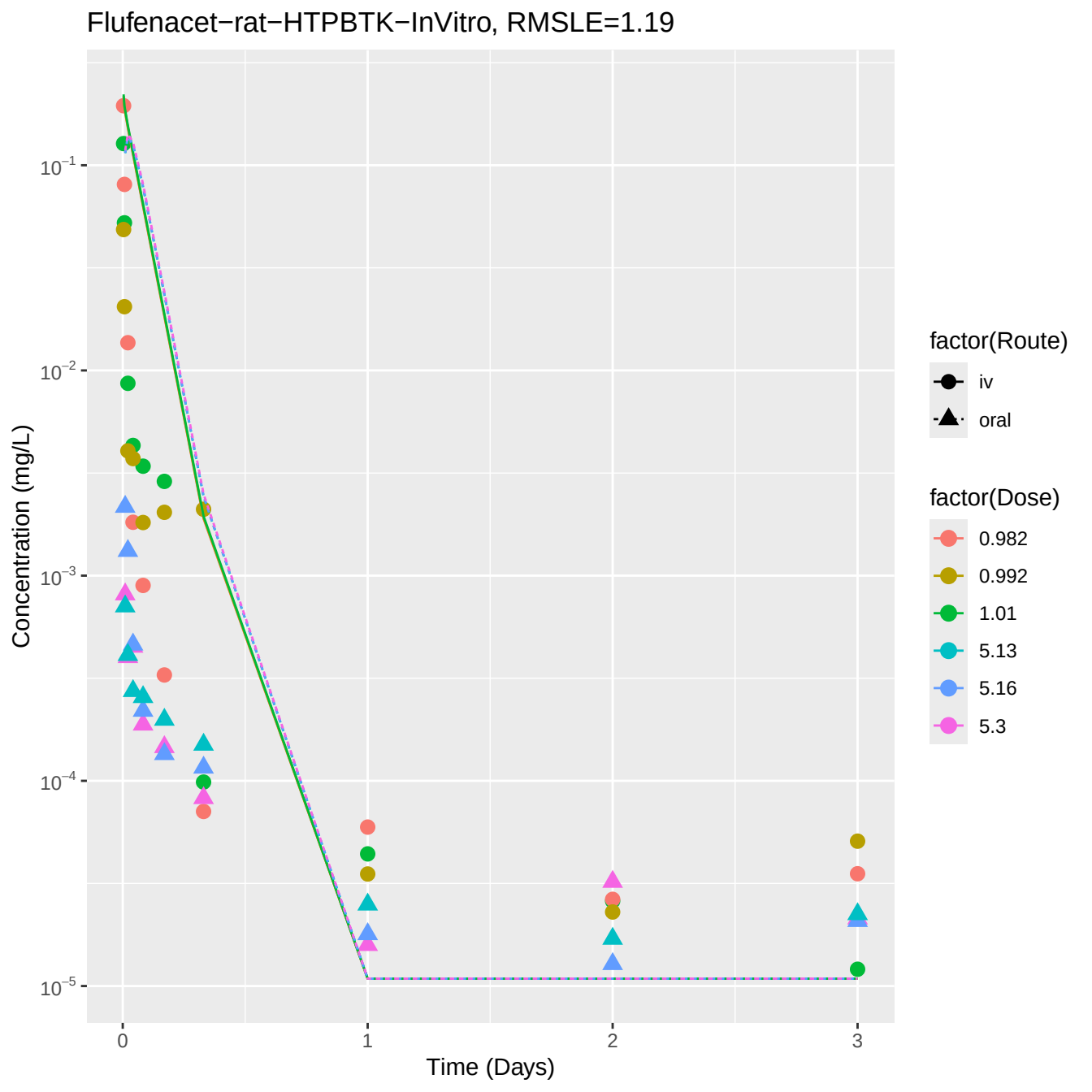

Imidacloprid−rat−HTPBTK−InVitro, RMSLE=1.2
Concentration (mg/L)
1
10^{-1}
10^{-2}
0
1
2
3
Time (Days)
factor(Route)
oral
factor(Dose)
5

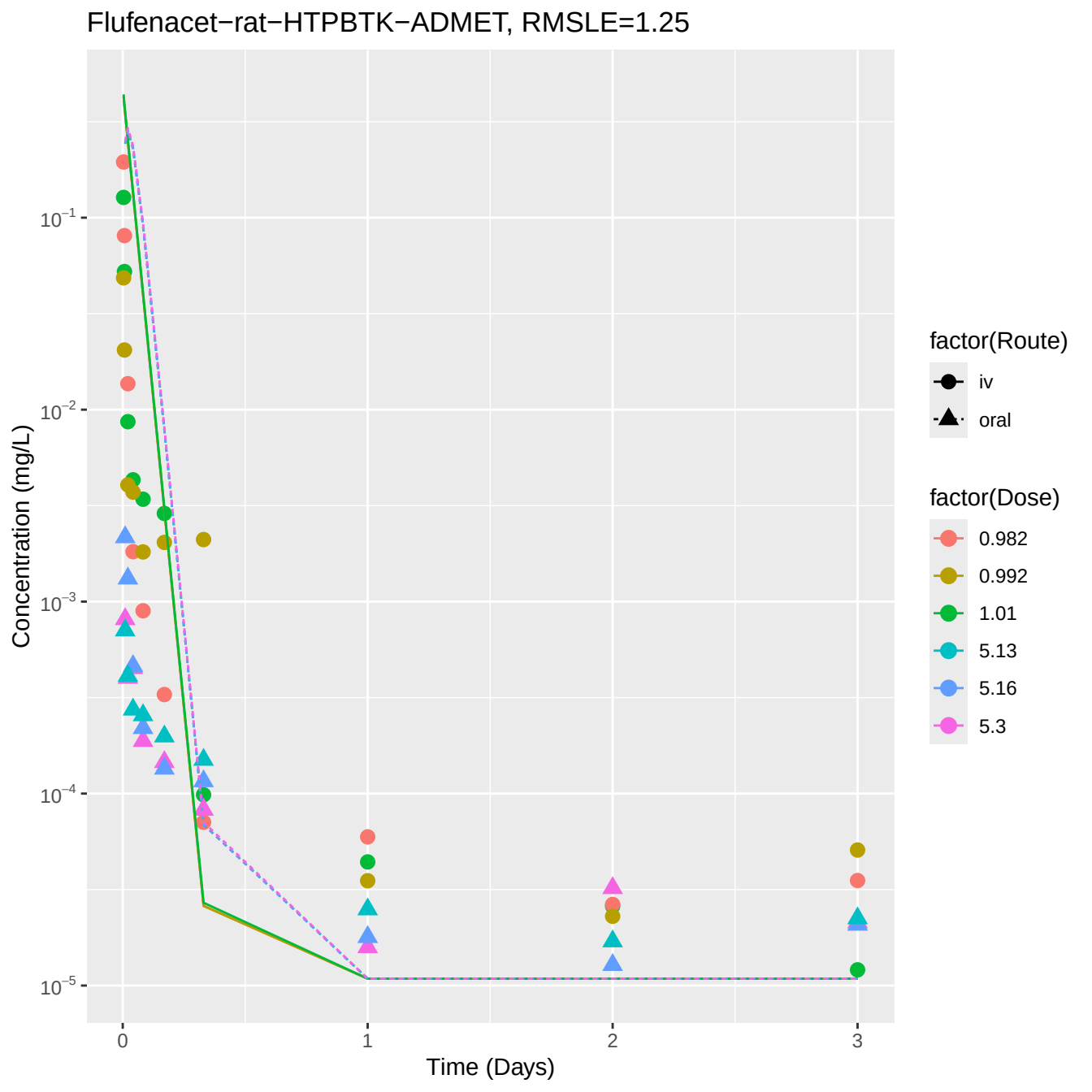

Imidacloprid−rat−HTPBTK−ADMET, RMSLE=1.12
Concentration (mg/L)
1
10^{-1}
10^{-2}
0
1
2
3
Time (Days)
factor(Route)
oral
factor(Dose)
5

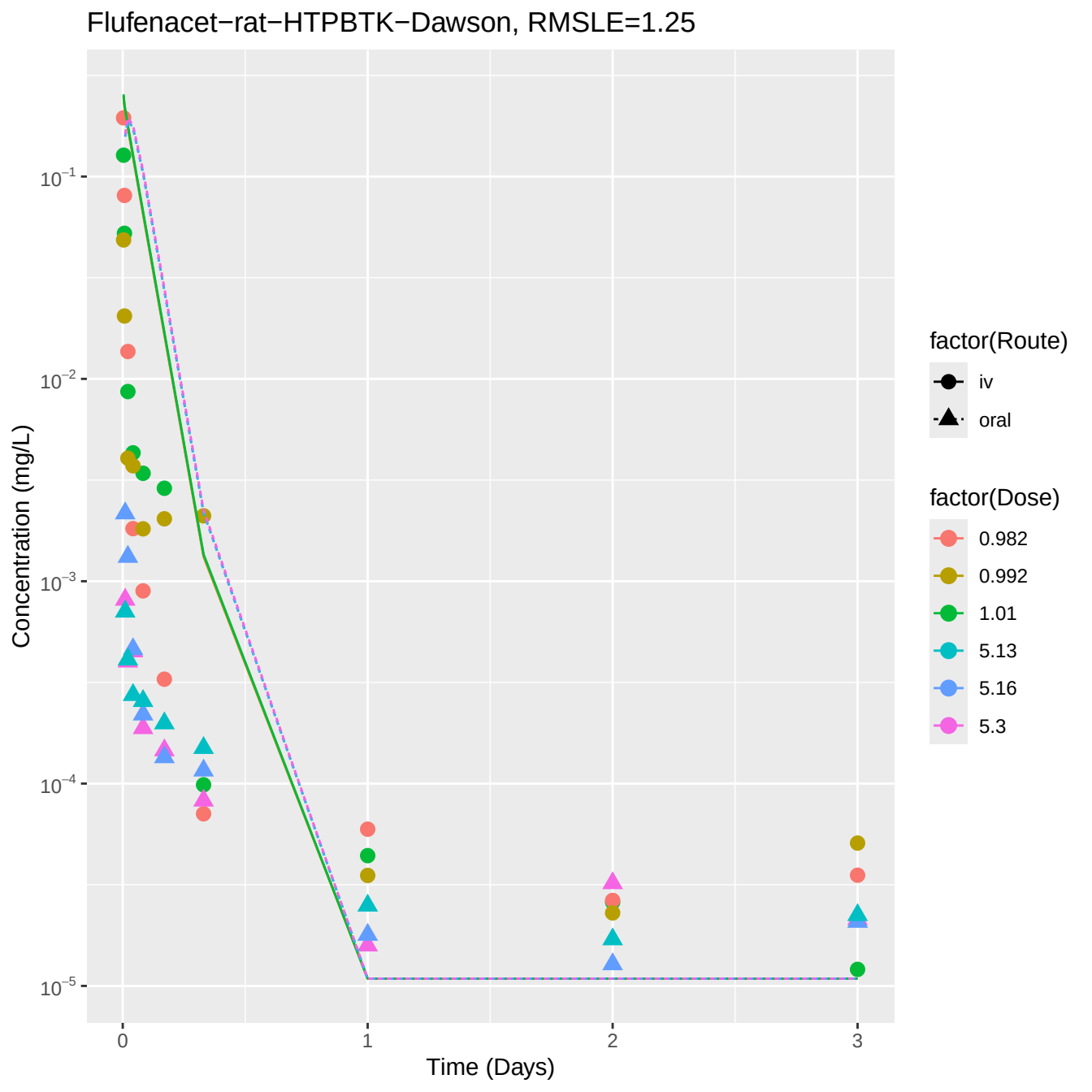

Imidacloprid−rat−HTPBTK−Dawson, RMSLE=1.38
Concentration (mg/L)
1
10^{-1}
10^{-2}
0
1
2
3
Time (Days)
factor(Route)
oral
factor(Dose)
5

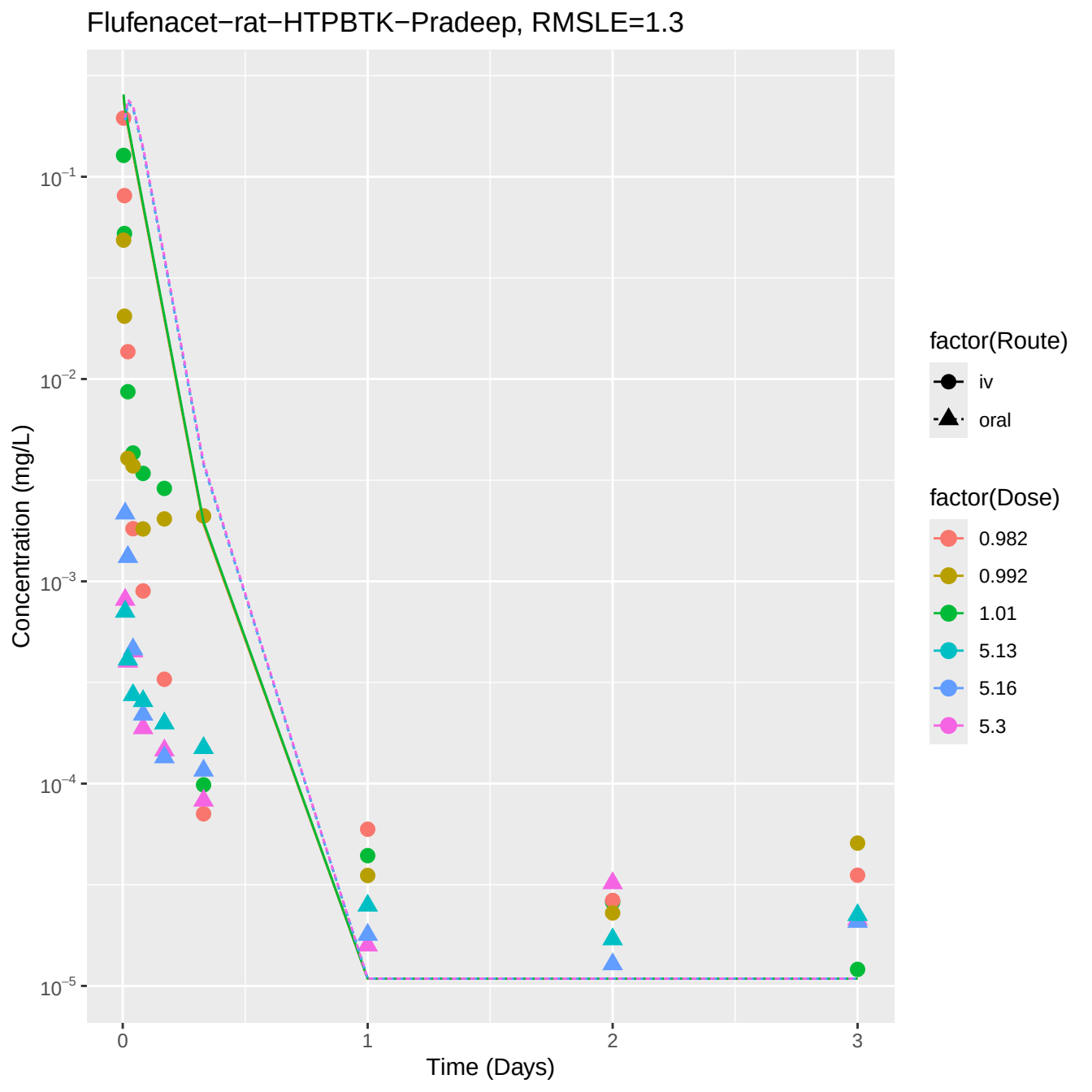

Imidacloprid−rat−HTPBTK−Pradeep, RMSLE=1.32
Concentration (mg/L)
1
10^{-1}
10^{-2}
0
1
2
3
Time (Days)
factor(Route)
oral
factor(Dose)
5

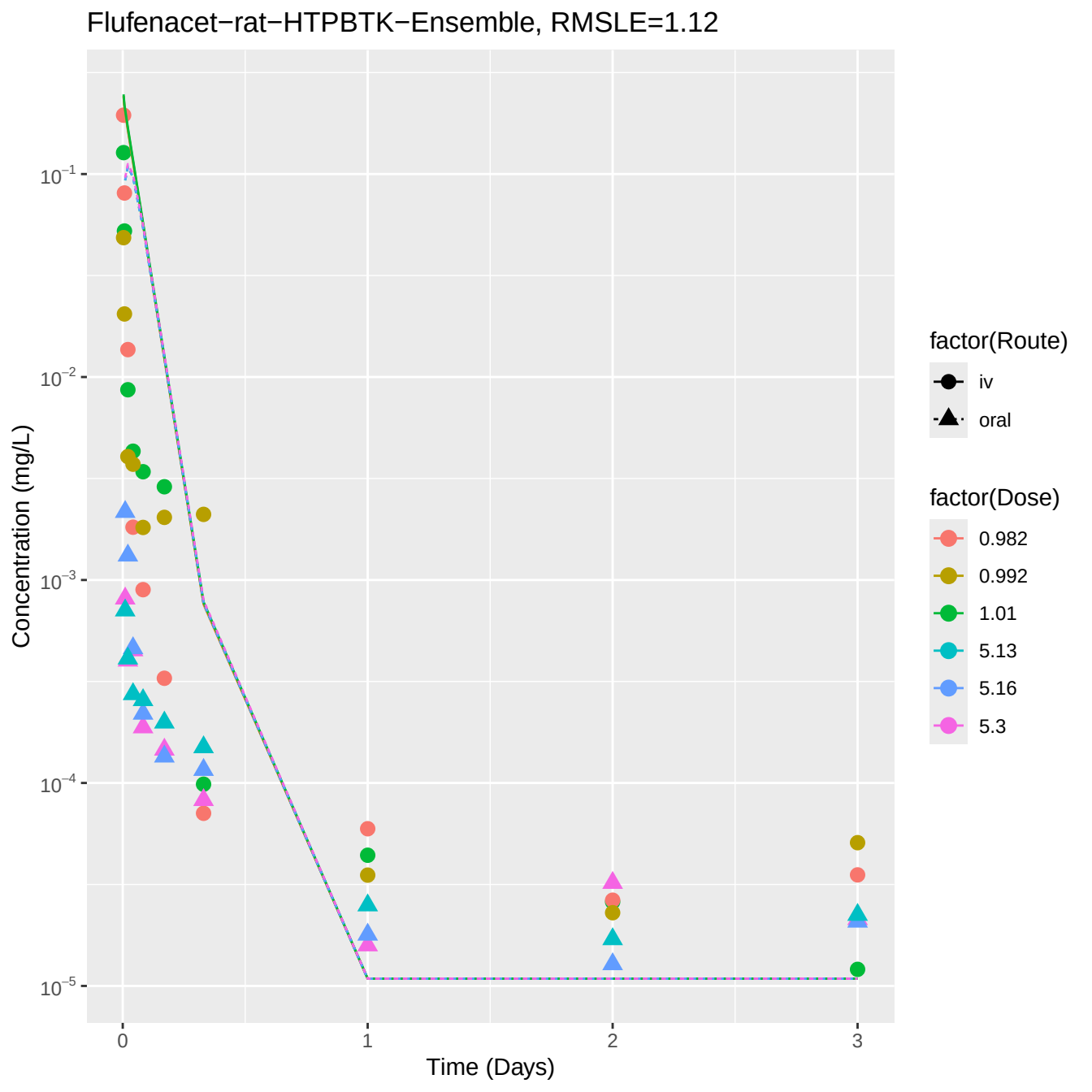

Imidacloprid−rat−HTPBTK−Ensemble, RMSLE=1.37
Concentration (mg/L)
1
10^{-1}
10^{-2}
0
1
2
3
Time (Days)
factor(Route)
oral
factor(Dose)
5

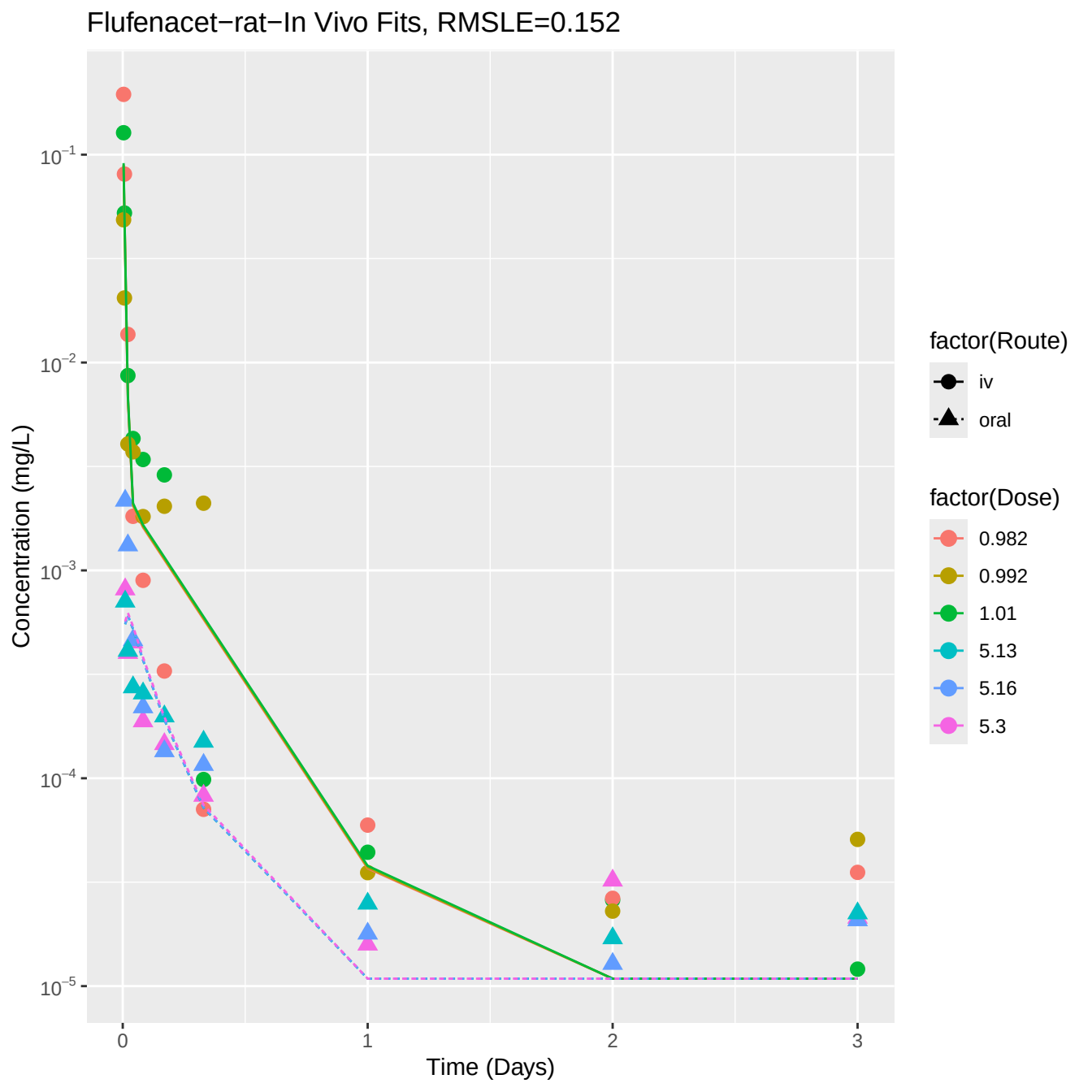

Flufenacet−rat−HTPBTK−InVitro, RMSLE=1.19
Concentration (mg/L)
10^{-1}
10^{-2}
10^{-3}
10^{-4}
10^{-5}
0
1
2
3
Time (Days)
factor(Route)
iv
oral
factor(Dose)
0.982
0.992
1.01
5.13
5.16
5.3

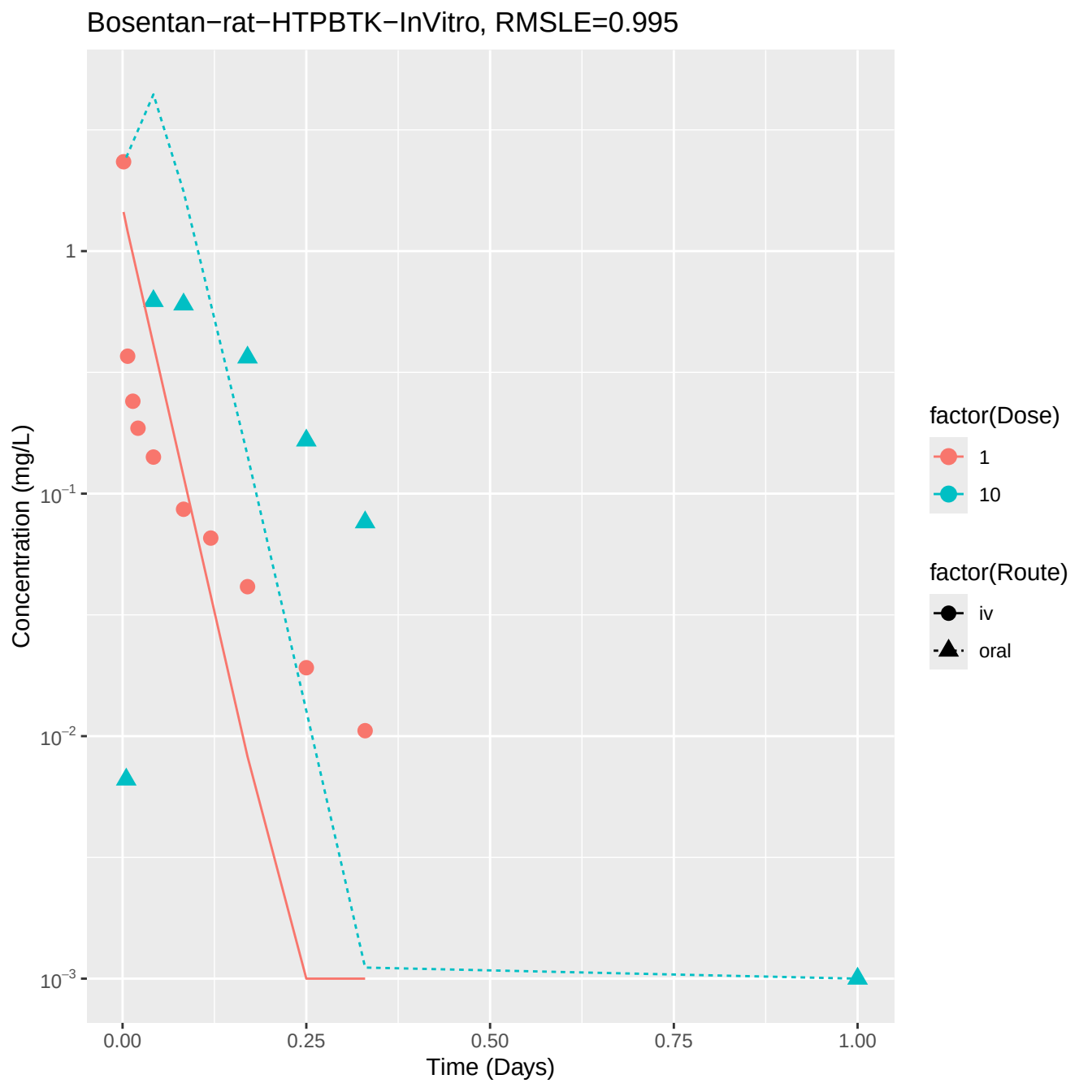

Flufenacet−rat−HTPBTK−ADMET, RMSLE=1.25
Concentration (mg/L)
10^{-1}
10^{-2}
10^{-3}
10^{-4}
10^{-5}
0
1
2
3
Time (Days)
factor(Route)
iv
oral
factor(Dose)
0.982
0.992
1.01
5.13
5.16
5.3

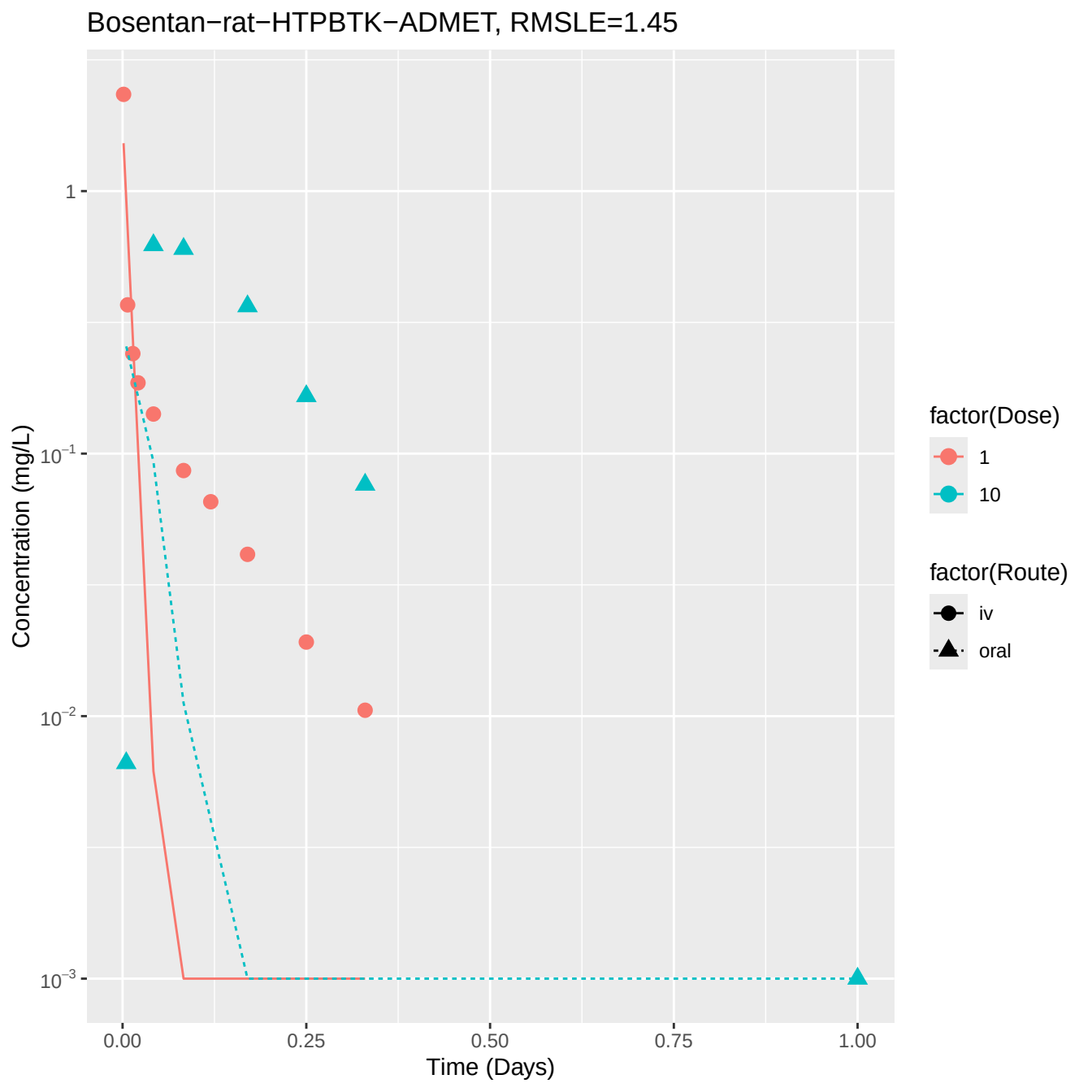

Flufenacet−rat−HTPBTK−Dawson, RMSLE=1.25
Concentration (mg/L)
10^{-1}
10^{-2}
10^{-3}
10^{-4}
10^{-5}
0
1
2
3
Time (Days)
factor(Route)
iv
oral
factor(Dose)
0.982
0.992
1.01
5.13
5.16
5.3

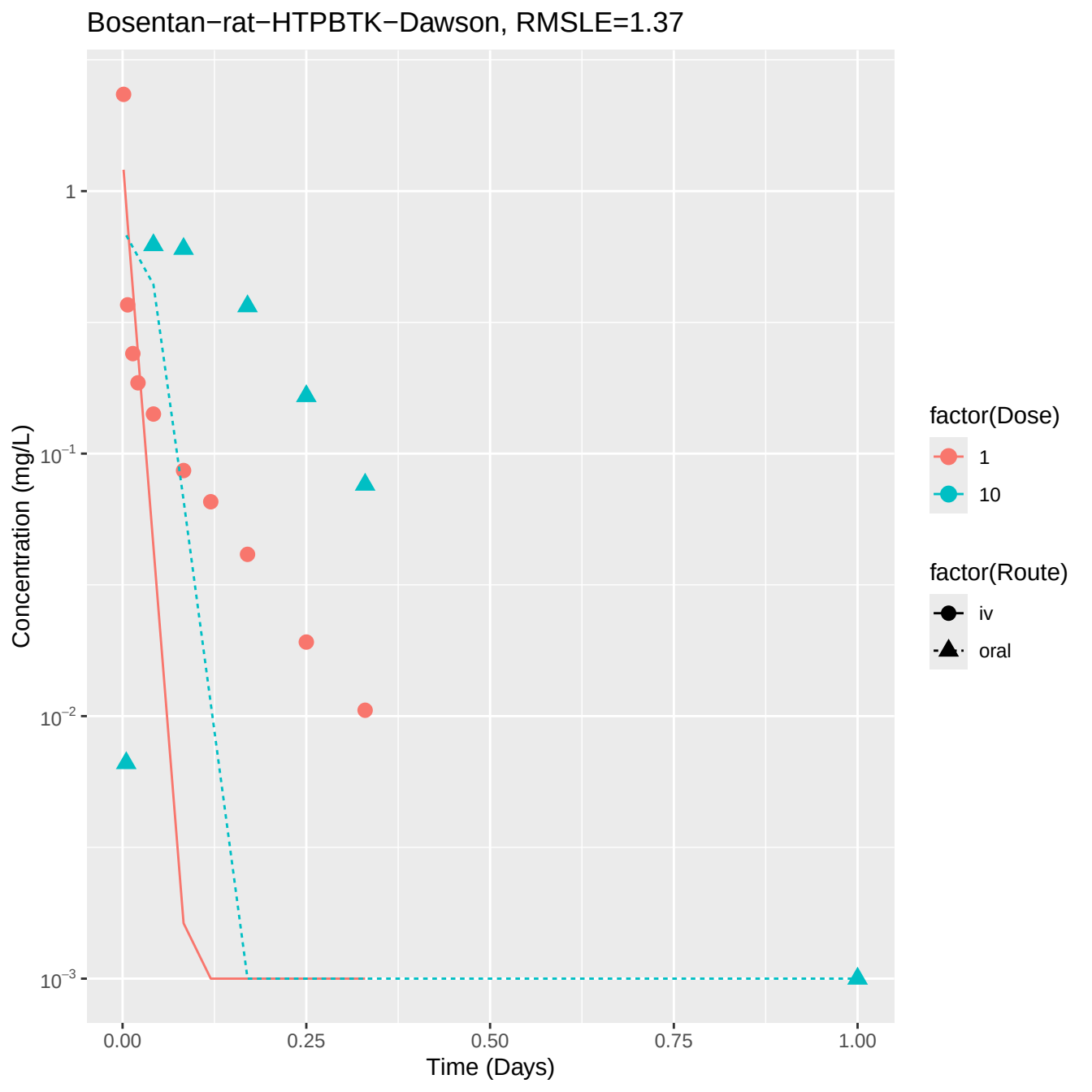

Flufenacet−rat−HTPBTK−Pradeep, RMSLE=1.3
Concentration (mg/L)
10^{-1}
10^{-2}
10^{-3}
10^{-4}
10^{-5}
0
1
2
3
Time (Days)
factor(Route)
iv
oral
factor(Dose)
0.982
0.992
1.01
5.13
5.16
5.3

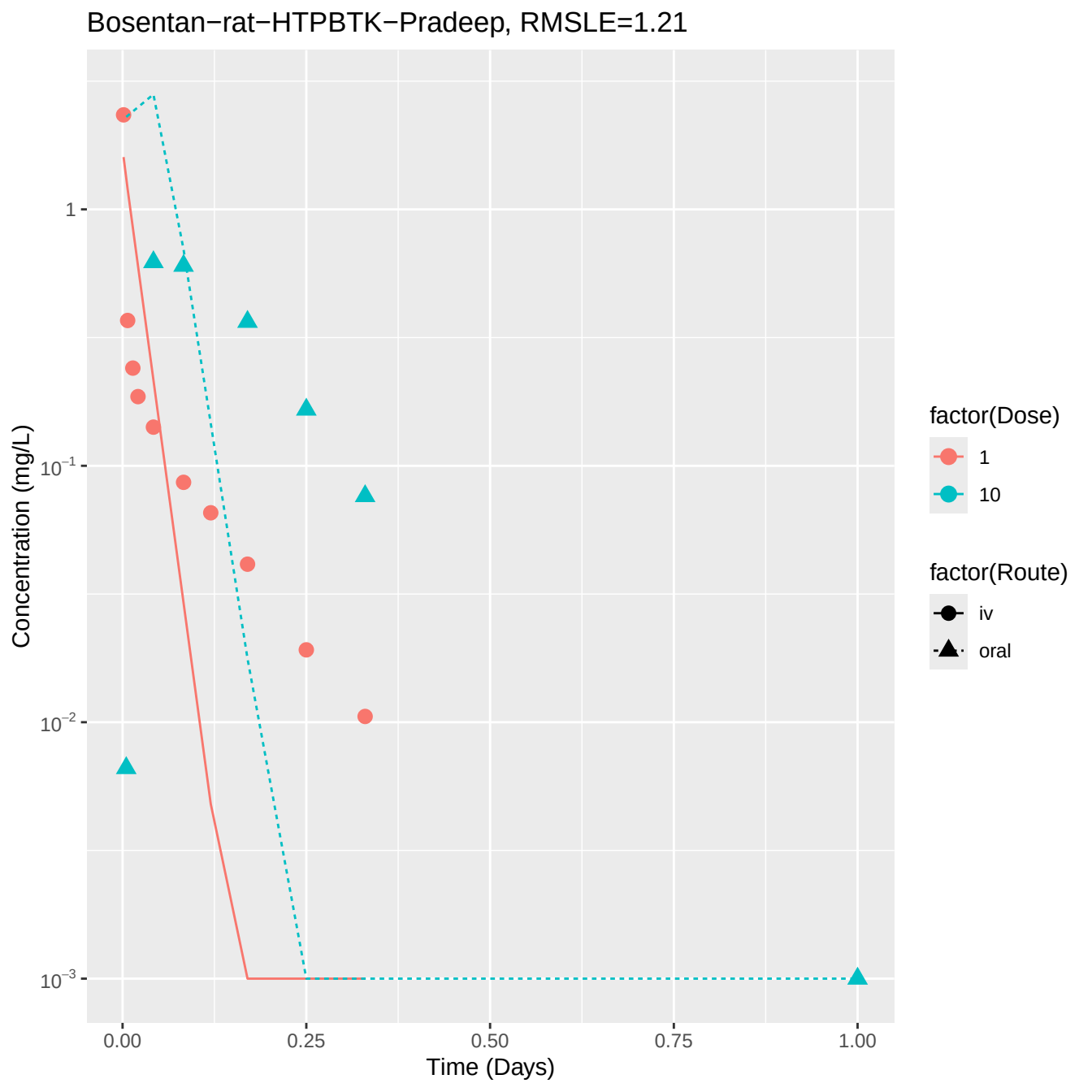

Flufenacet−rat−HTPBTK−Ensemble, RMSLE=1.12
Concentration (mg/L)
10^{-1}
10^{-2}
10^{-3}
10^{-4}
10^{-5}
0
1
2
3
Time (Days)
factor(Route)
iv
oral
factor(Dose)
0.982
0.992
1.01
5.13
5.16
5.3

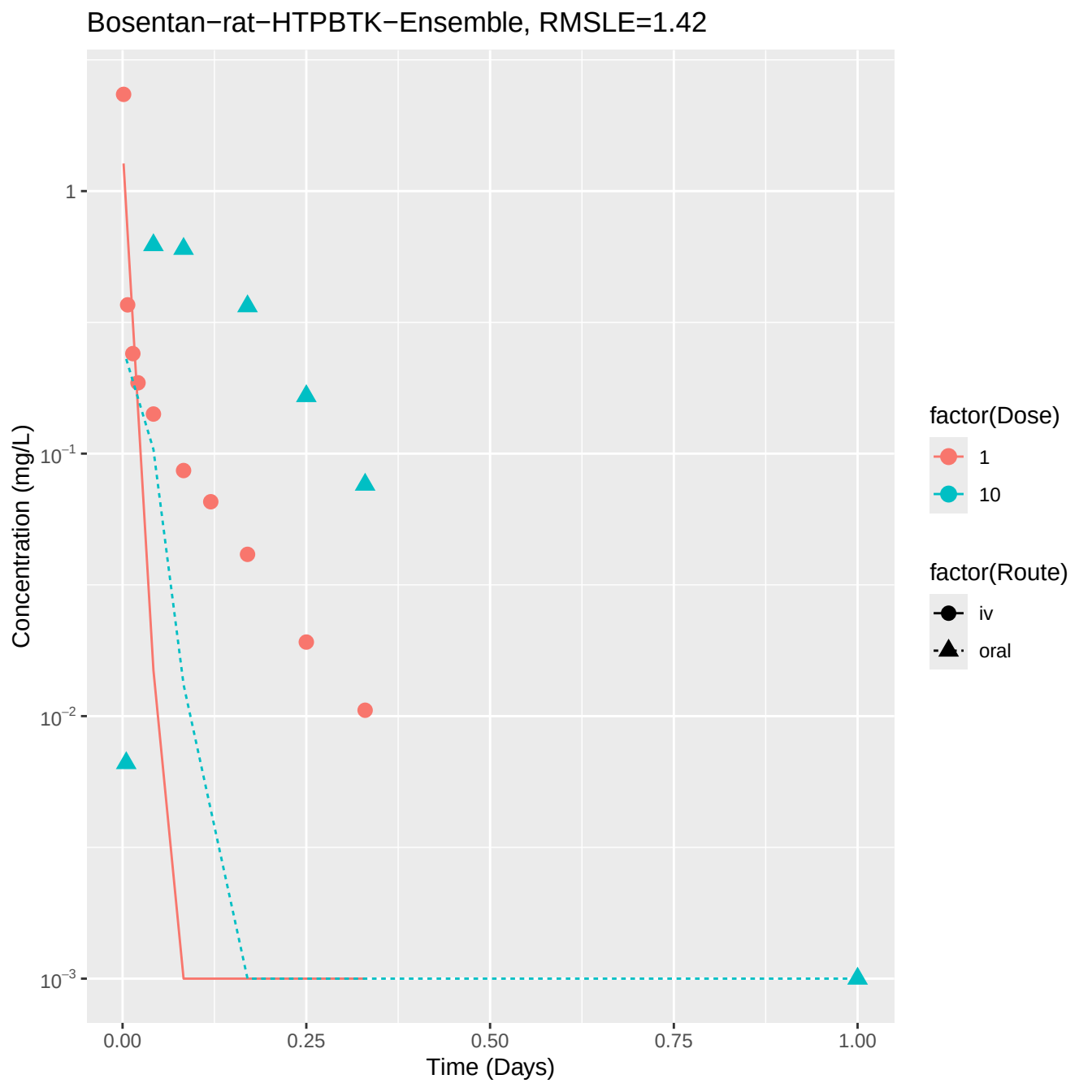

Flufenacet−rat−In Vivo Fits, RMSLE=0.152
Concentration (mg/L)
10^{-1}
10^{-2}
10^{-3}
10^{-4}
10^{-5}
0
1
2
3
Time (Days)
factor(Route)
iv
oral
factor(Dose)
0.982
0.992
1.01
5.13
5.16
5.3

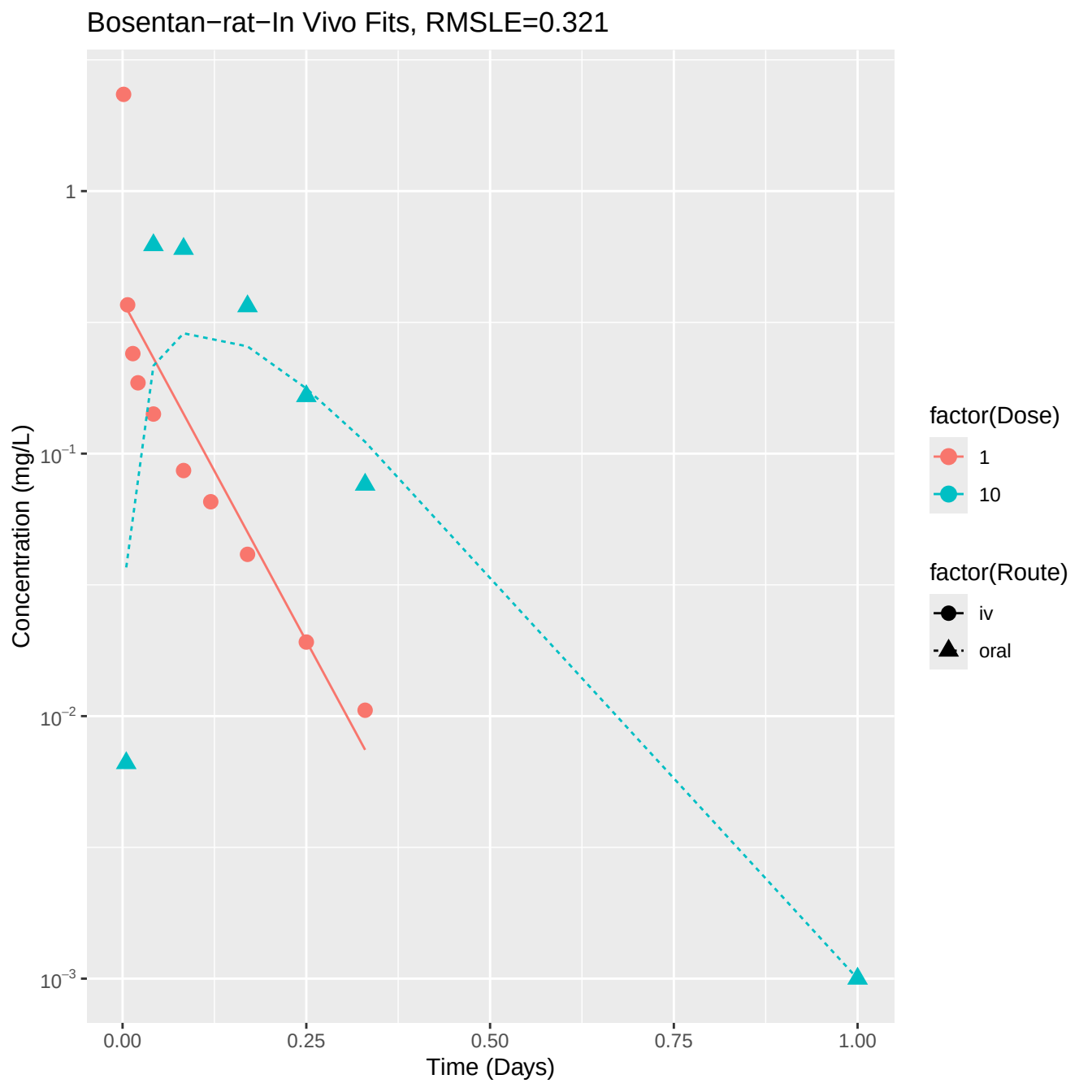

Bosentan−rat−HTPBTK−InVitro, RMSLE=0.995
Concentration (mg/L)
1
10^{-1}
10^{-2}
10^{-3}
0.00
0.25
0.50
0.75
1.00
Time (Days)
factor(Dose)
1
10
factor(Route)
iv
oral

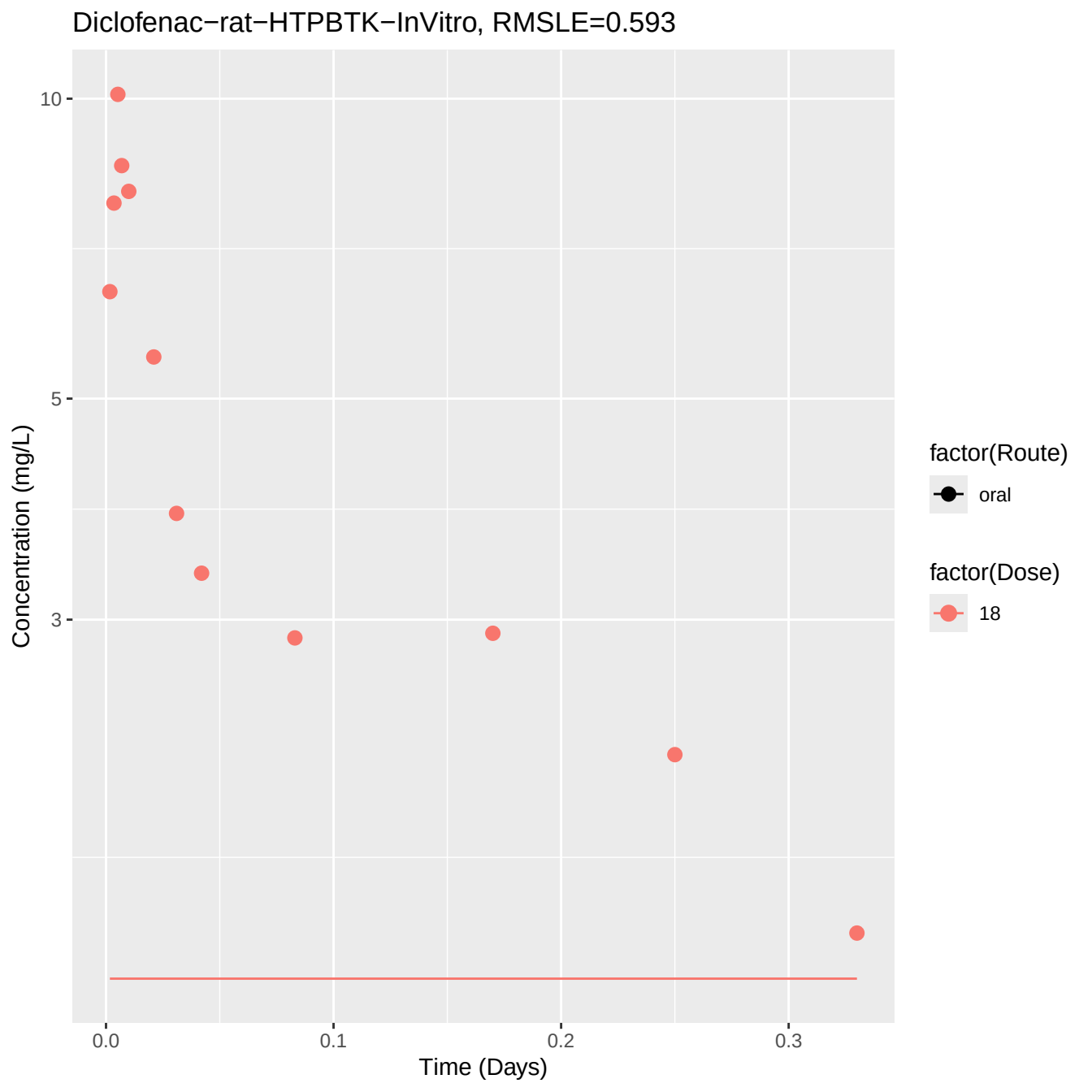

Bosentan−rat−HTPBTK−ADMET, RMSLE=1.45
Concentration (mg/L)
1
10^{-1}
10^{-2}
10^{-3}
0.00
0.25
0.50
0.75
1.00
Time (Days)
factor(Dose)
1
10
factor(Route)
iv
oral

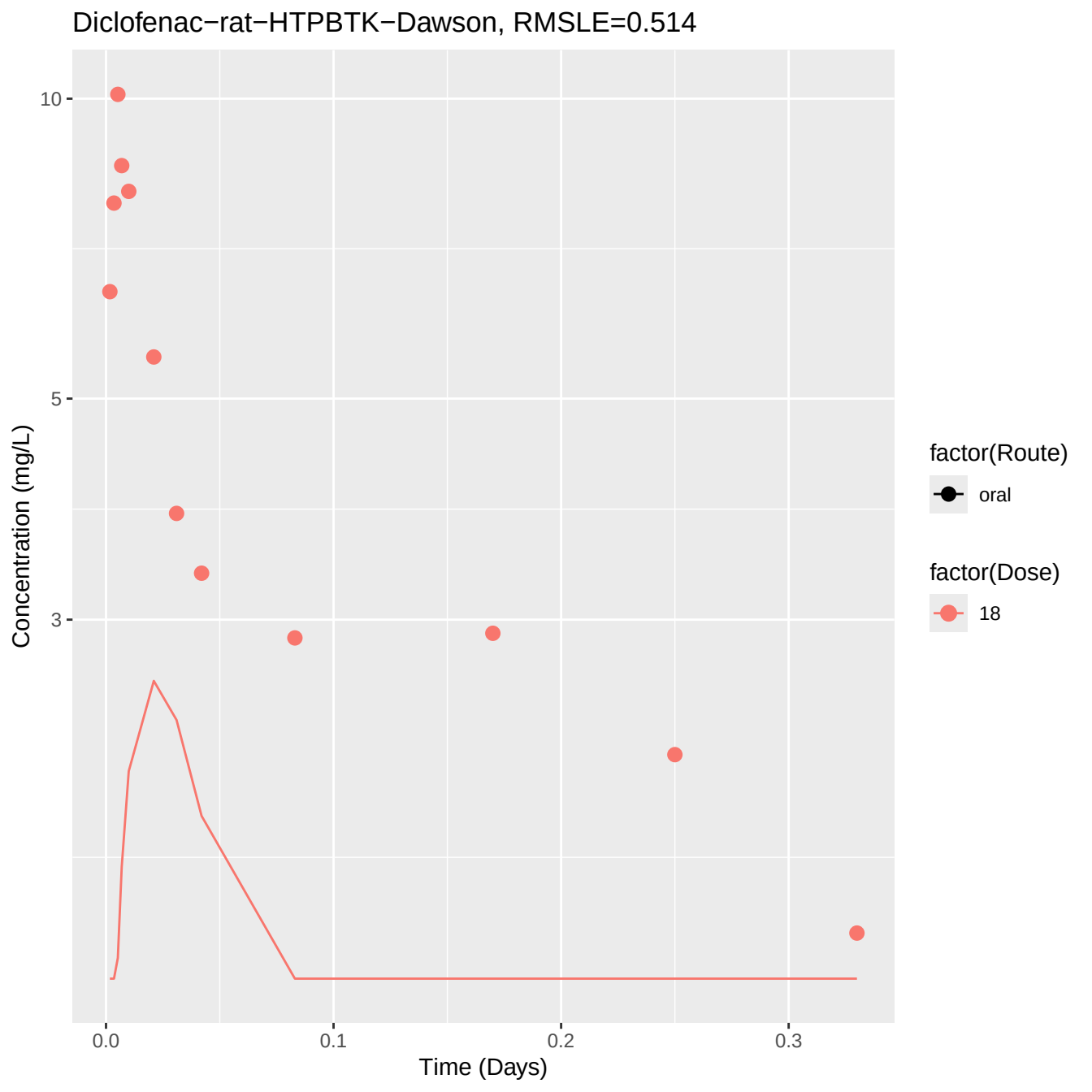

Bosentan−rat−HTPBTK−Dawson, RMSLE=1.37
Concentration (mg/L)
1
10^{-1}
10^{-2}
10^{-3}
0.00
0.25
0.50
0.75
1.00
Time (Days)
factor(Dose)
1
10
factor(Route)
iv
oral

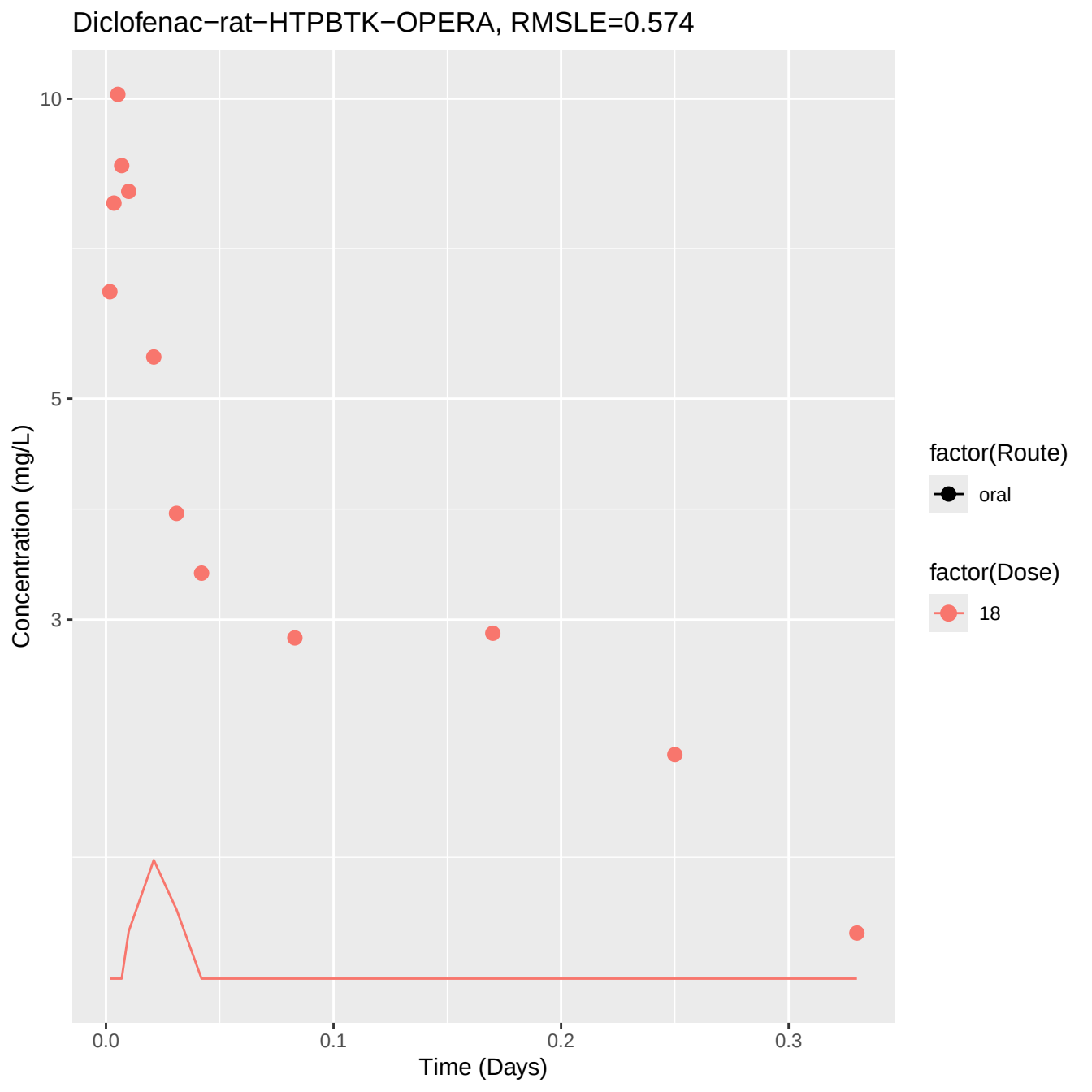

Bosentan−rat−HTPBTK−Pradeep, RMSLE=1.21
Concentration (mg/L)
1
10^{-1}
10^{-2}
10^{-3}
0.00
0.25
0.50
0.75
1.00
Time (Days)
factor(Dose)
1
10
factor(Route)
iv
oral

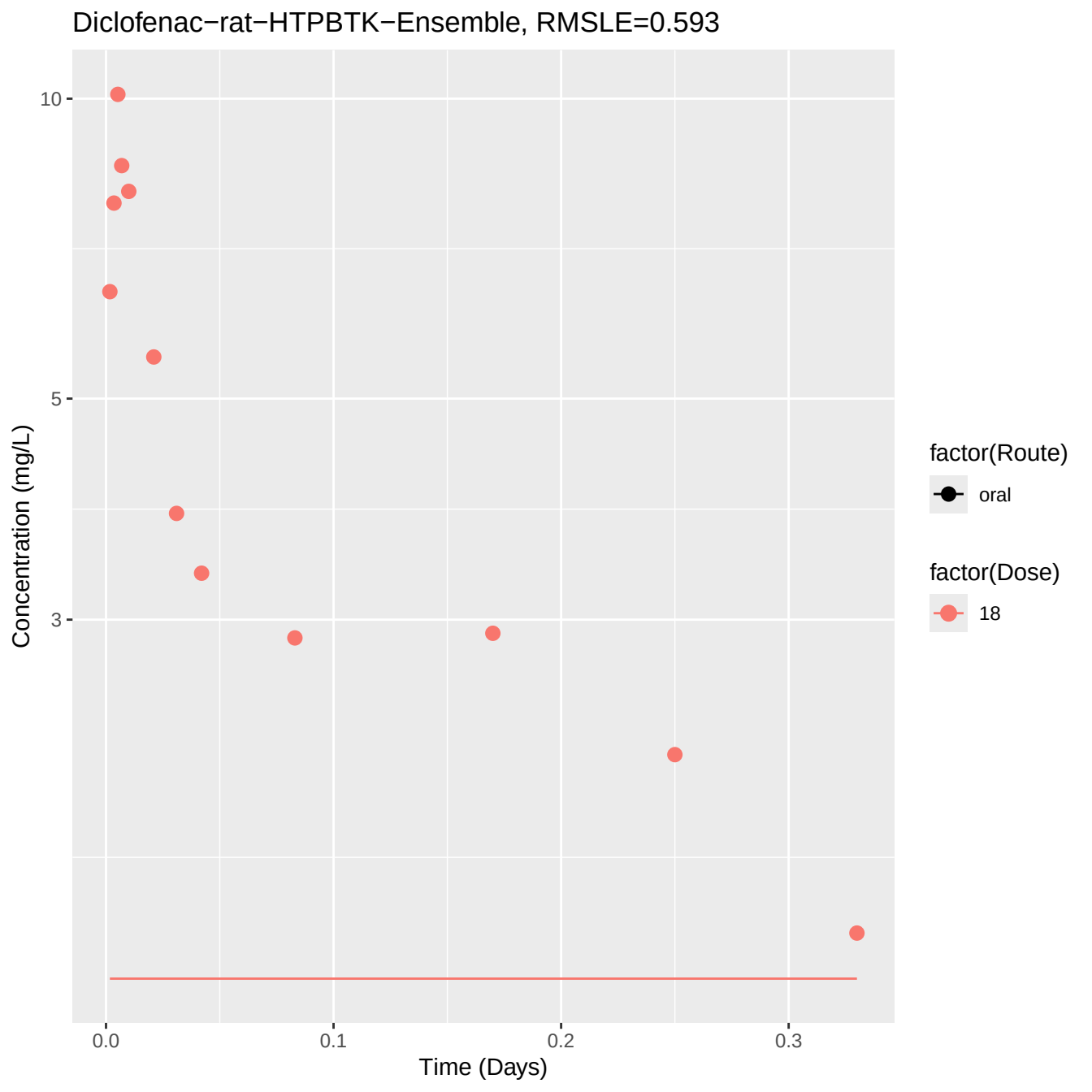

Bosentan−rat−HTPBTK−Ensemble, RMSLE=1.42
Concentration (mg/L)
1
10^{-1}
10^{-2}
10^{-3}
0.00
0.25
0.50
0.75
1.00
Time (Days)
factor(Dose)
1
10
factor(Route)
iv
oral

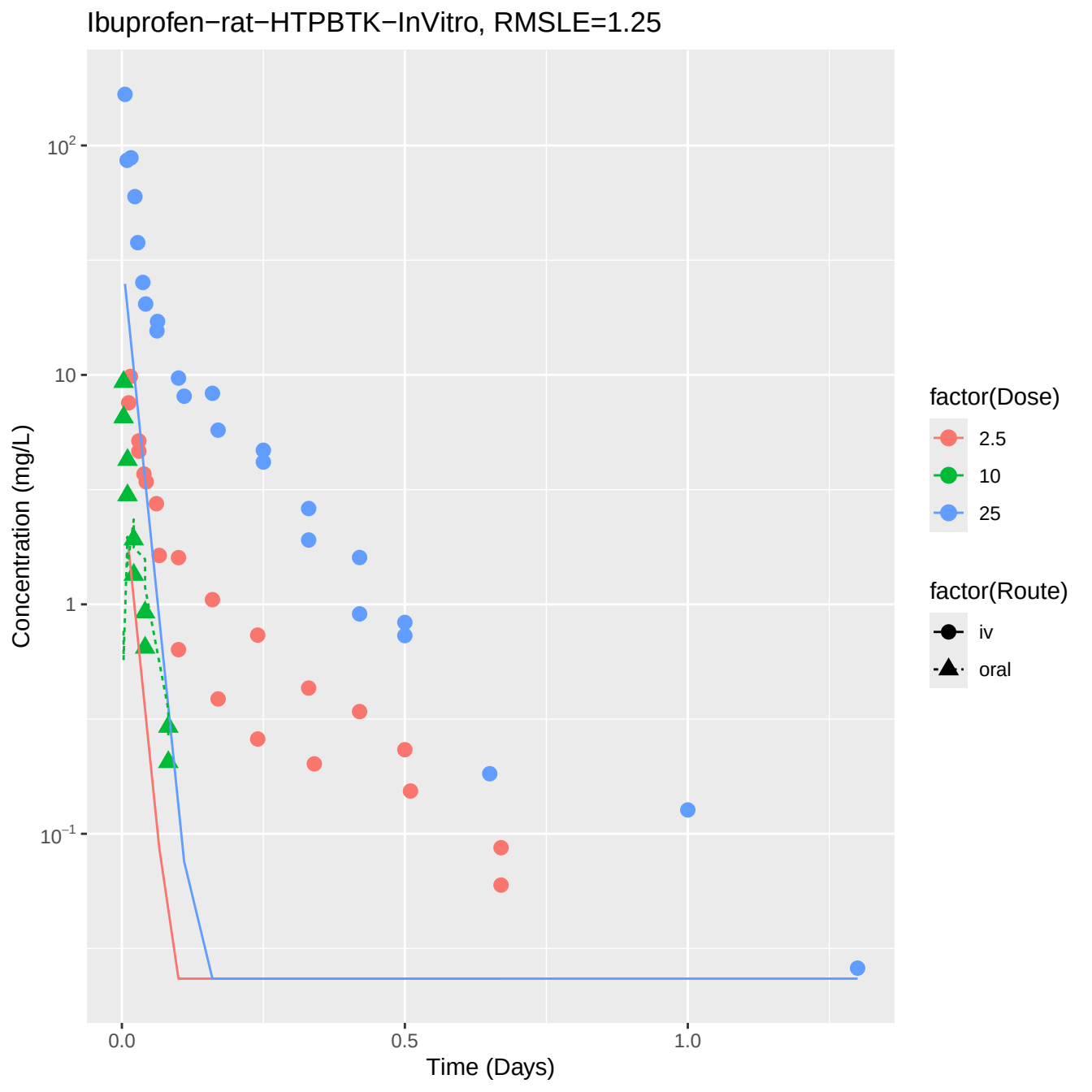

Bosentan−rat−In Vivo Fits, RMSLE=0.321
Concentration (mg/L)
1
10^{-1}
10^{-2}
10^{-3}
0.00
0.25
0.50
0.75
1.00
Time (Days)
factor(Dose)
1
10
factor(Route)
iv
oral

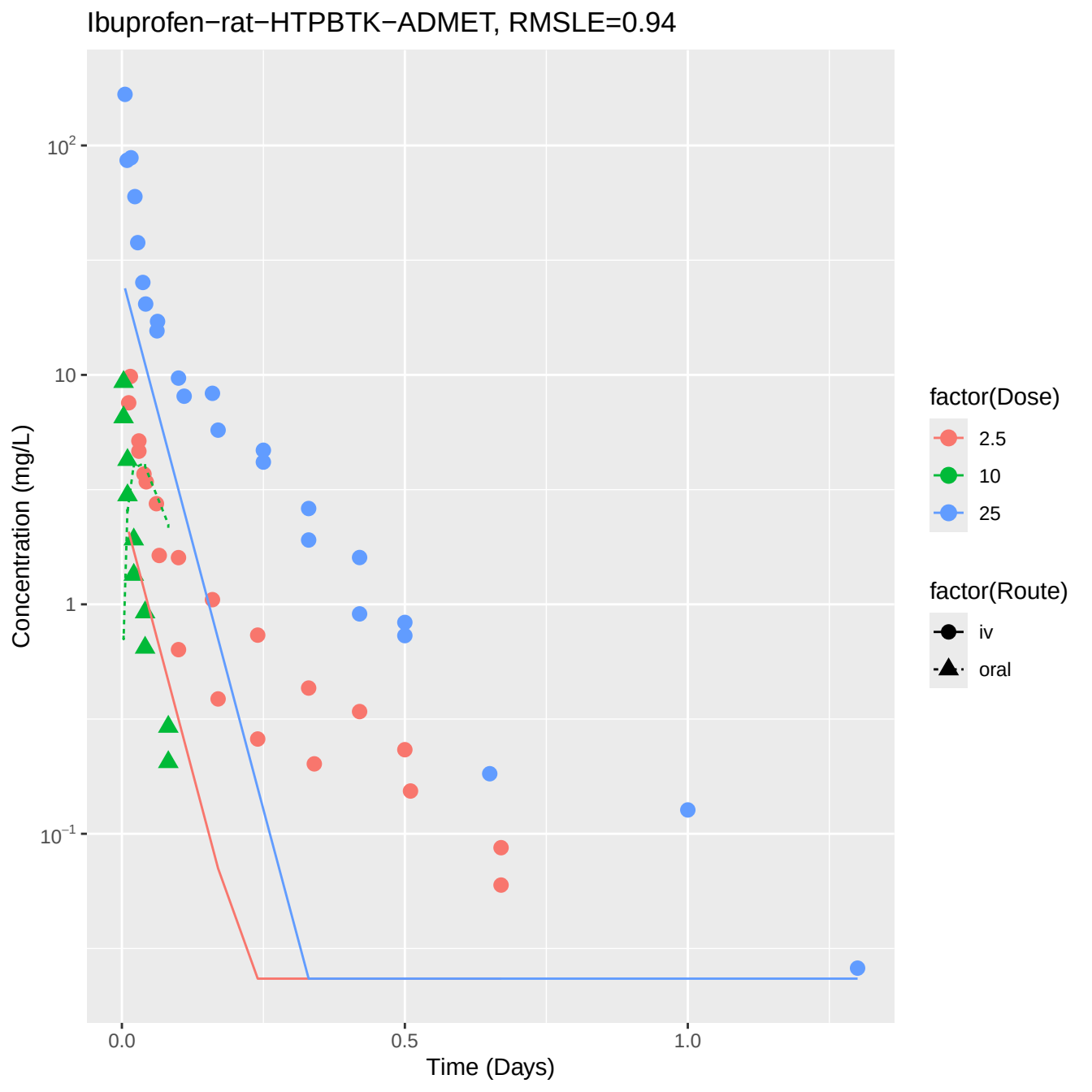

Diclofenac−rat−HTPBTK−InVitro, RMSLE=0.593
Concentration (mg/L)
10
5
3
0.0
0.1
0.2
0.3
Time (Days)
factor(Route)
oral
factor(Dose)
18

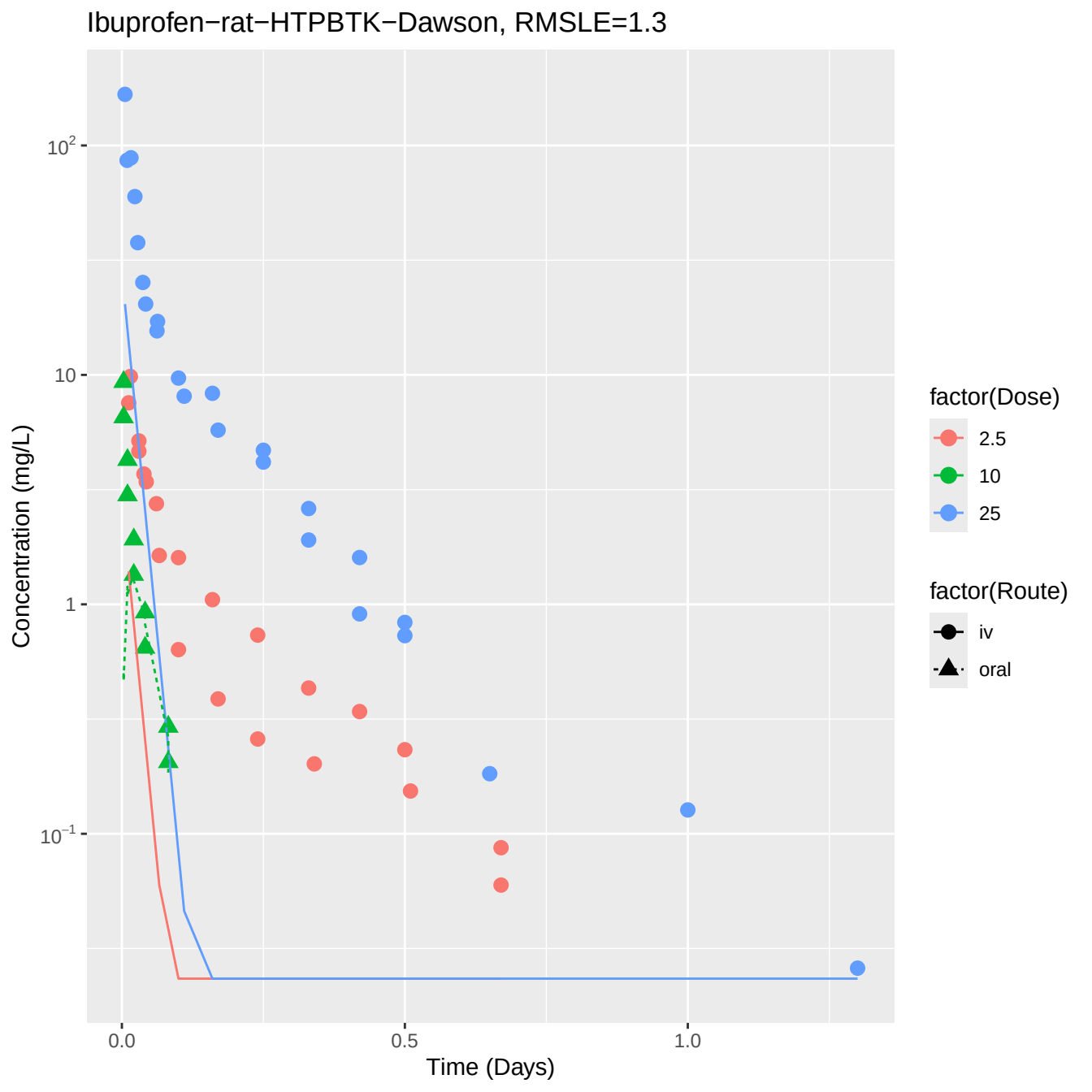

Diclofenac−rat−HTPBTK−Dawson, RMSLE=0.514
10
5
3
Concentration (mg/L)
0.0
0.1
0.2
0.3
Time (Days)
factor(Route)
oral
factor(Dose)
18

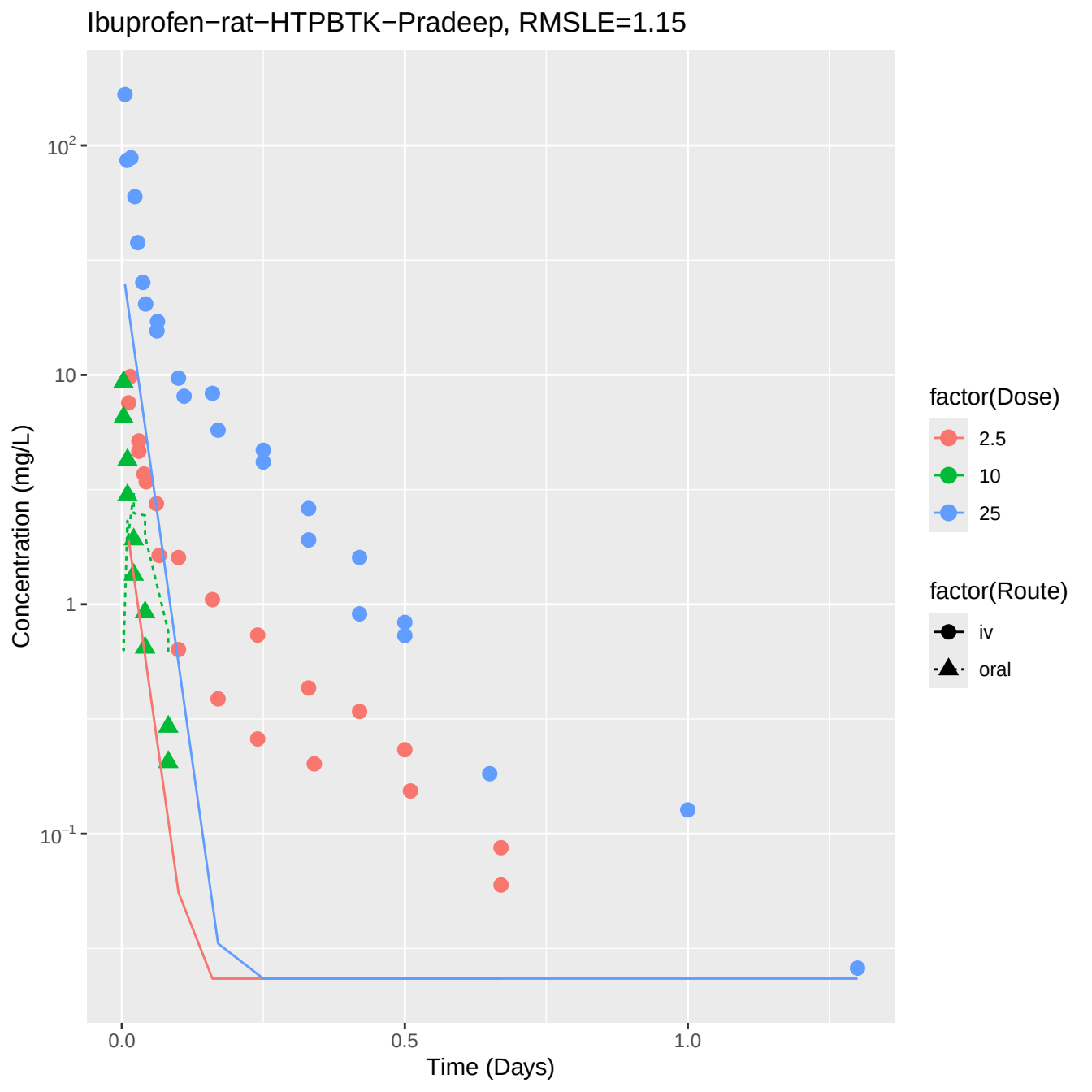

Diclofenac−rat−HTPBTK−OPERA, RMSLE=0.574
Concentration (mg/L)
10
5
3
0.0
0.1
0.2
0.3
Time (Days)
factor(Route)
oral
factor(Dose)
18

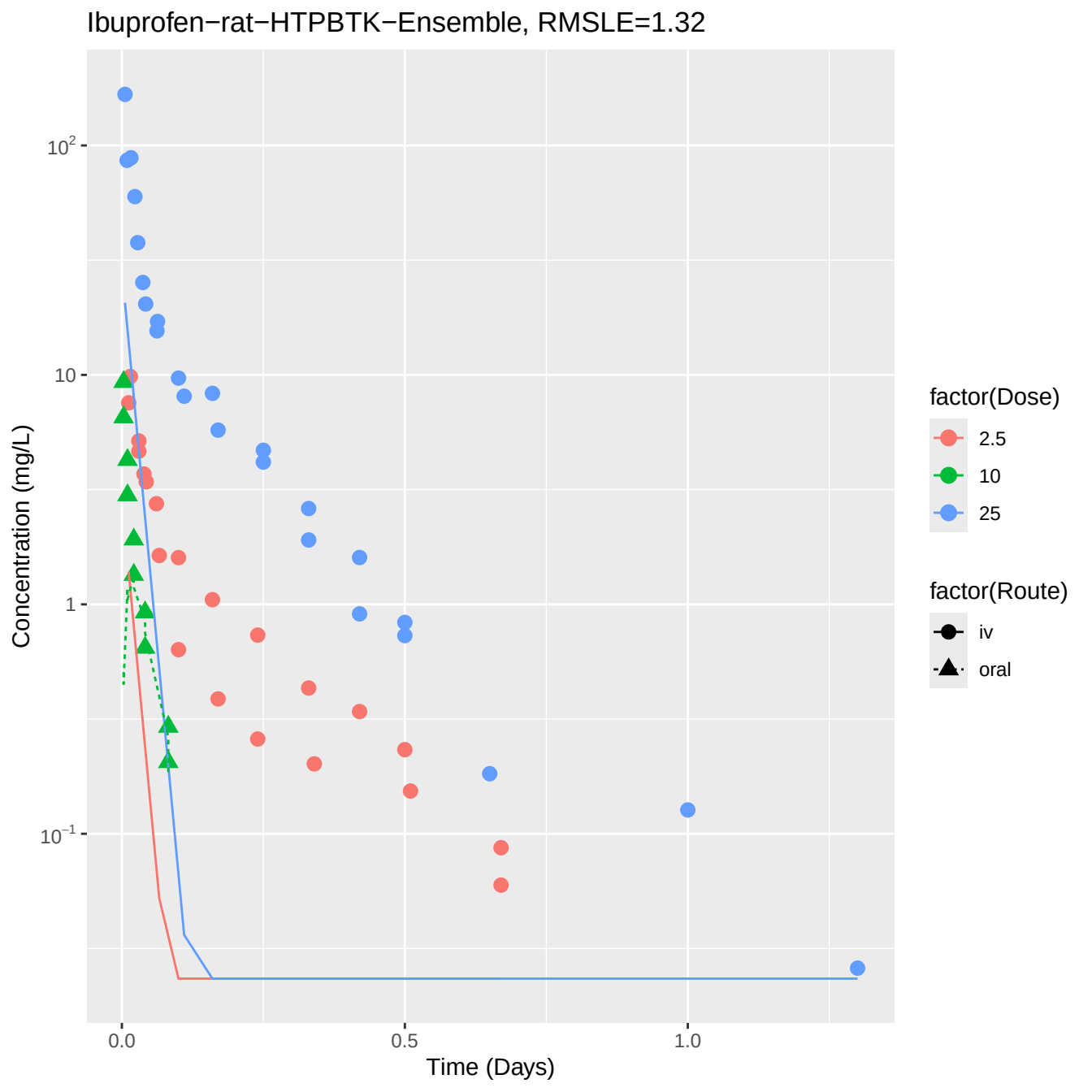

Diclofenac−rat−HTPBTK−Ensemble, RMSLE=0.593
Concentration (mg/L)
10
5
3
0.0
0.1
0.2
0.3
Time (Days)
factor(Route)
oral
factor(Dose)
18

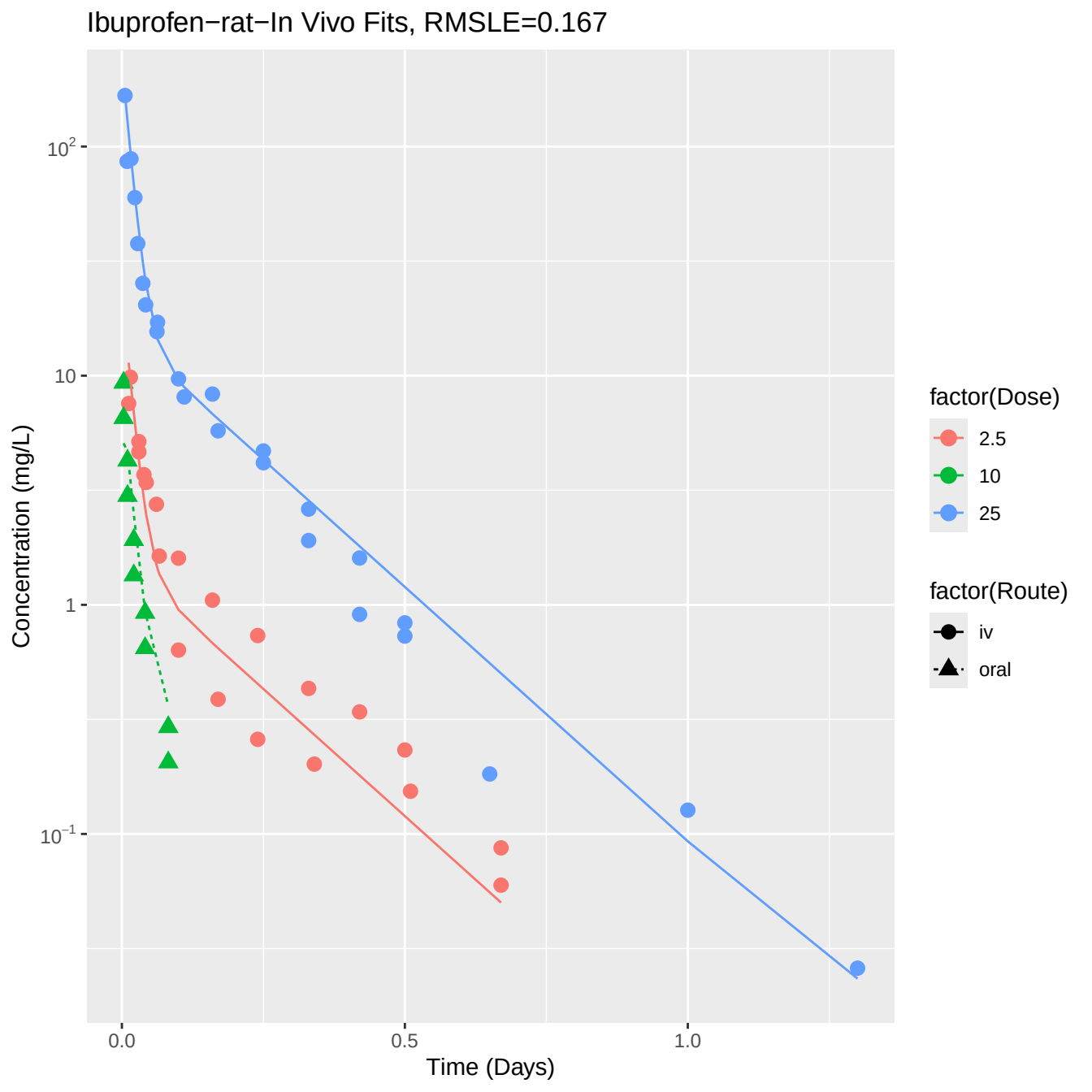

Ibuprofen−rat−HTPBTK−InVitro, RMSLE=1.25
Concentration (mg/L)
10^2
10
1
10^{-1}
0.0
0.5
1.0
Time (Days)
factor(Dose)
2.5
10
25
factor(Route)
iv
oral

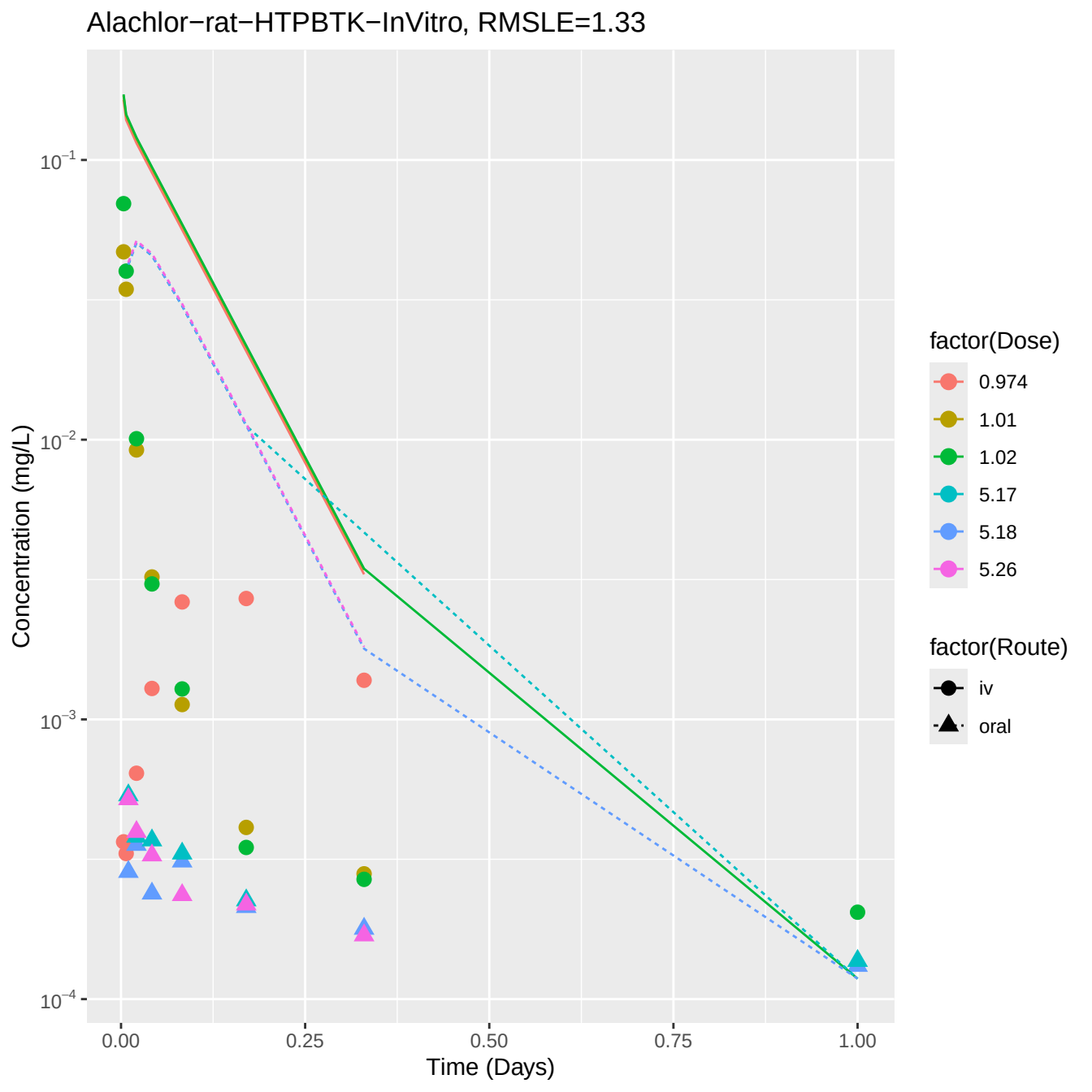

Ibuprofen−rat−HTPBTK−ADMET, RMSLE=0.94
Concentration (mg/L)
10^2
10
1
10^{-1}
0.0
0.5
1.0
Time (Days)
factor(Dose)
2.5
10
25
factor(Route)
iv
oral

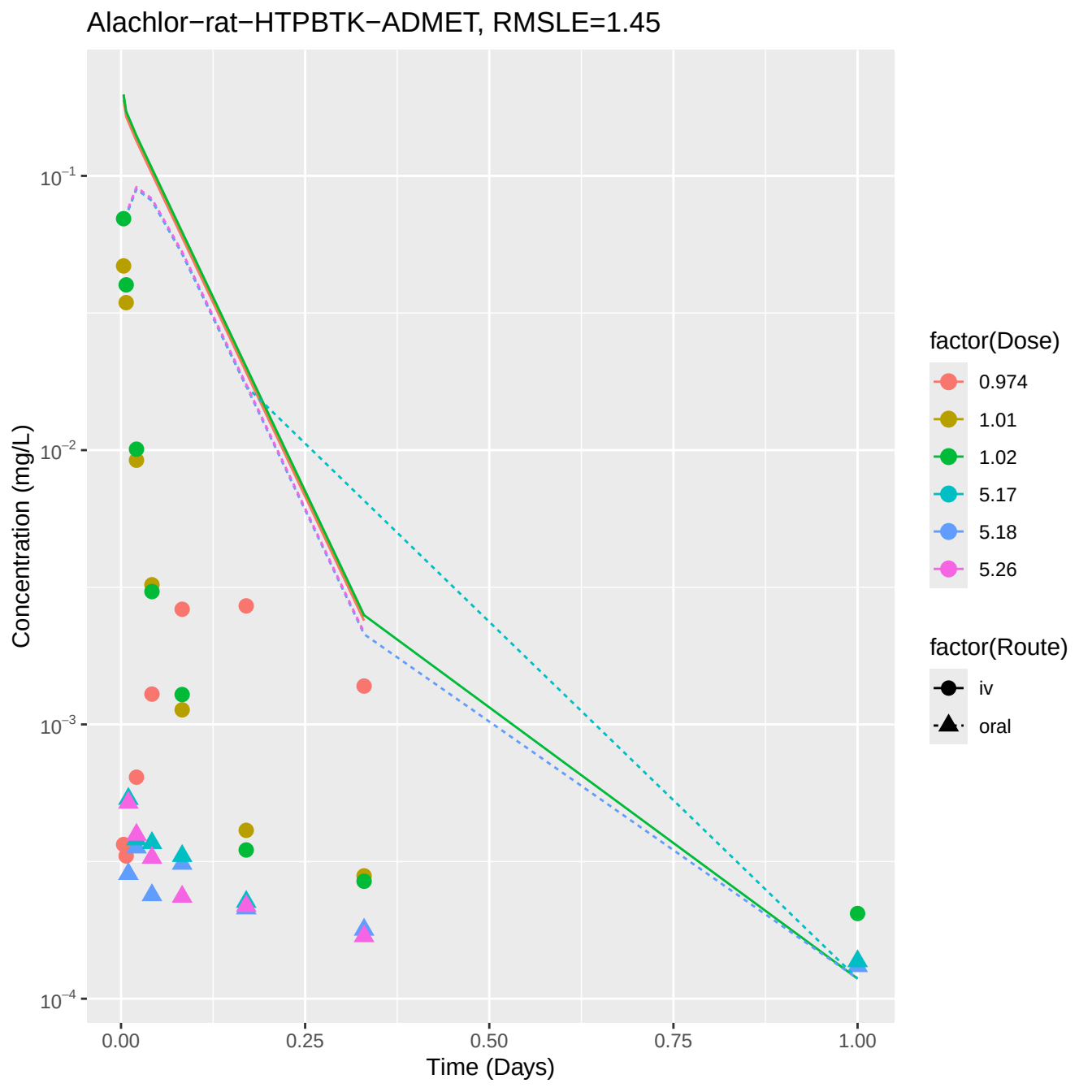

Ibuprofen−rat−HTPBTK−Dawson, RMSLE=1.3
Concentration (mg/L)
10^2
10
1
10^{-1}
0.0
0.5
1.0
Time (Days)
factor(Dose)
2.5
10
25
factor(Route)
iv
oral

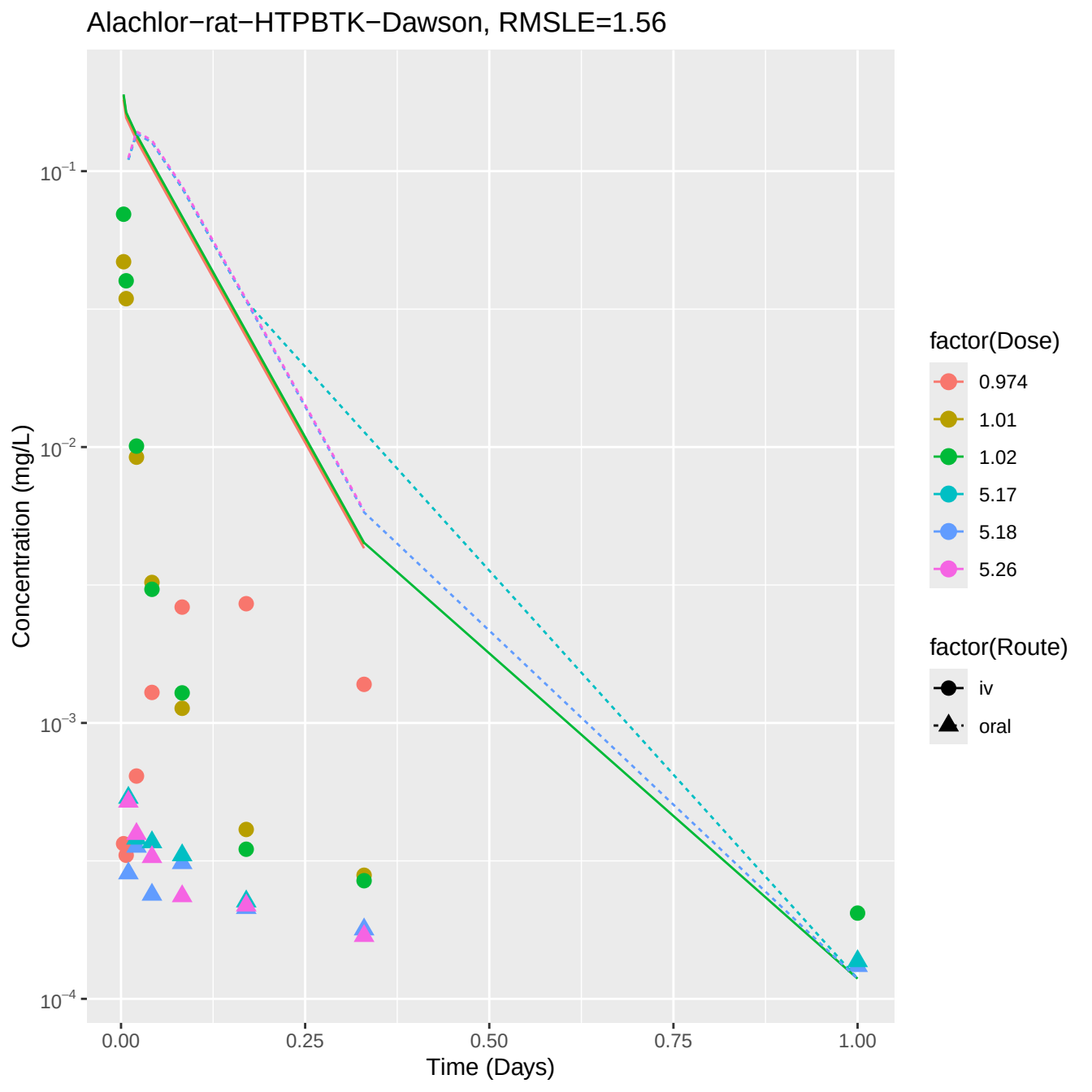

Ibuprofen−rat−HTPBTK−Pradeep, RMSLE=1.15
Concentration (mg/L)
10^2
10
1
10^{-1}
0.0
0.5
1.0
Time (Days)
factor(Dose)
2.5
10
25
factor(Route)
iv
oral

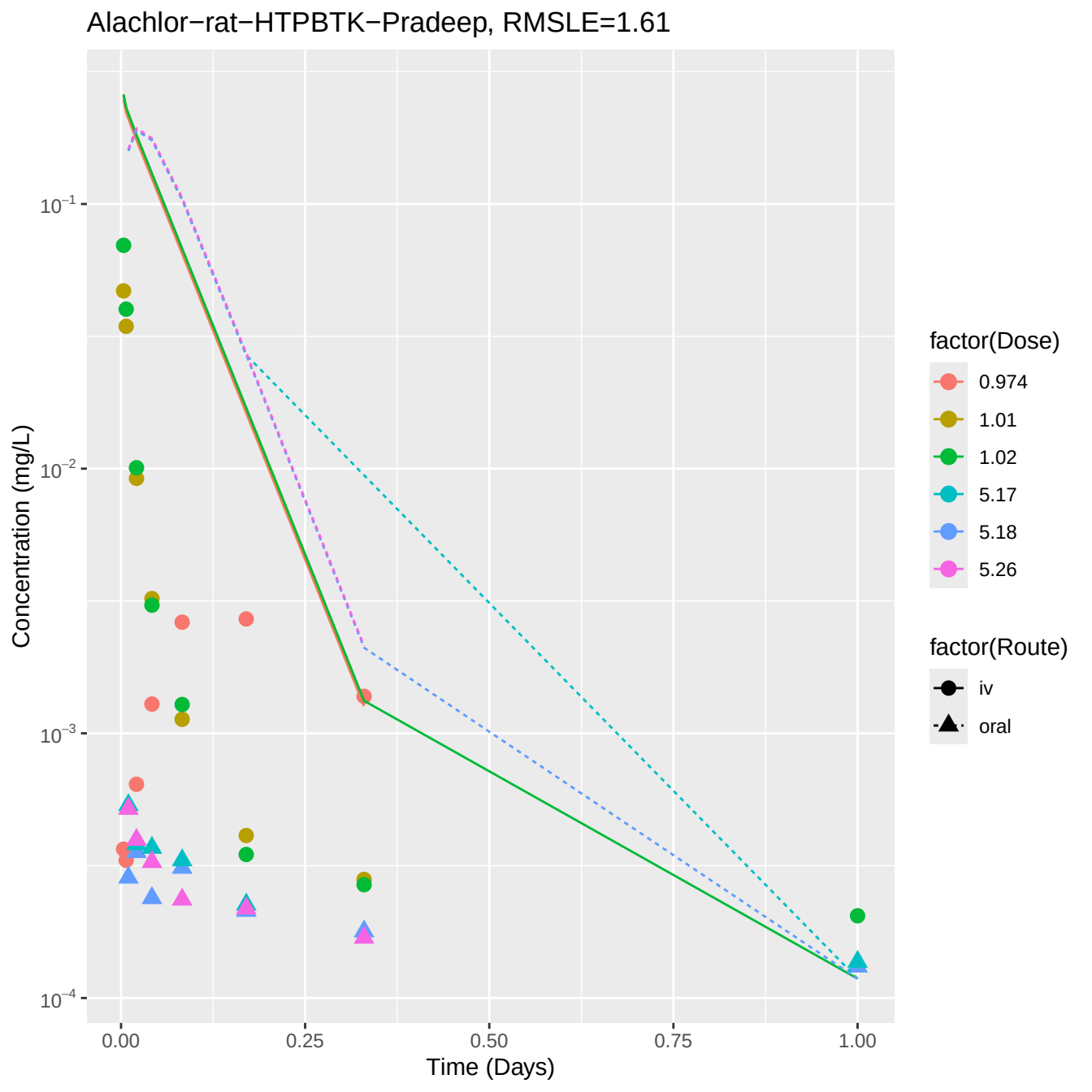

Ibuprofen−rat−HTPBTK−Ensemble, RMSLE=1.32
Concentration (mg/L)
10^2
10
1
10^{-1}
0.0
0.5
1.0
Time (Days)
factor(Dose)
2.5
10
25
factor(Route)
iv
oral

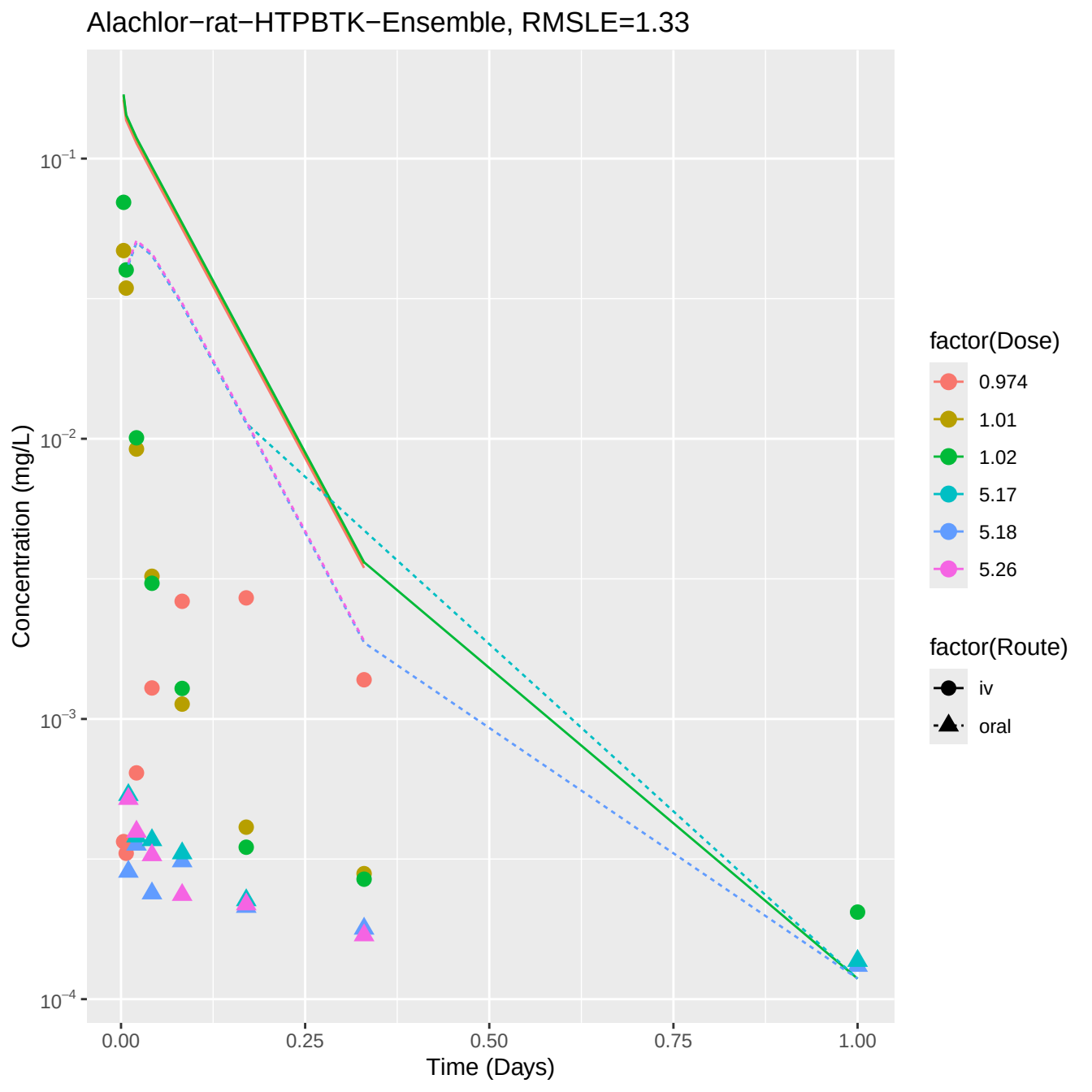

Ibuprofen−rat−In Vivo Fits, RMSLE=0.167
Concentration (mg/L)
10^2
10
1
10^{-1}
0.0
0.5
1.0
Time (Days)
factor(Dose)
2.5
10
25
factor(Route)
iv
oral

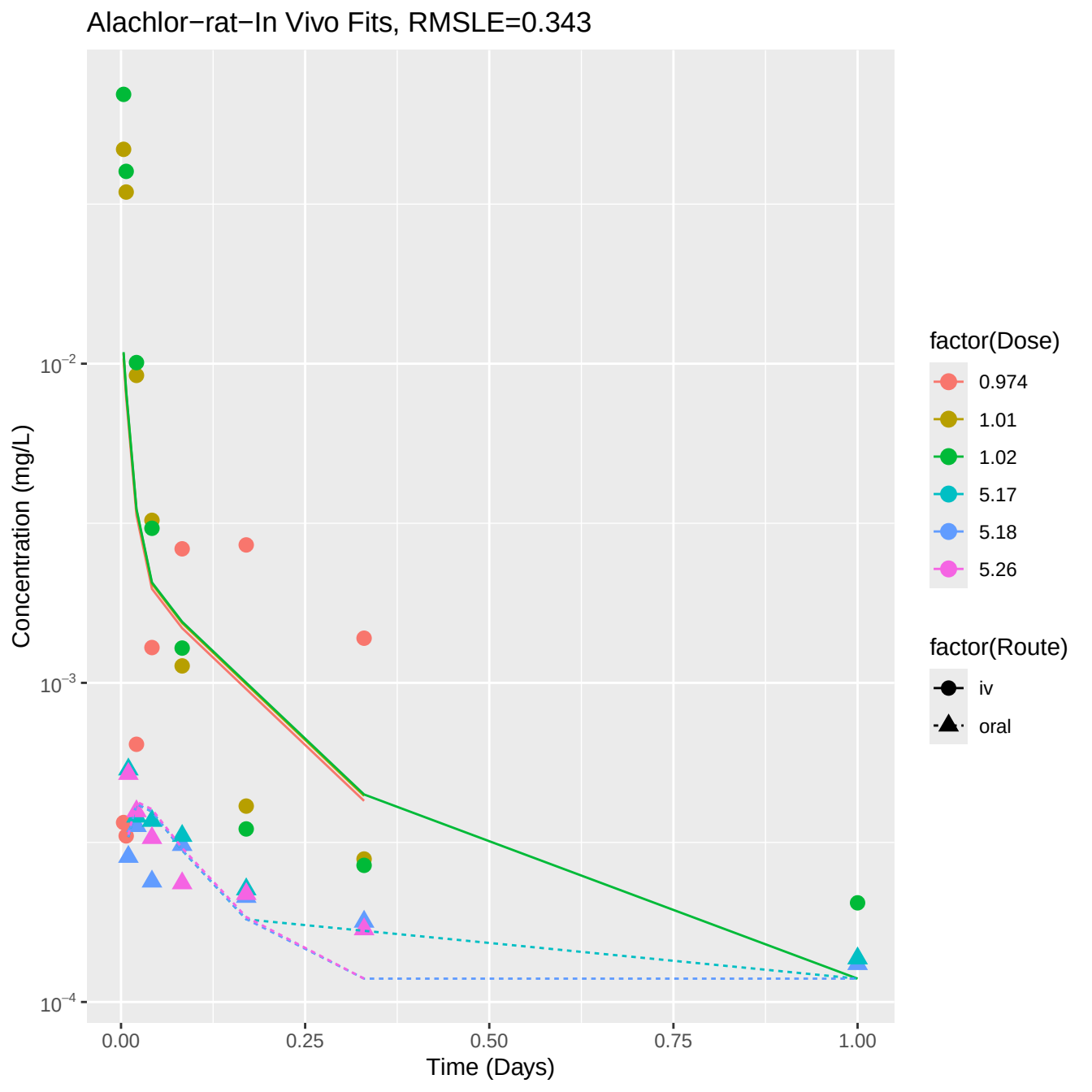

Alachlor−rat−HTPBTK−InVitro, RMSLE=1.33
Concentration (mg/L)
10^{-1}
10^{-2}
10^{-3}
10^{-4}
0.00
0.25
0.50
0.75
1.00
Time (Days)
factor(Dose)
0.974
1.01
1.02
5.17
5.18
5.26
factor(Route)
iv
oral

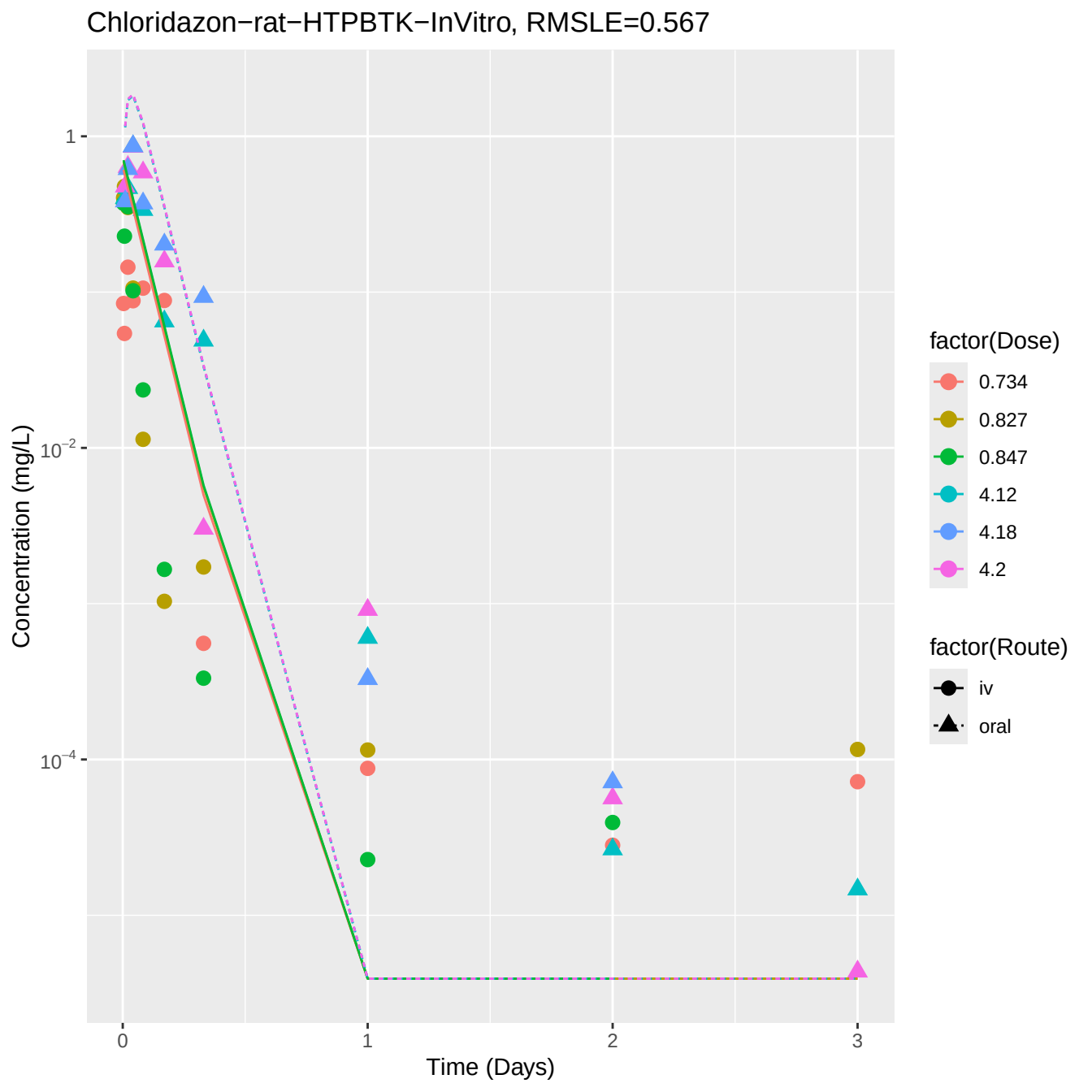

Alachlor−rat−HTPBTK−ADMET, RMSLE=1.45
Concentration (mg/L)
10^{-1}
10^{-2}
10^{-3}
10^{-4}
0.00
0.25
0.50
0.75
1.00
Time (Days)
factor(Dose)
0.974
1.01
1.02
5.17
5.18
5.26
factor(Route)
iv
oral

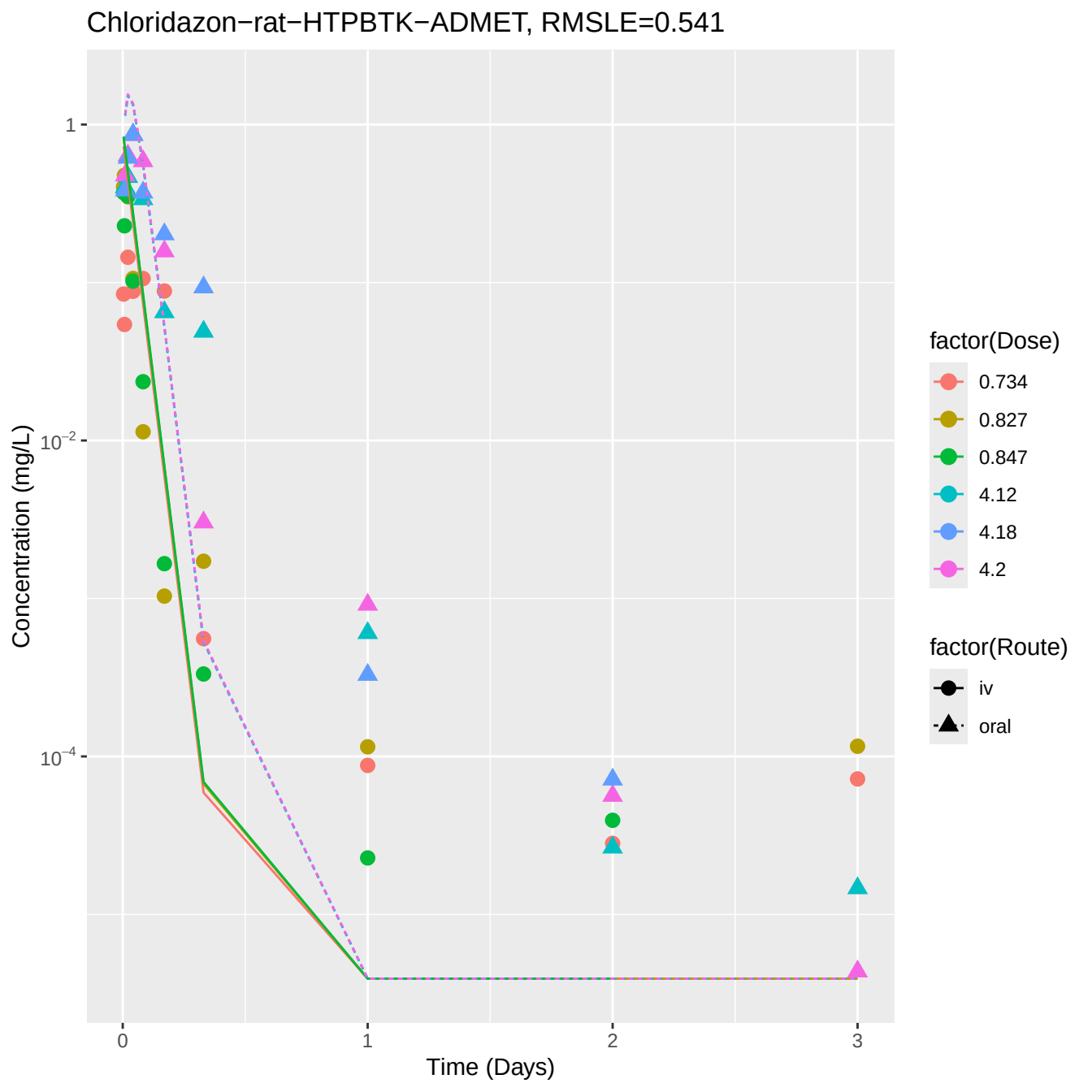

Alachlor−rat−HTPBTK−Dawson, RMSLE=1.56
Concentration (mg/L)
10^{-1}
10^{-2}
10^{-3}
10^{-4}
0.00
0.25
0.50
0.75
1.00
Time (Days)
factor(Dose)
0.974
1.01
1.02
5.17
5.18
5.26
factor(Route)
iv
oral

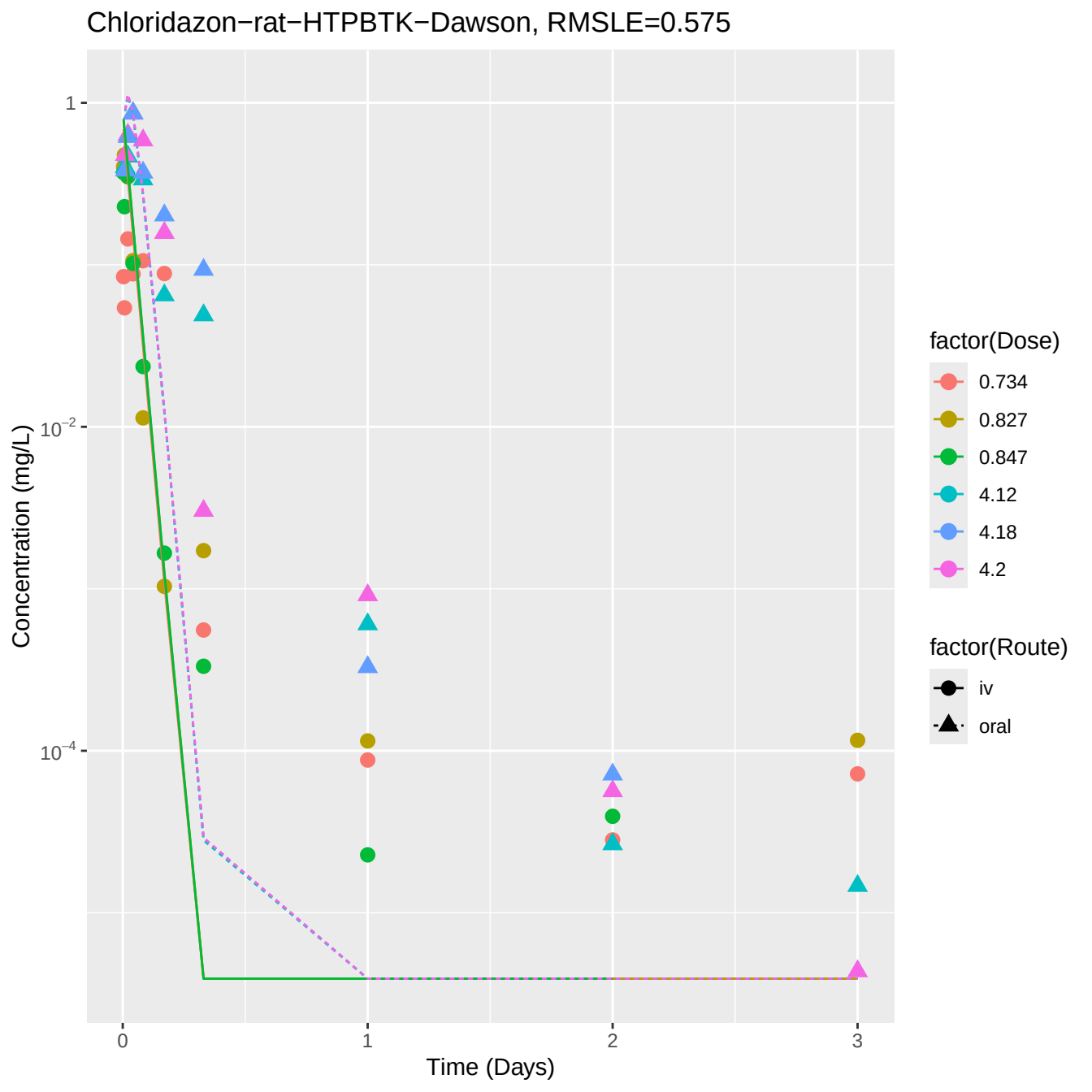

Alachlor−rat−HTPBTK−Pradeep, RMSLE=1.61
Concentration (mg/L)
10^{-1}
10^{-2}
10^{-3}
10^{-4}
0.00
0.25
0.50
0.75
1.00
Time (Days)
factor(Dose)
0.974
1.01
1.02
5.17
5.18
5.26
factor(Route)
iv
oral

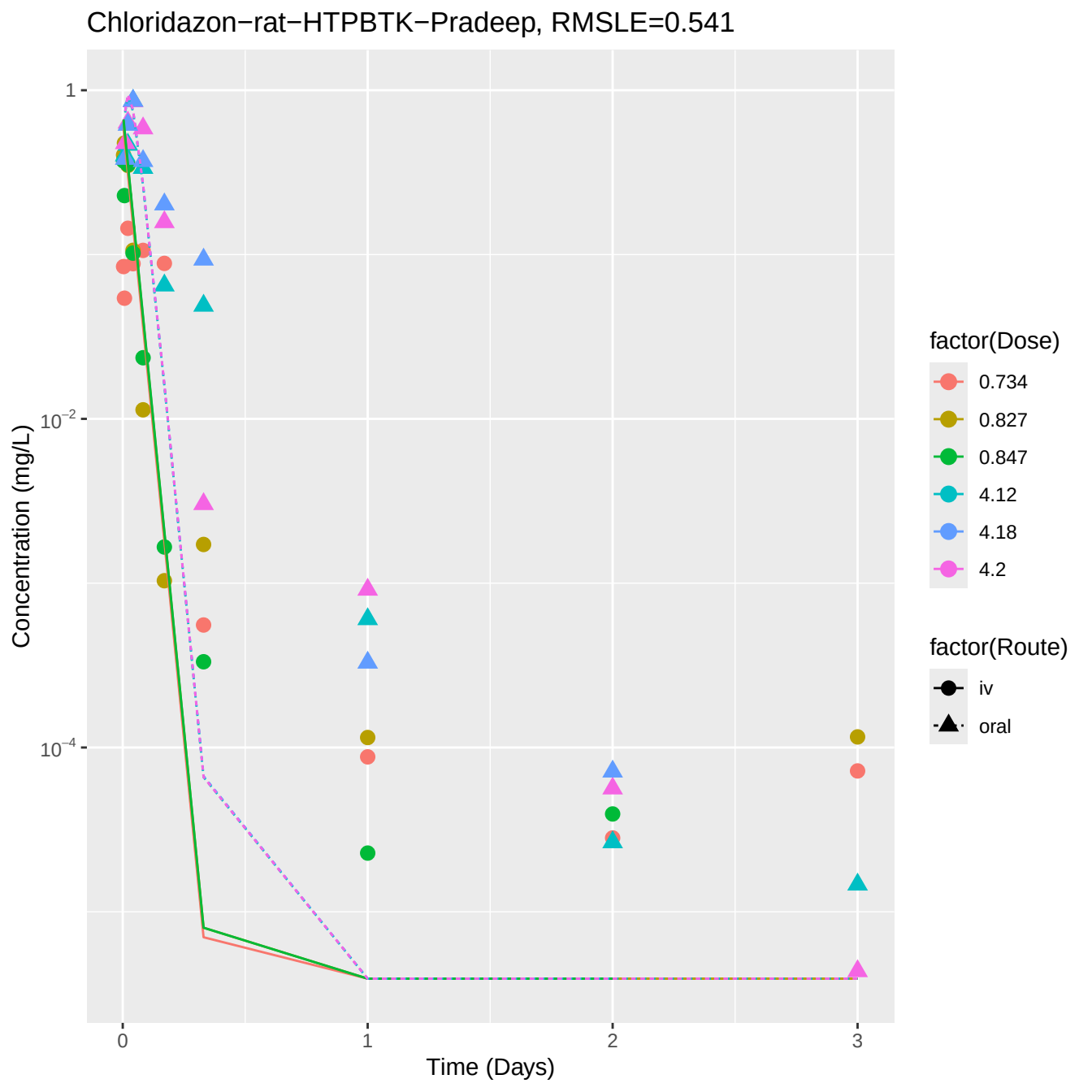

Alachlor−rat−HTPBTK−Ensemble, RMSLE=1.33
Concentration (mg/L)
10^{-1}
10^{-2}
10^{-3}
10^{-4}
0.00
0.25
0.50
0.75
1.00
Time (Days)
factor(Dose)
0.974
1.01
1.02
5.17
5.18
5.26
factor(Route)
iv
oral

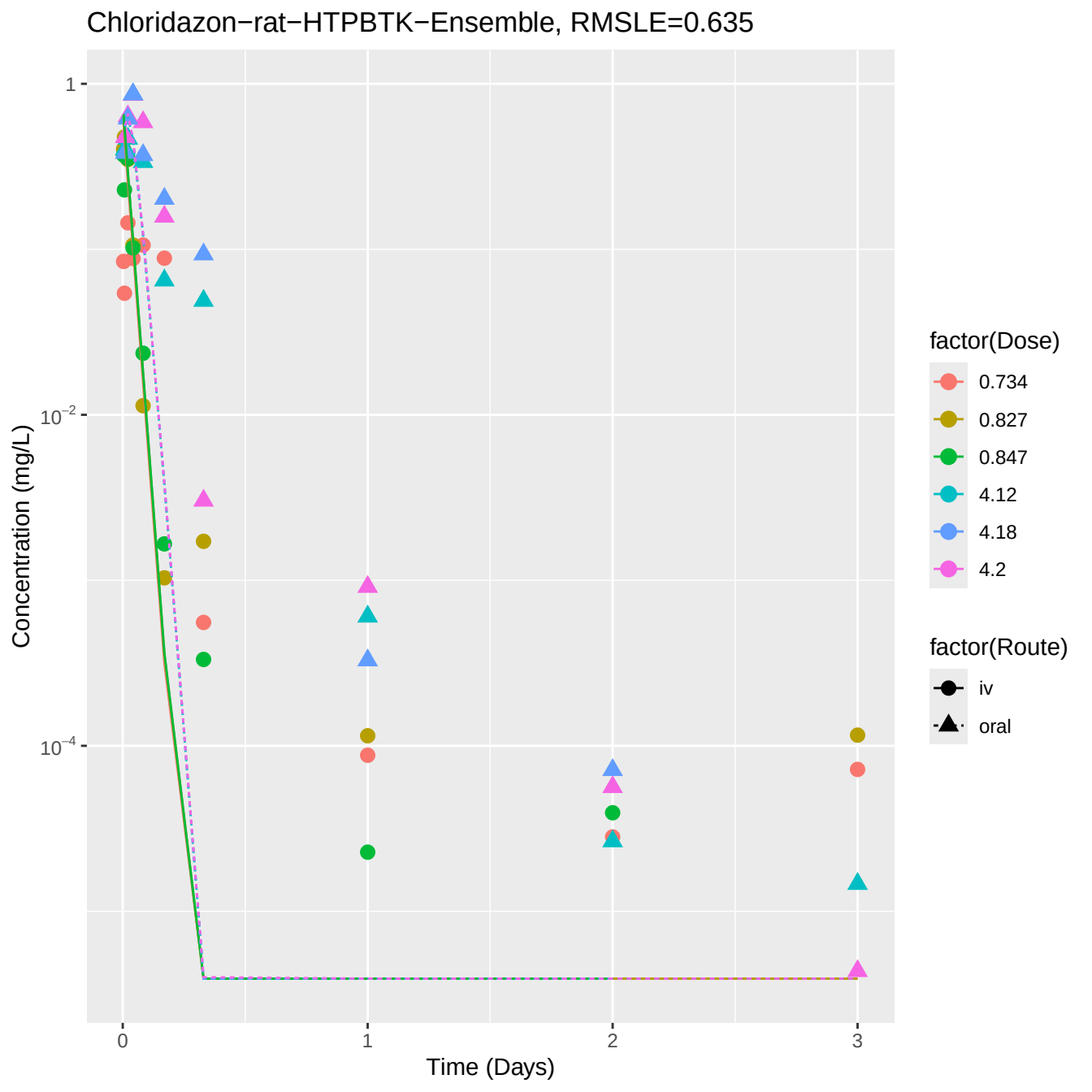

Alachlor−rat−In Vivo Fits, RMSLE=0.343
Concentration (mg/L)
10^{-2}
10^{-3}
10^{-4}
0.00
0.25
0.50
0.75
1.00
Time (Days)
factor(Dose)
0.974
1.01
1.02
5.17
5.18
5.26
factor(Route)
iv
oral

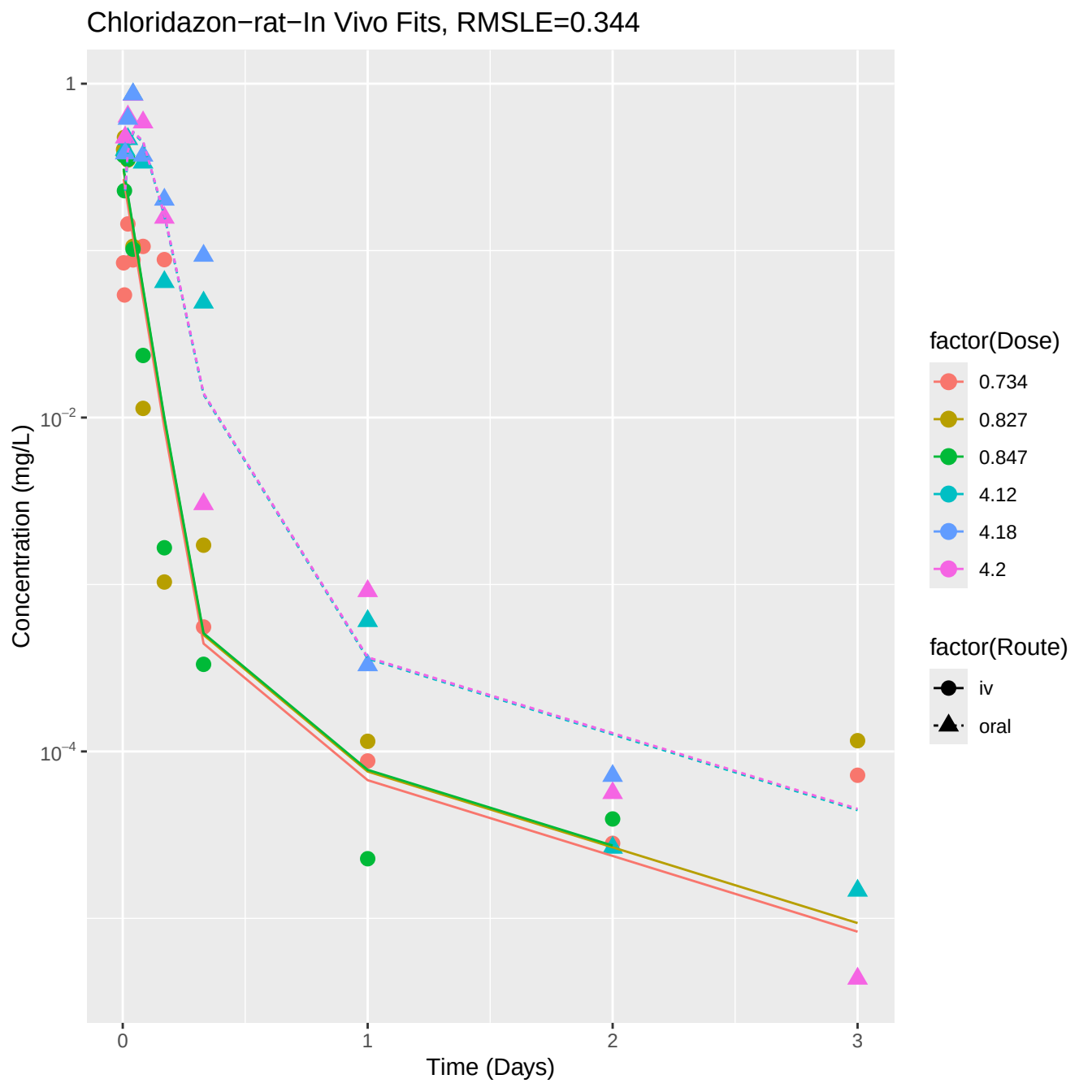

Chloridazon−rat−HTPBTK−InVitro, RMSLE=0.567
Concentration (mg/L)
1
10^{-2}
10^{-4}
0
1
2
3
Time (Days)
factor(Dose)
0.734
0.827
0.847
4.12
4.18
4.2
factor(Route)
iv
oral

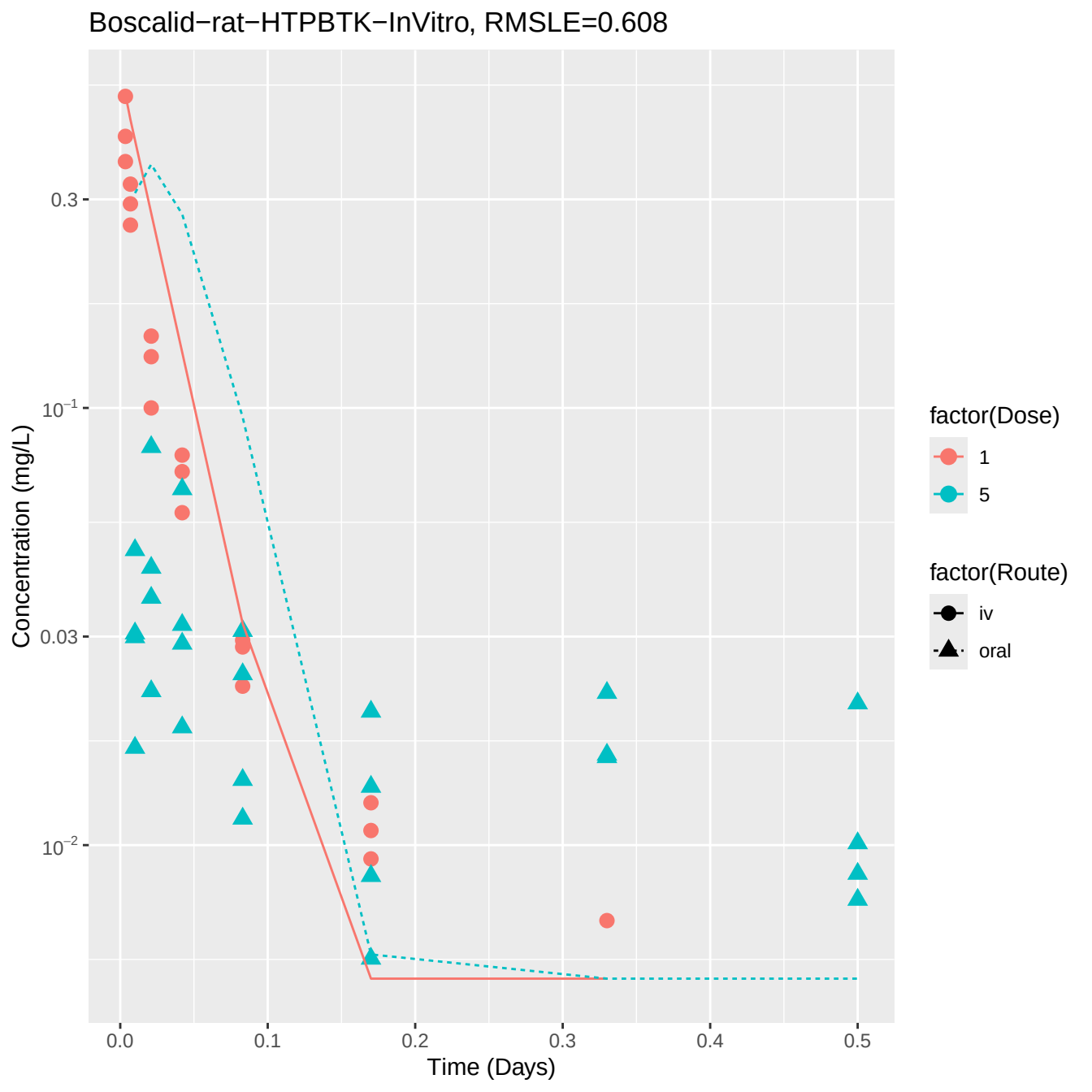

Chloridazon−rat−HTPBTK−ADMET, RMSLE=0.541
Concentration (mg/L)
1
10^{-2}
10^{-4}
0
1
2
3
Time (Days)
factor(Dose)
0.734
0.827
0.847
4.12
4.18
4.2
factor(Route)
iv
oral

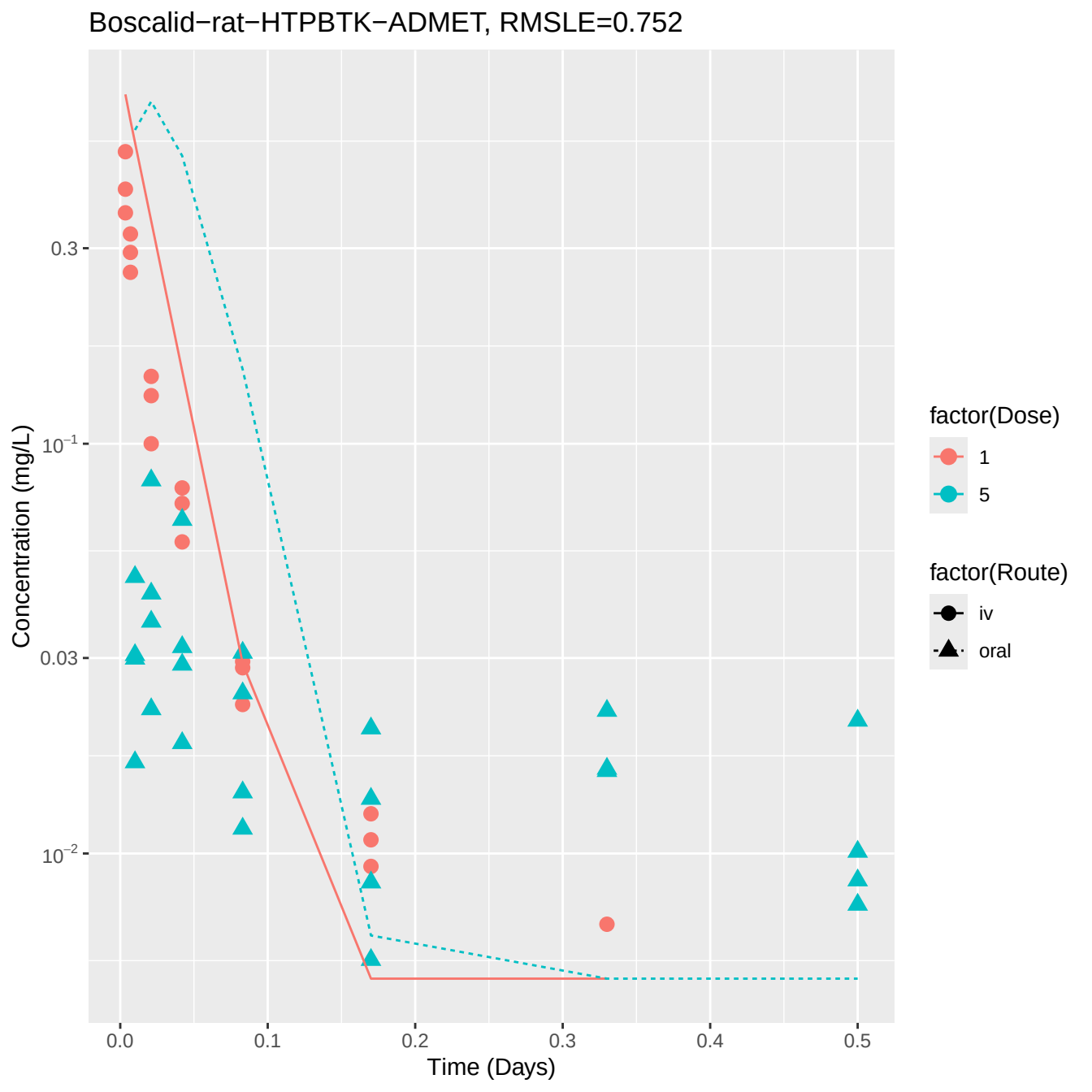

Chloridazon−rat−HTPBTK−Dawson, RMSLE=0.575
Concentration (mg/L)
1
10^{-2}
10^{-4}
0
1
2
3
Time (Days)
factor(Dose)
0.734
0.827
0.847
4.12
4.18
4.2
factor(Route)
iv
oral

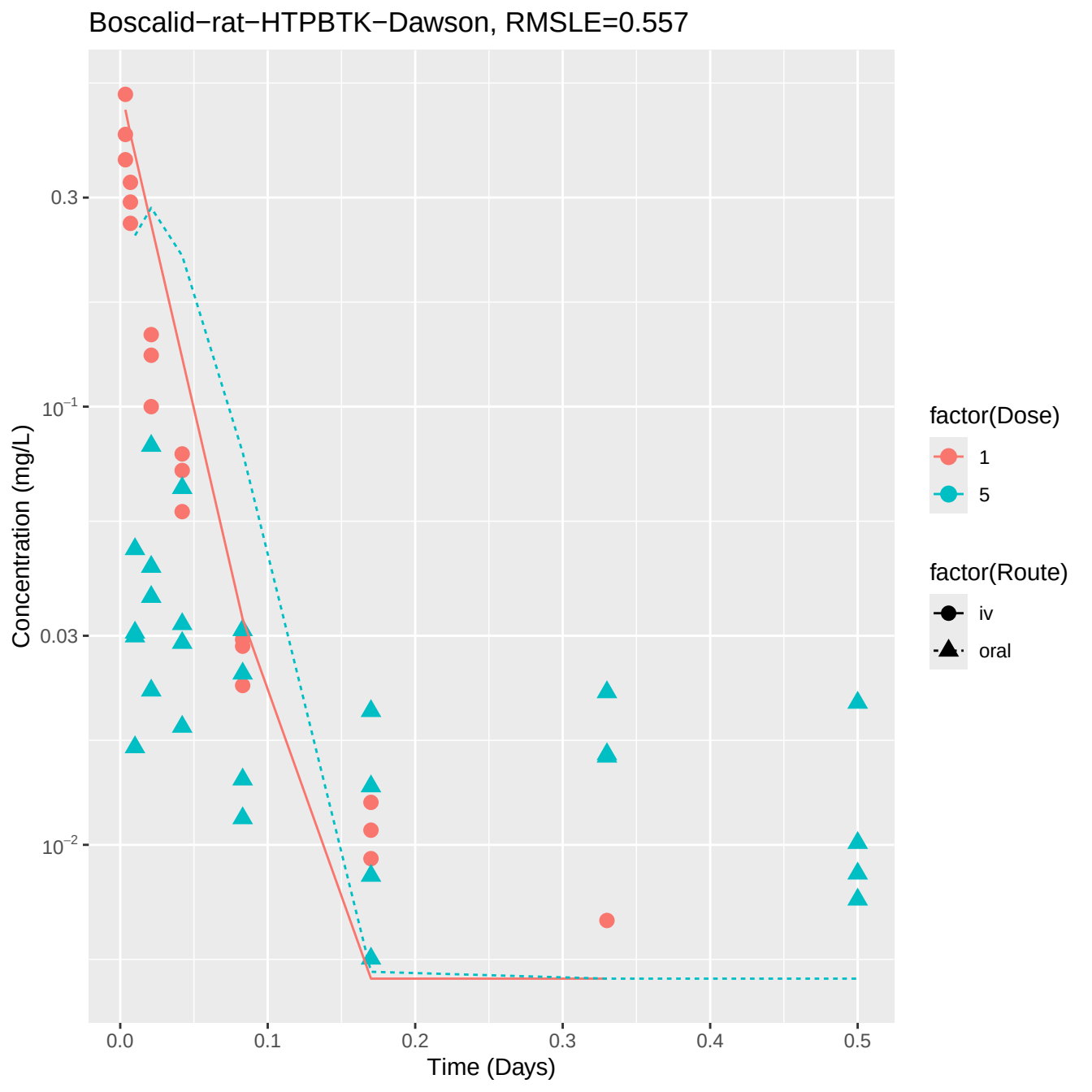

Chloridazon−rat−HTPBTK−Pradeep, RMSLE=0.541
Concentration (mg/L)
1
10^{-2}
10^{-4}
0
1
2
3
Time (Days)
factor(Dose)
0.734
0.827
0.847
4.12
4.18
4.2
factor(Route)
iv
oral

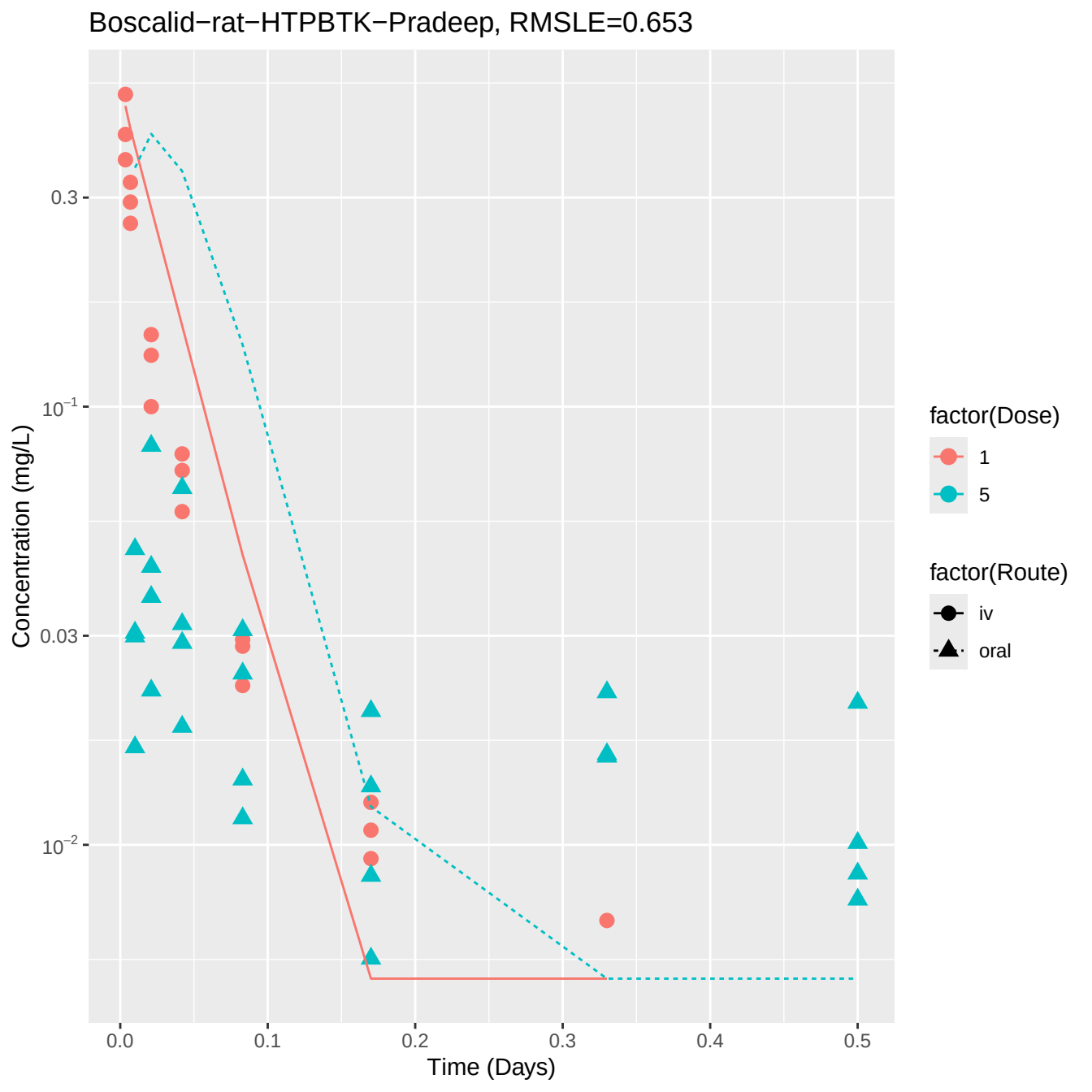

Chloridazon−rat−HTPBTK−Ensemble, RMSLE=0.635
Concentration (mg/L)
1
10^{-2}
10^{-4}
0
1
2
3
Time (Days)
factor(Dose)
0.734
0.827
0.847
4.12
4.18
4.2
factor(Route)
iv
oral

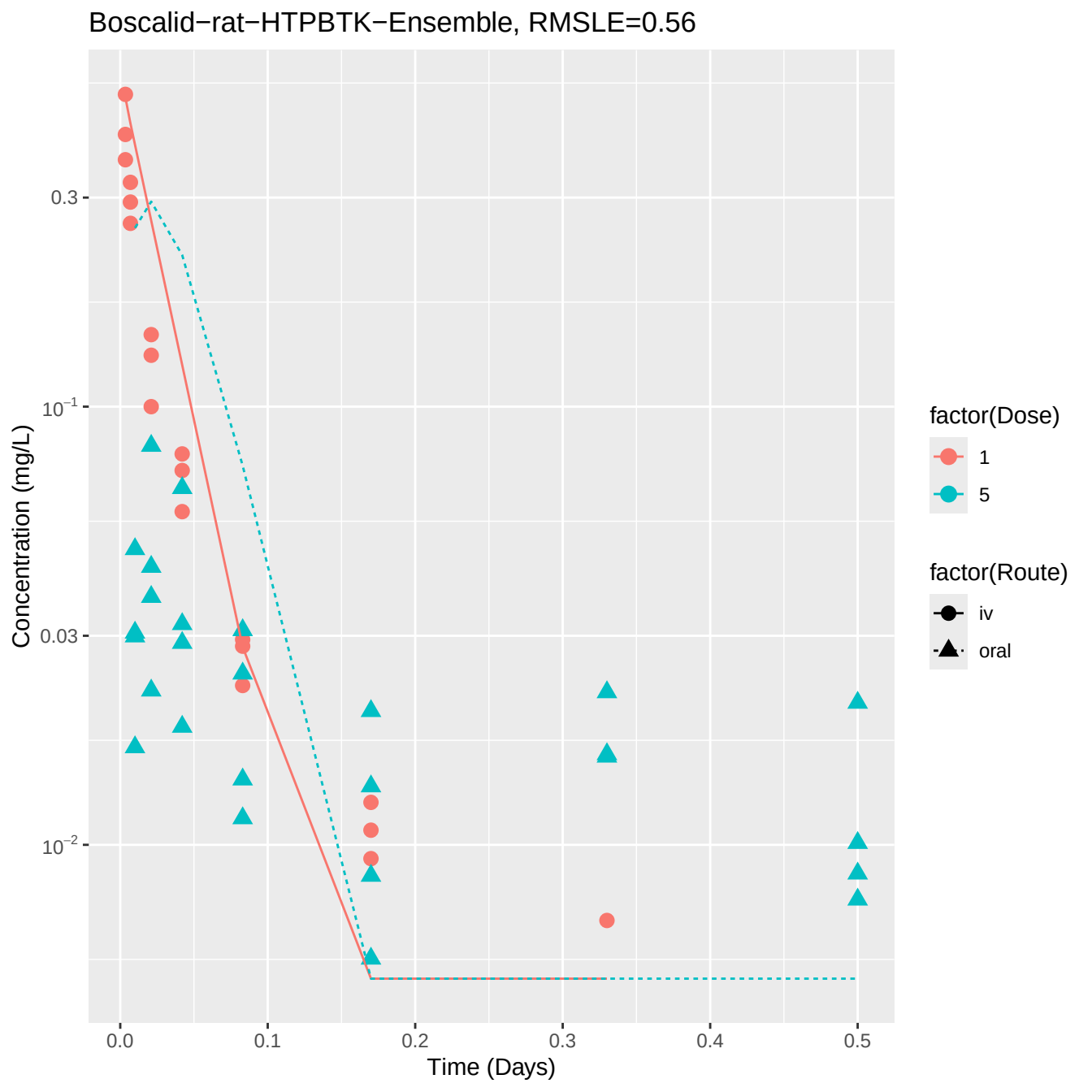

Chloridazon−rat−In Vivo Fits, RMSLE=0.344
Concentration (mg/L)
1
10^{-2}
10^{-4}
0
1
2
3
Time (Days)
factor(Dose)
0.734
0.827
0.847
4.12
4.18
4.2
factor(Route)
iv
oral

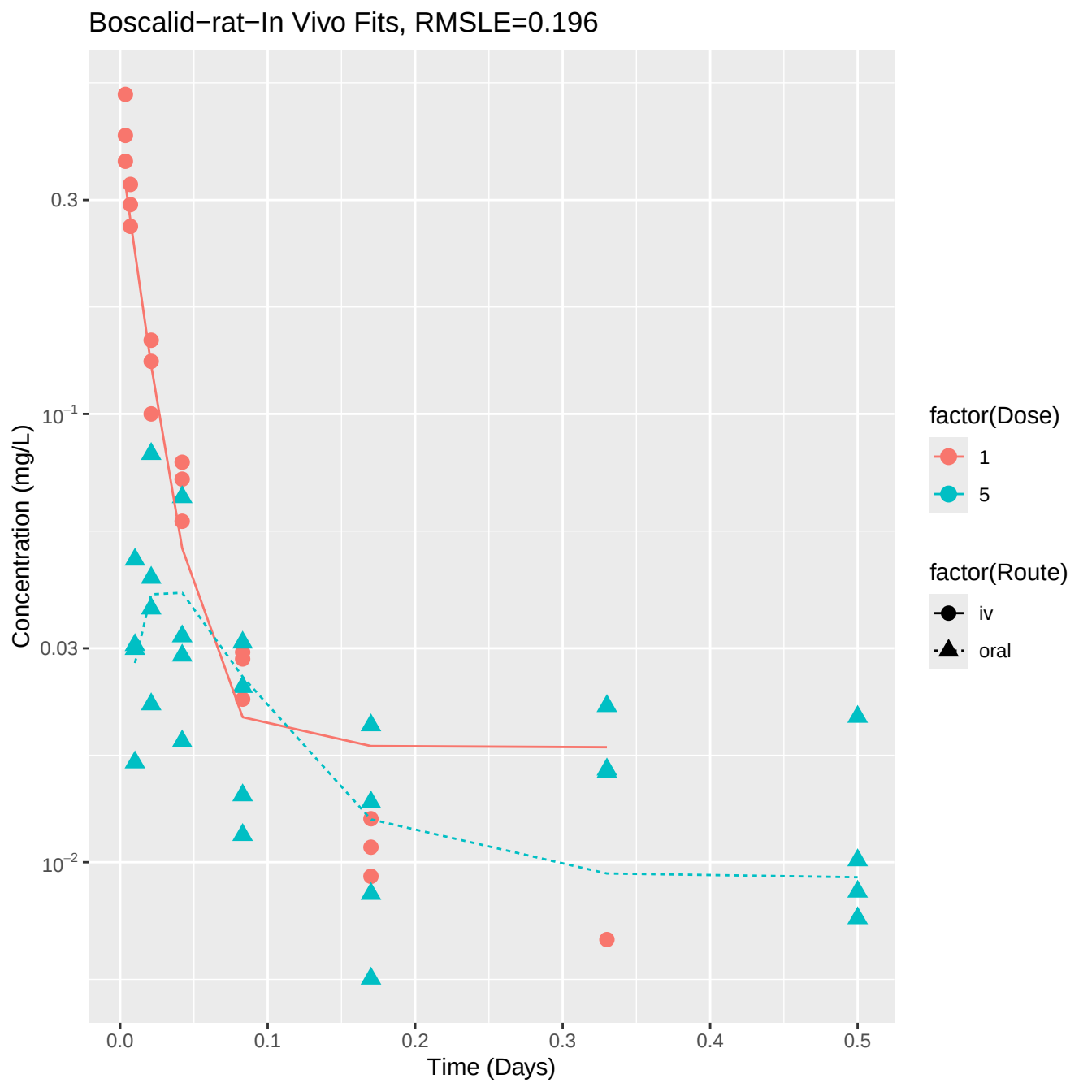

Boscalid−rat−HTPBTK−InVitro, RMSLE=0.608
Concentration (mg/L)
0.3
10^{-1}
0.03
10^{-2}
0.0
0.1
0.2
0.3
0.4
0.5
Time (Days)
factor(Dose)
1
5
factor(Route)
iv
oral

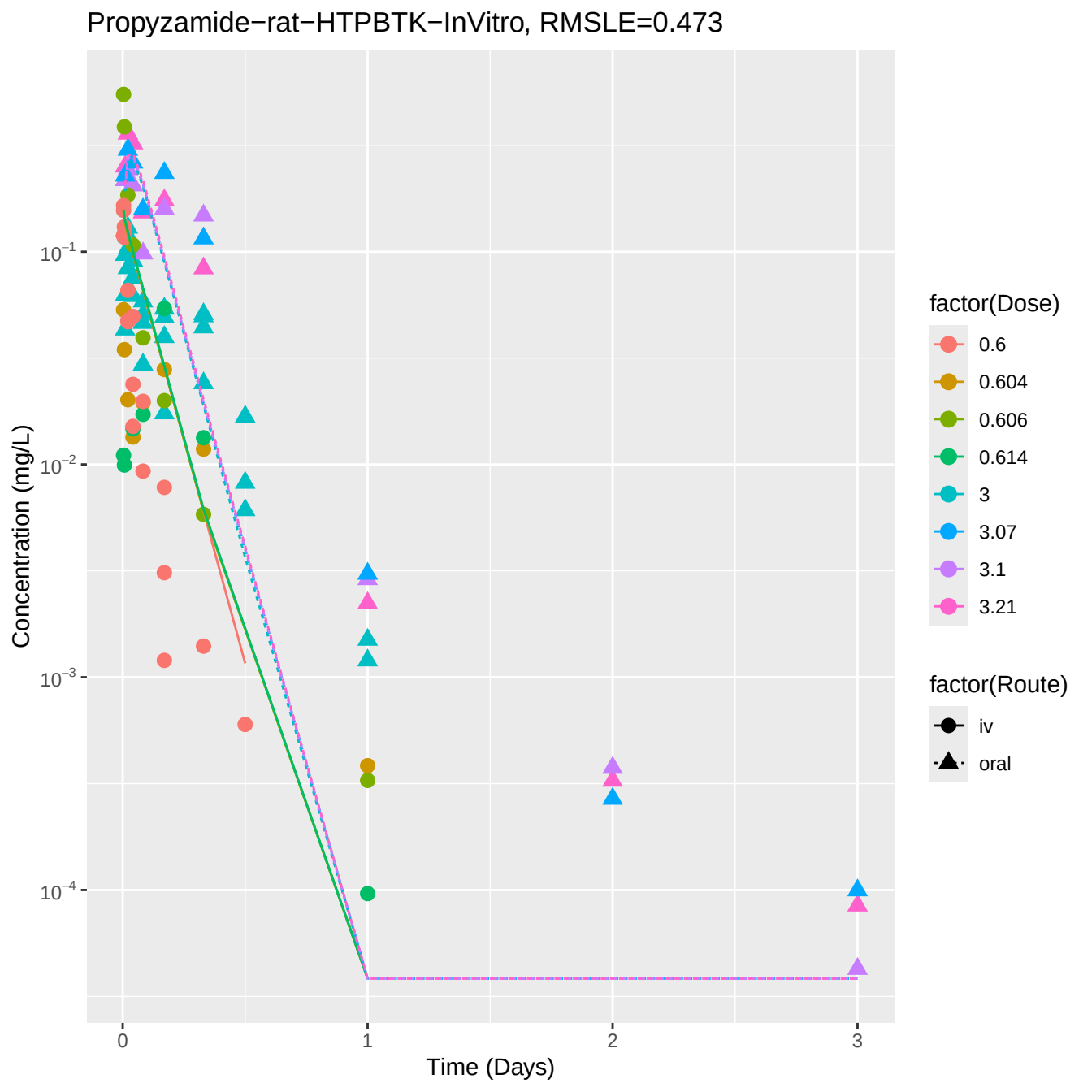

Boscalid−rat−HTPBTK−ADMET, RMSLE=0.752
Concentration (mg/L)
0.3
10^{-1}
0.03
10^{-2}
0.0
0.1
0.2
0.3
0.4
0.5
Time (Days)
factor(Dose)
1
5
factor(Route)
iv
oral

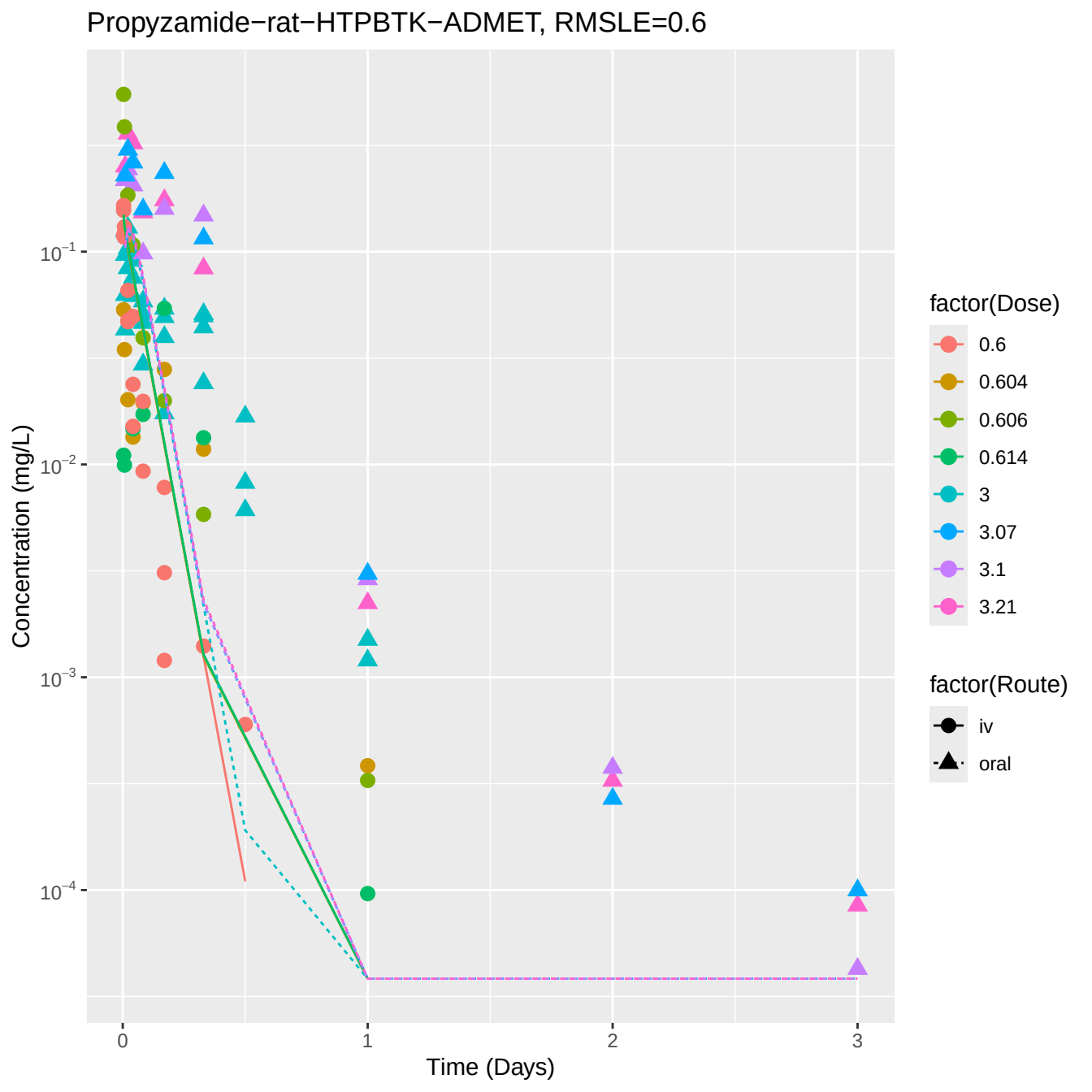

Boscalid−rat−HTPBTK−Dawson, RMSLE=0.557
Concentration (mg/L)
0.3
10^{-1}
0.03
10^{-2}
0.0
0.1
0.2
0.3
0.4
0.5
Time (Days)
factor(Dose)
1
5
factor(Route)
iv
oral

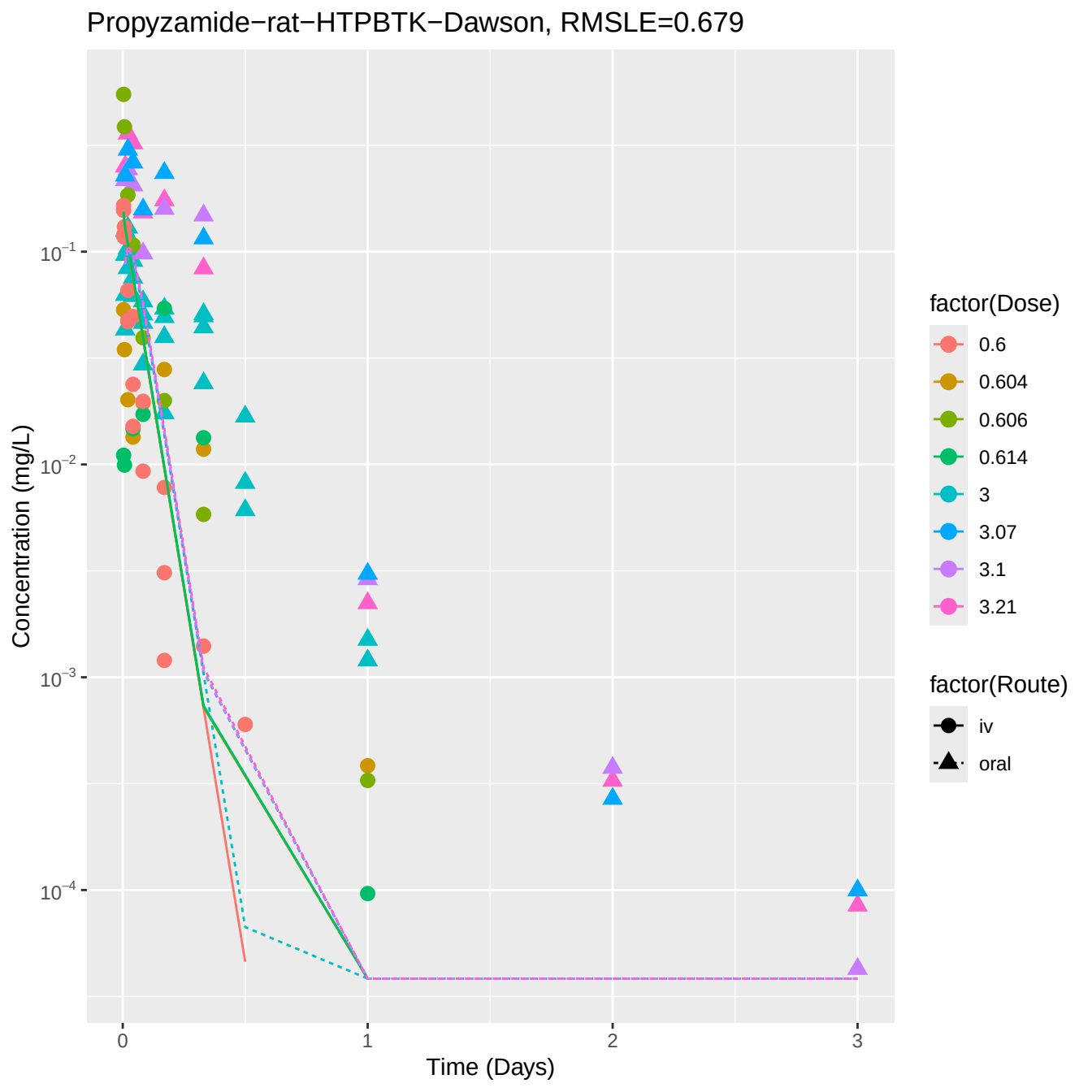

Boscalid−rat−HTPBTK−Pradeep, RMSLE=0.653
Concentration (mg/L)
0.3
10^{-1}
0.03
10^{-2}
0.0
0.1
0.2
0.3
0.4
0.5
Time (Days)
factor(Dose)
1
5
factor(Route)
iv
oral

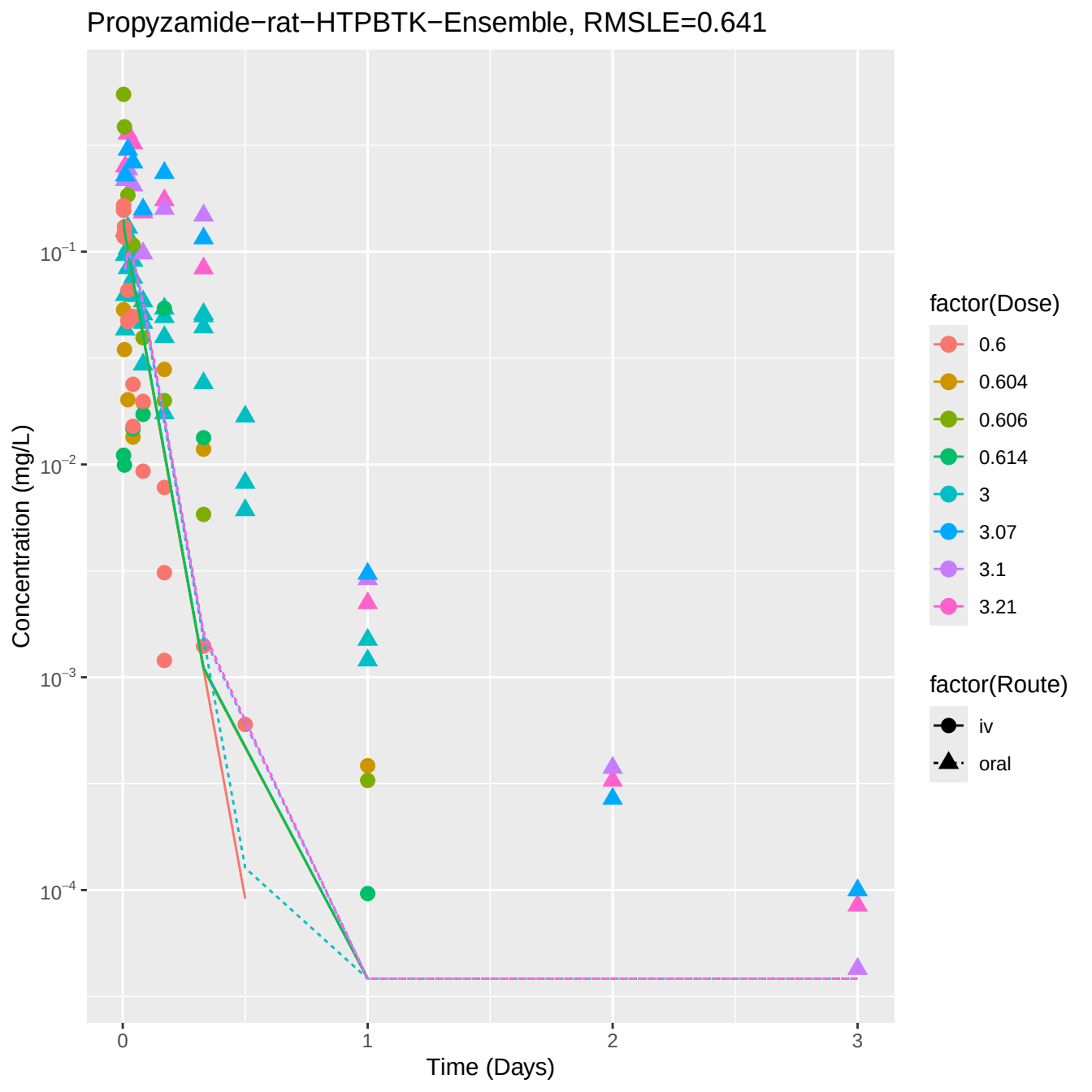

Boscalid−rat−HTPBTK−Ensemble, RMSLE=0.56
Concentration (mg/L)
0.3
10^{-1}
0.03
10^{-2}
0.0
0.1
0.2
0.3
0.4
0.5
Time (Days)
factor(Dose)
1
5
factor(Route)
iv
oral

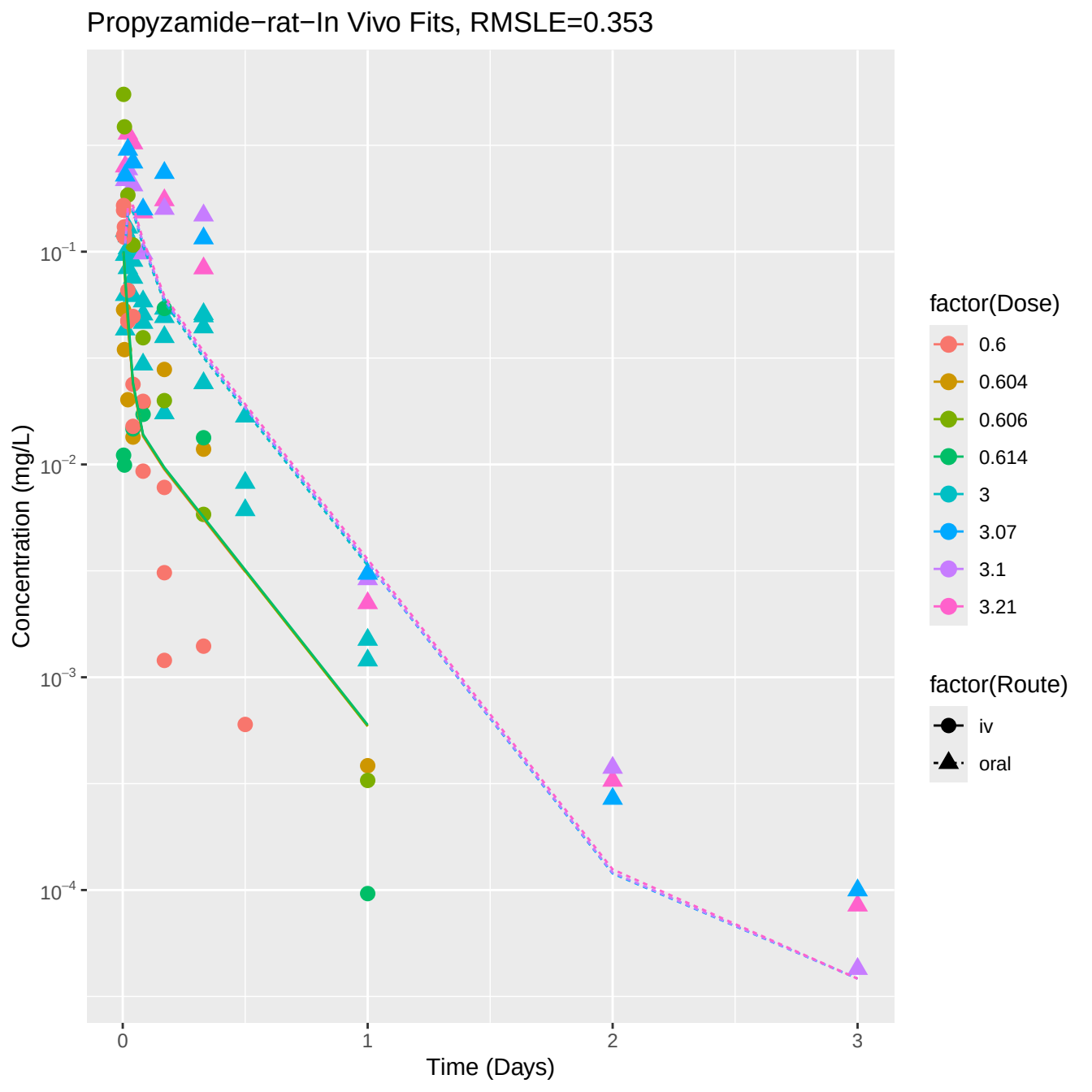

Boscalid−rat−In Vivo Fits, RMSLE=0.196
Concentration (mg/L)
0.3
10^{-1}
0.03
10^{-2}
0.0
0.1
0.2
0.3
0.4
0.5
Time (Days)
factor(Dose)
1
5
factor(Route)
iv
oral

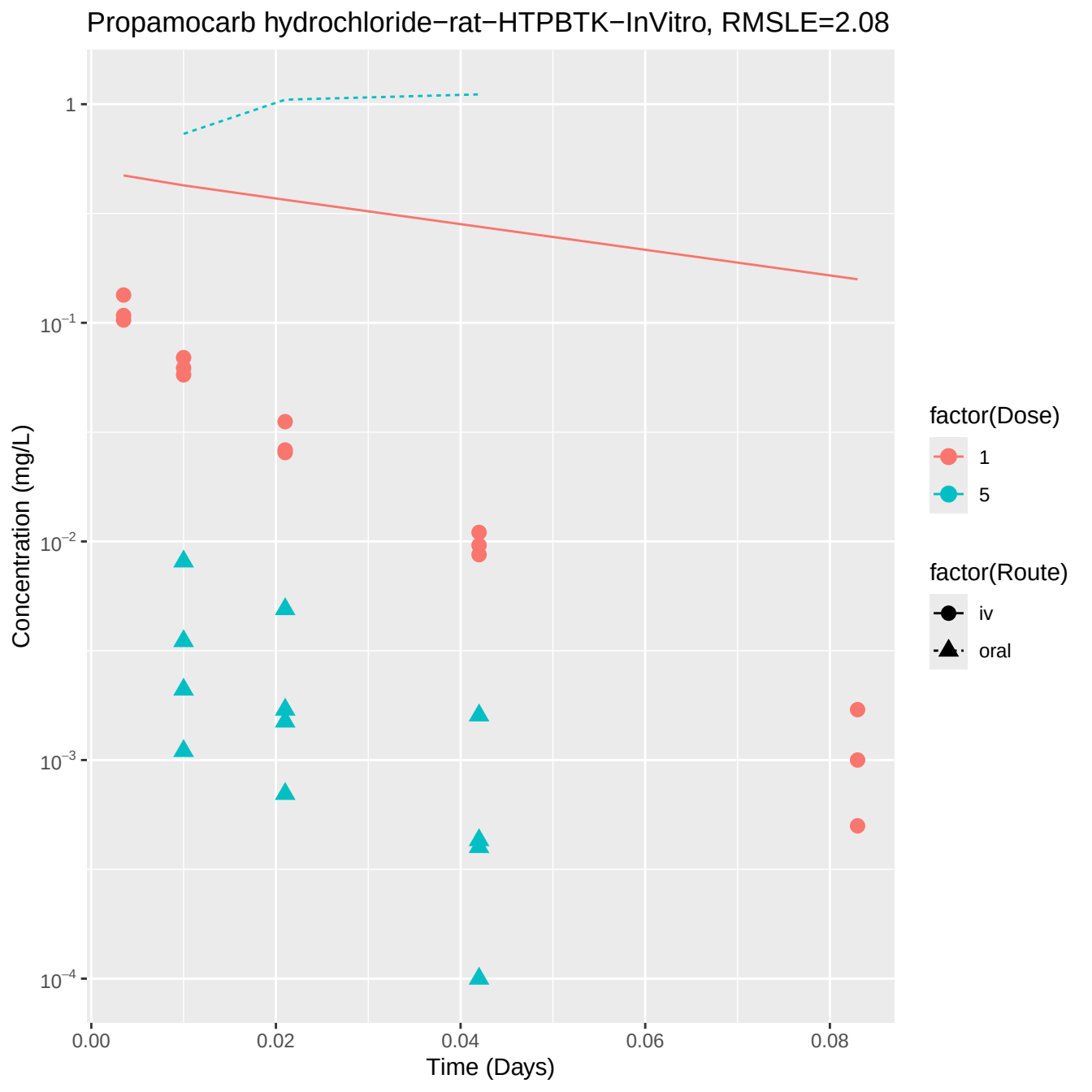

Propyzamide−rat−HTPBTK−InVitro, RMSLE=0.473
Concentration (mg/L)
10^{-1}
10^{-2}
10^{-3}
10^{-4}
0
1
2
3
Time (Days)
factor(Dose)
0.6
0.604
0.606
0.614
3
3.07
3.1
3.21
factor(Route)
iv
oral

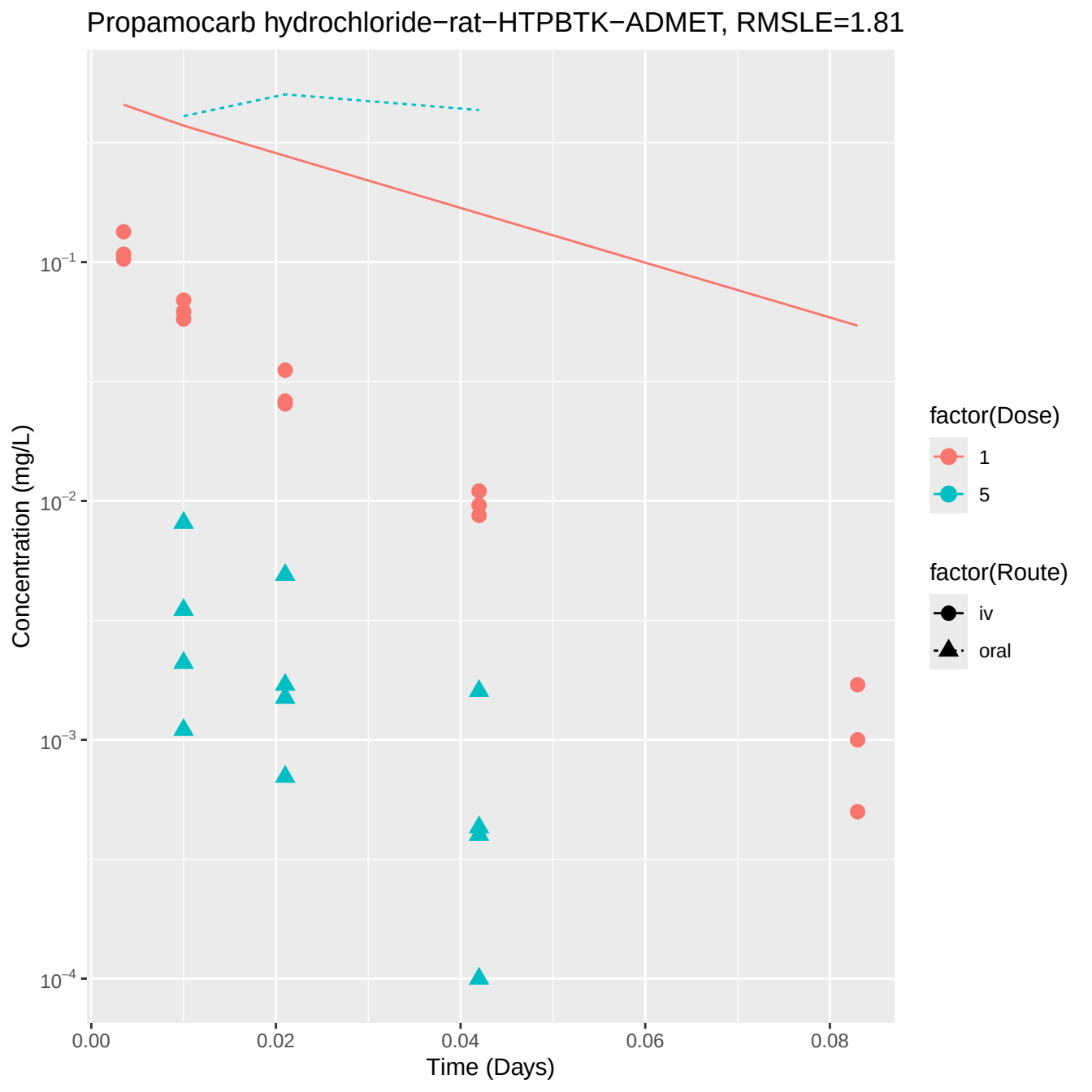

Propyzamide−rat−HTPBTK−ADMET, RMSLE=0.6
Concentration (mg/L)
10^{-1}
10^{-2}
10^{-3}
10^{-4}
0
1
2
3
Time (Days)
factor(Dose)
0.6
0.604
0.606
0.614
3
3.07
3.1
3.21
factor(Route)
iv
oral

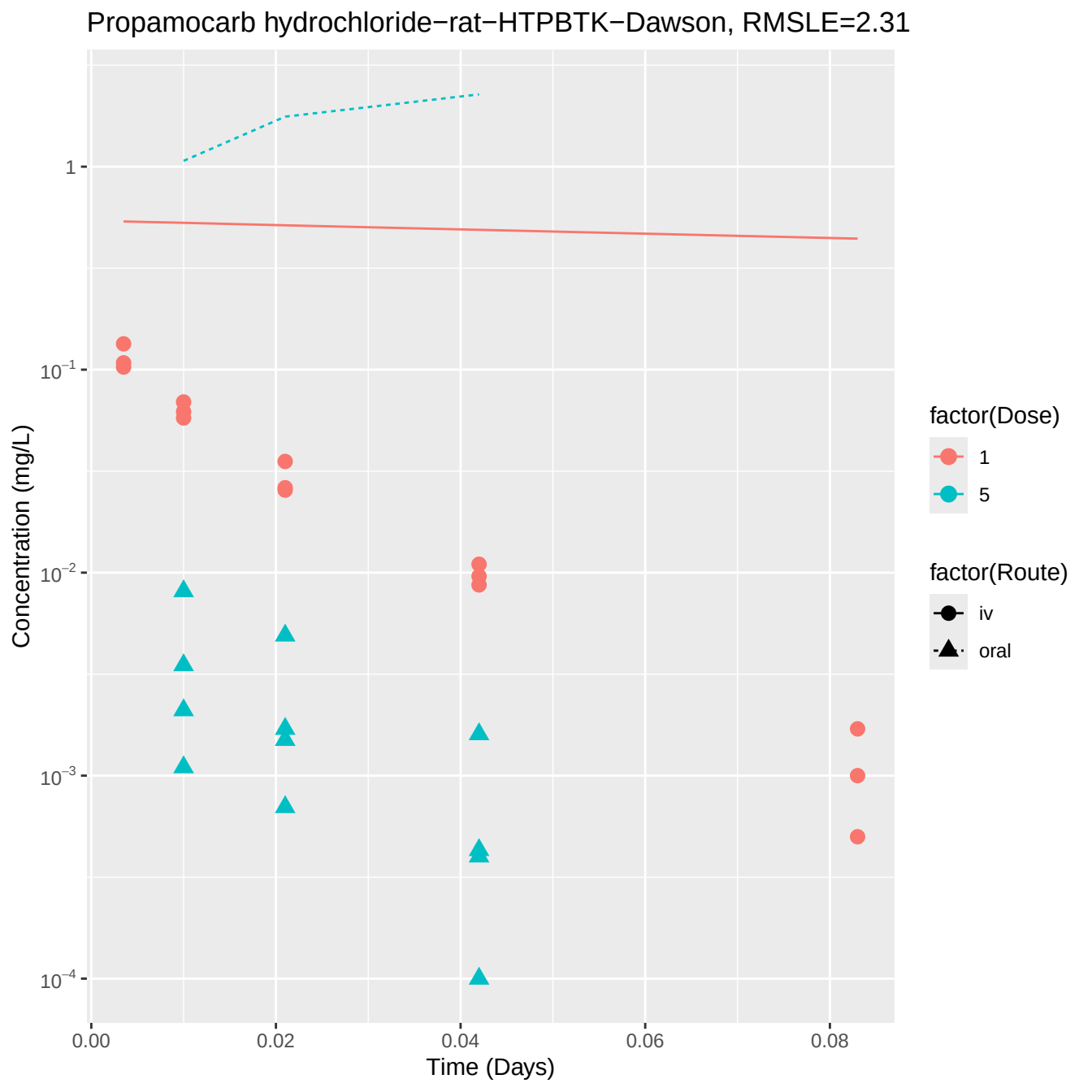

Propyzamide−rat−HTPBTK−Dawson, RMSLE=0.679
Concentration (mg/L)
10^{-1}
10^{-2}
10^{-3}
10^{-4}
0
1
2
3
Time (Days)
factor(Dose)
0.6
0.604
0.606
0.614
3
3.07
3.1
3.21
factor(Route)
iv
oral

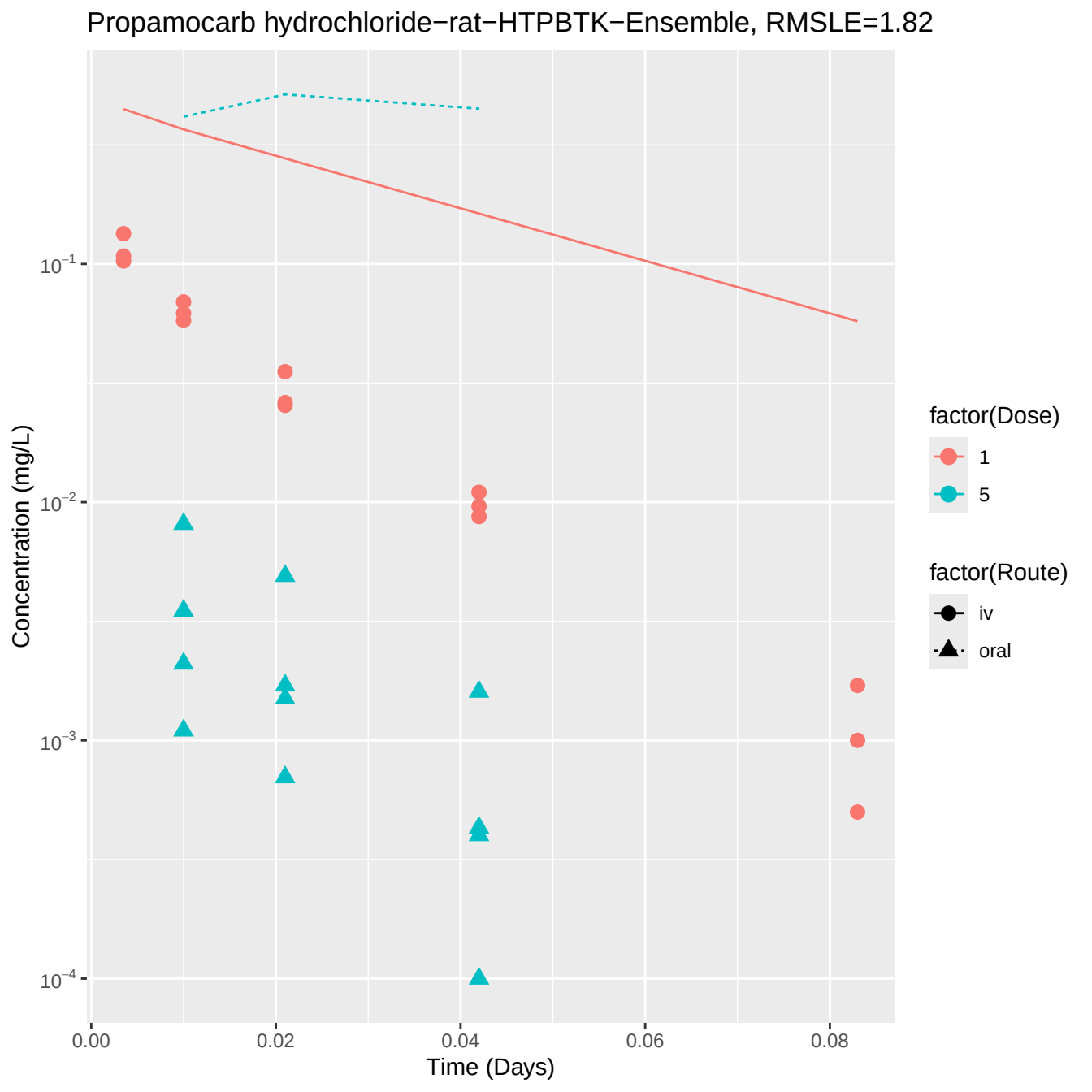

Propyzamide−rat−HTPBTK−Ensemble, RMSLE=0.641
Concentration (mg/L)
10^{-1}
10^{-2}
10^{-3}
10^{-4}
0
1
2
3
Time (Days)
factor(Dose)
0.6
0.604
0.606
0.614
3
3.07
3.1
3.21
factor(Route)
iv
oral

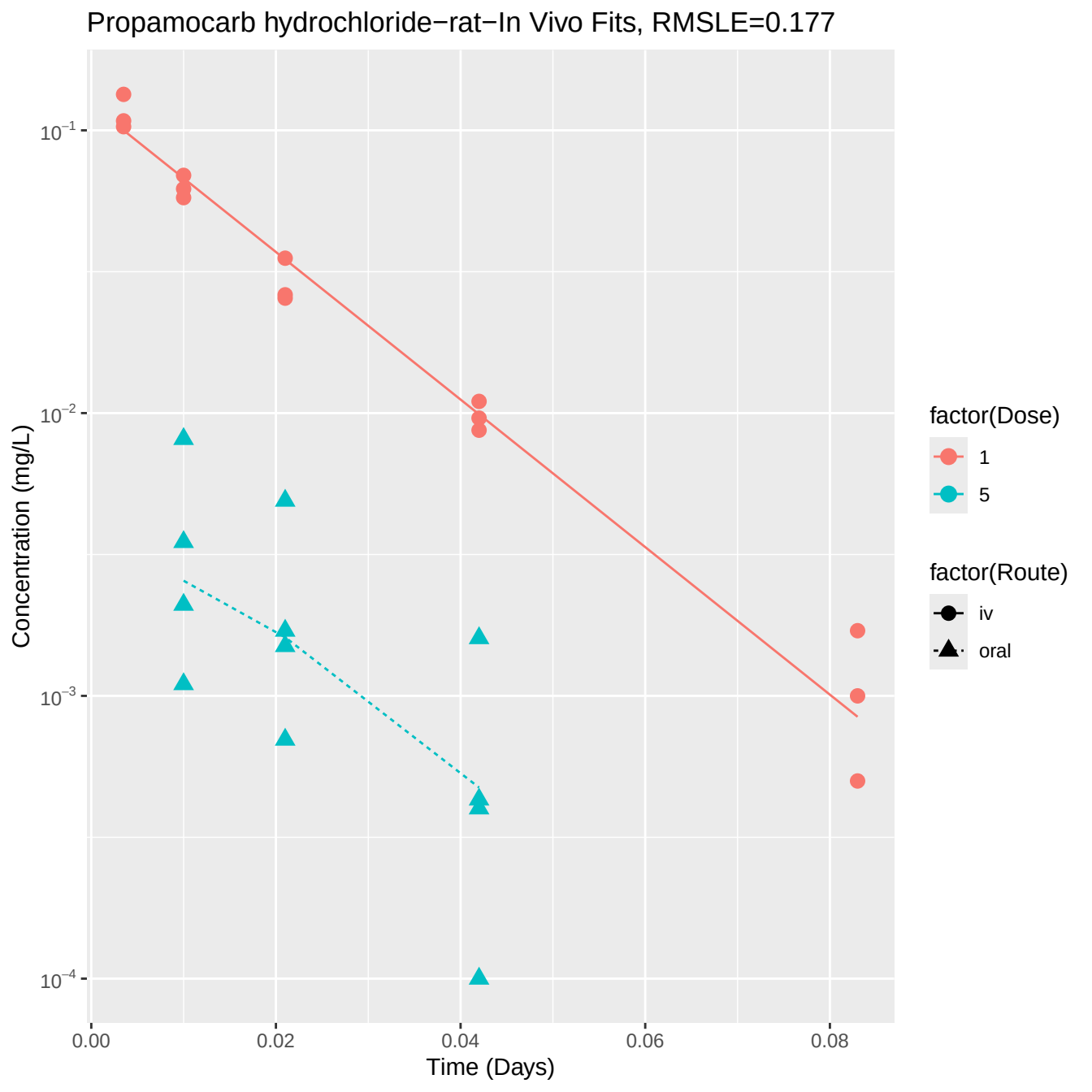

Propyzamide−rat−In Vivo Fits, RMSLE=0.353
Concentration (mg/L)
10^{-1}
10^{-2}
10^{-3}
10^{-4}
0
1
2
3
Time (Days)
factor(Dose)
0.6
0.604
0.606
0.614
3
3.07
3.1
3.21
factor(Route)
iv
oral

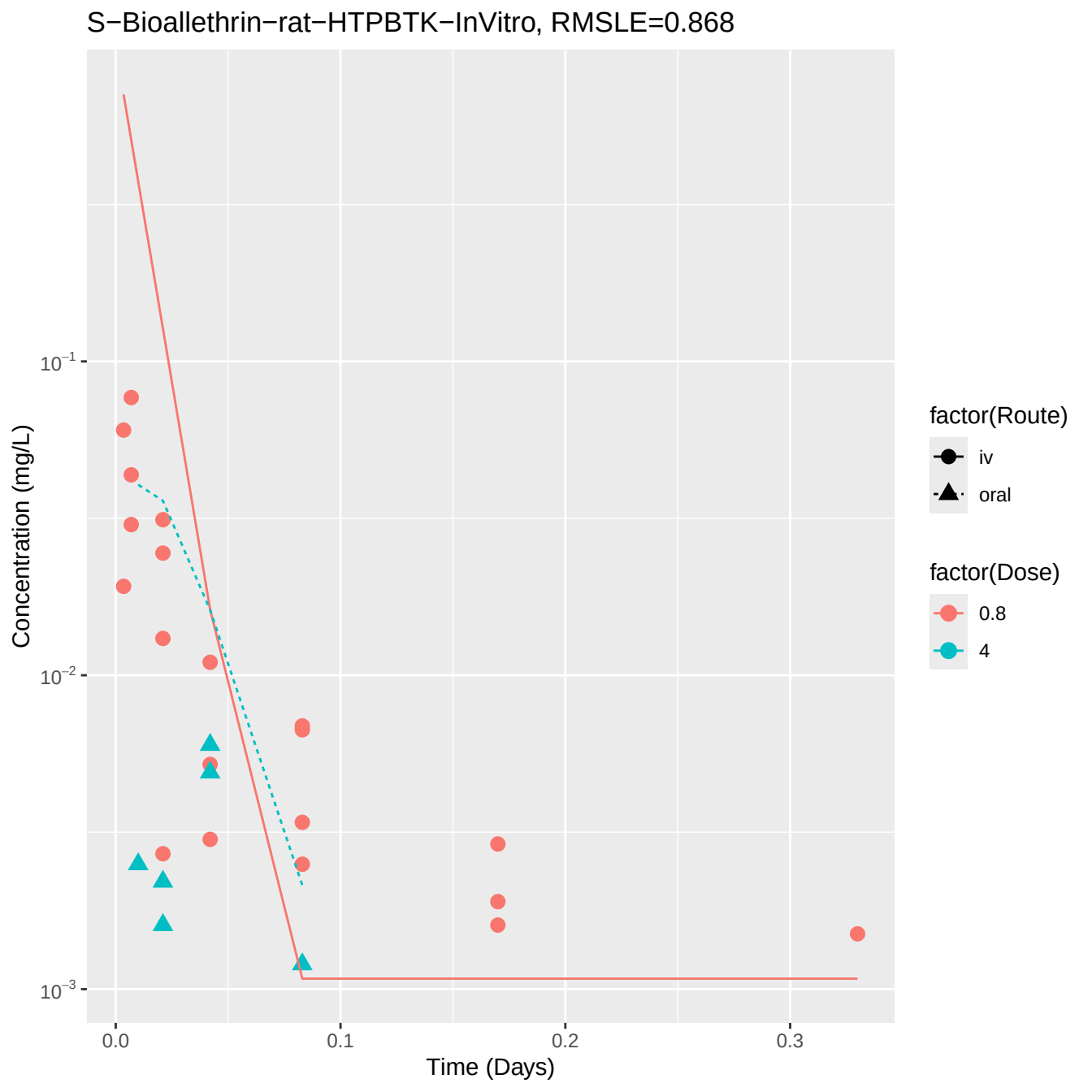

Propamocarb hydrochloride−rat−HTPBTK−InVitro, RMSLE=2.08
Concentration (mg/L)
1
10^{-1}
10^{-2}
10^{-3}
10^{-4}
0.00
0.02
0.04
0.06
0.08
Time (Days)
factor(Dose)
1
5
factor(Route)
iv
oral

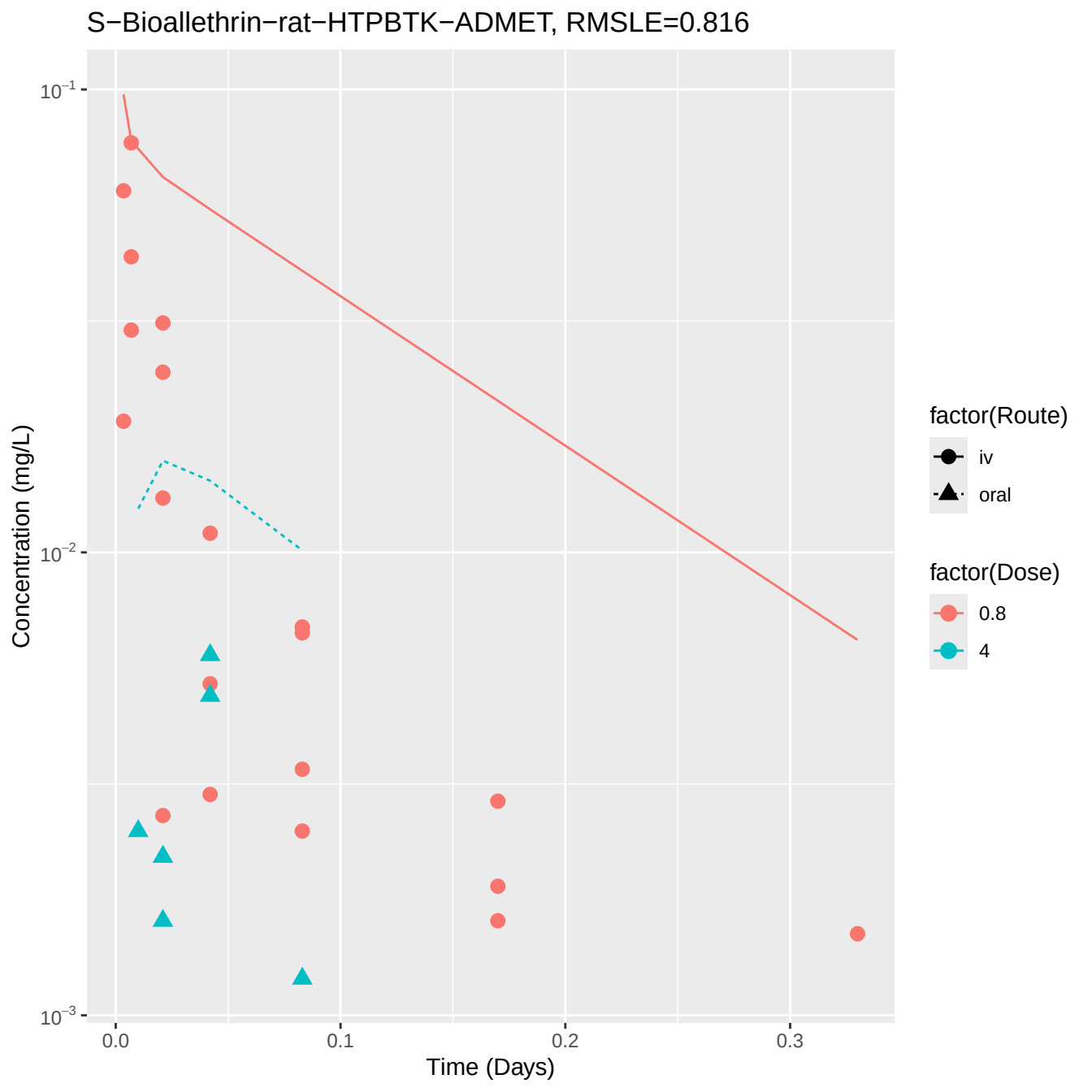

Propamocarb hydrochloride−rat−HTPBTK−ADMET, RMSLE=1.81
Concentration (mg/L)
10^{-1}
10^{-2}
10^{-3}
10^{-4}
0.00
0.02
0.04
0.06
0.08
Time (Days)
factor(Dose)
1
5
factor(Route)
iv
oral

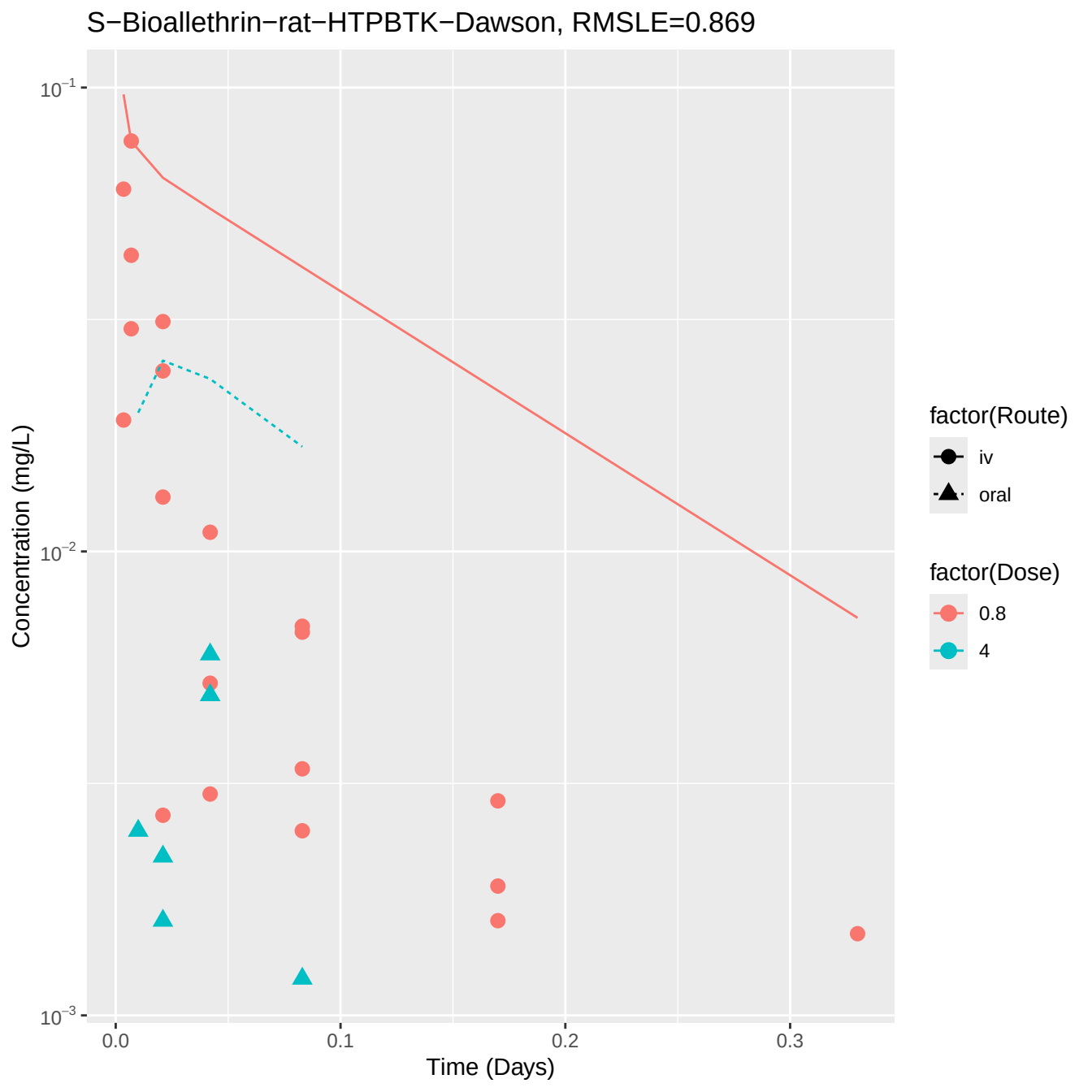

Propamocarb hydrochloride−rat−HTPBTK−Dawson, RMSLE=2.31
Concentration (mg/L)
1
10^{-1}
10^{-2}
10^{-3}
10^{-4}
0.00
0.02
0.04
0.06
0.08
Time (Days)
factor(Dose)
1
5
factor(Route)
iv
oral

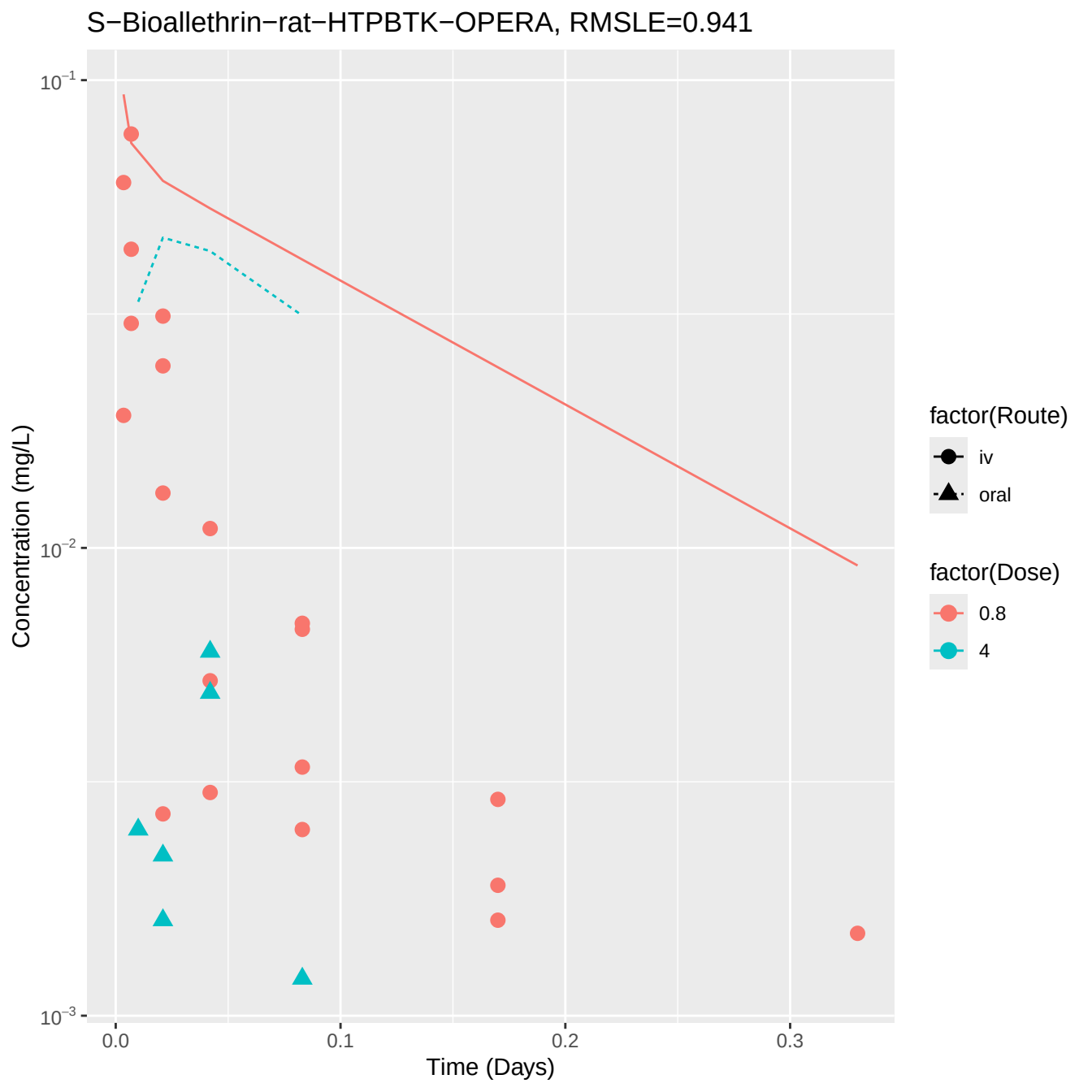

Propamocarb hydrochloride−rat−HTPBTK−Ensemble, RMSLE=1.82
Concentration (mg/L)
10^{-1}
10^{-2}
10^{-3}
10^{-4}
0.00
0.02
0.04
0.06
0.08
Time (Days)
factor(Dose)
1
5
factor(Route)
iv
oral

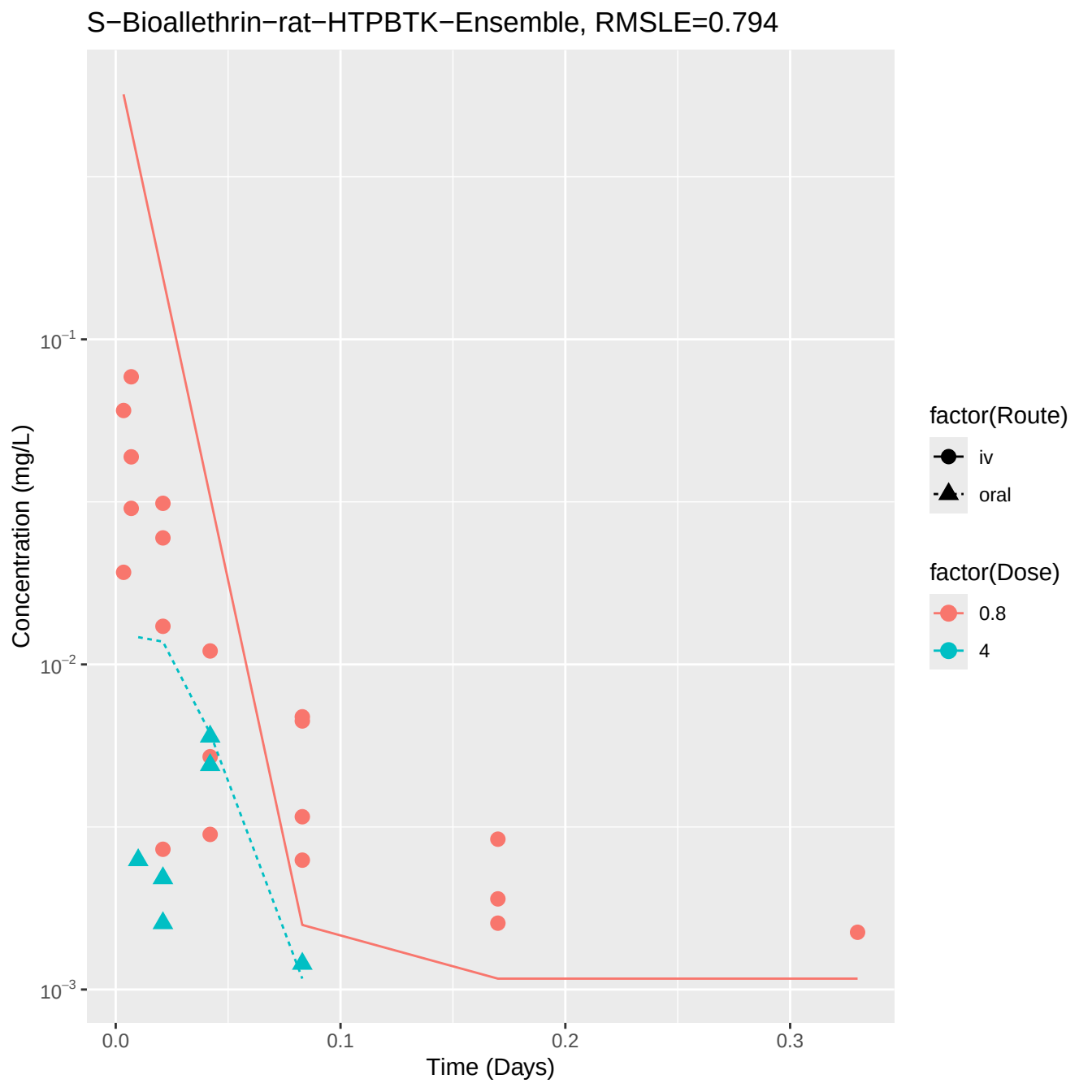

Propamocarb hydrochloride−rat−In Vivo Fits, RMSLE=0.177
Concentration (mg/L)
10^{-1}
10^{-2}
10^{-3}
10^{-4}
0.00
0.02
0.04
0.06
0.08
Time (Days)
factor(Dose)
1
5
factor(Route)
iv
oral

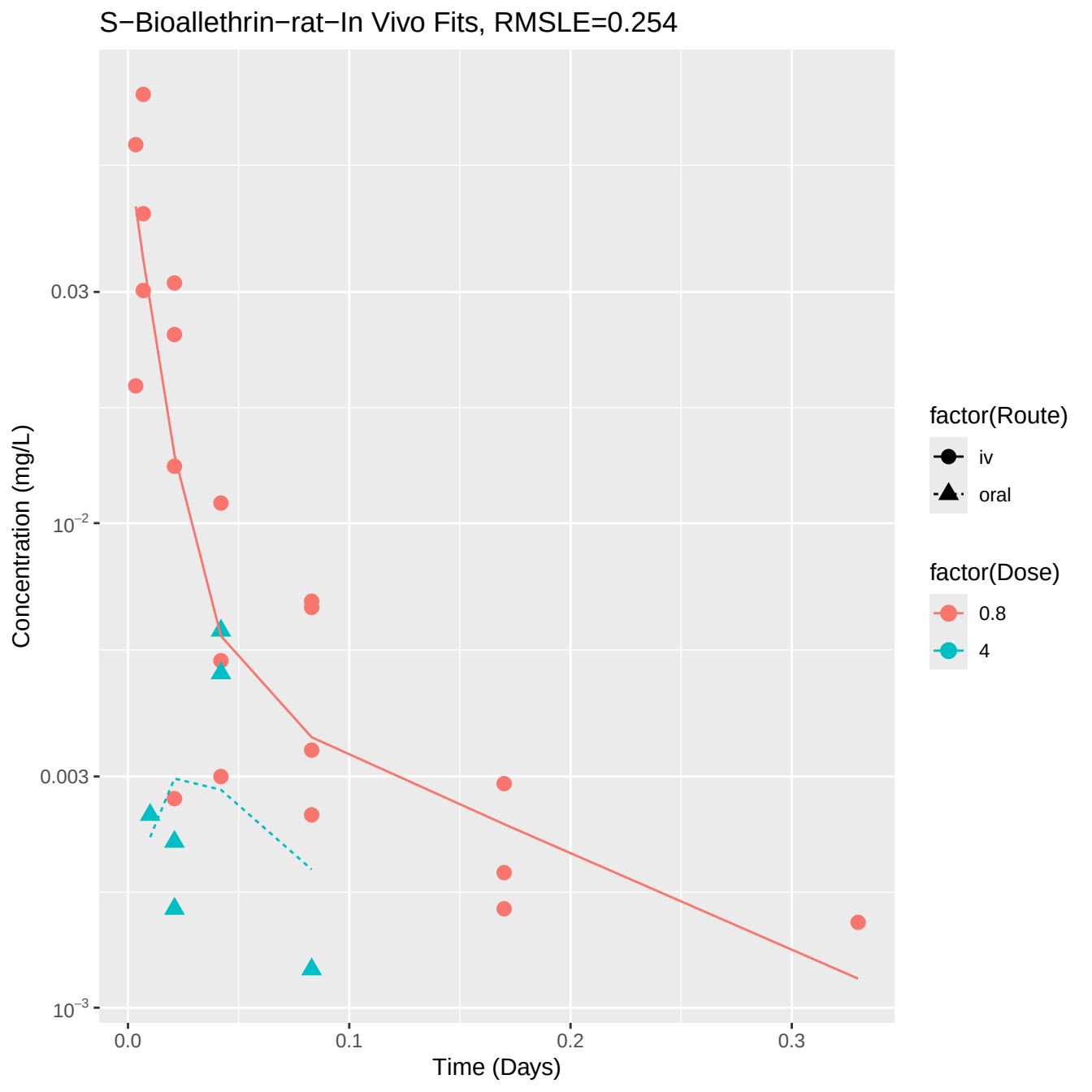

S−Bioallethrin−rat−HTPBTK−InVitro, RMSLE=0.868
Concentration (mg/L)
10^{-1}
10^{-2}
10^{-3}
0.0
0.1
0.2
0.3
Time (Days)
factor(Route)
iv
oral
factor(Dose)
0.8
4

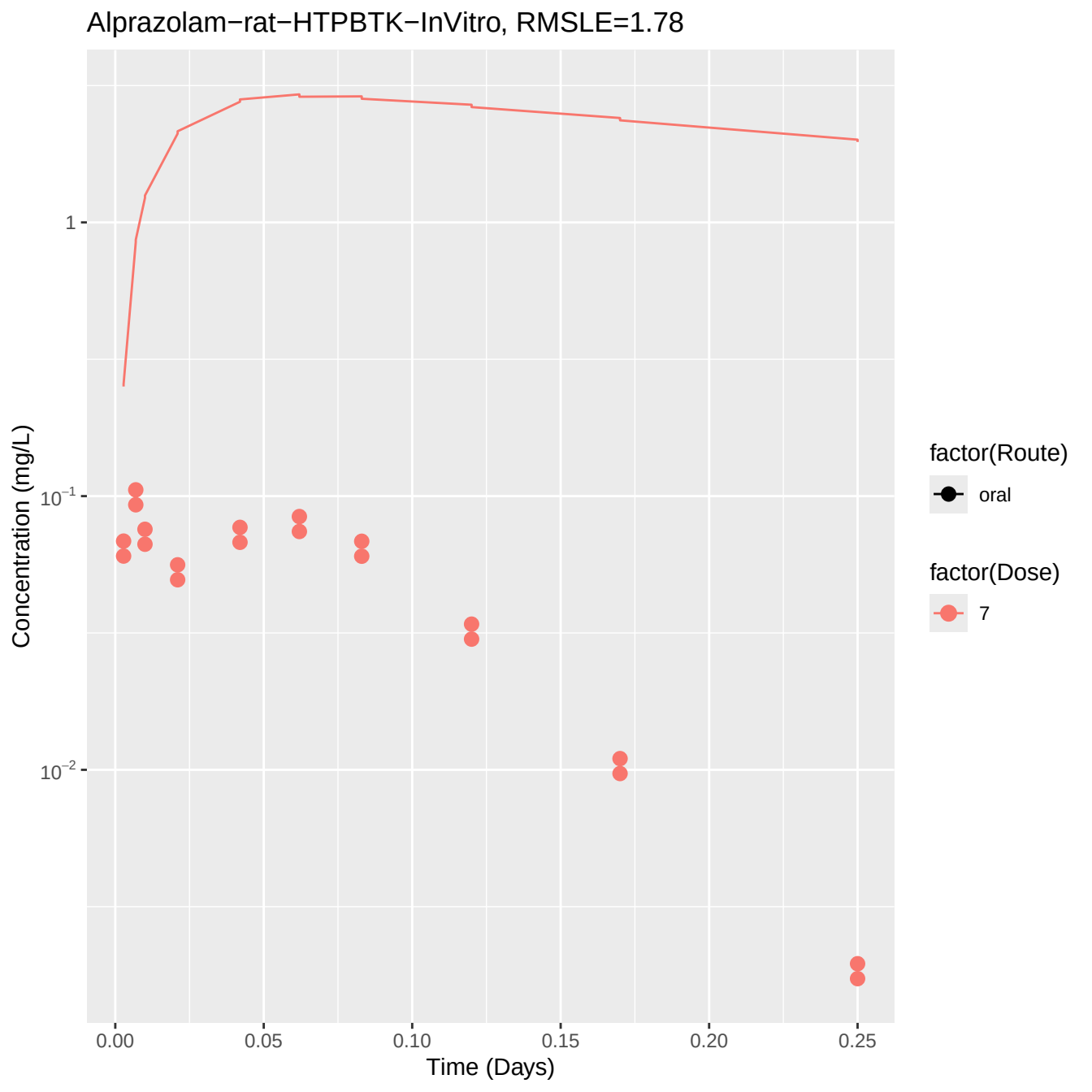

S−Bioallethrin−rat−HTPBTK−ADMET, RMSLE=0.816
Concentration (mg/L)
10^{-1}
10^{-2}
10^{-3}
0.0
0.1
0.2
0.3
Time (Days)
factor(Route)
iv
oral
factor(Dose)
0.8
4

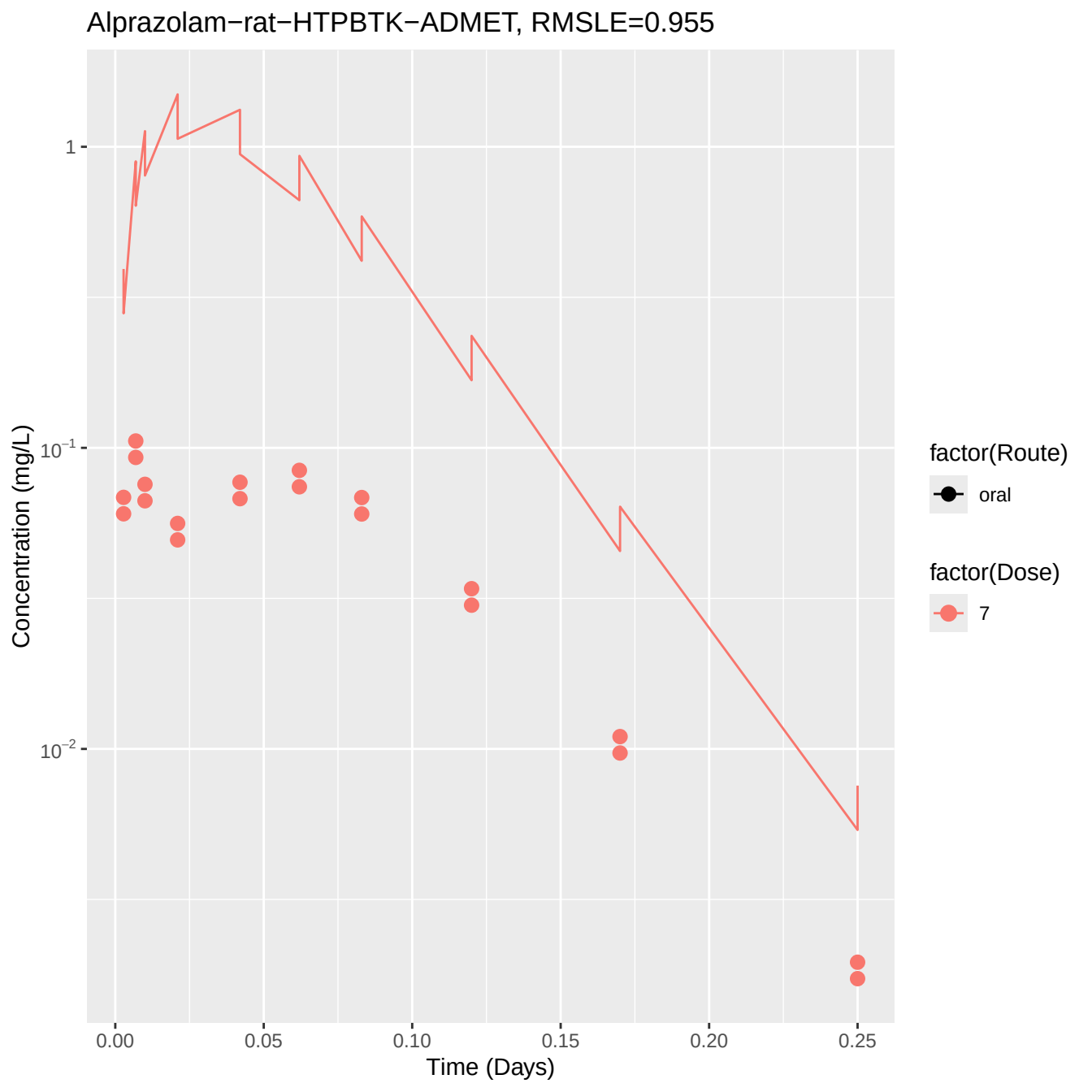

S−Bioallethrin−rat−HTPBTK−Dawson, RMSLE=0.869
Concentration (mg/L)
10^{-1}
10^{-2}
10^{-3}
0.0
0.1
0.2
0.3
Time (Days)
factor(Route)
iv
oral
factor(Dose)
0.8
4

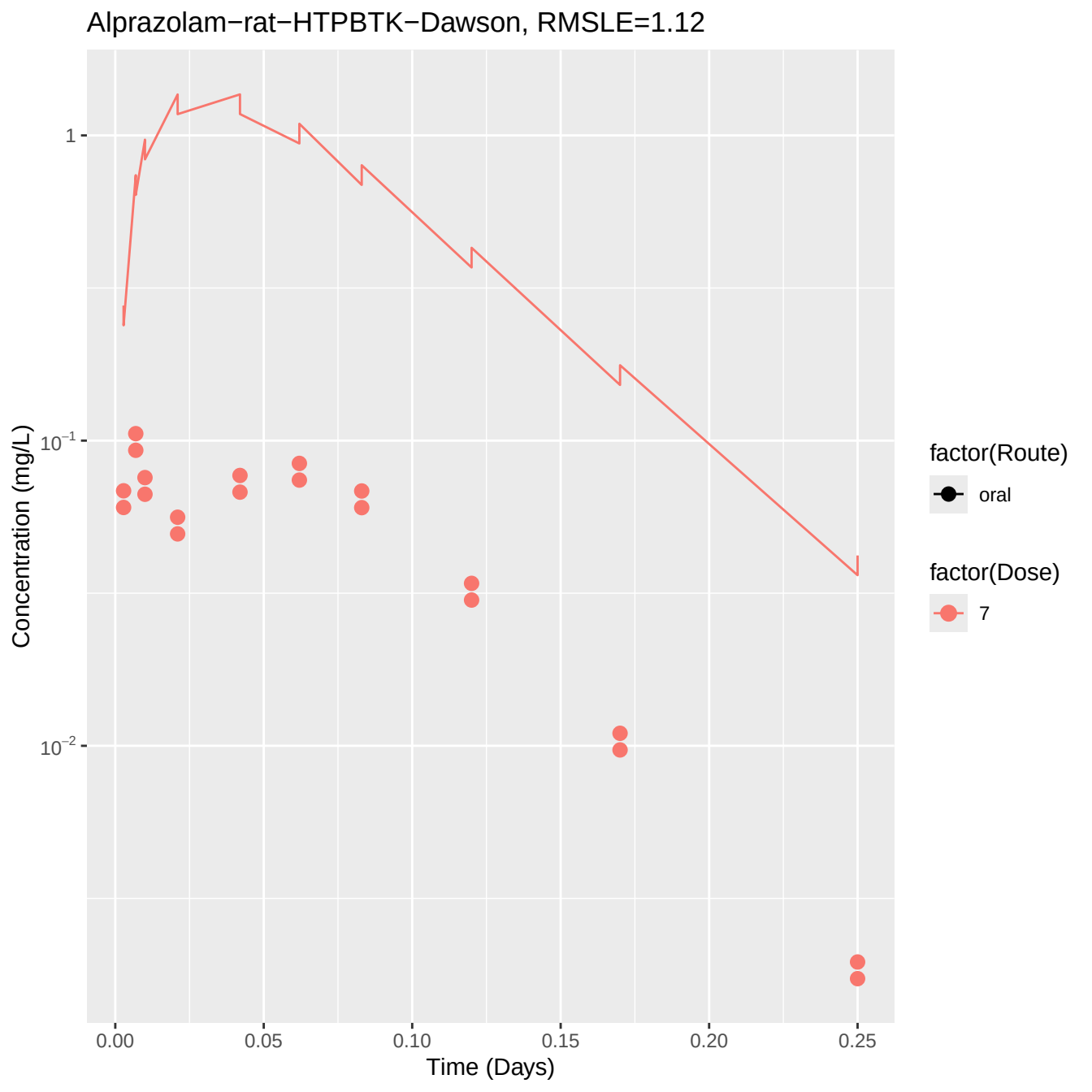

S−Bioallethrin−rat−HTPBTK−OPERA, RMSLE=0.941
Concentration (mg/L)
10^{-1}
10^{-2}
10^{-3}
0.0
0.1
0.2
0.3
Time (Days)
factor(Route)
iv
oral
factor(Dose)
0.8
4

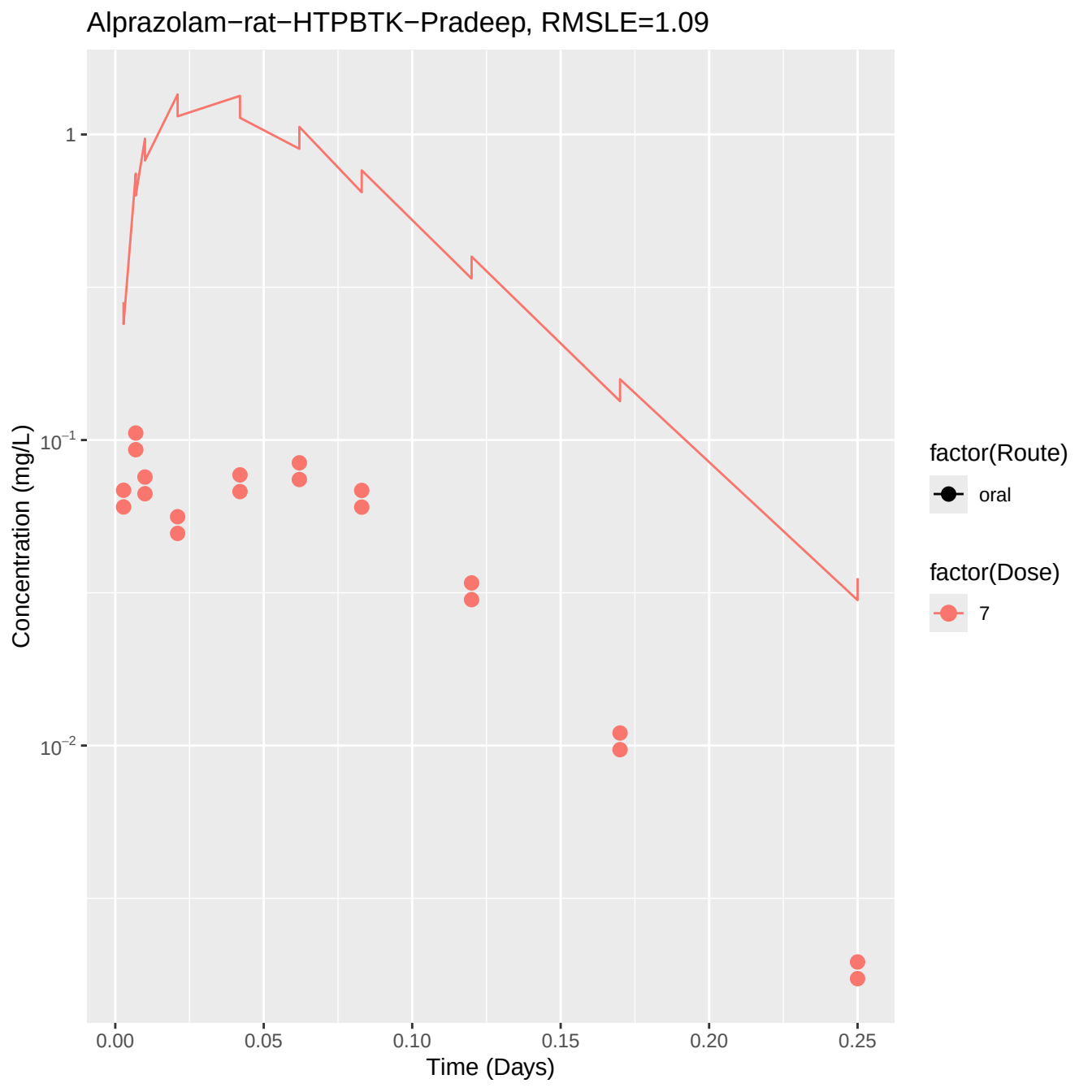

S−Bioallethrin−rat−HTPBTK−Ensemble, RMSLE=0.794
Concentration (mg/L)
10^{-1}
10^{-2}
10^{-3}
0.0
0.1
0.2
0.3
Time (Days)
factor(Route)
iv
oral
factor(Dose)
0.8
4

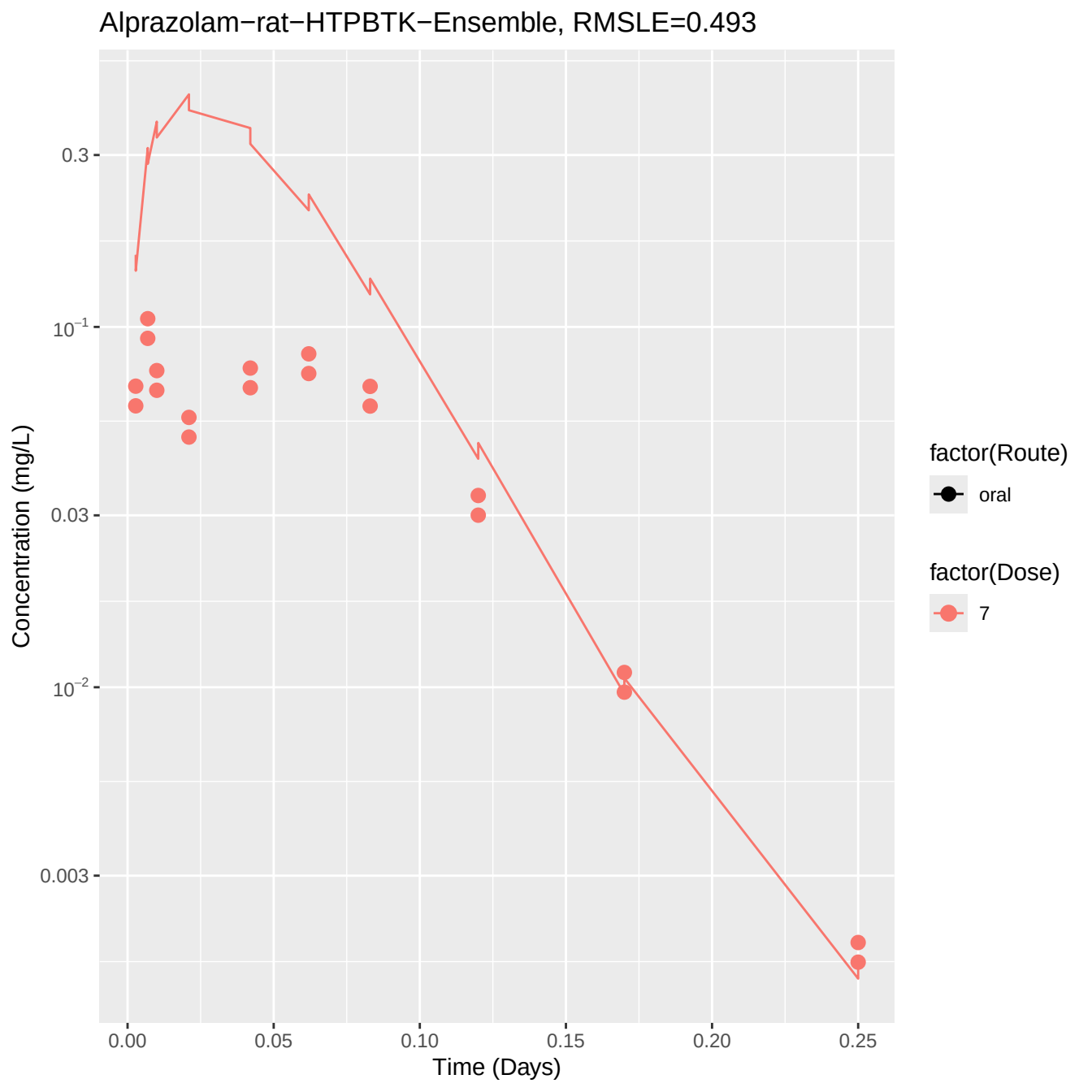

S−Bioallethrin−rat−In Vivo Fits, RMSLE=0.254
Concentration (mg/L)
0.03
10^{-2}
0.003
10^{-3}
0.0
0.1
0.2
0.3
Time (Days)
factor(Route)
iv
oral
factor(Dose)
0.8
4

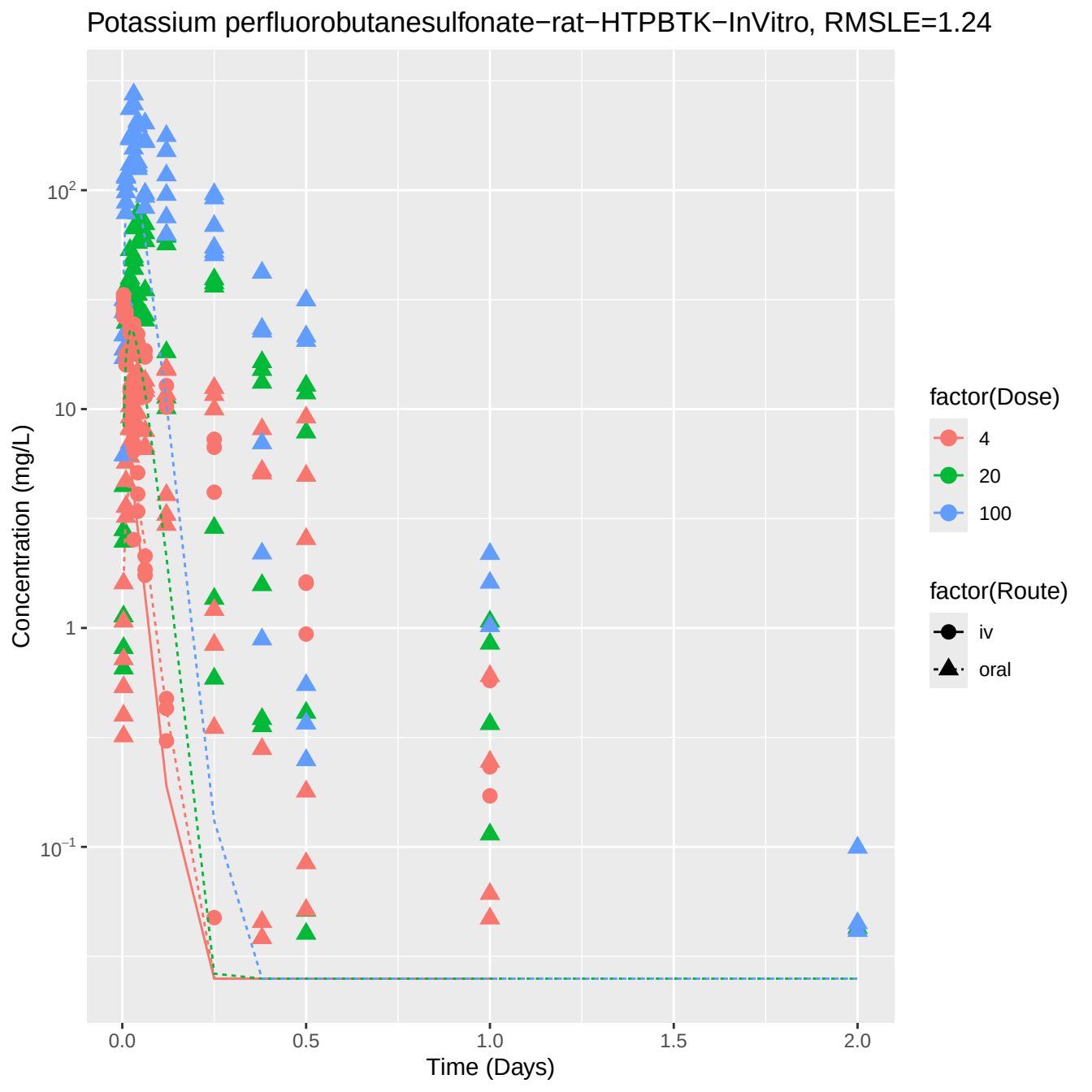

Alprazolam−rat−HTPBTK−InVitro, RMSLE=1.78
Concentration (mg/L)
1
10^{-1}
10^{-2}
0.00
0.05
0.10
0.15
0.20
0.25
Time (Days)
factor(Route)
oral
factor(Dose)
7

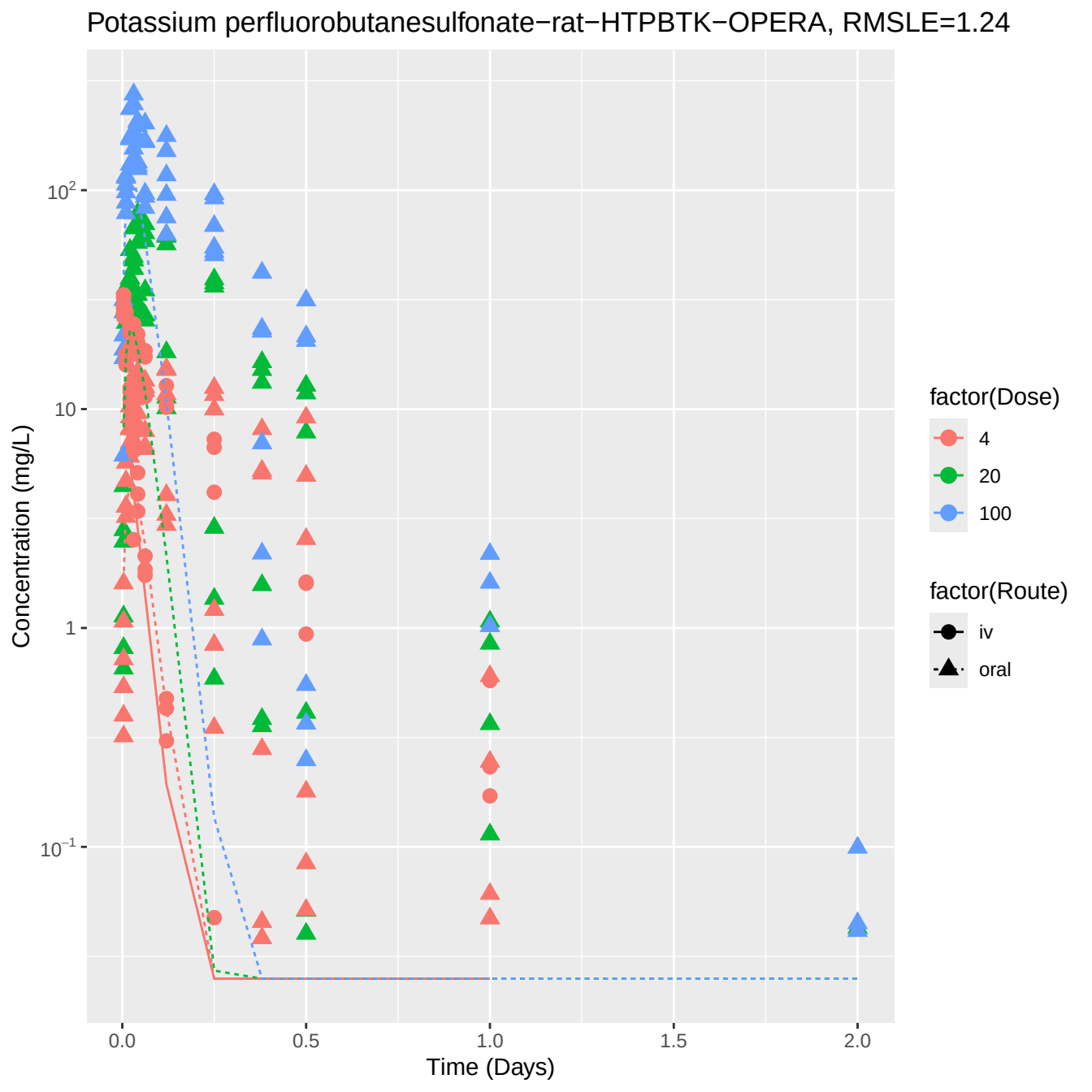

Alprazolam−rat−HTPBTK−ADMET, RMSLE=0.955
Concentration (mg/L)
1
10^{-1}
10^{-2}
0.00
0.05
0.10
0.15
0.20
0.25
Time (Days)
factor(Route)
oral
factor(Dose)
7

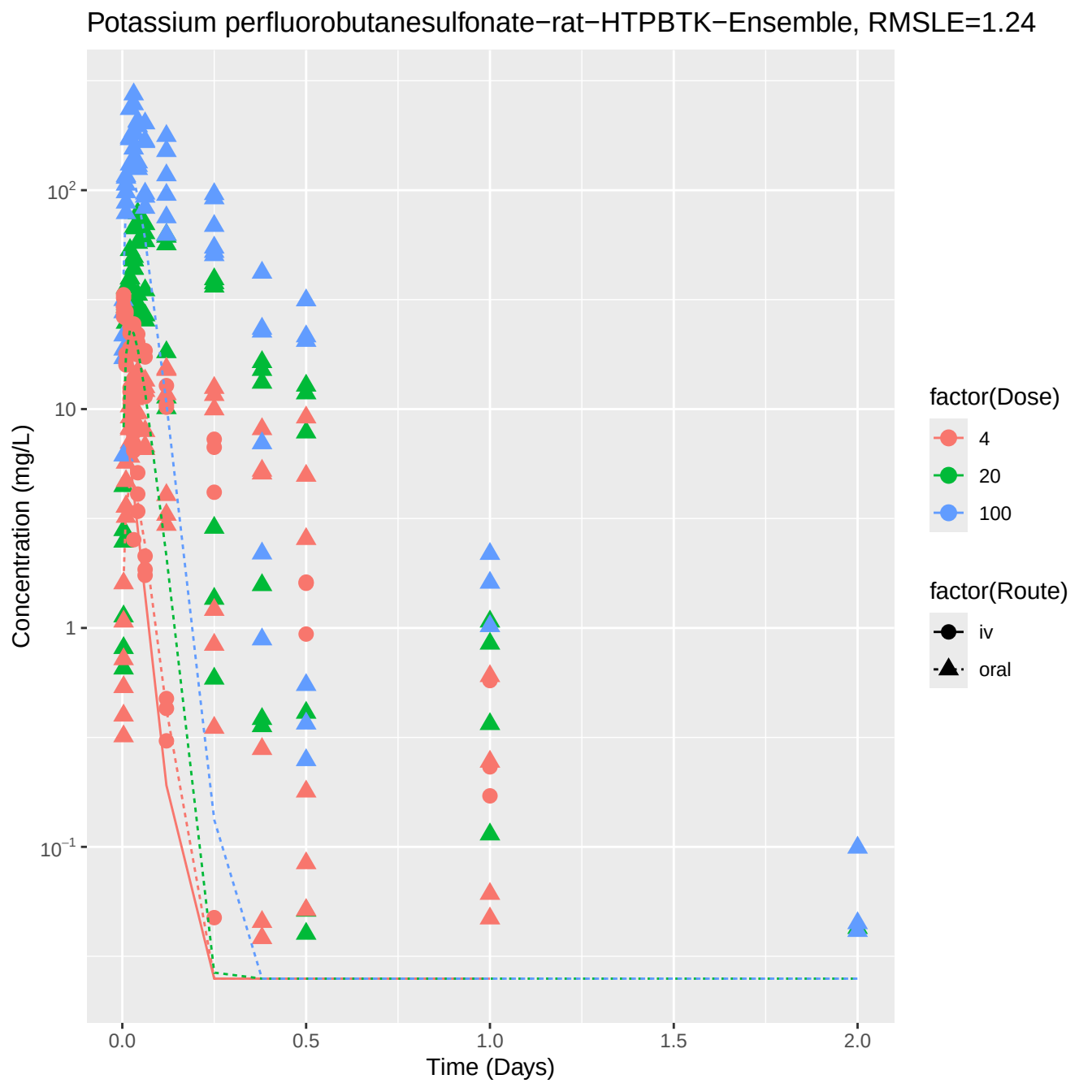

Alprazolam−rat−HTPBTK−Dawson, RMSLE=1.12
Concentration (mg/L)
1
10^{-1}
10^{-2}
0.00
0.05
0.10
0.15
0.20
0.25
Time (Days)
factor(Route)
oral
factor(Dose)
7

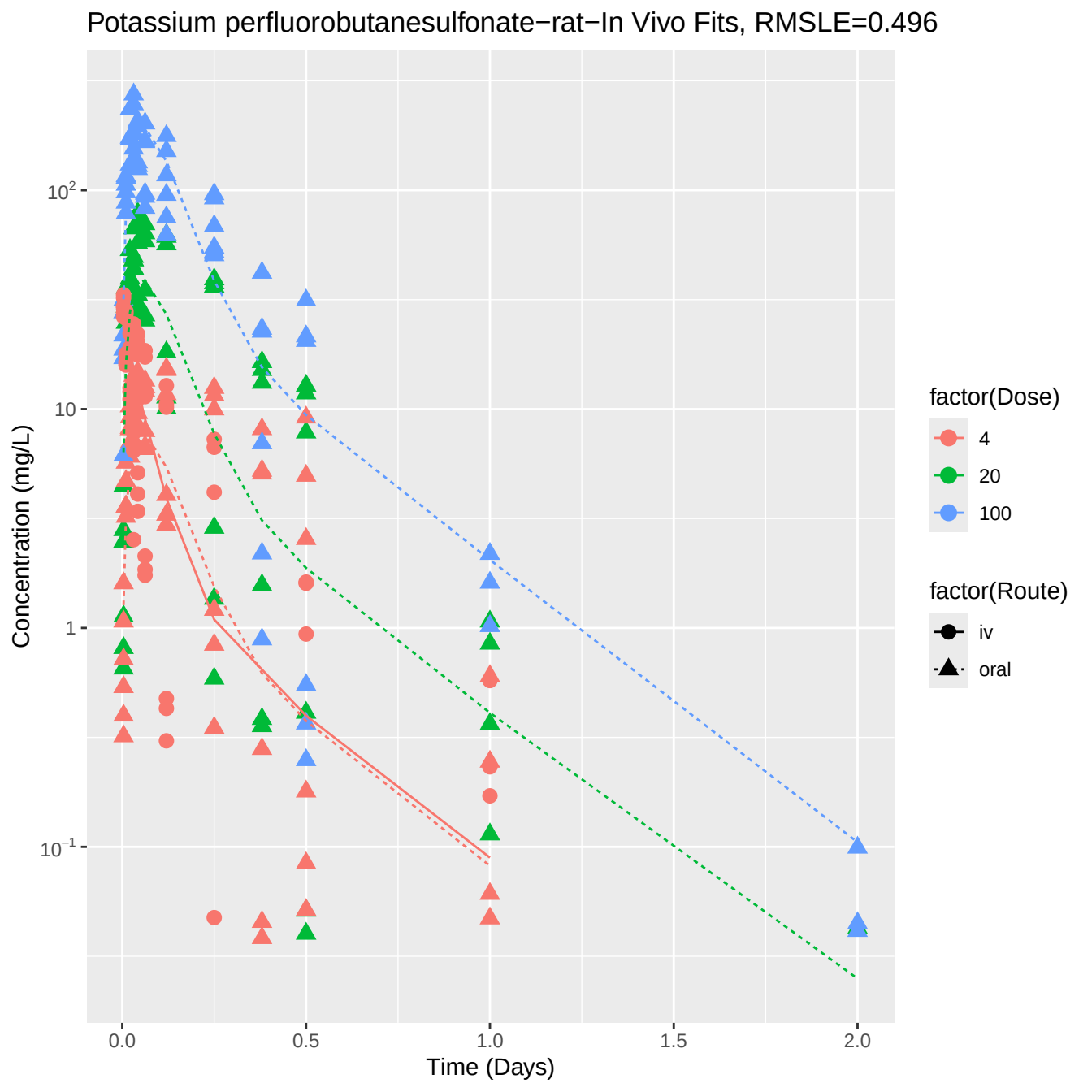

Alprazolam−rat−HTPBTK−Pradeep, RMSLE=1.09
Concentration (mg/L)
1
10^{-1}
10^{-2}
0.00
0.05
0.10
0.15
0.20
0.25
Time (Days)
factor(Route)
oral
factor(Dose)
7

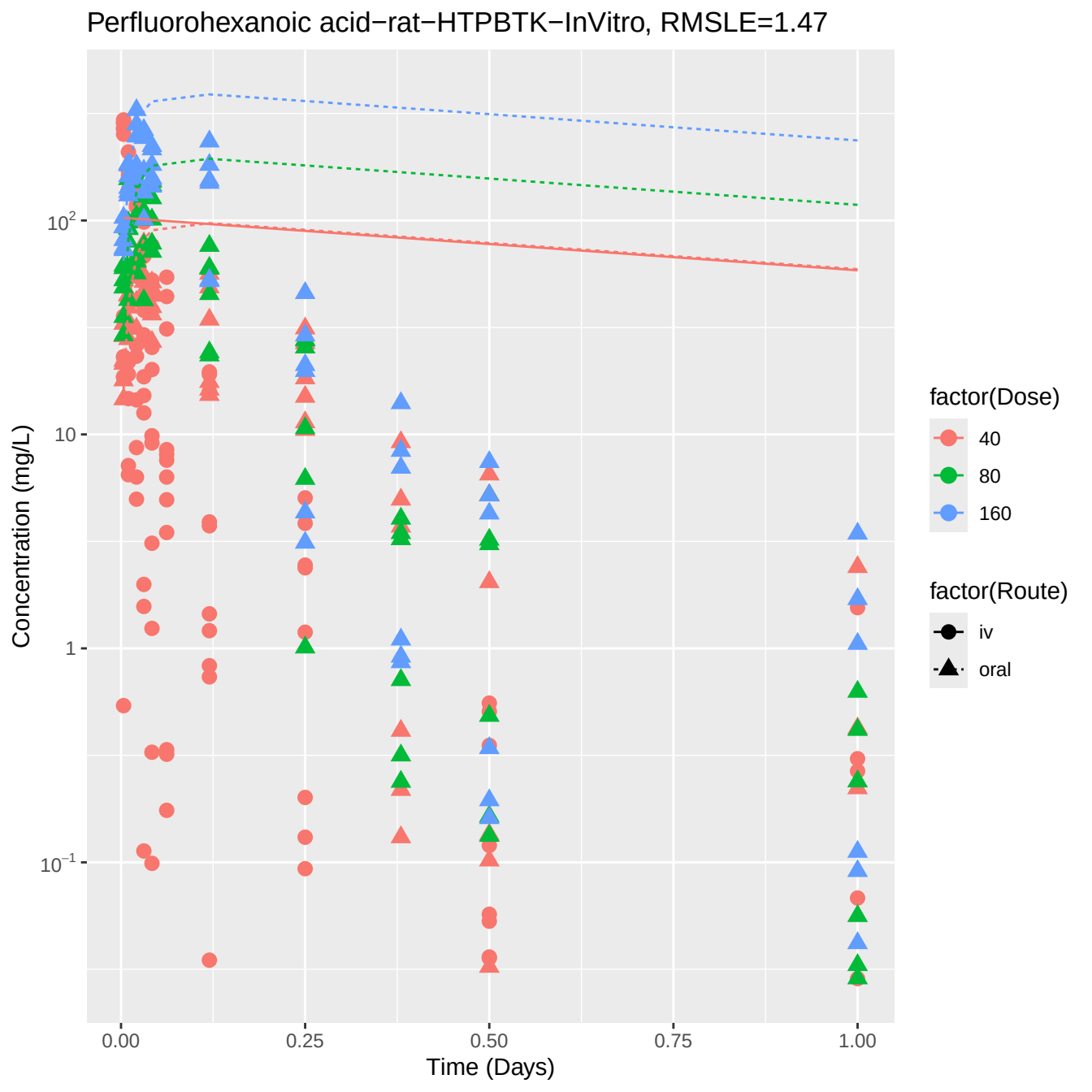

Alprazolam−rat−HTPBTK−Ensemble, RMSLE=0.493
Concentration (mg/L)
0.3
10^{-1}
0.03
10^{-2}
0.003
0.00
0.05
0.10
0.15
0.20
0.25
Time (Days)
factor(Route)
oral
factor(Dose)
7

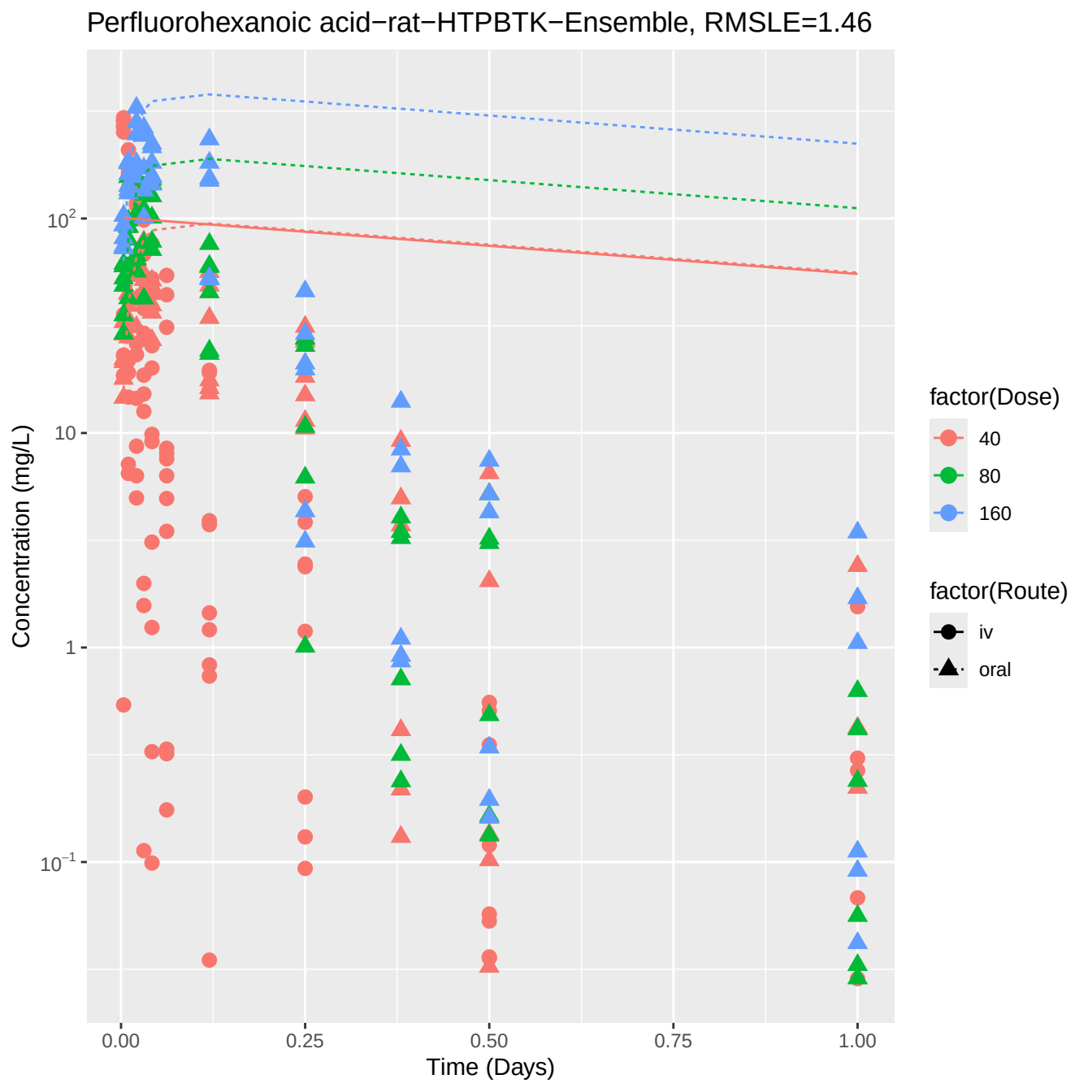

Potassium perfluorobutanesulfonate−rat−HTPBTK−InVitro, RMSLE=1.24
Concentration (mg/L)
10^2
10
1
10^{-1}
0.0
0.5
1.0
1.5
2.0
Time (Days)
factor(Dose)
4
20
100
factor(Route)
iv
oral

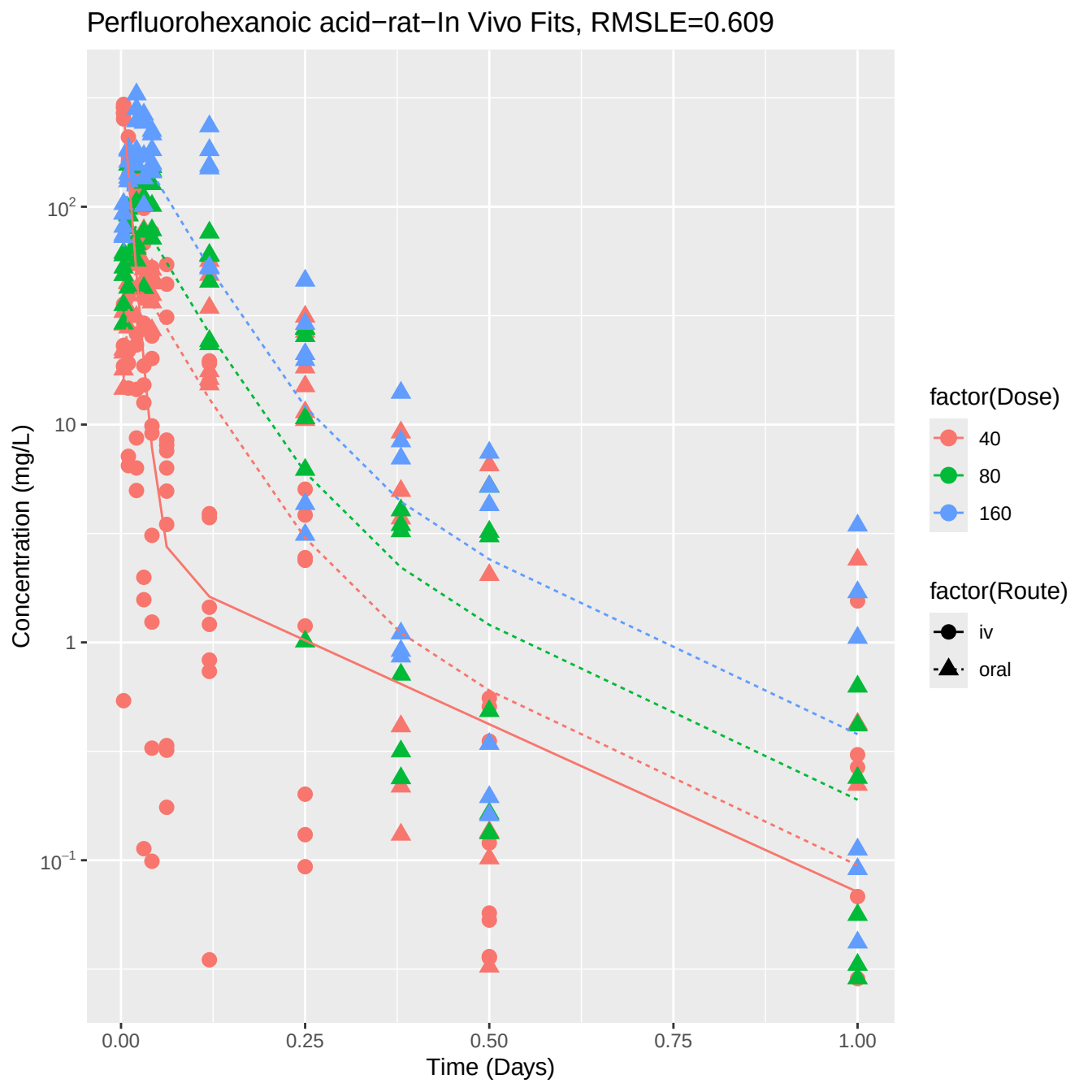

Potassium perfluorobutanesulfonate−rat−HTPBTK−OPERA, RMSLE=1.24
Concentration (mg/L)
10^2
10
1
10^{-1}
0.0
0.5
1.0
1.5
2.0
Time (Days)
factor(Dose)
4
20
100
factor(Route)
iv
oral

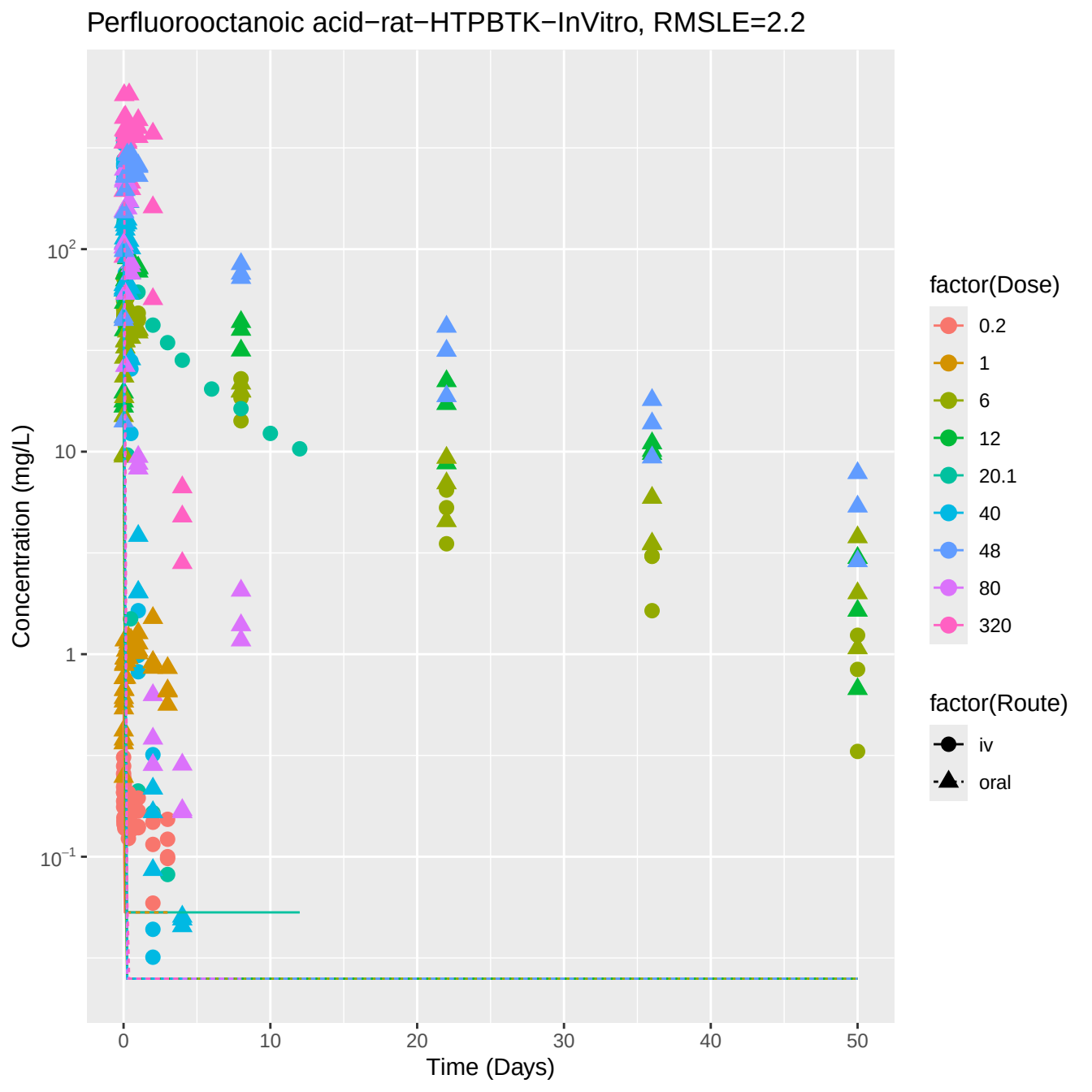

Potassium perfluorobutanesulfonate−rat−HTPBTK−Ensemble, RMSLE=1.24
Concentration (mg/L)
10^2
10
1
10^{-1}
0.0
0.5
1.0
1.5
2.0
Time (Days)
factor(Dose)
4
20
100
factor(Route)
iv
oral

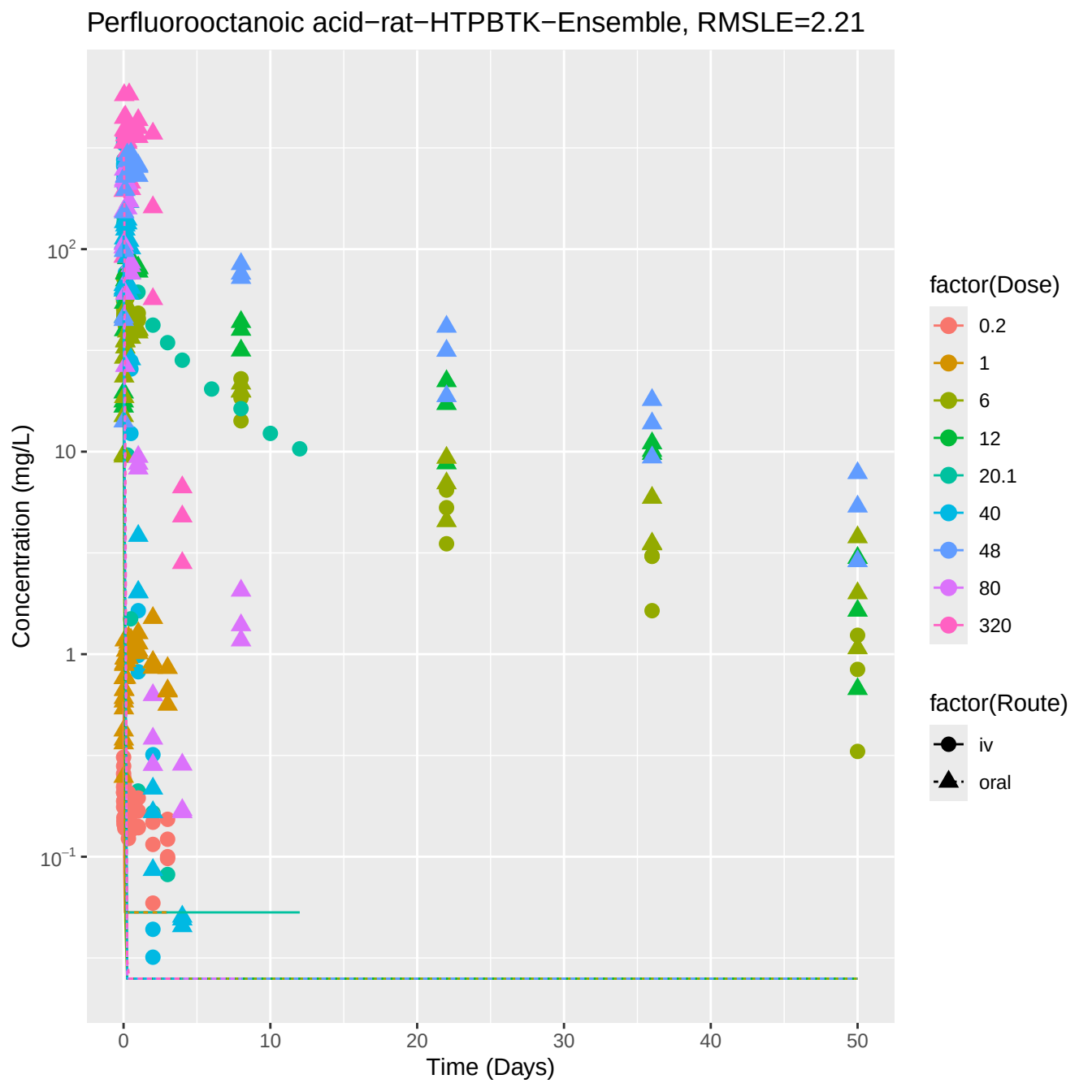

Potassium perfluorobutanesulfonate−rat−In Vivo Fits, RMSLE=0.496
Concentration (mg/L)
10^2
10
1
10^{-1}
0.0
0.5
1.0
1.5
2.0
Time (Days)
factor(Dose)
4
20
100
factor(Route)
iv
oral

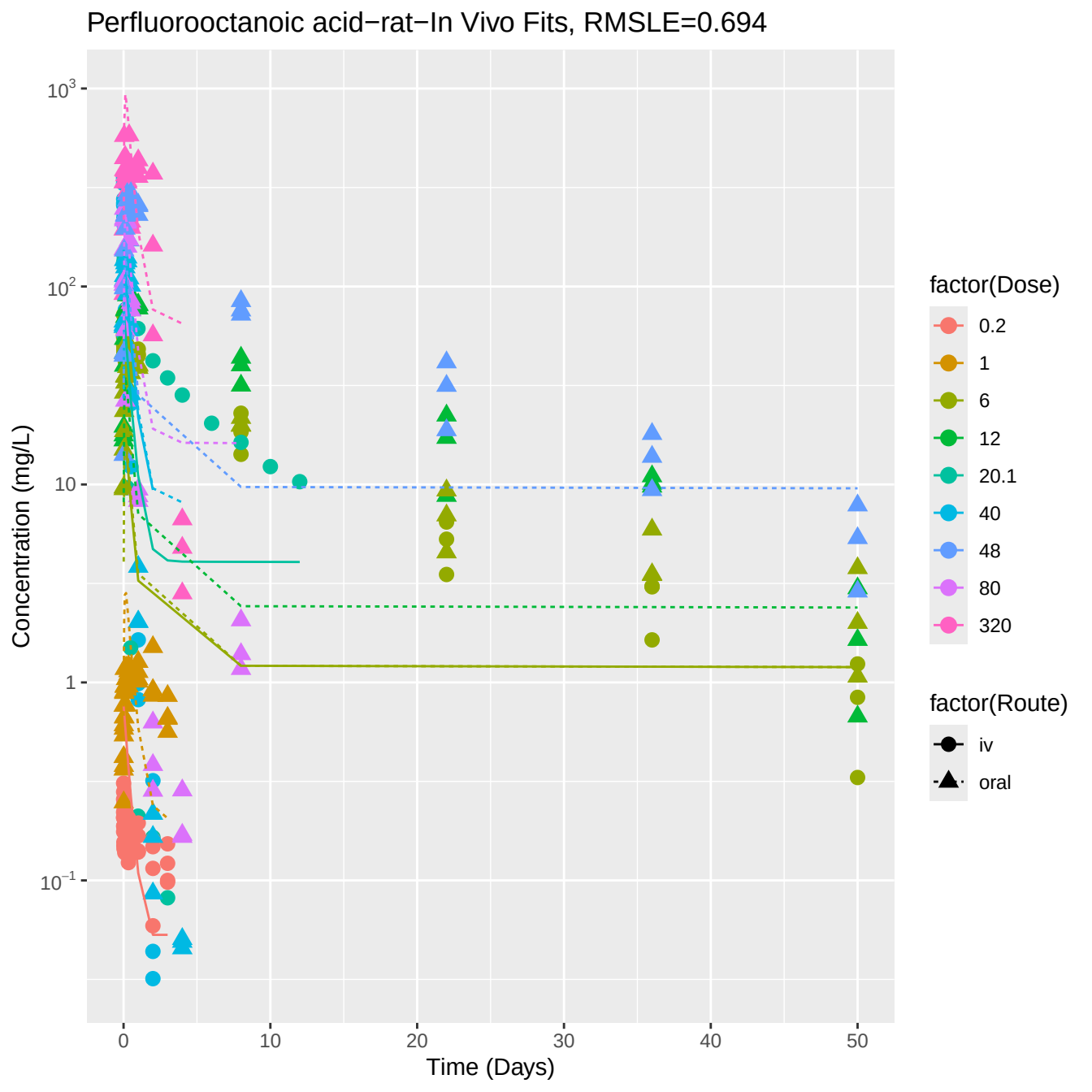

Perfluorohexanoic acid−rat−HTPBTK−InVitro, RMSLE=1.47
Concentration (mg/L)
10^2
10
1
10^{-1}
0.00
0.25
0.50
0.75
1.00
Time (Days)
factor(Dose)
40
80
160
factor(Route)
iv
oral

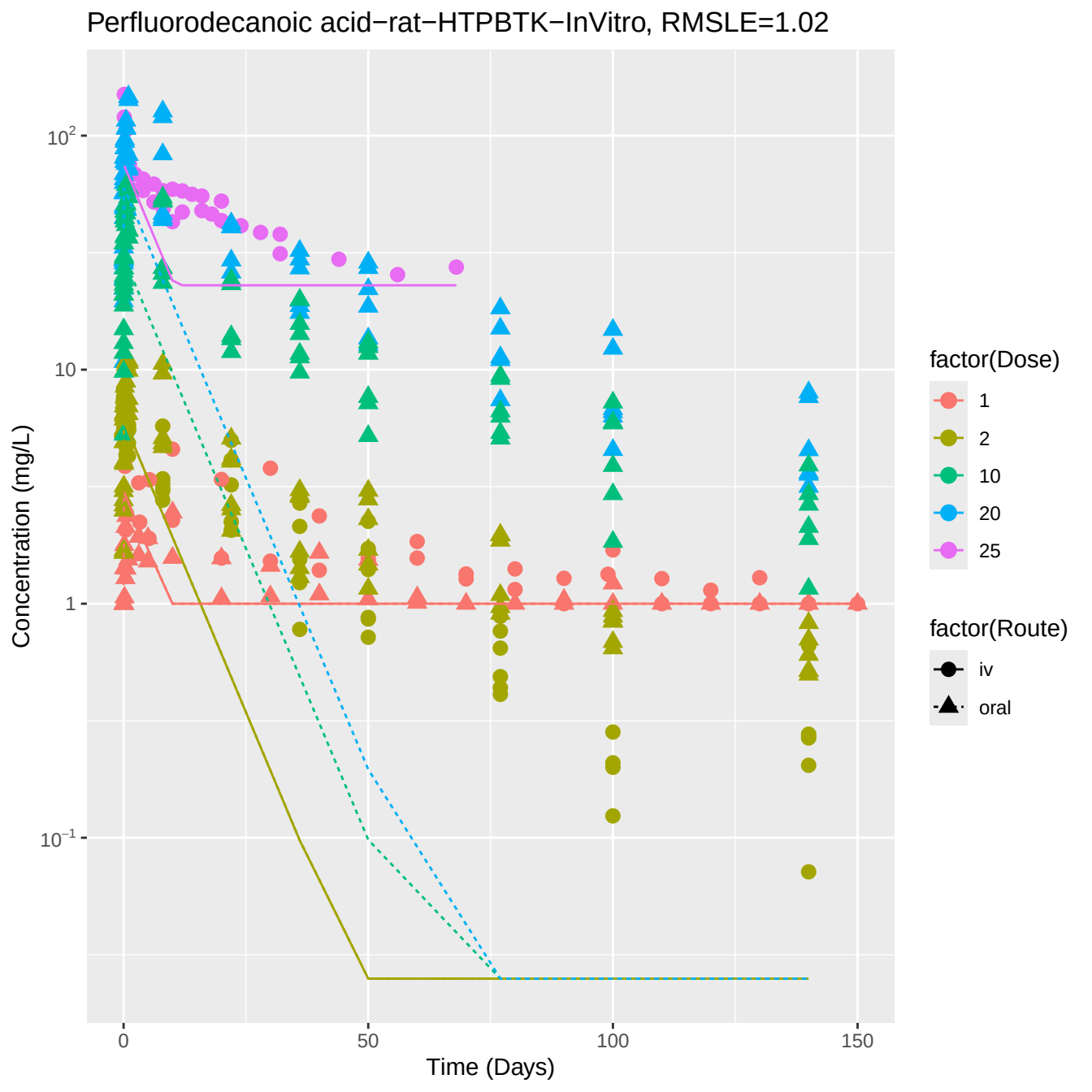

Perfluorohexanoic acid−rat−HTPBTK−Ensemble, RMSLE=1.46
Concentration (mg/L)
10^2
10
1
10^{-1}
0.00
0.25
0.50
0.75
1.00
Time (Days)
factor(Dose)
40
80
160
factor(Route)
iv
oral

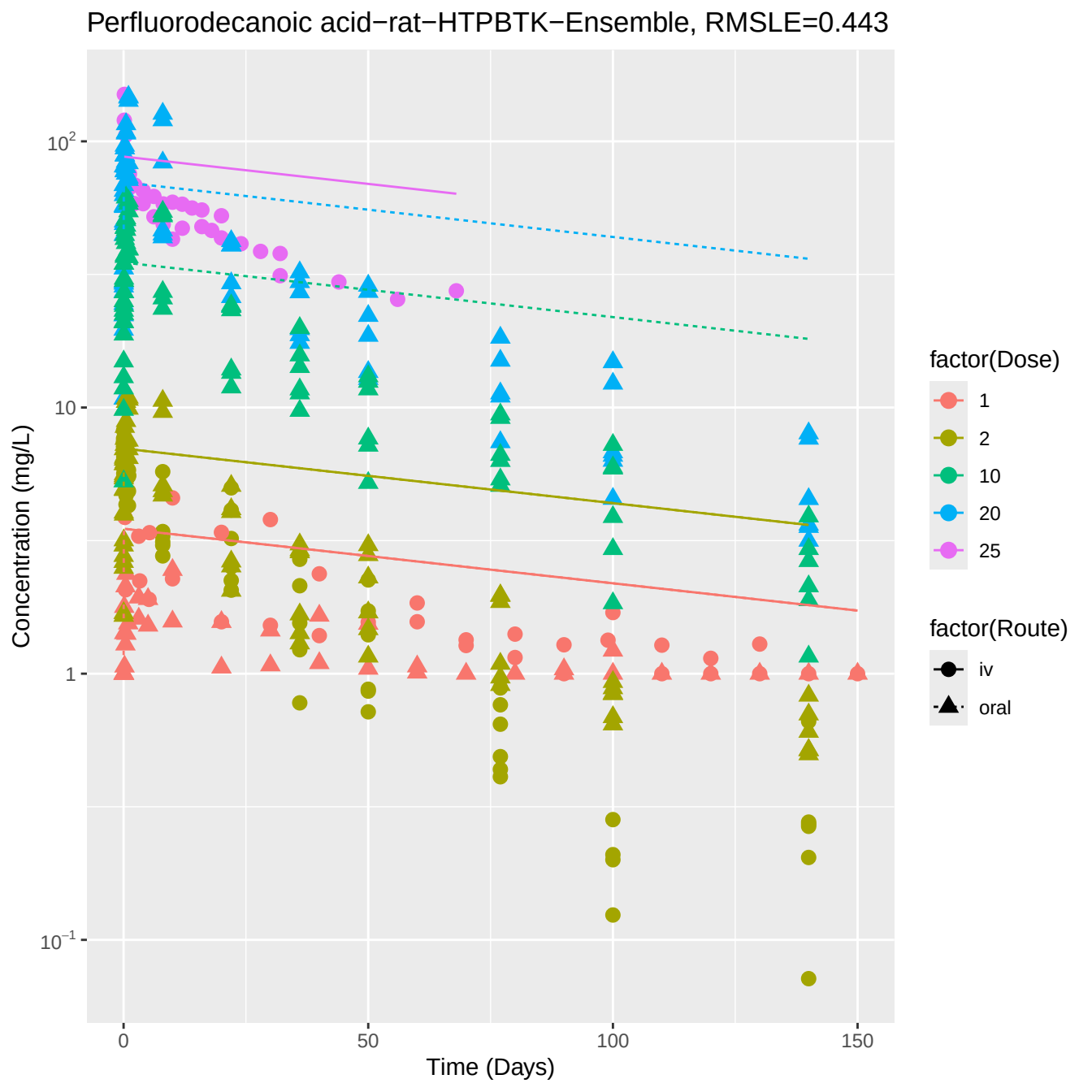

Perfluorohexanoic acid−rat−In Vivo Fits, RMSLE=0.609
Concentration (mg/L)
10^2
10
1
10^{-1}
0.00
0.25
0.50
0.75
1.00
Time (Days)
factor(Dose)
40
80
160
factor(Route)
iv
oral

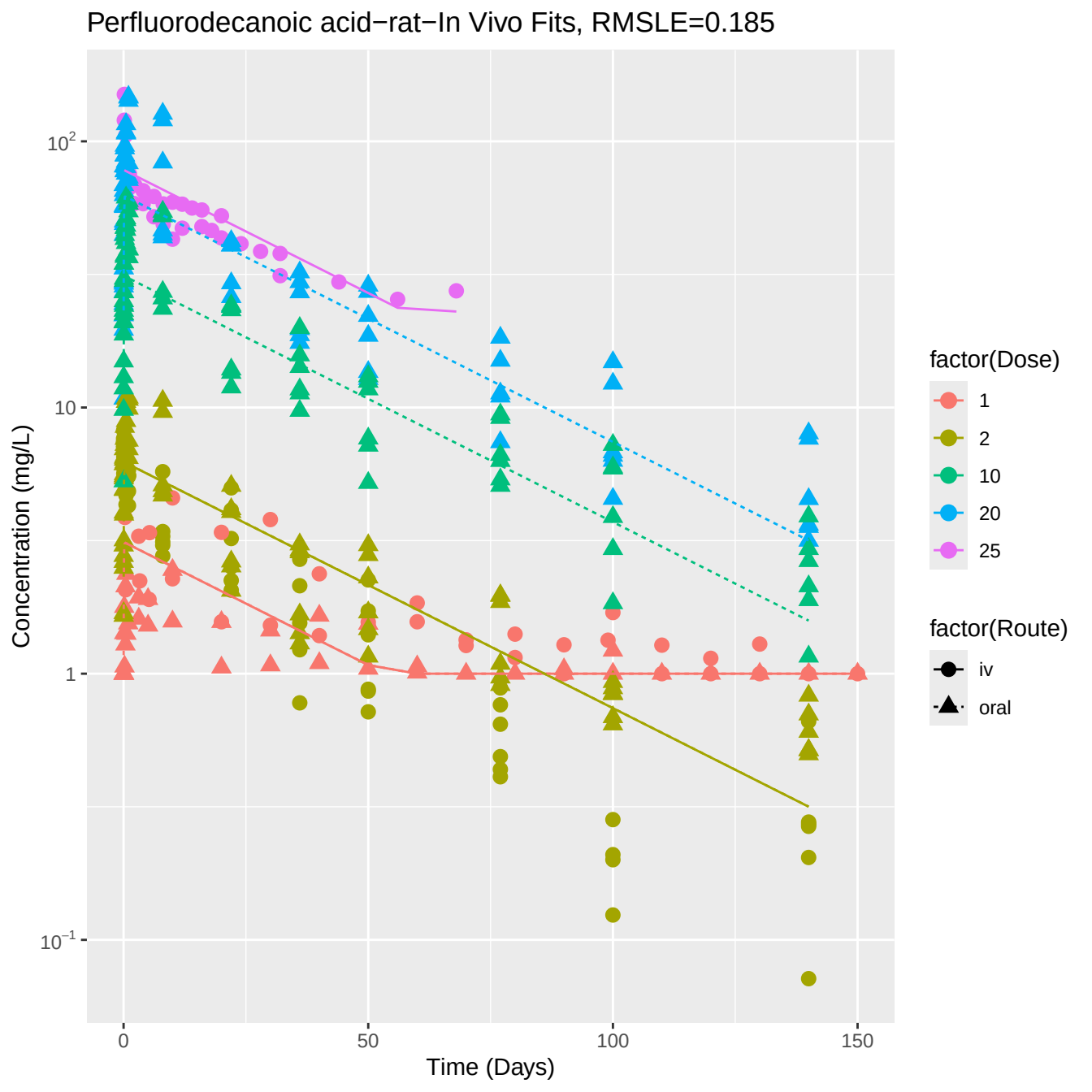

Perfluorooctanoic acid−rat−HTPBTK−InVitro, RMSLE=2.2
Concentration (mg/L)
10^2
10
1
10^{-1}
0
10
20
30
40
50
Time (Days)
factor(Dose)
0.2
1
6
12
20.1
40
48
80
320
factor(Route)
iv
oral

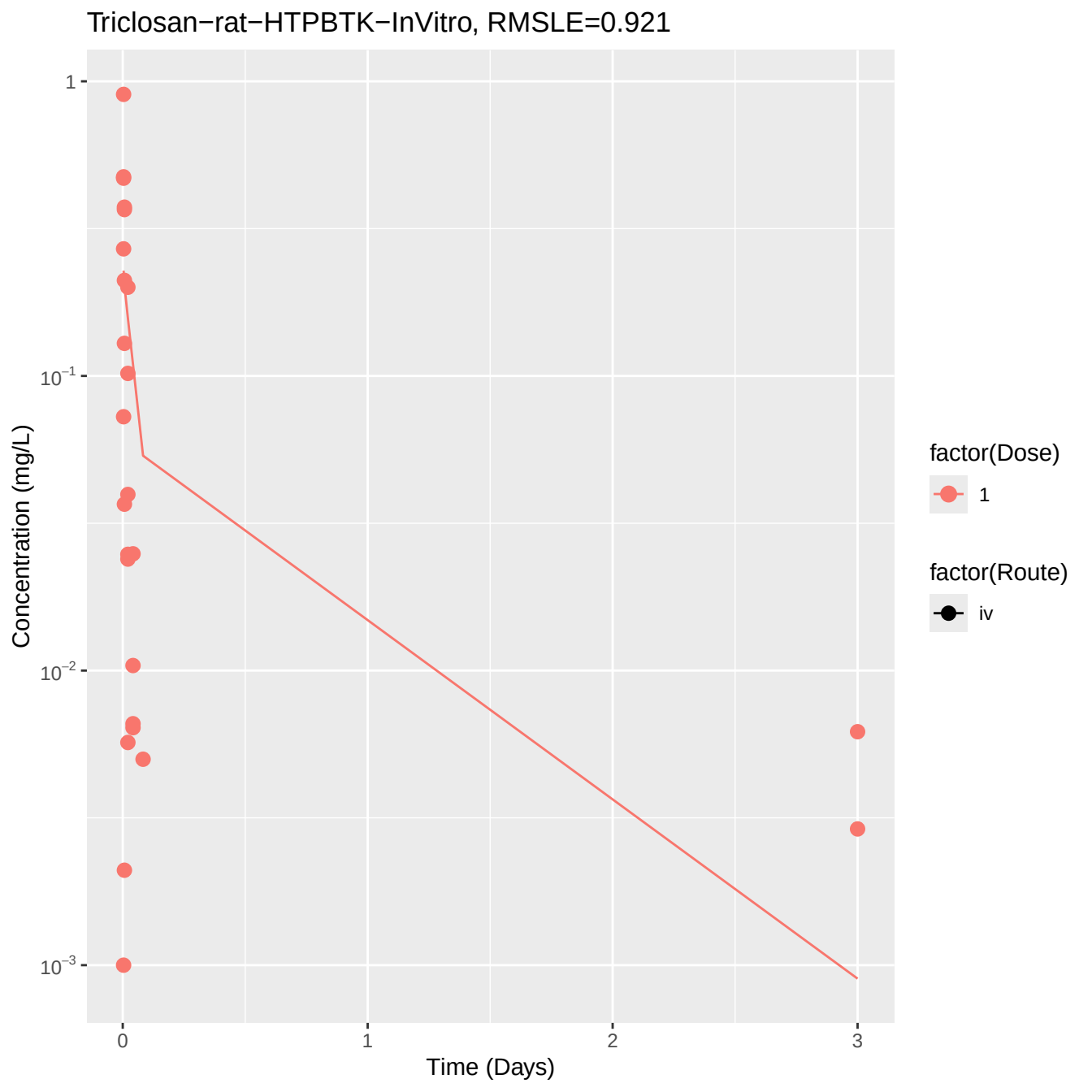

Perfluorooctanoic acid−rat−HTPBTK−Ensemble, RMSLE=2.21
Concentration (mg/L)
10^2
10
1
10^{-1}
0
10
20
30
40
50
Time (Days)
factor(Dose)
0.2
1
6
12
20.1
40
48
80
320
factor(Route)
iv
oral

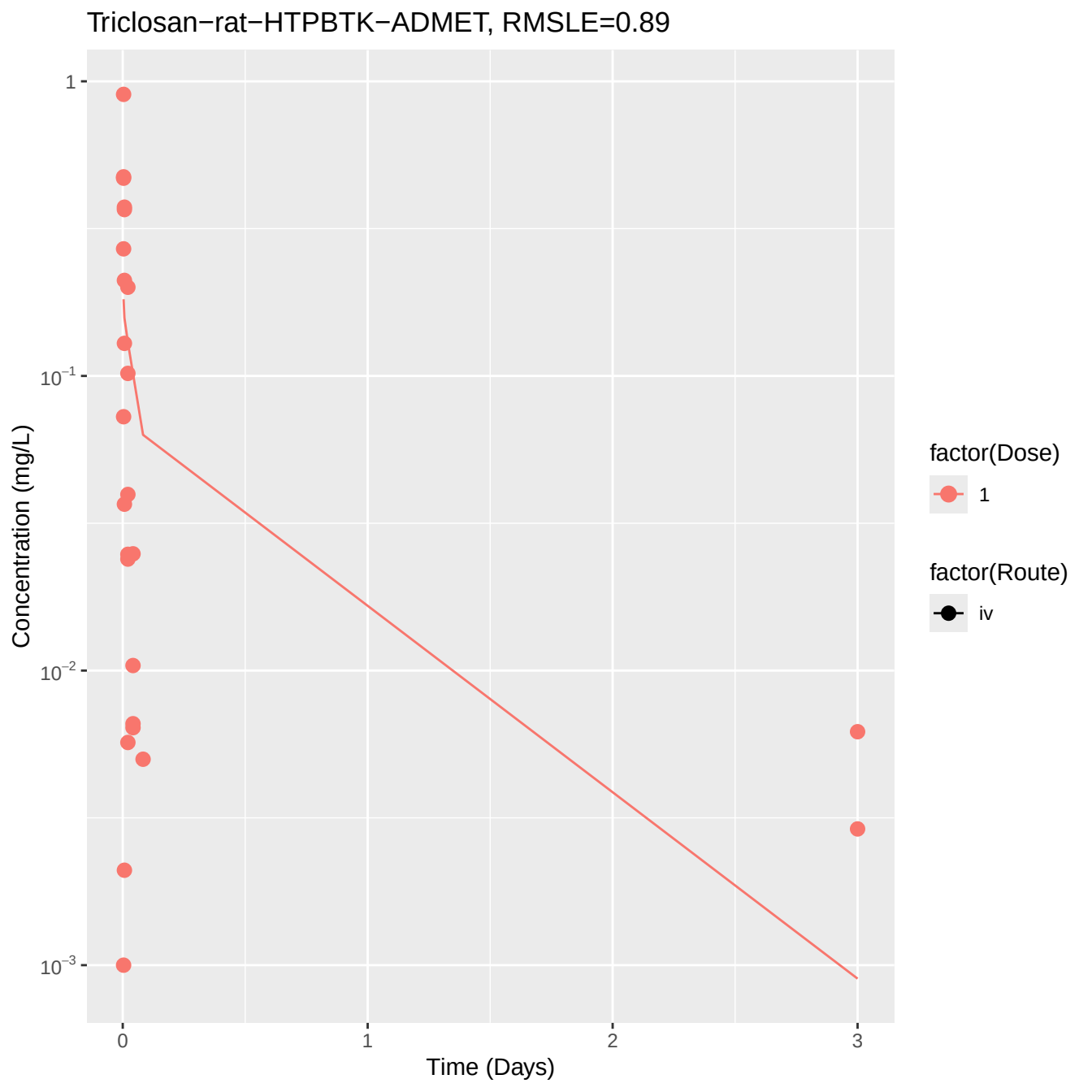

Perfluorooctanoic acid−rat−In Vivo Fits, RMSLE=0.694
Concentration (mg/L)
10^3
10^2
10
1
10^{-1}
0
10
20
30
40
50
Time (Days)
factor(Dose)
0.2
1
6
12
20.1
40
48
80
320
factor(Route)
iv
oral

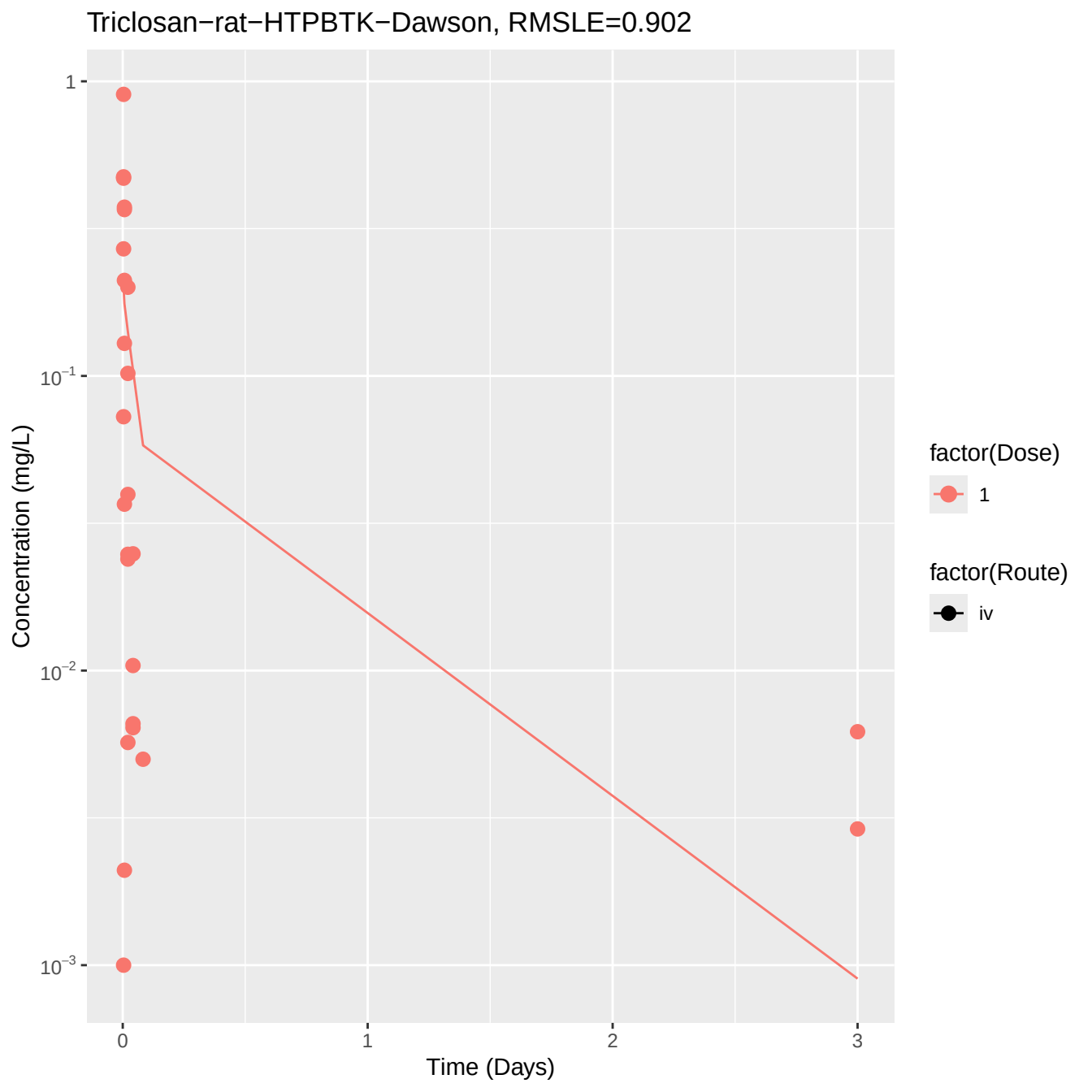

Perfluorodecanoic acid−rat−HTPBTK−InVitro, RMSLE=1.02
Concentration (mg/L)
10^2
10
1
10^{-1}
0
50
100
150
Time (Days)
factor(Dose)
1
2
10
20
25
factor(Route)
iv
oral

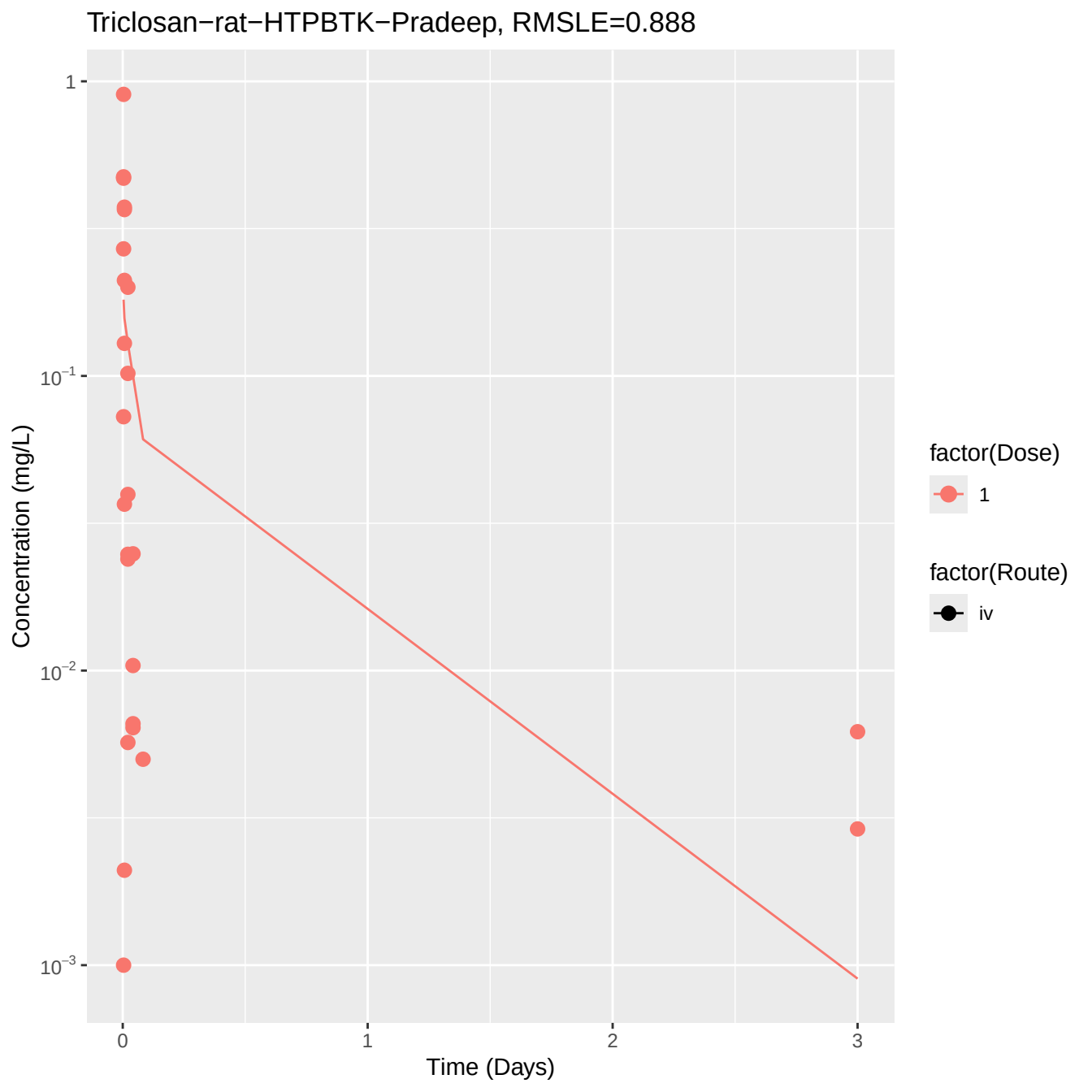

Perfluorodecanoic acid−rat−HTPBTK−Ensemble, RMSLE=0.443
Concentration (mg/L)
10^2
10
1
10^{-1}
0
50
100
150
Time (Days)
factor(Dose)
1
2
10
20
25
factor(Route)
iv
oral

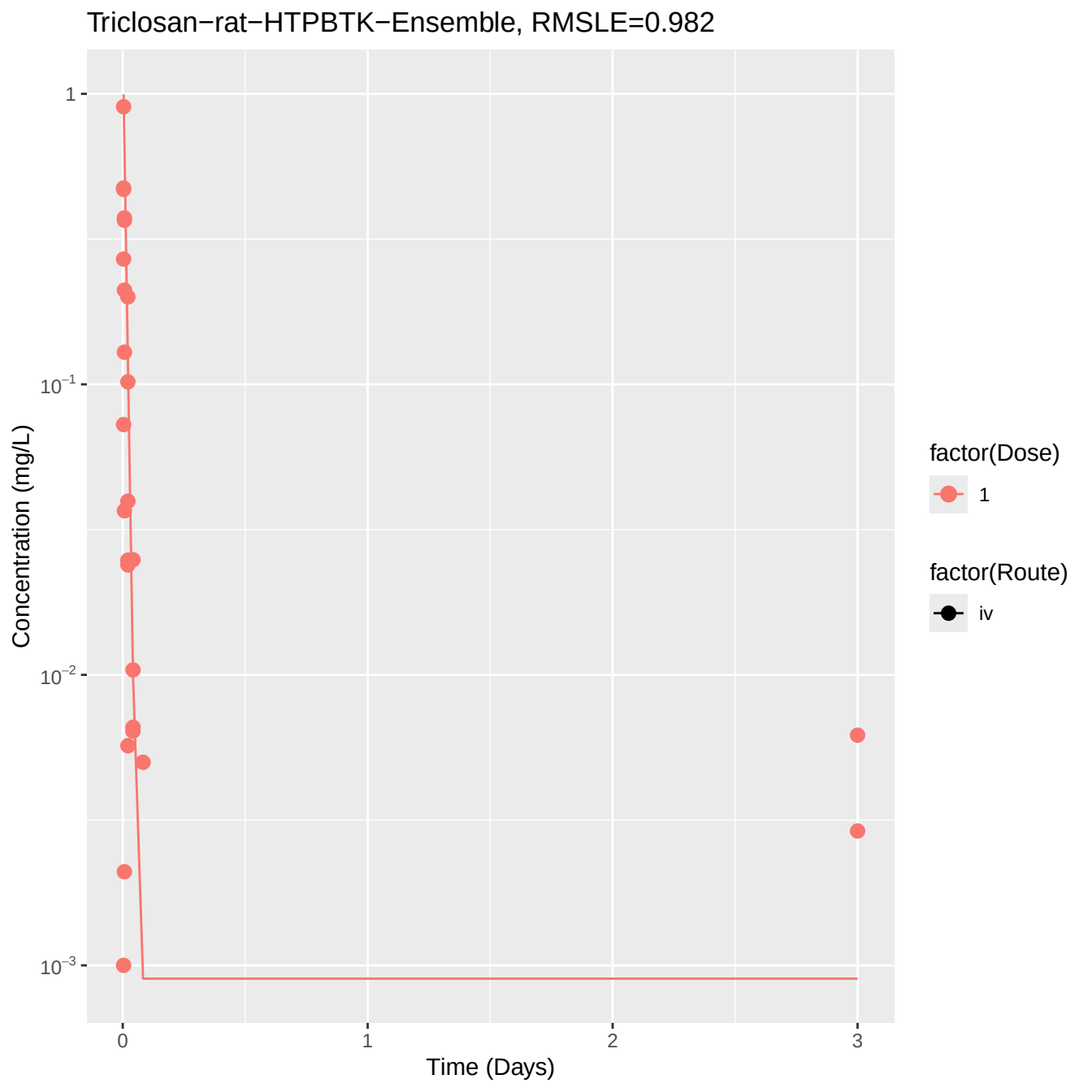

Perfluorodecanoic acid−rat−In Vivo Fits, RMSLE=0.185
Concentration (mg/L)
10^2
10
1
10^{-1}
0
50
100
150
Time (Days)
factor(Dose)
1
2
10
20
25
factor(Route)
iv
oral

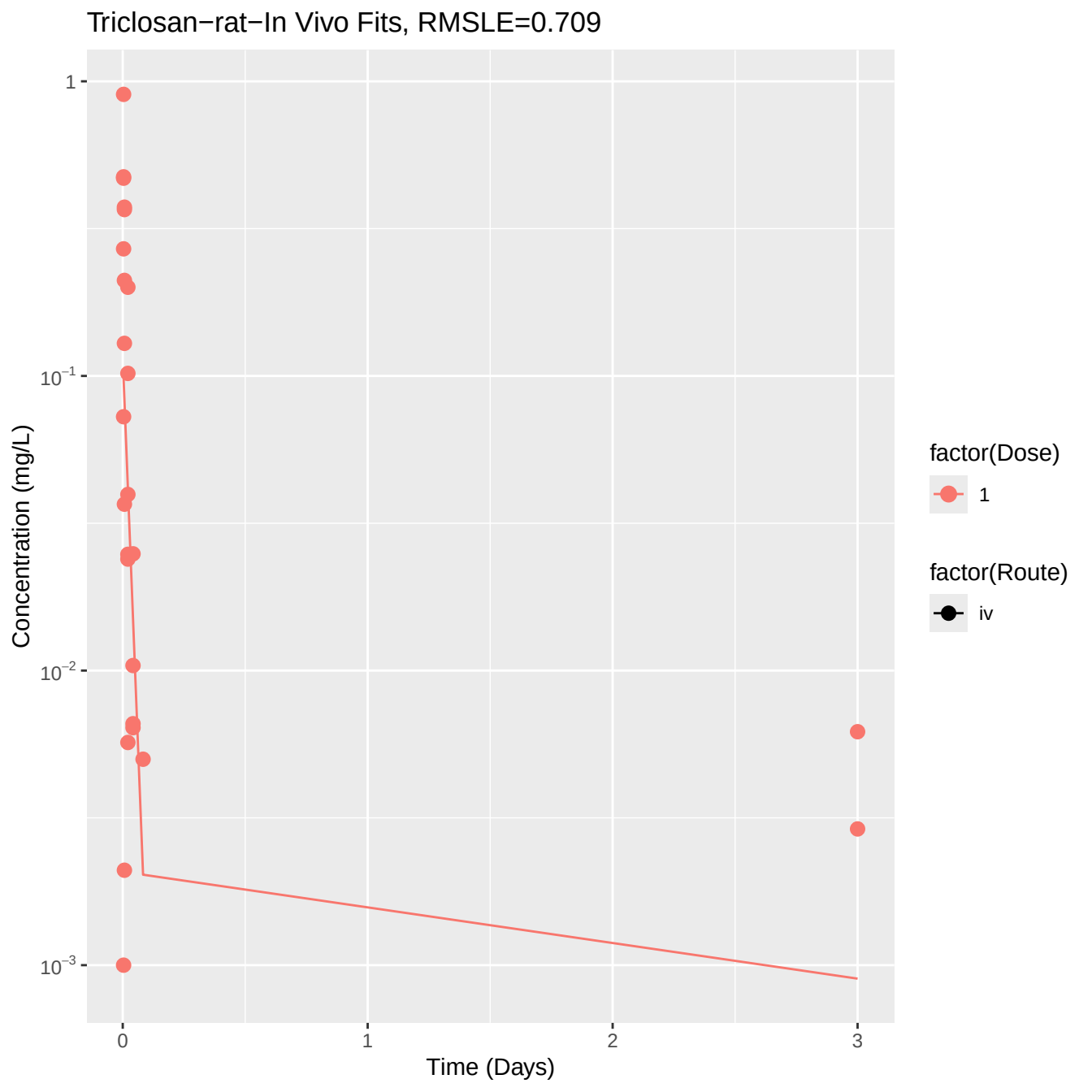

Triclosan−rat−HTPBTK−InVitro, RMSLE=0.921
Concentration (mg/L)
1
10^{-1}
10^{-2}
10^{-3}
0
1
2
3
Time (Days)
factor(Dose)
1
factor(Route)
iv

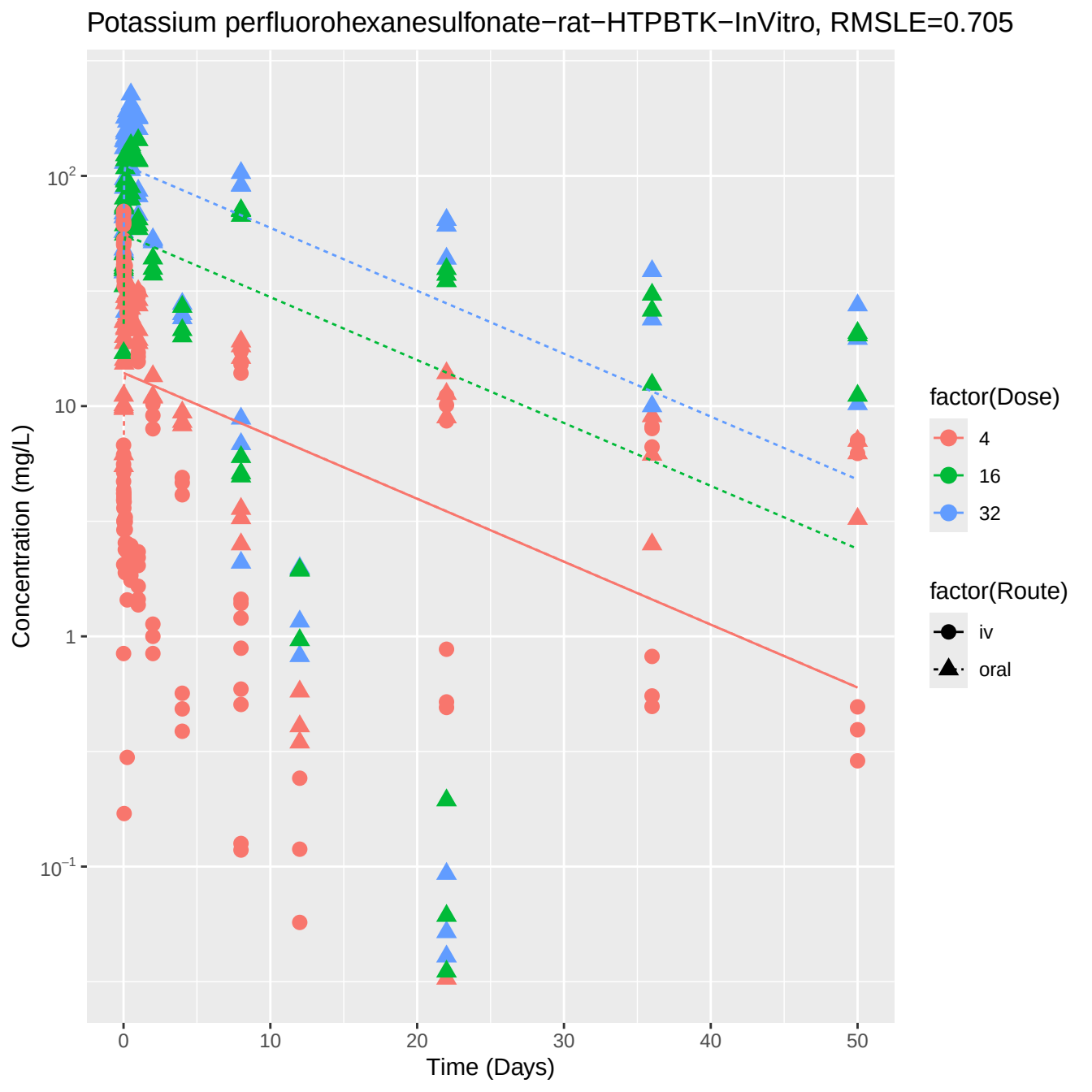

Triclosan−rat−HTPBTK−ADMET, RMSLE=0.89
Concentration (mg/L)
1
10^{-1}
10^{-2}
10^{-3}
0
1
2
3
Time (Days)
factor(Dose)
1
factor(Route)
iv

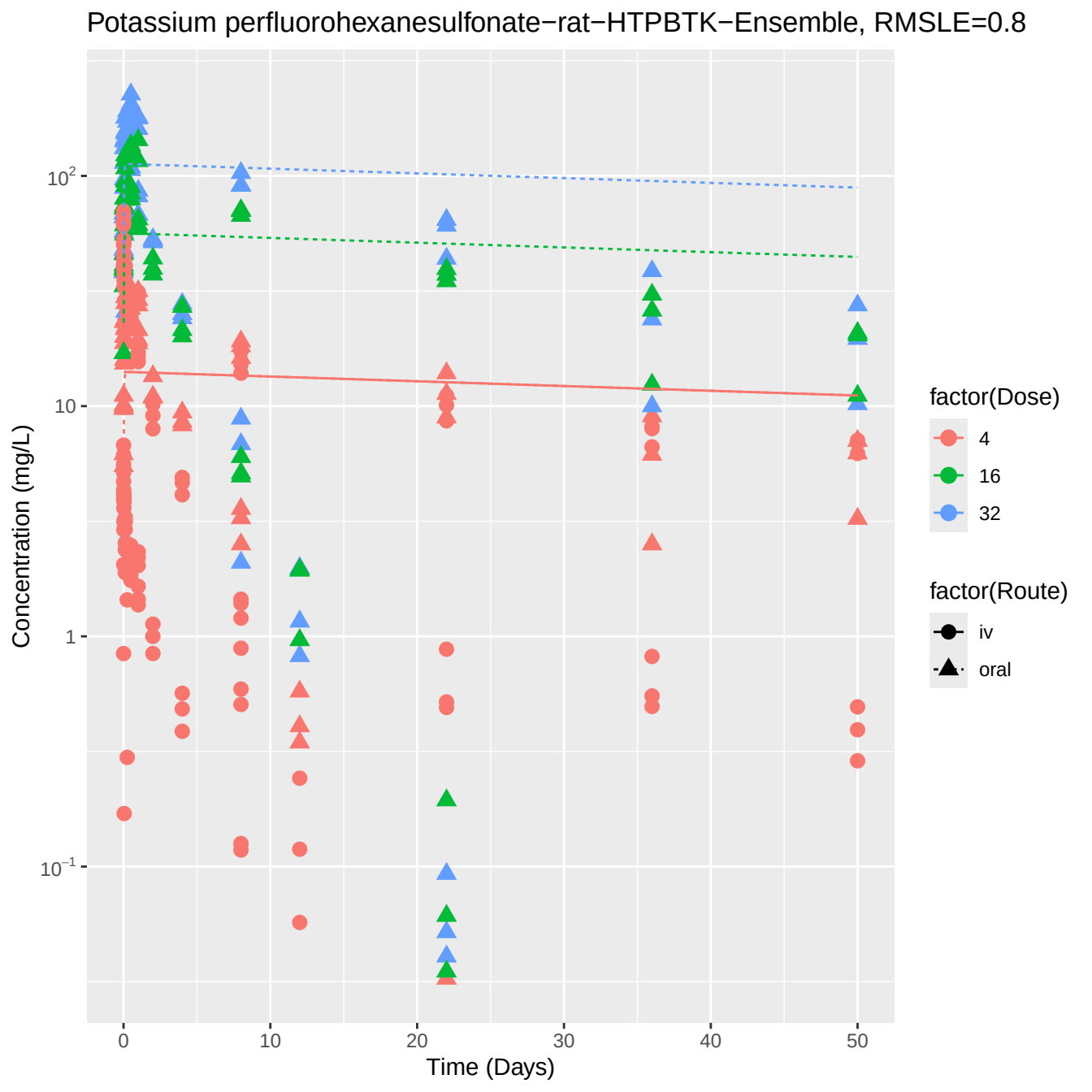

Triclosan−rat−HTPBTK−Dawson, RMSLE=0.902
Concentration (mg/L)
1
10^{-1}
10^{-2}
10^{-3}
0
1
2
3
Time (Days)
factor(Dose)
1
factor(Route)
iv

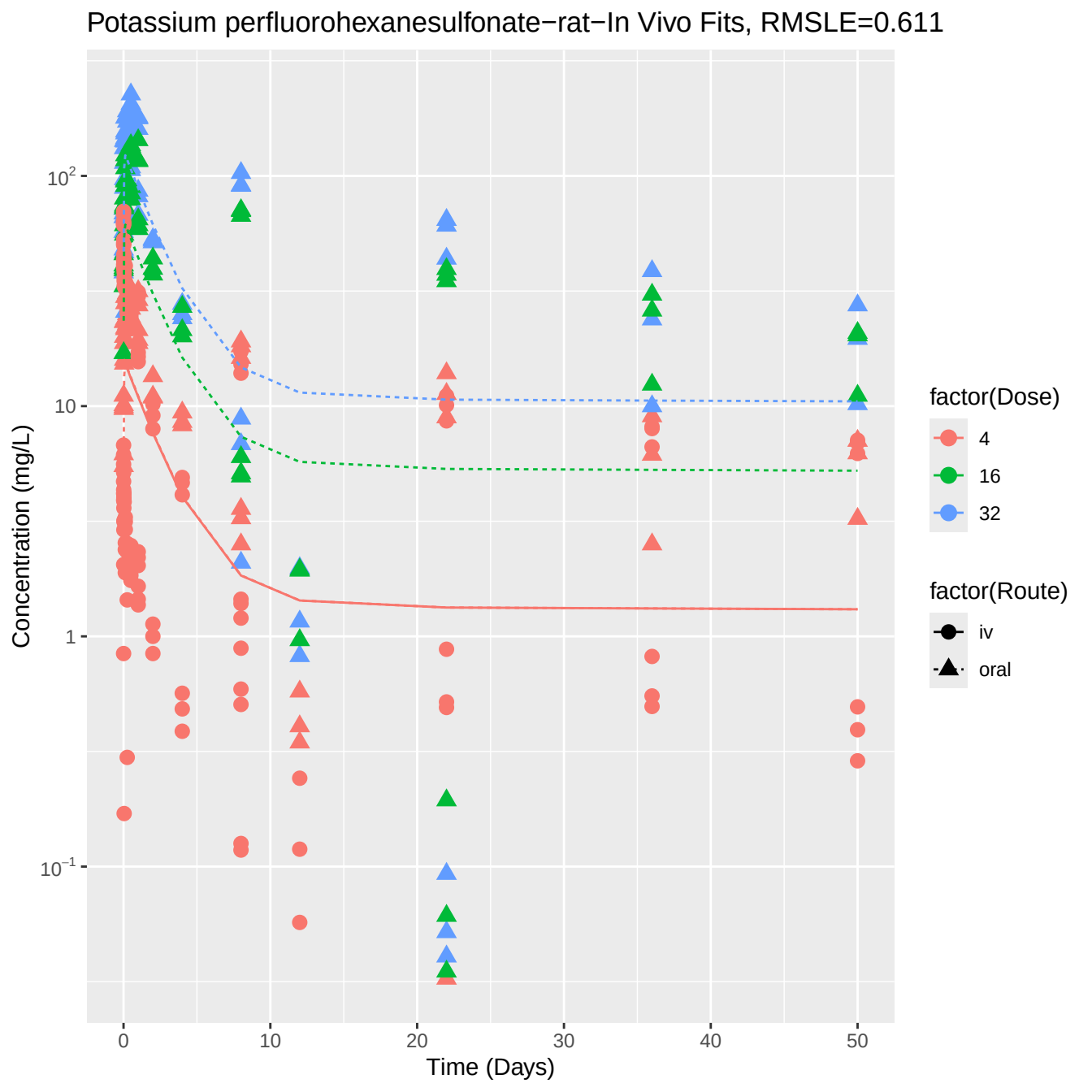

Triclosan−rat−HTPBTK−Pradeep, RMSLE=0.888
Concentration (mg/L)
1
10^{-1}
10^{-2}
10^{-3}
0
1
2
3
Time (Days)
factor(Dose)
1
factor(Route)
iv

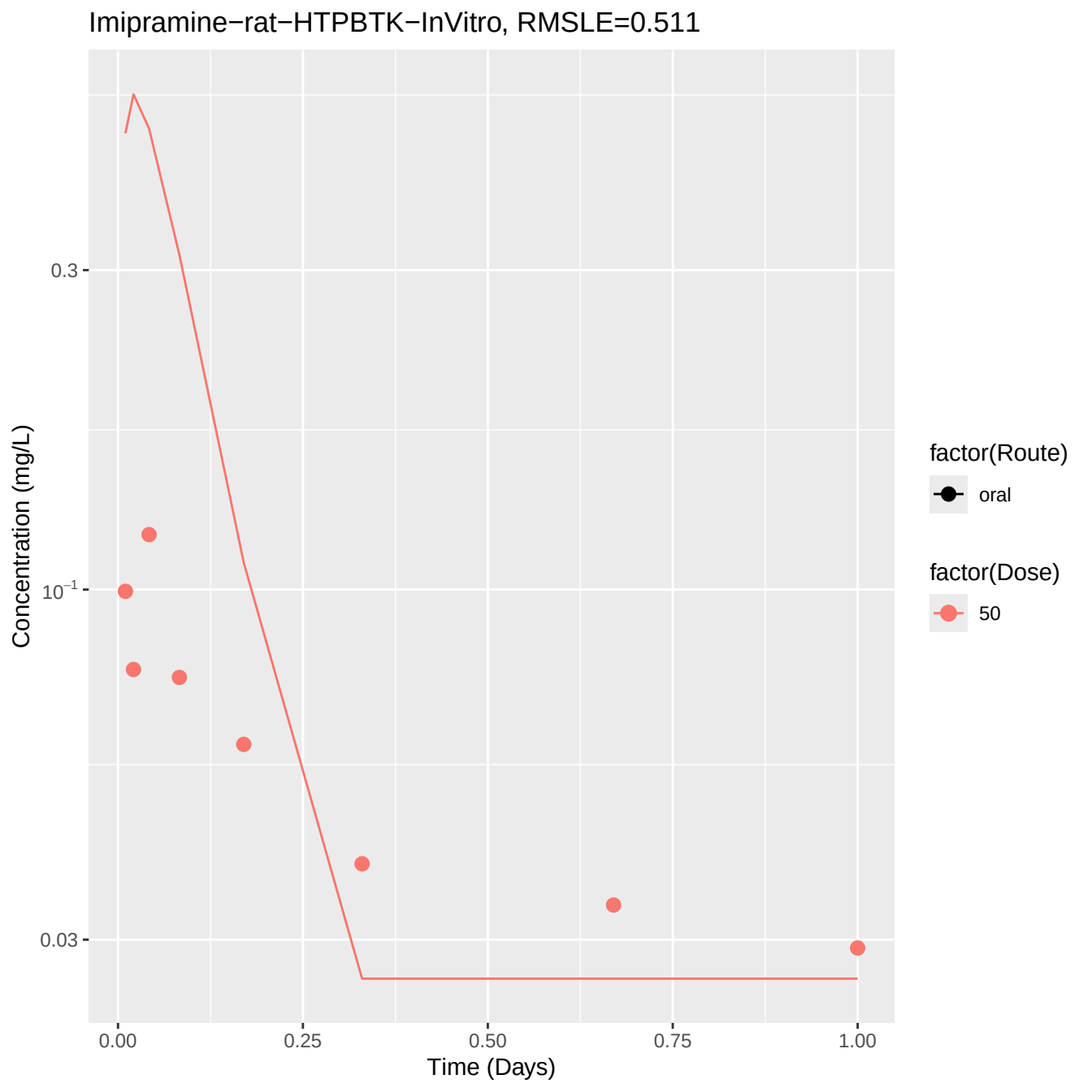

Triclosan−rat−HTPBTK−Ensemble, RMSLE=0.982
Concentration (mg/L)
1
10^{-1}
10^{-2}
10^{-3}
0
1
2
3
Time (Days)
factor(Dose)
1
factor(Route)
iv

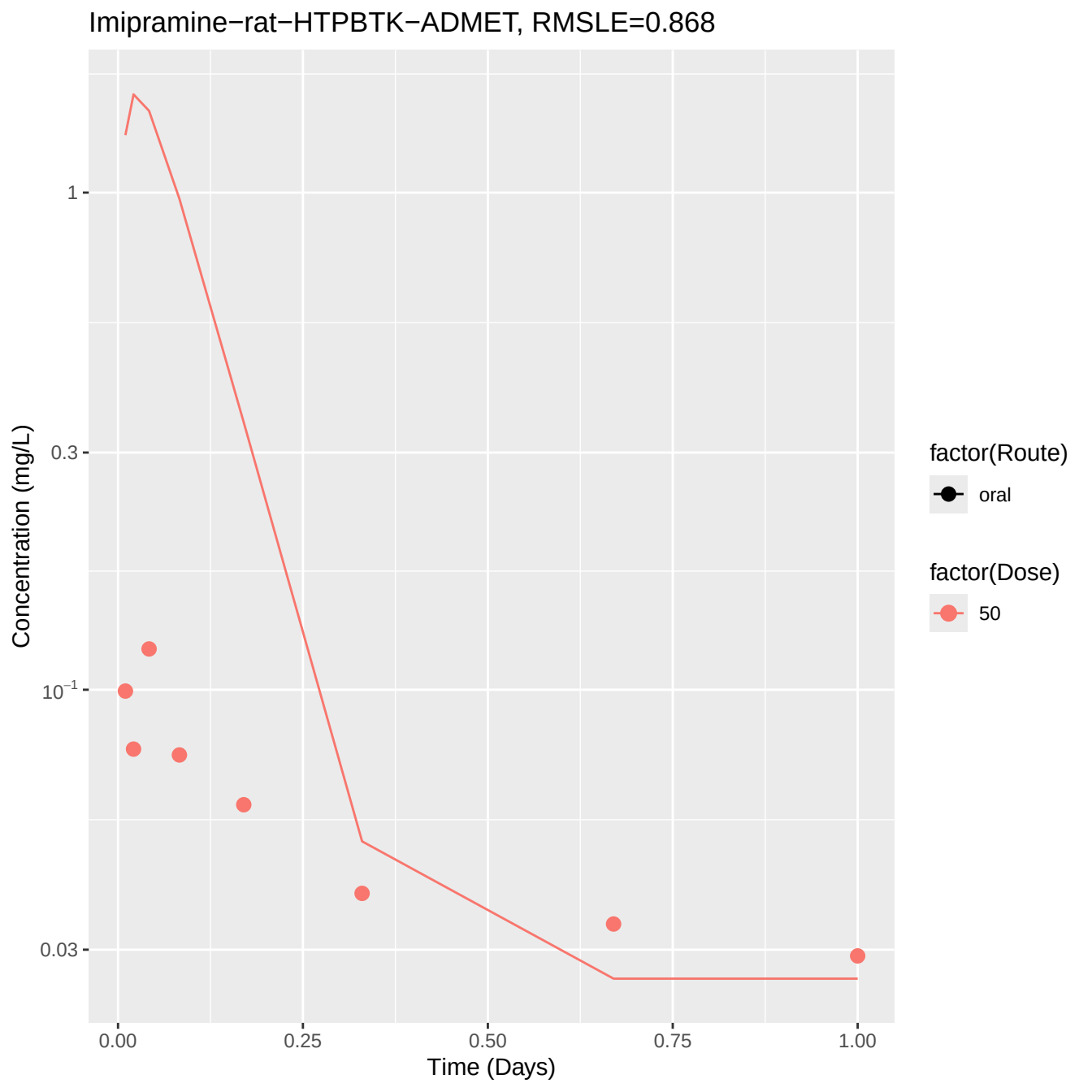

Triclosan−rat−In Vivo Fits, RMSLE=0.709
Concentration (mg/L)
1
10^{-1}
10^{-2}
10^{-3}
0
1
2
3
Time (Days)
factor(Dose)
1
factor(Route)
iv

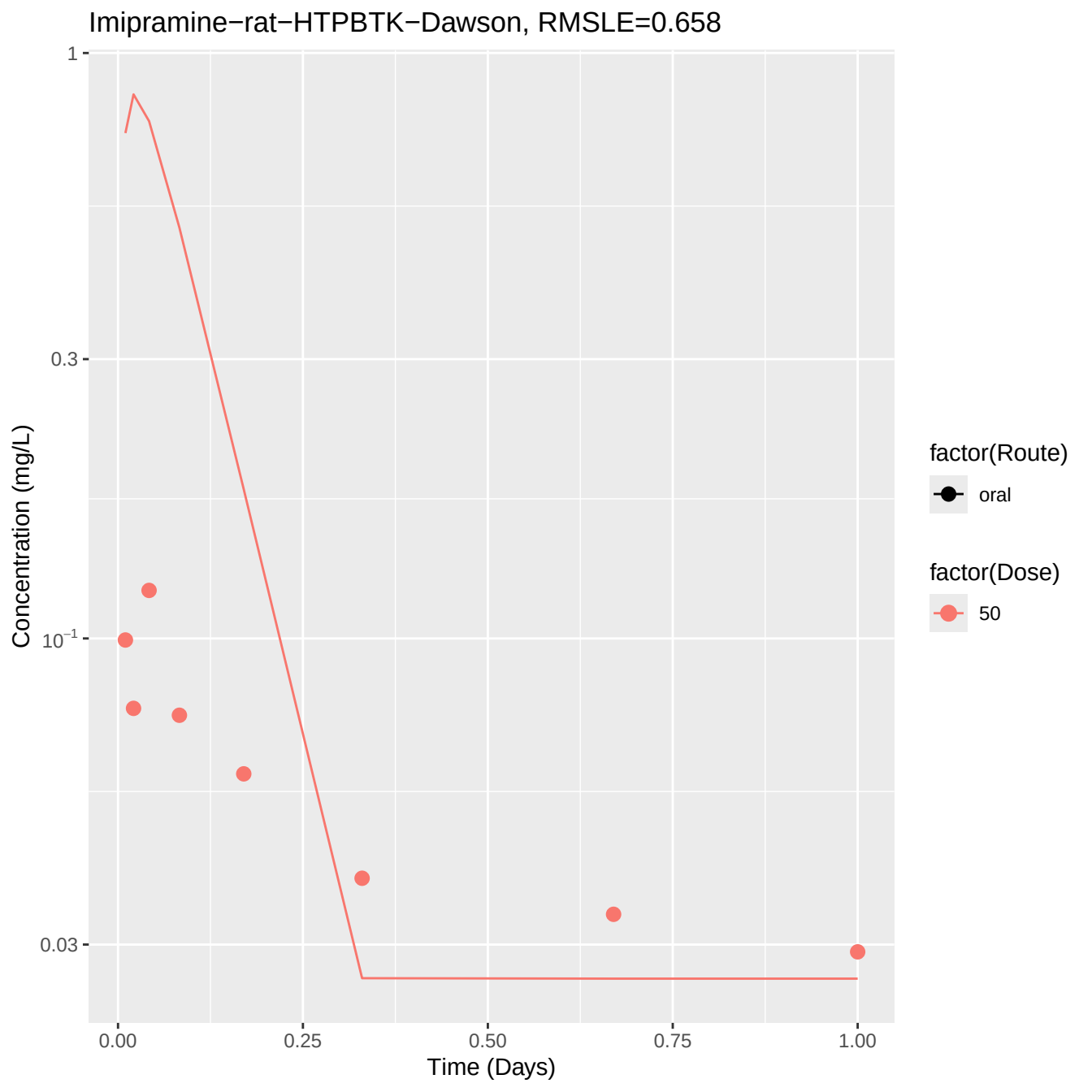

Potassium perfluorohexanesulfonate−rat−HTPBTK−InVitro, RMSLE=0.705
Concentration (mg/L)
10^2
10
1
10^{-1}
0
10
20
30
40
50
Time (Days)
factor(Dose)
4
16
32
factor(Route)
iv
oral

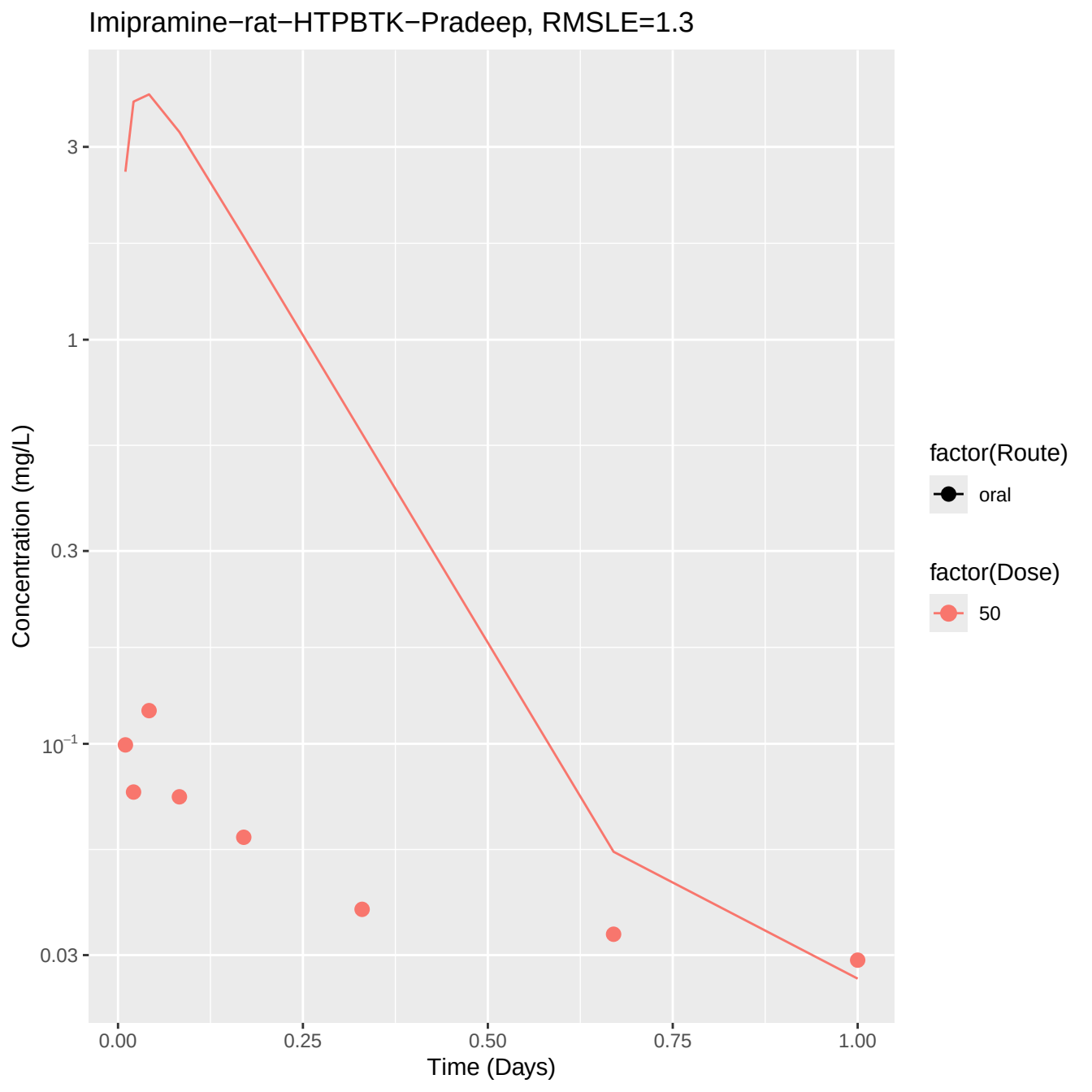

Potassium perfluorohexanesulfonate−rat−HTPBTK−Ensemble, RMSLE=0.8
Concentration (mg/L)
10^2
10
1
10^{-1}
0
10
20
30
40
50
Time (Days)
factor(Dose)
4
16
32
factor(Route)
iv
oral

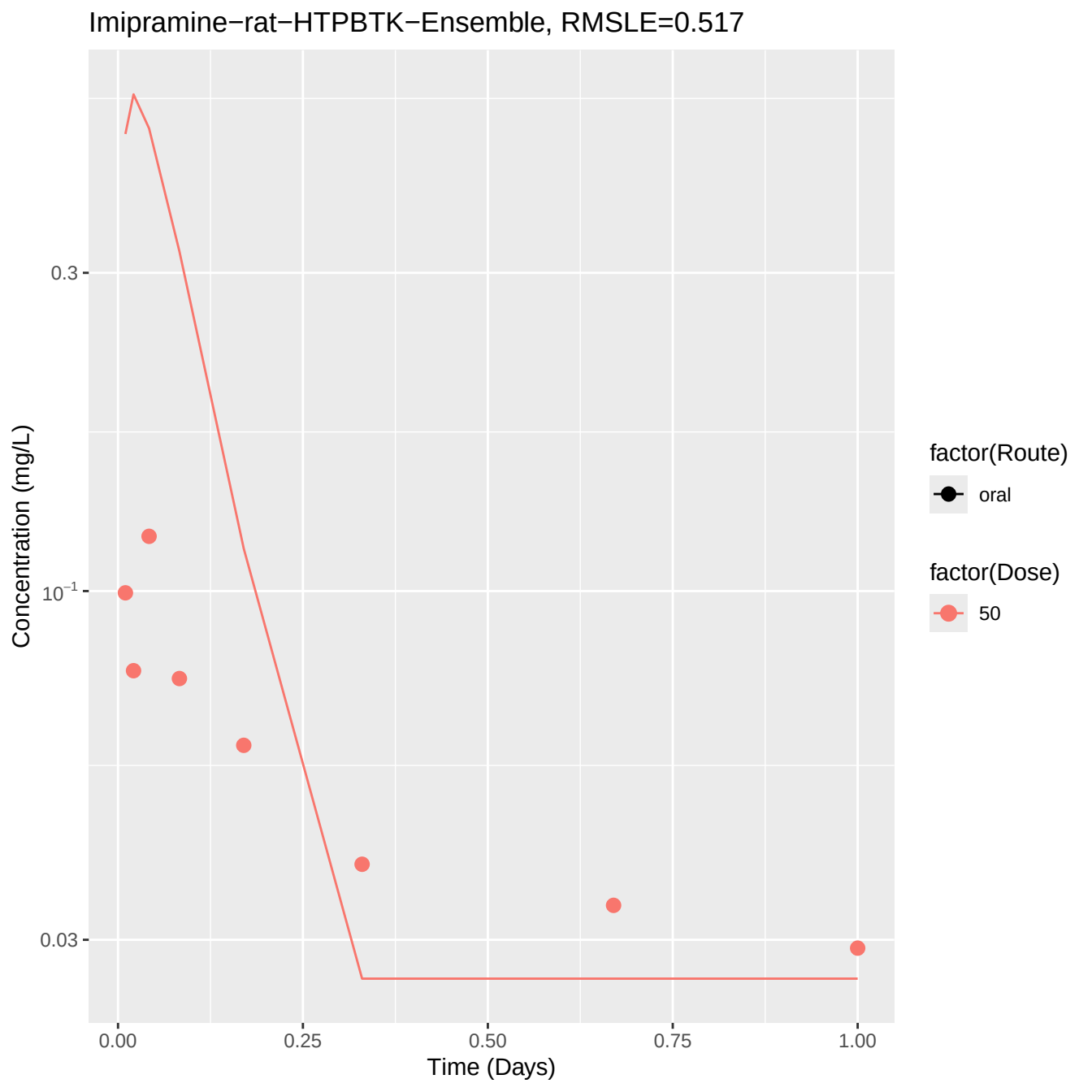

Potassium perfluorohexanesulfonate−rat−In Vivo Fits, RMSLE=0.611
Concentration (mg/L)
10^2
10
1
10^{-1}
0
10
20
30
40
50
Time (Days)
factor(Dose)
4
16
32
factor(Route)
iv
oral

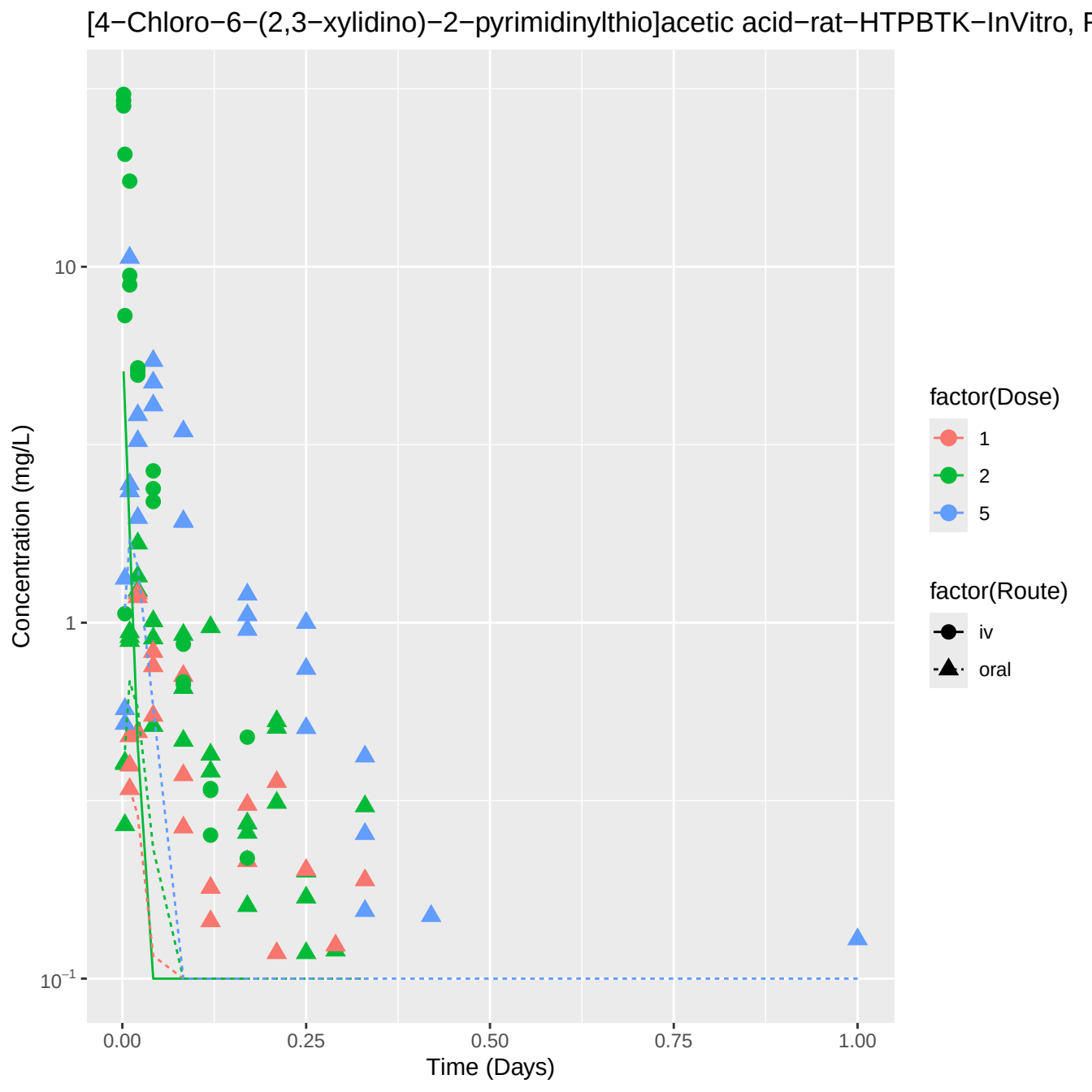

Imipramine−rat−HTPBTK−InVitro, RMSLE=0.511
Concentration (mg/L)
0.3
10^{-1}
0.03
0.00
0.25
0.50
0.75
1.00
Time (Days)
factor(Route)
oral
factor(Dose)
50

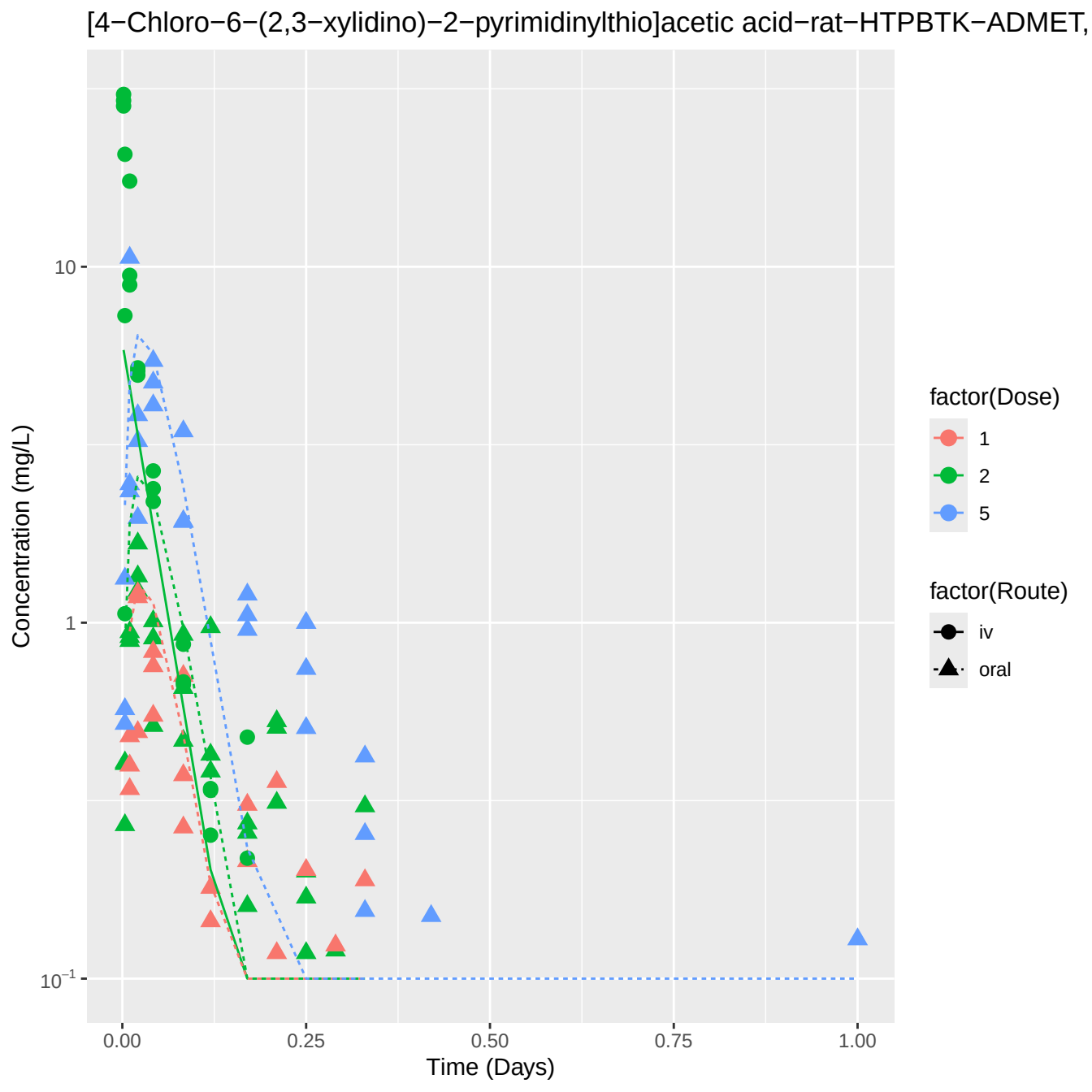

Imipramine−rat−HTPBTK−ADMET, RMSLE=0.868
Concentration (mg/L)
1
0.3
10^{-1}
0.03
0.00
0.25
0.50
0.75
1.00
Time (Days)
factor(Route)
oral
factor(Dose)
50

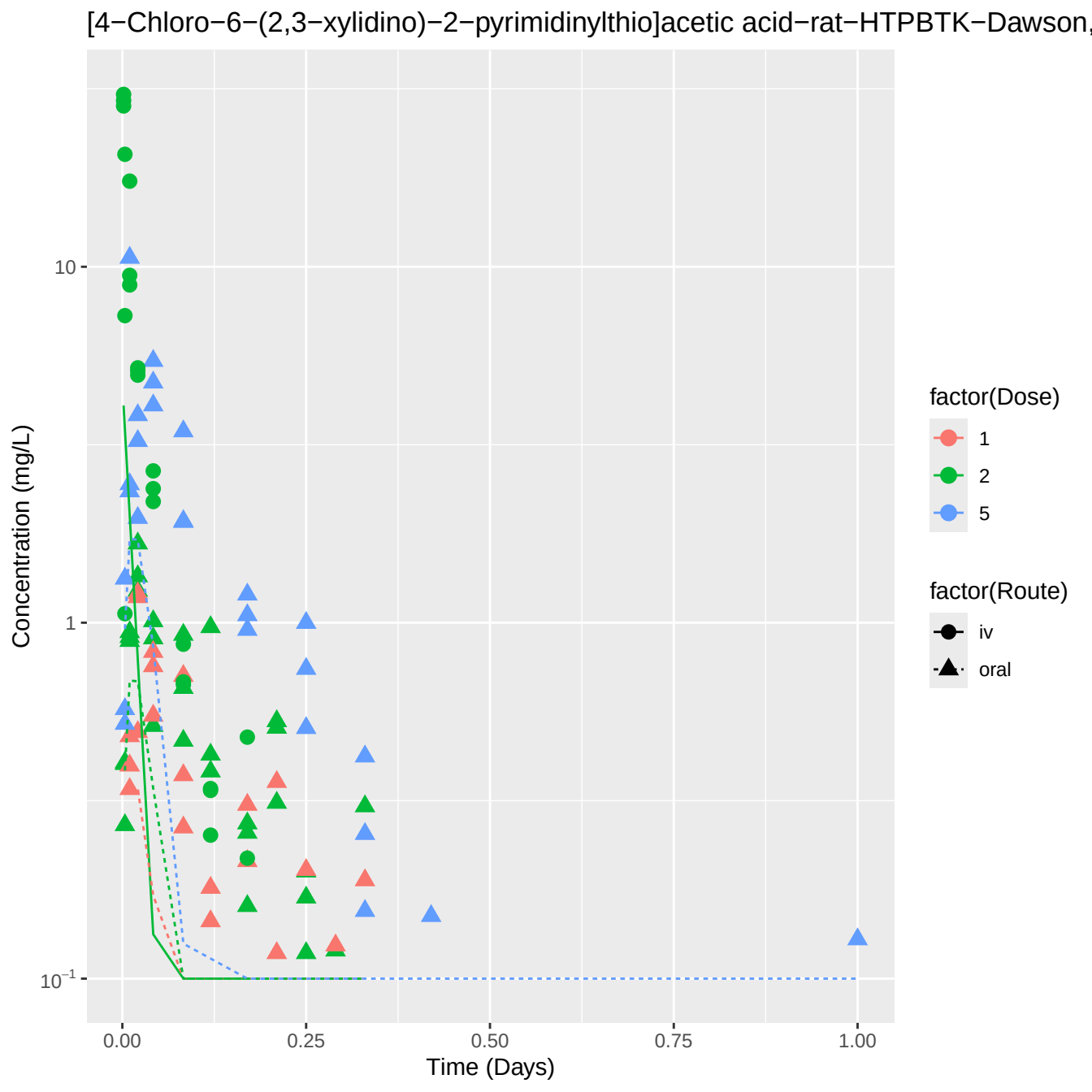

Imipramine−rat−HTPBTK−Dawson, RMSLE=0.658
Concentration (mg/L)
1
0.3
10^{-1}
0.03
0.00
0.25
0.50
0.75
1.00
Time (Days)
factor(Route)
oral
factor(Dose)
50

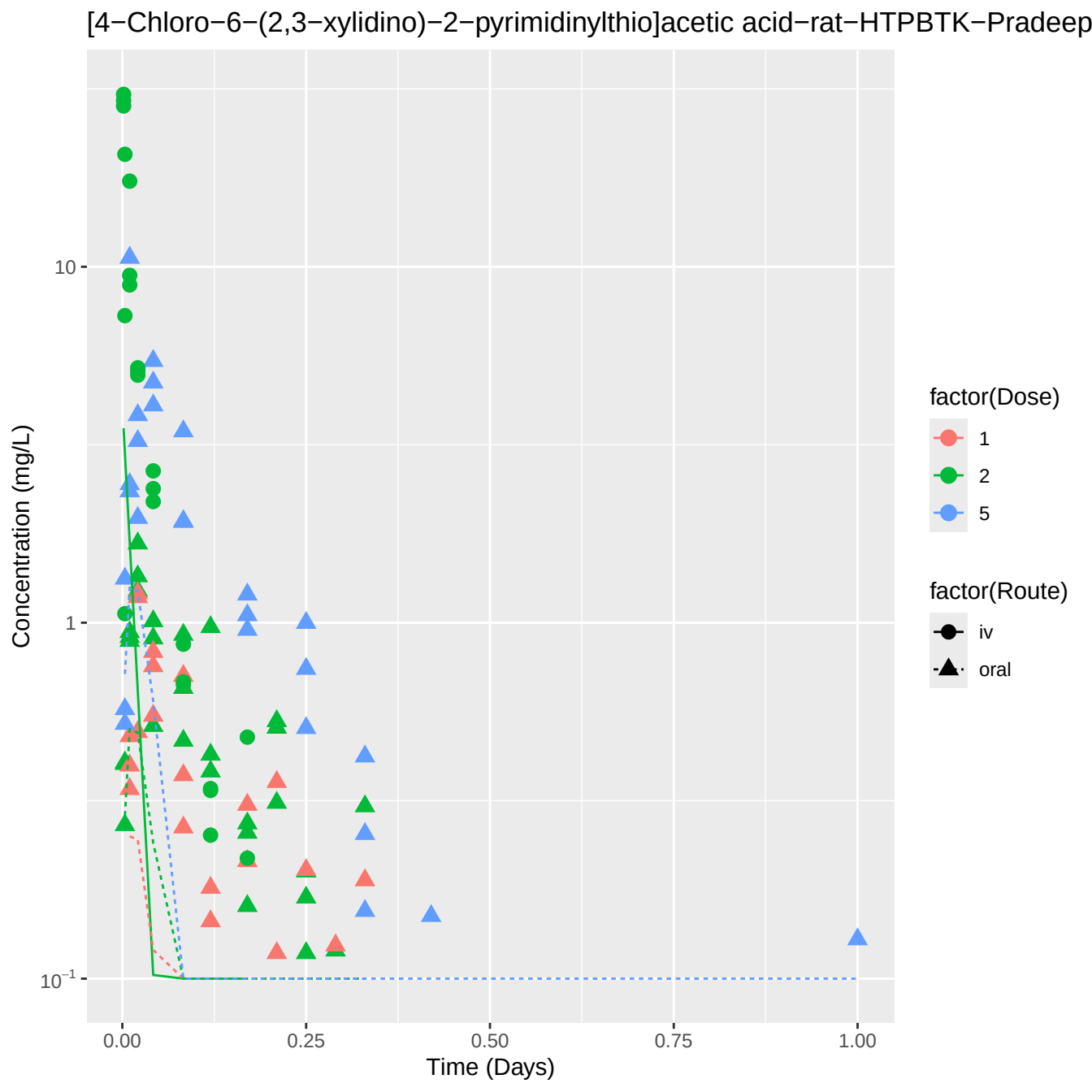

Imipramine−rat−HTPBTK−Pradeep, RMSLE=1.3
Concentration (mg/L)
3
1
0.3
10−1
0.03
0.00
0.25
0.50
0.75
1.00
Time (Days)
factor(Route)
oral
factor(Dose)
50

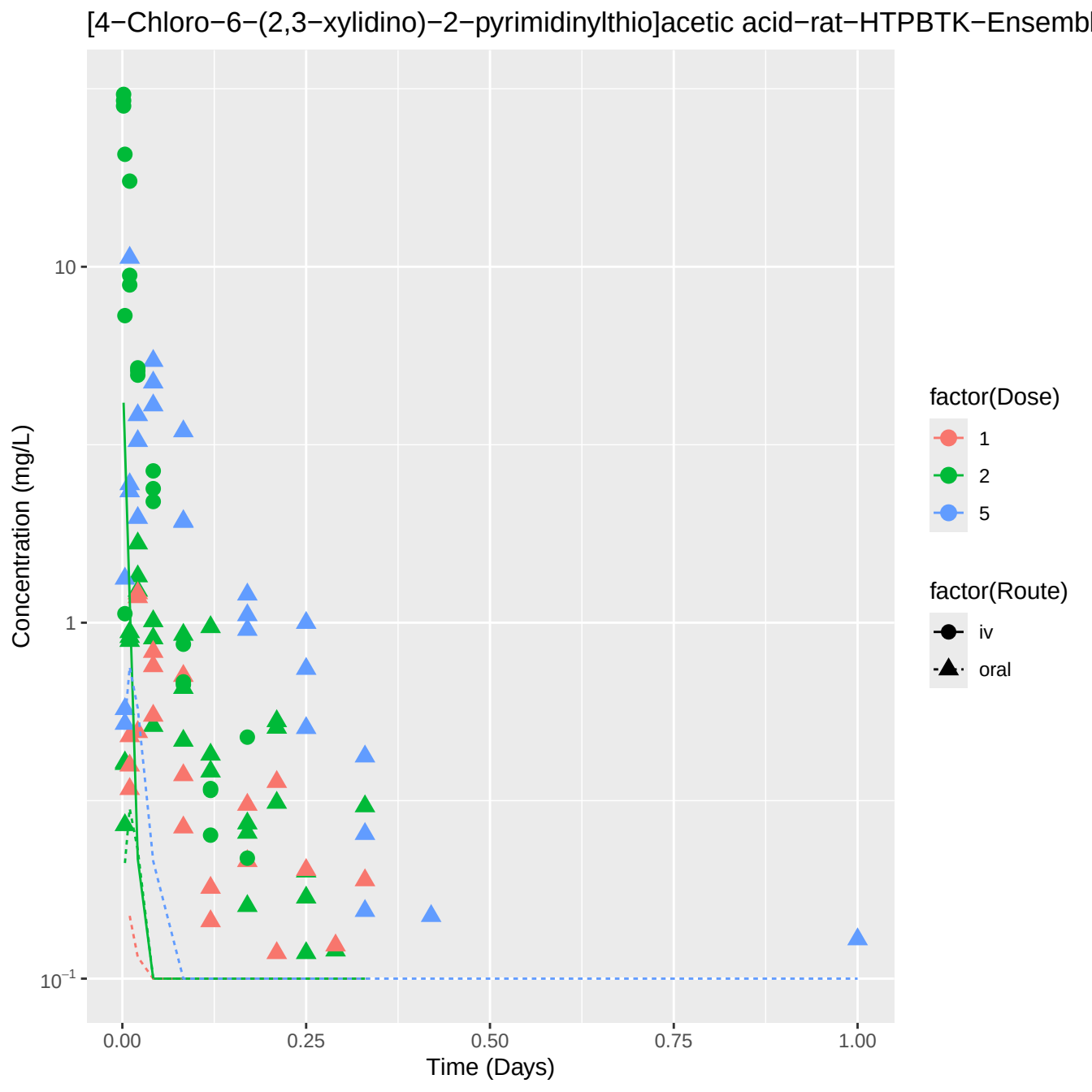

Imipramine−rat−HTPBTK−Ensemble, RMSLE=0.517
Concentration (mg/L)
0.3
10^{-1}
0.03
0.00
0.25
0.50
0.75
1.00
Time (Days)
factor(Route)
oral
factor(Dose)
50

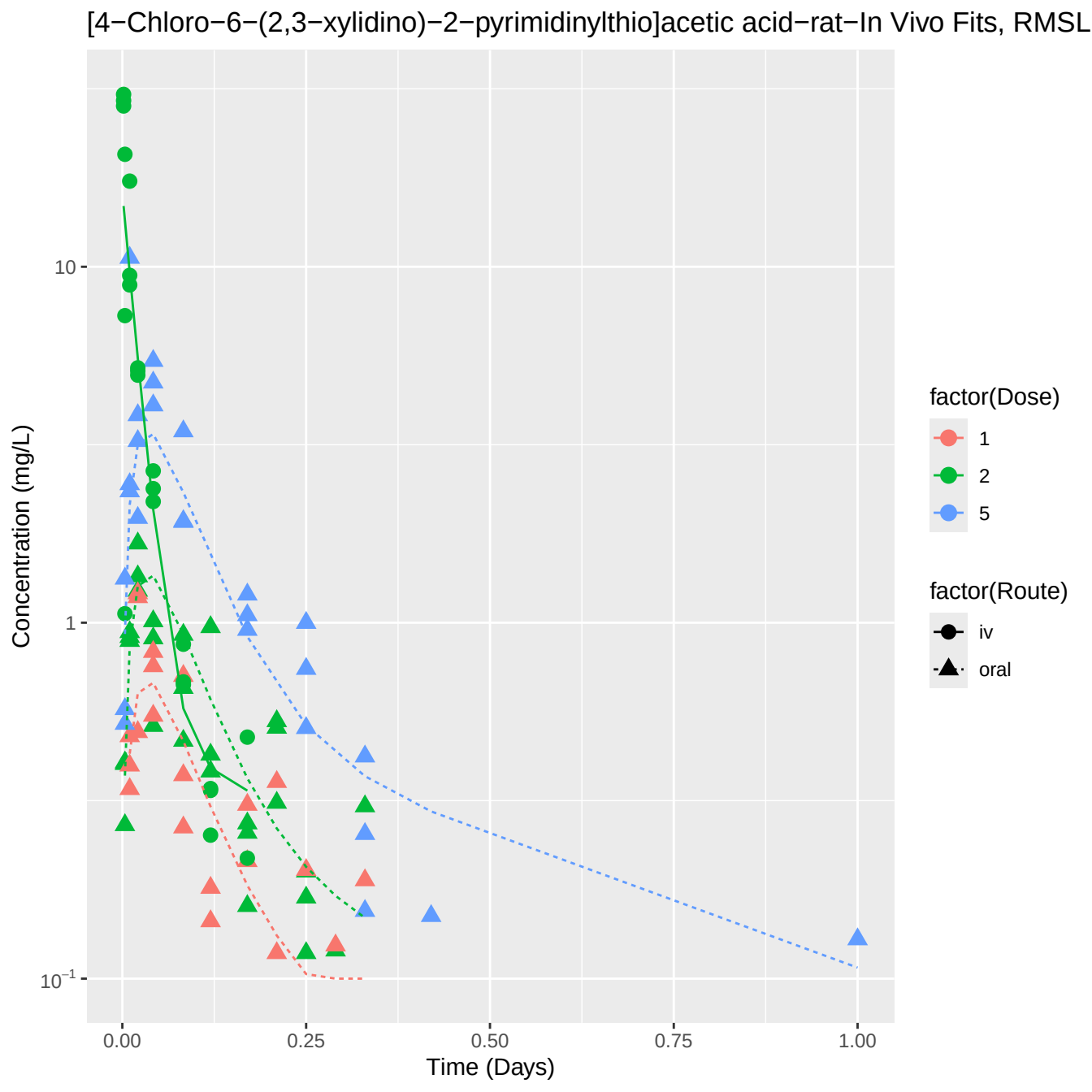

[4−Chloro−6−(2,3−xylidino)−2−pyrimidinylthio]acetic acid−rat−HTPBTK−InVitro, P
Concentration (mg/L)
10
1
10^{-1}
0.00
0.25
0.50
0.75
1.00
Time (Days)
factor(Dose)
1
2
5
factor(Route)
iv
oral

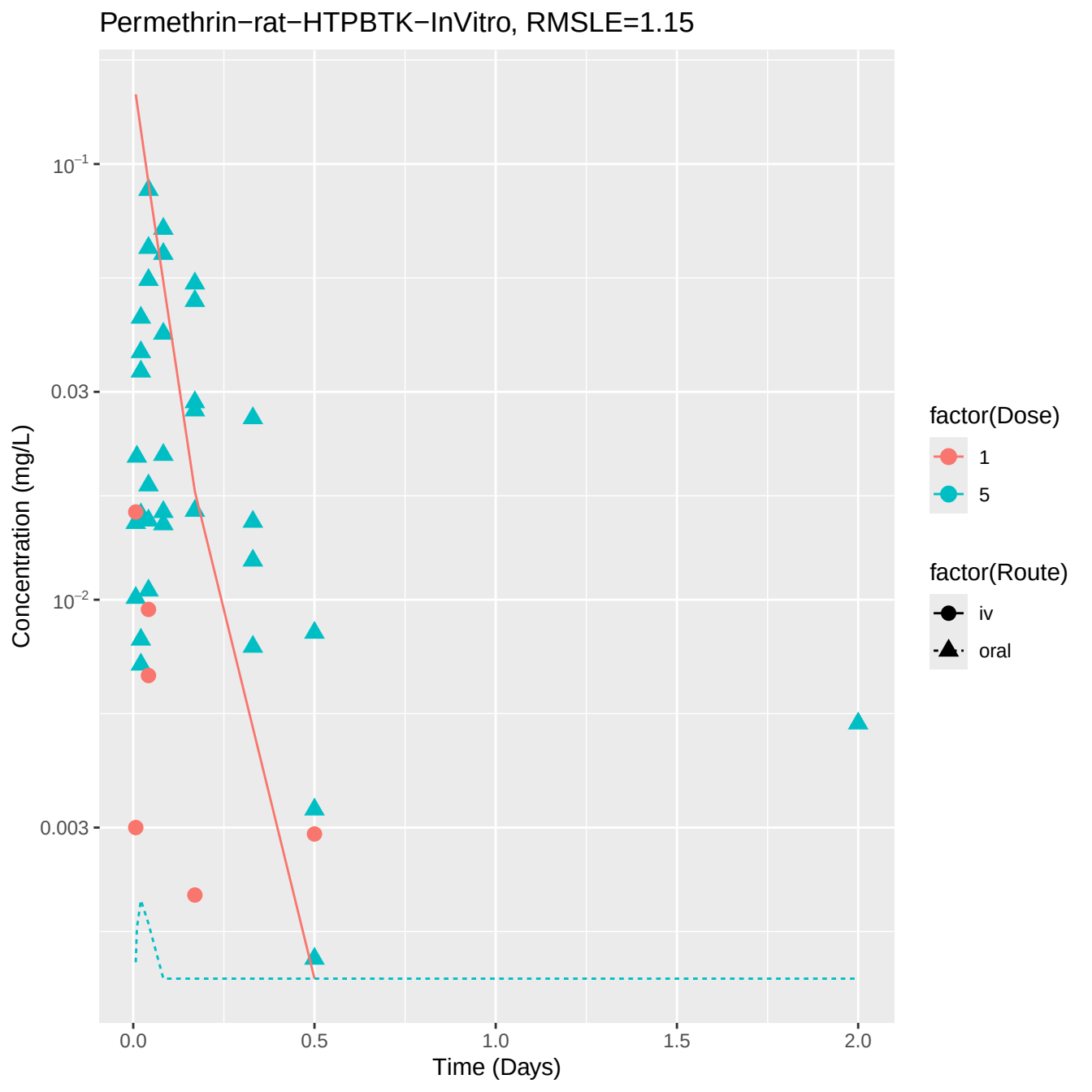

[4−Chloro−6−(2,3−xylidino)−2−pyrimidinylthio]acetic acid−rat−HTPBTK−ADMET,
Concentration (mg/L)
10
1
10^{-1}
0.00
0.25
0.50
0.75
1.00
Time (Days)
factor(Dose)
1
2
5
factor(Route)
iv
oral

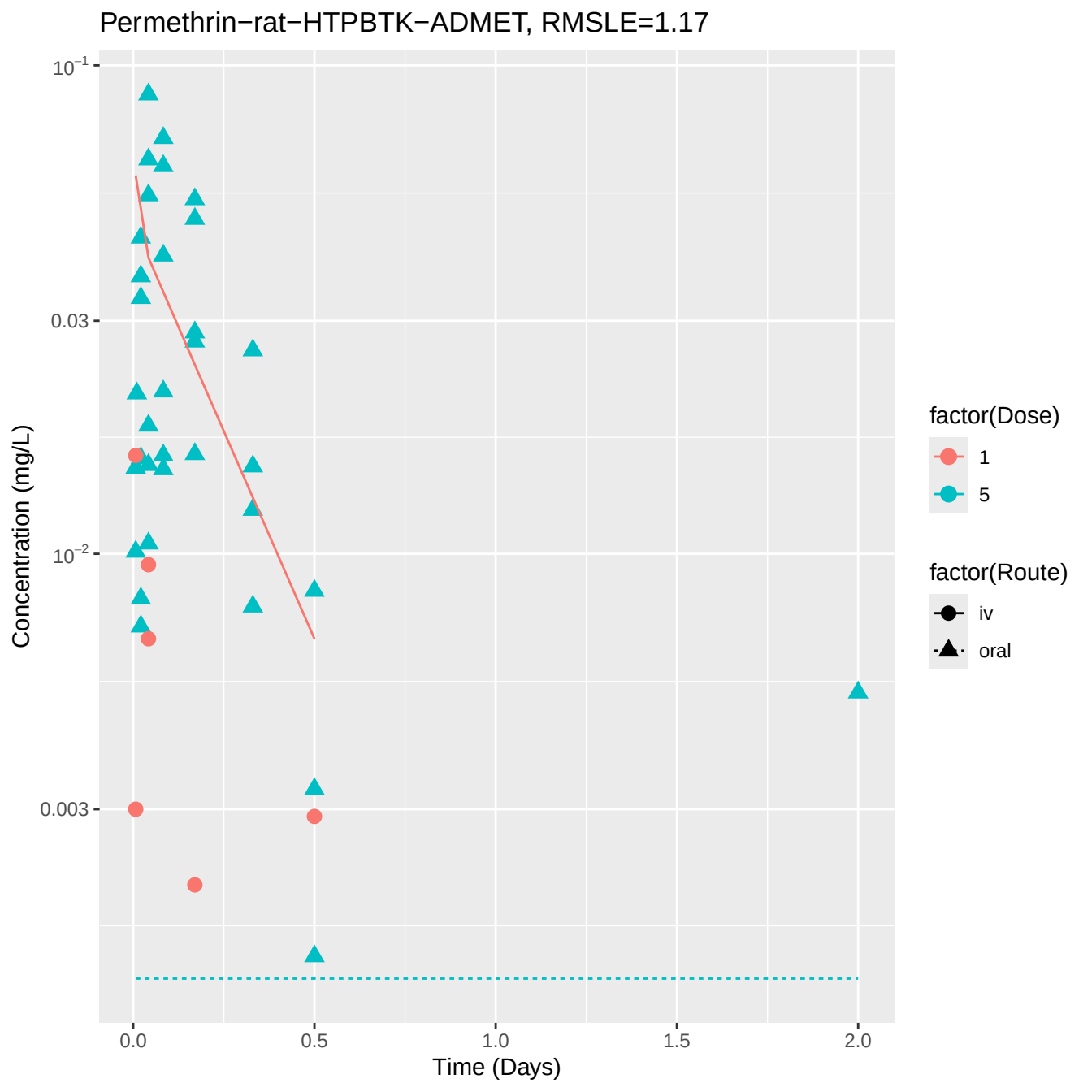

[4−Chloro−6−(2,3−xylidino)−2−pyrimidinylthio]acetic acid−rat−HTPBTK−Dawson,
Concentration (mg/L)
10
1
10^{-1}
0.00
0.25
0.50
0.75
1.00
Time (Days)
factor(Dose)
1
2
5
factor(Route)
iv
oral

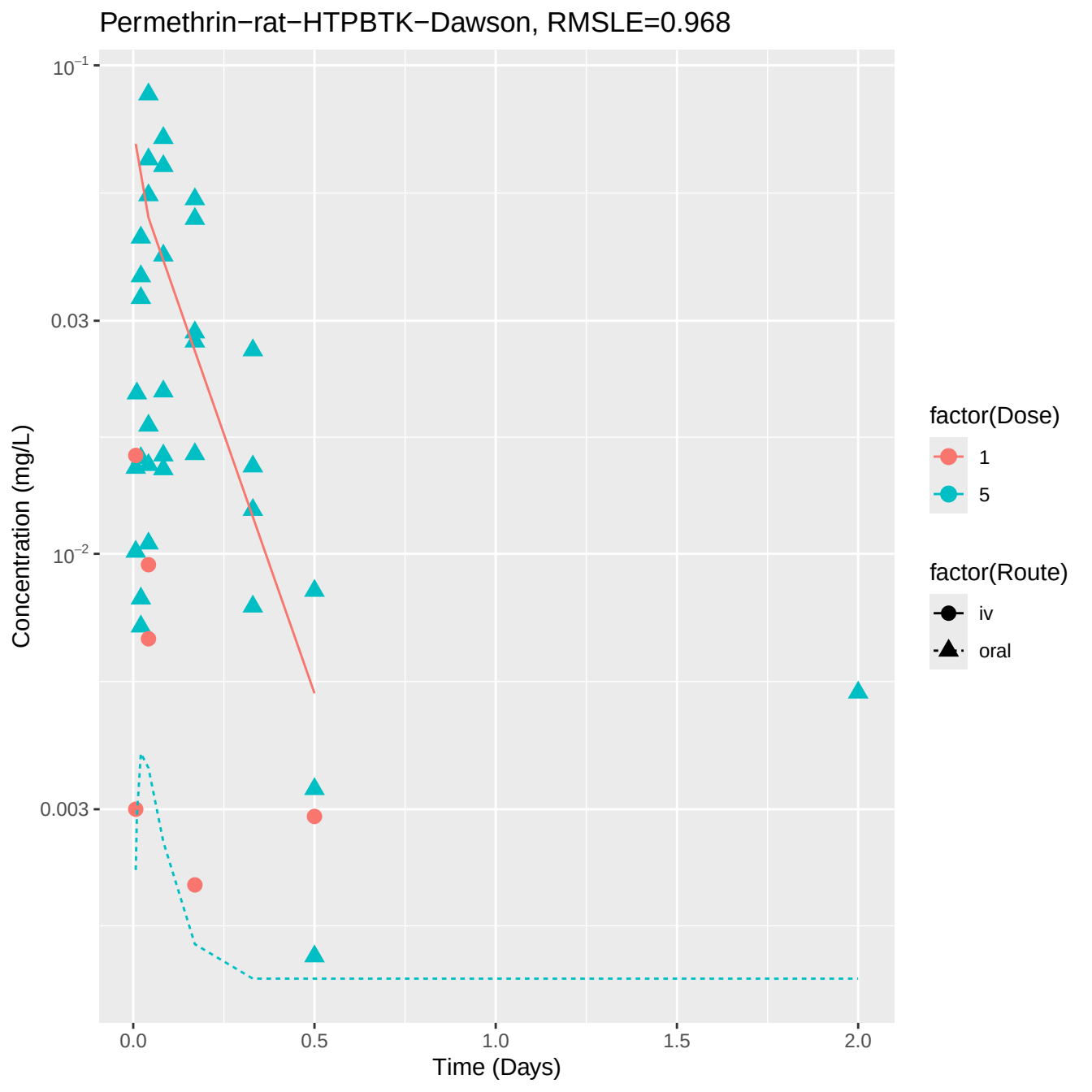

[4−Chloro−6−(2,3−xylidino)−2−pyrimidinylthio]acetic acid−rat−HTPBTK−Pradeep
Concentration (mg/L)
10
1
10^{-1}
0.00
0.25
0.50
0.75
1.00
Time (Days)
factor(Dose)
1
2
5
factor(Route)
iv
oral

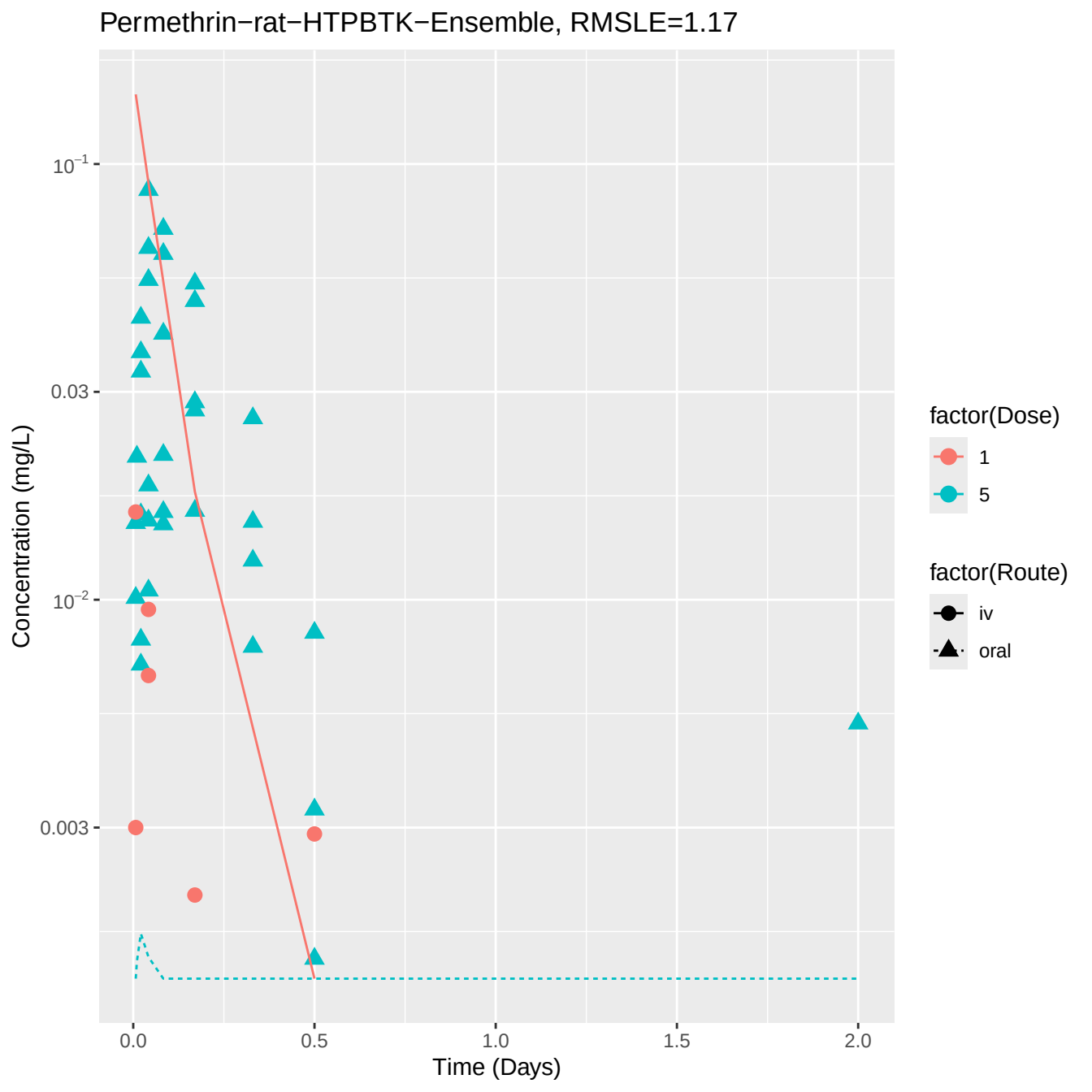

[4−Chloro−6−(2,3−xylidino)−2−pyrimidinylthio]acetic acid−rat−HTPBTK−Ensembl
Concentration (mg/L)
10
1
10^{-1}
0.00
0.25
0.50
0.75
1.00
Time (Days)
factor(Dose)
1
2
5
factor(Route)
iv
oral

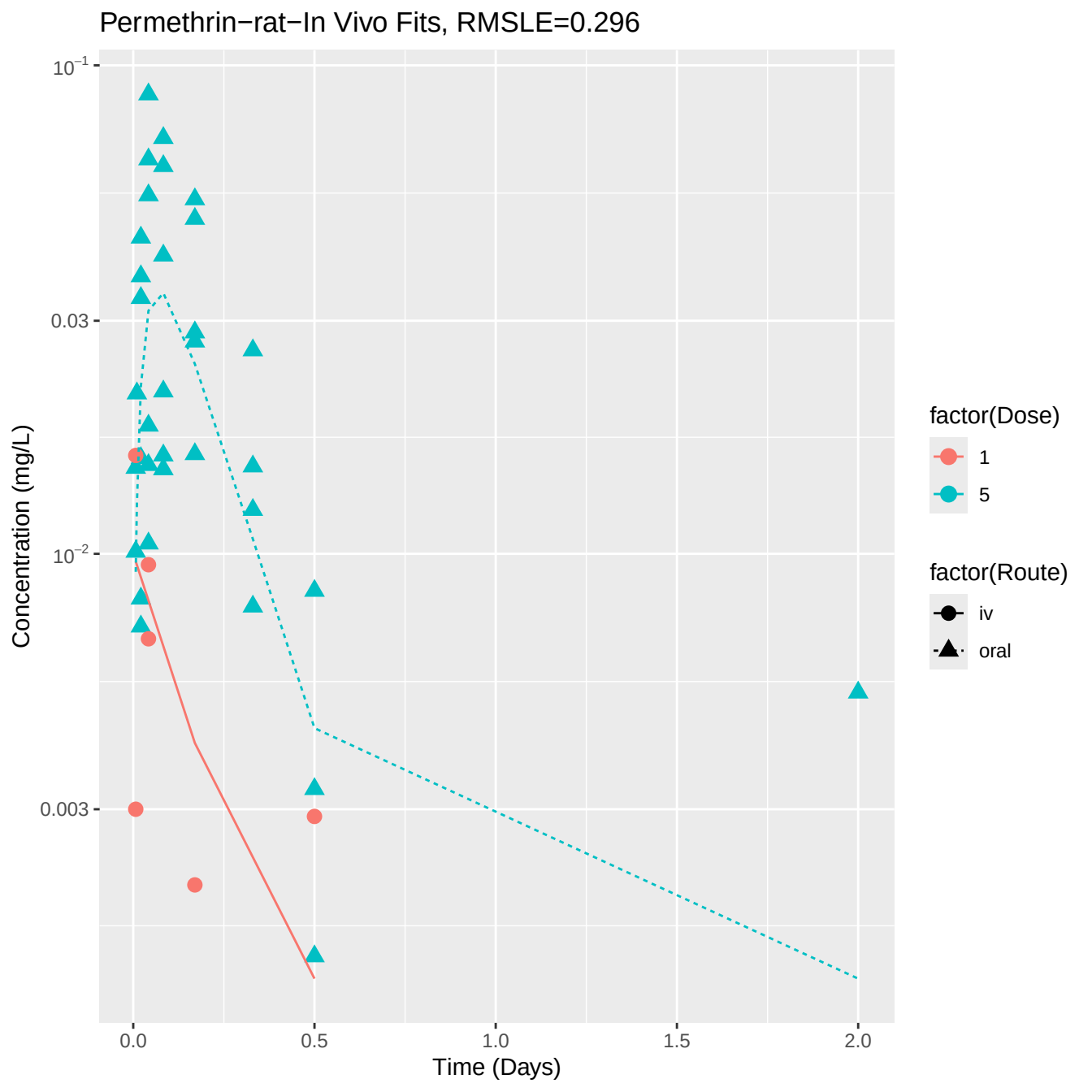

[4−Chloro−6−(2,3−xylidino)−2−pyrimidinylthio]acetic acid−rat−In Vivo Fits, RMSL
Concentration (mg/L)
10
1
10^{-1}
0.00
0.25
0.50
0.75
1.00
Time (Days)
factor(Dose)
1
2
5
factor(Route)
iv
oral

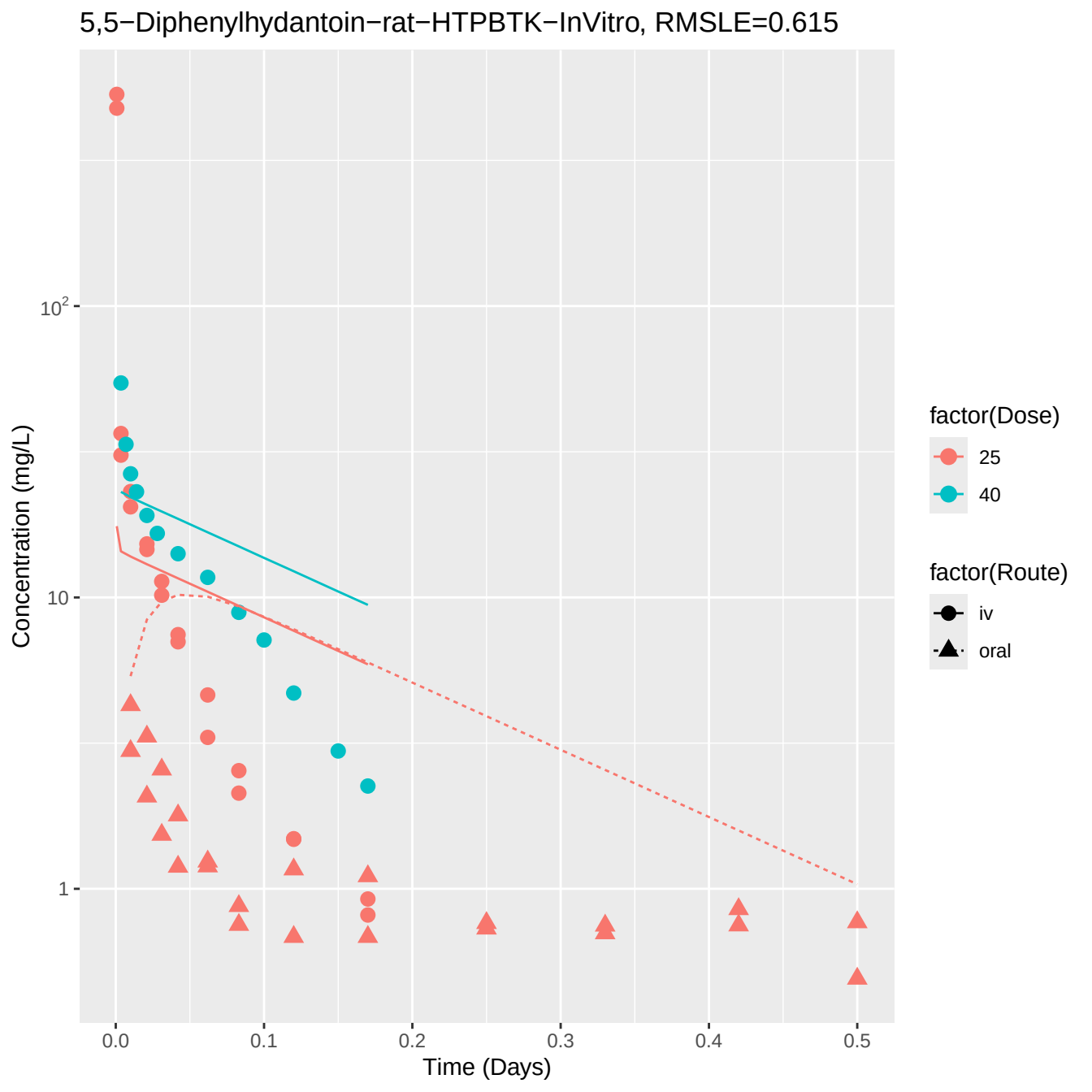

Permethrin−rat−HTPBTK−InVitro, RMSLE=1.15
Concentration (mg/L)
10^{-1}
0.03
10^{-2}
0.003
0.0
0.5
1.0
1.5
2.0
Time (Days)
factor(Dose)
1
5
factor(Route)
iv
oral

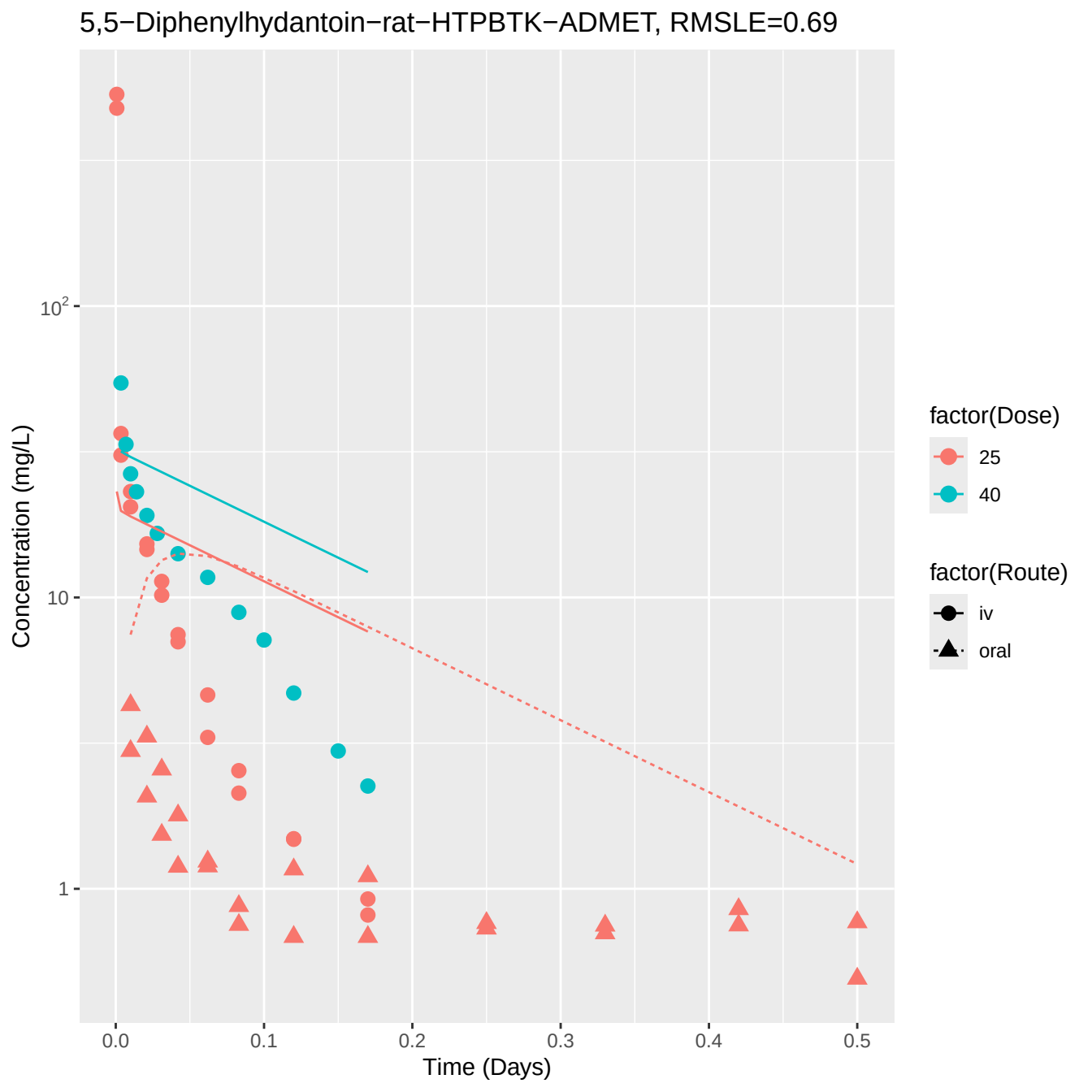

Permethrin−rat−HTPBTK−ADMET, RMSLE=1.17
Concentration (mg/L)
10^−1
0.03
10^−2
0.003
0.0
0.5
1.0
1.5
2.0
Time (Days)
factor(Dose)
1
5
factor(Route)
iv
oral

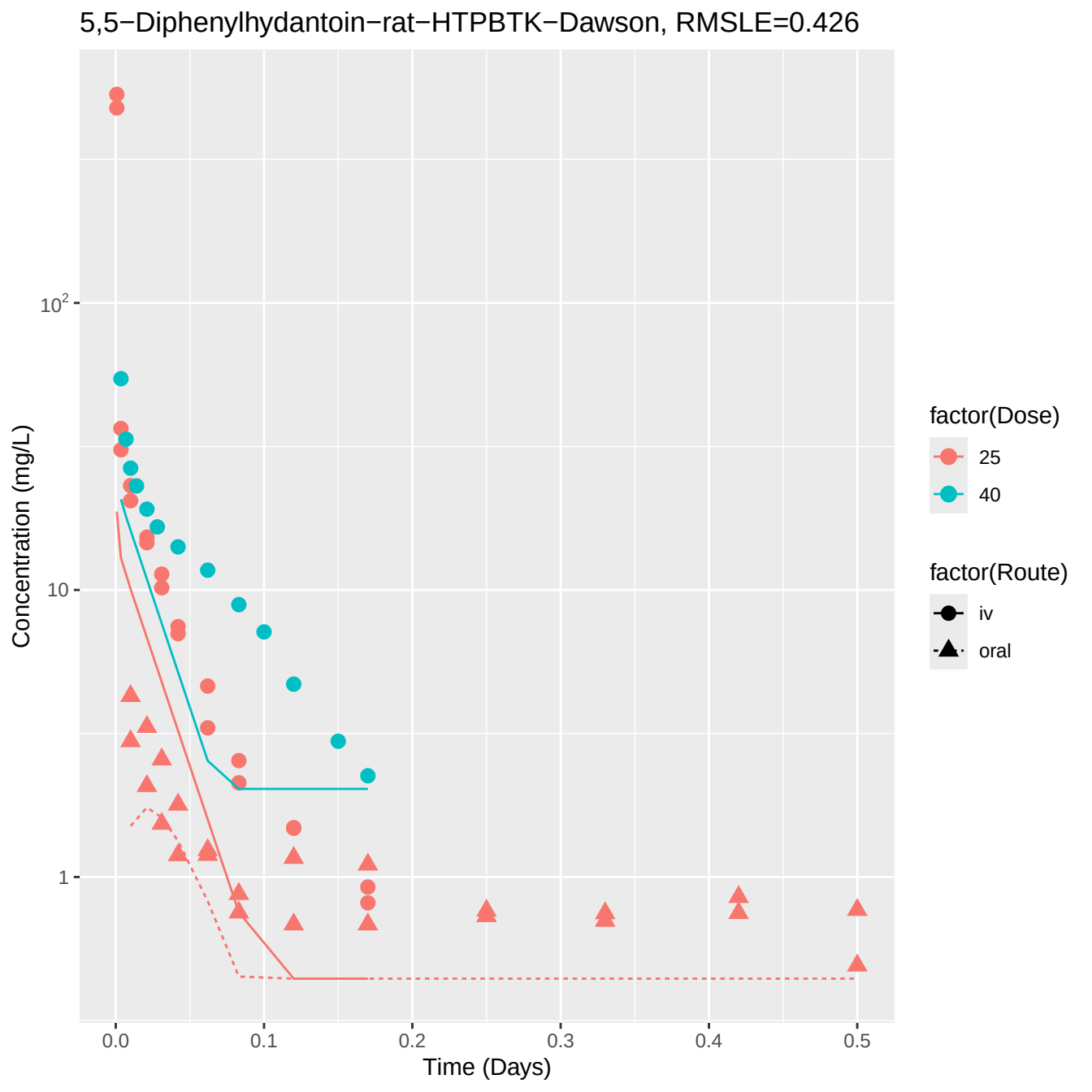

Permethrin−rat−HTPBTK−Dawson, RMSLE=0.968
Concentration (mg/L)
10^−1
0.03
10^−2
0.003
0.0
0.5
1.0
1.5
2.0
Time (Days)
factor(Dose)
1
5
factor(Route)
iv
oral

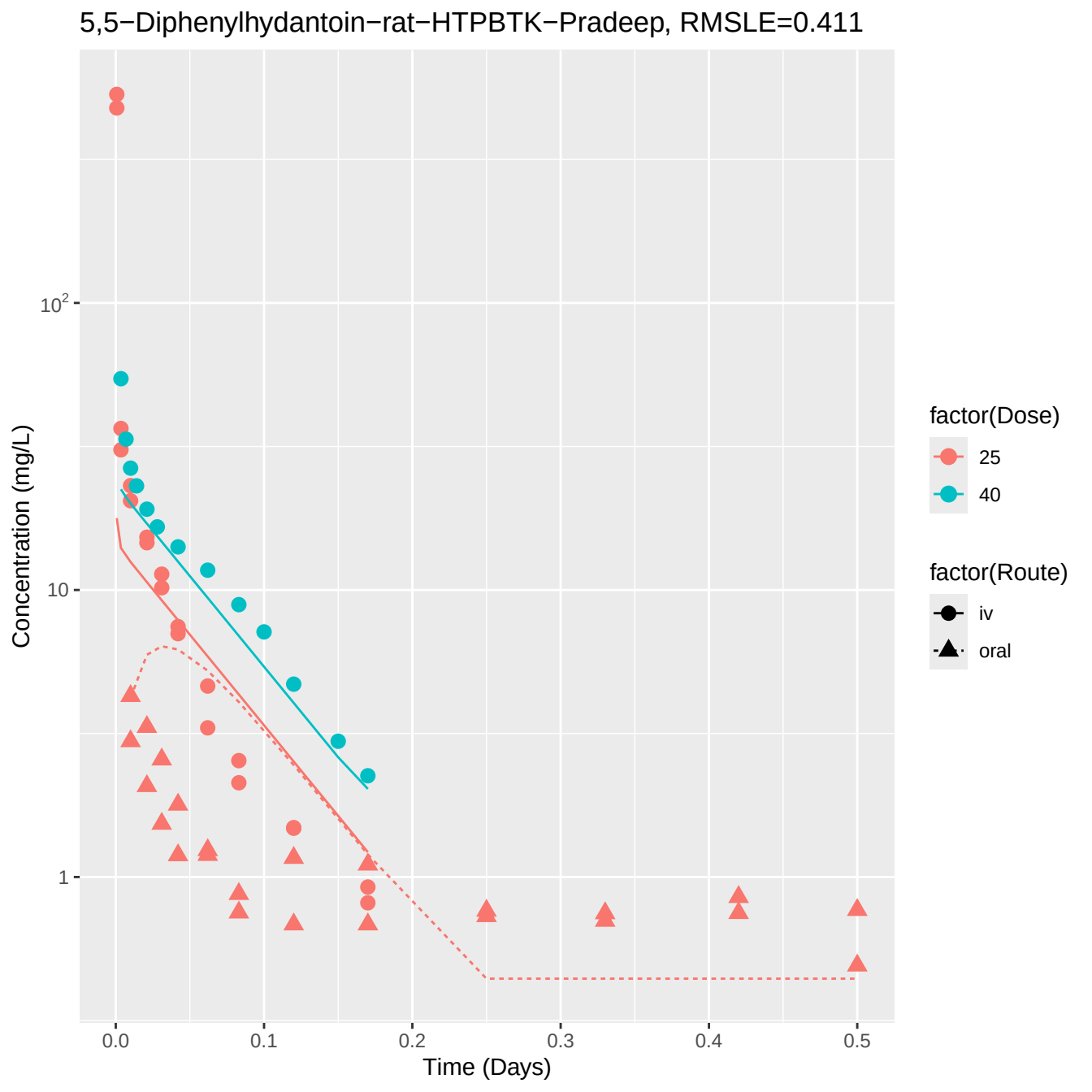

Permethrin−rat−HTPBTK−Ensemble, RMSLE=1.17
Concentration (mg/L)
10^{-1}
0.03
10^{-2}
0.003
0.0
0.5
1.0
1.5
2.0
Time (Days)
factor(Dose)
1
5
factor(Route)
iv
oral

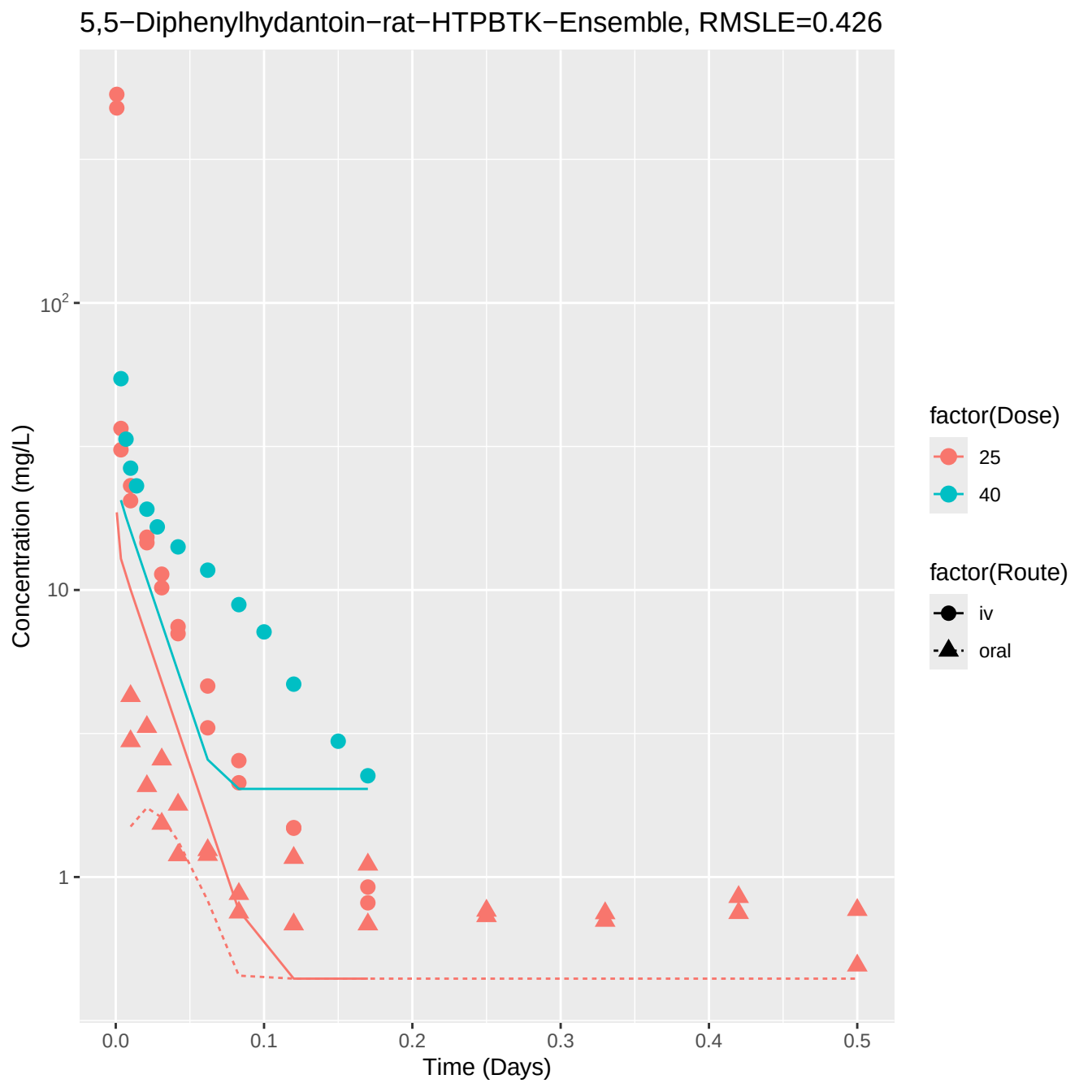

Permethrin−rat−In Vivo Fits, RMSLE=0.296
Concentration (mg/L)
10^−1
0.03
10^−2
0.003
0.0
0.5
1.0
1.5
2.0
Time (Days)
factor(Dose)
1
5
factor(Route)
iv
oral

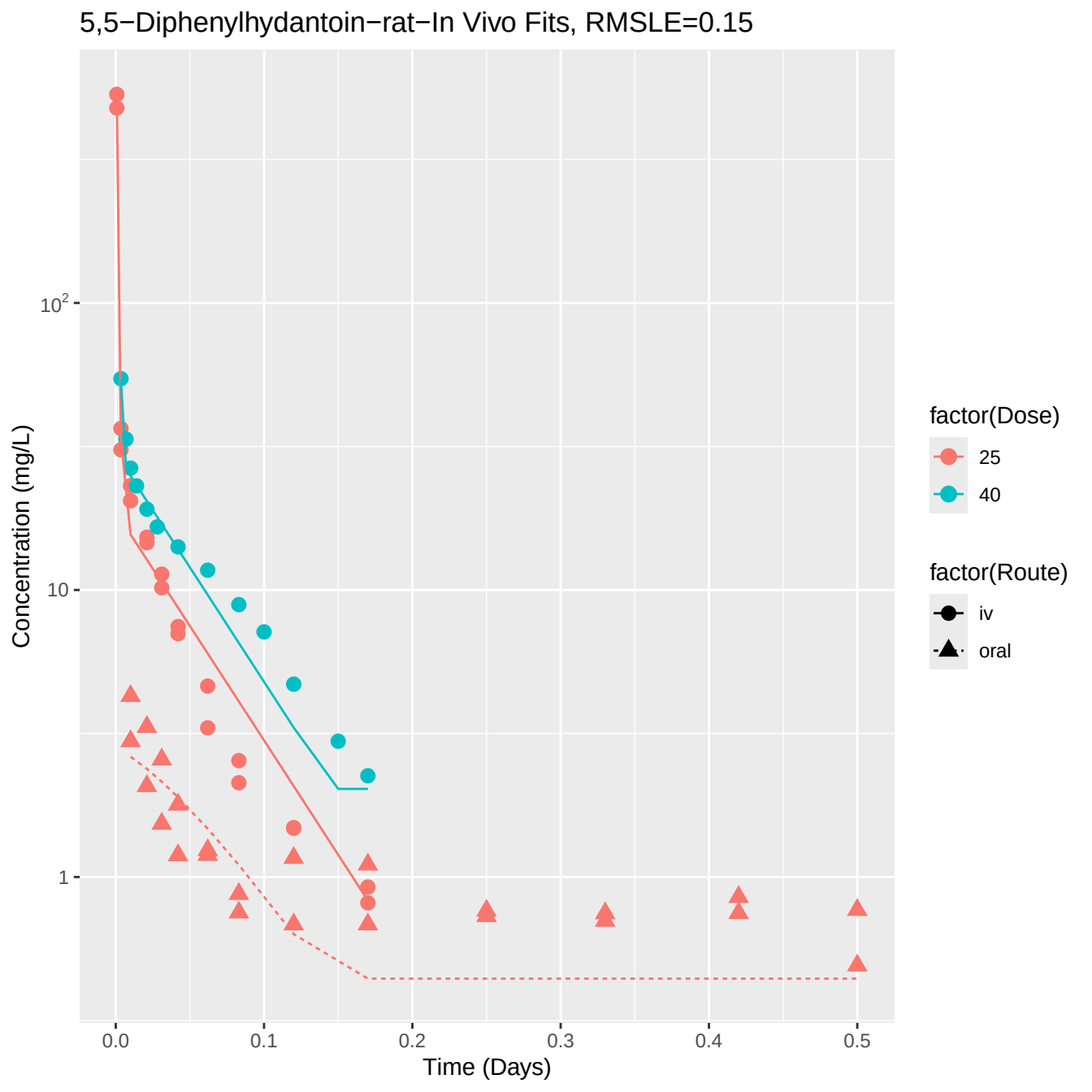

5,5−Diphenylhydantoin−rat−HTPBTK−InVitro, RMSLE=0.615
Concentration (mg/L)
10^2
10
1
0.0
0.1
0.2
0.3
0.4
0.5
Time (Days)
factor(Dose)
25
40
factor(Route)
iv
oral

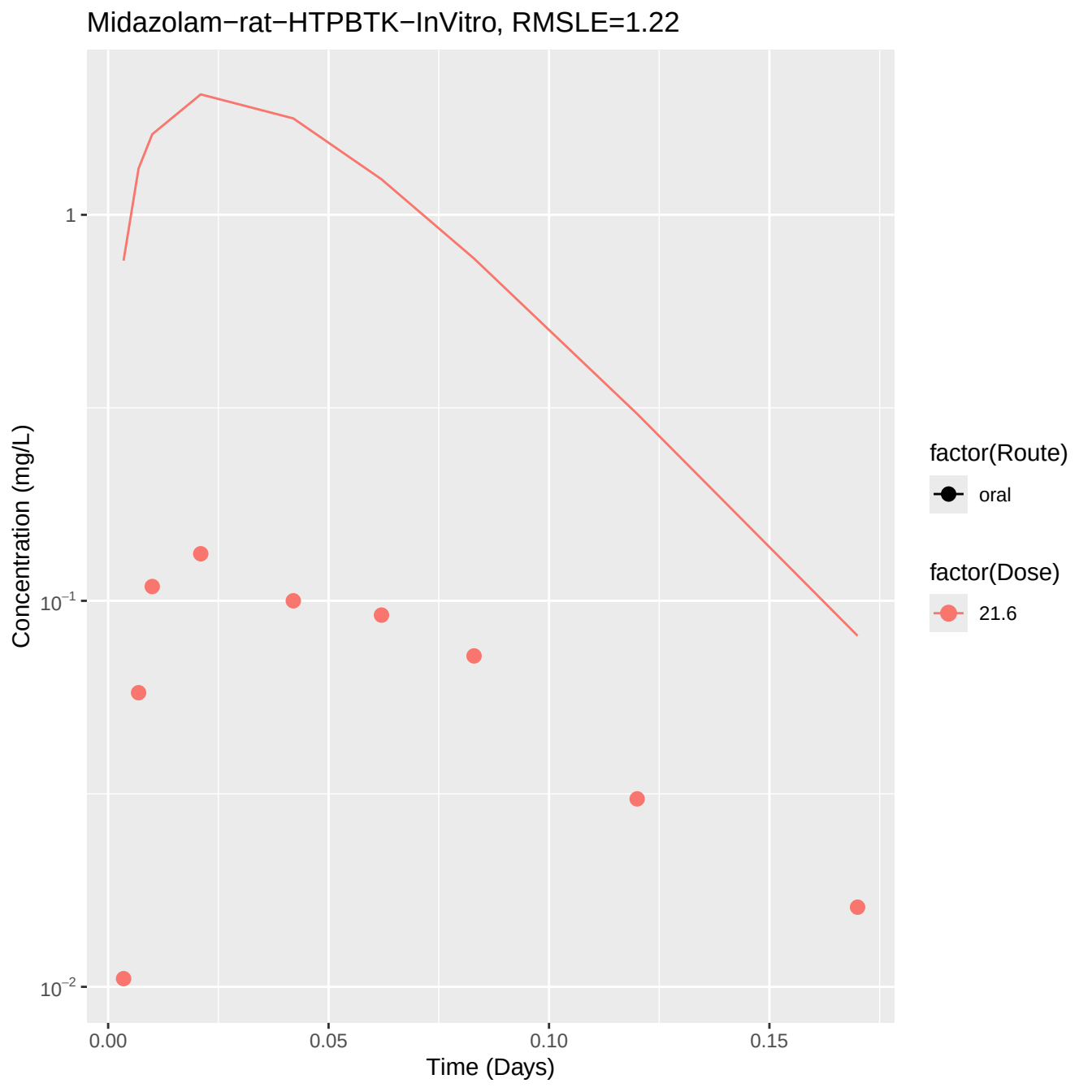

5,5−Diphenylhydantoin−rat−HTPBTK−ADMET, RMSLE=0.69
Concentration (mg/L)
10^2
10
1
0.0
0.1
0.2
0.3
0.4
0.5
Time (Days)
factor(Dose)
25
40
factor(Route)
iv
oral

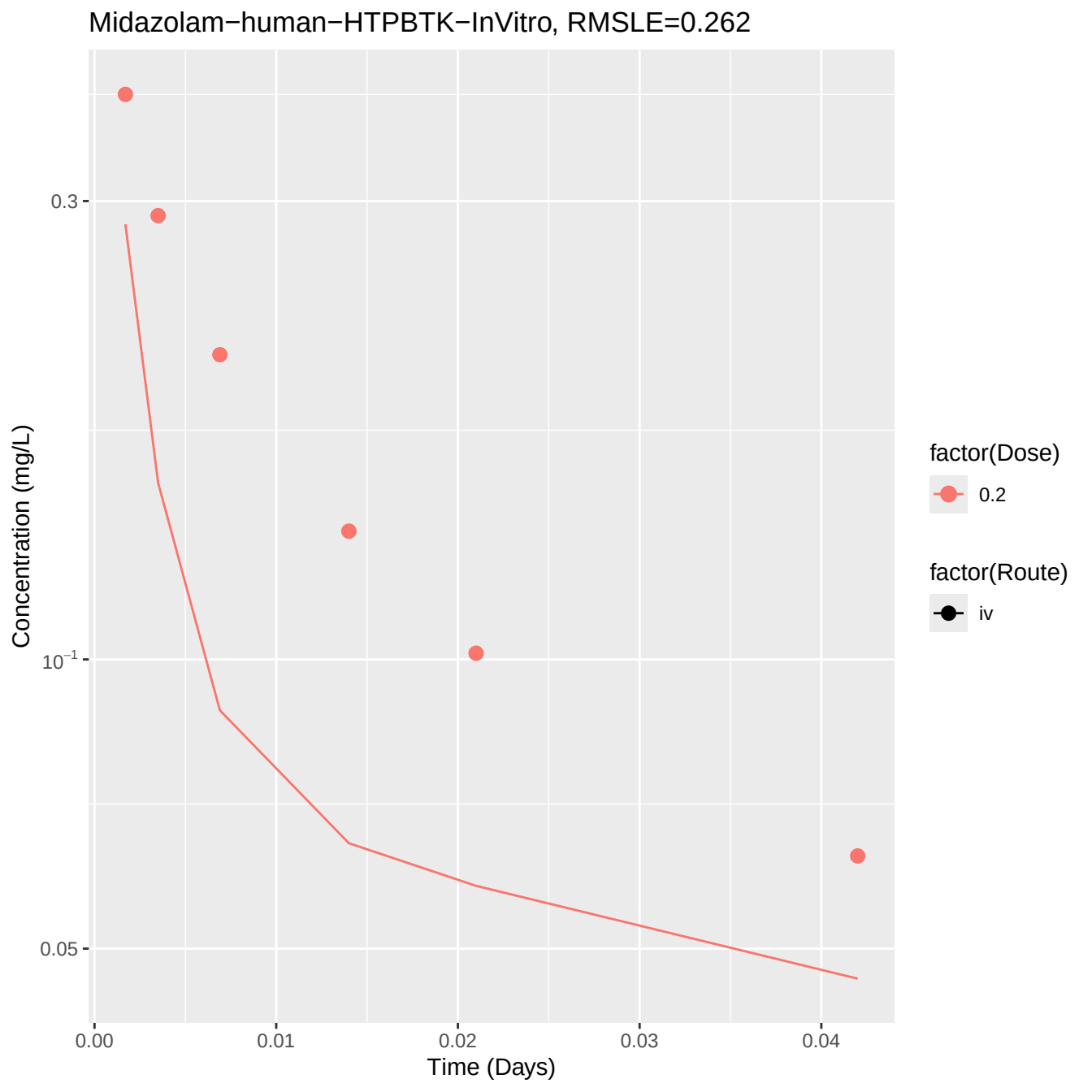

5,5−Diphenylhydantoin−rat−HTPBTK−Dawson, RMSLE=0.426
Concentration (mg/L)
10^2
10
1
0.0
0.1
0.2
0.3
0.4
0.5
Time (Days)
factor(Dose)
25
40
factor(Route)
iv
oral

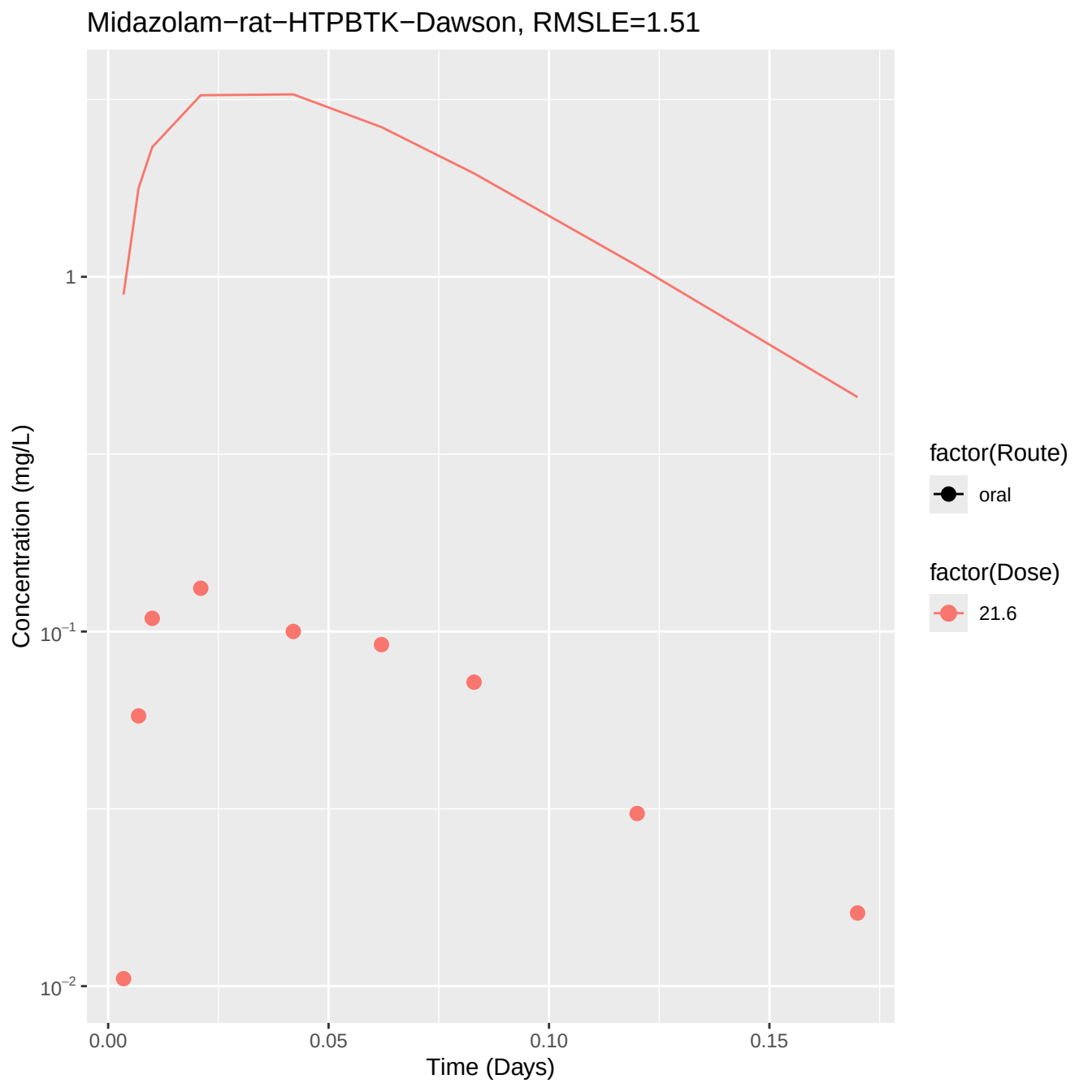

5,5−Diphenylhydantoin−rat−HTPBTK−Pradeep, RMSLE=0.411
Concentration (mg/L)
10^2
10
1
0.0
0.1
0.2
0.3
0.4
0.5
Time (Days)
factor(Dose)
25
40
factor(Route)
iv
oral

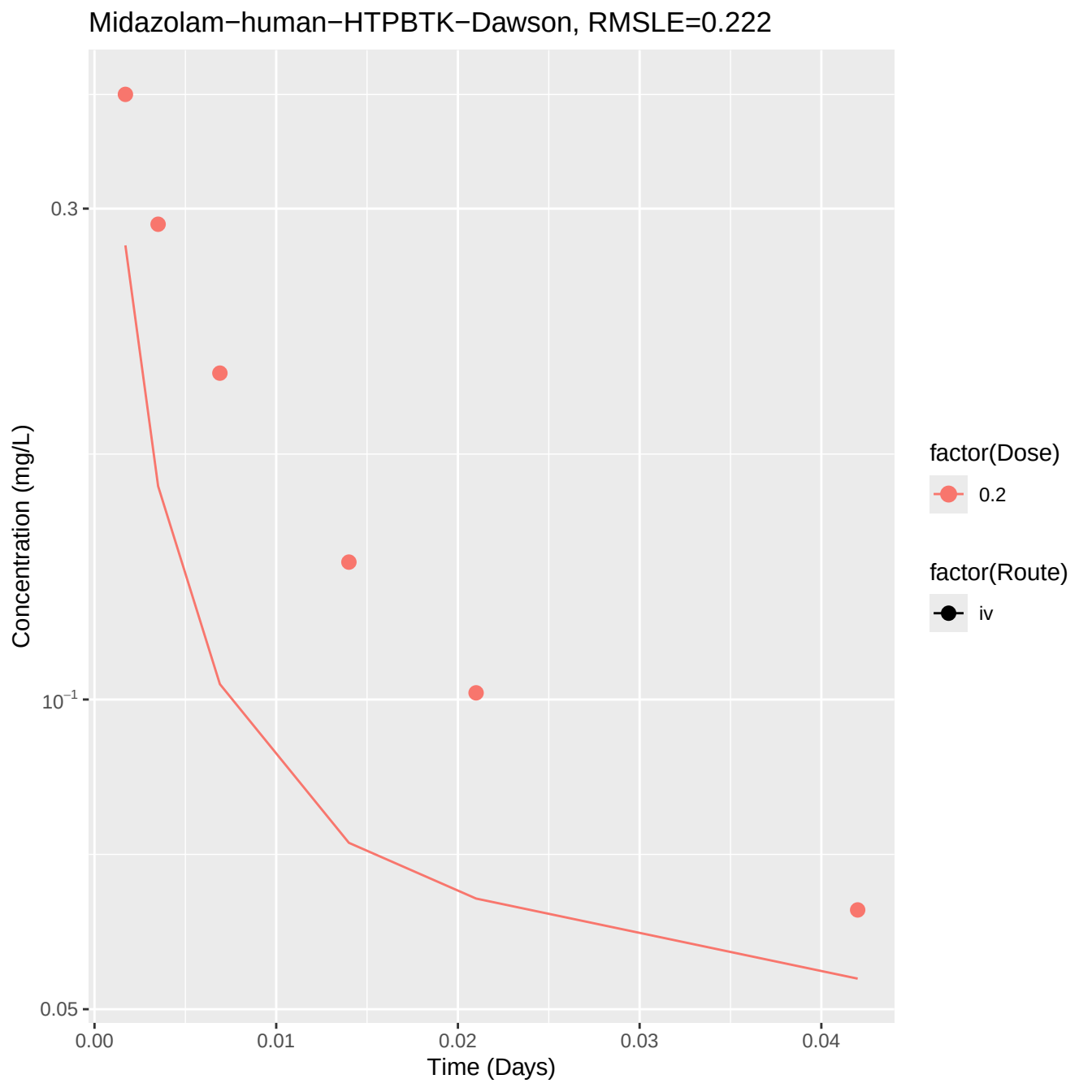

5,5−Diphenylhydantoin−rat−HTPBTK−Ensemble, RMSLE=0.426
Concentration (mg/L)
10^2
10
1
0.0
0.1
0.2
0.3
0.4
0.5
Time (Days)
factor(Dose)
25
40
factor(Route)
iv
oral

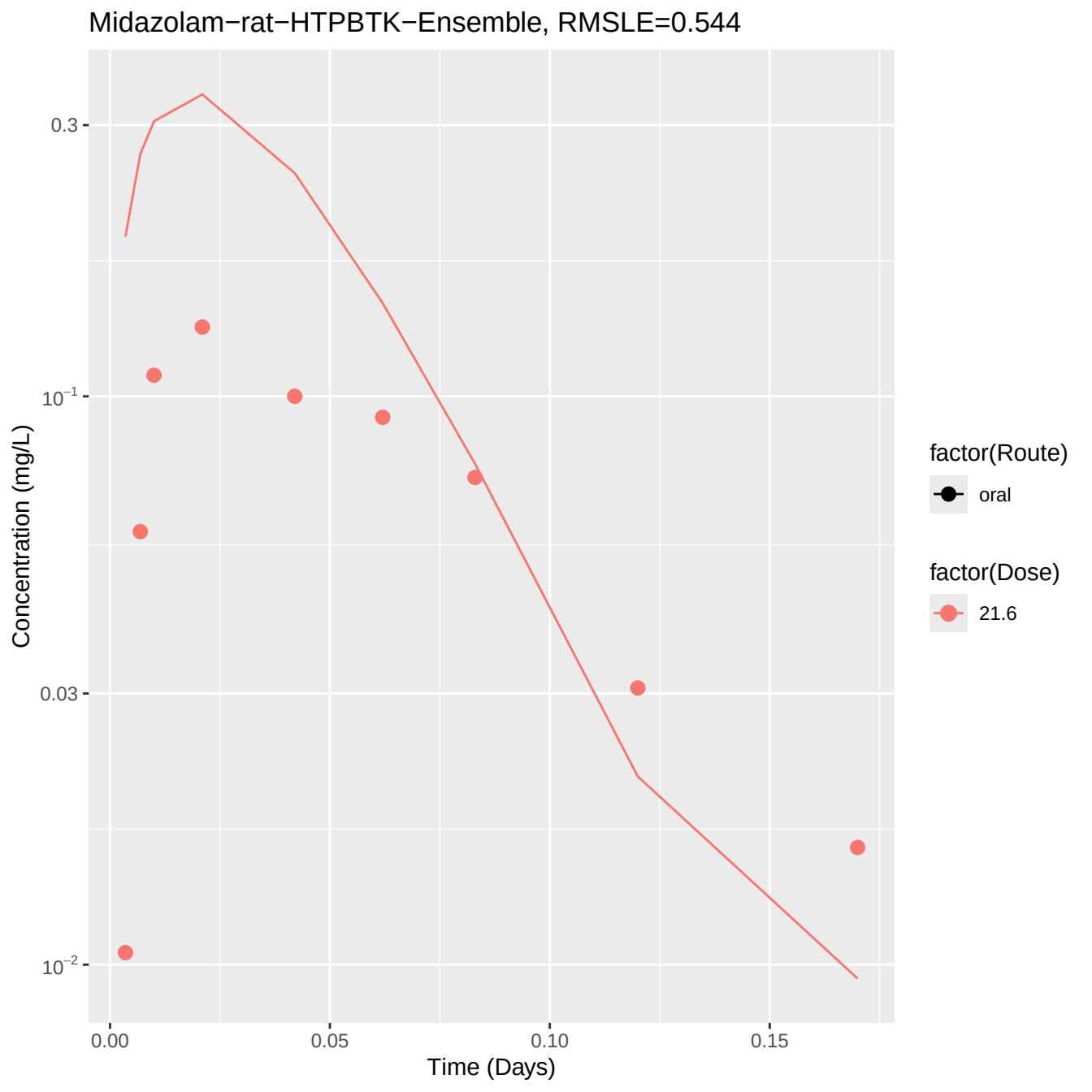

5,5−Diphenylhydantoin−rat−In Vivo Fits, RMSLE=0.15
Concentration (mg/L)
10^2
10
1
0.0
0.1
0.2
0.3
0.4
0.5
Time (Days)
factor(Dose)
25
40
factor(Route)
iv
oral

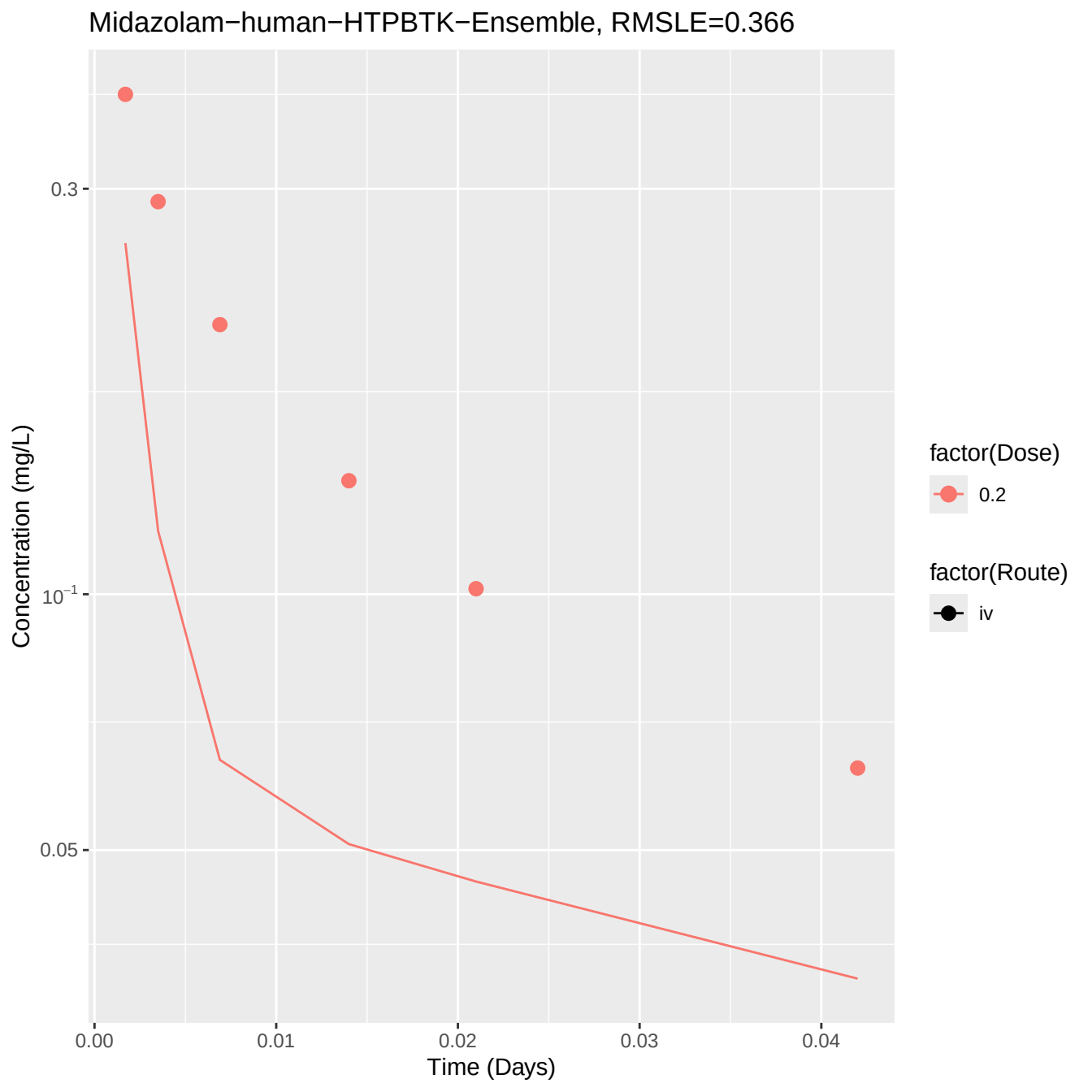

Midazolam−rat−HTPBTK−InVitro, RMSLE=1.22
Concentration (mg/L)
1
10^{-1}
10^{-2}
0.00
0.05
0.10
0.15
Time (Days)
factor(Route)
oral
factor(Dose)
21.6

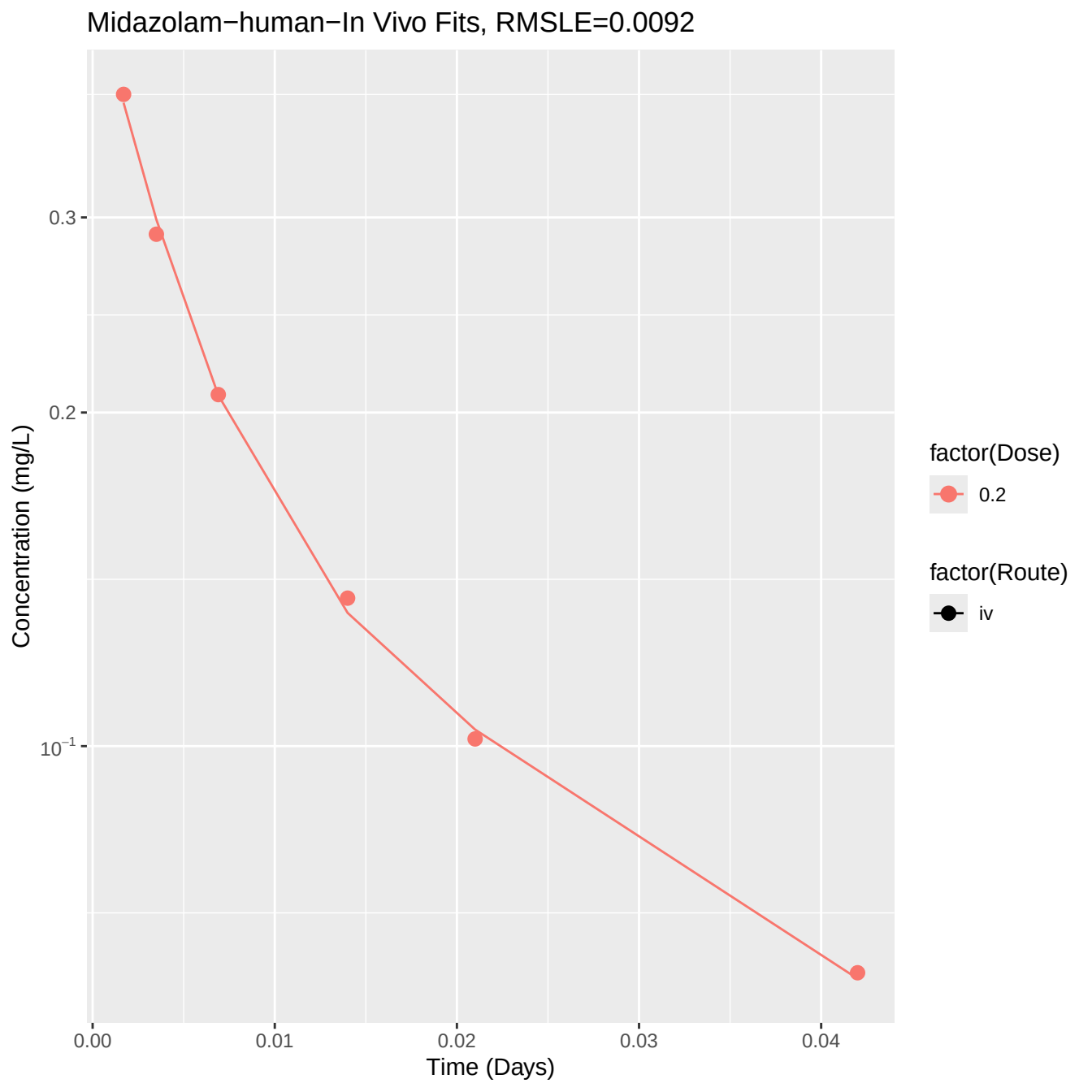

Midazolam−human−HTPBTK−InVitro, RMSLE=0.262
Concentration (mg/L)
0.3
10^{-1}
0.05
0.00
0.01
0.02
0.03
0.04
Time (Days)
factor(Dose)
0.2
factor(Route)
iv

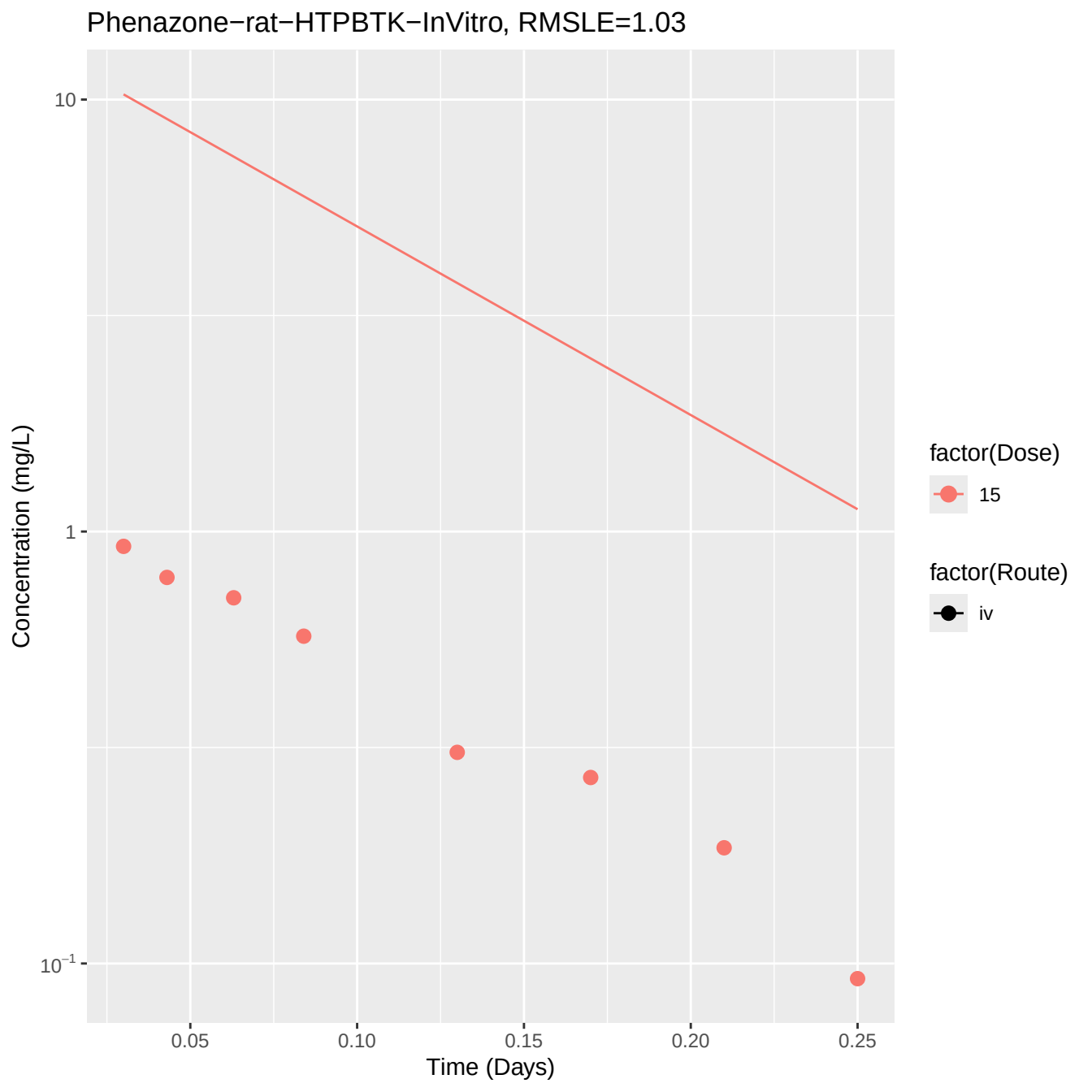

Midazolam−rat−HTPBTK−Dawson, RMSLE=1.51
Concentration (mg/L)
1
10^{-1}
10^{-2}
0.00
0.05
0.10
0.15
Time (Days)
factor(Route)
oral
factor(Dose)
21.6

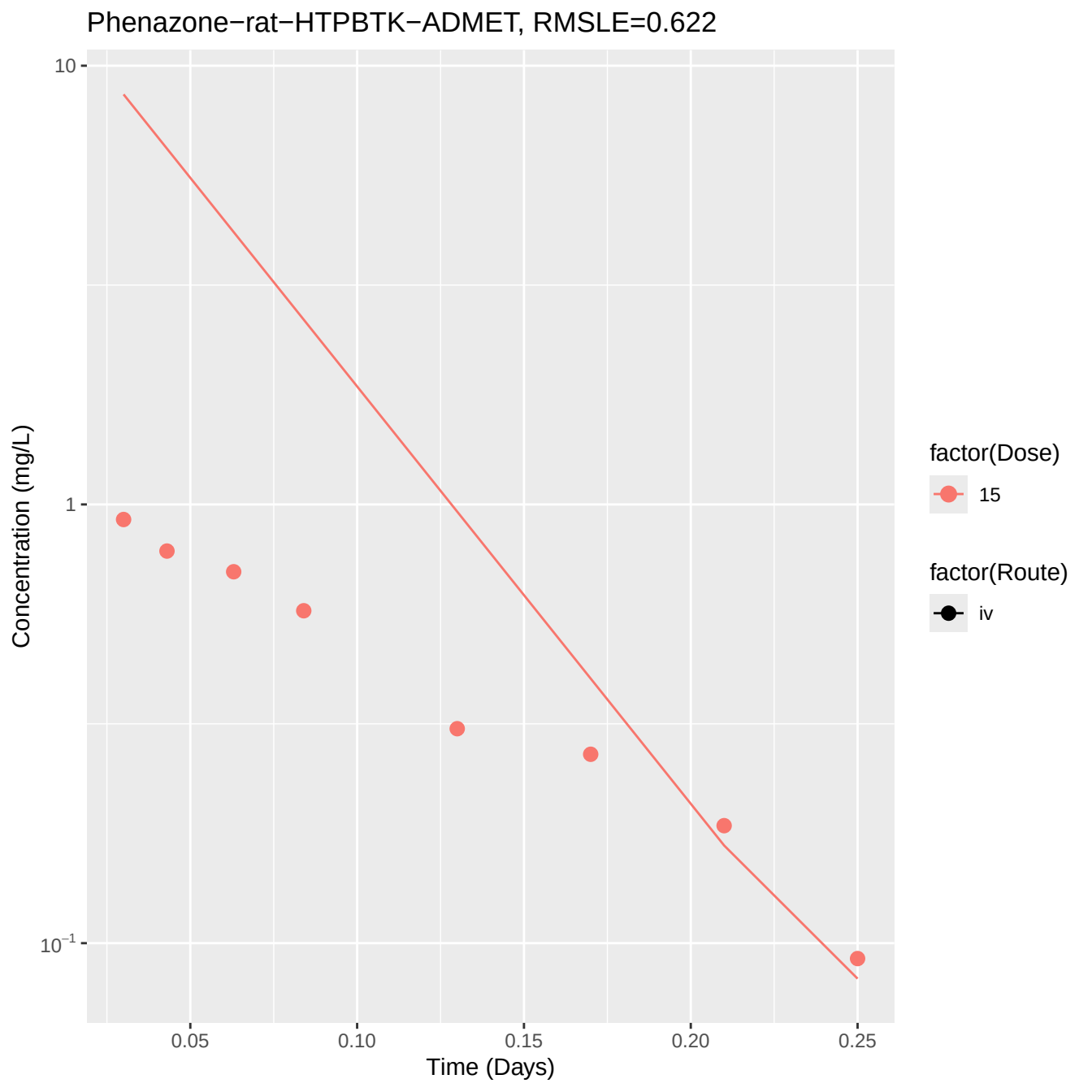

Midazolam−human−HTPBTK−Dawson, RMSLE=0.222
Concentration (mg/L)
0.3
10^{-1}
0.05
0.00
0.01
0.02
0.03
0.04
Time (Days)
factor(Dose)
0.2
factor(Route)
iv

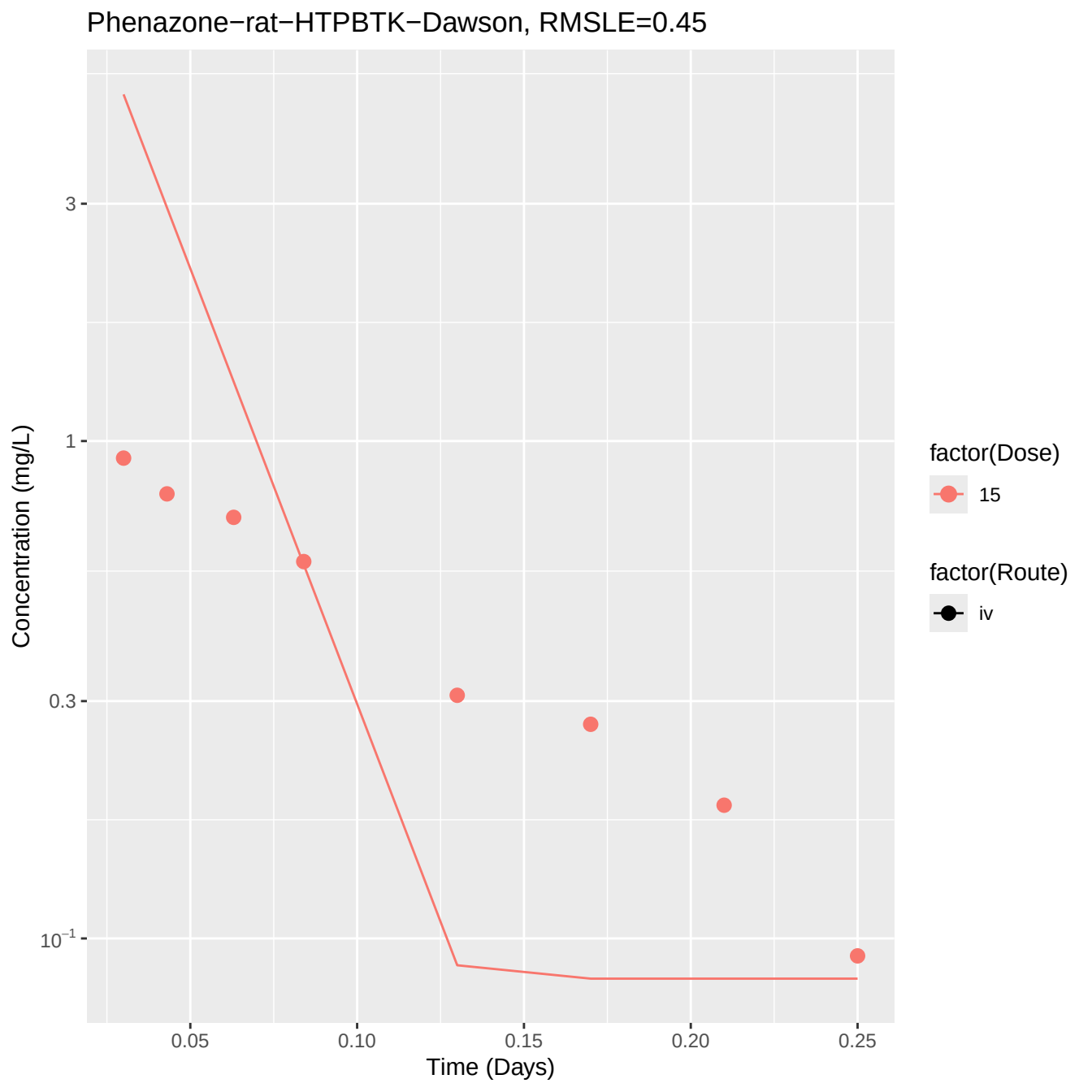

Midazolam−rat−HTPBTK−Ensemble, RMSLE=0.544
0.3
10^{-1}
0.03
10^{-2}
Concentration (mg/L)
0.00
0.05
0.10
0.15
Time (Days)
factor(Route)
oral
factor(Dose)
21.6

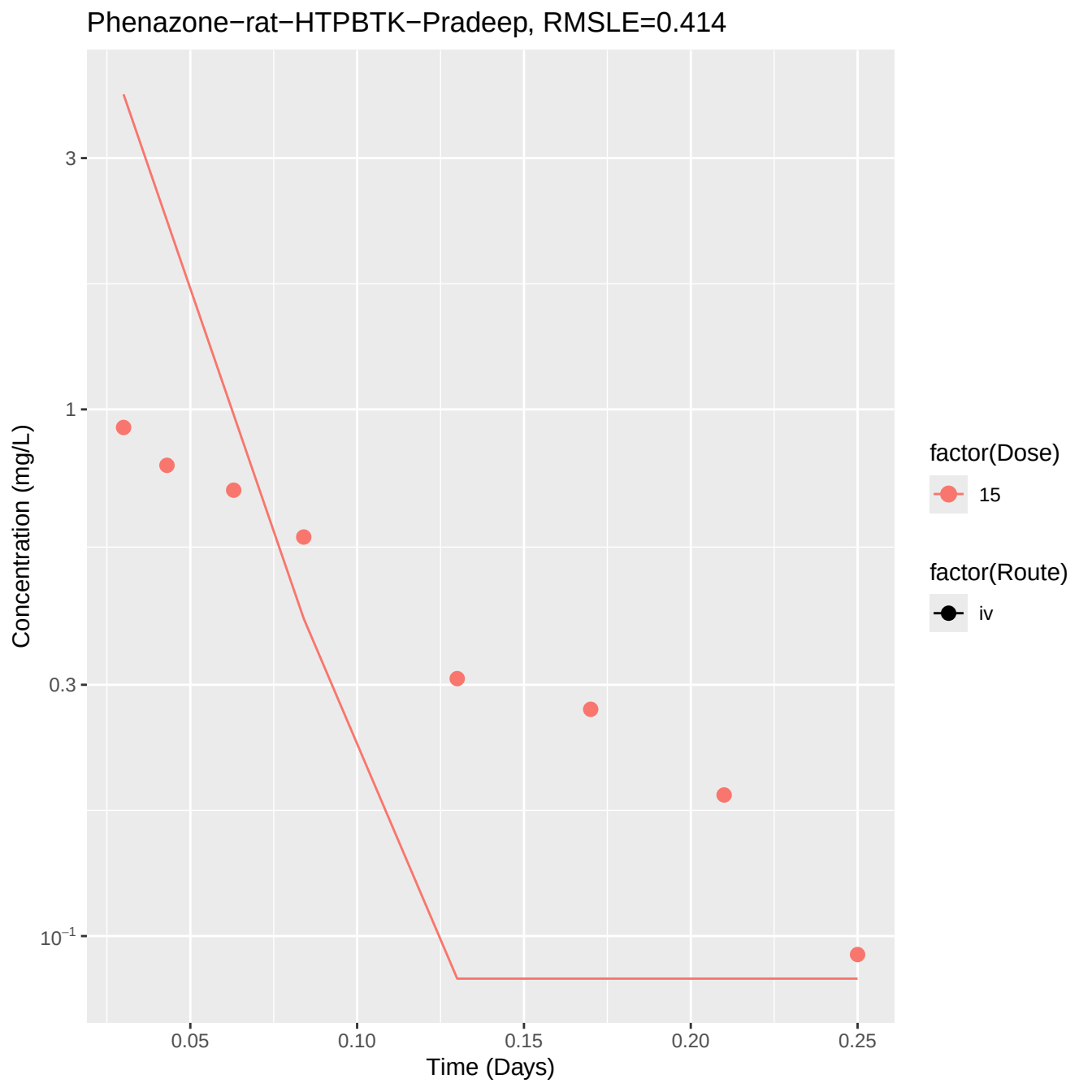

Midazolam−human−HTPBTK−Ensemble, RMSLE=0.366
Concentration (mg/L)
0.3
10^{-1}
0.05
0.00
0.01
0.02
0.03
0.04
Time (Days)
factor(Dose)
0.2
factor(Route)
iv

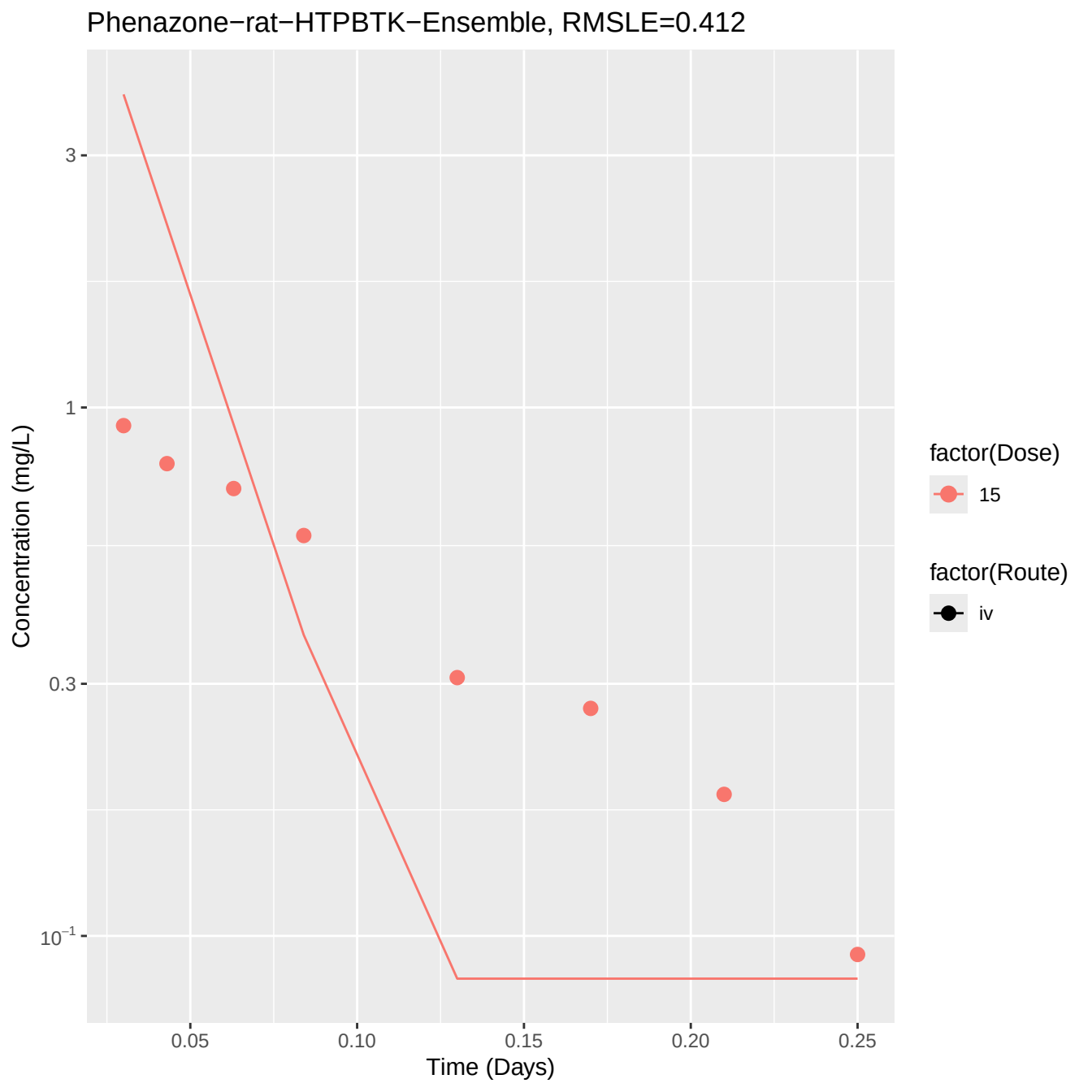

Midazolam−human−In Vivo Fits, RMSLE=0.0092
Concentration (mg/L)
0.3
0.2
10^{-1}
0.00
0.01
0.02
0.03
0.04
Time (Days)
factor(Dose)
0.2
factor(Route)
iv

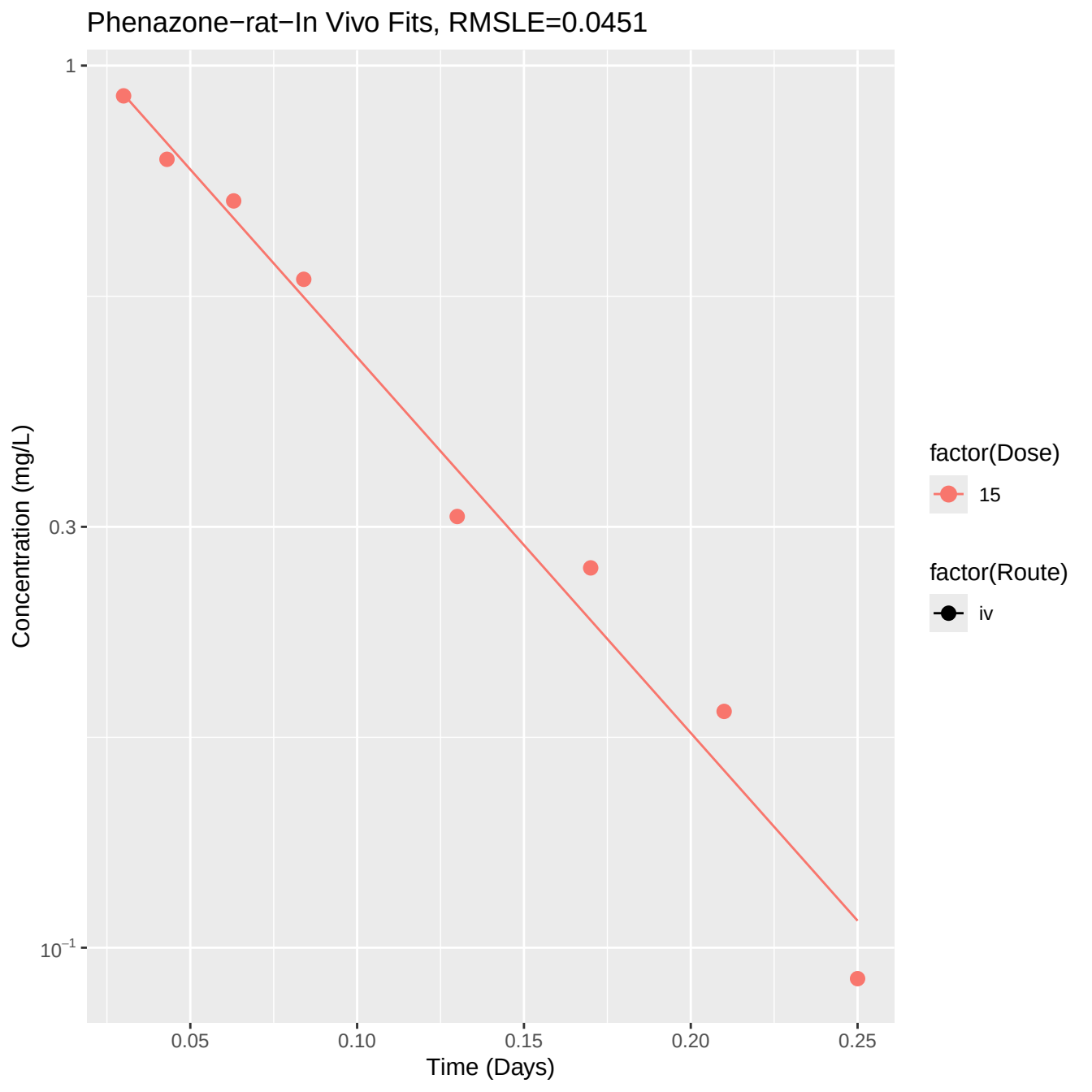

Phenazone−rat−HTPBTK−InVitro, RMSLE=1.03
10
1
10^{-1}
Concentration (mg/L)
0.05
0.10
0.15
0.20
0.25
Time (Days)
factor(Dose)
15
factor(Route)
iv

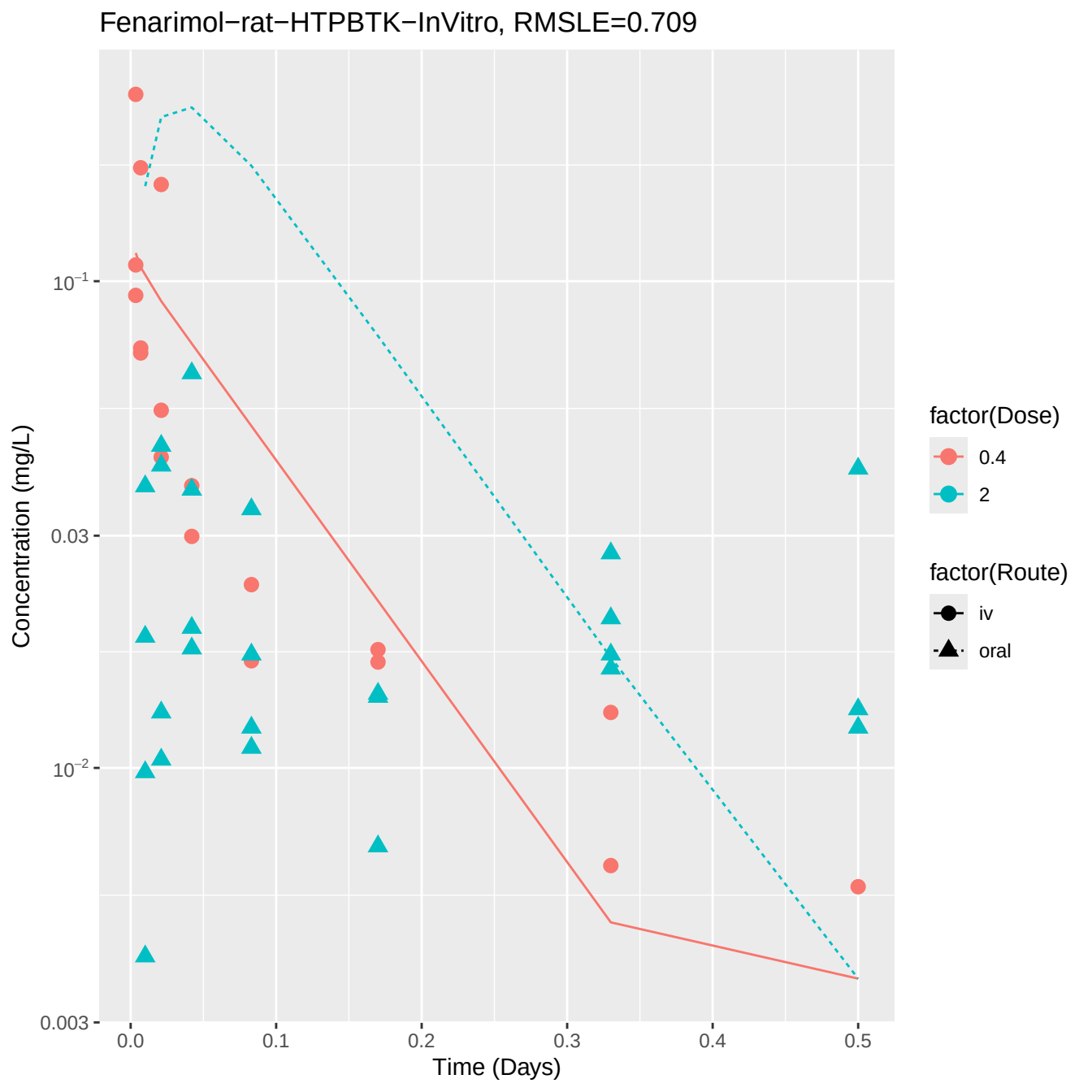

Phenazone−rat−HTPBTK−ADMET, RMSLE=0.622
10
1
10^{-1}
Concentration (mg/L)
0.05
0.10
0.15
0.20
0.25
Time (Days)
factor(Dose)
15
factor(Route)
iv

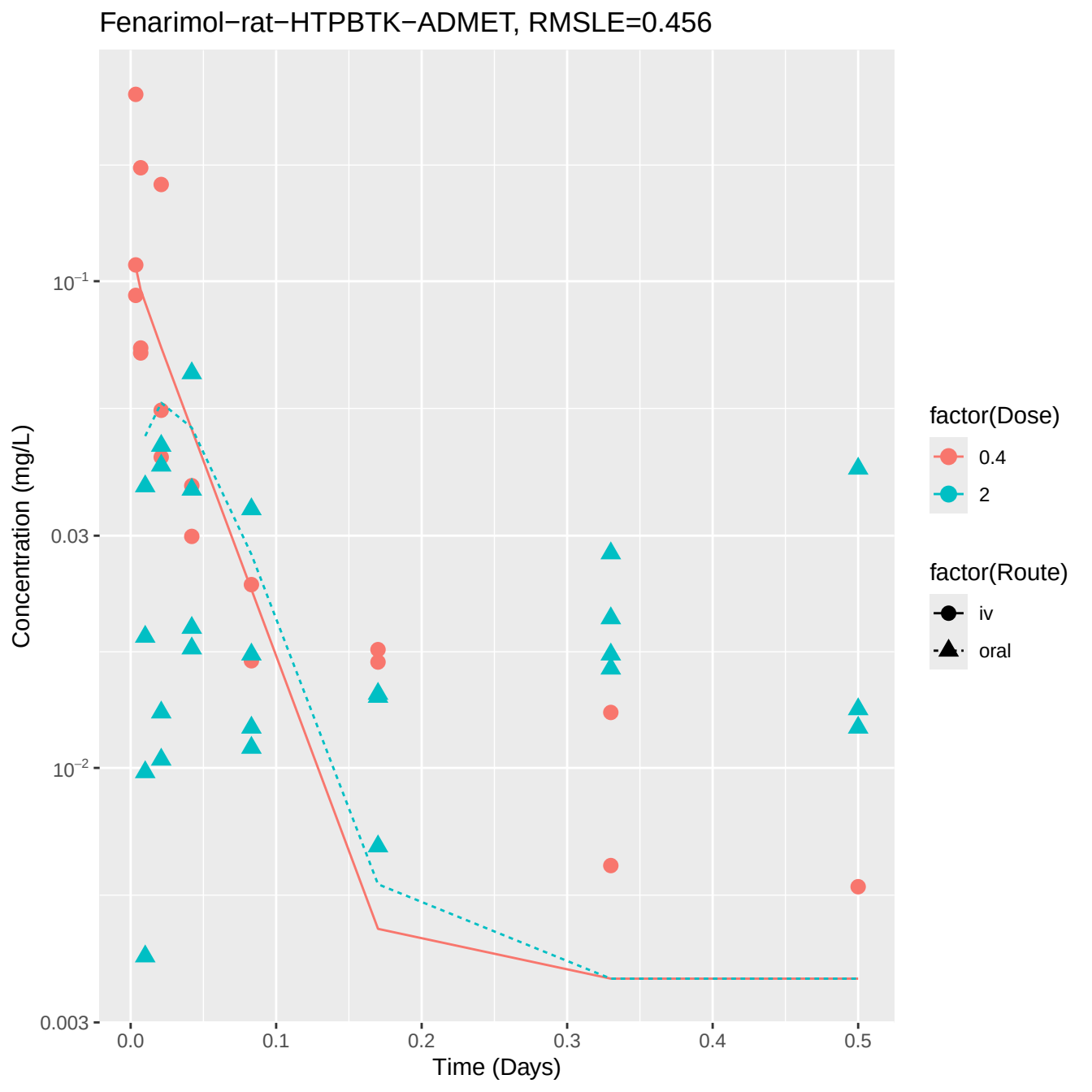

Phenazone−rat−HTPBTK−Dawson, RMSLE=0.45
Concentration (mg/L)
3
1
0.3
10^{-1}
0.05
0.10
0.15
0.20
0.25
Time (Days)
factor(Dose)
15
factor(Route)
iv

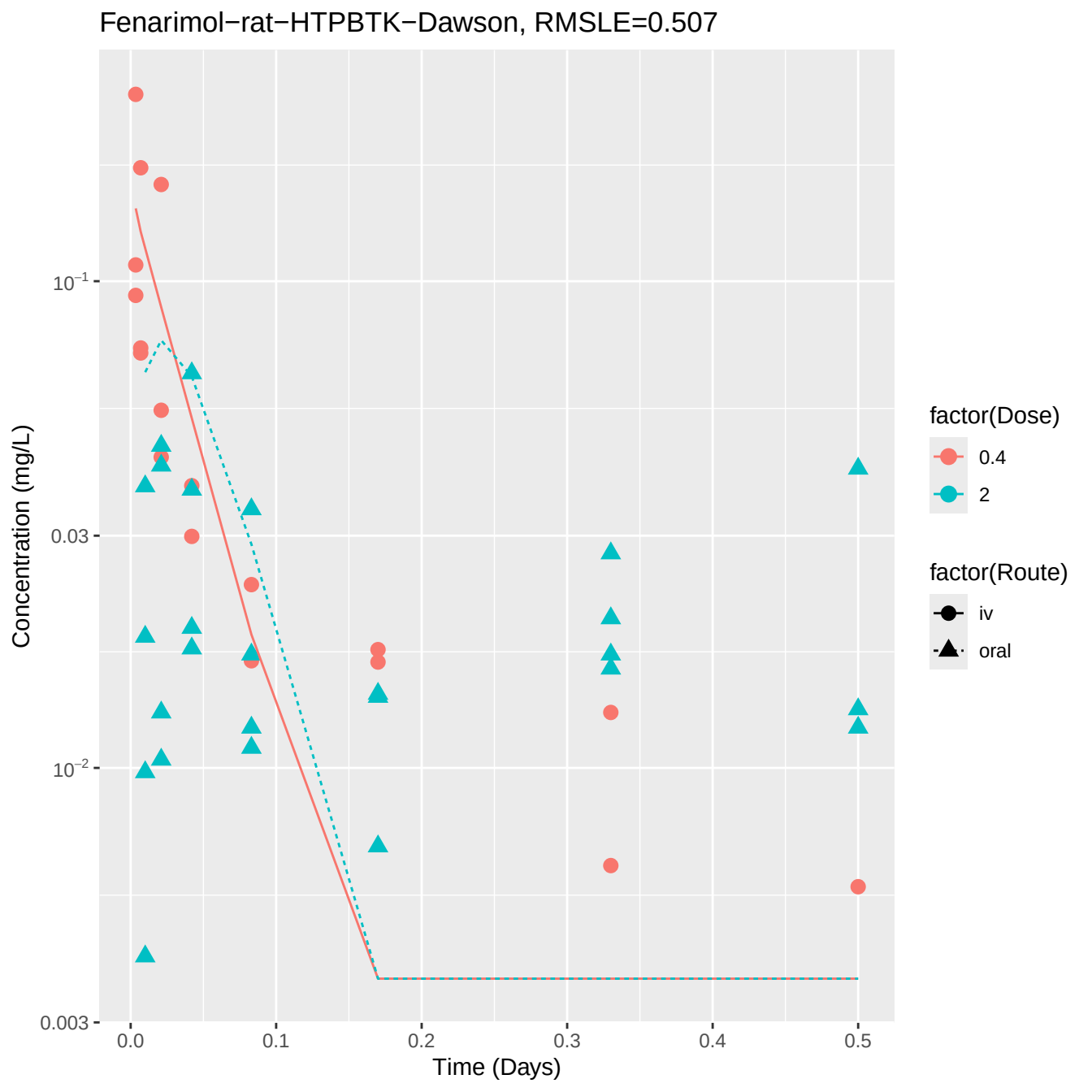

Phenazone−rat−HTPBTK−Pradeep, RMSLE=0.414
Concentration (mg/L)
3
1
0.3
10^{-1}
0.05
0.10
0.15
0.20
0.25
Time (Days)
factor(Dose)
15
factor(Route)
iv

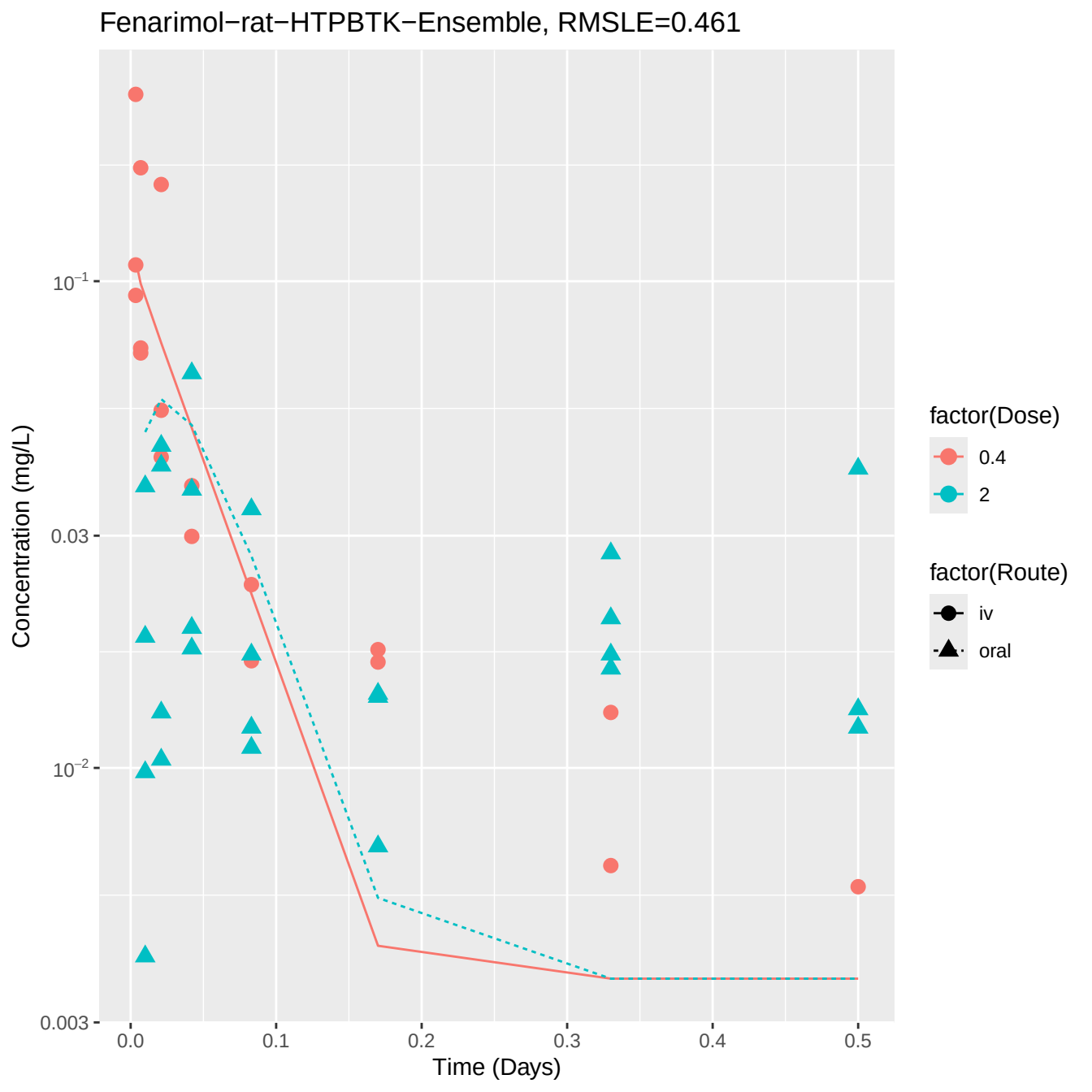

Phenazone−rat−HTPBTK−Ensemble, RMSLE=0.412
Concentration (mg/L)
3
1
0.3
10^{-1}
0.05
0.10
0.15
0.20
0.25
Time (Days)
factor(Dose)
15
factor(Route)
iv

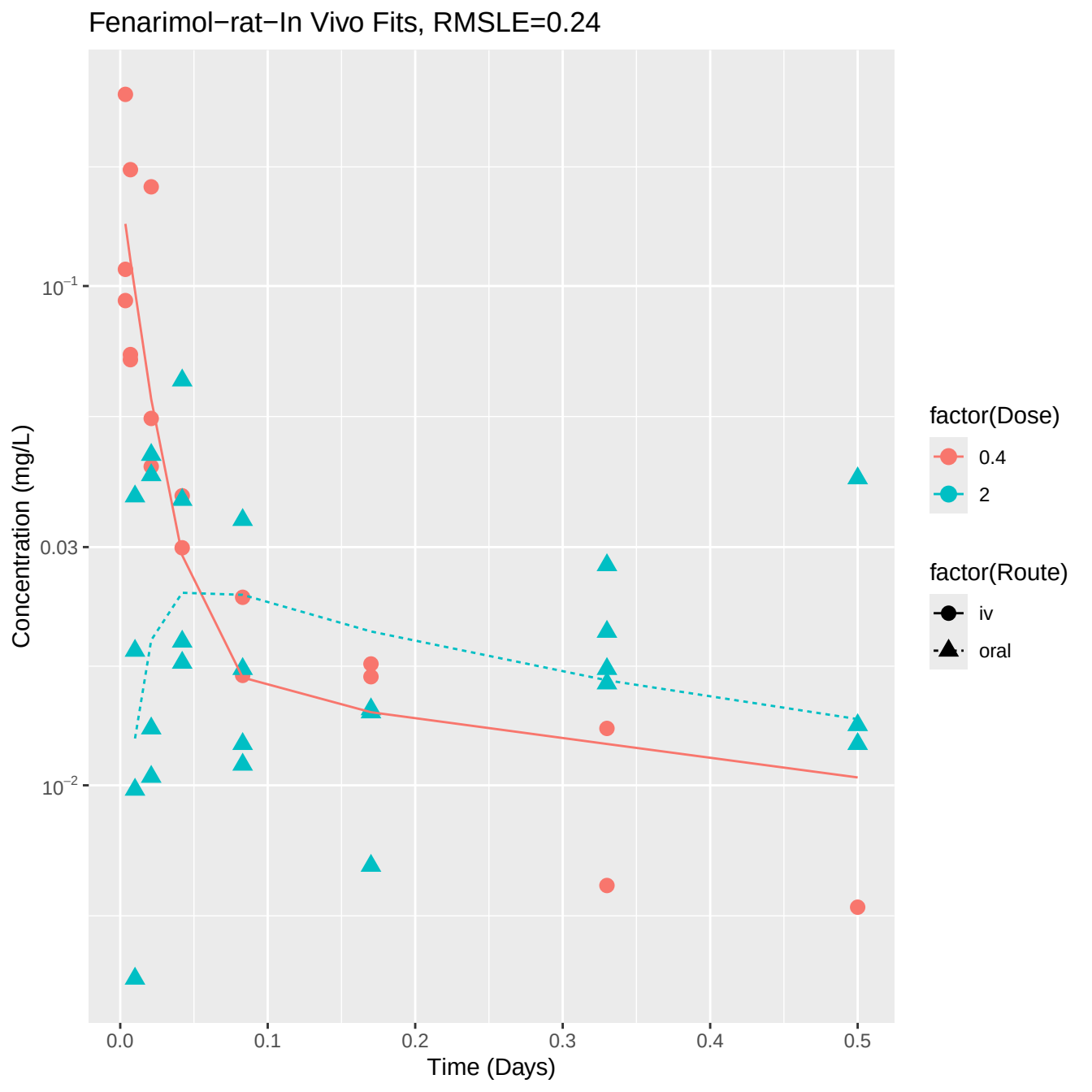

Phenazone−rat−In Vivo Fits, RMSLE=0.0451
Concentration (mg/L)
1
0.3
10^{-1}
0.05
0.10
0.15
0.20
0.25
Time (Days)
factor(Dose)
15
factor(Route)
iv

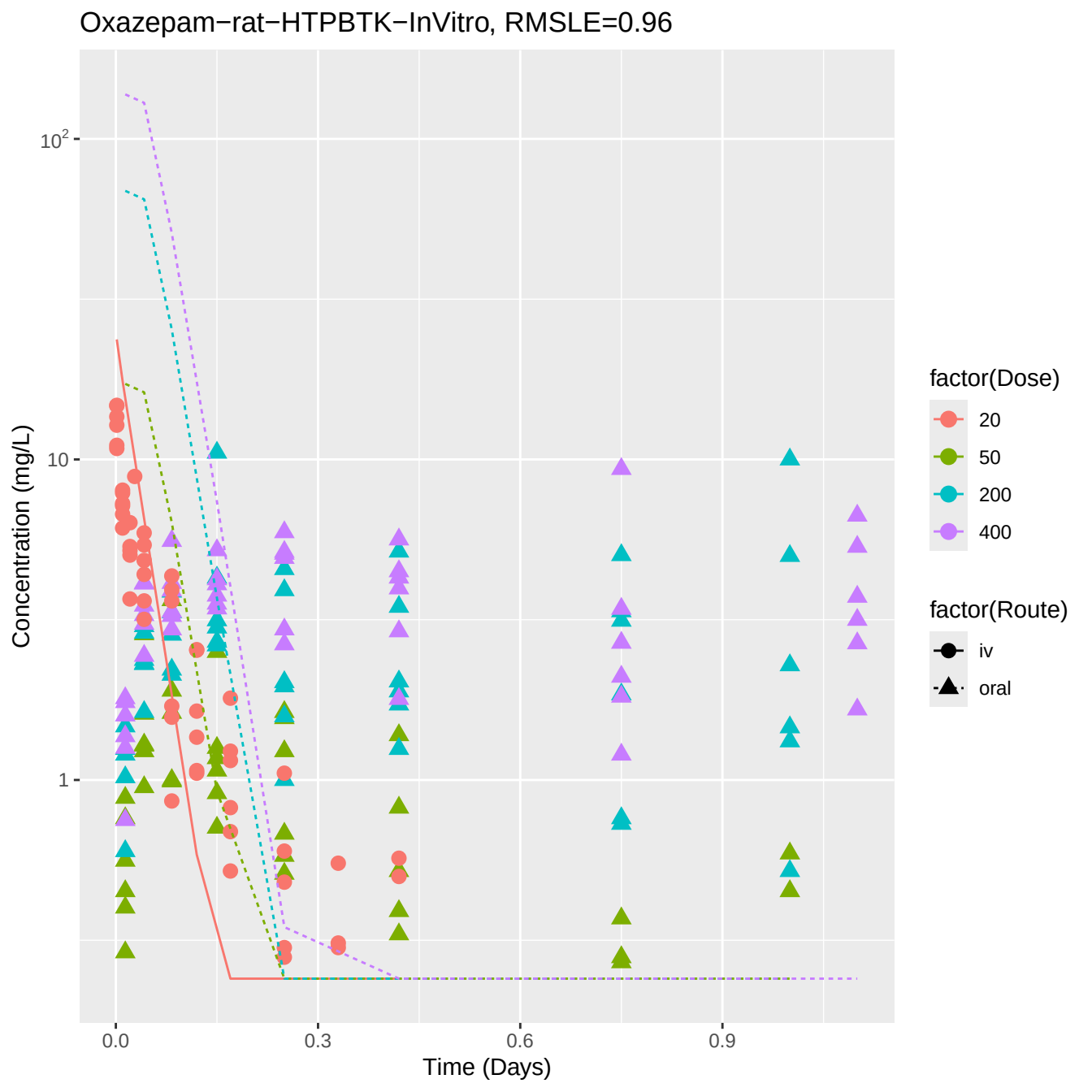

Fenarimol−rat−HTPBTK−InVitro, RMSLE=0.709
Concentration (mg/L)
10^{-1}
0.03
10^{-2}
0.003
0.0
0.1
0.2
0.3
0.4
0.5
Time (Days)
factor(Dose)
0.4
2
factor(Route)
iv
oral

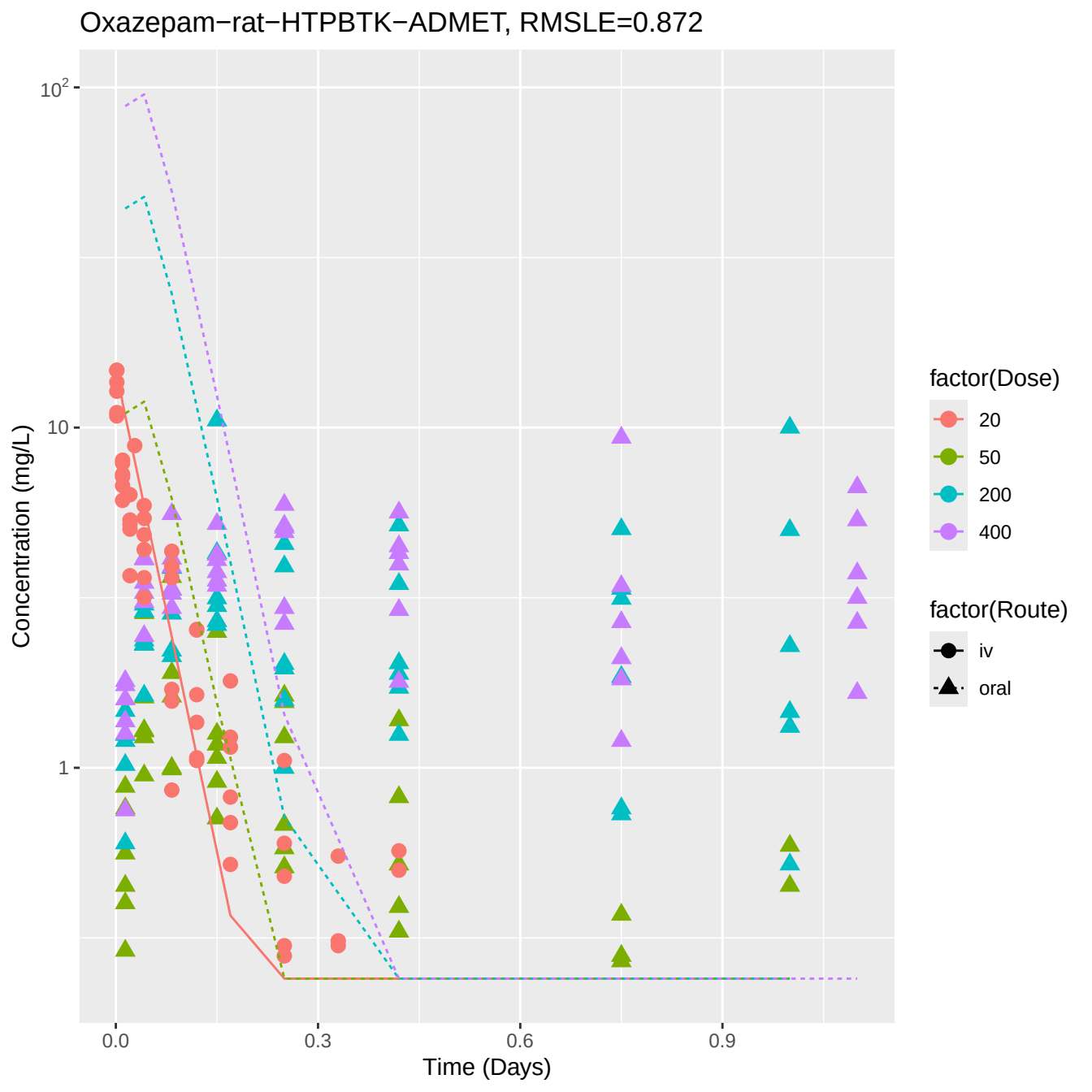

Fenarimol−rat−HTPBTK−ADMET, RMSLE=0.456
Concentration (mg/L)
10^−1
0.03
10^−2
0.003
0.0
0.1
0.2
0.3
0.4
0.5
Time (Days)
factor(Dose)
0.4
2
factor(Route)
iv
oral

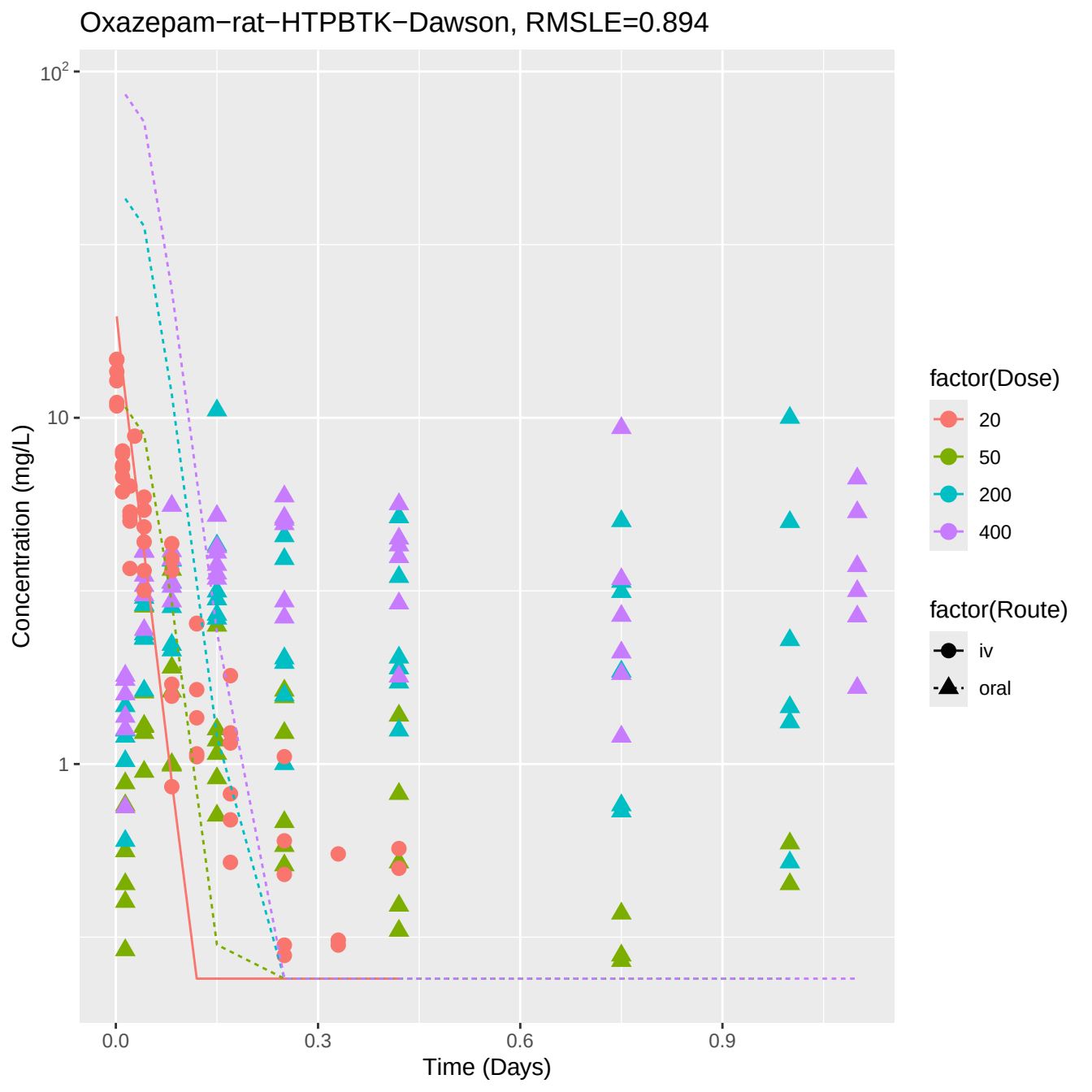

Fenarimol−rat−HTPBTK−Dawson, RMSLE=0.507
Concentration (mg/L)
10^{-1}
0.03
10^{-2}
0.003
0.0
0.1
0.2
0.3
0.4
0.5
Time (Days)
factor(Dose)
0.4
2
factor(Route)
iv
oral

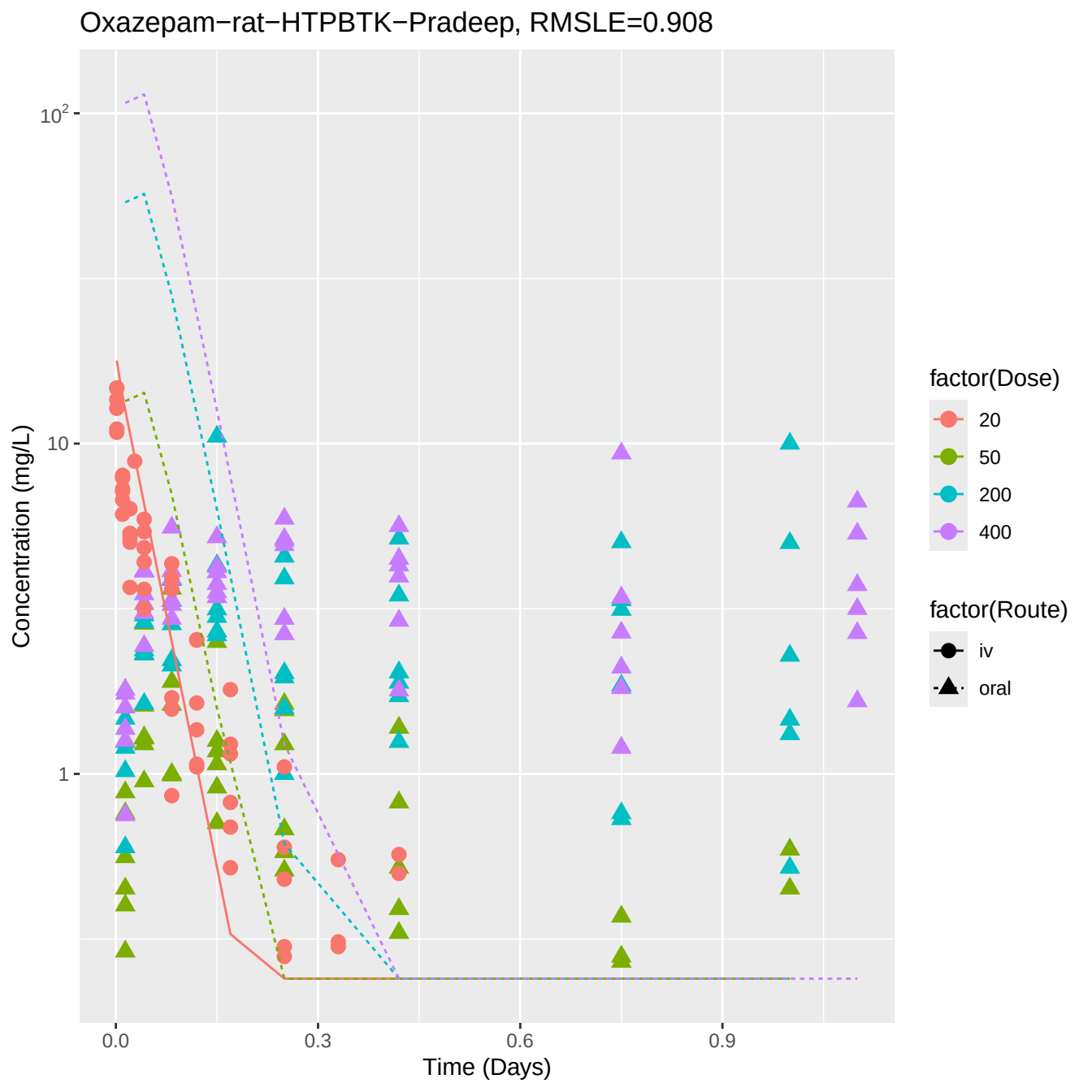

Fenarimol−rat−HTPBTK−Ensemble, RMSLE=0.461
Concentration (mg/L)
10^{-1}
0.03
10^{-2}
0.003
0.0
0.1
0.2
0.3
0.4
0.5
Time (Days)
factor(Dose)
0.4
2
factor(Route)
iv
oral

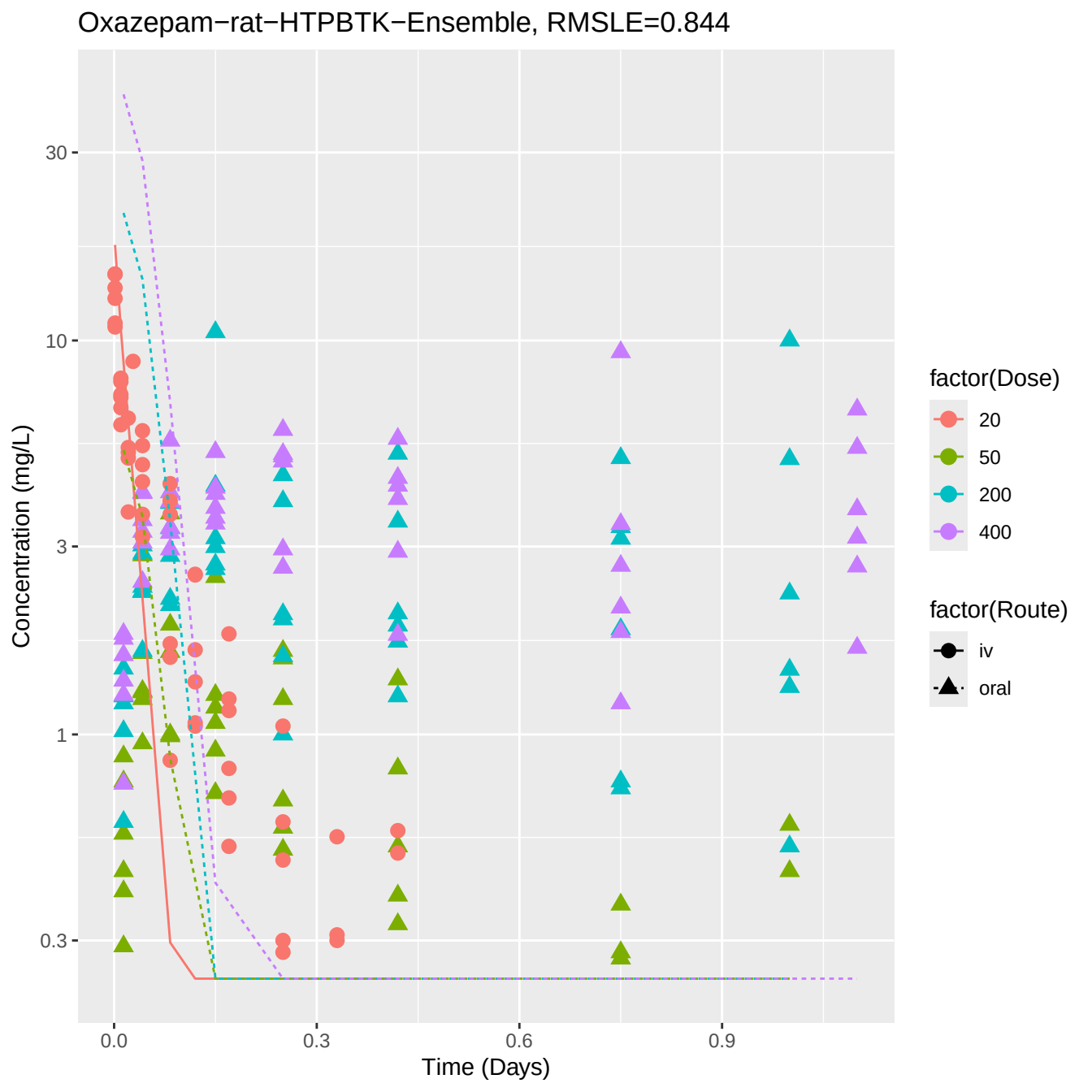

Fenarimol−rat−In Vivo Fits, RMSLE=0.24
Concentration (mg/L)
10^{-1}
0.03
10^{-2}
0.0
0.1
0.2
0.3
0.4
0.5
Time (Days)
factor(Dose)
0.4
2
factor(Route)
iv
oral

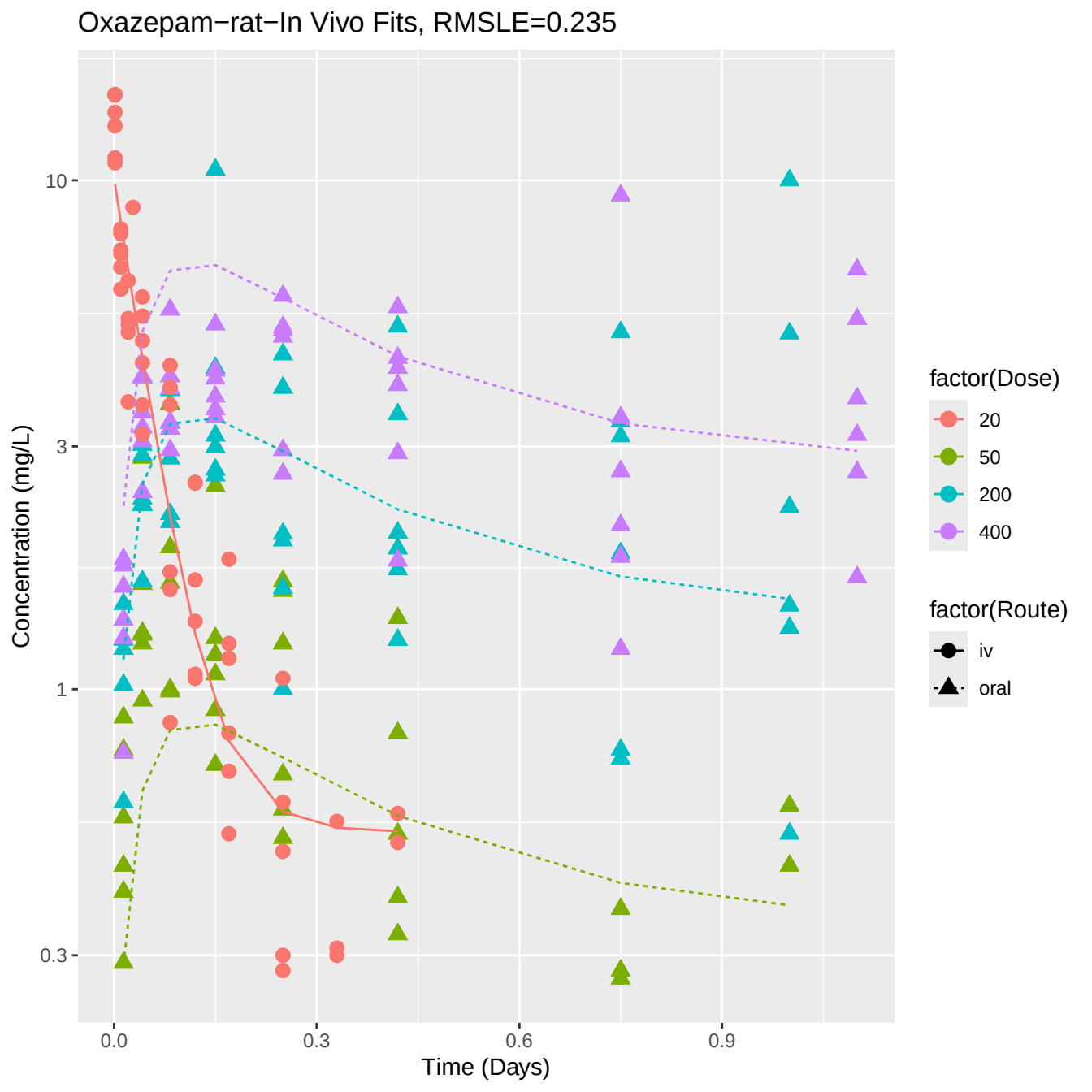

Oxazepam−rat−HTPBTK−InVitro, RMSLE=0.96
Concentration (mg/L)
10^2
10
1
0.0
0.3
0.6
0.9
Time (Days)
factor(Dose)
20
50
200
400
factor(Route)
iv
oral

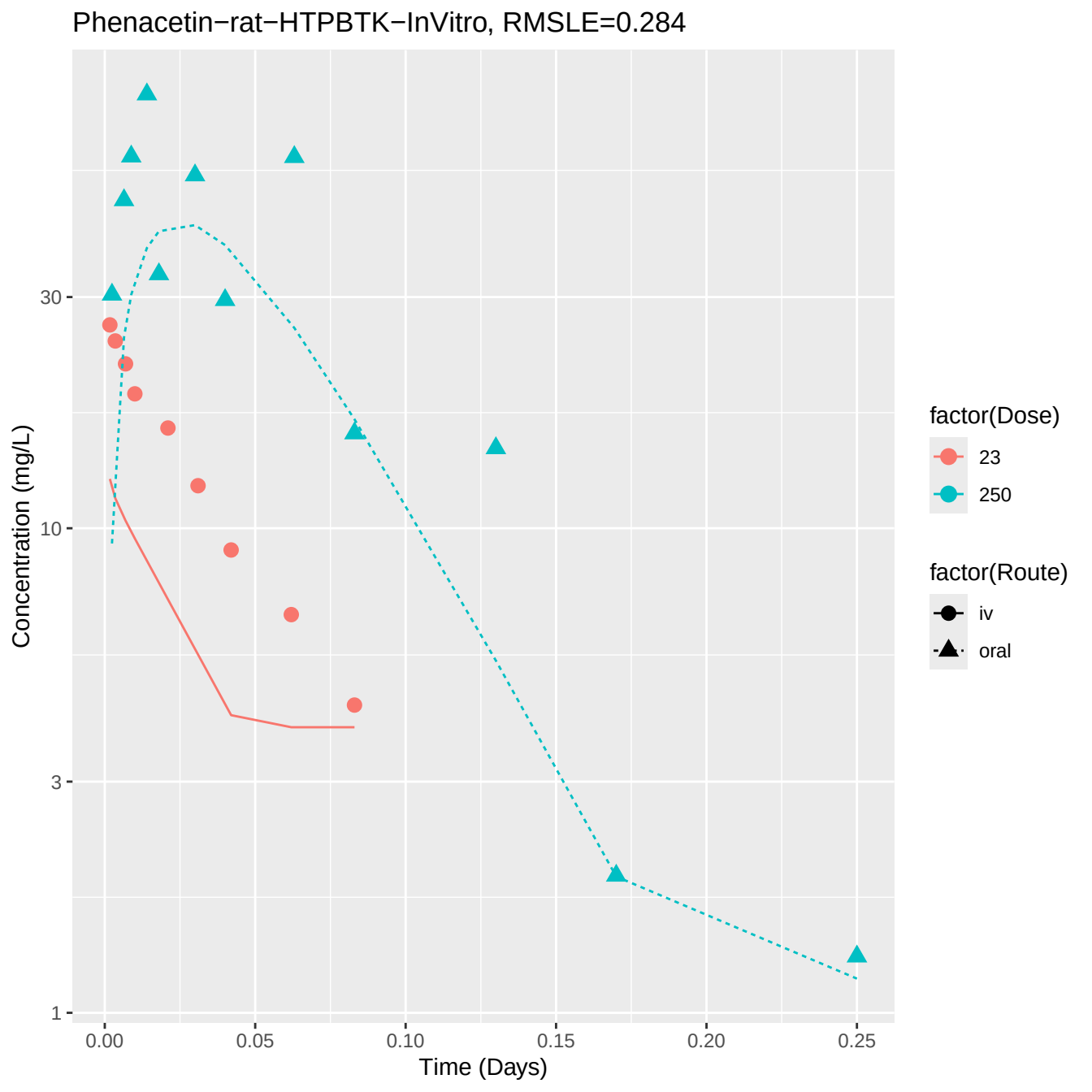

Oxazepam−rat−HTPBTK−ADMET, RMSLE=0.872
10^2
10
1
Concentration (mg/L)
0.0
0.3
0.6
0.9
Time (Days)
factor(Dose)
20
50
200
400
factor(Route)
iv
oral

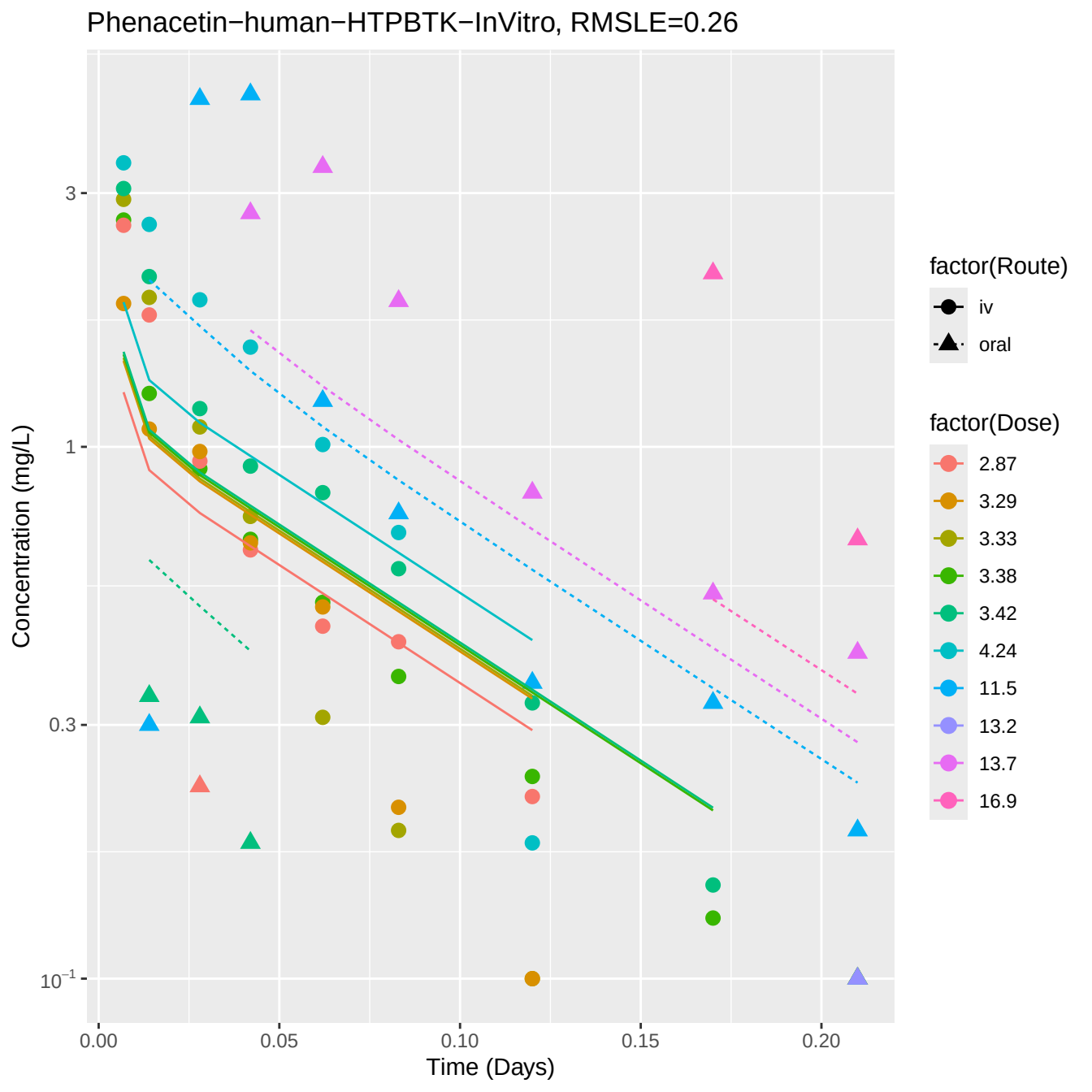

Oxazepam−rat−HTPBTK−Dawson, RMSLE=0.894
10^2
10
1
Concentration (mg/L)
0.0
0.3
0.6
0.9
Time (Days)
factor(Dose)
20
50
200
400
factor(Route)
iv
oral

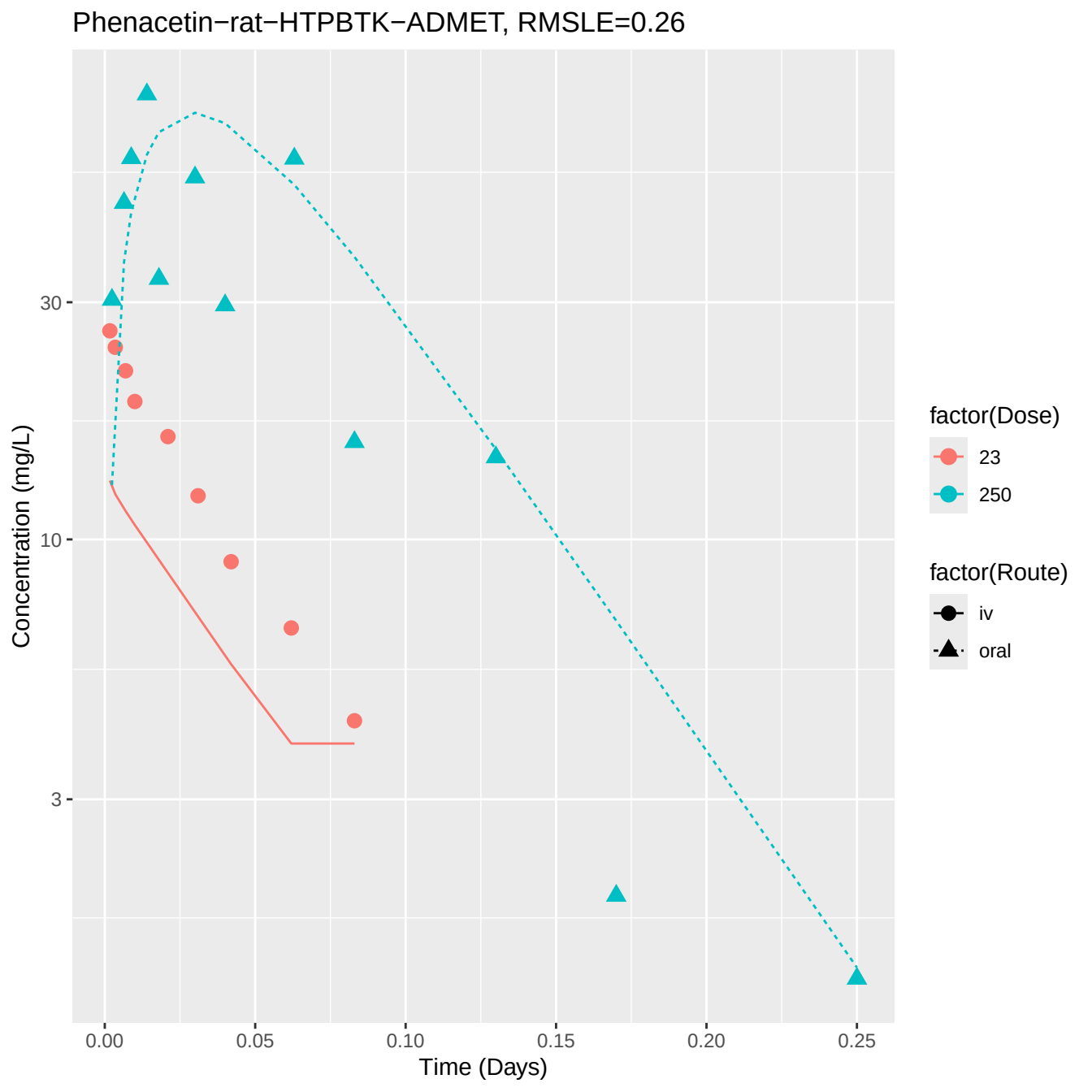

Oxazepam−rat−HTPBTK−Pradeep, RMSLE=0.908
Concentration (mg/L)
10^2
10
1
0.0
0.3
0.6
0.9
Time (Days)
factor(Dose)
20
50
200
400
factor(Route)
iv
oral

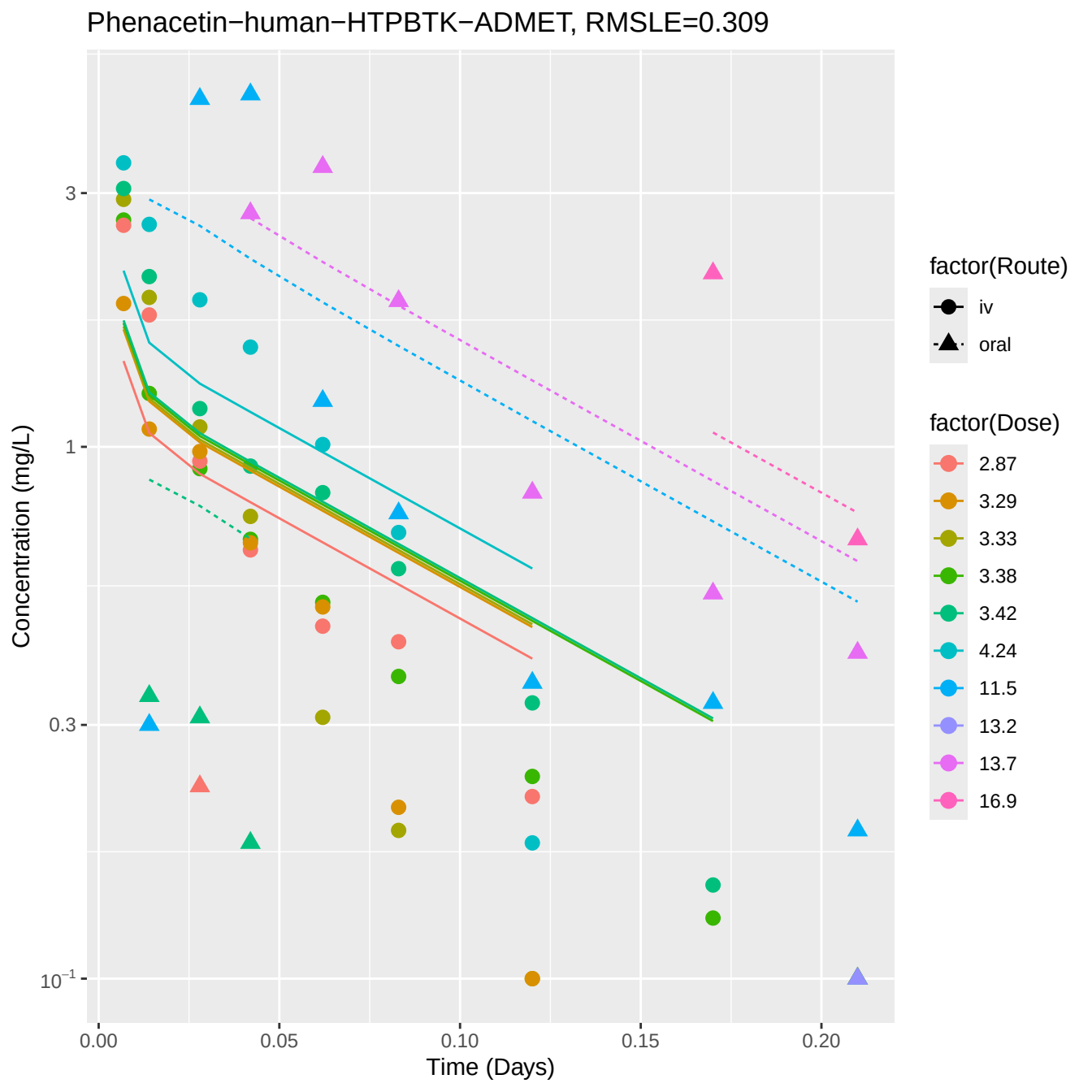

Oxazepam−rat−HTPBTK−Ensemble, RMSLE=0.844
Concentration (mg/L)
30
10
3
1
0.3
0.0
0.3
0.6
0.9
Time (Days)
factor(Dose)
20
50
200
400
factor(Route)
iv
oral

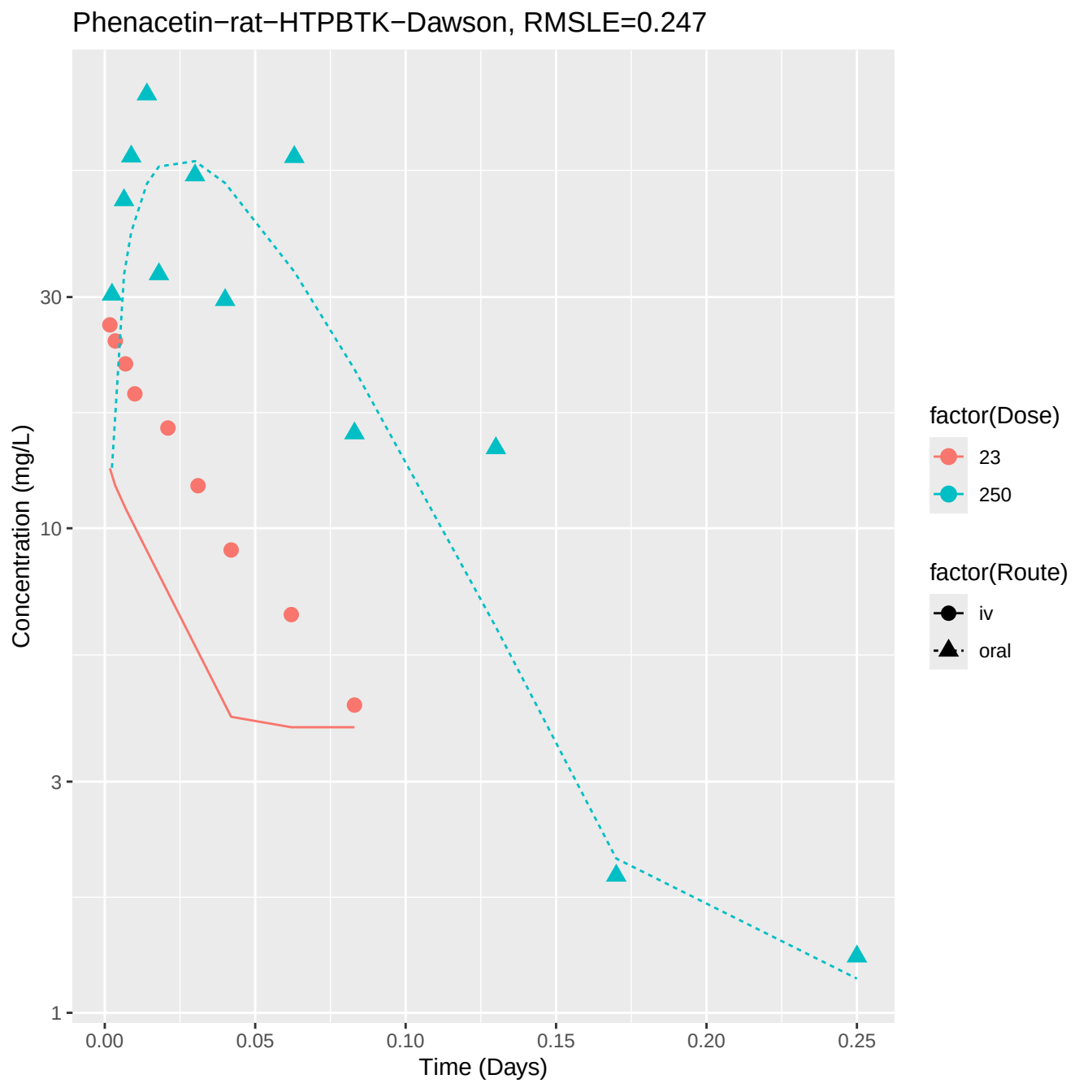

Oxazepam−rat−In Vivo Fits, RMSLE=0.235
Concentration (mg/L)
10
3
1
0.3
0.0
0.3
0.6
0.9
Time (Days)
factor(Dose)
20
50
200
400
factor(Route)
iv
oral

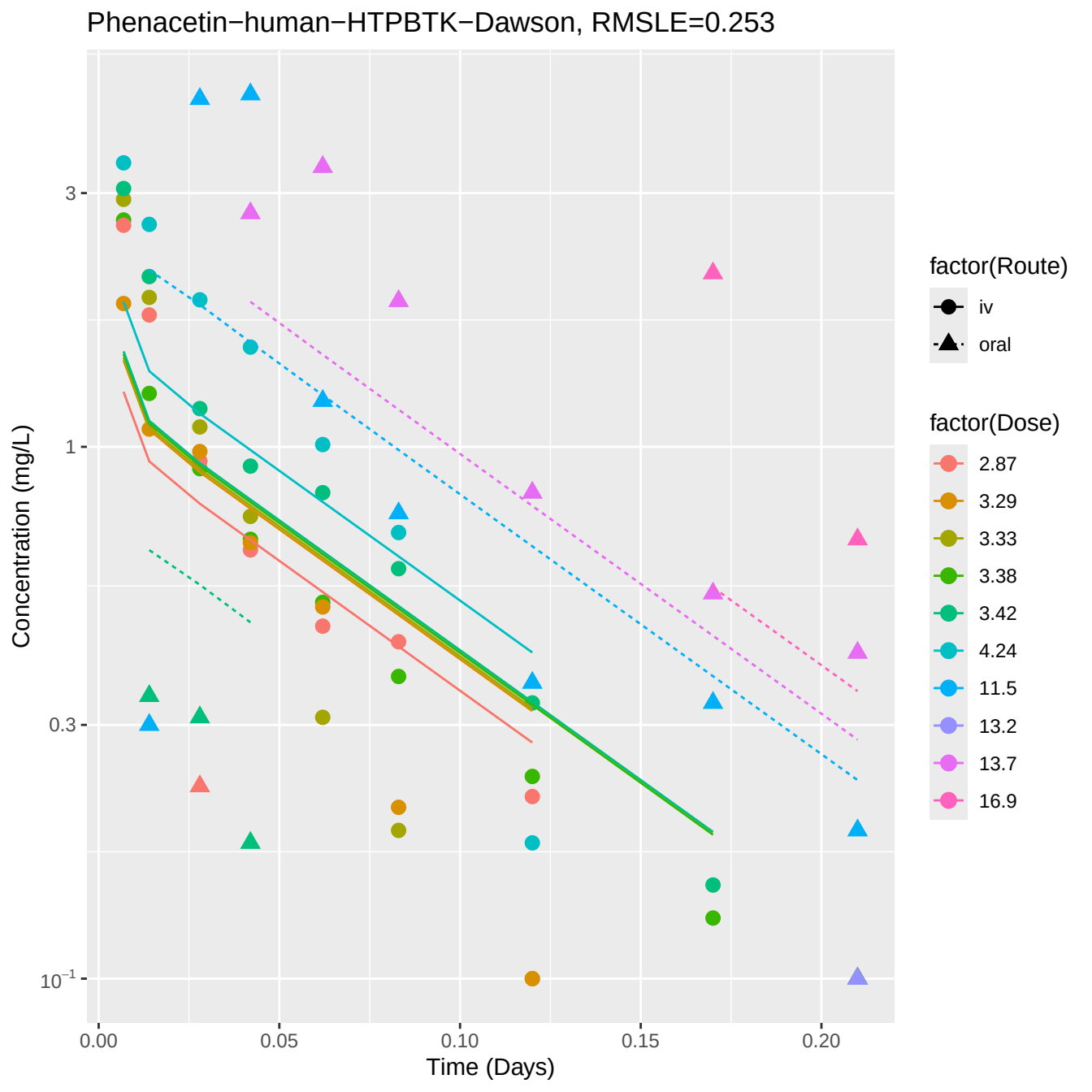

Phenacetin−rat−HTPBTK−InVitro, RMSLE=0.284
Concentration (mg/L)
30
10
3
1
0.00
0.05
0.10
0.15
0.20
0.25
Time (Days)
factor(Dose)
23
250
factor(Route)
iv
oral

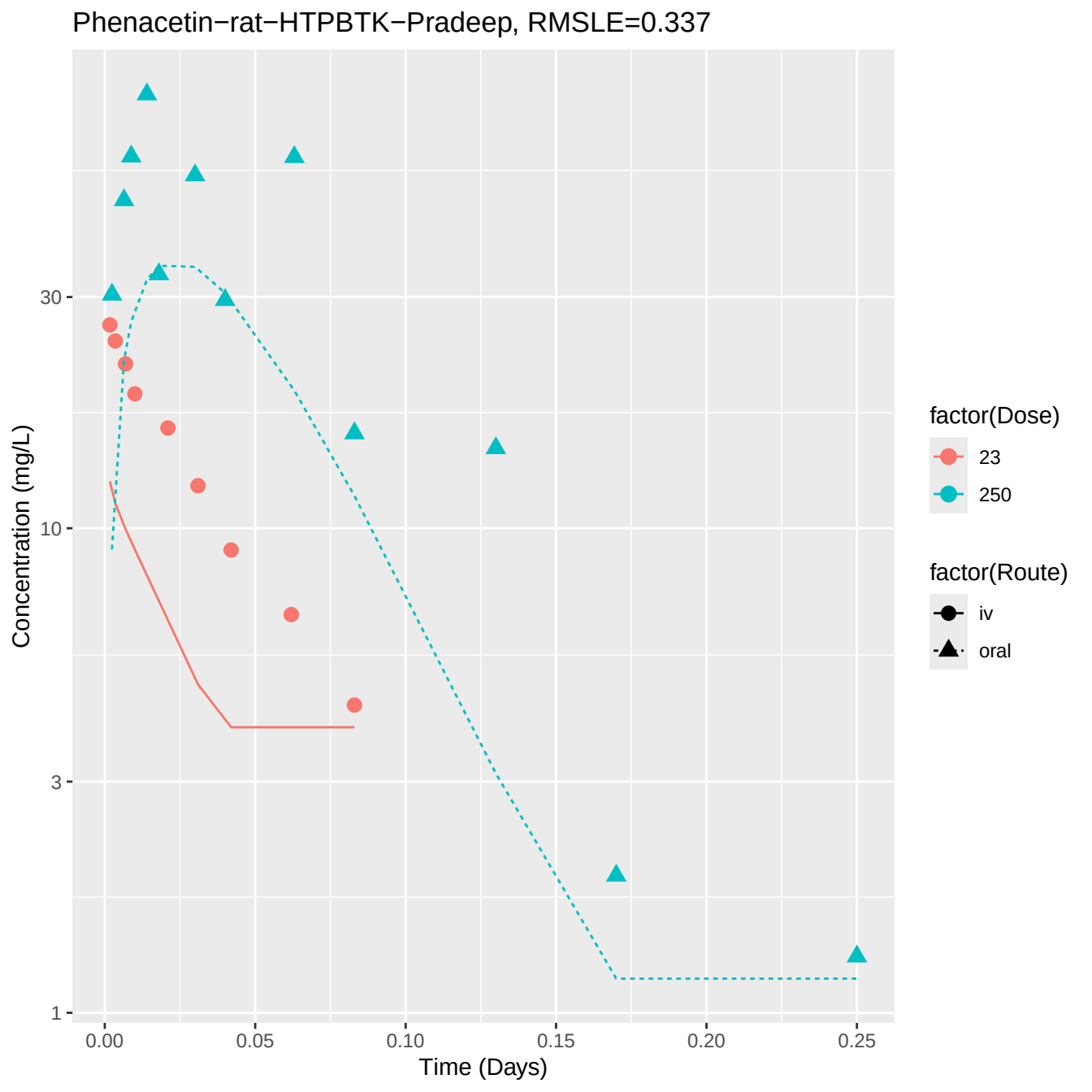

Phenacetin−human−HTPBTK−InVitro, RMSLE=0.26
Concentration (mg/L)
3
1
0.3
10^{-1}
0.00
0.05
0.10
0.15
0.20
Time (Days)
factor(Route)
iv
oral
factor(Dose)
2.87
3.29
3.33
3.38
3.42
4.24
11.5
13.2
13.7
16.9

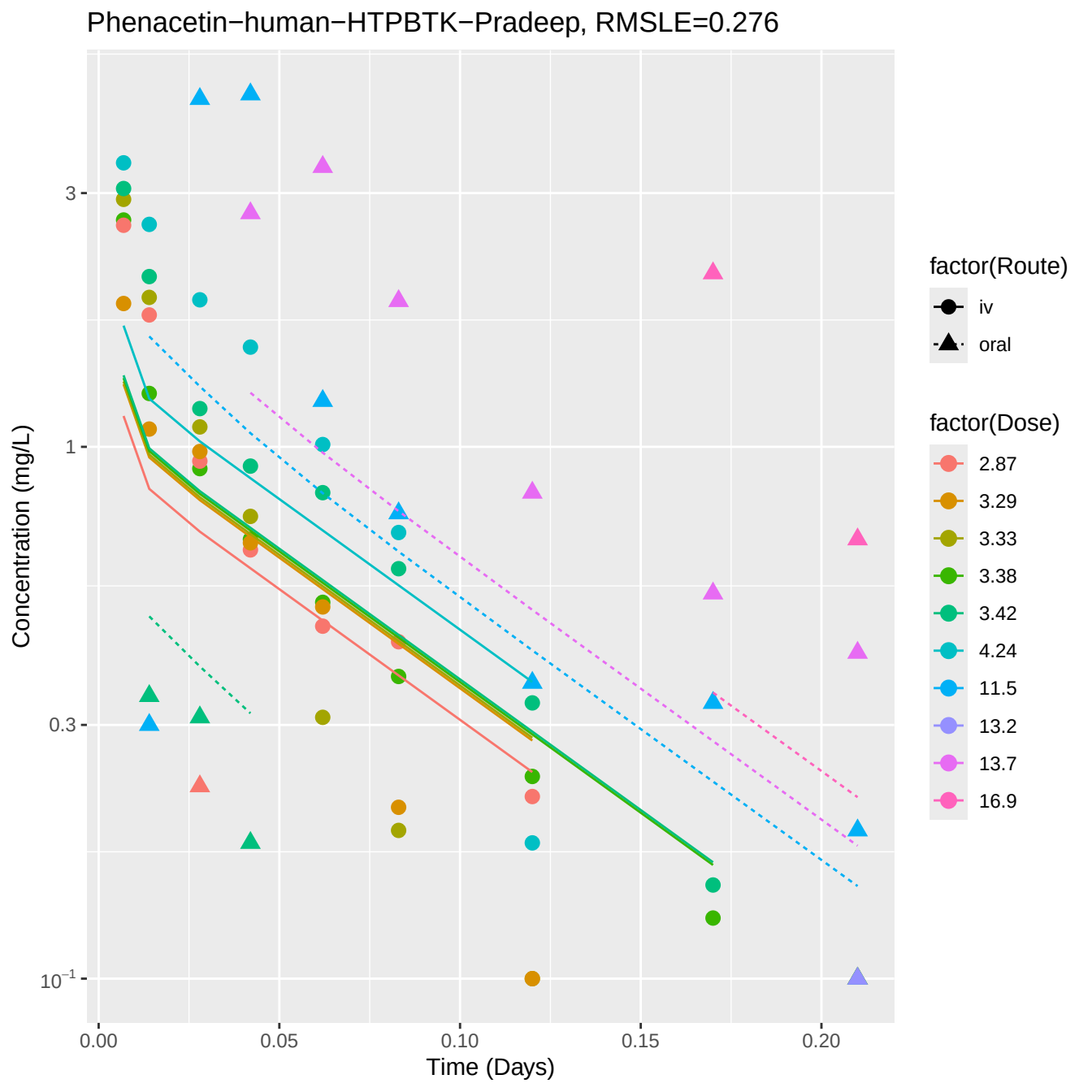

Phenacetin−rat−HTPBTK−ADMET, RMSLE=0.26
Concentration (mg/L)
30
10
3
0.00
0.05
0.10
0.15
0.20
0.25
Time (Days)
factor(Dose)
23
250
factor(Route)
iv
oral

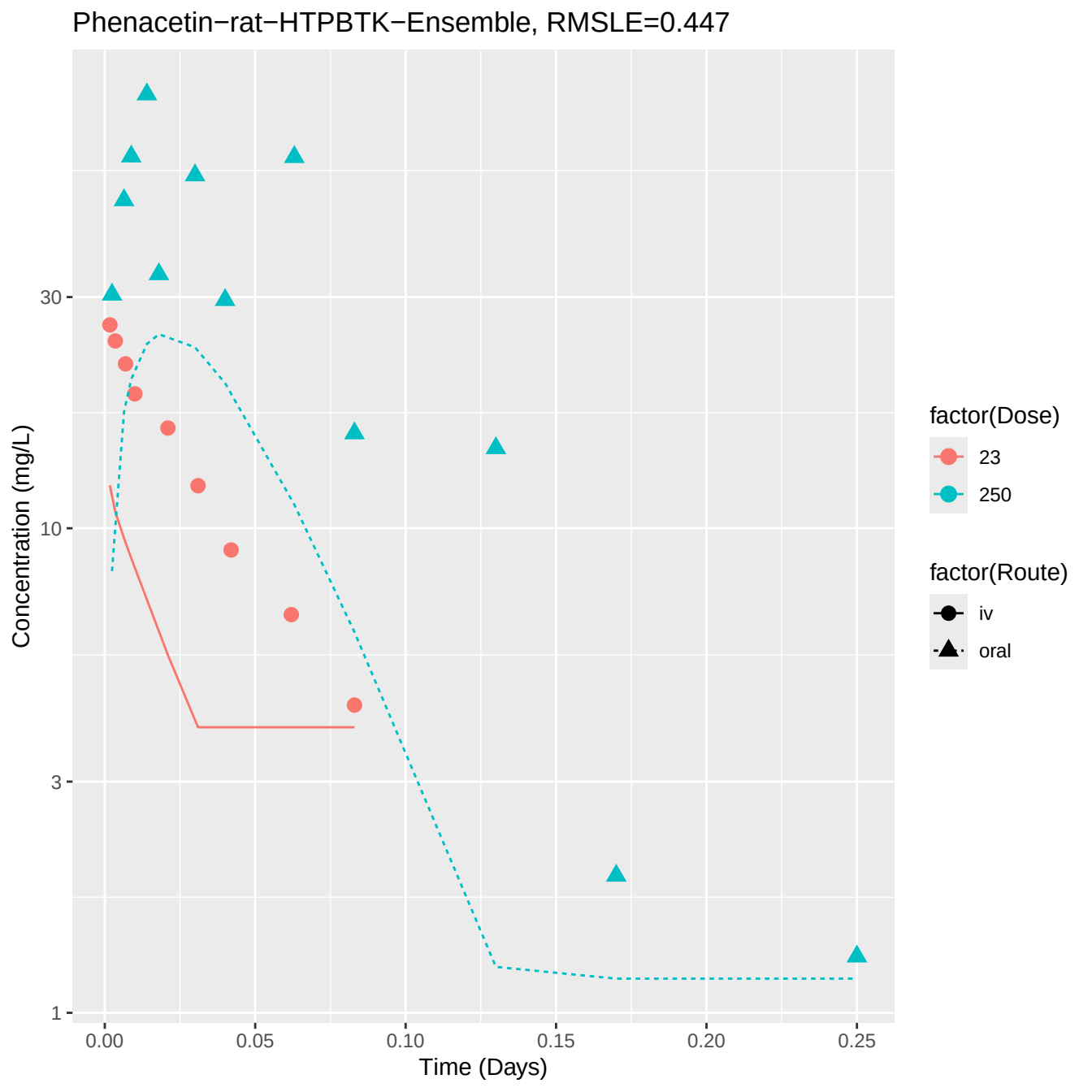

Phenacetin−human−HTPBTK−ADMET, RMSLE=0.309
Concentration (mg/L)
3
1
0.3
10^{-1}
0.00
0.05
0.10
0.15
0.20
Time (Days)
factor(Route)
iv
oral
factor(Dose)
2.87
3.29
3.33
3.38
3.42
4.24
11.5
13.2
13.7
16.9

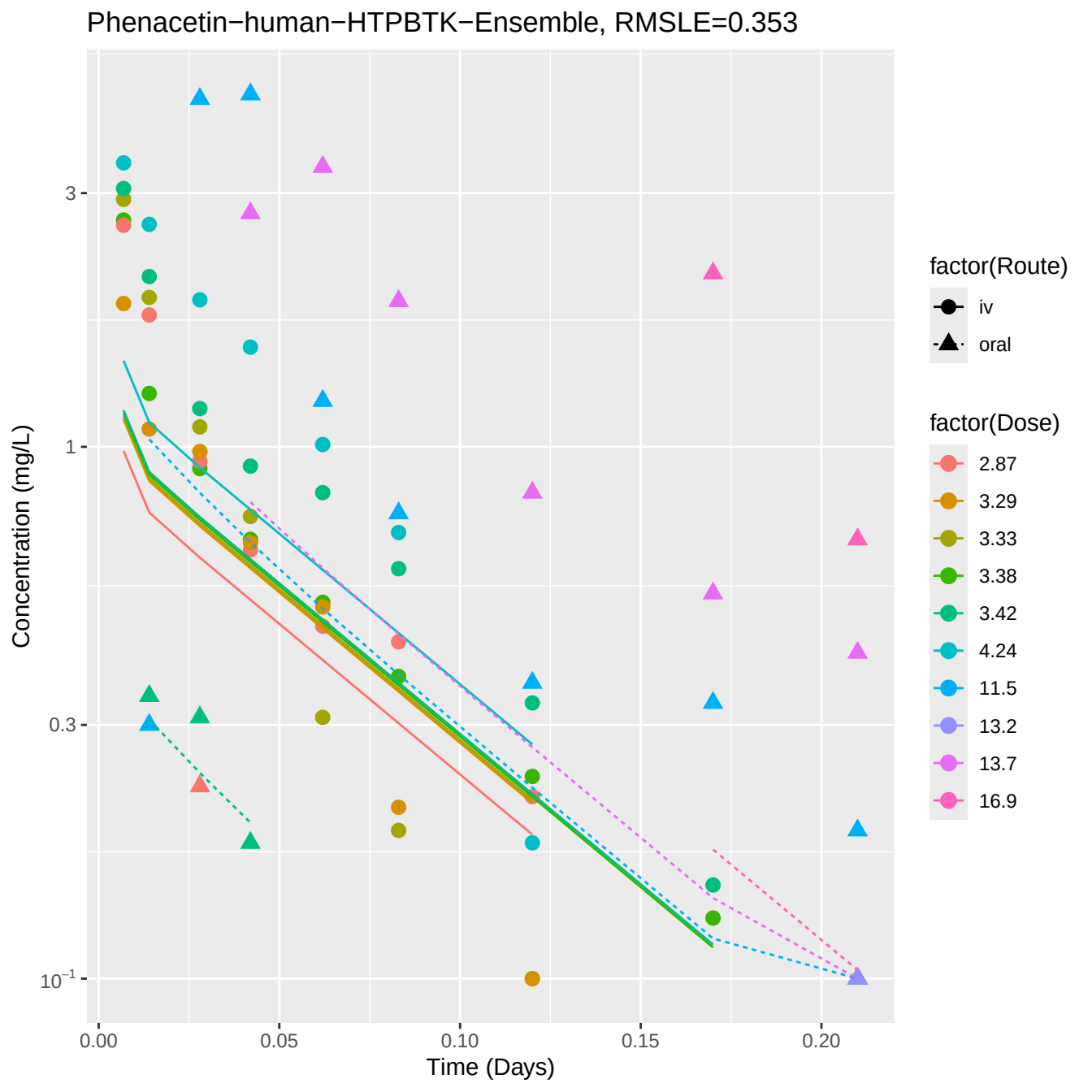

Phenacetin−rat−HTPBTK−Dawson, RMSLE=0.247
Concentration (mg/L)
30
10
3
1
0.00
0.05
0.10
0.15
0.20
0.25
Time (Days)
factor(Dose)
23
250
factor(Route)
iv
oral

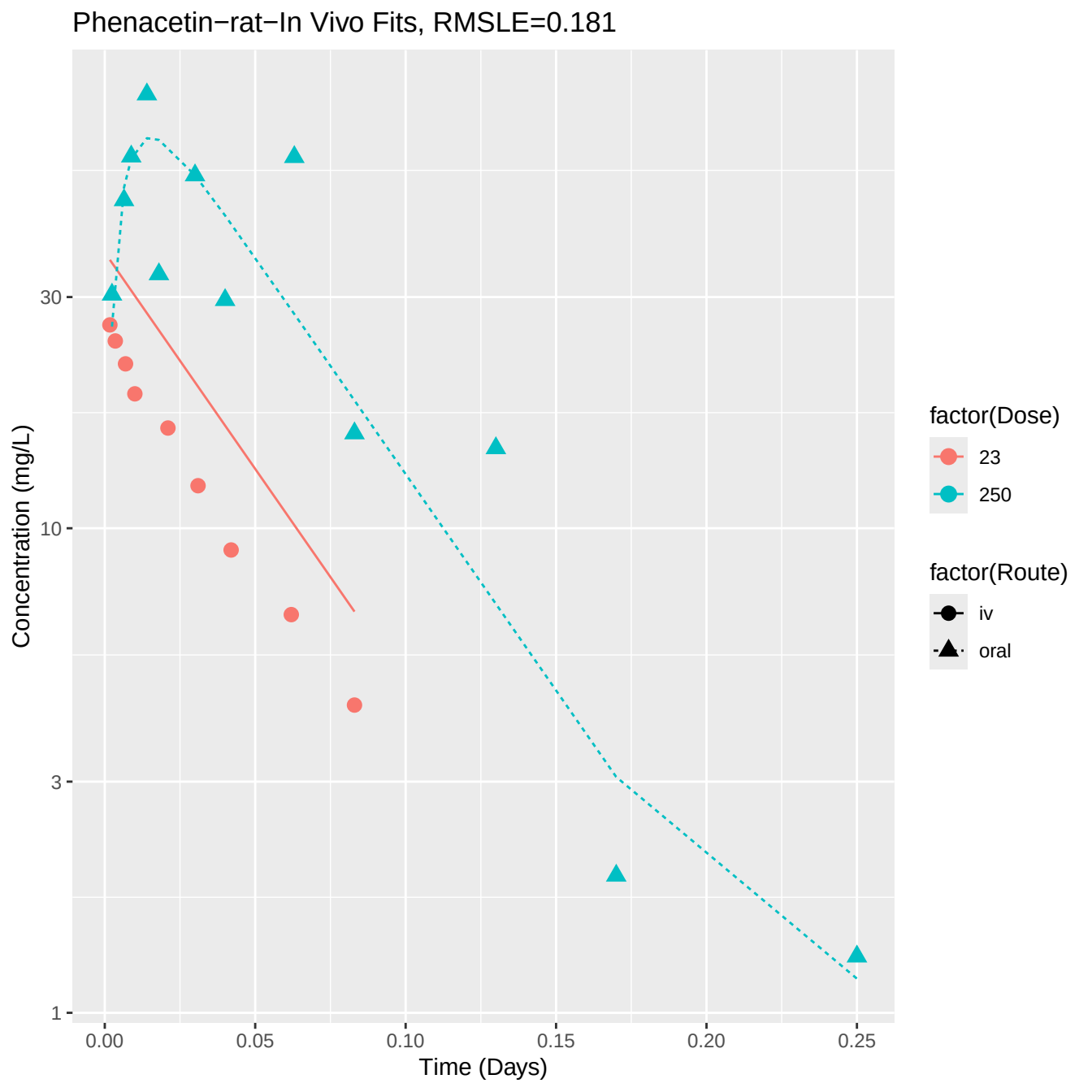

Phenacetin−human−HTPBTK−Dawson, RMSLE=0.253
Concentration (mg/L)
3
1
0.3
10^{-1}
0.00
0.05
0.10
0.15
0.20
Time (Days)
factor(Route)
iv
oral
factor(Dose)
2.87
3.29
3.33
3.38
3.42
4.24
11.5
13.2
13.7
16.9

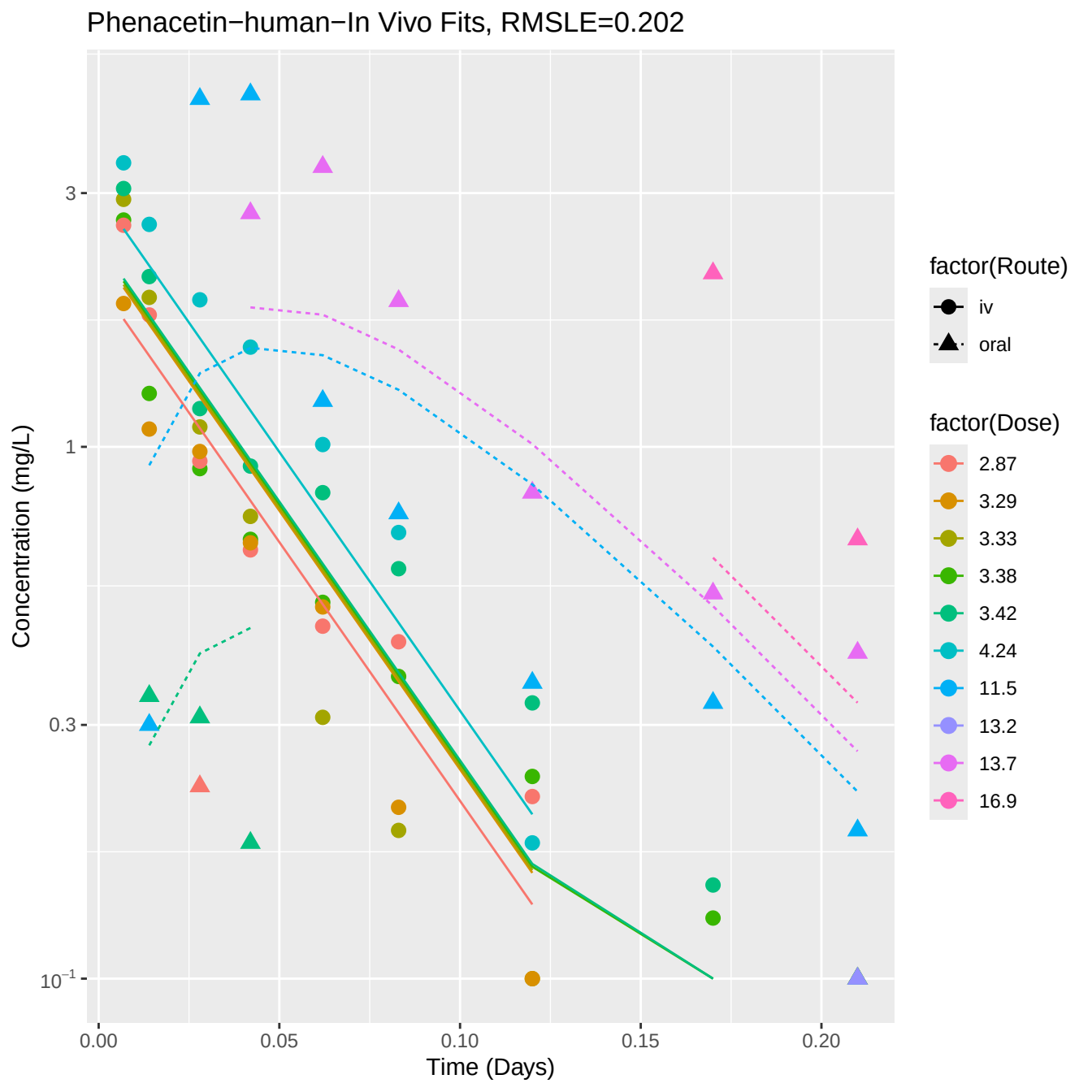

Phenacetin−rat−HTPBTK−Pradeep, RMSLE=0.337
Concentration (mg/L)
30
10
3
1
0.00
0.05
0.10
0.15
0.20
0.25
Time (Days)
factor(Dose)
23
250
factor(Route)
iv
oral

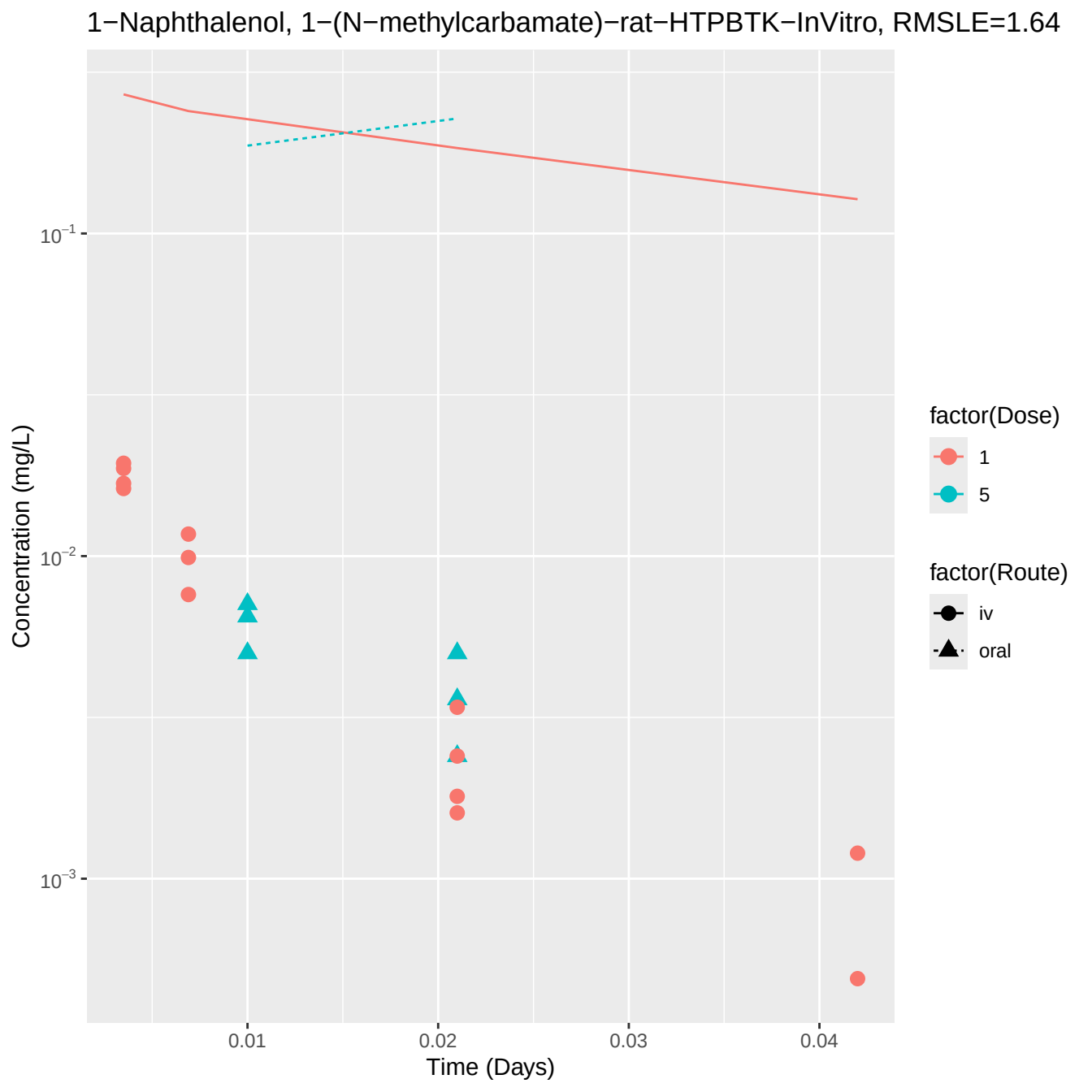

Phenacetin−human−HTPBTK−Pradeep, RMSLE=0.276
Concentration (mg/L)
3
1
0.3
10^{-1}
0.00
0.05
0.10
0.15
0.20
Time (Days)
factor(Route)
iv
oral
factor(Dose)
2.87
3.29
3.33
3.38
3.42
4.24
11.5
13.2
13.7
16.9

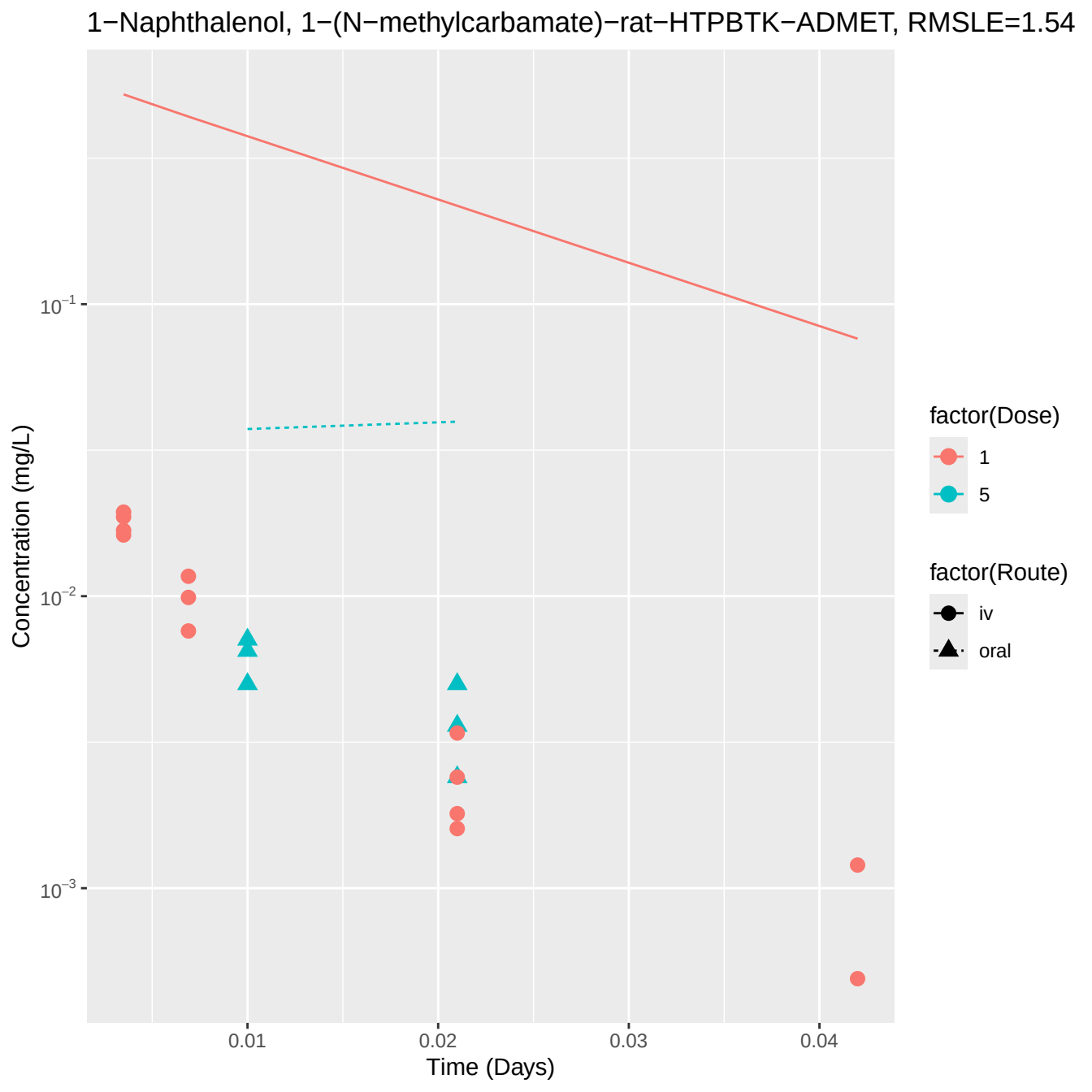

Phenacetin−rat−HTPBTK−Ensemble, RMSLE=0.447
Concentration (mg/L)
30
10
3
1
0.00
0.05
0.10
0.15
0.20
0.25
Time (Days)
factor(Dose)
23
250
factor(Route)
iv
oral

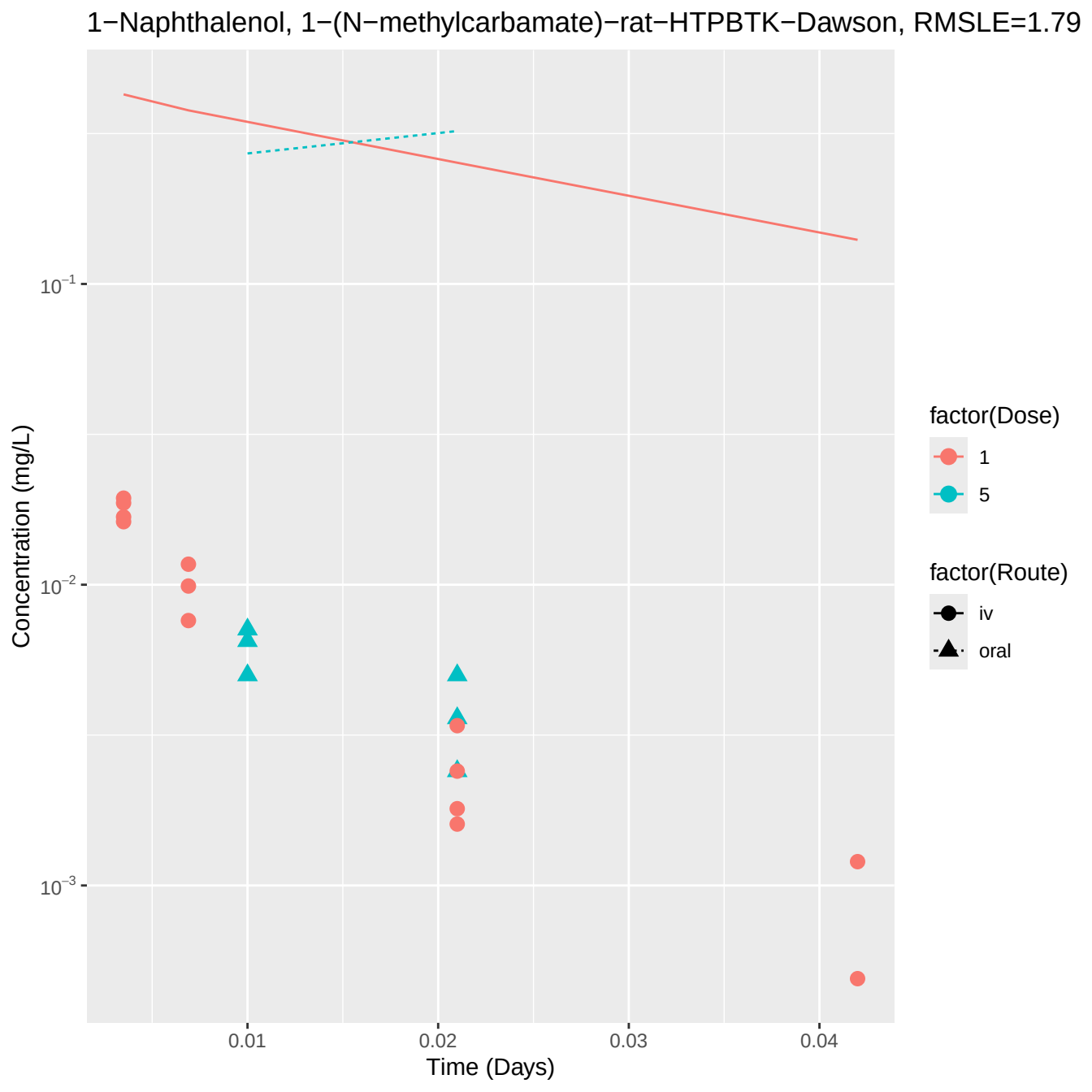

Phenacetin−human−HTPBTK−Ensemble, RMSLE=0.353
Concentration (mg/L)
3
1
0.3
10^{-1}
0.00
0.05
0.10
0.15
0.20
Time (Days)
factor(Route)
iv
oral
factor(Dose)
2.87
3.29
3.33
3.38
3.42
4.24
11.5
13.2
13.7
16.9

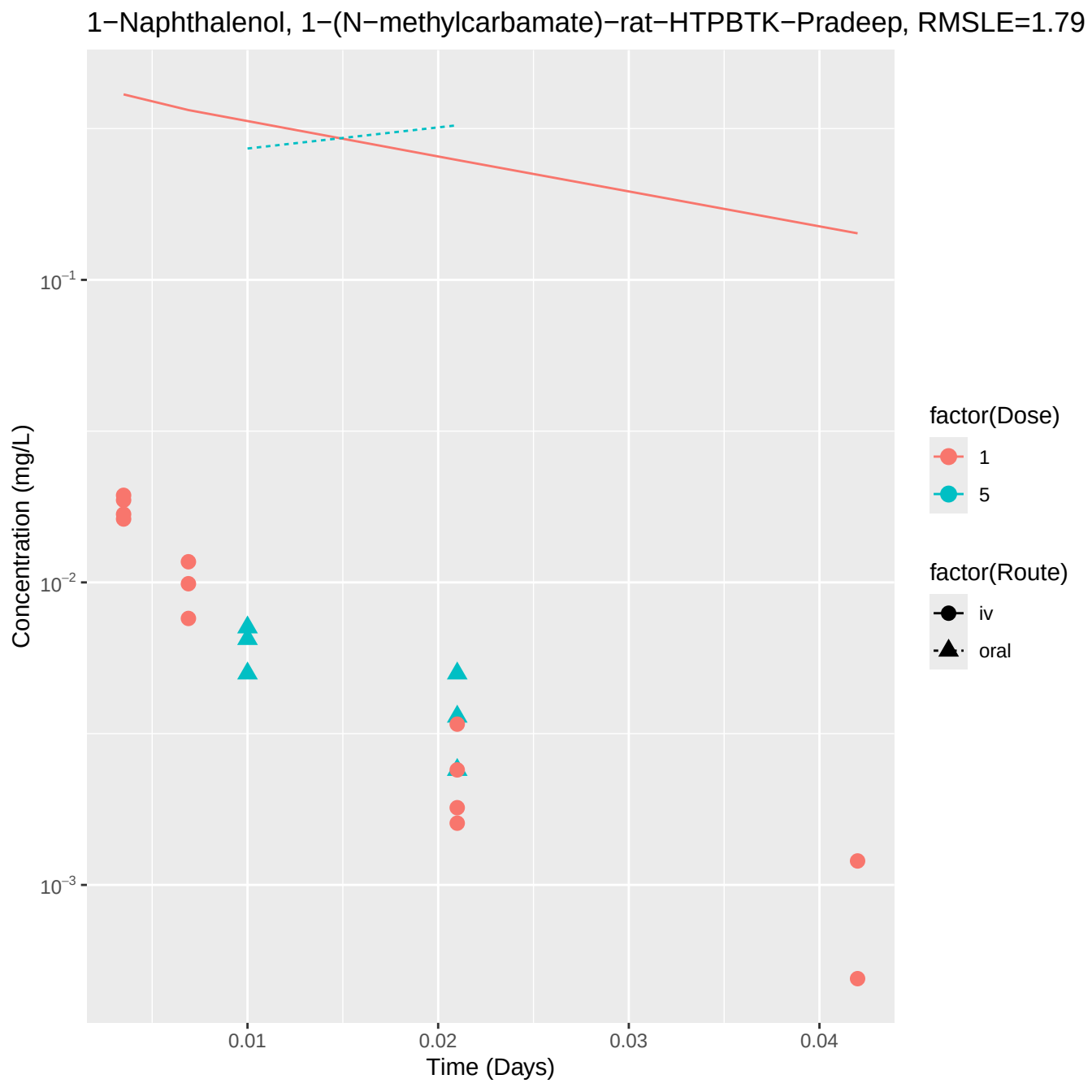

Phenacetin−rat−In Vivo Fits, RMSLE=0.181
Concentration (mg/L)
30
10
3
1
0.00
0.05
0.10
0.15
0.20
0.25
Time (Days)
factor(Dose)
23
250
factor(Route)
iv
oral

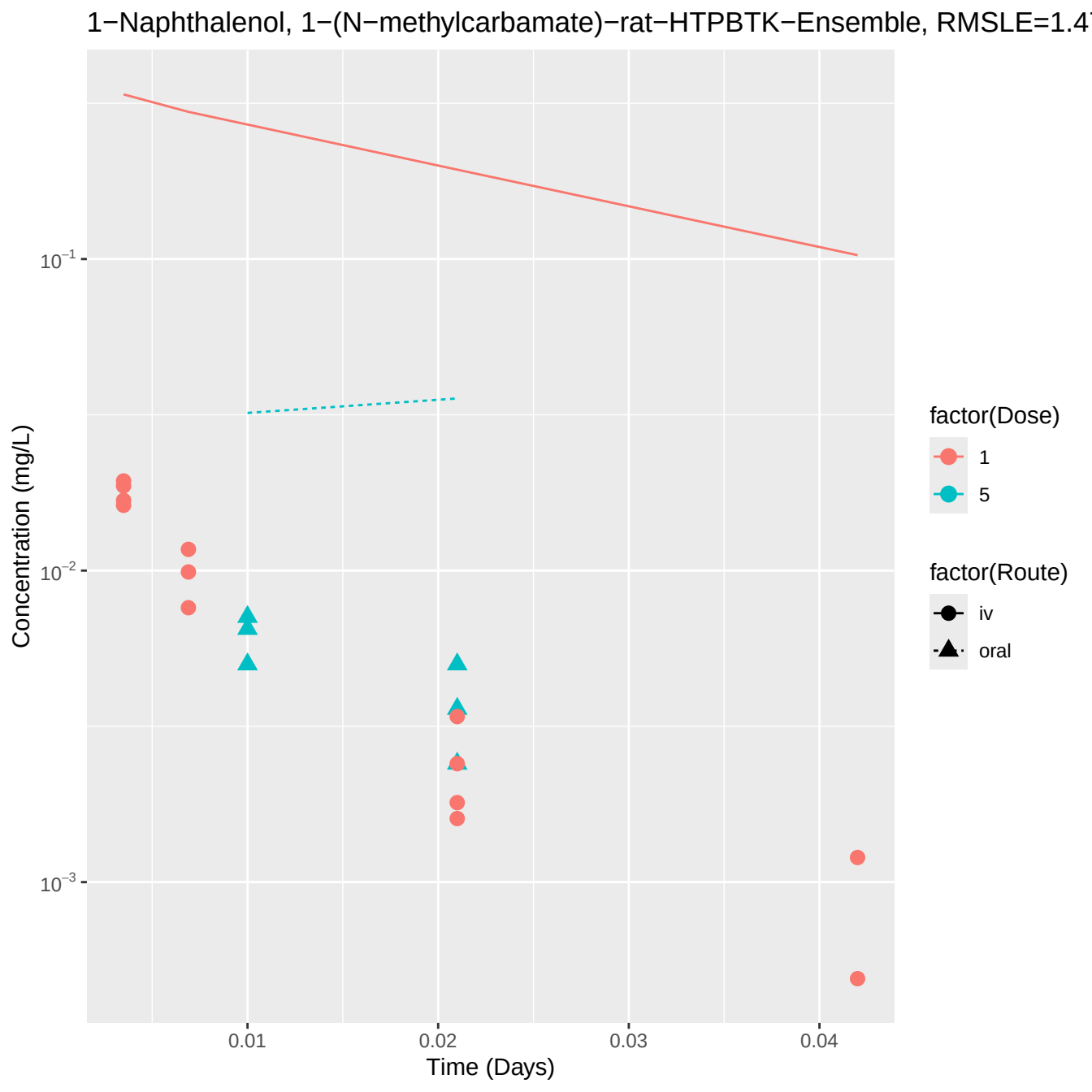

Phenacetin−human−In Vivo Fits, RMSLE=0.202
Concentration (mg/L)
3
1
0.3
10^{-1}
0.00
0.05
0.10
0.15
0.20
Time (Days)
factor(Route)
iv
oral
factor(Dose)
2.87
3.29
3.33
3.38
3.42
4.24
11.5
13.2
13.7
16.9

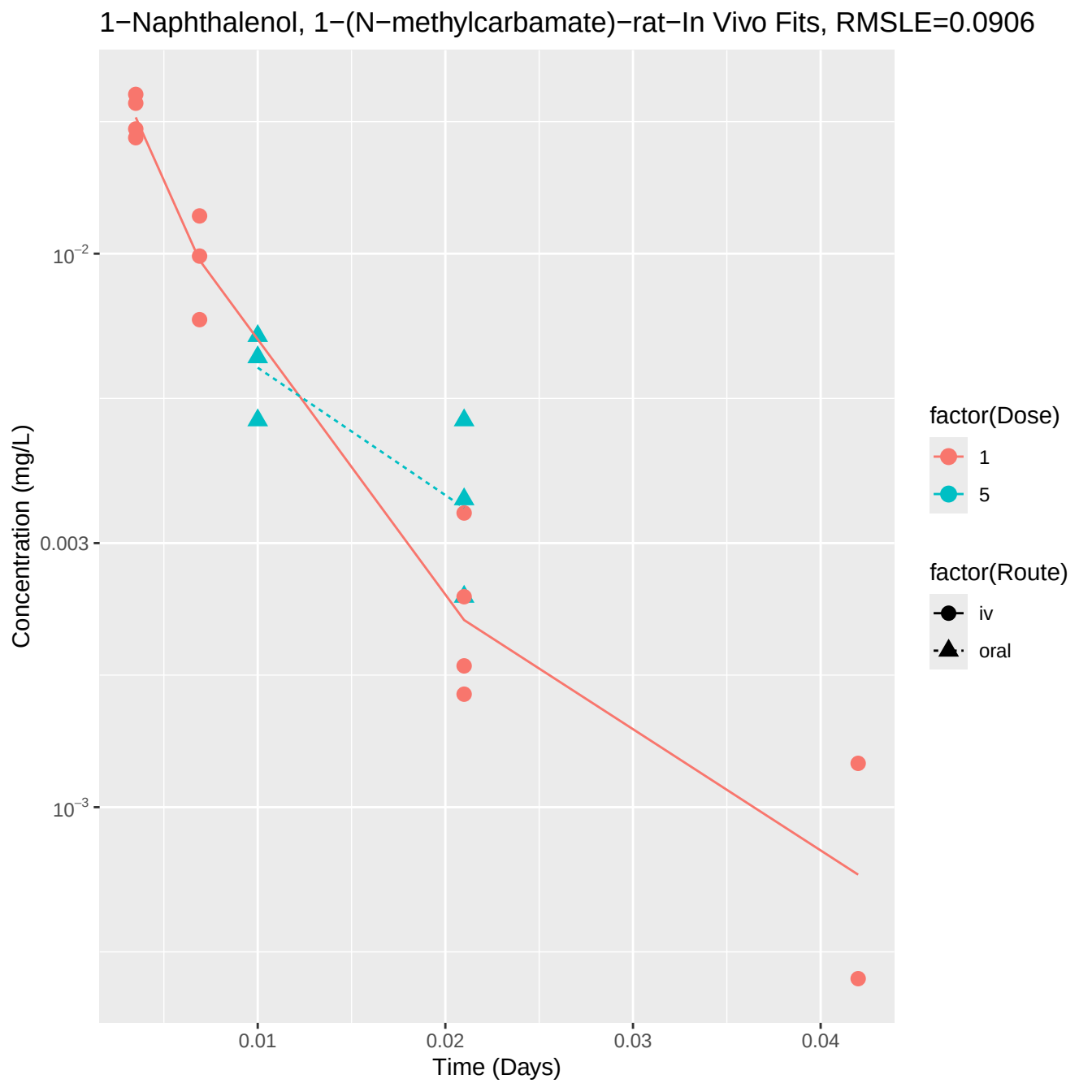

1−Naphthalenol, 1−(N−methylcarbamate)−rat−HTPBTK−InVitro, RMSLE=1.64
Concentration (mg/L)
10^{-1}
10^{-2}
10^{-3}
0.01
0.02
0.03
0.04
Time (Days)
factor(Dose)
1
5
factor(Route)
iv
oral

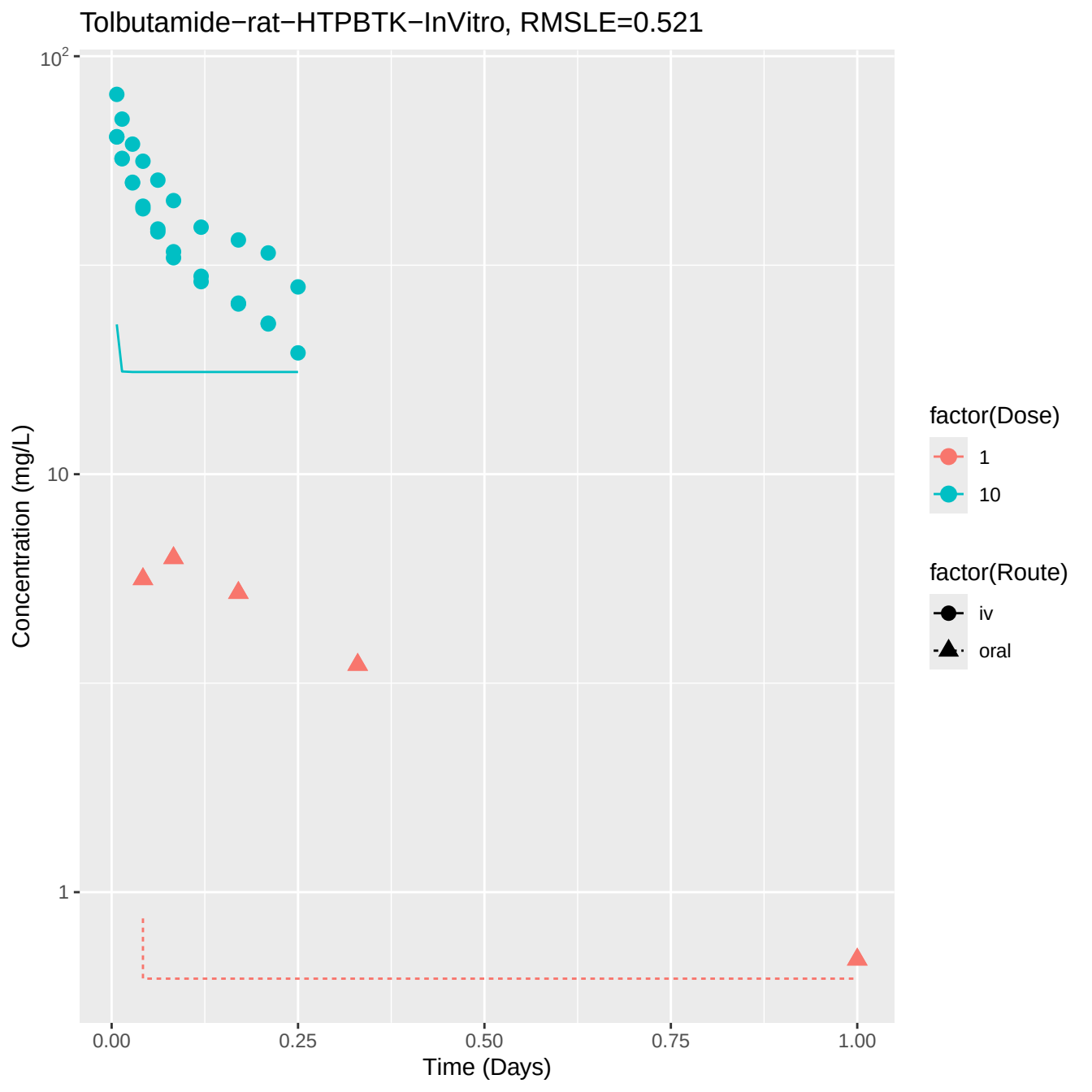

1−Naphthalenol, 1−(N−methylcarbamate)−rat−HTPBTK−ADMET, RMSLE=1.54
Concentration (mg/L)
10^{-1}
10^{-2}
10^{-3}
0.01
0.02
0.03
0.04
Time (Days)
factor(Dose)
1
5
factor(Route)
iv
oral

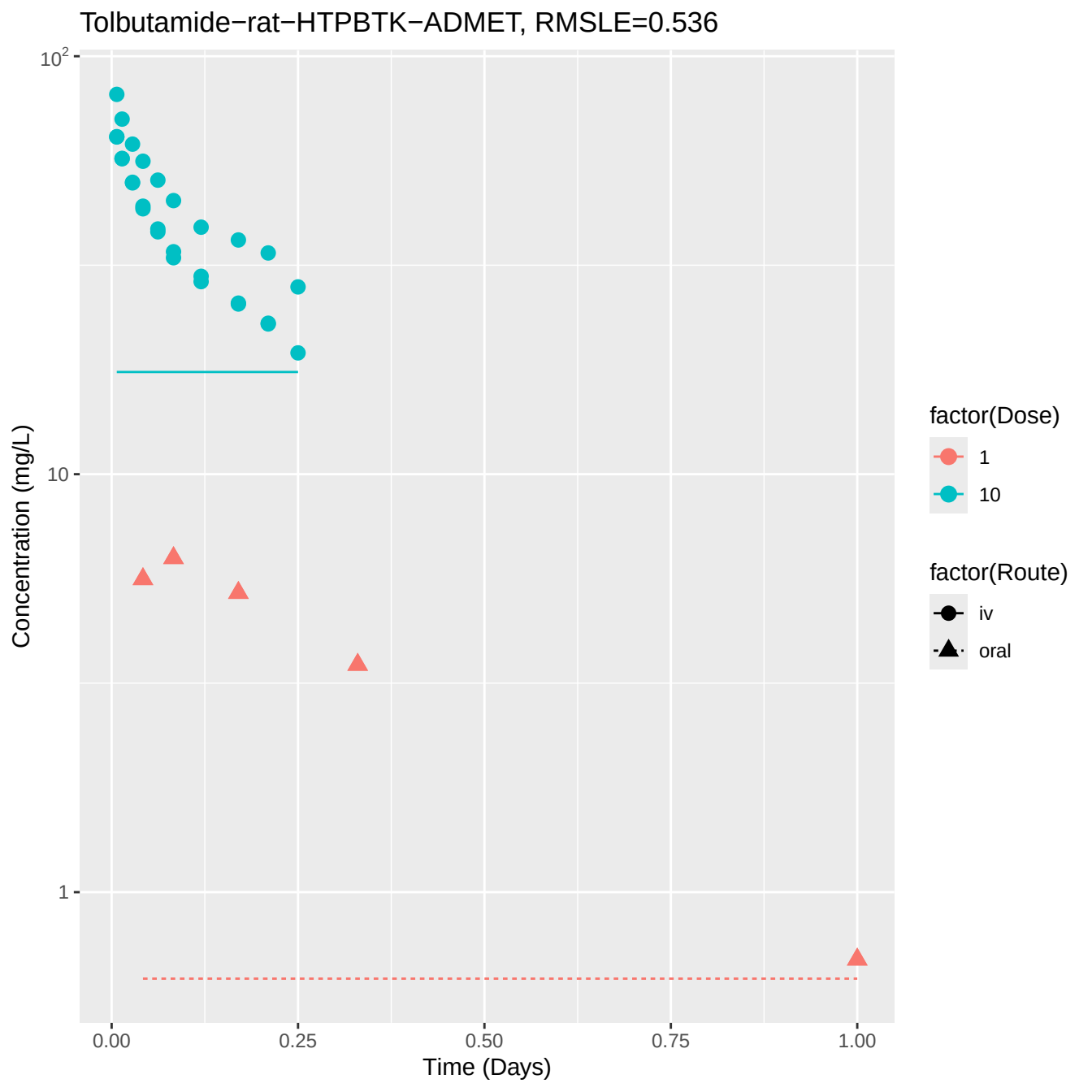

1−Naphthalenol, 1−(N−methylcarbamate)−rat−HTPBTK−Dawson, RMSLE=1.79
Concentration (mg/L)
10^{-1}
10^{-2}
10^{-3}
0.01
0.02
0.03
0.04
Time (Days)
factor(Dose)
1
5
factor(Route)
iv
oral

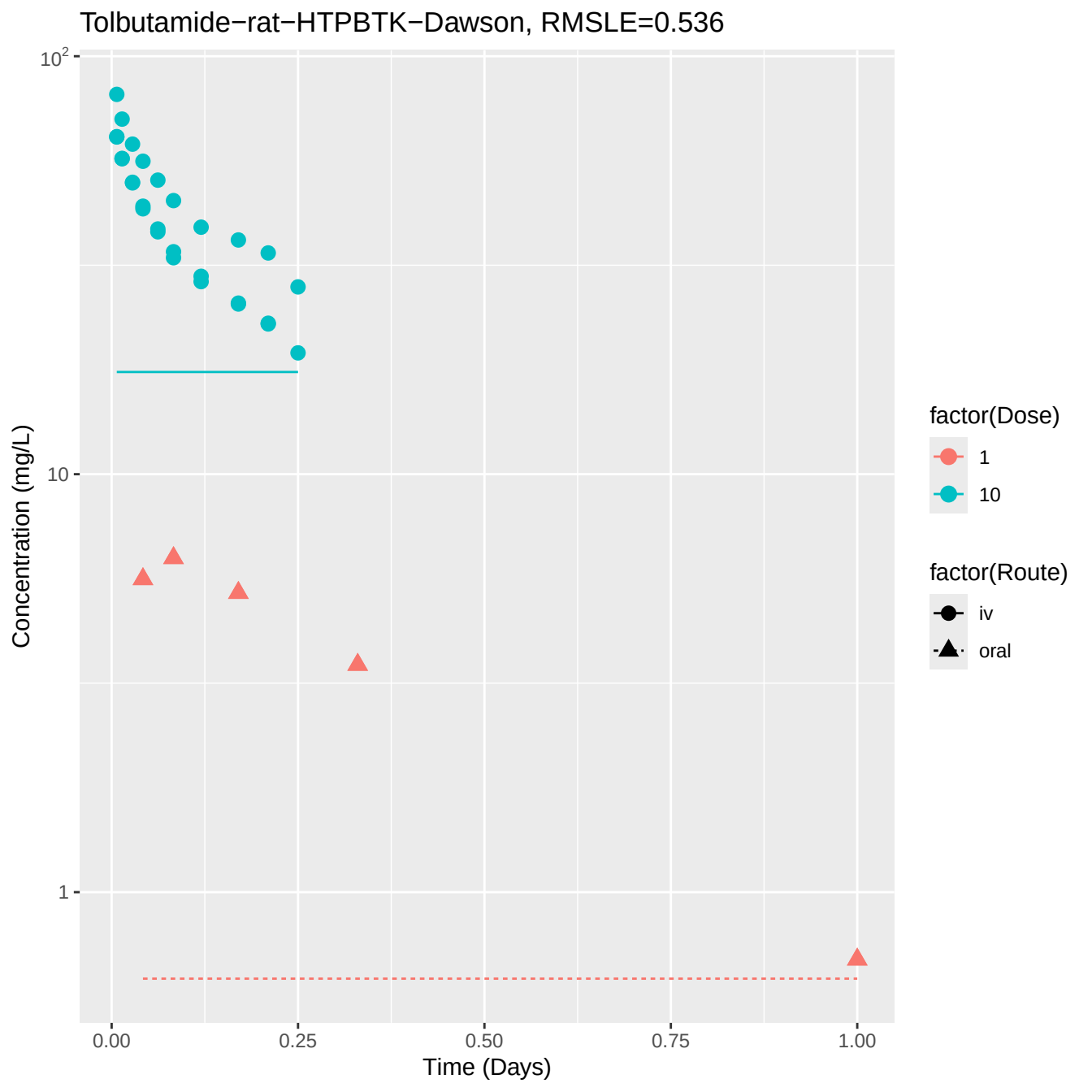

1−Naphthalenol, 1−(N−methylcarbamate)−rat−HTPBTK−Pradeep, RMSLE=1.79
Concentration (mg/L)
10^{-1}
10^{-2}
10^{-3}
0.01
0.02
0.03
0.04
Time (Days)
factor(Dose)
1
5
factor(Route)
iv
oral

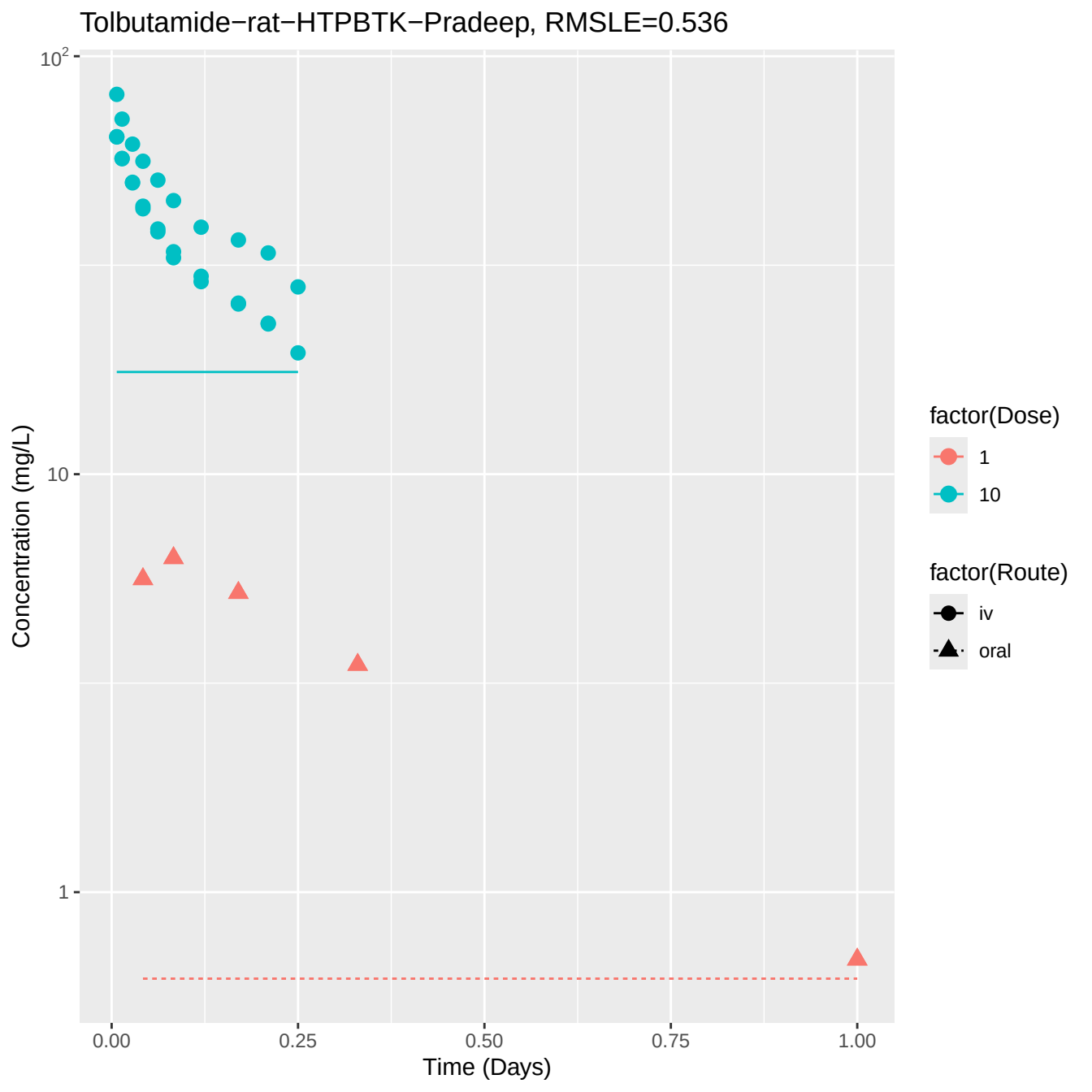

1−Naphthalenol, 1−(N−methylcarbamate)−rat−HTPBTK−Ensemble, RMSLE=1.47
Concentration (mg/L)
10^{-1}
10^{-2}
10^{-3}
0.01
0.02
0.03
0.04
Time (Days)
factor(Dose)
1
5
factor(Route)
iv
oral

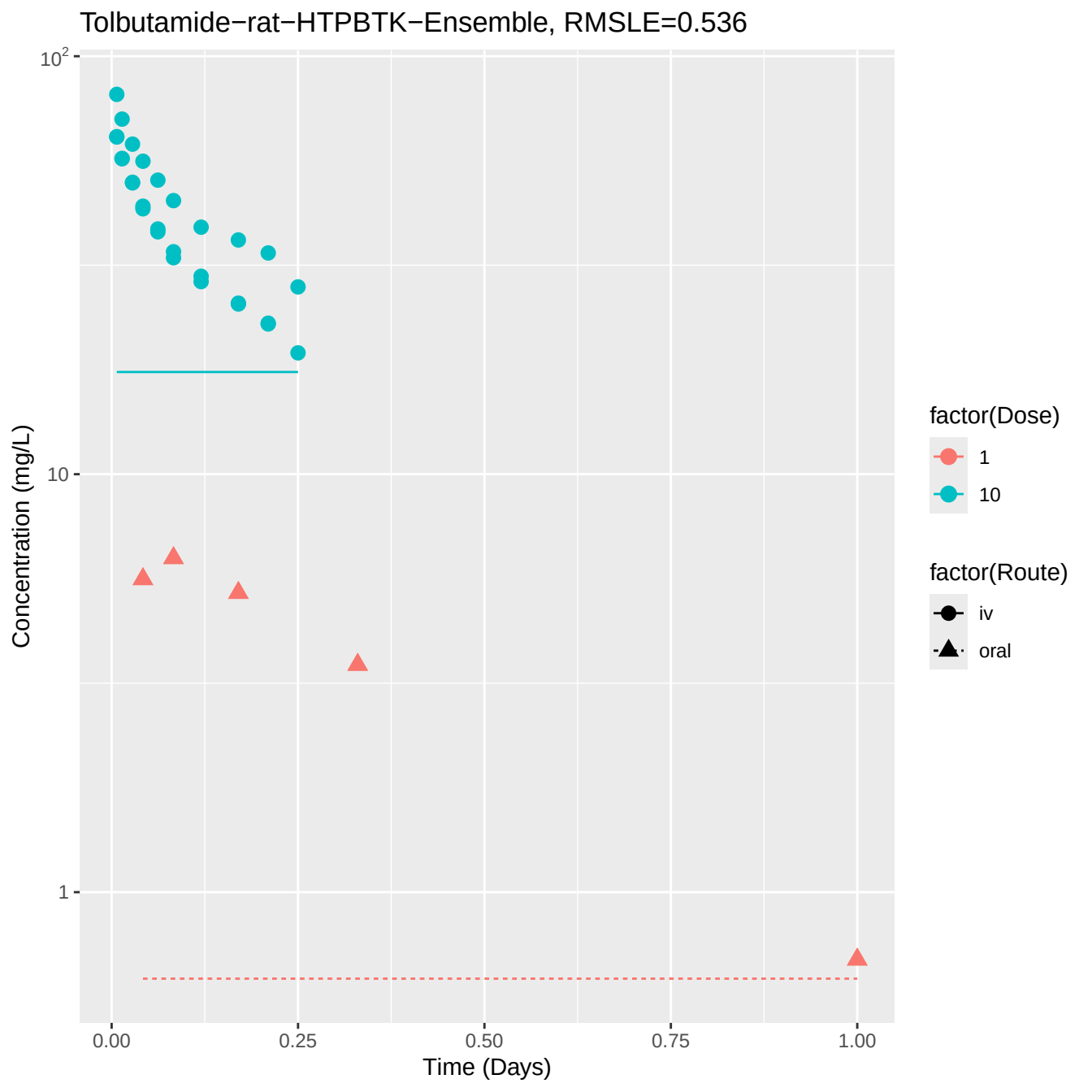

1−Naphthalenol, 1−(N−methylcarbamate)−rat−In Vivo Fits, RMSLE=0.0906
Concentration (mg/L)
10^{-2}
0.003
10^{-3}
0.01
0.02
0.03
0.04
Time (Days)
factor(Dose)
1
5
factor(Route)
iv
oral

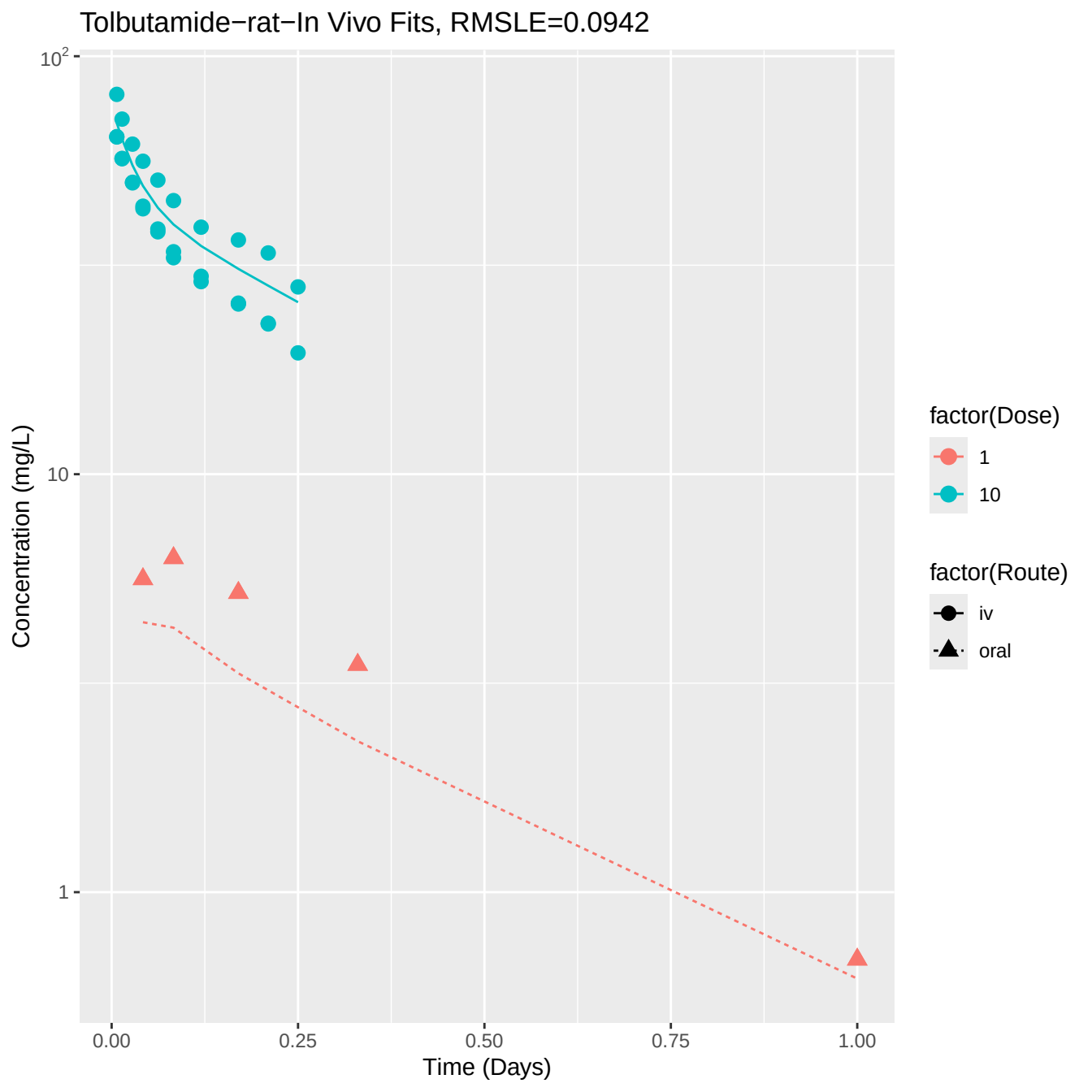

Tolbutamide−rat−HTPBTK−InVitro, RMSLE=0.521
10^2
10
1
Concentration (mg/L)
0.00
0.25
0.50
0.75
1.00
Time (Days)
factor(Dose)
1
10
factor(Route)
iv
oral

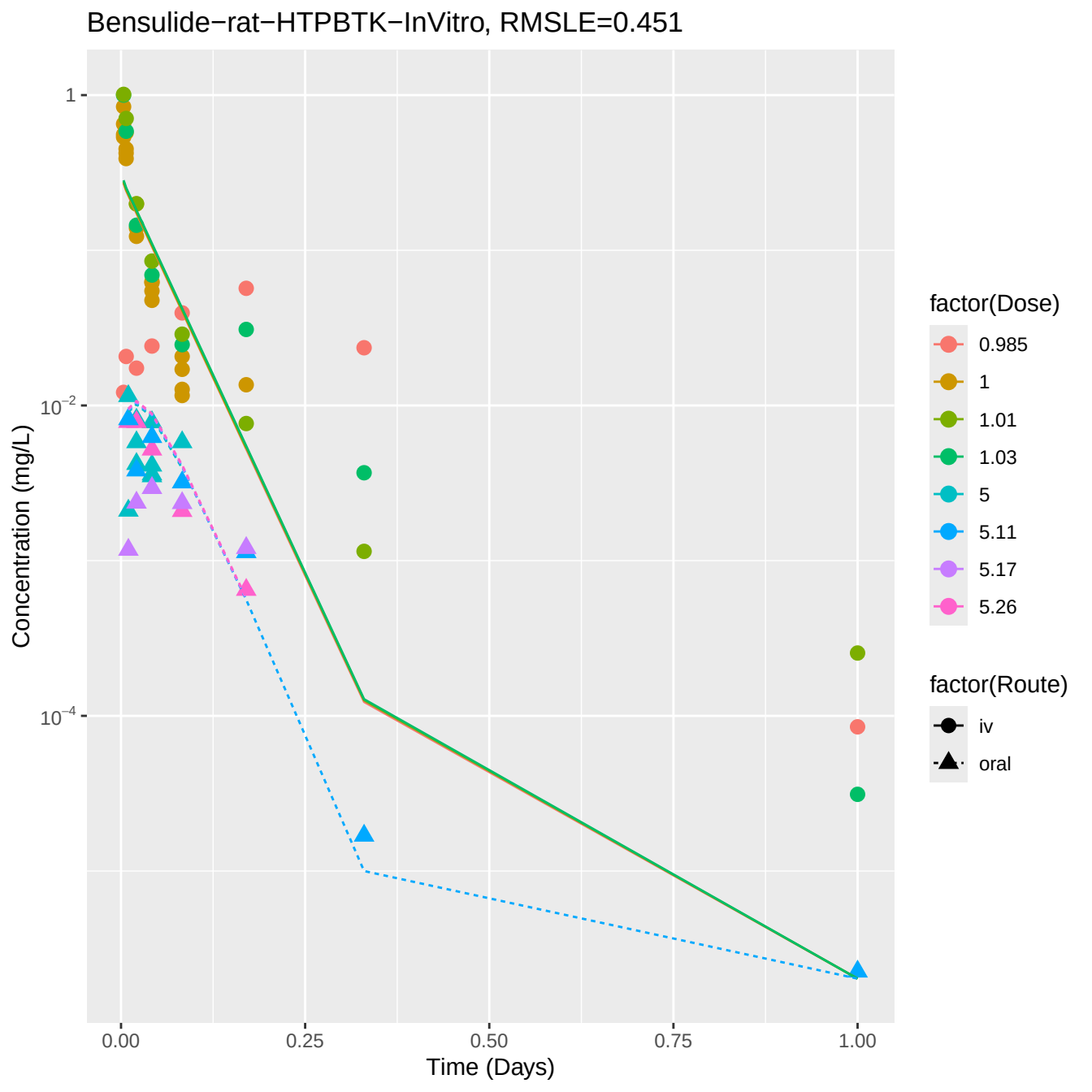

Tolbutamide−rat−HTPBTK−ADMET, RMSLE=0.536
10^2
10
1
Concentration (mg/L)
0.00
0.25
0.50
0.75
1.00
Time (Days)
factor(Dose)
1
10
factor(Route)
iv
oral

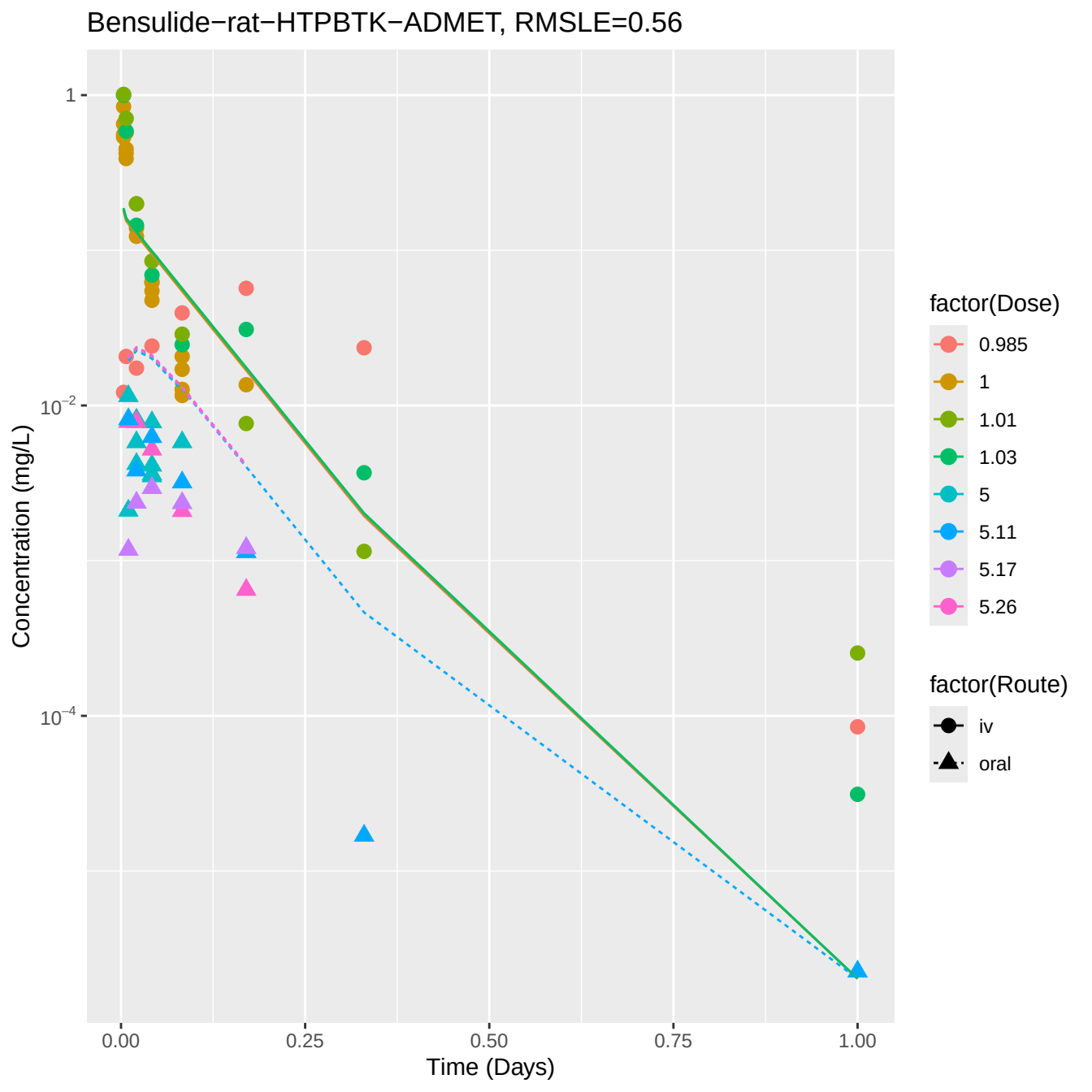

Tolbutamide−rat−HTPBTK−Dawson, RMSLE=0.536
10^2
10
1
Concentration (mg/L)
0.00
0.25
0.50
0.75
1.00
Time (Days)
factor(Dose)
1
10
factor(Route)
iv
oral

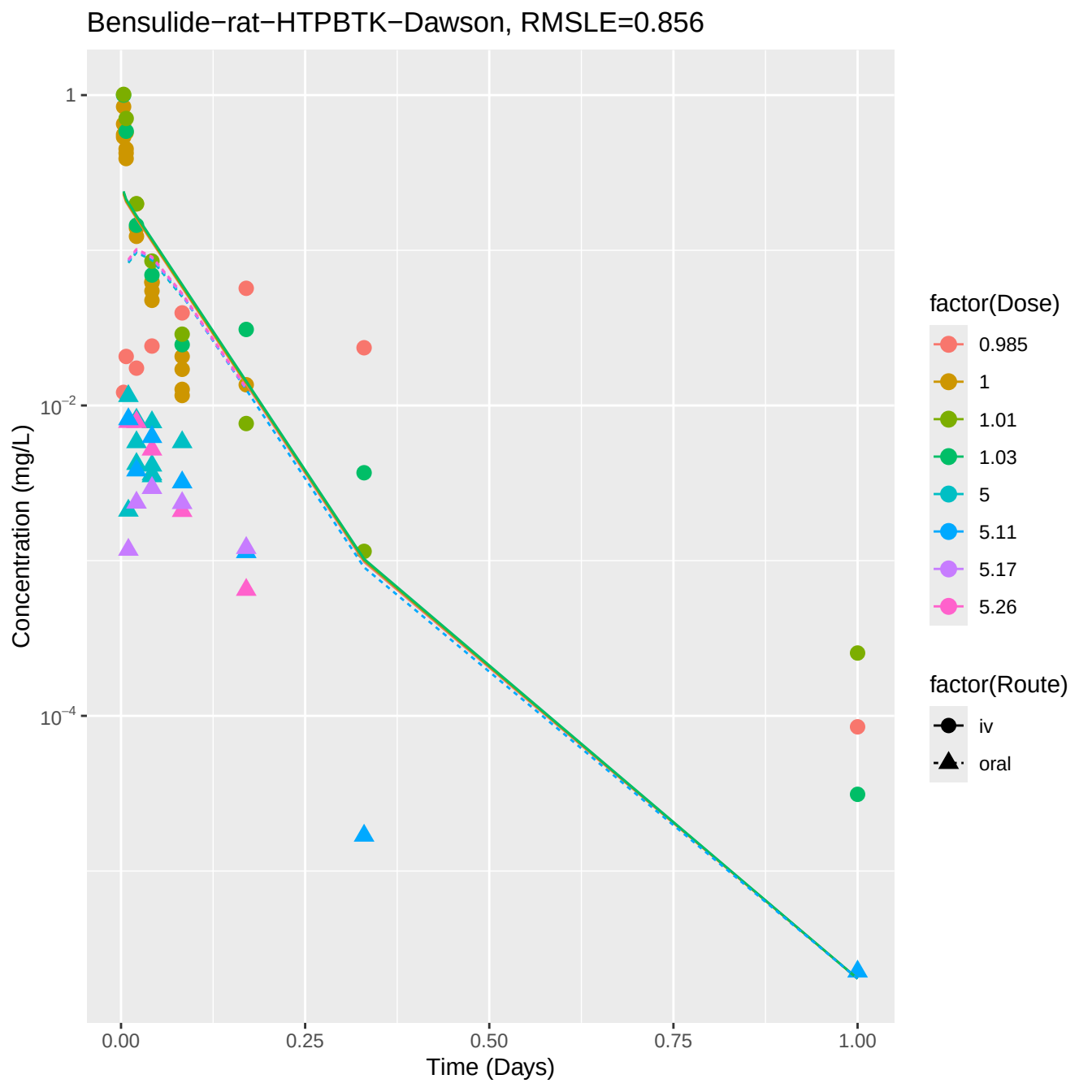

Tolbutamide−rat−HTPBTK−Pradeep, RMSLE=0.536
10^2
10
1
Concentration (mg/L)
0.00
0.25
0.50
0.75
1.00
Time (Days)
factor(Dose)
1
10
factor(Route)
iv
oral

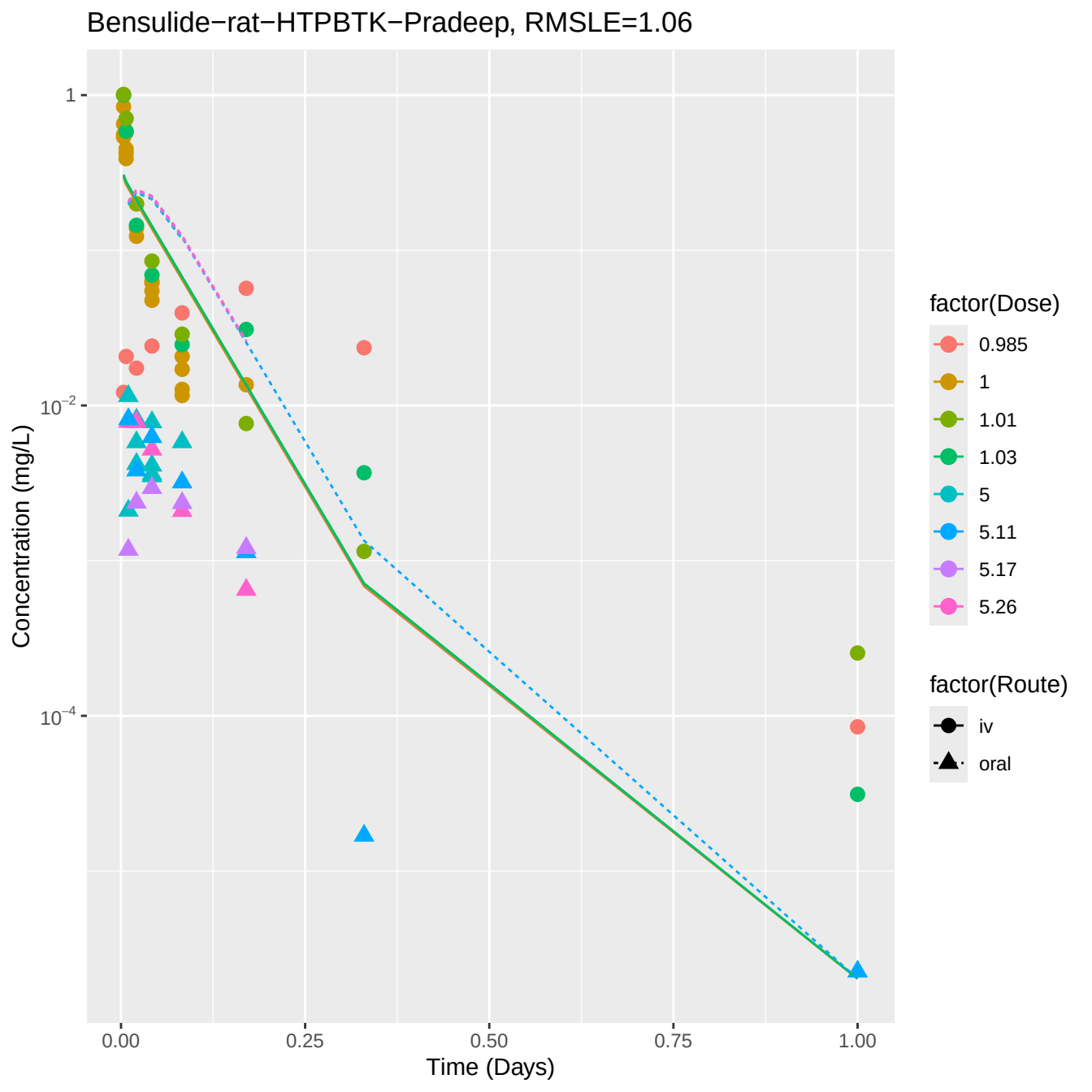

Tolbutamide−rat−HTPBTK−Ensemble, RMSLE=0.536
10^2
10
1
Concentration (mg/L)
0.00
0.25
0.50
0.75
1.00
Time (Days)
factor(Dose)
1
10
factor(Route)
iv
oral

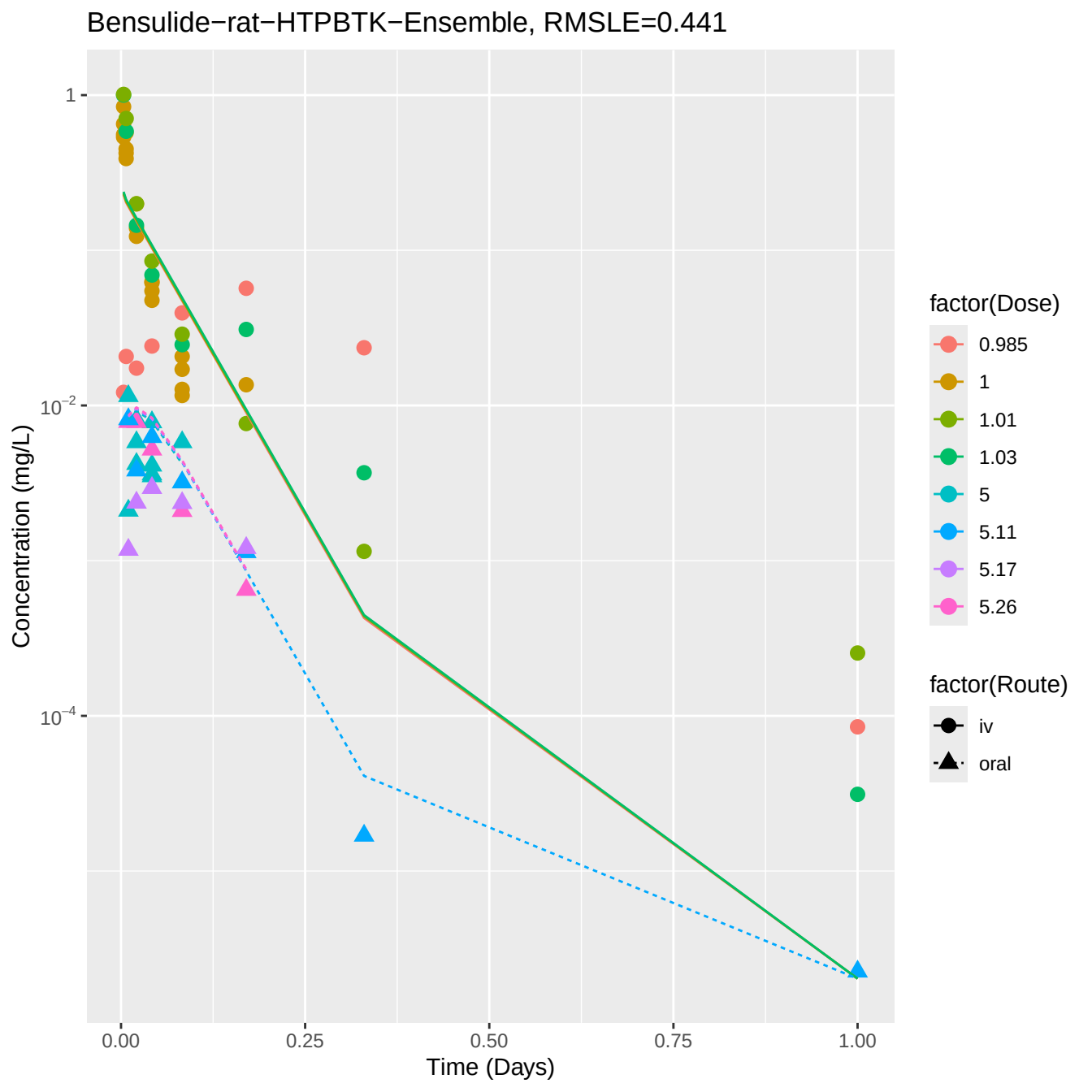

Tolbutamide−rat−In Vivo Fits, RMSLE=0.0942
10^2
10
1
Concentration (mg/L)
0.00
0.25
0.50
0.75
1.00
Time (Days)
factor(Dose)
1
10
factor(Route)
iv
oral

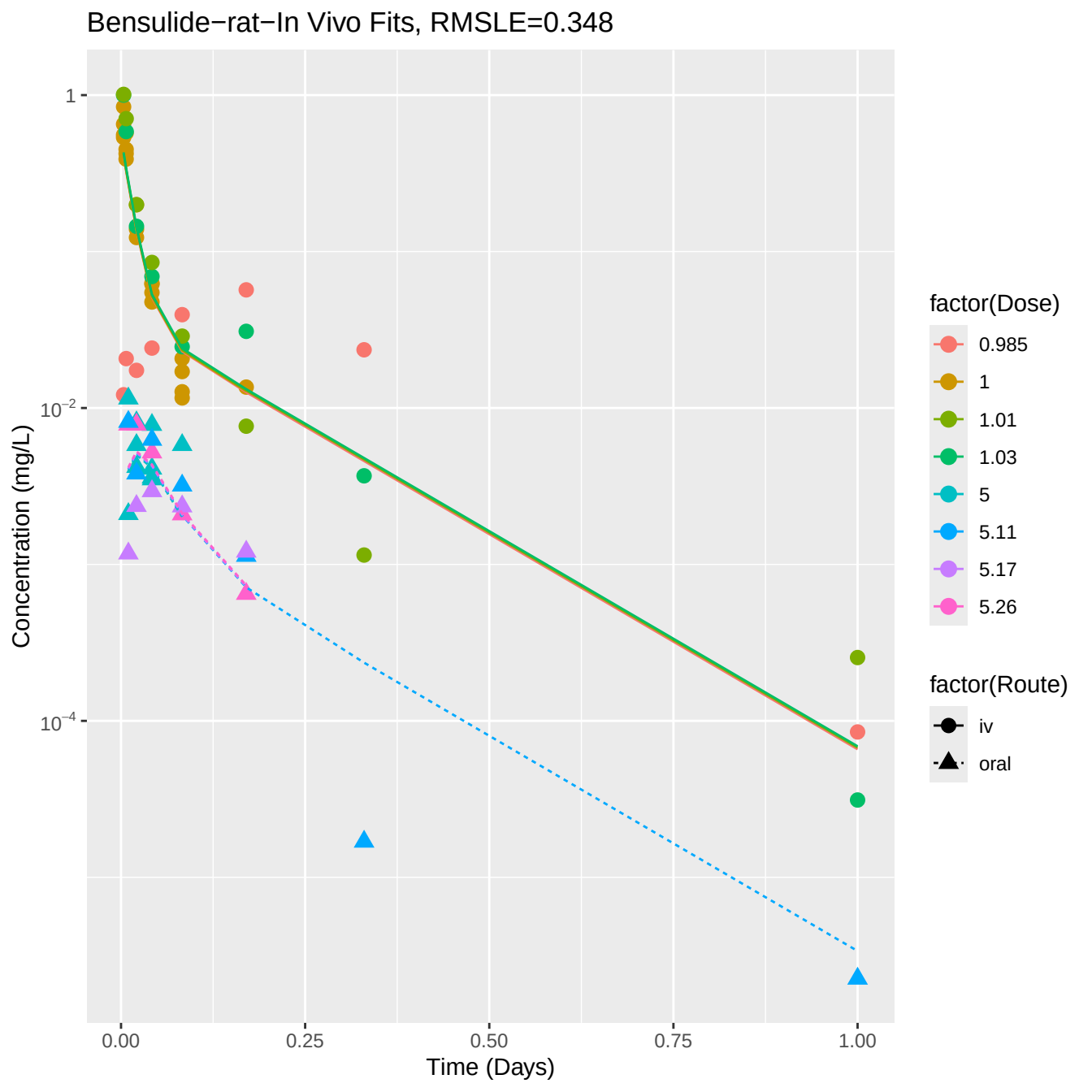

Bensulide−rat−HTPBTK−InVitro, RMSLE=0.451
1
10^{-2}
10^{-4}
Concentration (mg/L)
0.00
0.25
0.50
0.75
1.00
Time (Days)
factor(Dose)
0.985
1
1.01
1.03
5
5.11
5.17
5.26
factor(Route)
iv
oral

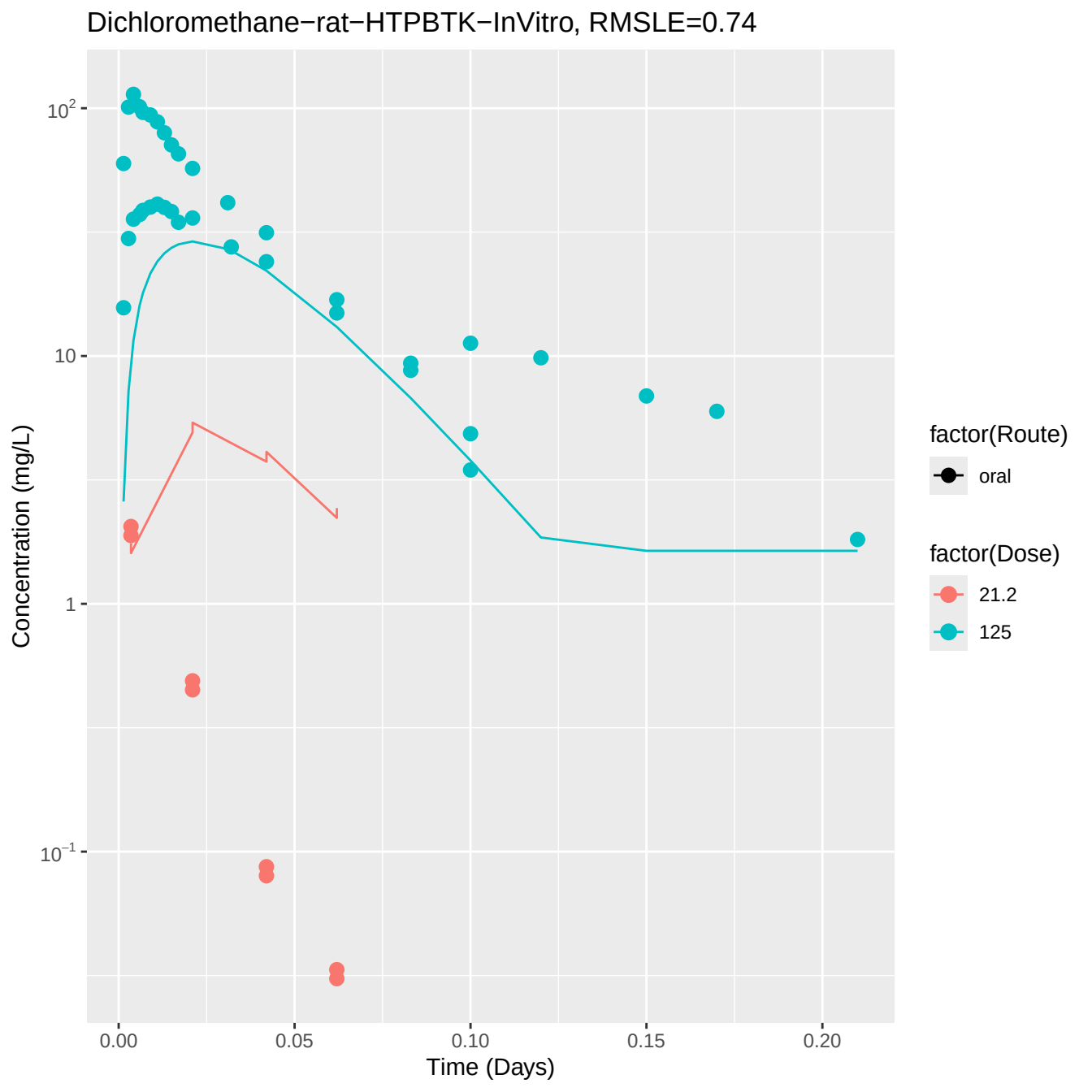

Bensulide−rat−HTPBTK−ADMET, RMSLE=0.56
Concentration (mg/L)
1
10^{-2}
10^{-4}
0.00
0.25
0.50
0.75
1.00
Time (Days)
factor(Dose)
0.985
1
1.01
1.03
5
5.11
5.17
5.26
factor(Route)
iv
oral

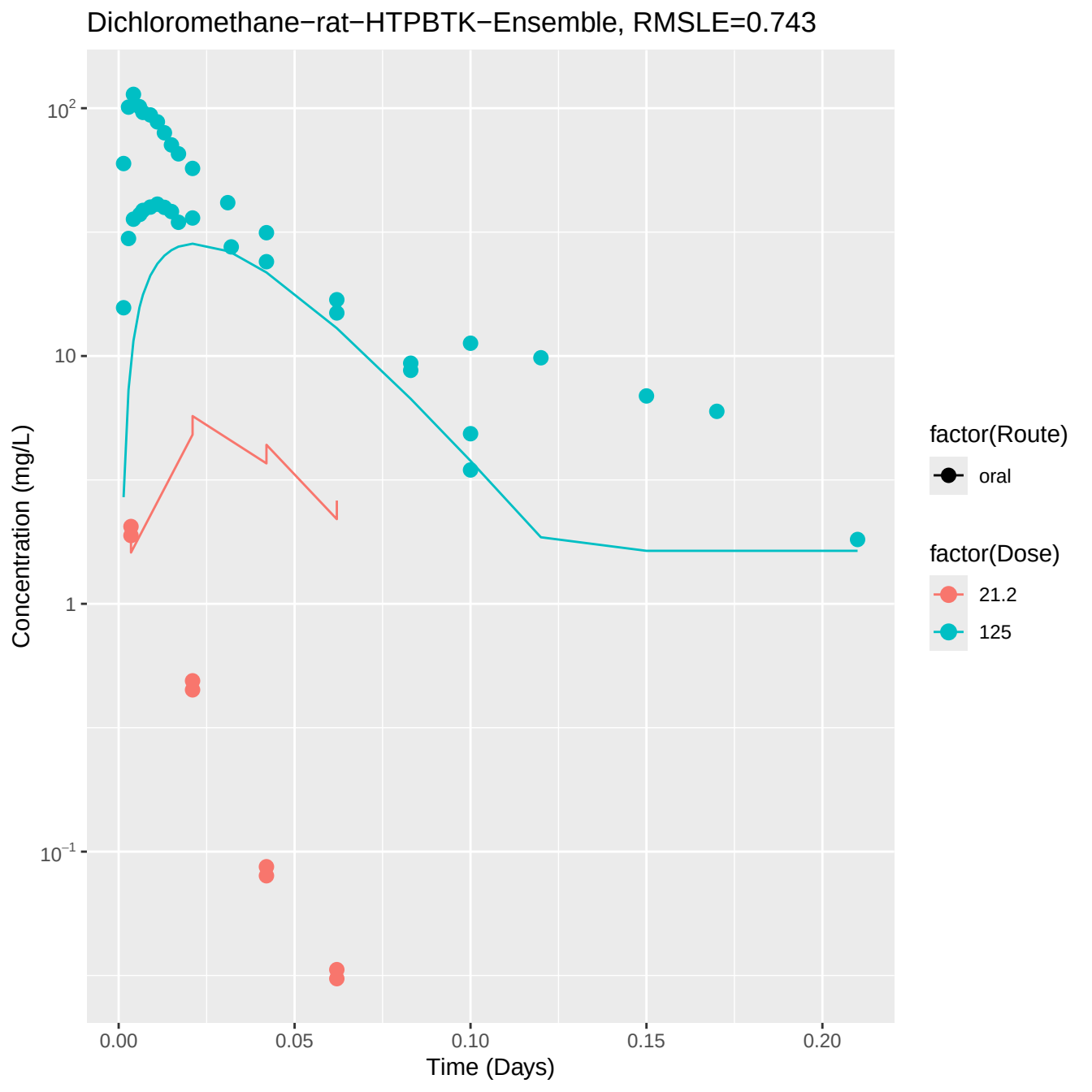

Bensulide−rat−HTPBTK−Dawson, RMSLE=0.856
Concentration (mg/L)
1
10^{-2}
10^{-4}
0.00
0.25
0.50
0.75
1.00
Time (Days)
factor(Dose)
0.985
1
1.01
1.03
5
5.11
5.17
5.26
factor(Route)
iv
oral

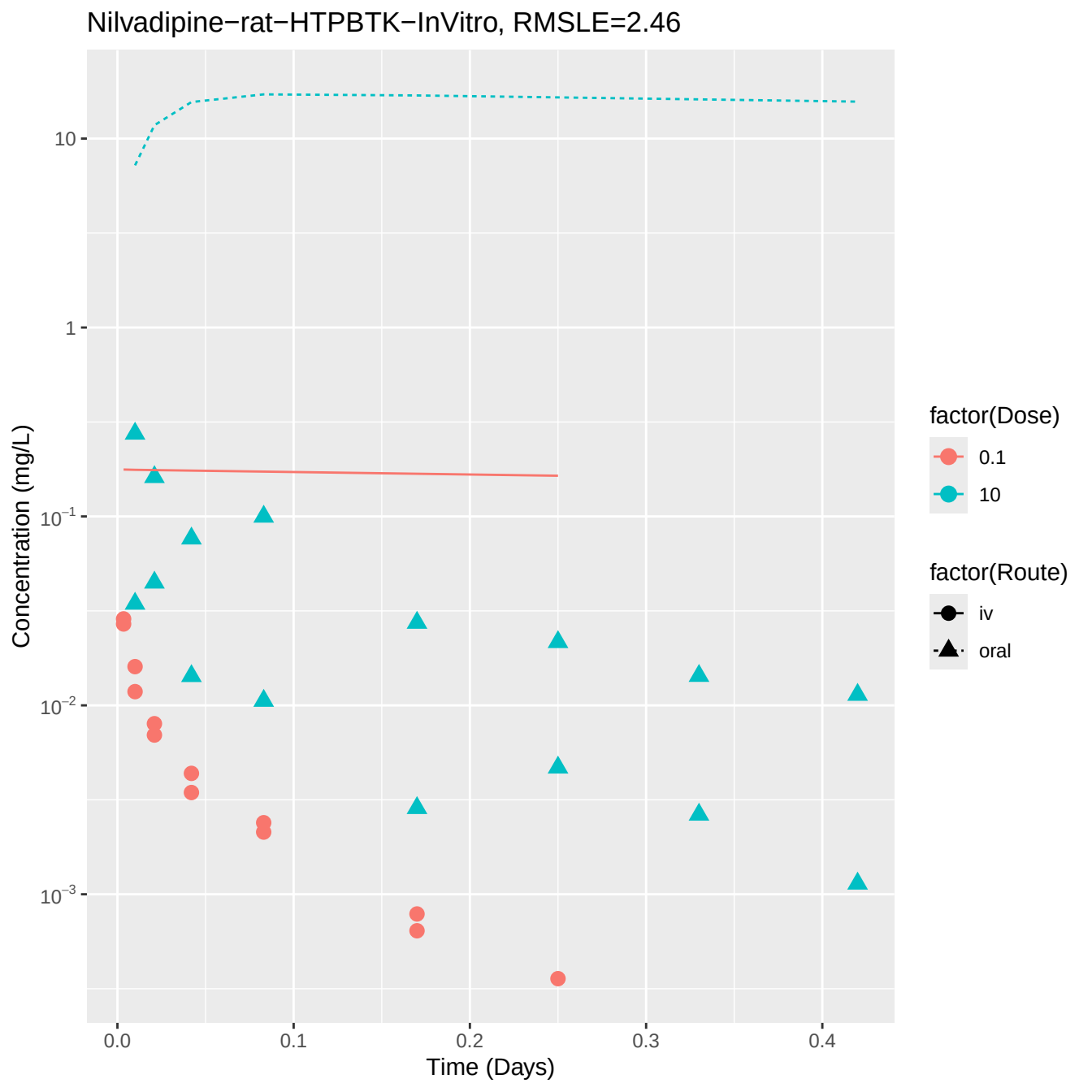

Bensulide−rat−HTPBTK−Pradeep, RMSLE=1.06
Concentration (mg/L)
1
10^{-2}
10^{-4}
0.00
0.25
0.50
0.75
1.00
Time (Days)
factor(Dose)
0.985
1
1.01
1.03
5
5.11
5.17
5.26
factor(Route)
iv
oral

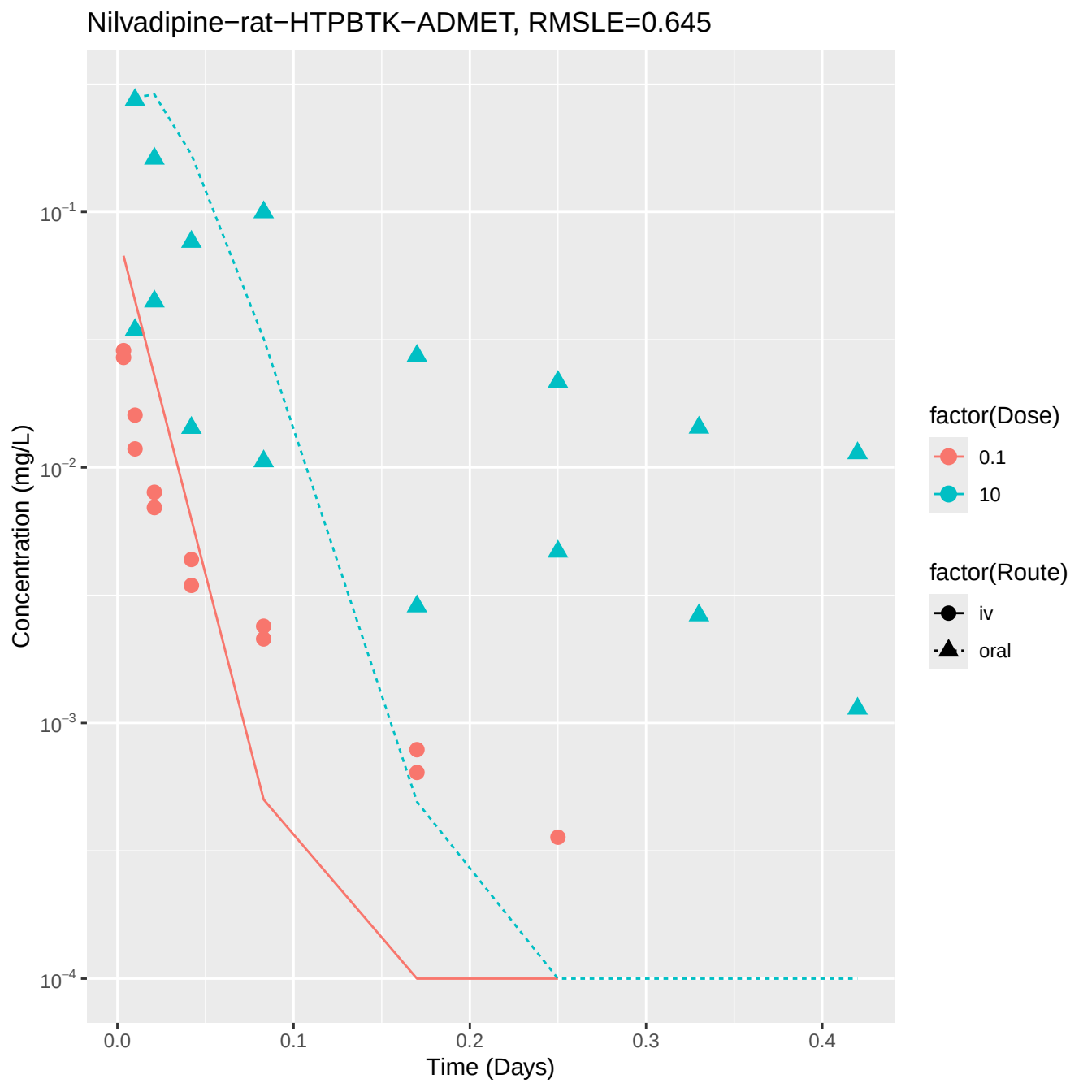

Bensulide−rat−HTPBTK−Ensemble, RMSLE=0.441
Concentration (mg/L)
1
10^{-2}
10^{-4}
0.00
0.25
0.50
0.75
1.00
Time (Days)
factor(Dose)
0.985
1
1.01
1.03
5
5.11
5.17
5.26
factor(Route)
iv
oral

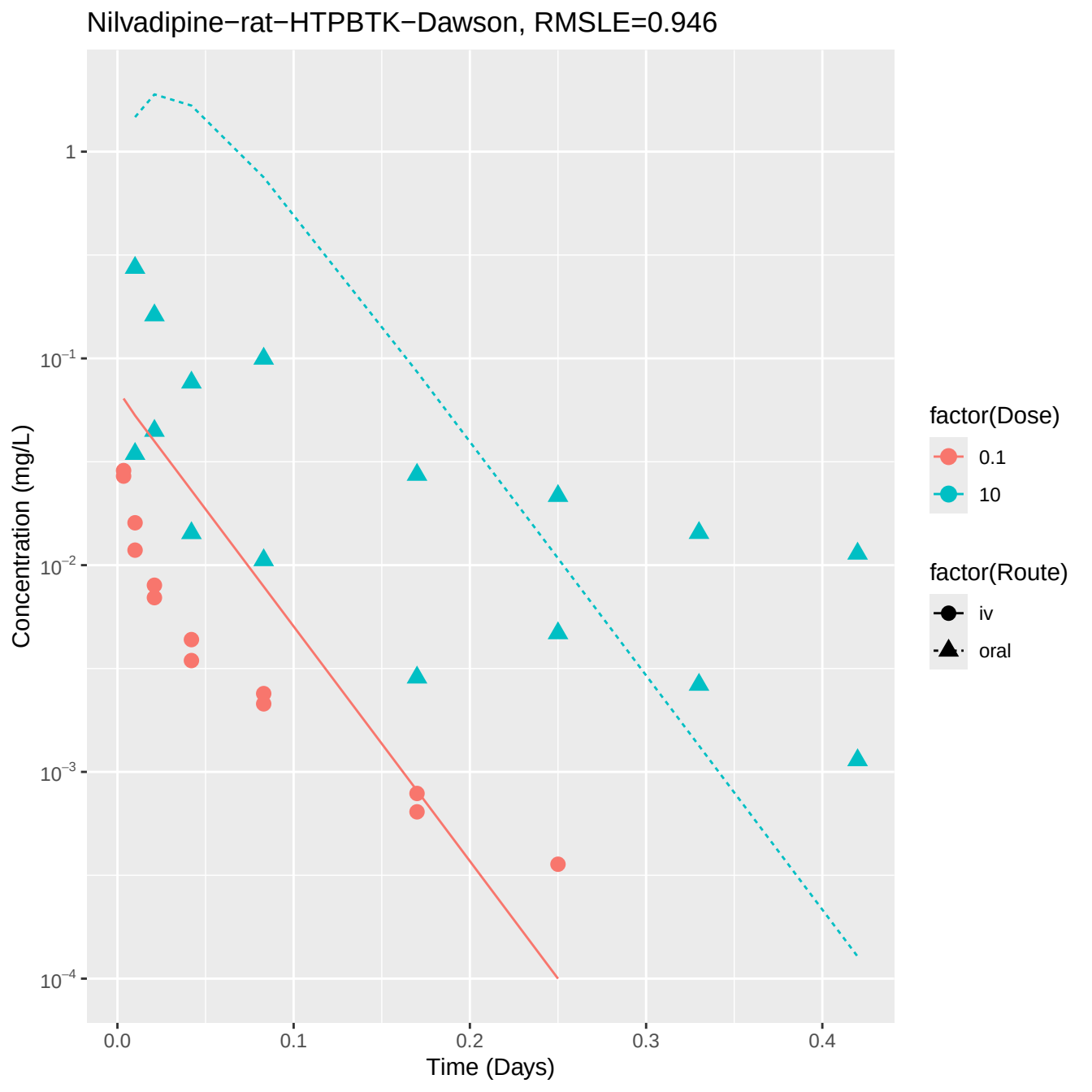

Bensulide−rat−In Vivo Fits, RMSLE=0.348
Concentration (mg/L)
1
10^{-2}
10^{-4}
0.00
0.25
0.50
0.75
1.00
Time (Days)
factor(Dose)
0.985
1
1.01
1.03
5
5.11
5.17
5.26
factor(Route)
iv
oral

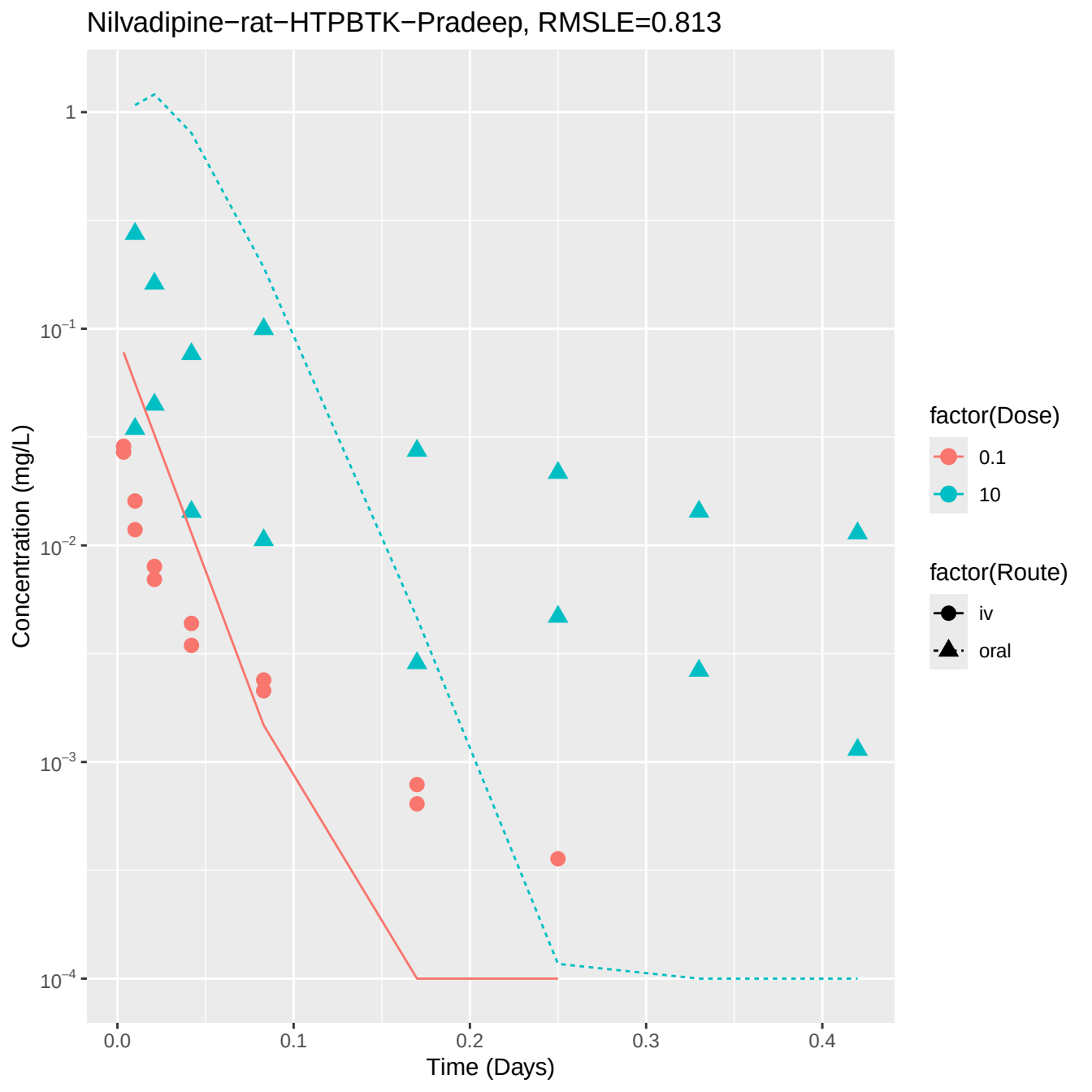

Dichloromethane−rat−HTPBTK−InVitro, RMSLE=0.74
Concentration (mg/L)
10^2
10
1
10^{-1}
0.00
0.05
0.10
0.15
0.20
Time (Days)
factor(Route)
oral
factor(Dose)
21.2
125

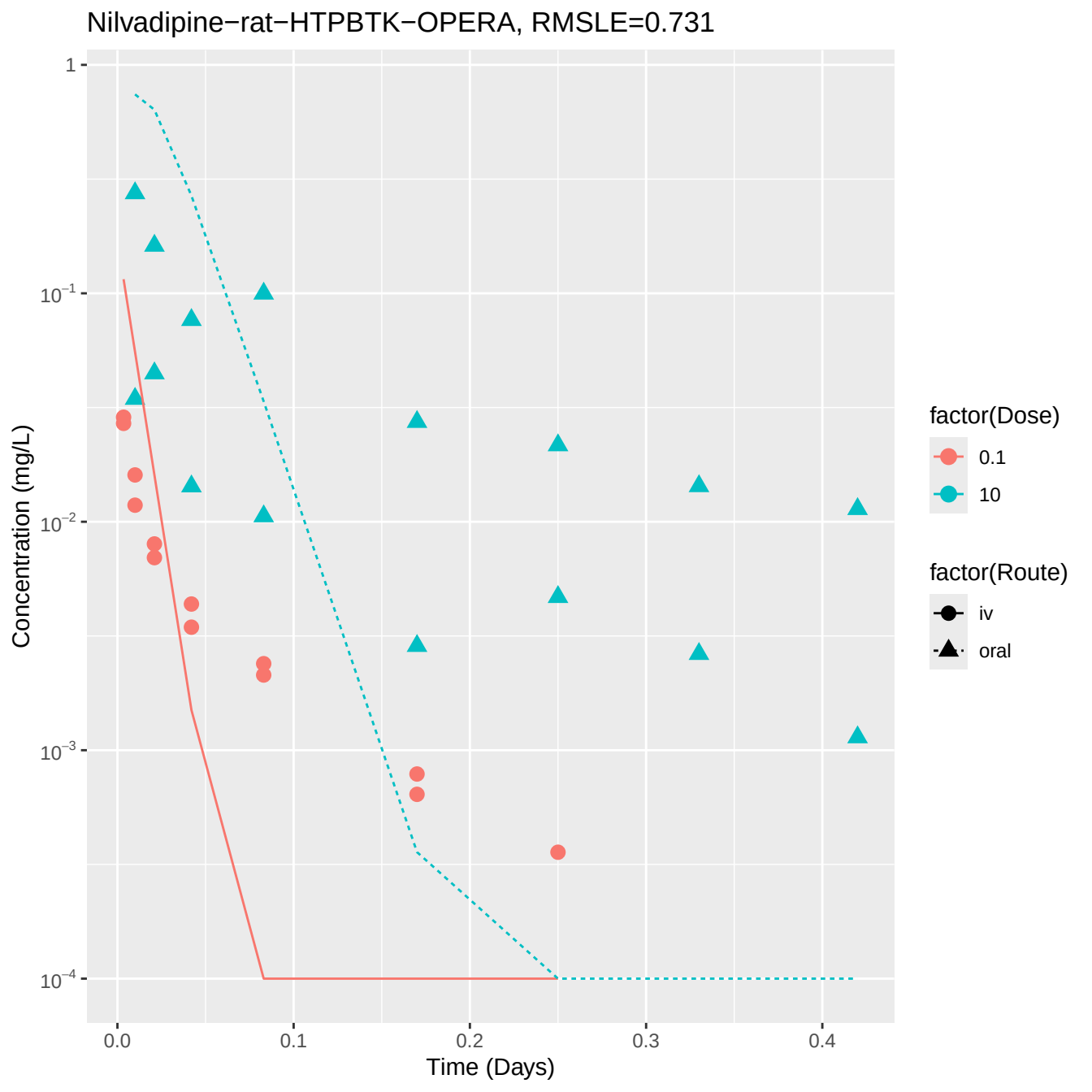

Dichloromethane−rat−HTPBTK−Ensemble, RMSLE=0.743
Concentration (mg/L)
10^2
10
1
10^{-1}
0.00
0.05
0.10
0.15
0.20
Time (Days)
factor(Route)
oral
factor(Dose)
21.2
125

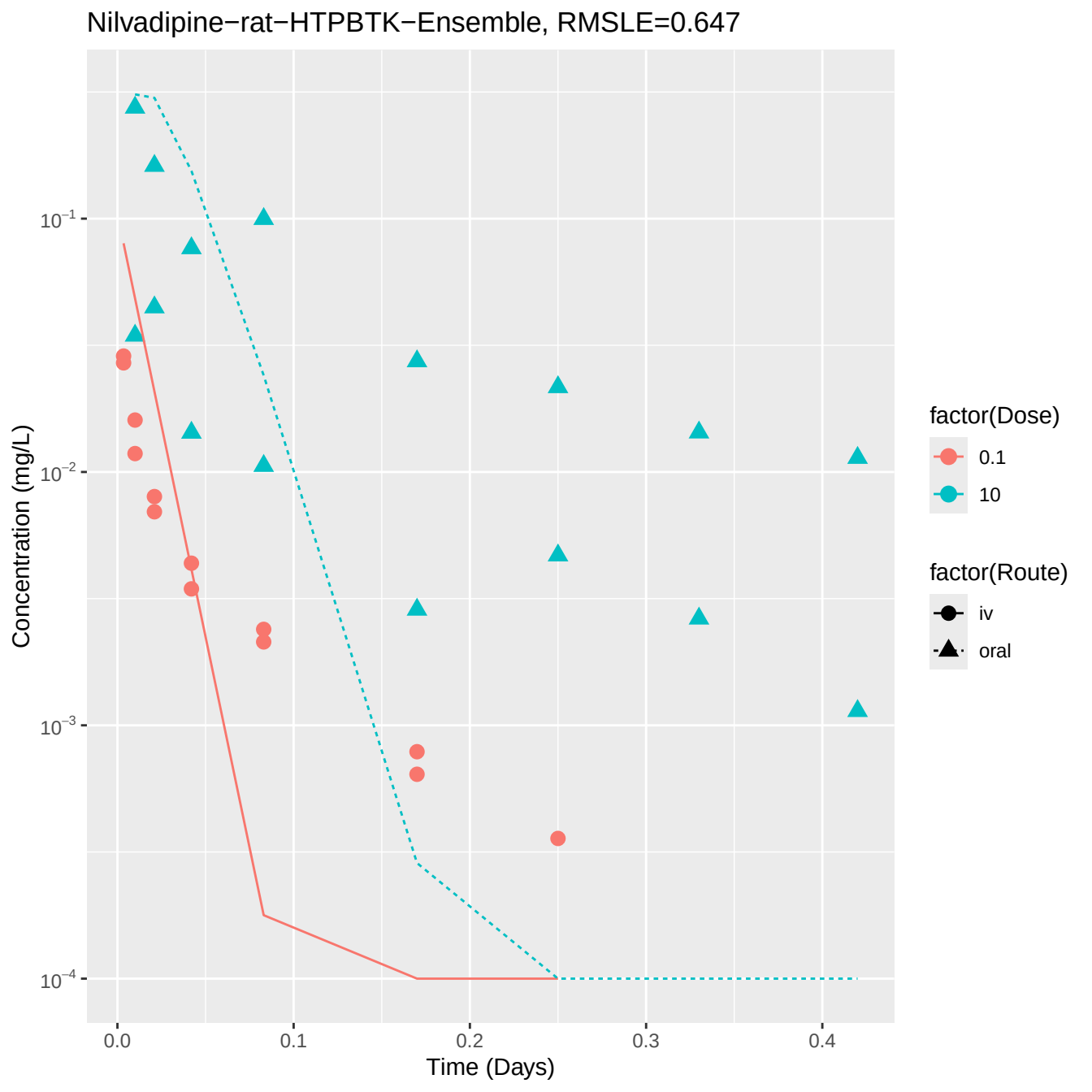

Nilvadipine−rat−HTPBTK−InVitro, RMSLE=2.46
Concentration (mg/L)
10
1
10^{-1}
10^{-2}
10^{-3}
0.0
0.1
0.2
0.3
0.4
Time (Days)
factor(Dose)
0.1
10
factor(Route)
iv
oral

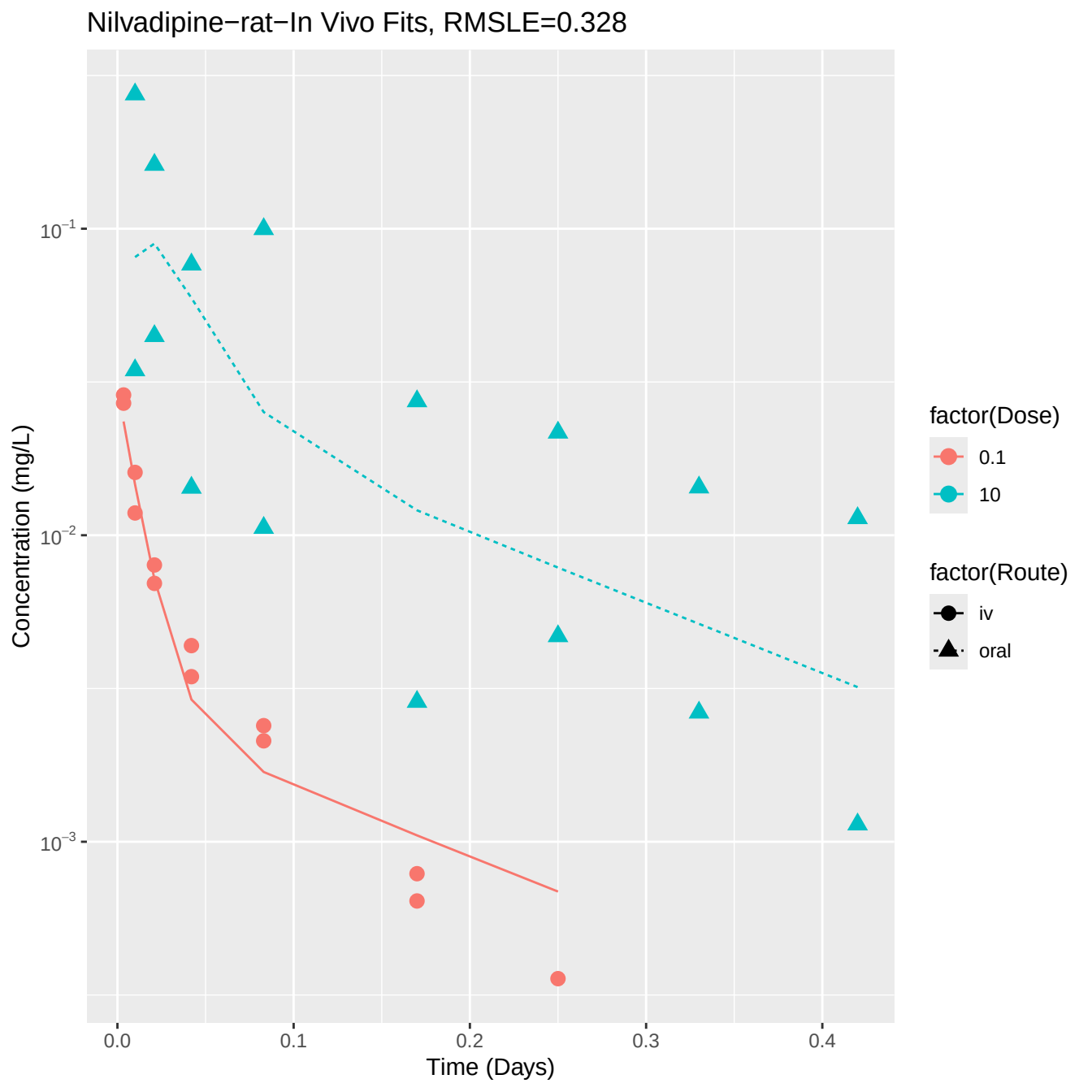

Nilvadipine−rat−HTPBTK−ADMET, RMSLE=0.645
Concentration (mg/L)
10^{-1}
10^{-2}
10^{-3}
10^{-4}
0.0
0.1
0.2
0.3
0.4
Time (Days)
factor(Dose)
0.1
10
factor(Route)
iv
oral

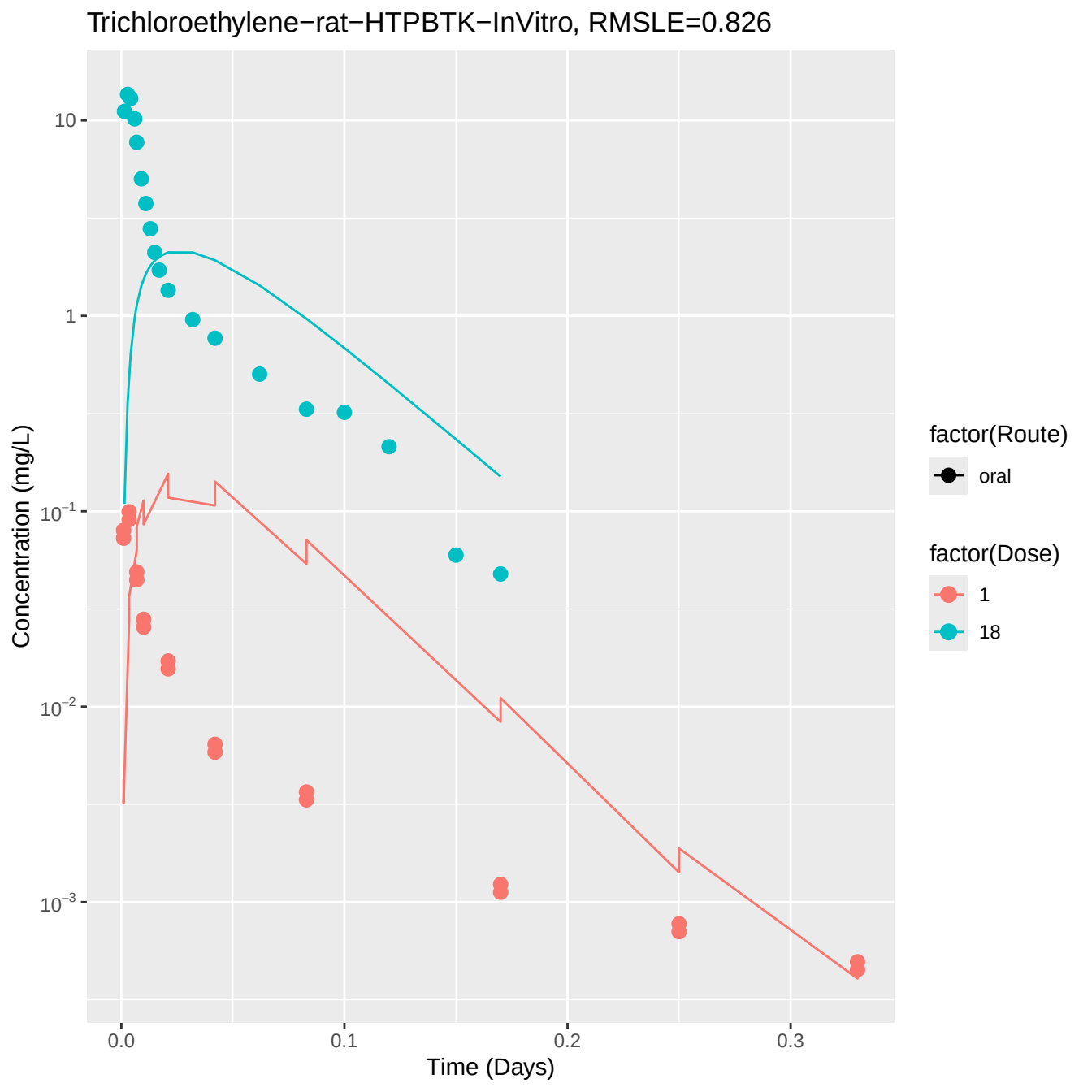

Nilvadipine−rat−HTPBTK−Dawson, RMSLE=0.946
Concentration (mg/L)
1
10^{-1}
10^{-2}
10^{-3}
10^{-4}
0.0
0.1
0.2
0.3
0.4
Time (Days)
factor(Dose)
0.1
10
factor(Route)
iv
oral

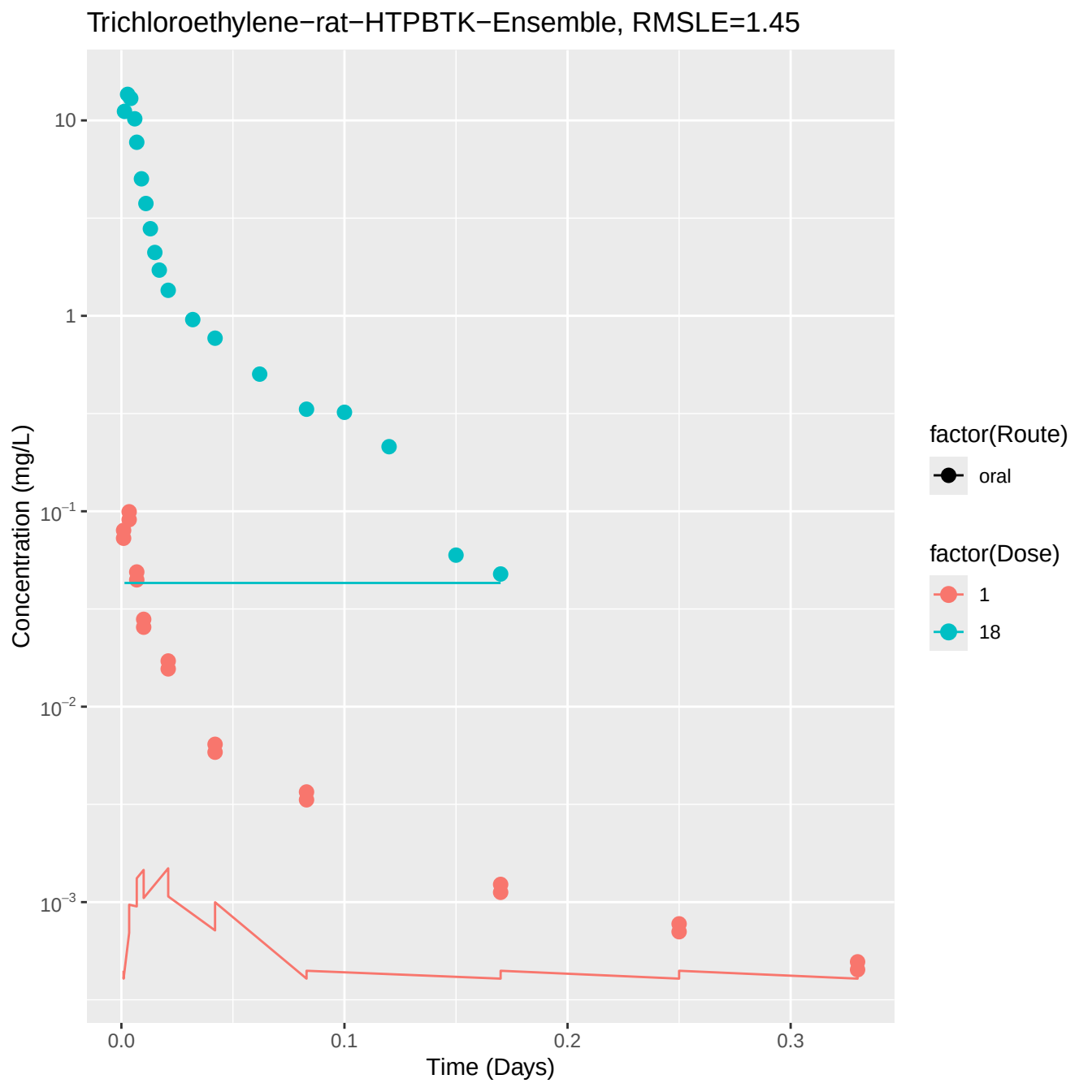

Nilvadipine−rat−HTPBTK−Pradeep, RMSLE=0.813
Concentration (mg/L)
1
10^{-1}
10^{-2}
10^{-3}
10^{-4}
0.0
0.1
0.2
0.3
0.4
Time (Days)
factor(Dose)
0.1
10
factor(Route)
iv
oral

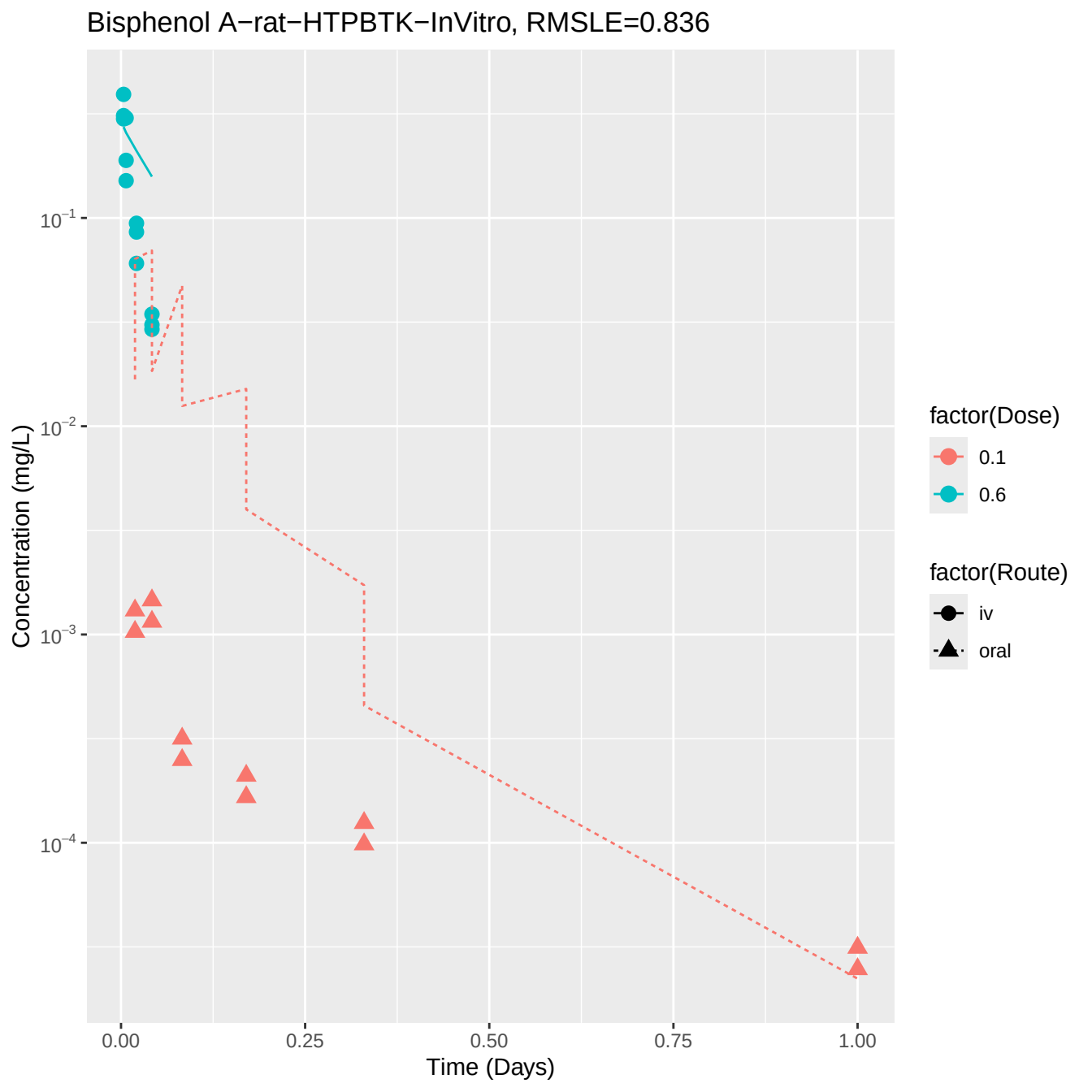

Nilvadipine−rat−HTPBTK−OPERA, RMSLE=0.731
Concentration (mg/L)
1
10^{-1}
10^{-2}
10^{-3}
10^{-4}
0.0
0.1
0.2
0.3
0.4
Time (Days)
factor(Dose)
0.1
10
factor(Route)
iv
oral

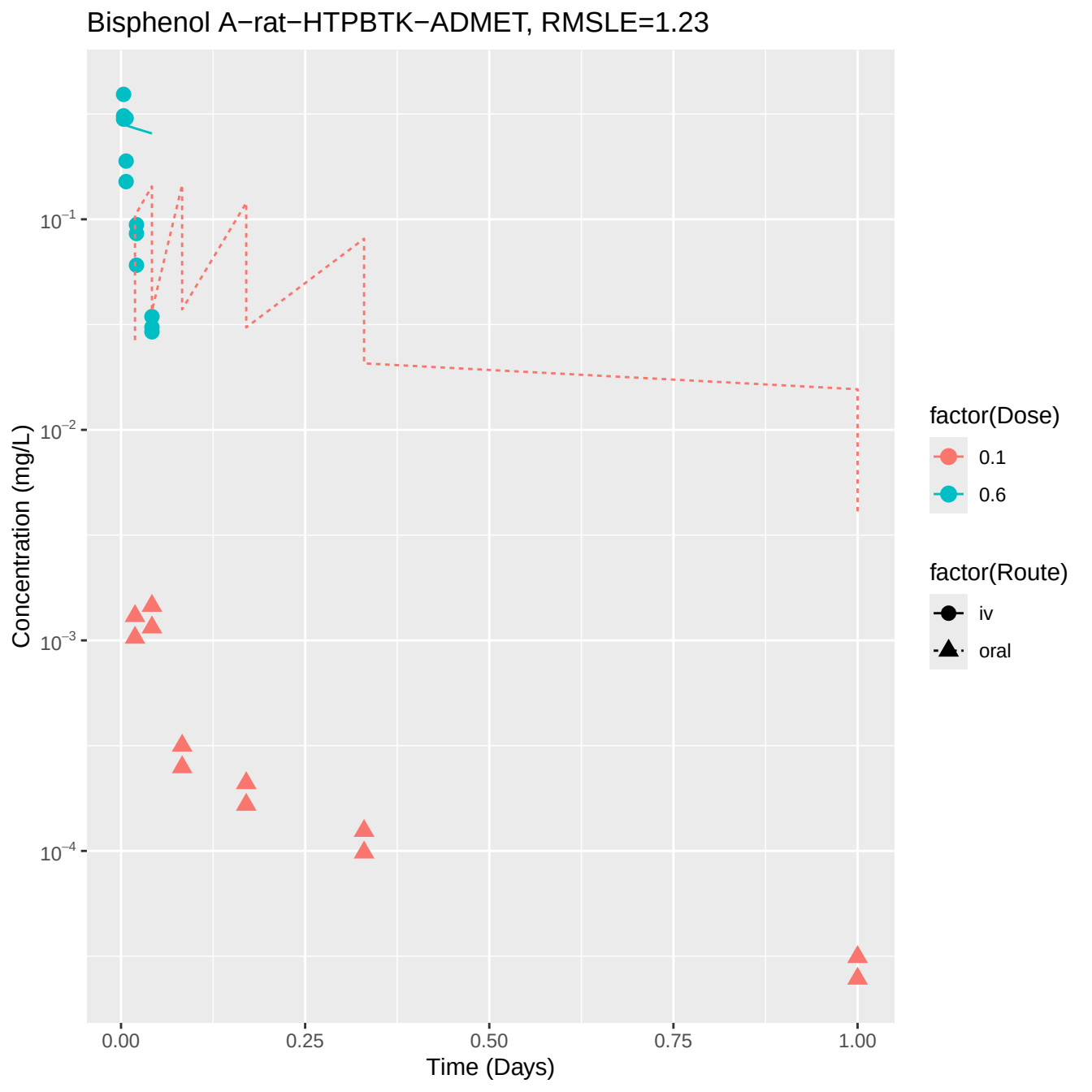

Nilvadipine−rat−HTPBTK−Ensemble, RMSLE=0.647
Concentration (mg/L)
10^{-1}
10^{-2}
10^{-3}
10^{-4}
0.0
0.1
0.2
0.3
0.4
Time (Days)
factor(Dose)
0.1
10
factor(Route)
iv
oral

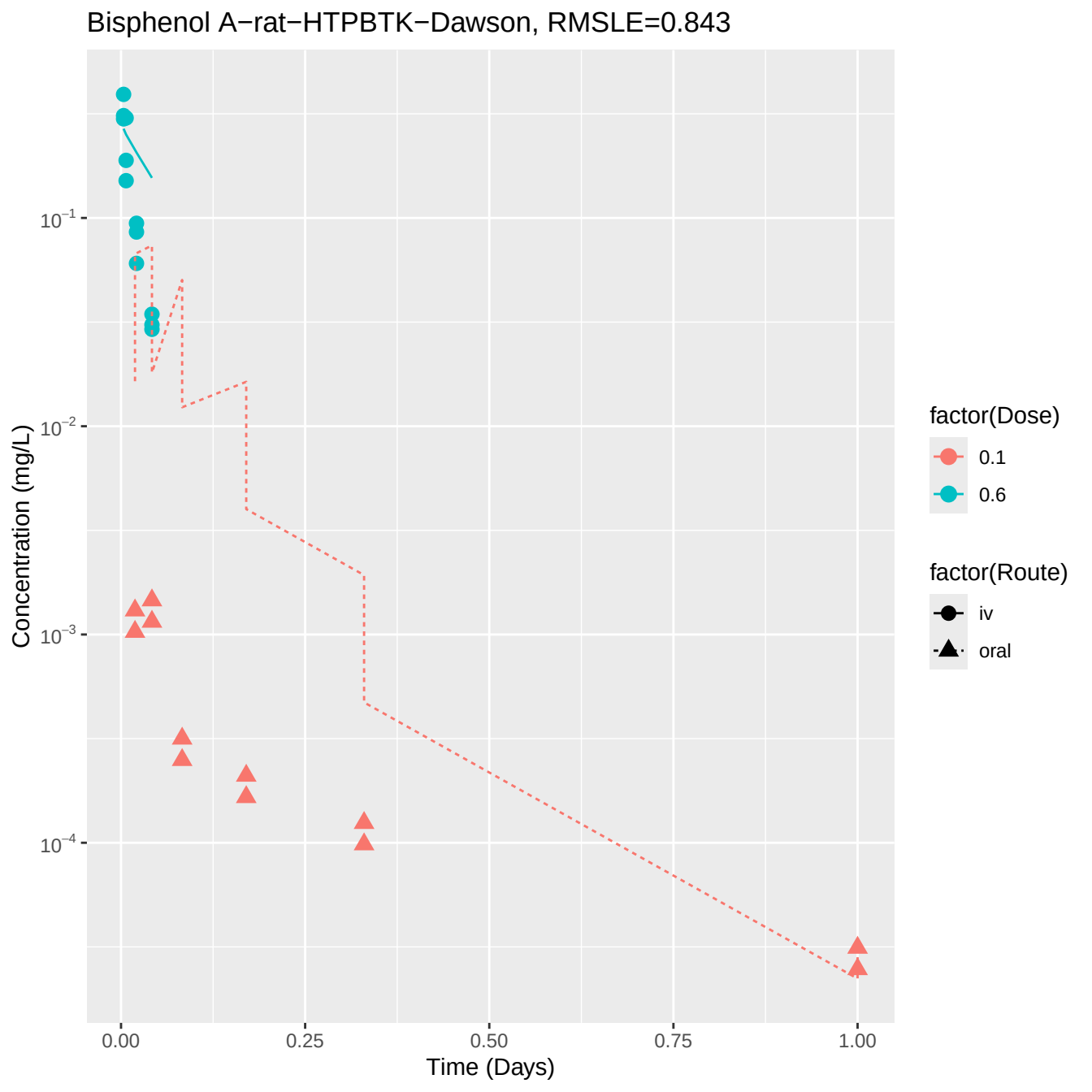

Nilvadipine−rat−In Vivo Fits, RMSLE=0.328
Concentration (mg/L)
10^{-1}
10^{-2}
10^{-3}
0.0
0.1
0.2
0.3
0.4
Time (Days)
factor(Dose)
0.1
10
factor(Route)
iv
oral

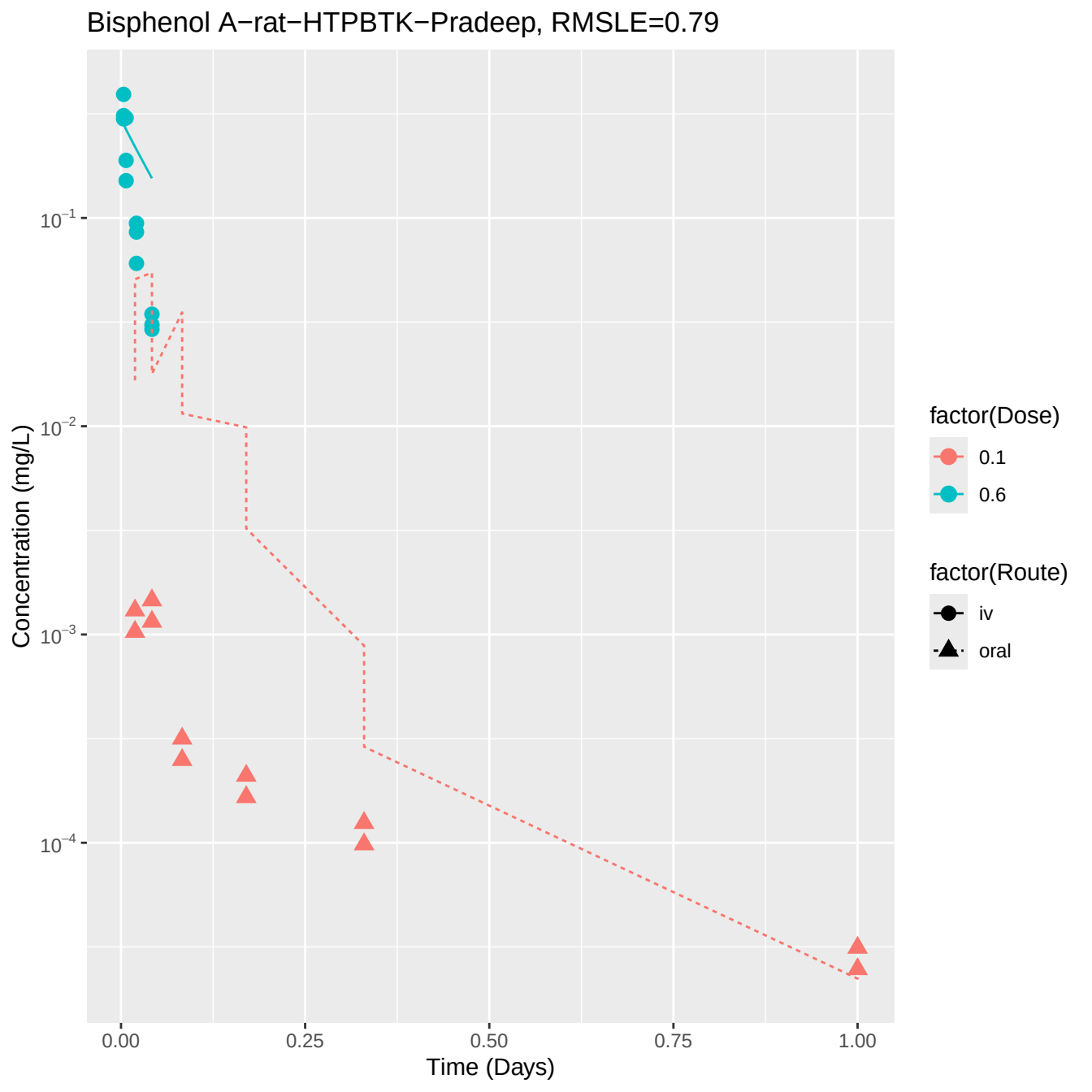

Trichloroethylene−rat−HTPBTK−InVitro, RMSLE=0.826
Concentration (mg/L)
10
1
10^{-1}
10^{-2}
10^{-3}
0.0
0.1
0.2
0.3
Time (Days)
factor(Route)
oral
factor(Dose)
1
18

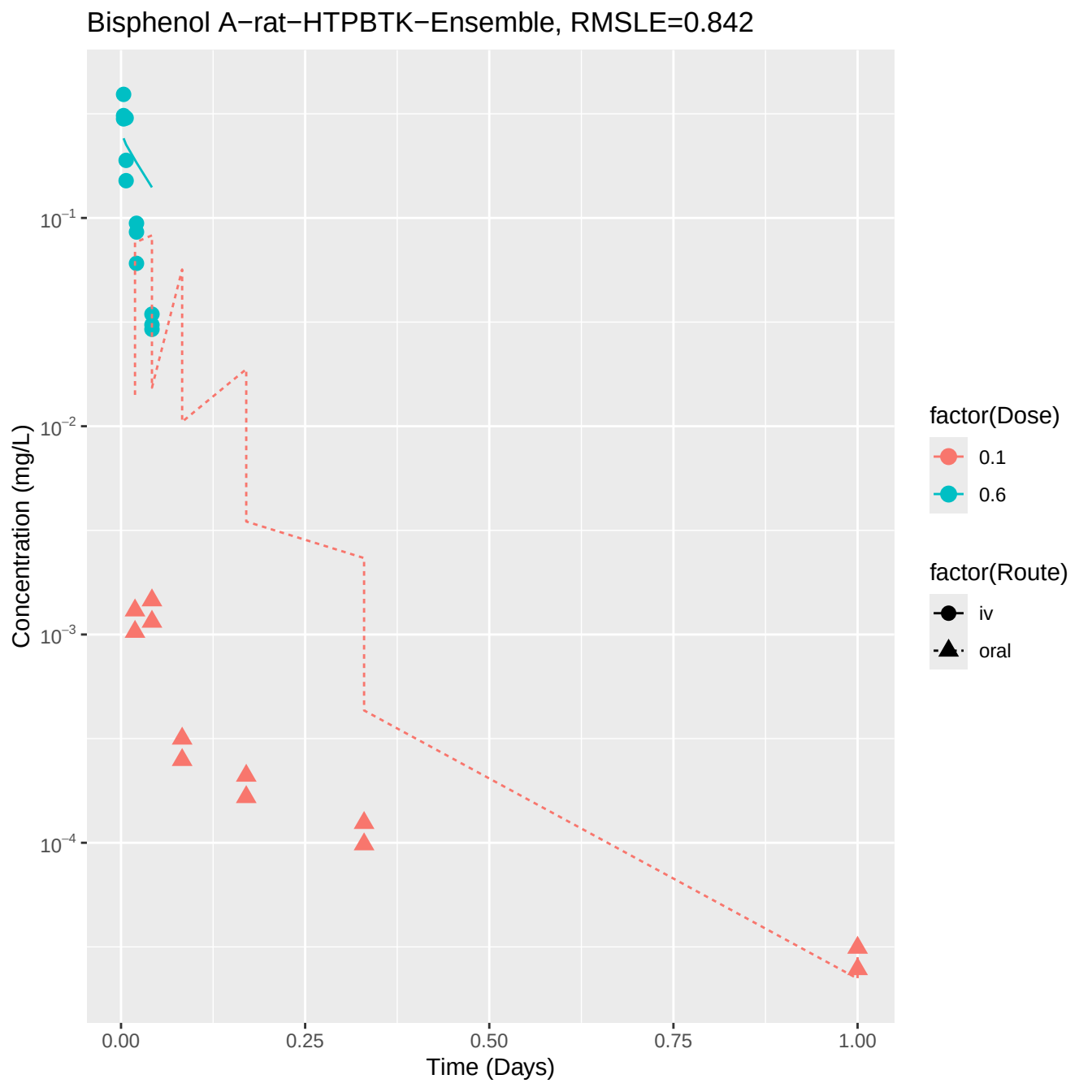

Trichloroethylene−rat−HTPBTK−Ensemble, RMSLE=1.45
Concentration (mg/L)
10
1
10^{-1}
10^{-2}
10^{-3}
0.0
0.1
0.2
0.3
Time (Days)
factor(Route)
oral
factor(Dose)
1
18

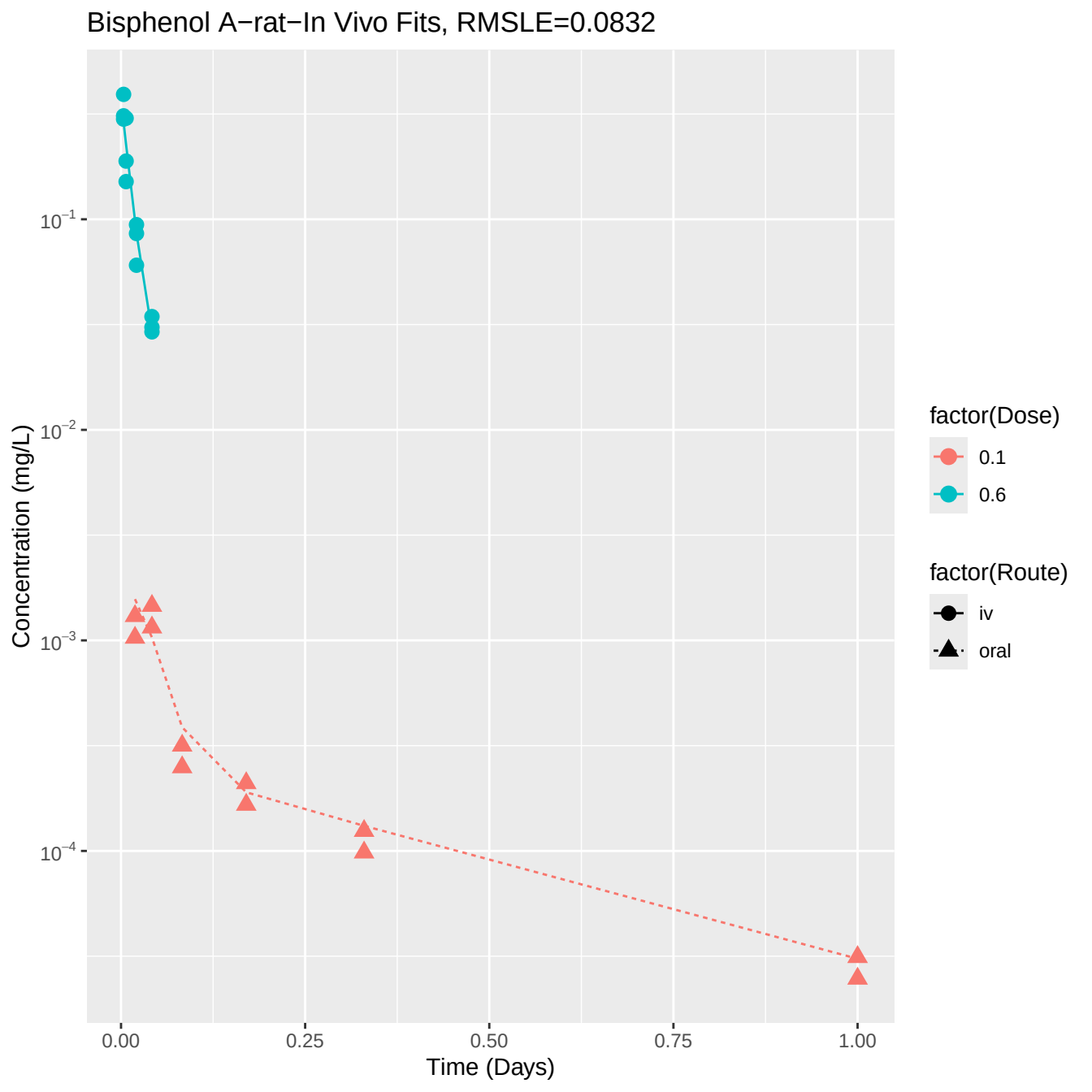

Bisphenol A−rat−HTPBTK−InVitro, RMSLE=0.836
Concentration (mg/L)
10^{-1}
10^{-2}
10^{-3}
10^{-4}
0.00
0.25
0.50
0.75
1.00
Time (Days)
factor(Dose)
0.1
0.6
factor(Route)
iv
oral

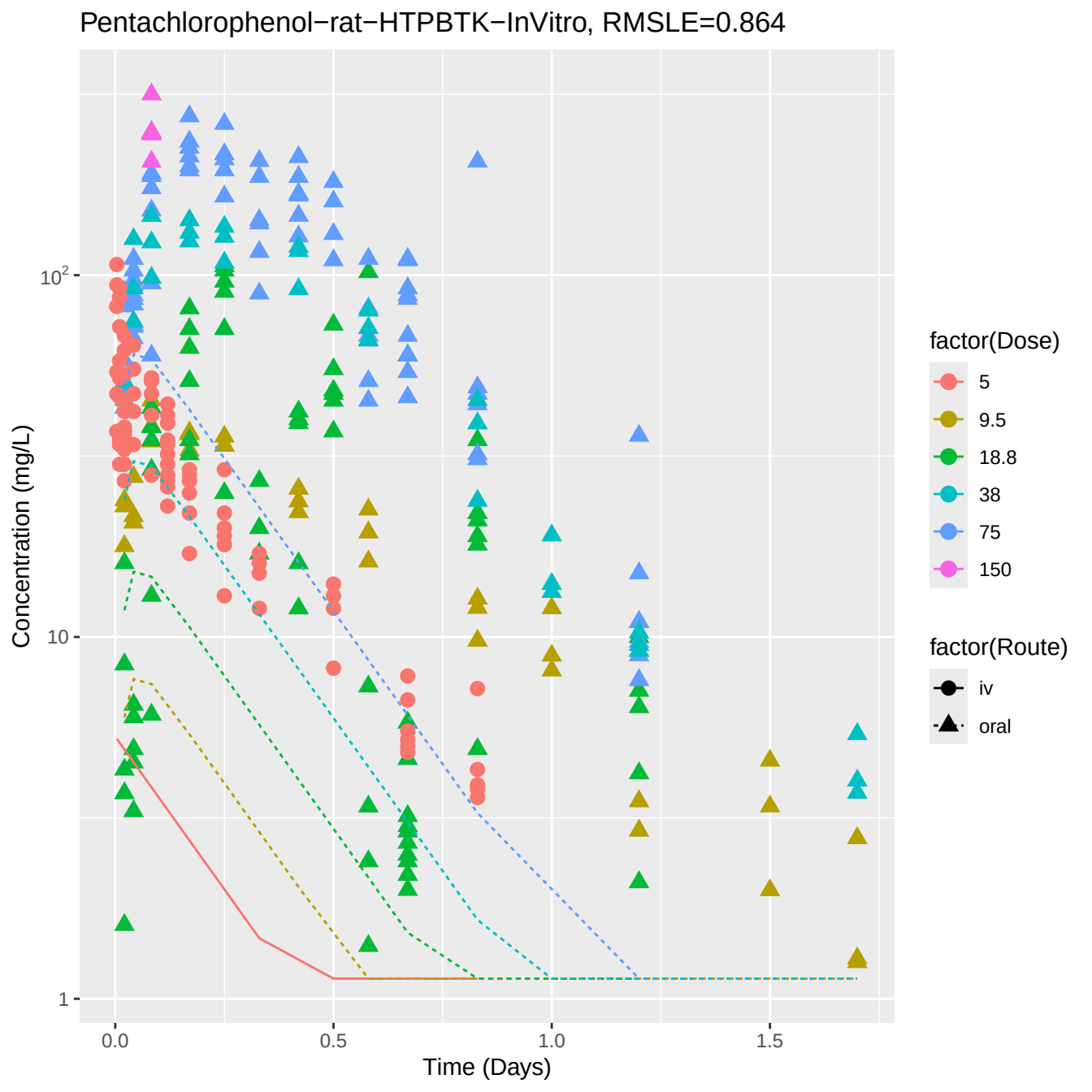

Bisphenol A−rat−HTPBTK−ADMET, RMSLE=1.23
Concentration (mg/L)
10^{-1}
10^{-2}
10^{-3}
10^{-4}
0.00
0.25
0.50
0.75
1.00
Time (Days)
factor(Dose)
0.1
0.6
factor(Route)
iv
oral

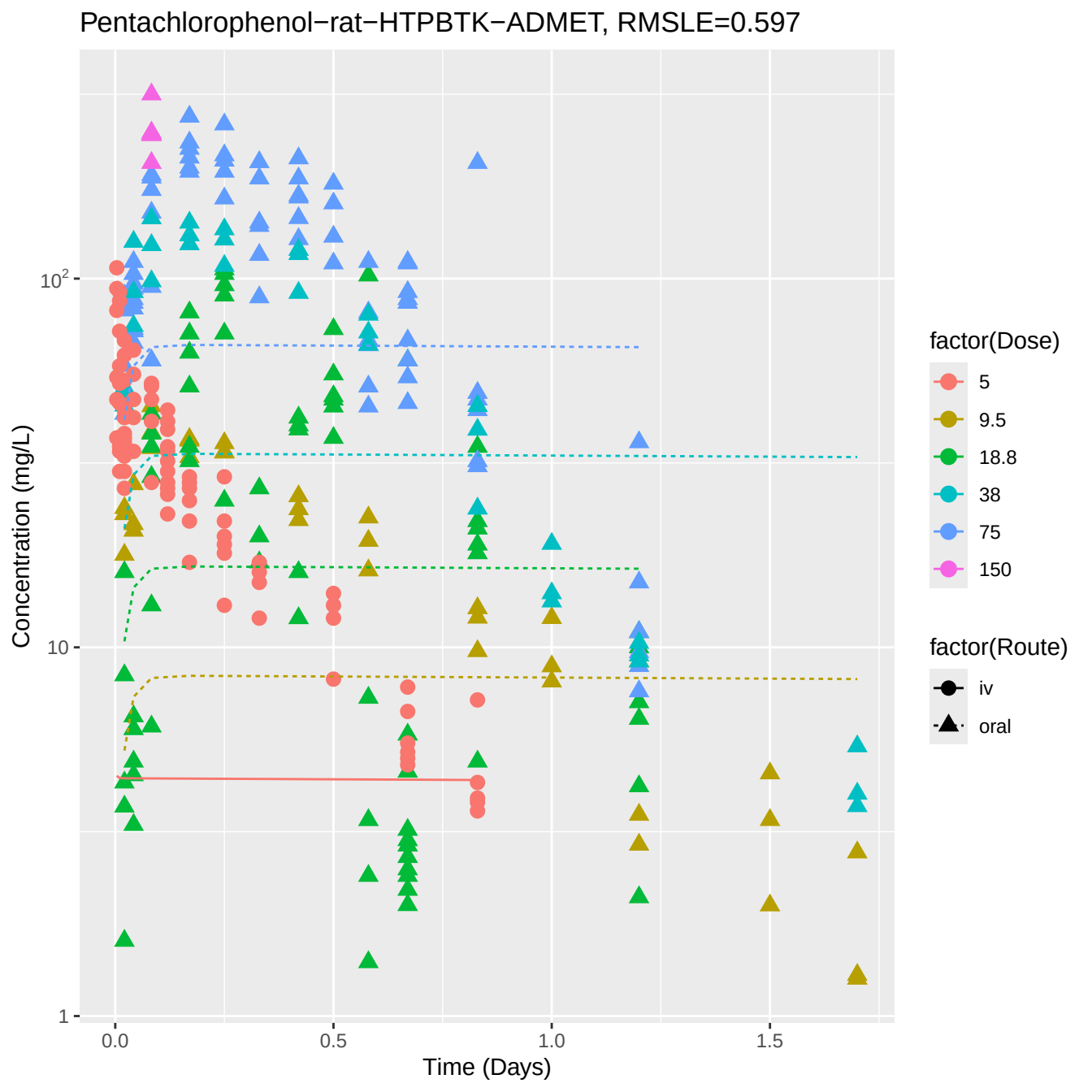

Bisphenol A−rat−HTPBTK−Dawson, RMSLE=0.843
Concentration (mg/L)
10^{-1}
10^{-2}
10^{-3}
10^{-4}
0.00
0.25
0.50
0.75
1.00
Time (Days)
factor(Dose)
0.1
0.6
factor(Route)
iv
oral

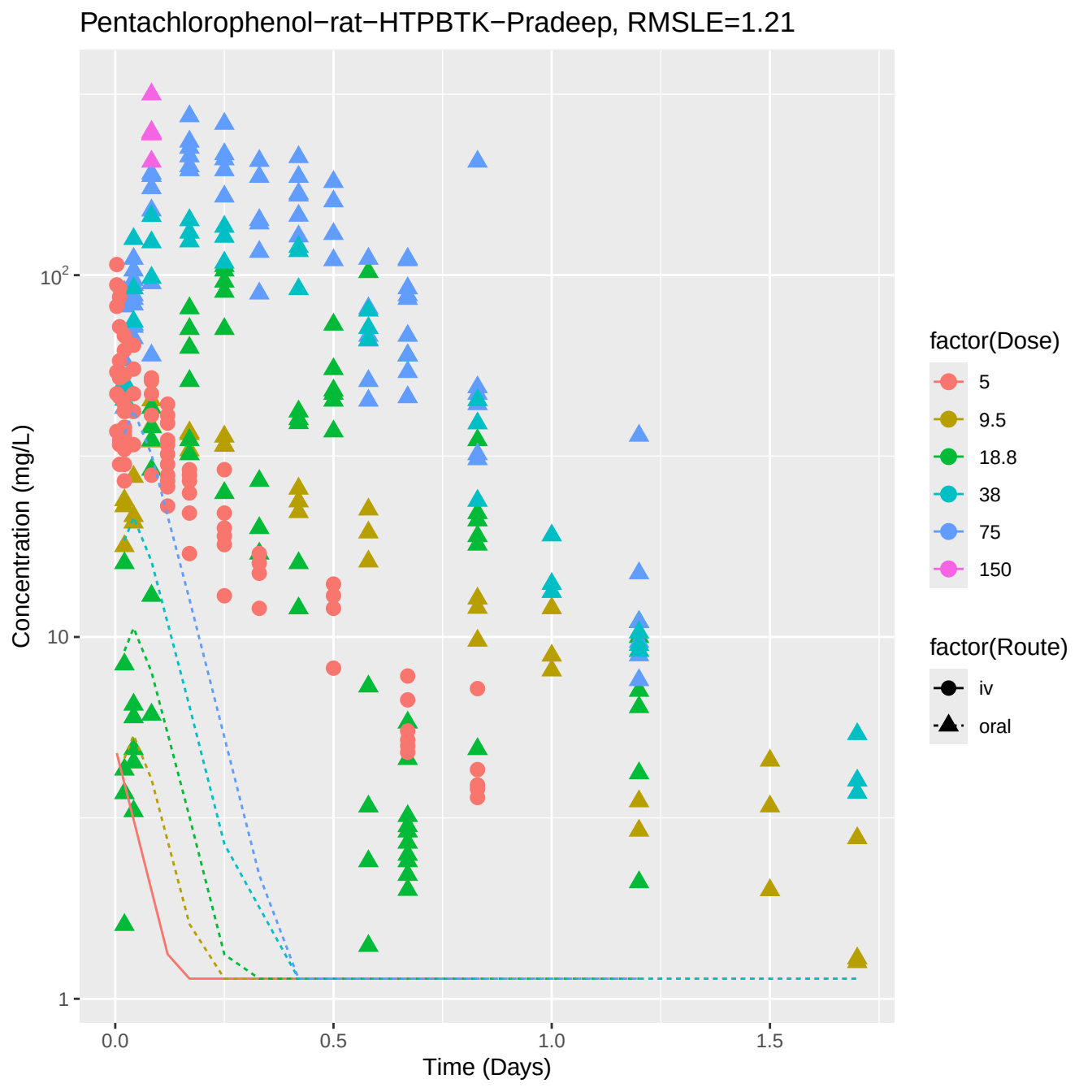

Bisphenol A−rat−HTPBTK−Pradeep, RMSLE=0.79
Concentration (mg/L)
10^{-1}
10^{-2}
10^{-3}
10^{-4}
0.00
0.25
0.50
0.75
1.00
Time (Days)
factor(Dose)
0.1
0.6
factor(Route)
iv
oral

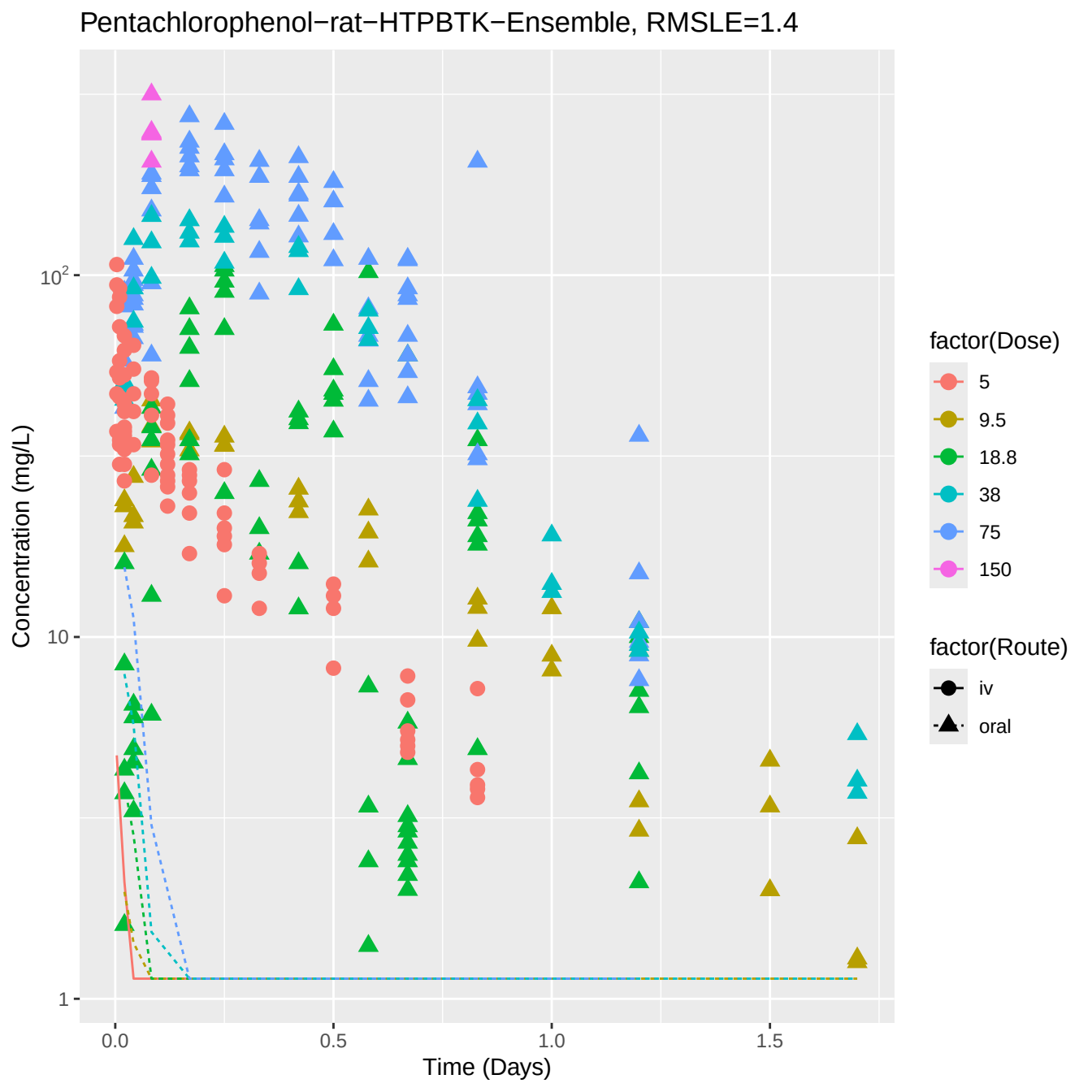

Bisphenol A−rat−HTPBTK−Ensemble, RMSLE=0.842
Concentration (mg/L)
10^{-1}
10^{-2}
10^{-3}
10^{-4}
0.00
0.25
0.50
0.75
1.00
Time (Days)
factor(Dose)
0.1
0.6
factor(Route)
iv
oral

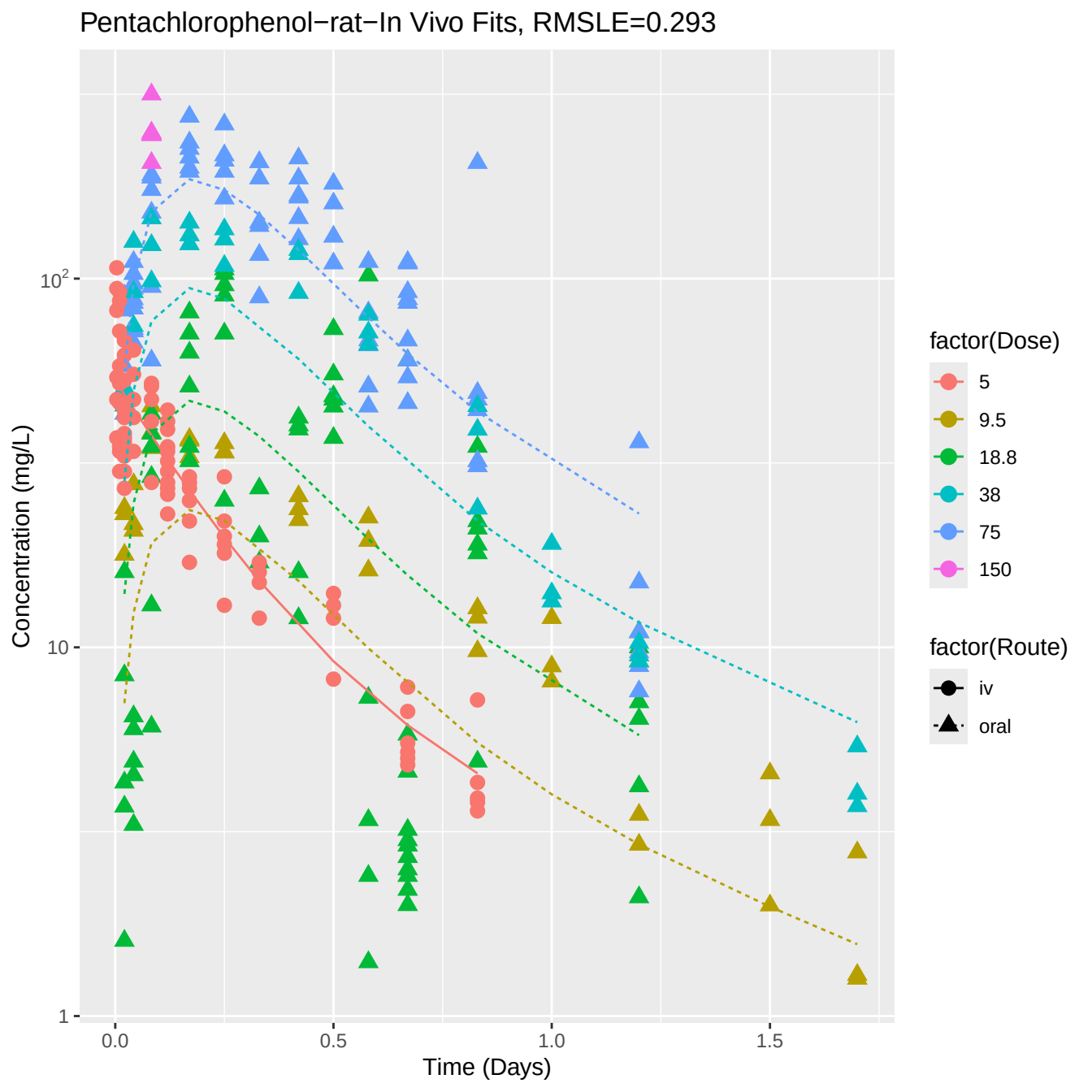

Bisphenol A−rat−In Vivo Fits, RMSLE=0.0832
Concentration (mg/L)
10^{-1}
10^{-2}
10^{-3}
10^{-4}
0.00
0.25
0.50
0.75
1.00
Time (Days)
factor(Dose)
0.1
0.6
factor(Route)
iv
oral

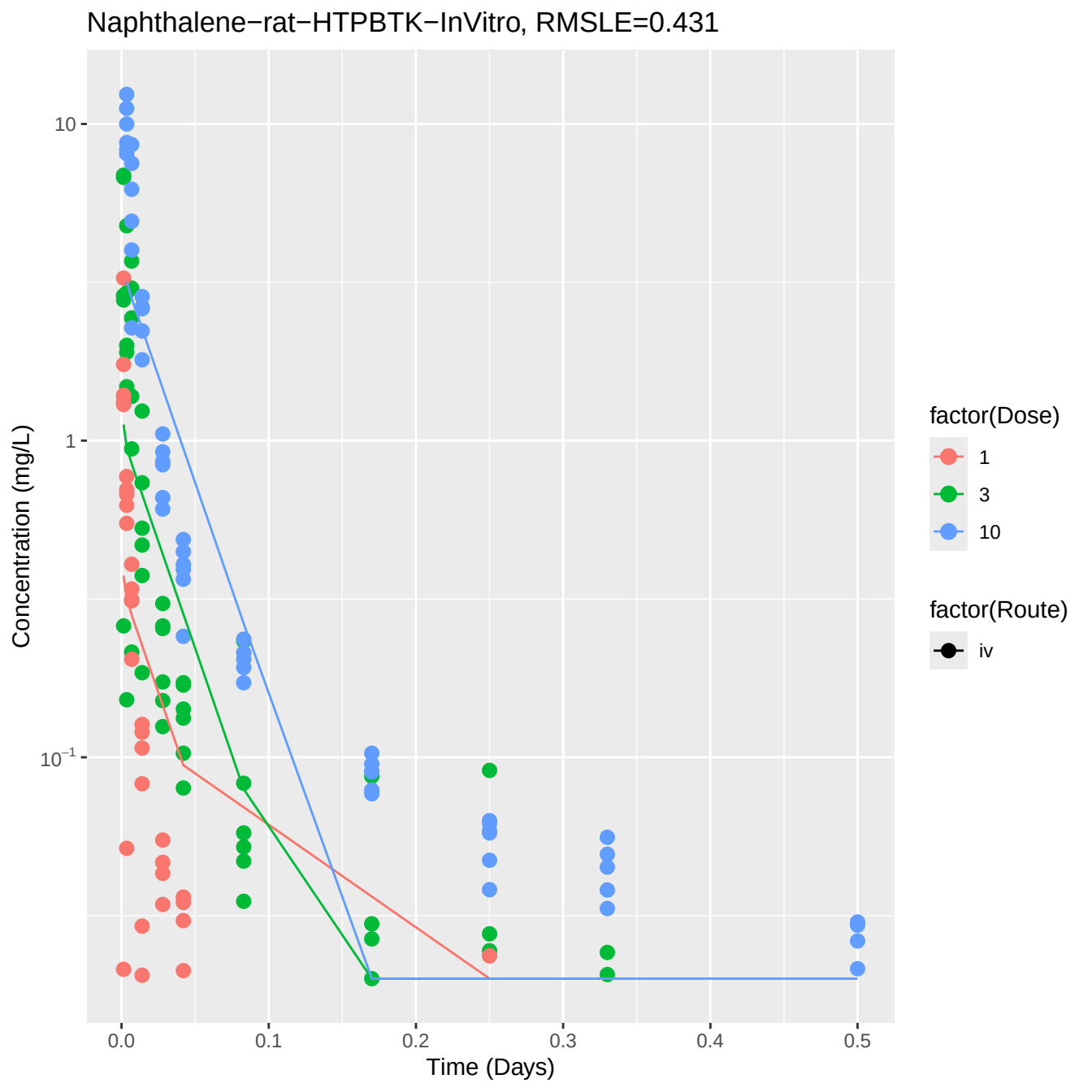

Pentachlorophenol−rat−HTPBTK−InVitro, RMSLE=0.864
Concentration (mg/L)
10^2
10
1
0.0
0.5
1.0
1.5
Time (Days)
factor(Dose)
5
9.5
18.8
38
75
150
factor(Route)
iv
oral

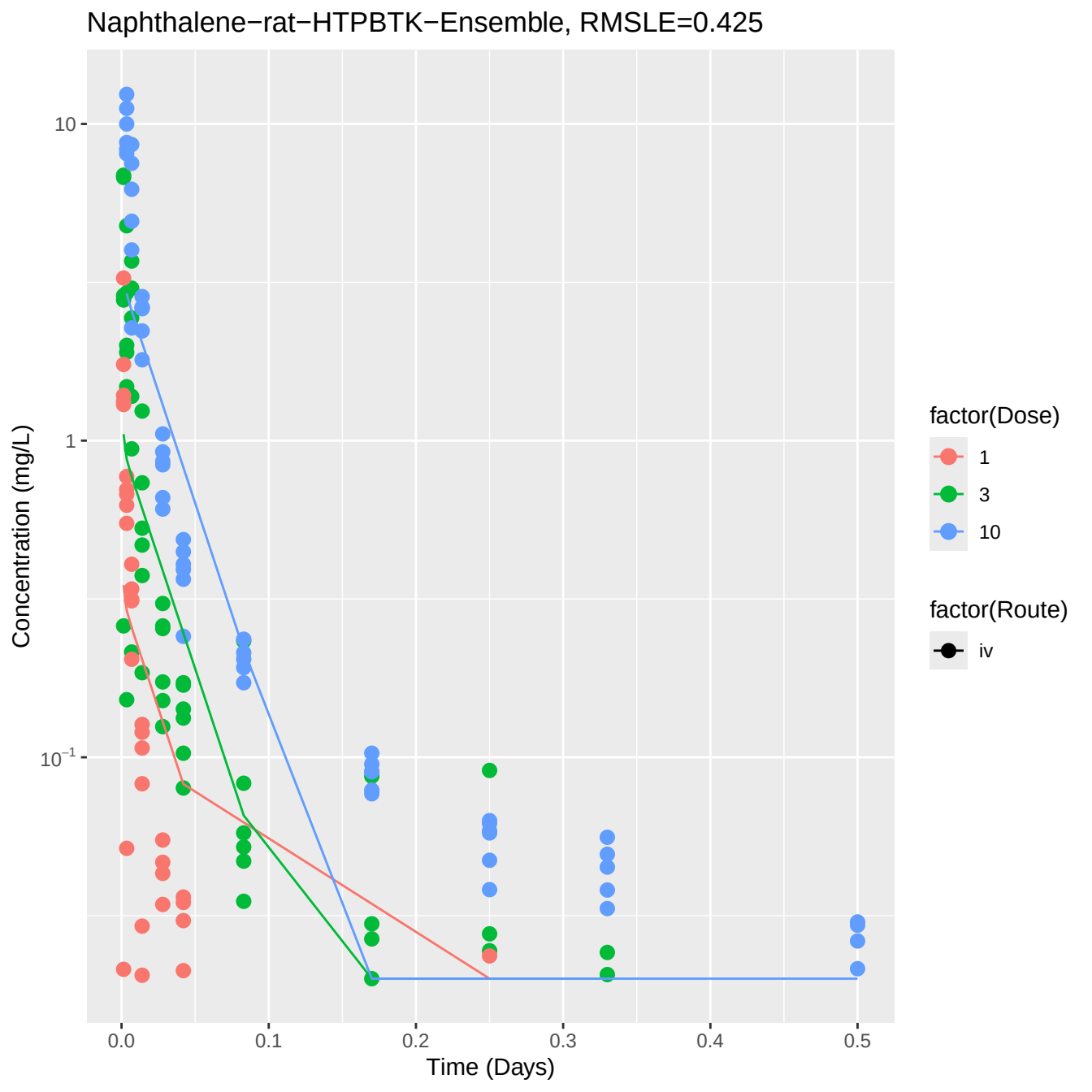

Pentachlorophenol−rat−HTPBTK−ADMET, RMSLE=0.597
Concentration (mg/L)
10^2
10
1
0.0
0.5
1.0
1.5
Time (Days)
factor(Dose)
5
9.5
18.8
38
75
150
factor(Route)
iv
oral

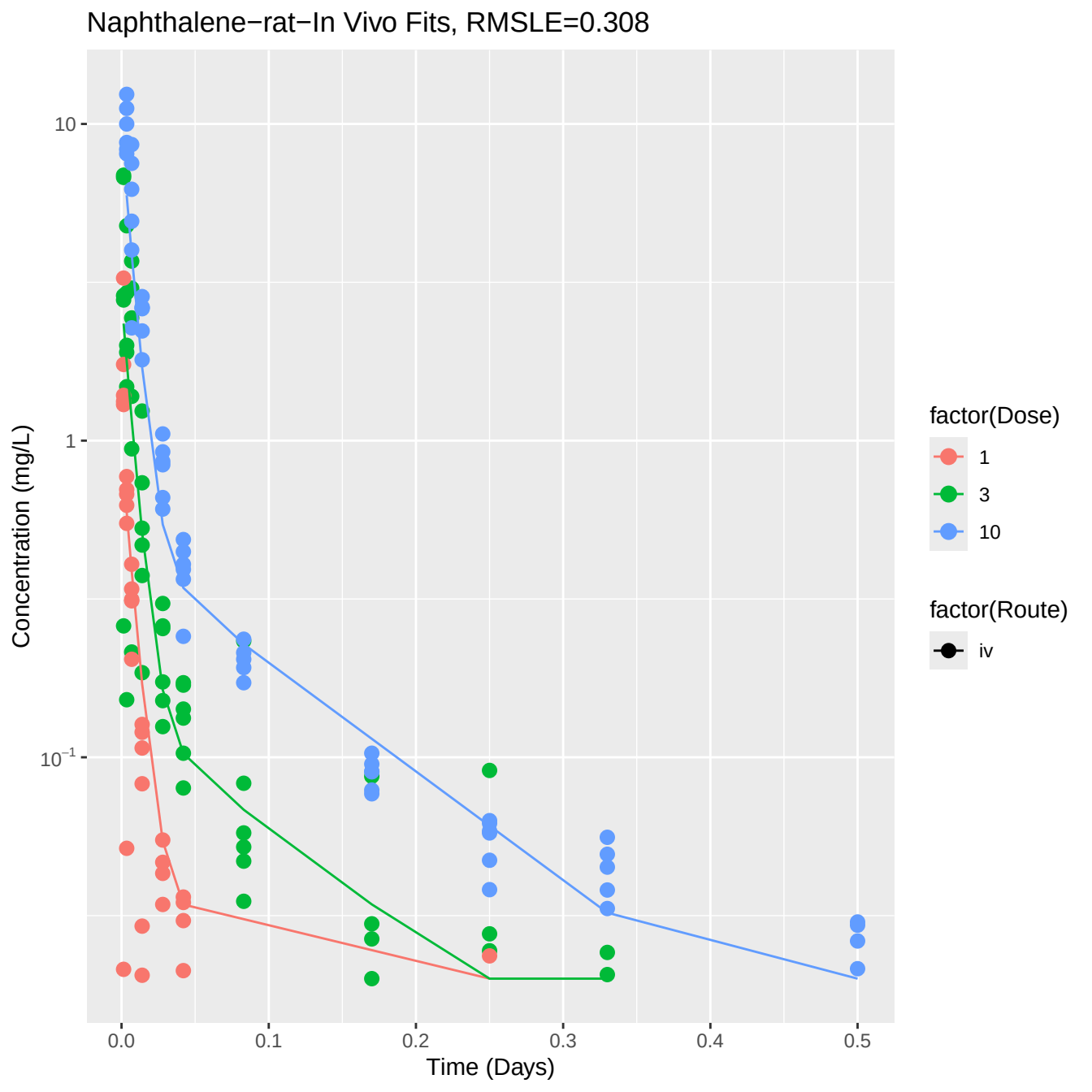

Pentachlorophenol−rat−HTPBTK−Pradeep, RMSLE=1.21
Concentration (mg/L)
10^2
10
1
0.0
0.5
1.0
1.5
Time (Days)
factor(Dose)
5
9.5
18.8
38
75
150
factor(Route)
iv
oral


Pentachlorophenol−rat−HTPBTK−Ensemble, RMSLE=1.4
Concentration (mg/L)
10^2
10
1
0.0
0.5
1.0
1.5
Time (Days)
factor(Dose)
5
9.5
18.8
38
75
150
factor(Route)
iv
oral

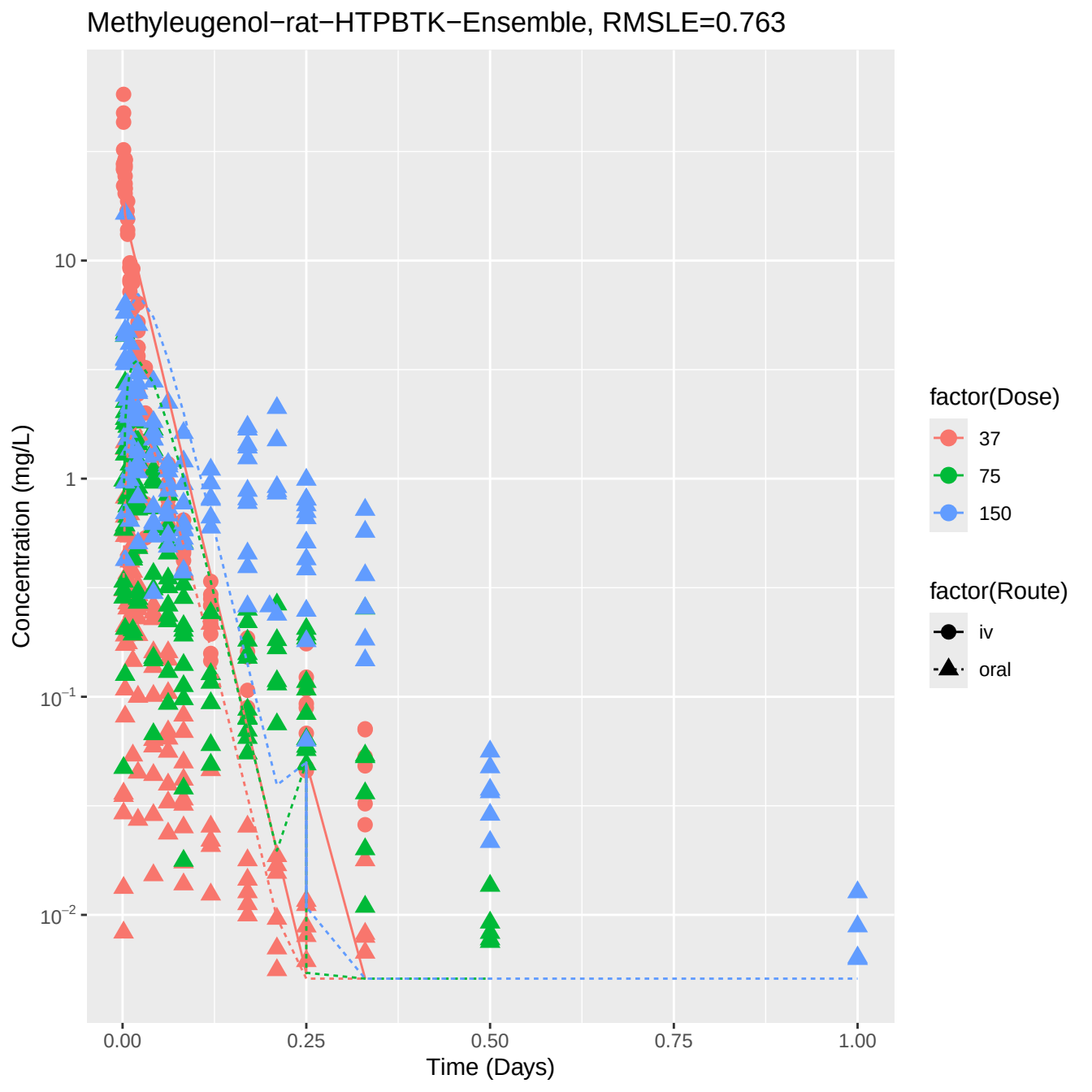

Pentachlorophenol−rat−In Vivo Fits, RMSLE=0.293
Concentration (mg/L)
10^2
10
1
0.0
0.5
1.0
1.5
Time (Days)
factor(Dose)
5
9.5
18.8
38
75
150
factor(Route)
iv
oral

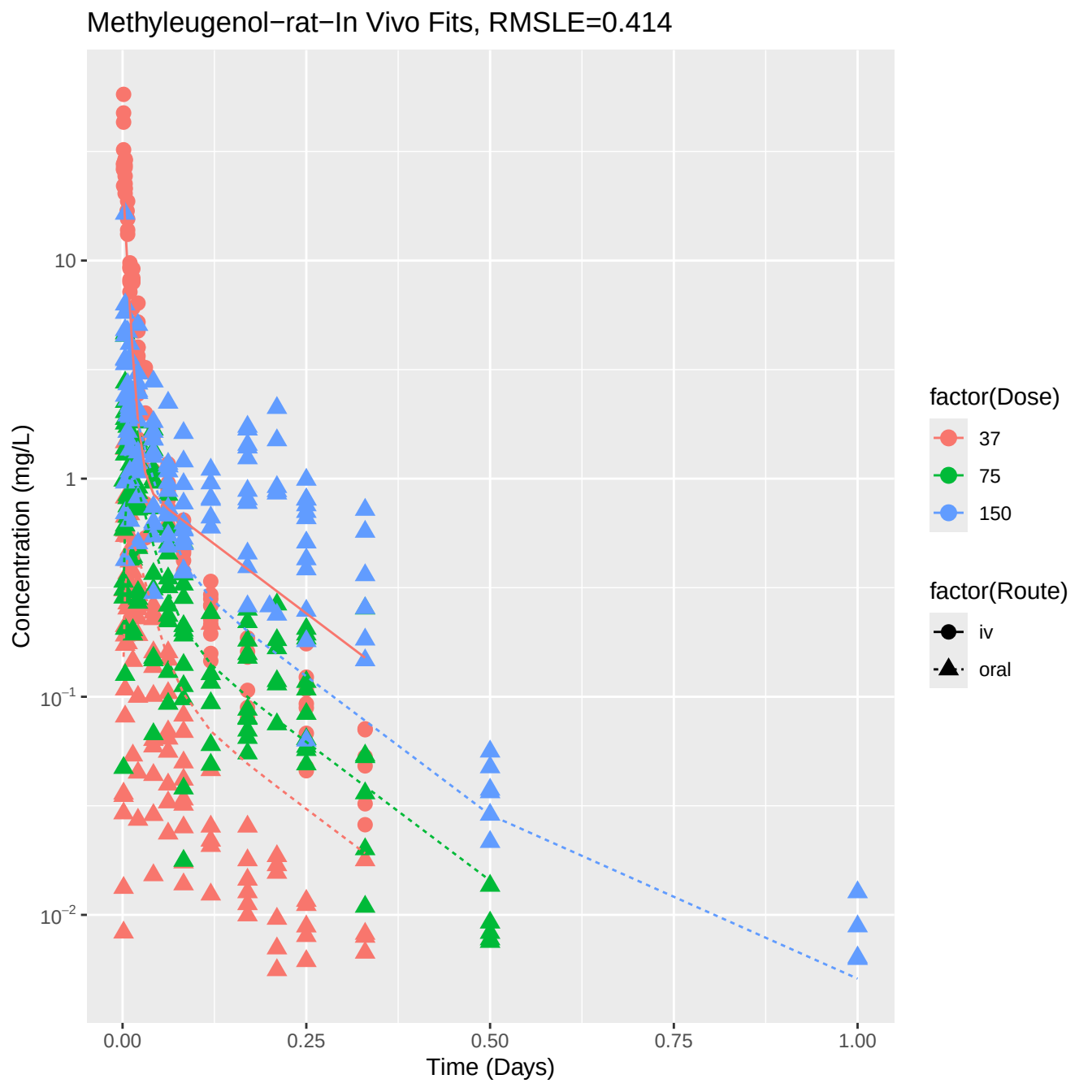

Naphthalene−rat−HTPBTK−InVitro, RMSLE=0.431
Concentration (mg/L)
10
1
10^{-1}
0.0
0.1
0.2
0.3
0.4
0.5
Time (Days)
factor(Dose)
1
3
10
factor(Route)
iv

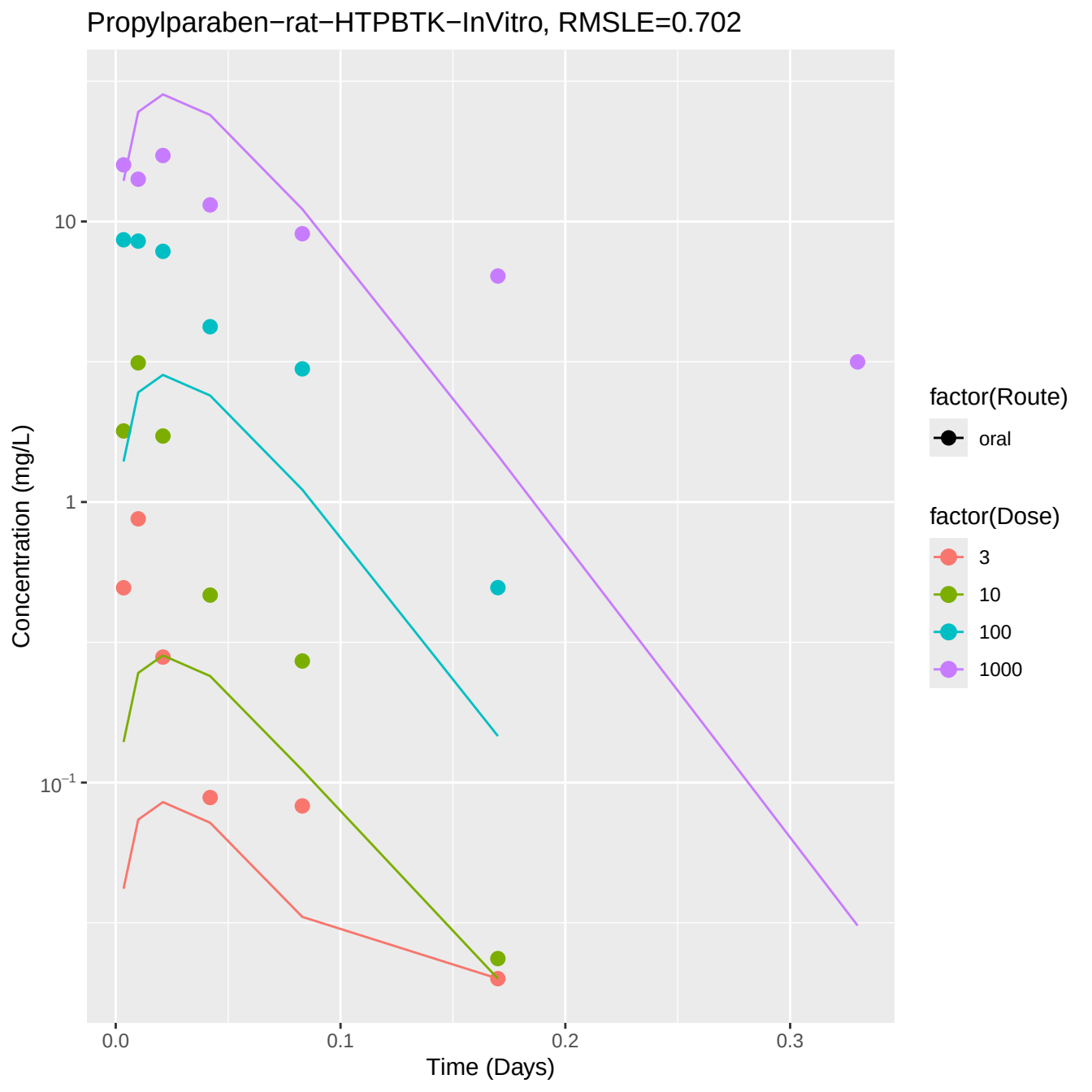

Naphthalene−rat−HTPBTK−Ensemble, RMSLE=0.425
Concentration (mg/L)
10
1
10^{-1}
0.0
0.1
0.2
0.3
0.4
0.5
Time (Days)
factor(Dose)
1
3
10
factor(Route)
iv

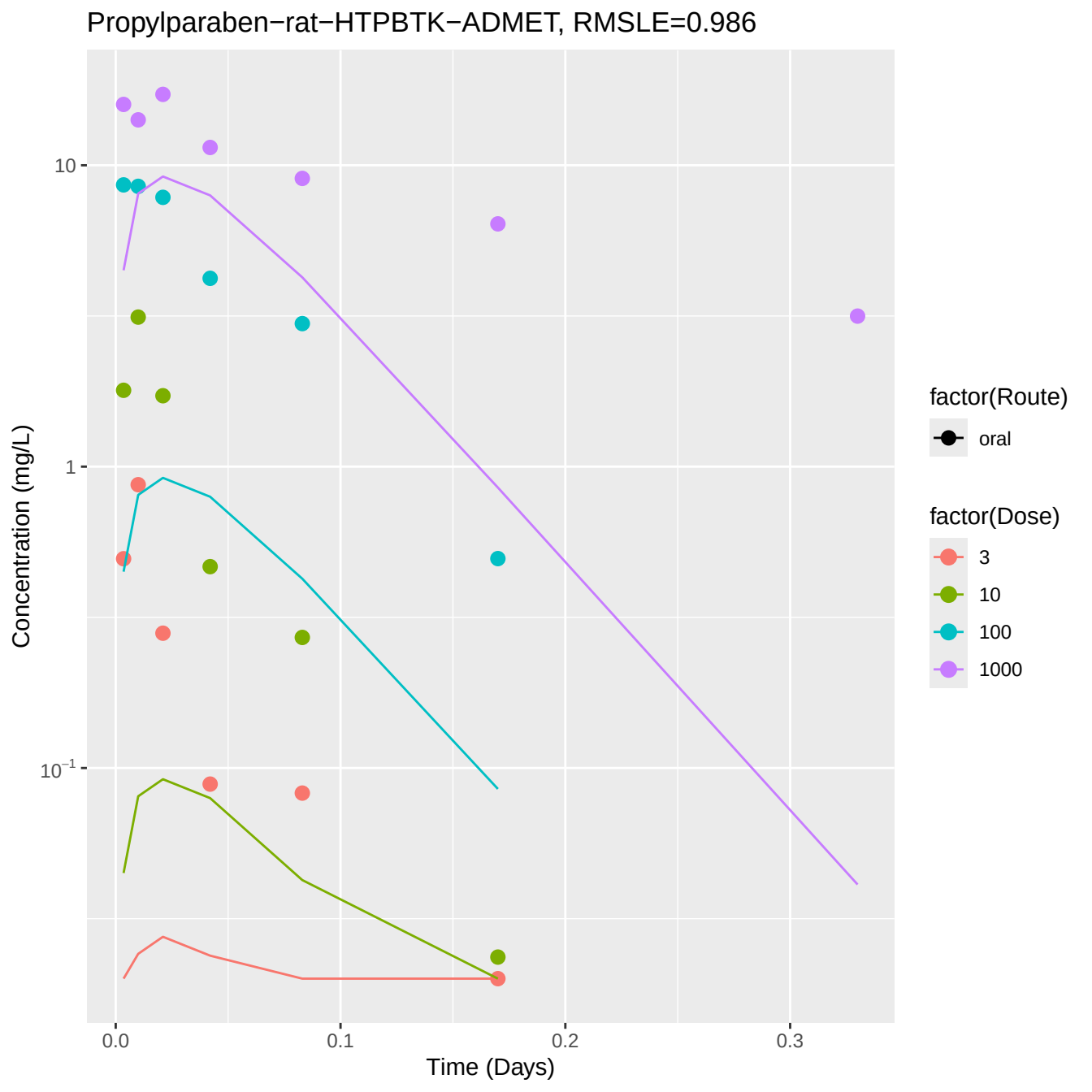

Naphthalene−rat−In Vivo Fits, RMSLE=0.308
Concentration (mg/L)
10
1
10^{-1}
0.0
0.1
0.2
0.3
0.4
0.5
Time (Days)
factor(Dose)
1
3
10
factor(Route)
iv

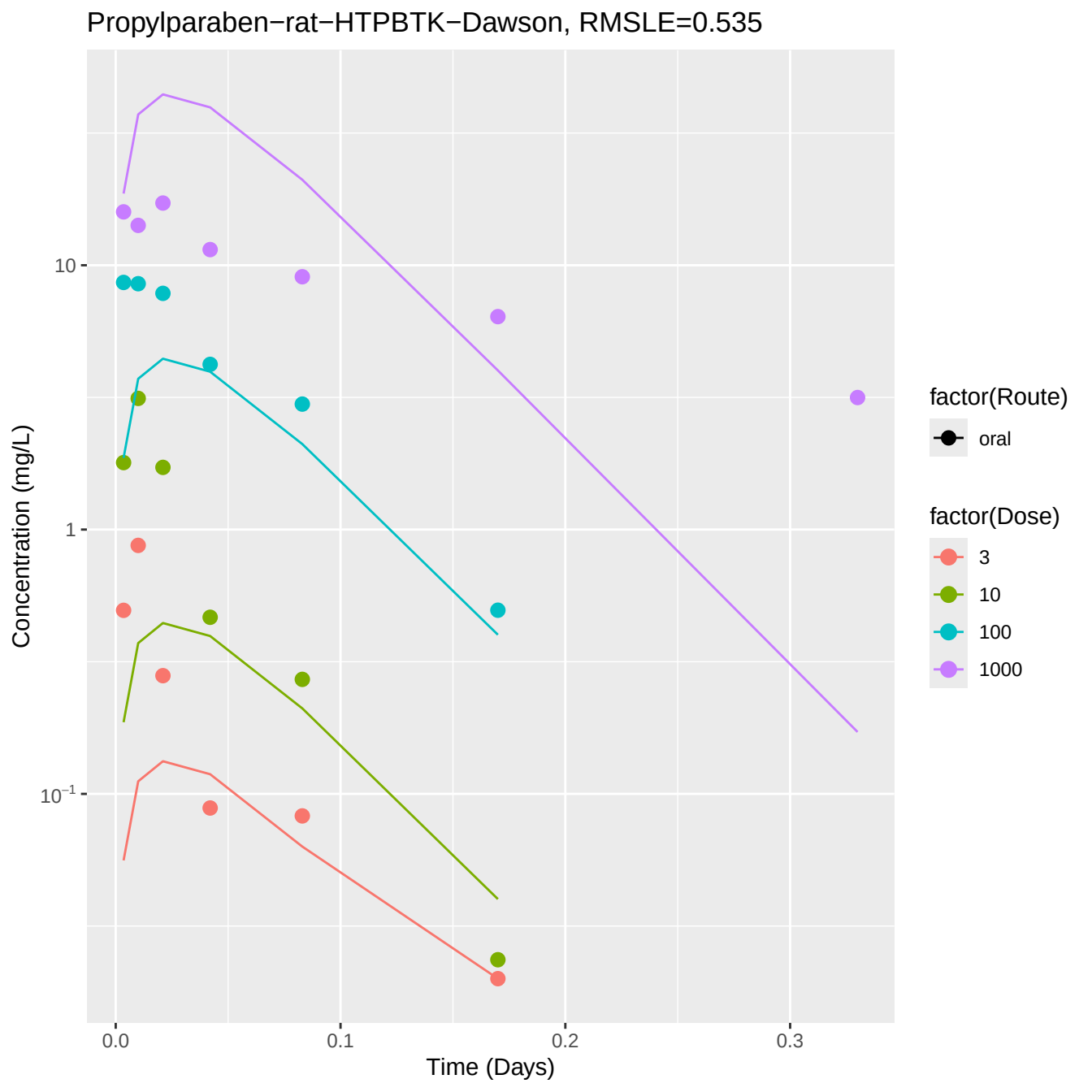

Methyleugenol−rat−HTPBTK−InVitro, RMSLE=0.78
Concentration (mg/L)
10
1
10^{-1}
10^{-2}
0.00
0.25
0.50
0.75
1.00
Time (Days)
factor(Dose)
37
75
150
factor(Route)
iv
oral

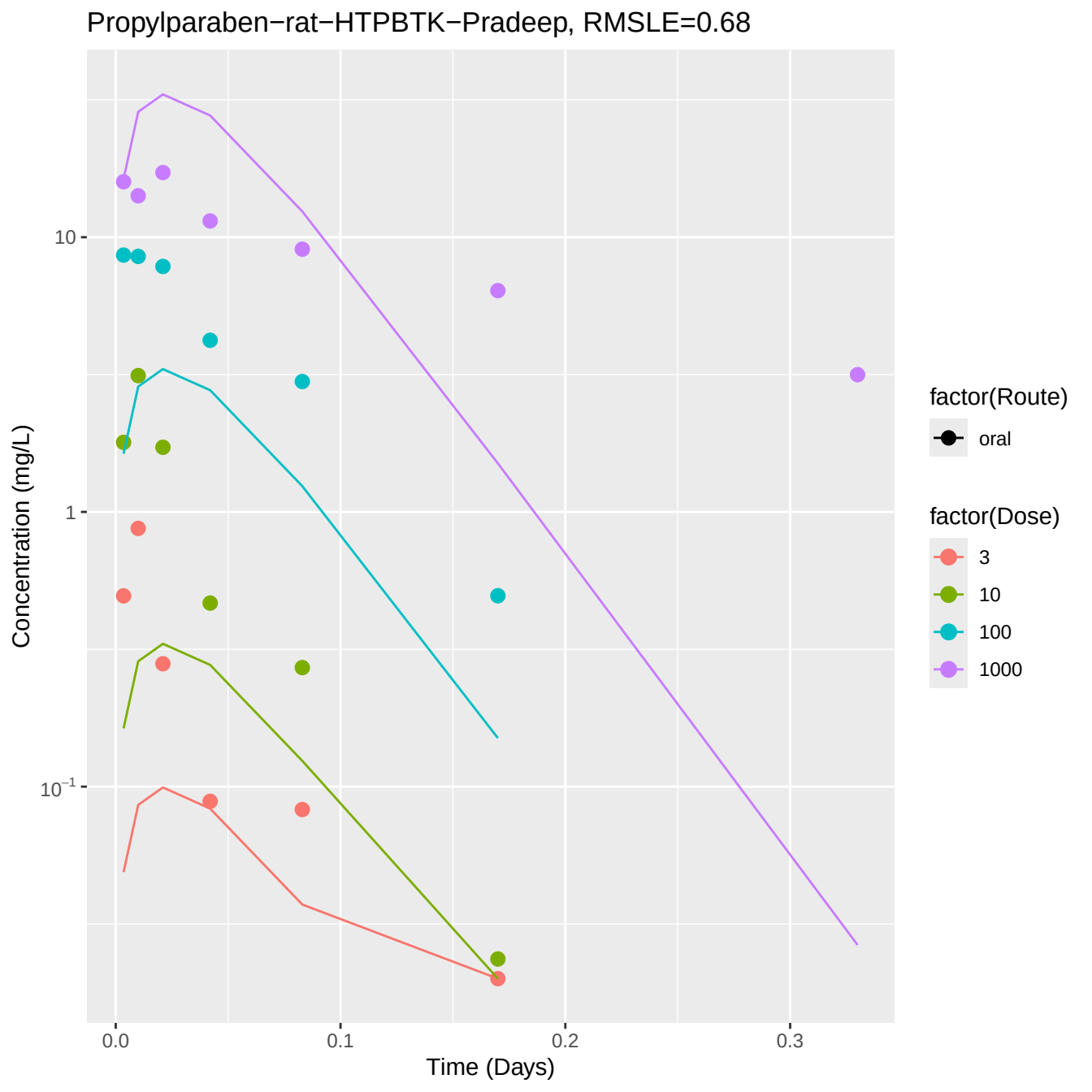

Methyleugenol−rat−HTPBTK−Ensemble, RMSLE=0.763
Concentration (mg/L)
10
1
10^{-1}
10^{-2}
0.00
0.25
0.50
0.75
1.00
Time (Days)
factor(Dose)
37
75
150
factor(Route)
iv
oral

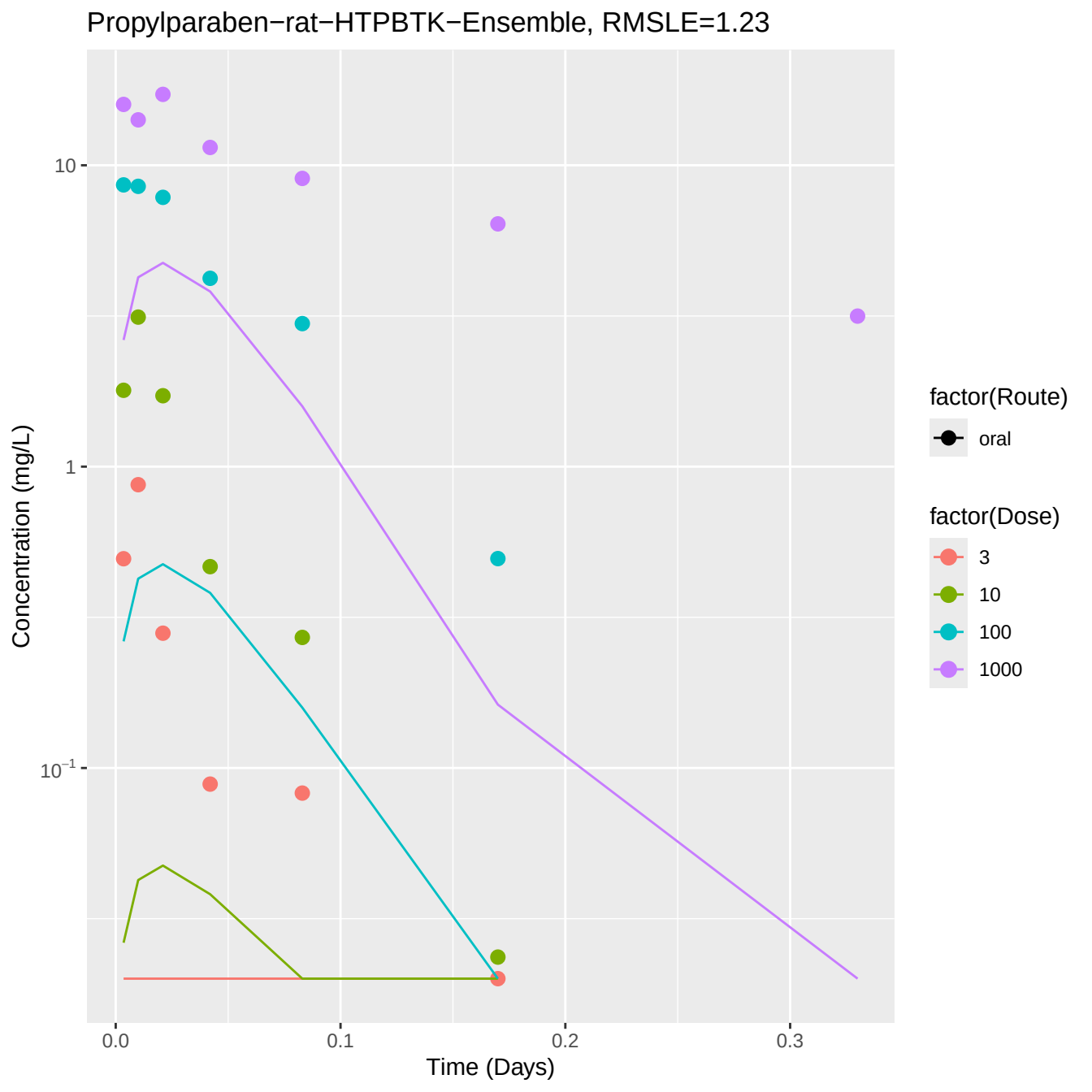

Methyleugenol−rat−In Vivo Fits, RMSLE=0.414
Concentration (mg/L)
10
1
10^{-1}
10^{-2}
0.00
0.25
0.50
0.75
1.00
Time (Days)
factor(Dose)
37
75
150
factor(Route)
iv
oral

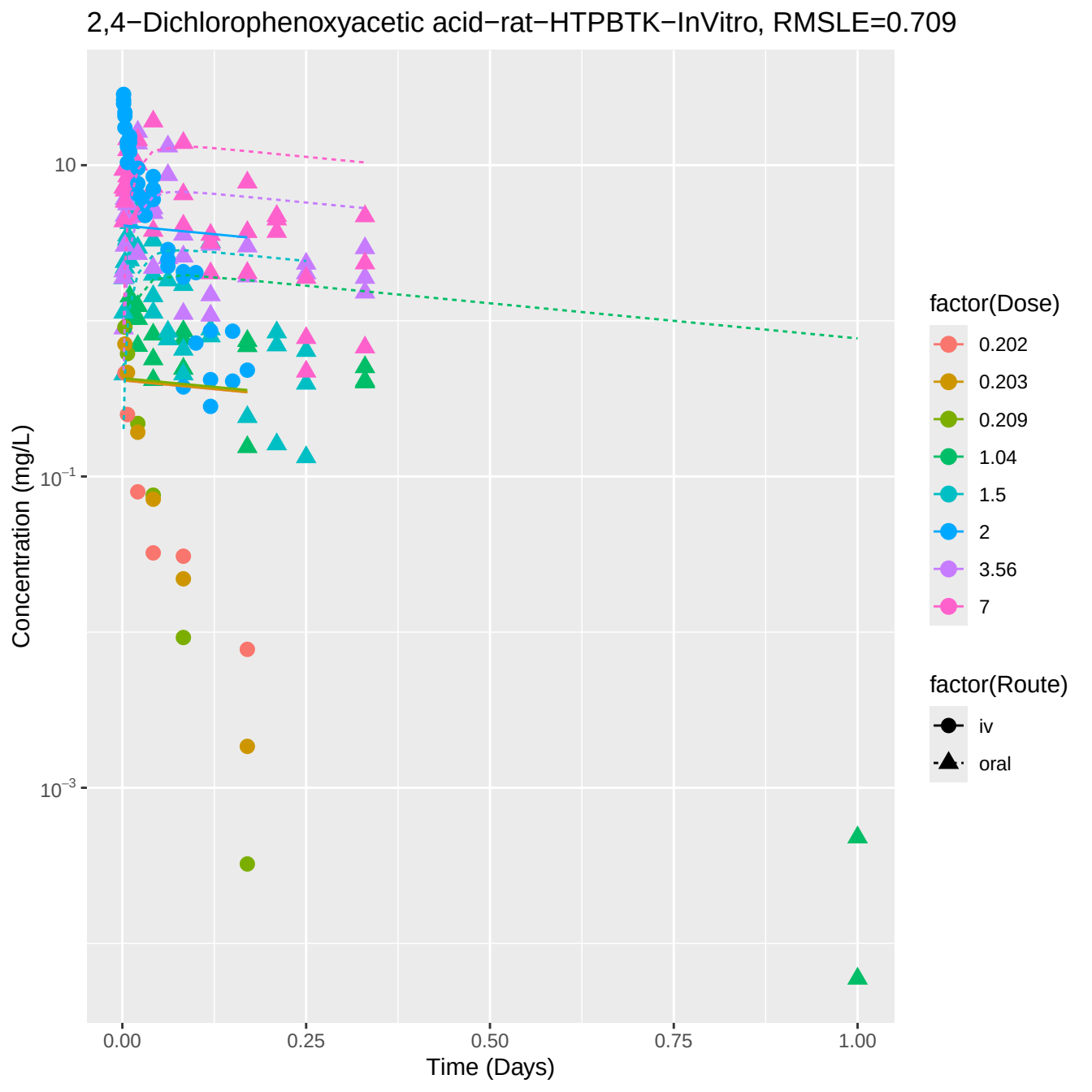

Propylparaben−rat−HTPBTK−InVitro, RMSLE=0.702
Concentration (mg/L)
10
1
10^{-1}
0.0
0.1
0.2
0.3
Time (Days)
factor(Route)
oral
factor(Dose)
3
10
100
1000

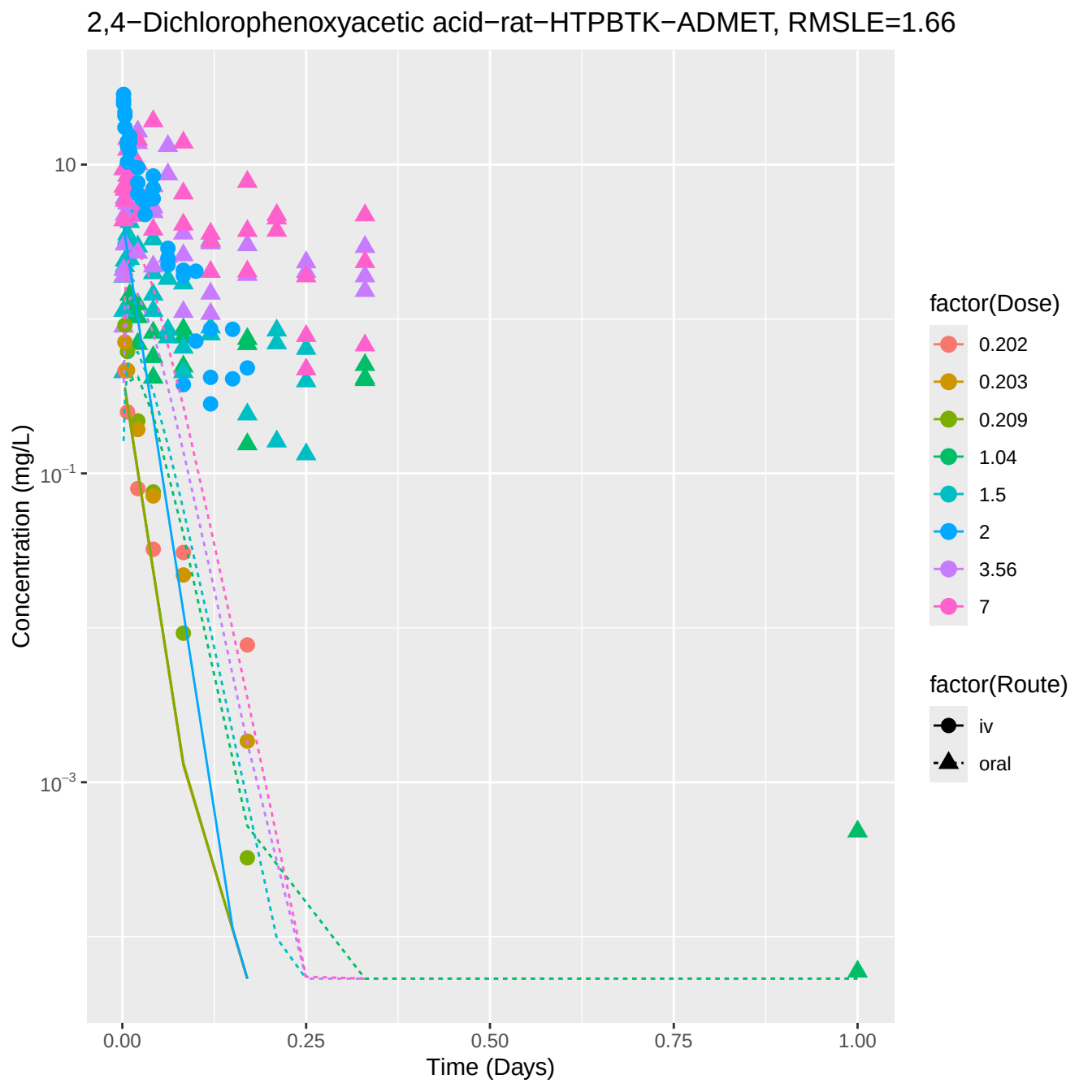

Propylparaben−rat−HTPBTK−ADMET, RMSLE=0.986
Concentration (mg/L)
10
1
10^{-1}
0.0
0.1
0.2
0.3
Time (Days)
factor(Route)
oral
factor(Dose)
3
10
100
1000

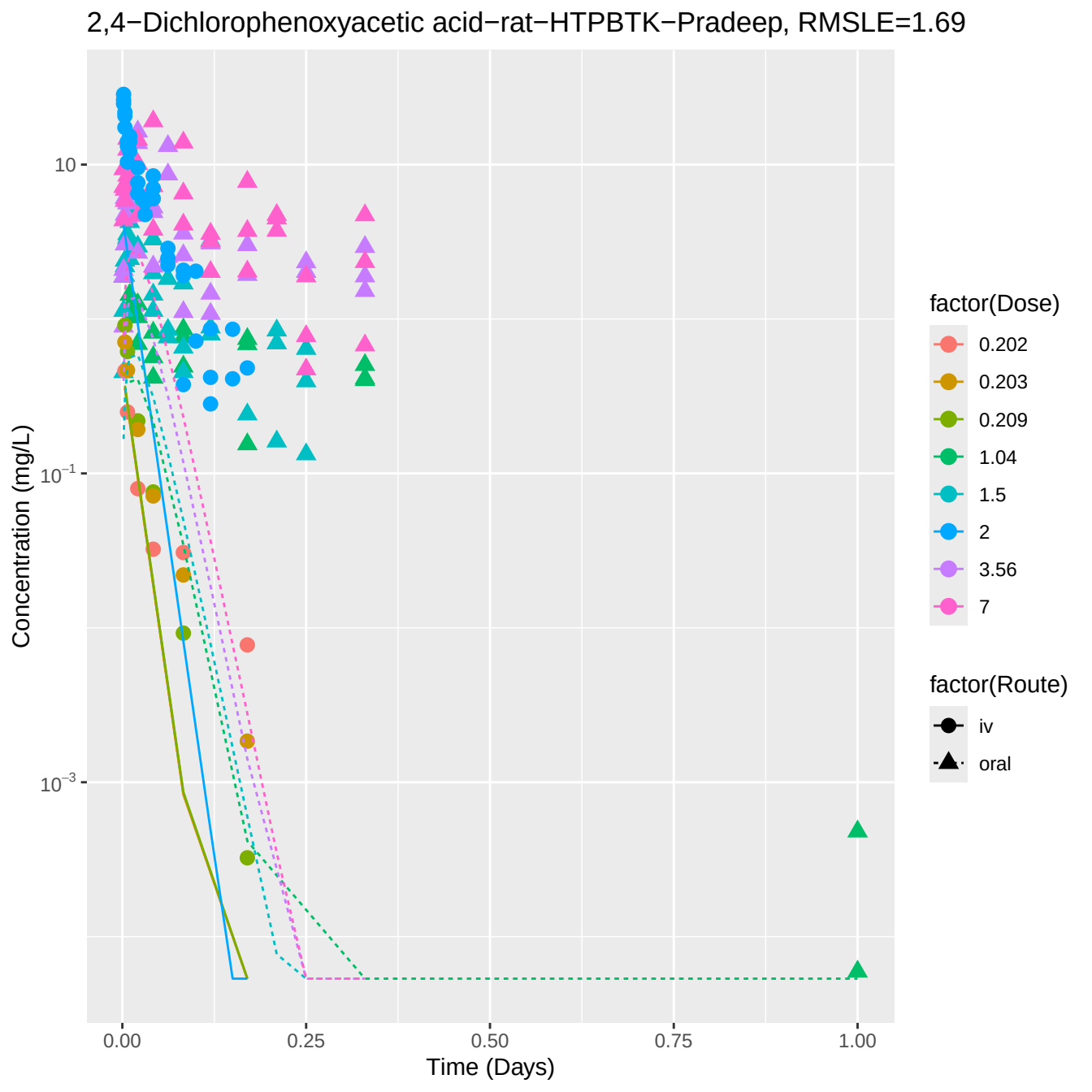

Propylparaben−rat−HTPBTK−Dawson, RMSLE=0.535
Concentration (mg/L)
10
1
10^{-1}
0.0
0.1
0.2
0.3
Time (Days)
factor(Route)
oral
factor(Dose)
3
10
100
1000

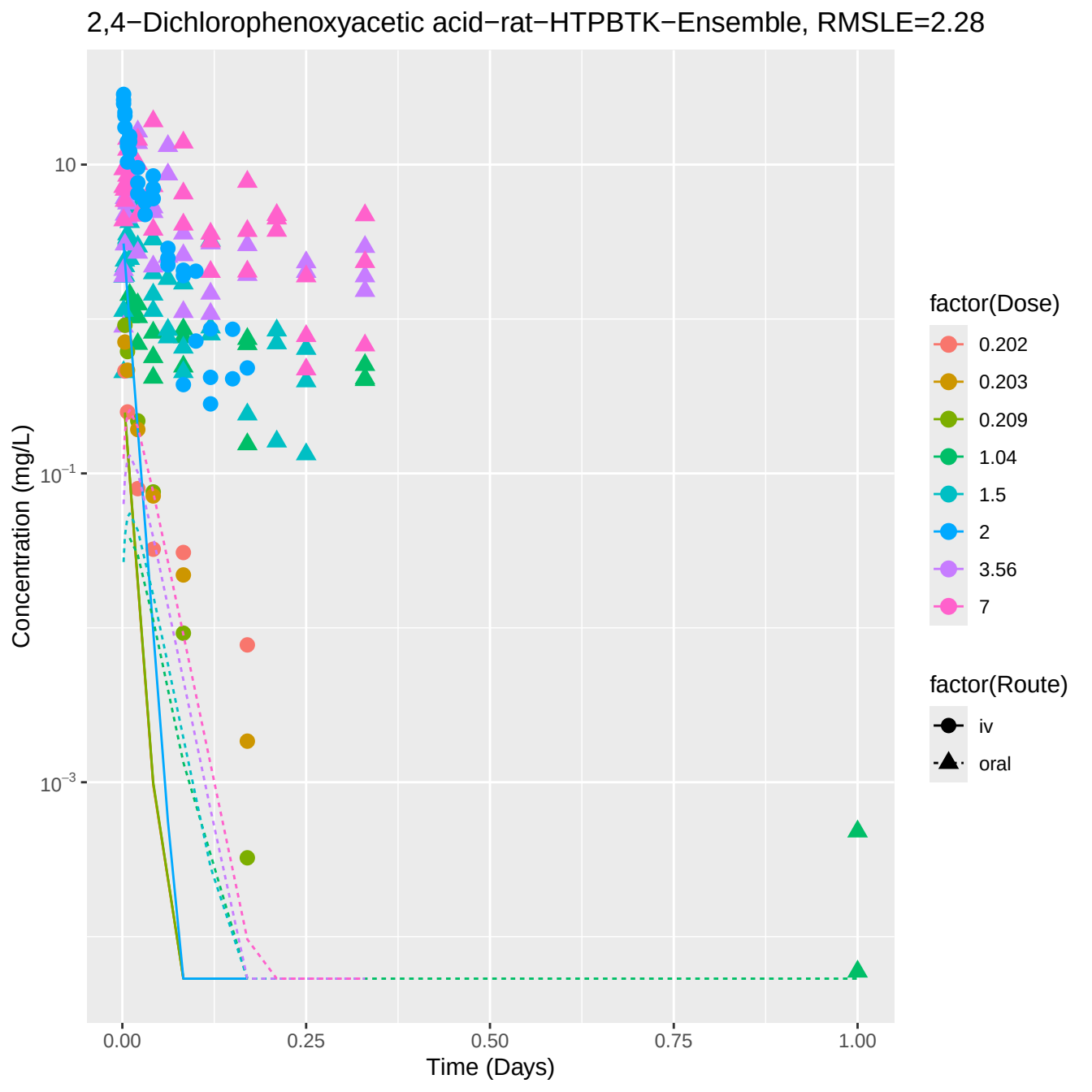

Propylparaben−rat−HTPBTK−Pradeep, RMSLE=0.68
Concentration (mg/L)
10
1
10^{-1}
0.0
0.1
0.2
0.3
Time (Days)
factor(Route)
oral
factor(Dose)
3
10
100
1000

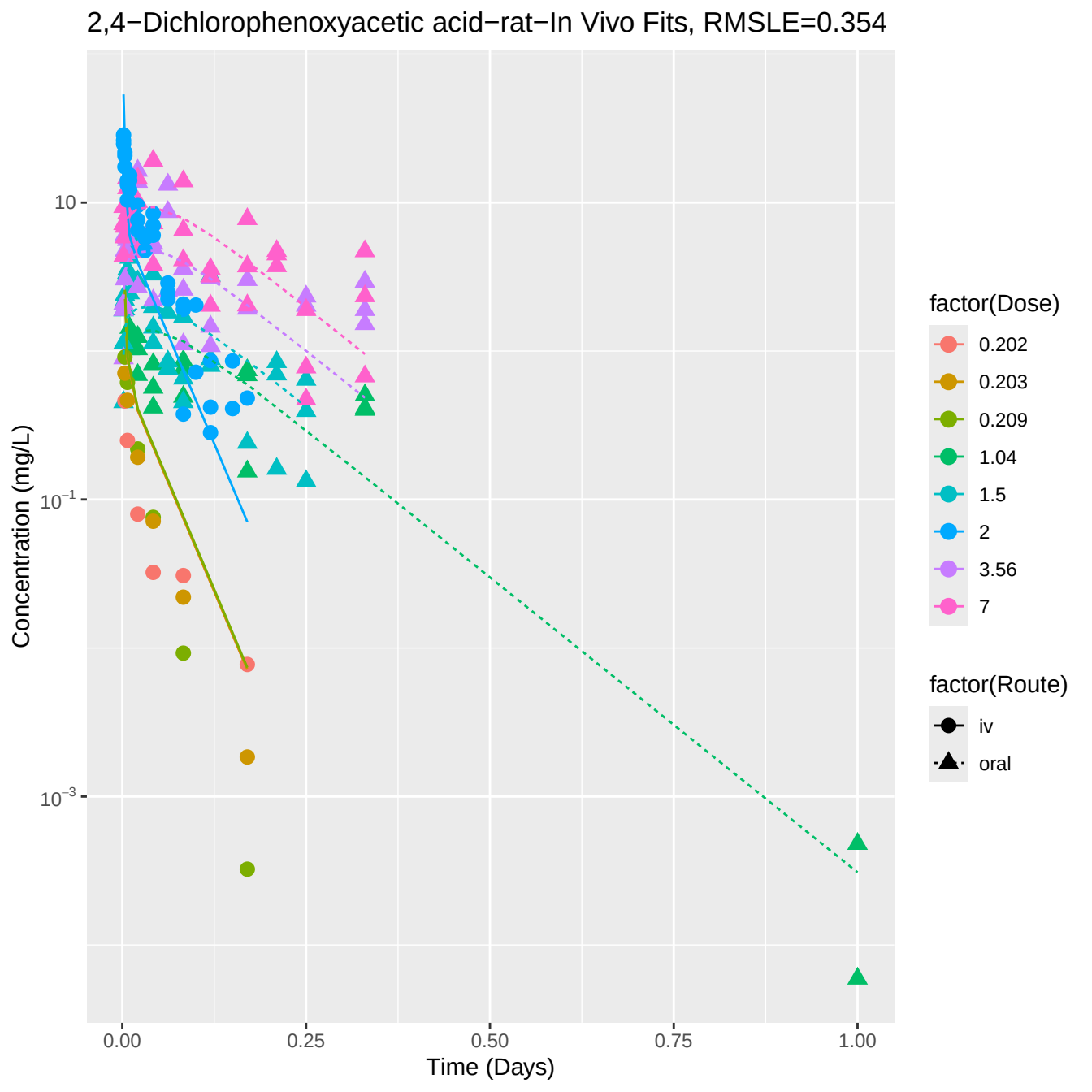

Propylparaben−rat−HTPBTK−Ensemble, RMSLE=1.23
Concentration (mg/L)
10
1
10^{-1}
0.0
0.1
0.2
0.3
Time (Days)
factor(Route)
oral
factor(Dose)
3
10
100
1000

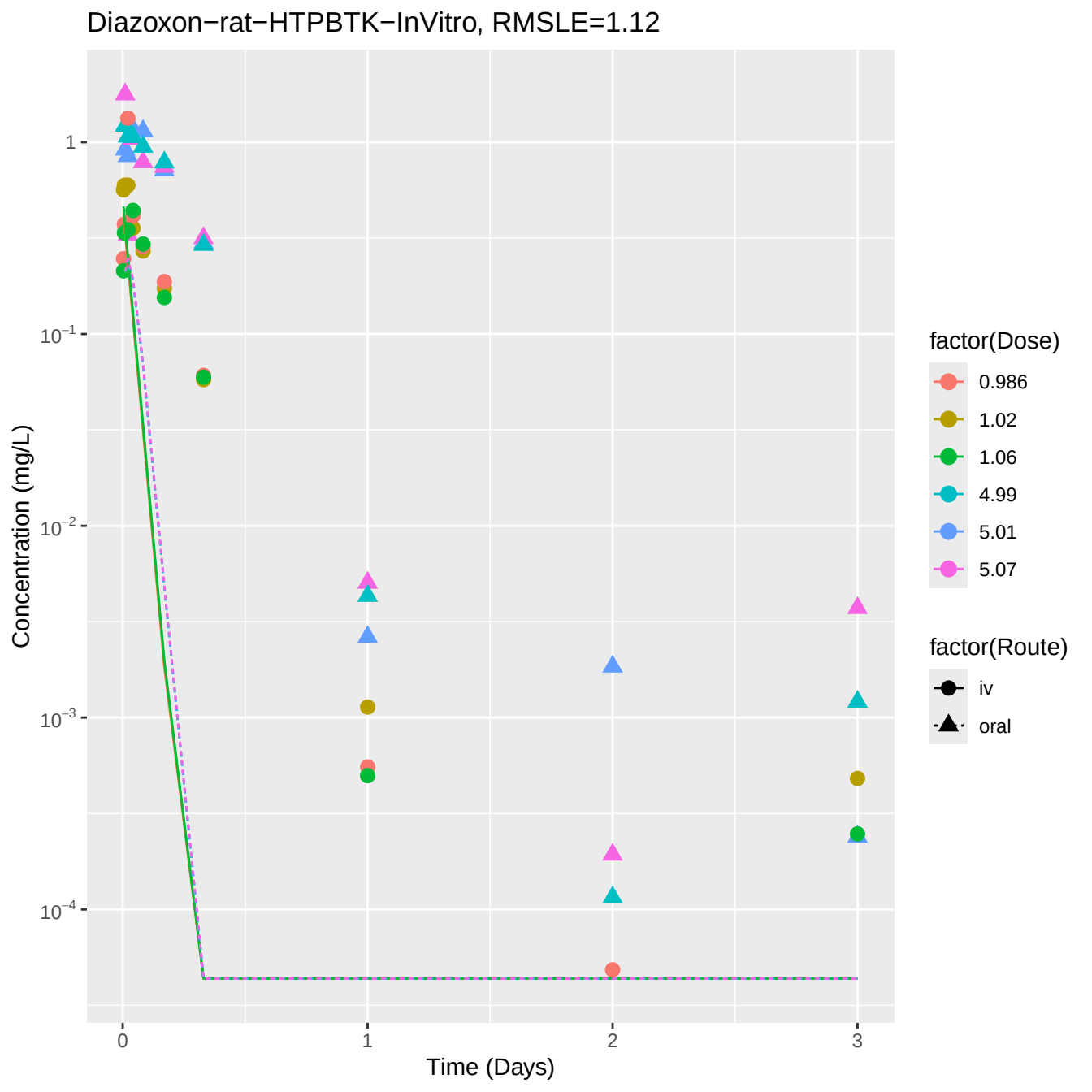

2,4−Dichlorophenoxyacetic acid−rat−HTPBTK−InVitro, RMSLE=0.709
Concentration (mg/L)
10
10^{-1}
10^{-3}
0.00
0.25
0.50
0.75
1.00
Time (Days)
factor(Dose)
0.202
0.203
0.209
1.04
1.5
2
3.56
7
factor(Route)
iv
oral

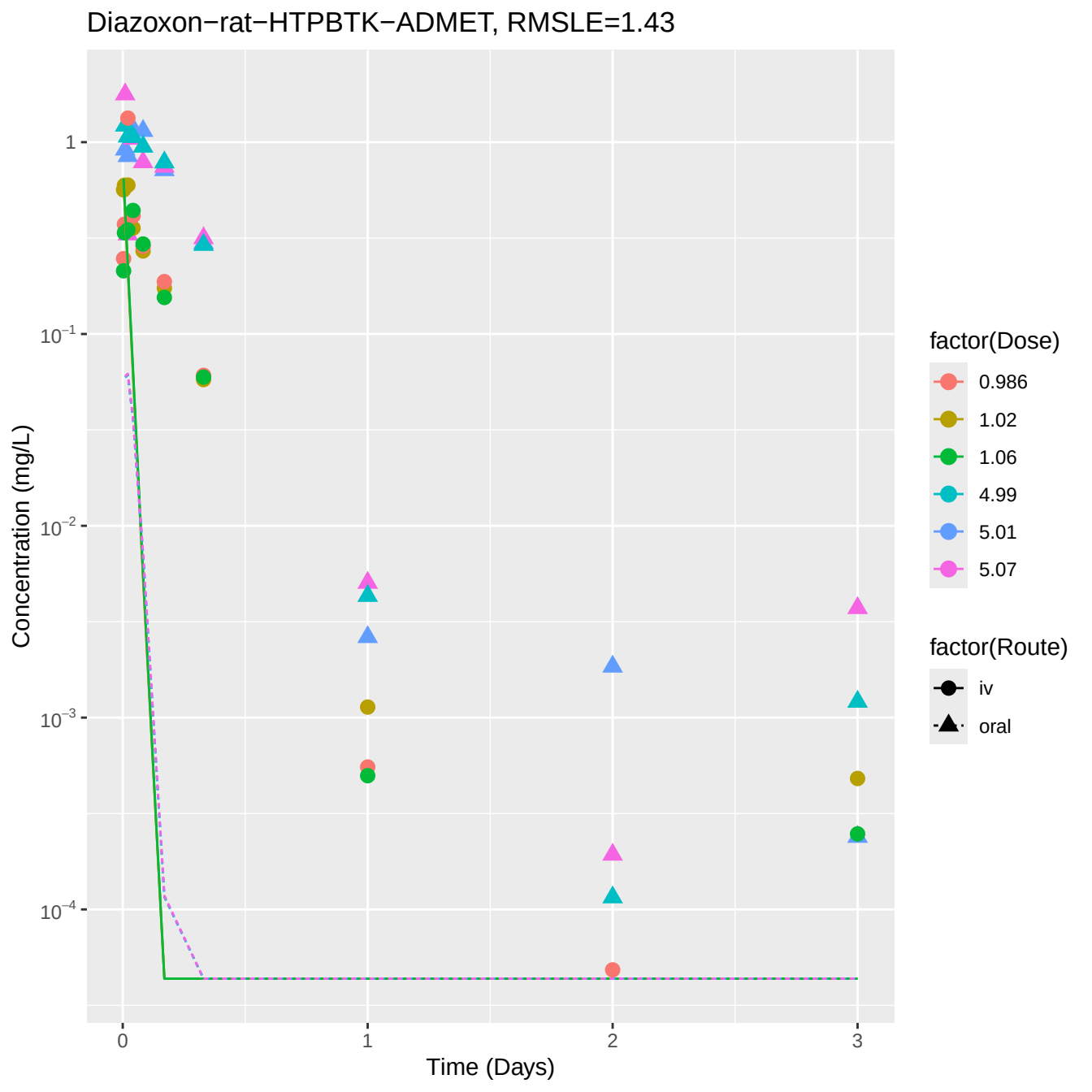

2,4−Dichlorophenoxyacetic acid−rat−HTPBTK−ADMET, RMSLE=1.66
Concentration (mg/L)
10
10^{-1}
10^{-3}
0.00
0.25
0.50
0.75
1.00
Time (Days)
factor(Dose)
0.202
0.203
0.209
1.04
1.5
2
3.56
7
factor(Route)
iv
oral

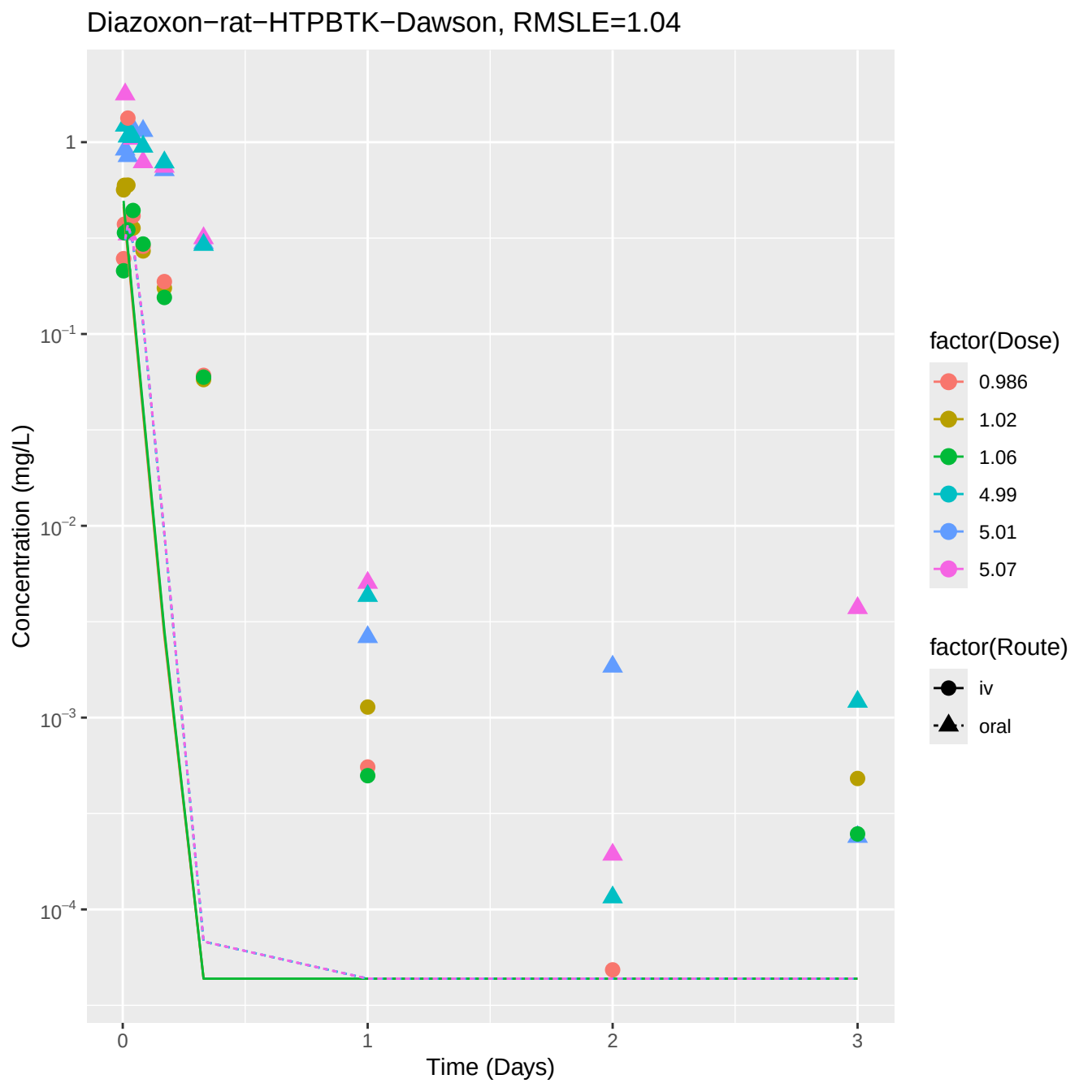

2,4−Dichlorophenoxyacetic acid−rat−HTPBTK−Pradeep, RMSLE=1.69
Concentration (mg/L)
10
10^{-1}
10^{-3}
0.00
0.25
0.50
0.75
1.00
Time (Days)
factor(Dose)
0.202
0.203
0.209
1.04
1.5
2
3.56
7
factor(Route)
iv
oral

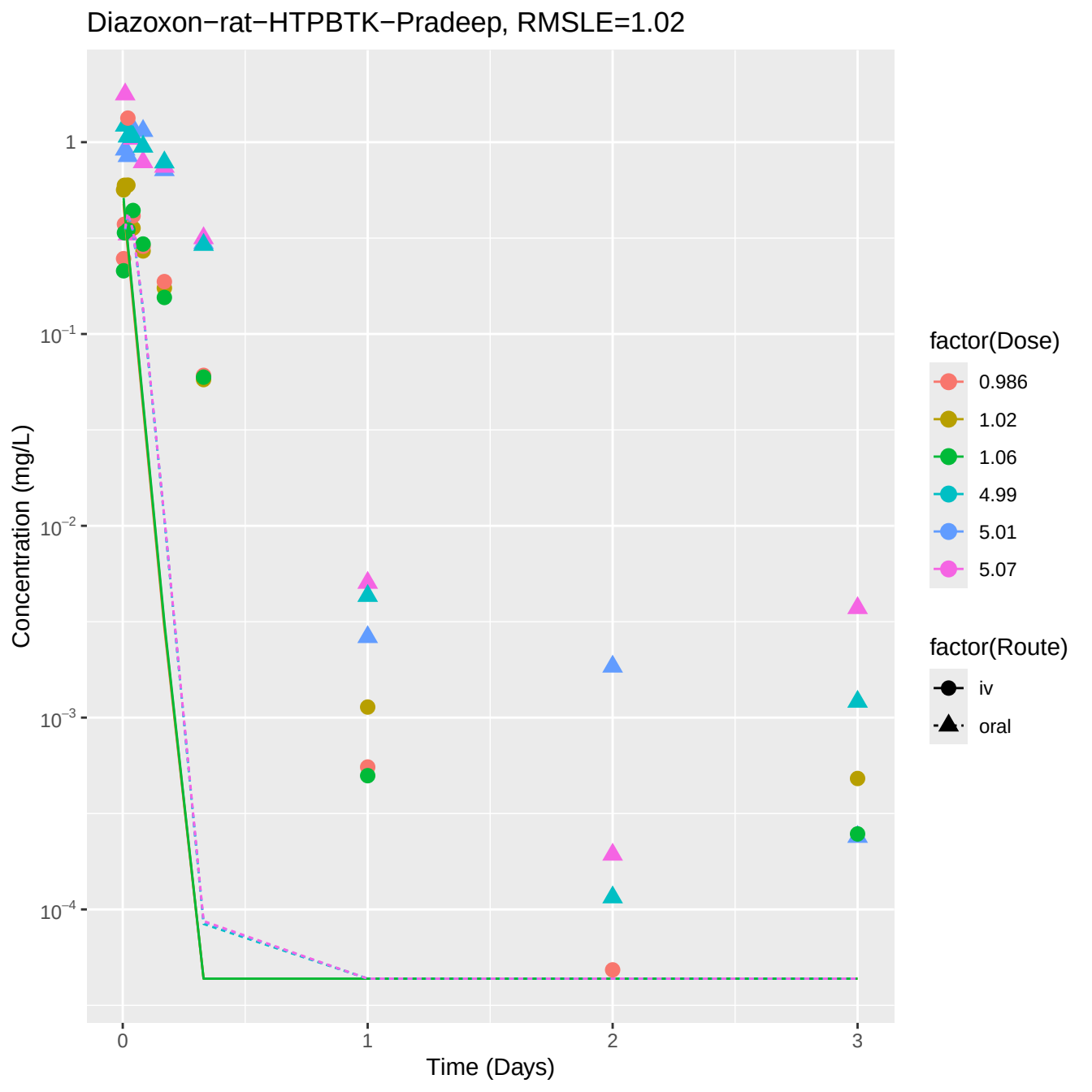

2,4−Dichlorophenoxyacetic acid−rat−HTPBTK−Ensemble, RMSLE=2.28
Concentration (mg/L)
10
10^{-1}
10^{-3}
0.00
0.25
0.50
0.75
1.00
Time (Days)
factor(Dose)
0.202
0.203
0.209
1.04
1.5
2
3.56
7
factor(Route)
iv
oral

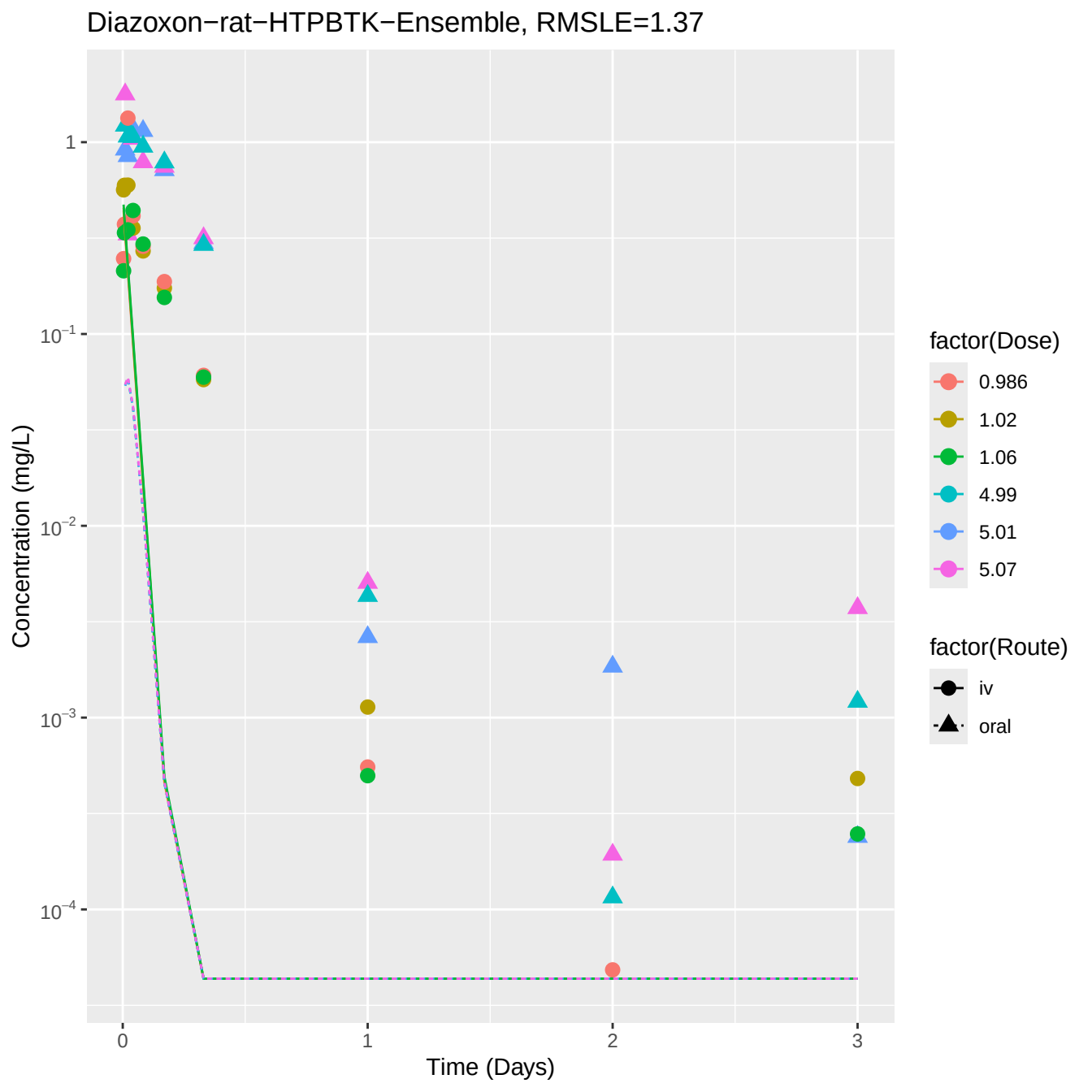

2,4−Dichlorophenoxyacetic acid−rat−In Vivo Fits, RMSLE=0.354
Concentration (mg/L)
10
10^{-1}
10^{-3}
0.00
0.25
0.50
0.75
1.00
Time (Days)
factor(Dose)
0.202
0.203
0.209
1.04
1.5
2
3.56
7
factor(Route)
iv
oral

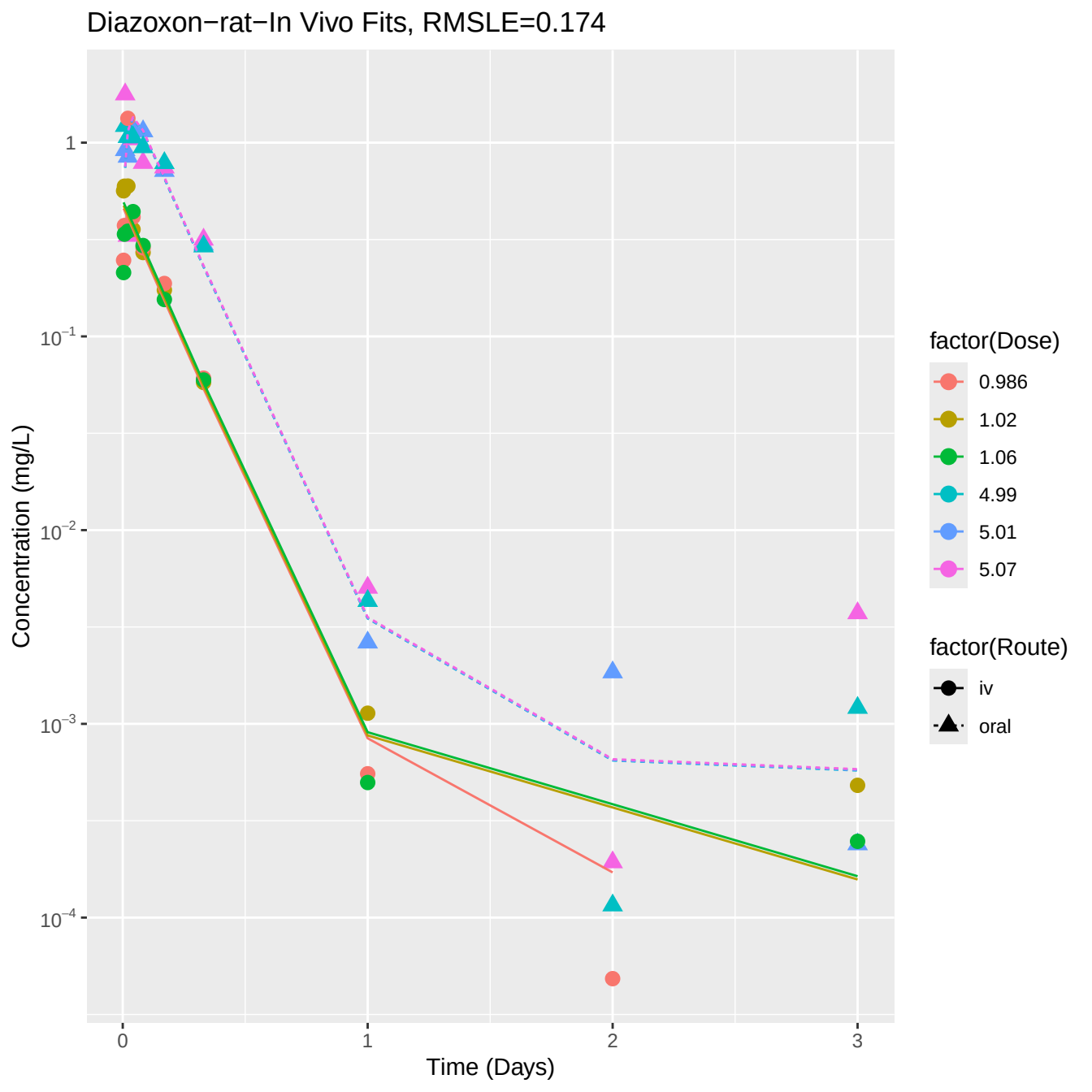

Diazoxon−rat−HTPBTK−InVitro, RMSLE=1.12
Concentration (mg/L)
1
10^{-1}
10^{-2}
10^{-3}
10^{-4}
0
1
2
3
Time (Days)
factor(Dose)
0.986
1.02
1.06
4.99
5.01
5.07
factor(Route)
iv
oral

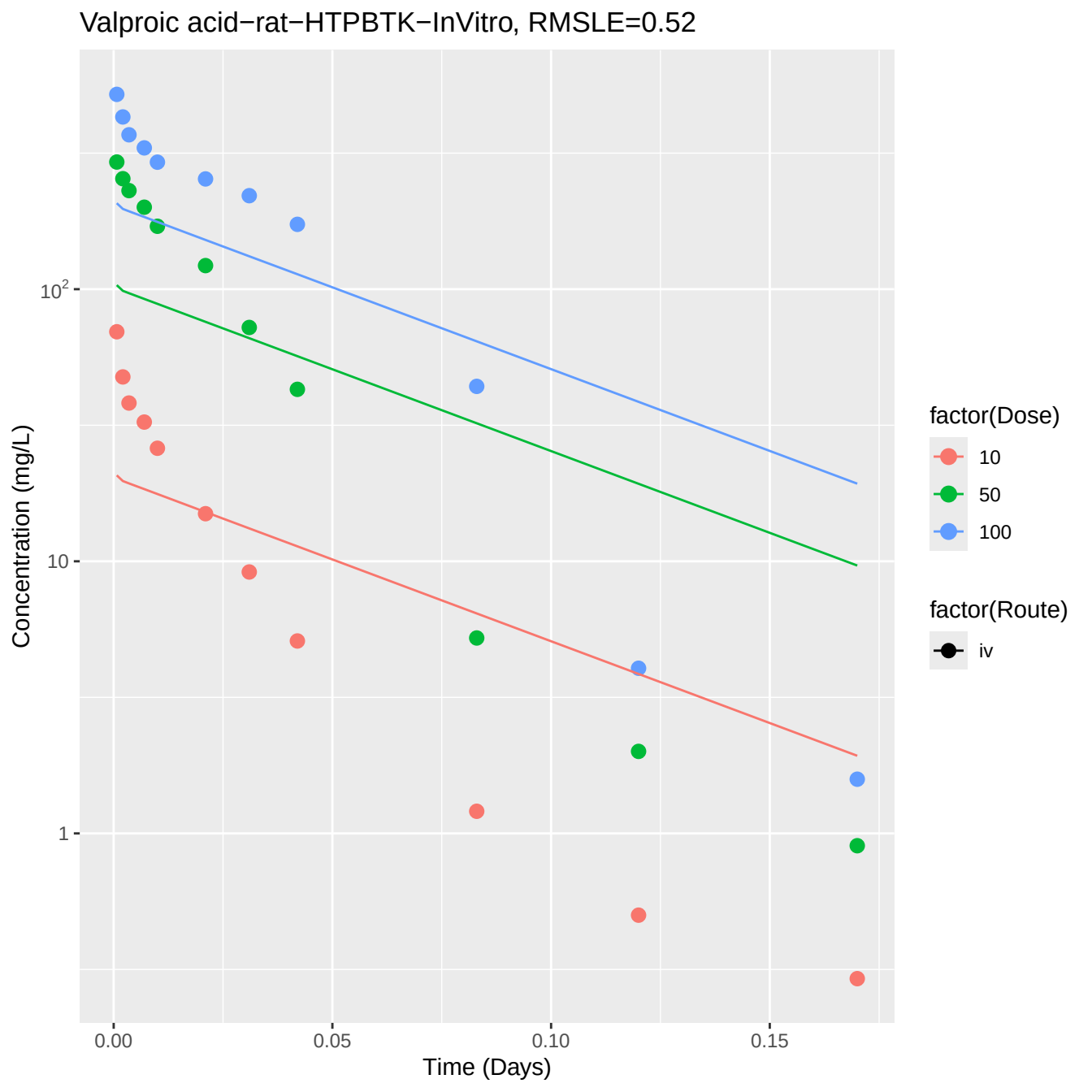

Diazoxon−rat−HTPBTK−ADMET, RMSLE=1.43
Concentration (mg/L)
1
10^{-1}
10^{-2}
10^{-3}
10^{-4}
0
1
2
3
Time (Days)
factor(Dose)
0.986
1.02
1.06
4.99
5.01
5.07
factor(Route)
iv
oral

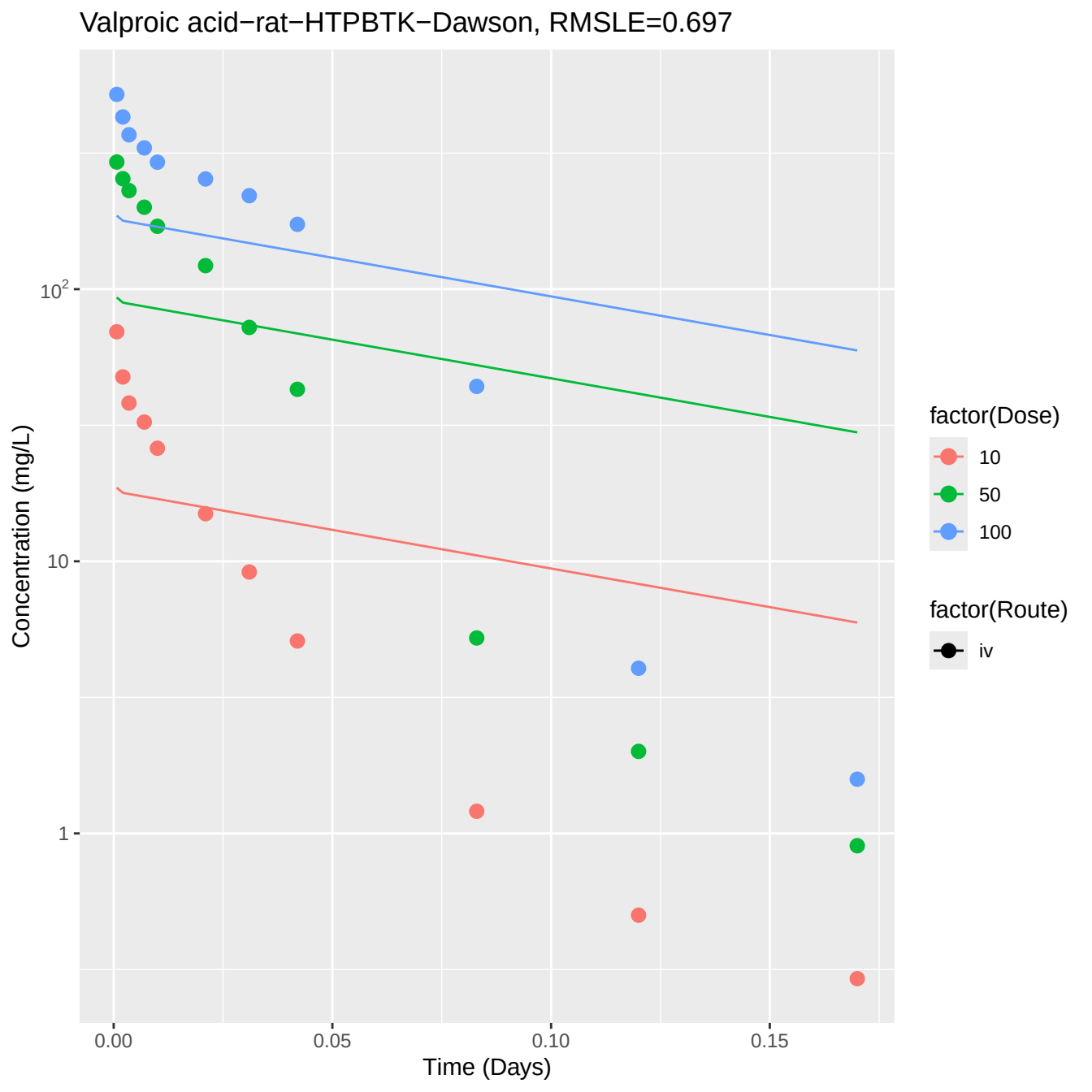

Diazoxon−rat−HTPBTK−Dawson, RMSLE=1.04
Concentration (mg/L)
1
10^{-1}
10^{-2}
10^{-3}
10^{-4}
0
1
2
3
Time (Days)
factor(Dose)
0.986
1.02
1.06
4.99
5.01
5.07
factor(Route)
iv
oral

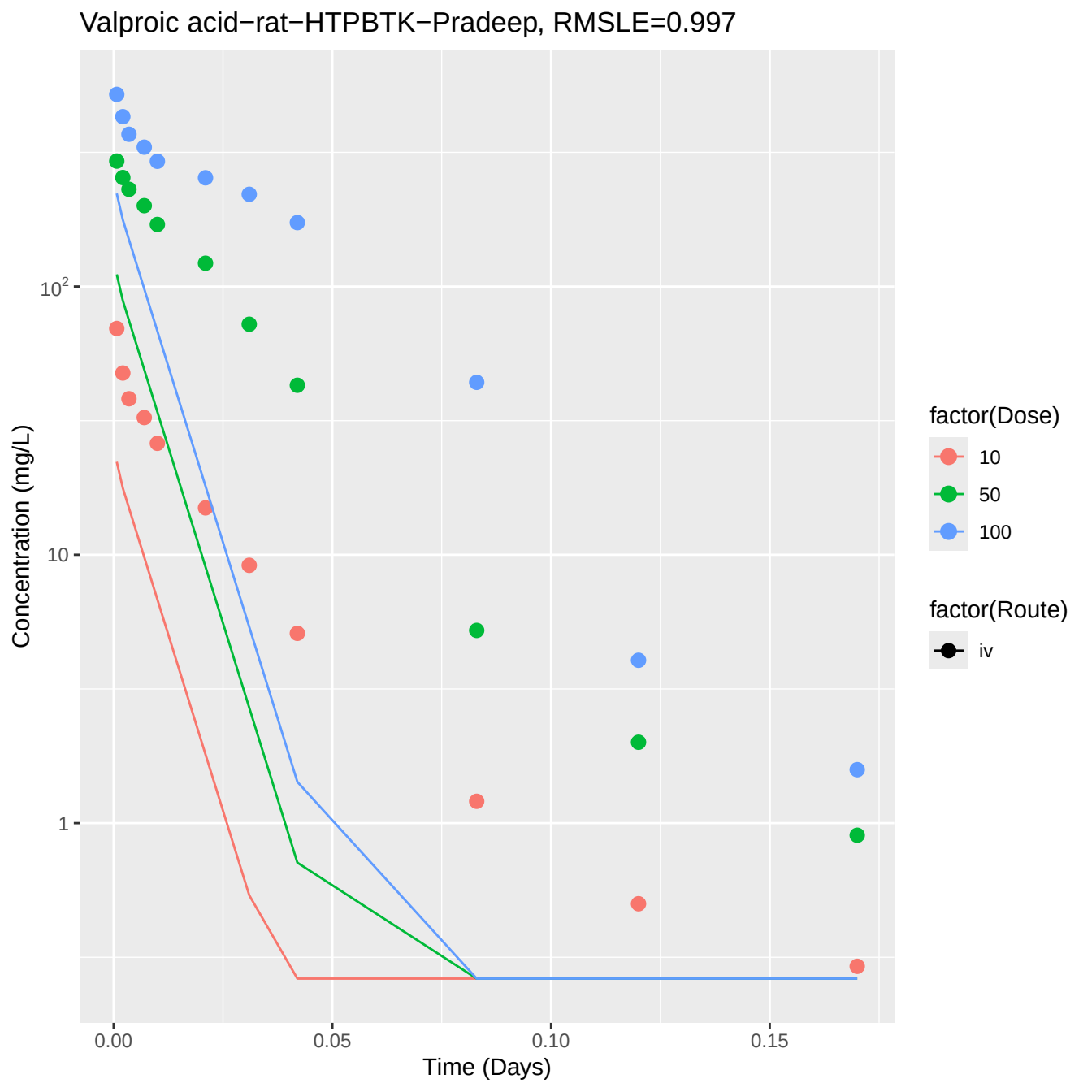

Diazoxon−rat−HTPBTK−Pradeep, RMSLE=1.02
Concentration (mg/L)
1
10^{-1}
10^{-2}
10^{-3}
10^{-4}
0
1
2
3
Time (Days)
factor(Dose)
0.986
1.02
1.06
4.99
5.01
5.07
factor(Route)
iv
oral

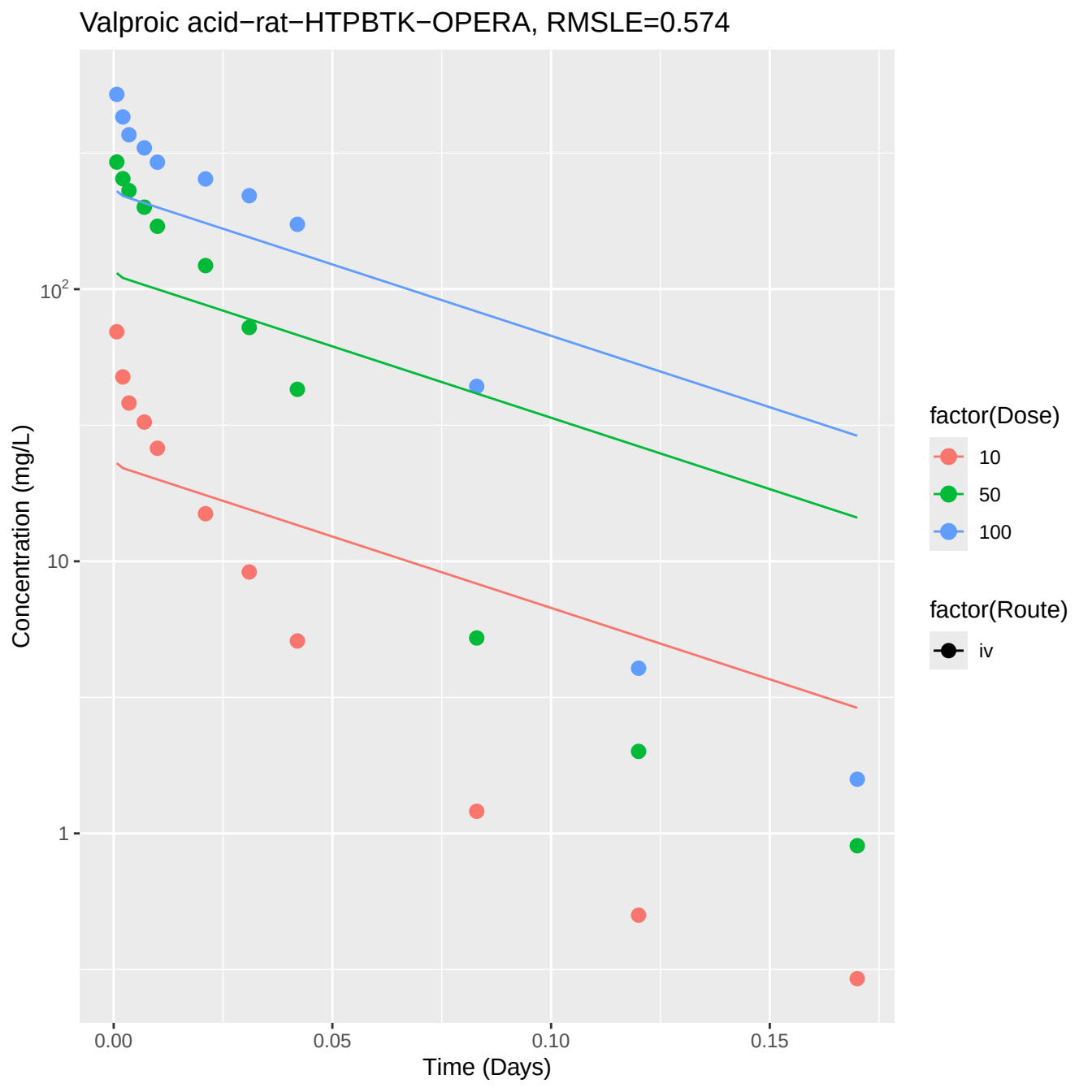

Diazoxon−rat−HTPBTK−Ensemble, RMSLE=1.37
Concentration (mg/L)
1
10^{-1}
10^{-2}
10^{-3}
10^{-4}
0
1
2
3
Time (Days)
factor(Dose)
0.986
1.02
1.06
4.99
5.01
5.07
factor(Route)
iv
oral

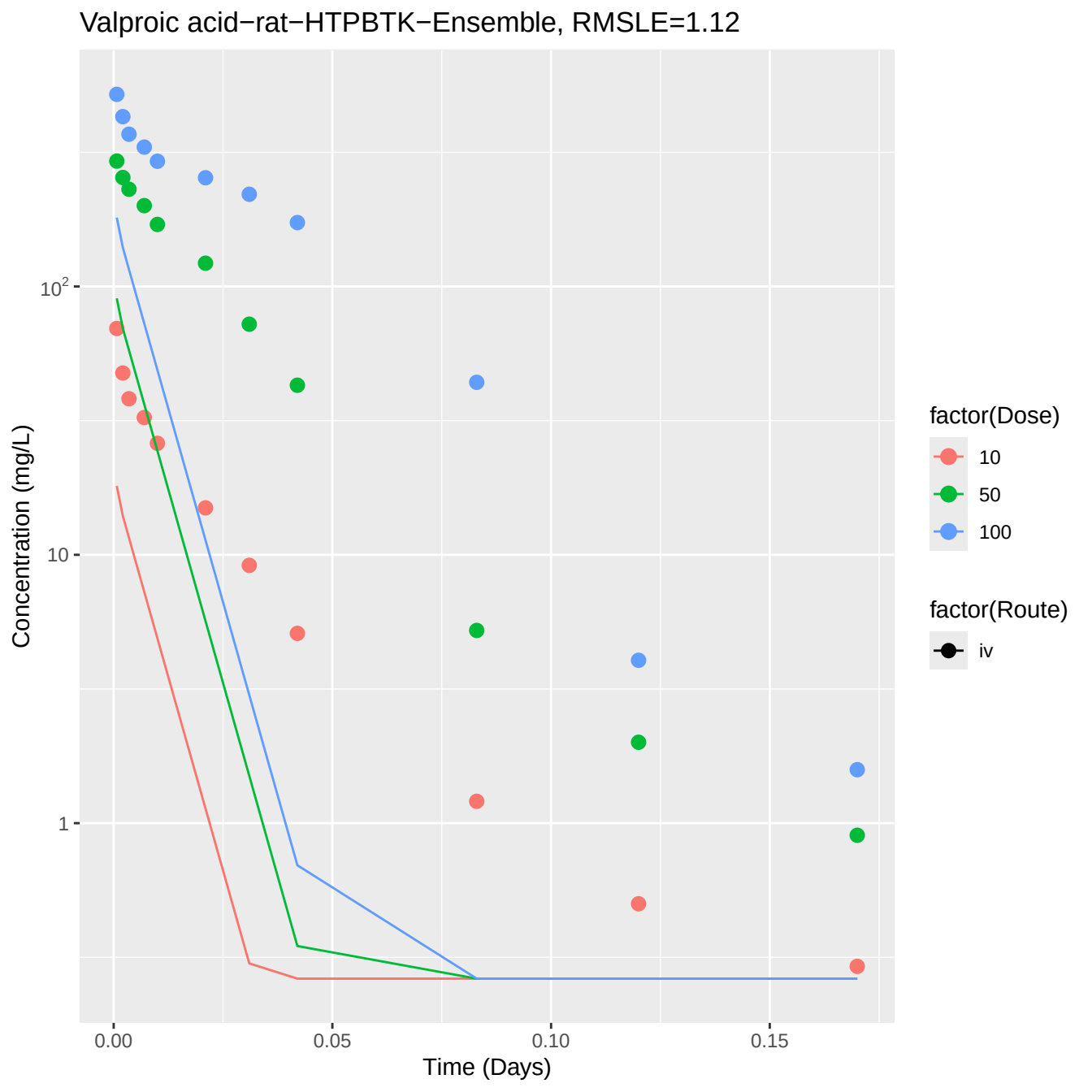

Diazoxon−rat−In Vivo Fits, RMSLE=0.174
Concentration (mg/L)
1
10^{-1}
10^{-2}
10^{-3}
10^{-4}
0
1
2
3
Time (Days)
factor(Dose)
0.986
1.02
1.06
4.99
5.01
5.07
factor(Route)
iv
oral

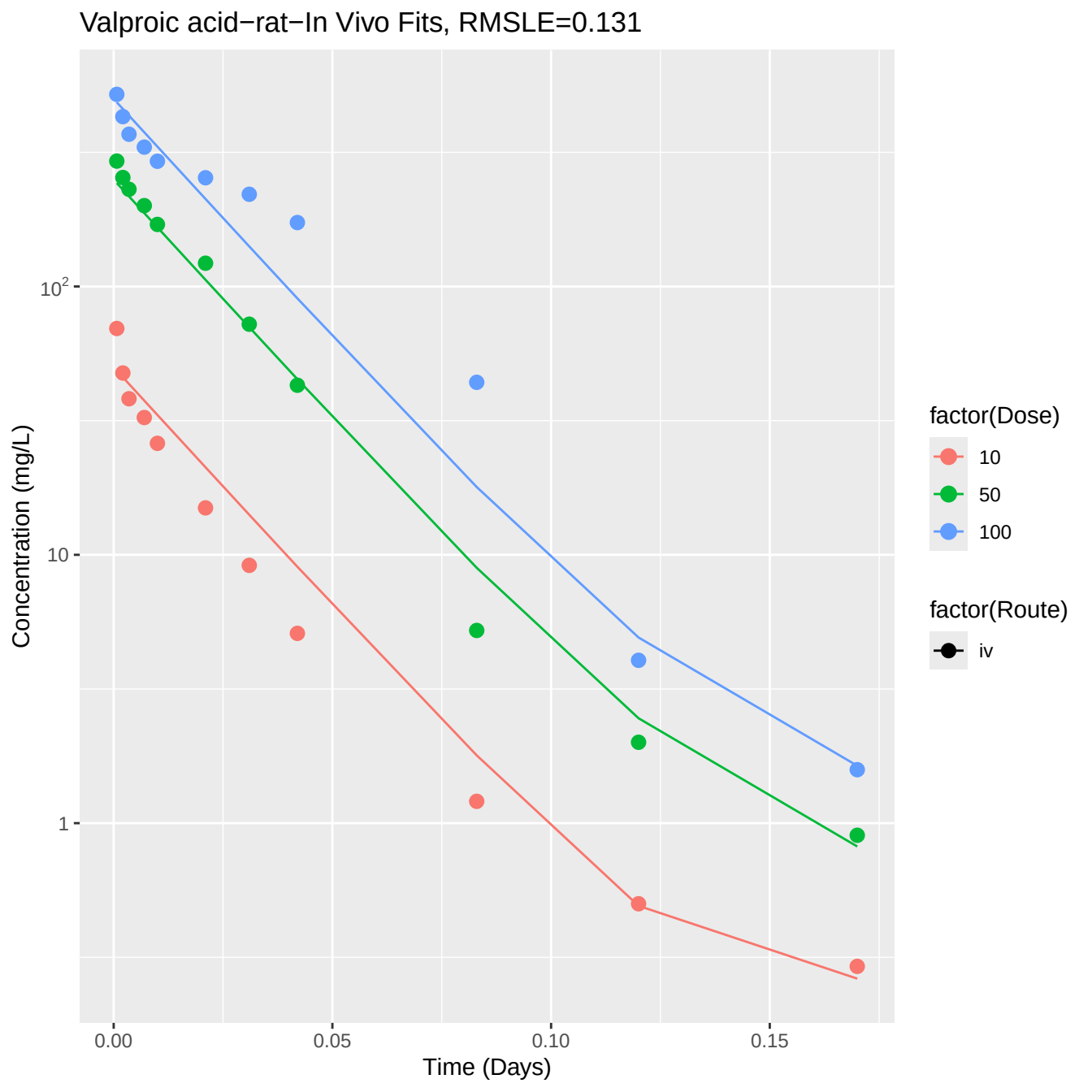

Valproic acid−rat−HTPBTK−InVitro, RMSLE=0.52
Concentration (mg/L)
10^2
10
1
0.00
0.05
0.10
0.15
Time (Days)
factor(Dose)
10
50
100
factor(Route)
iv

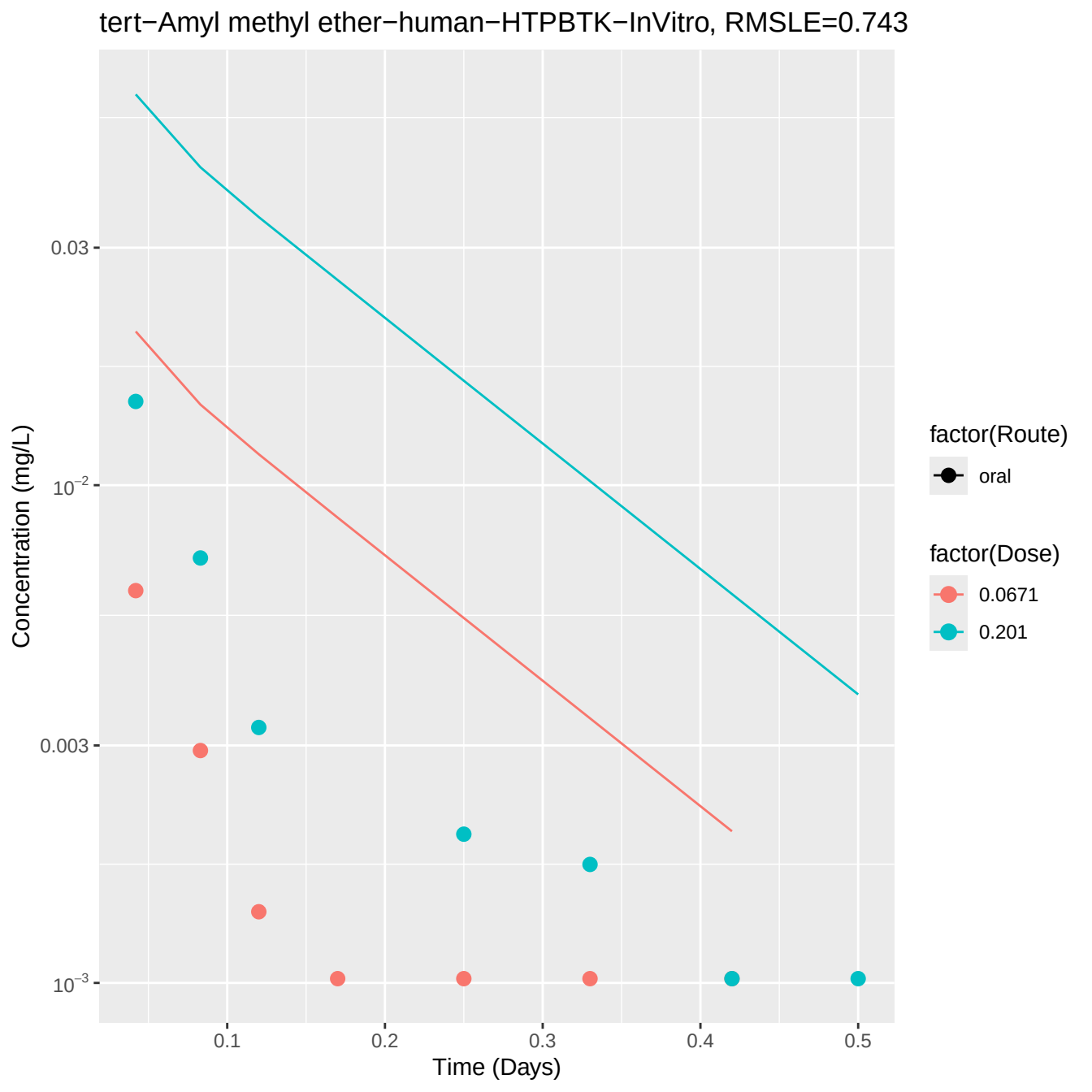

Valproic acid−rat−HTPBTK−Dawson, RMSLE=0.697
Concentration (mg/L)
10^2
10
1
0.00
0.05
0.10
0.15
Time (Days)
factor(Dose)
10
50
100
factor(Route)
iv

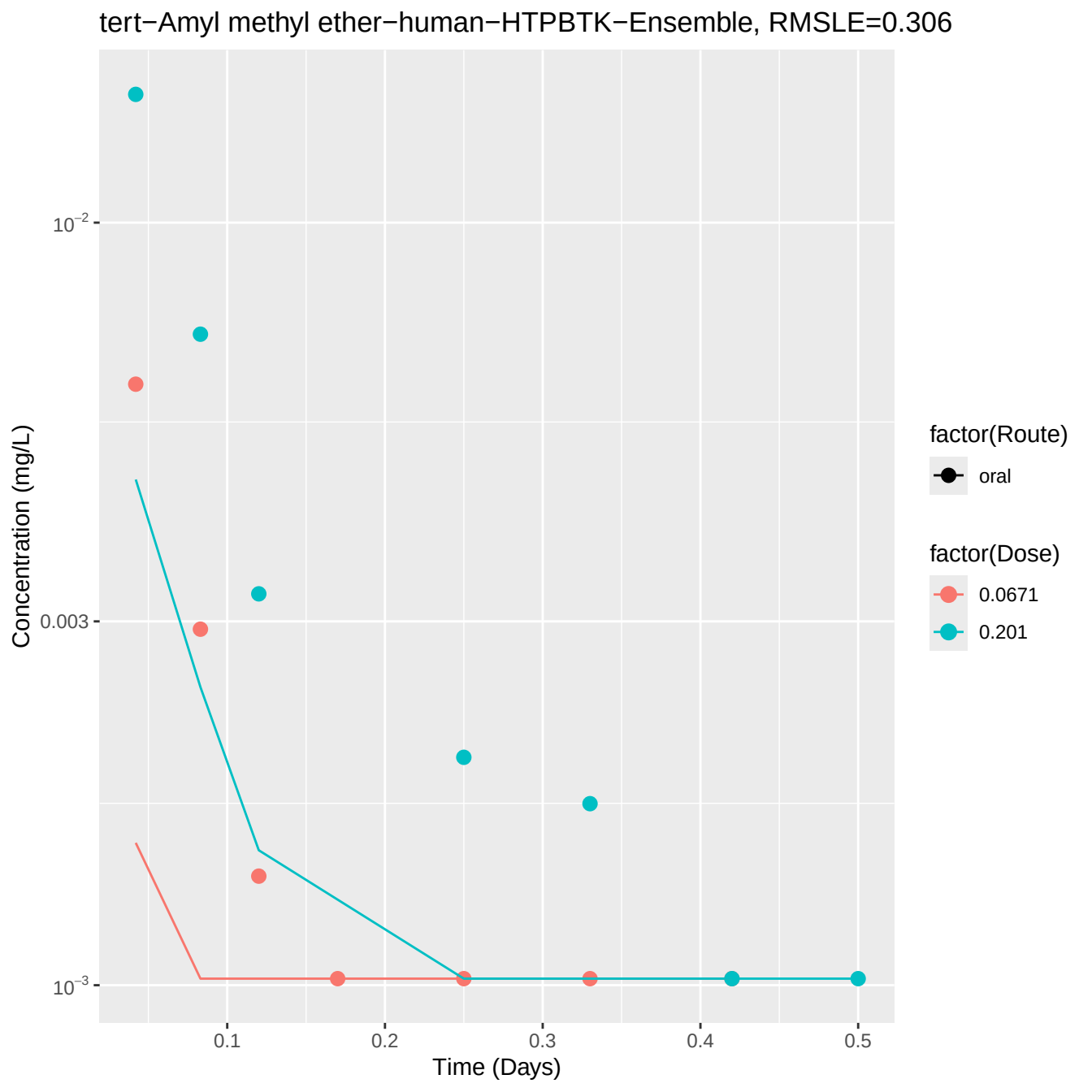

Valproic acid−rat−HTPBTK−Pradeep, RMSLE=0.997
Concentration (mg/L)
10^2
10
1
0.00
0.05
0.10
0.15
Time (Days)
factor(Dose)
10
50
100
factor(Route)
iv

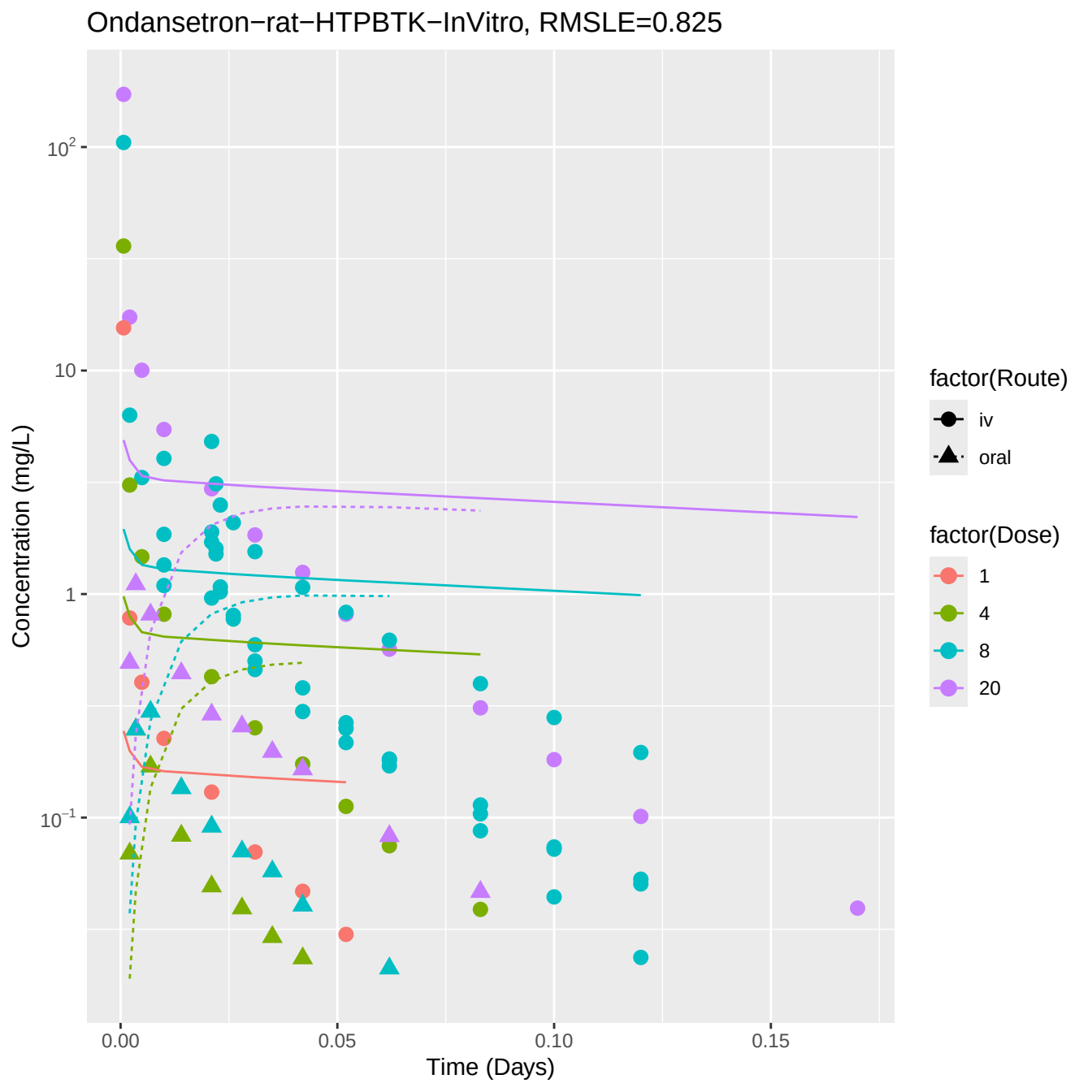

Valproic acid−rat−HTPBTK−OPERA, RMSLE=0.574
Concentration (mg/L)
10^2
10
1
0.00
0.05
0.10
0.15
Time (Days)
factor(Dose)
10
50
100
factor(Route)
iv

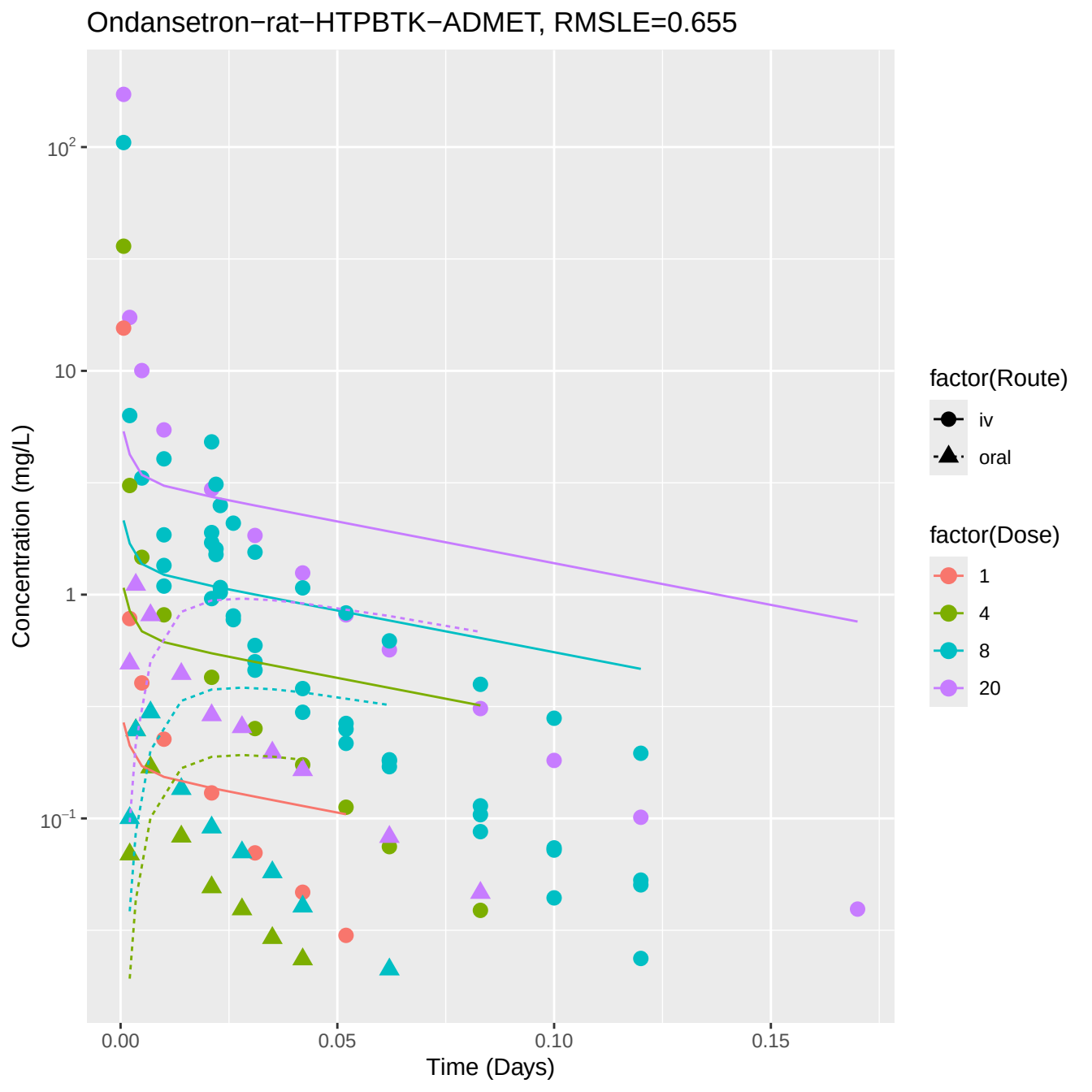

Valproic acid−rat−HTPBTK−Ensemble, RMSLE=1.12
Concentration (mg/L)
10^2
10
1
0.00
0.05
0.10
0.15
Time (Days)
factor(Dose)
10
50
100
factor(Route)
iv

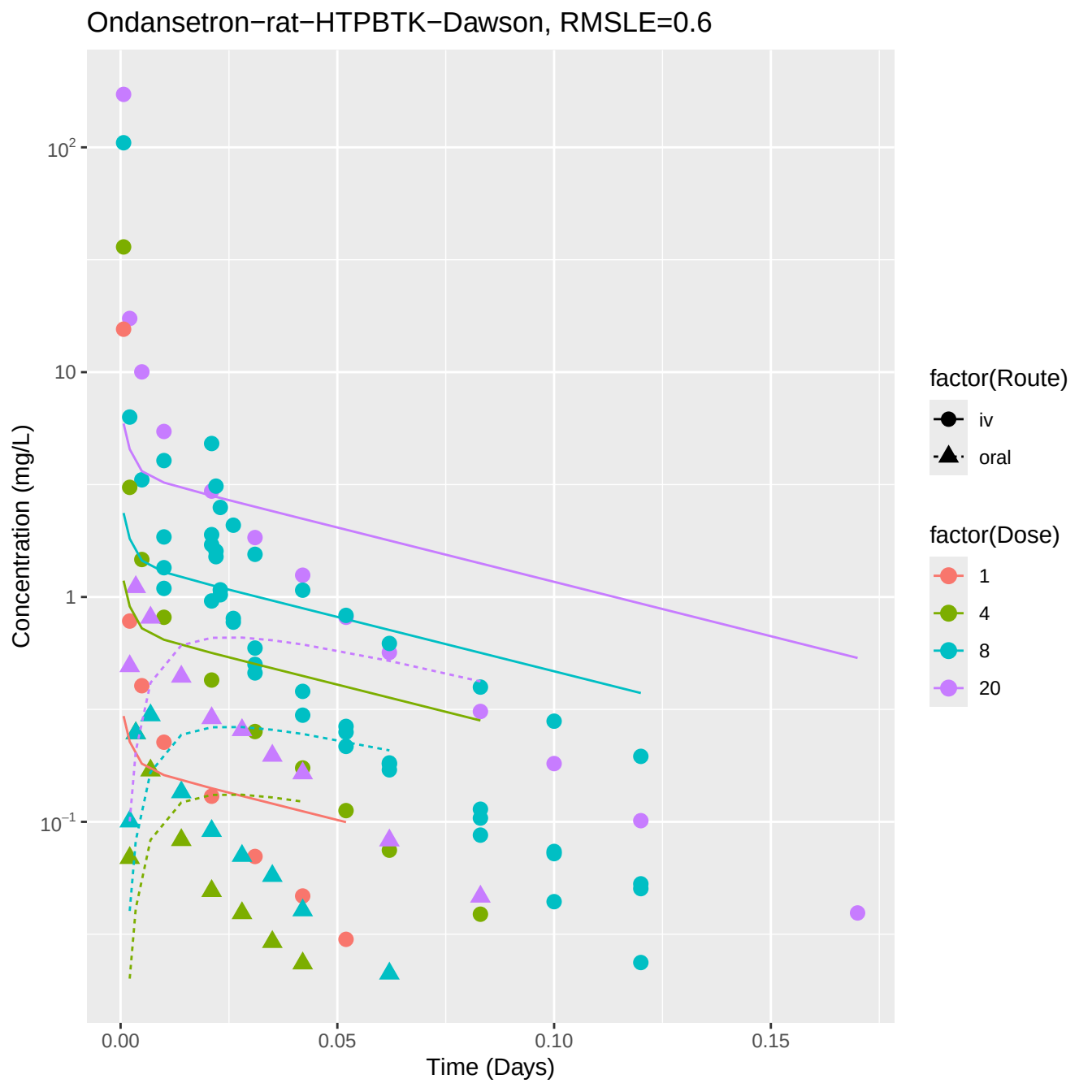

Valproic acid−rat−In Vivo Fits, RMSLE=0.131
Concentration (mg/L)
10^2
10
1
0.00
0.05
0.10
0.15
Time (Days)
factor(Dose)
10
50
100
factor(Route)
iv

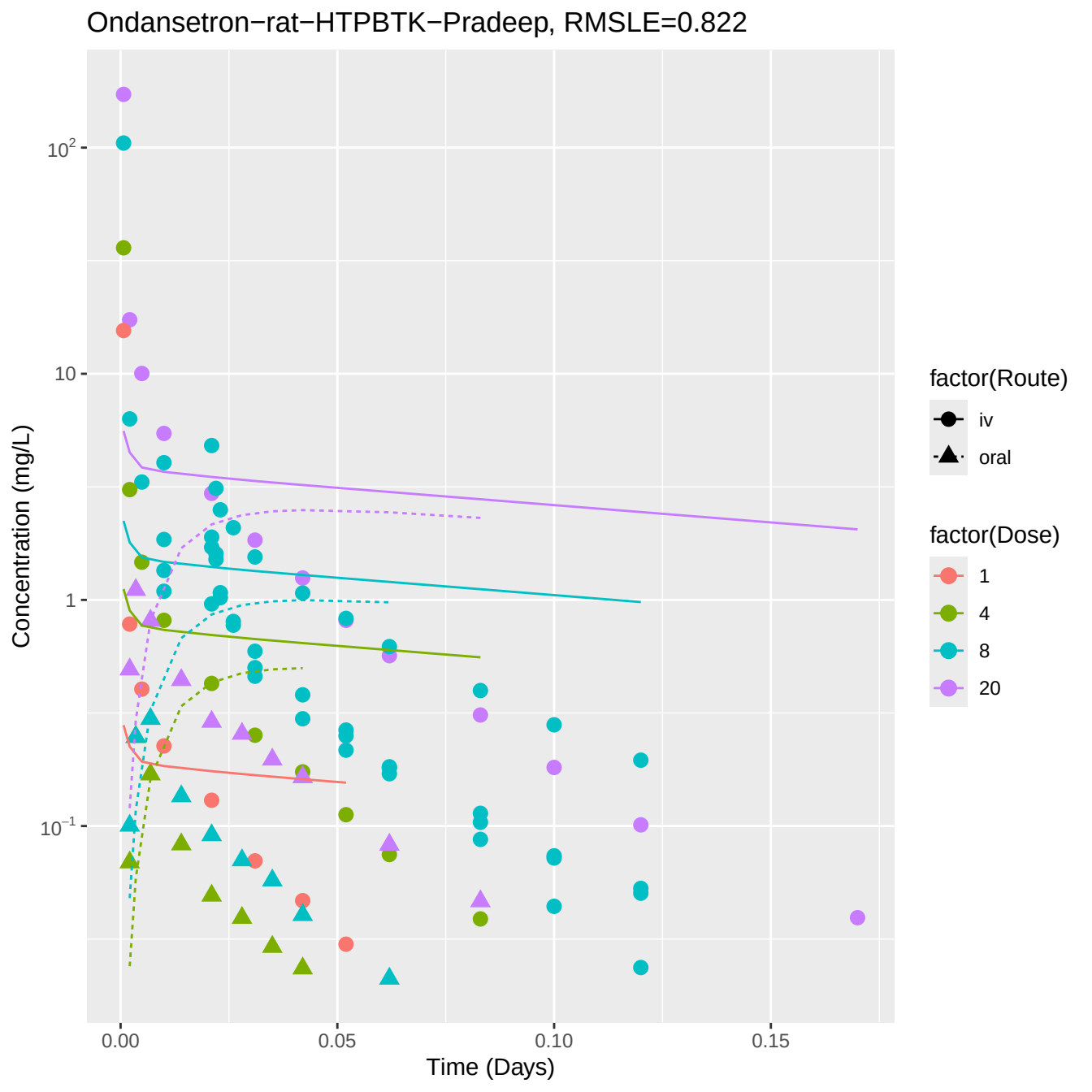

tert−Amyl methyl ether−human−HTPBTK−InVitro, RMSLE=0.743
Concentration (mg/L)
0.03
10^{-2}
0.003
10^{-3}
0.1
0.2
0.3
0.4
0.5
Time (Days)
factor(Route)
oral
factor(Dose)
0.0671
0.201

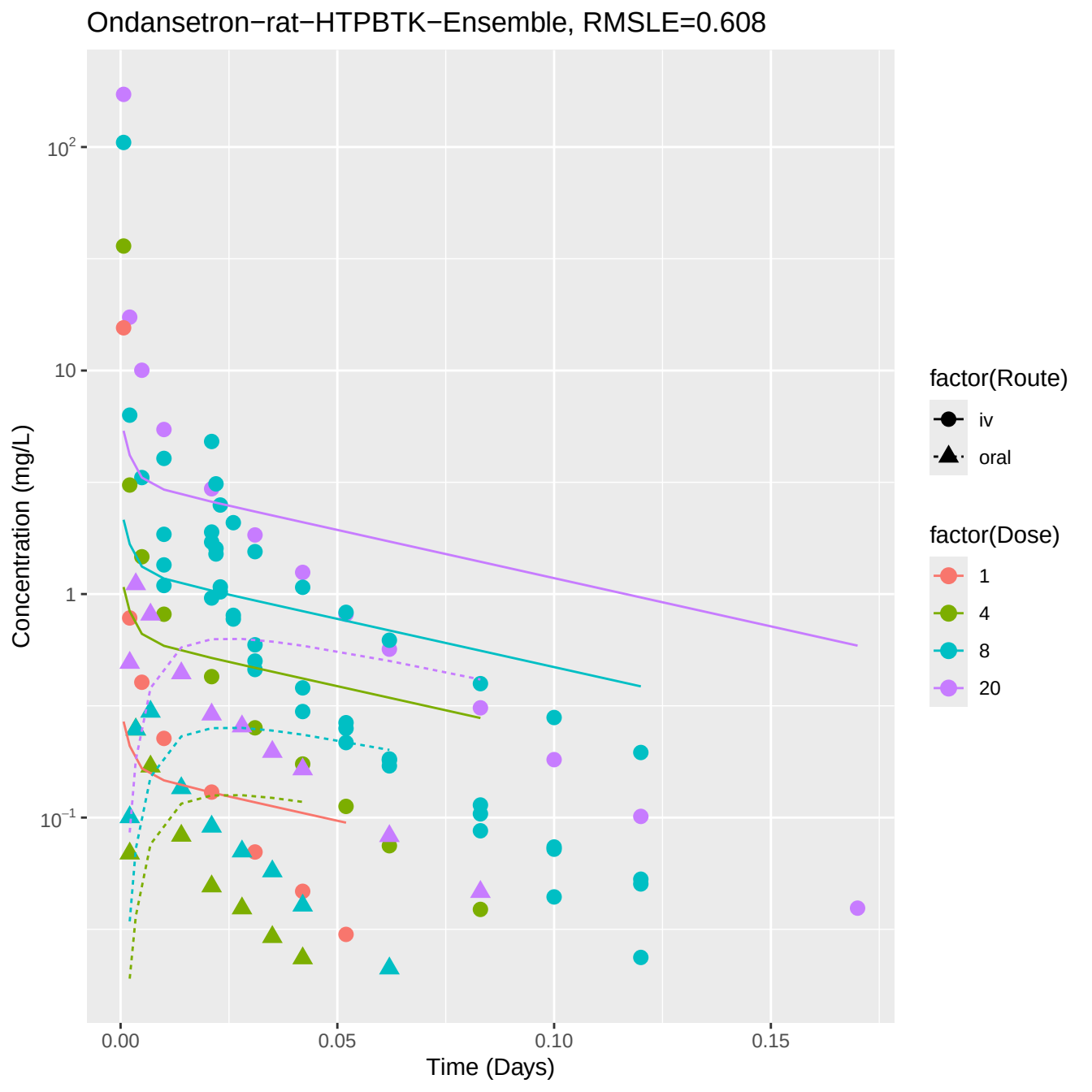

tert−Amyl methyl ether−human−HTPBTK−Ensemble, RMSLE=0.306
Concentration (mg/L)
10^{-2}
0.003
10^{-3}
0.1
0.2
0.3
0.4
0.5
Time (Days)
factor(Route)
oral
factor(Dose)
0.0671
0.201

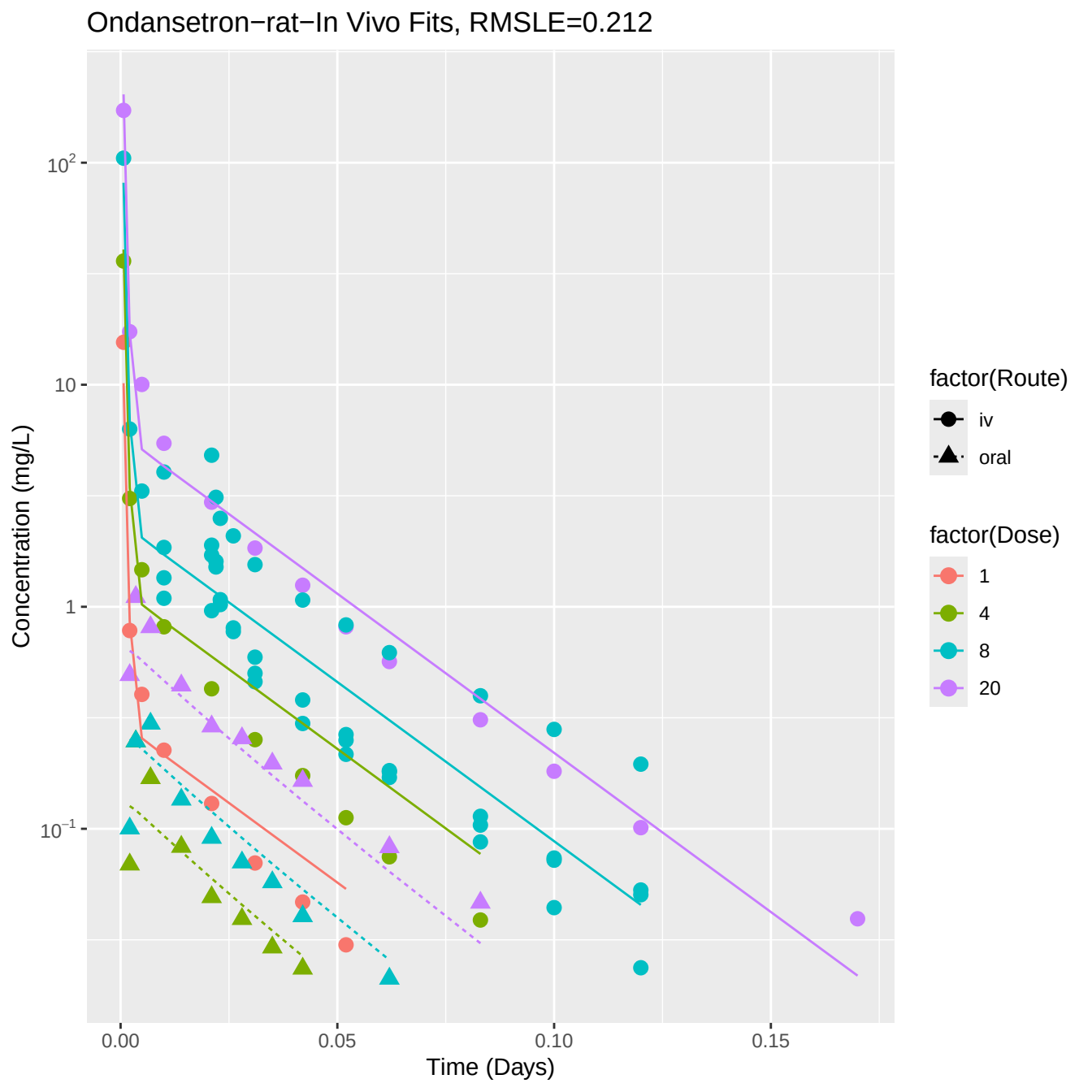

Ondansetron−rat−HTPBTK−InVitro, RMSLE=0.825
Concentration (mg/L)
10^2
10
1
10^{-1}
0.00
0.05
0.10
0.15
Time (Days)
factor(Route)
iv
oral
factor(Dose)
1
4
8
20

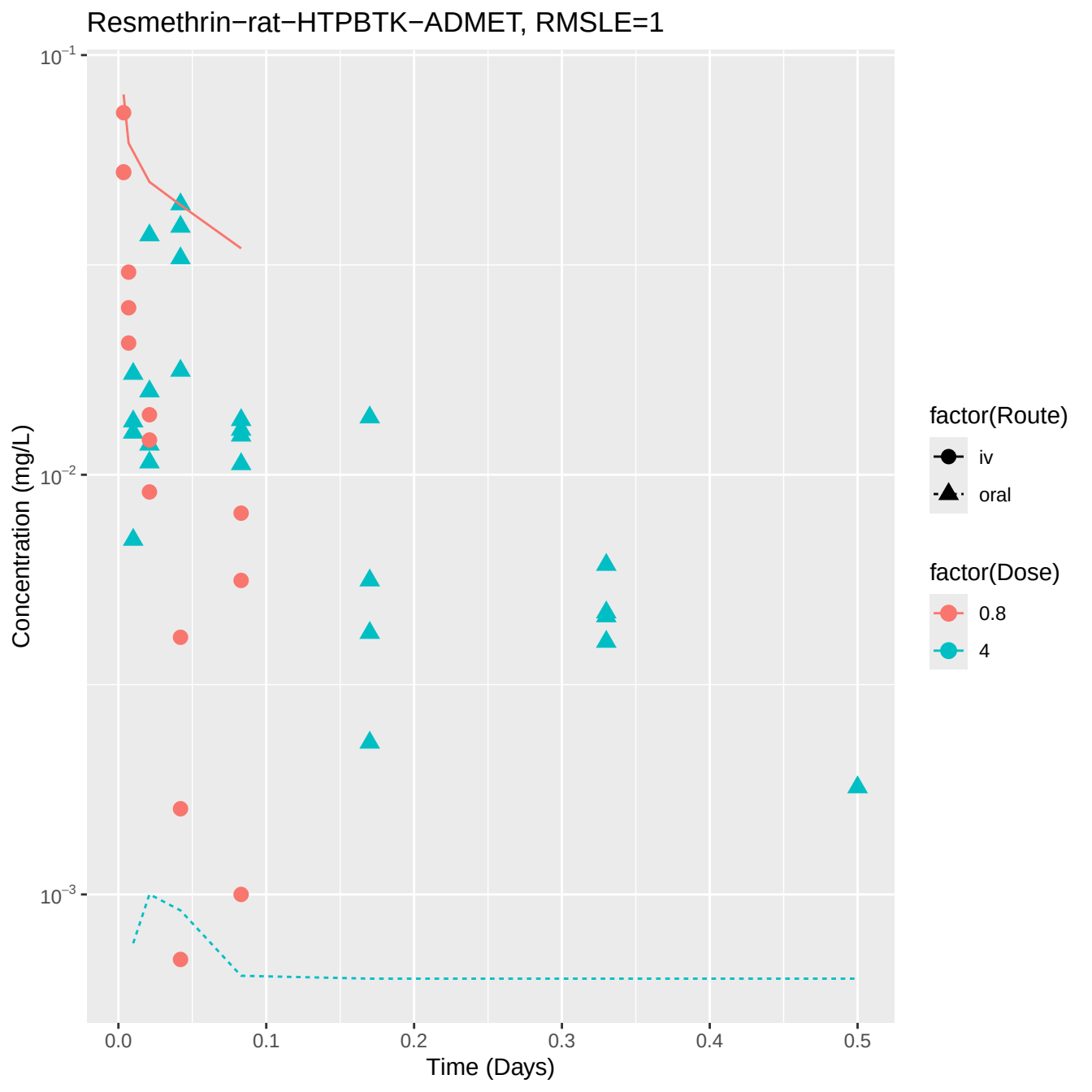

Ondansetron−rat−HTPBTK−ADMET, RMSLE=0.655
Concentration (mg/L)
10^2
10
1
10^{-1}
0.00
0.05
0.10
0.15
Time (Days)
factor(Route)
iv
oral
factor(Dose)
1
4
8
20

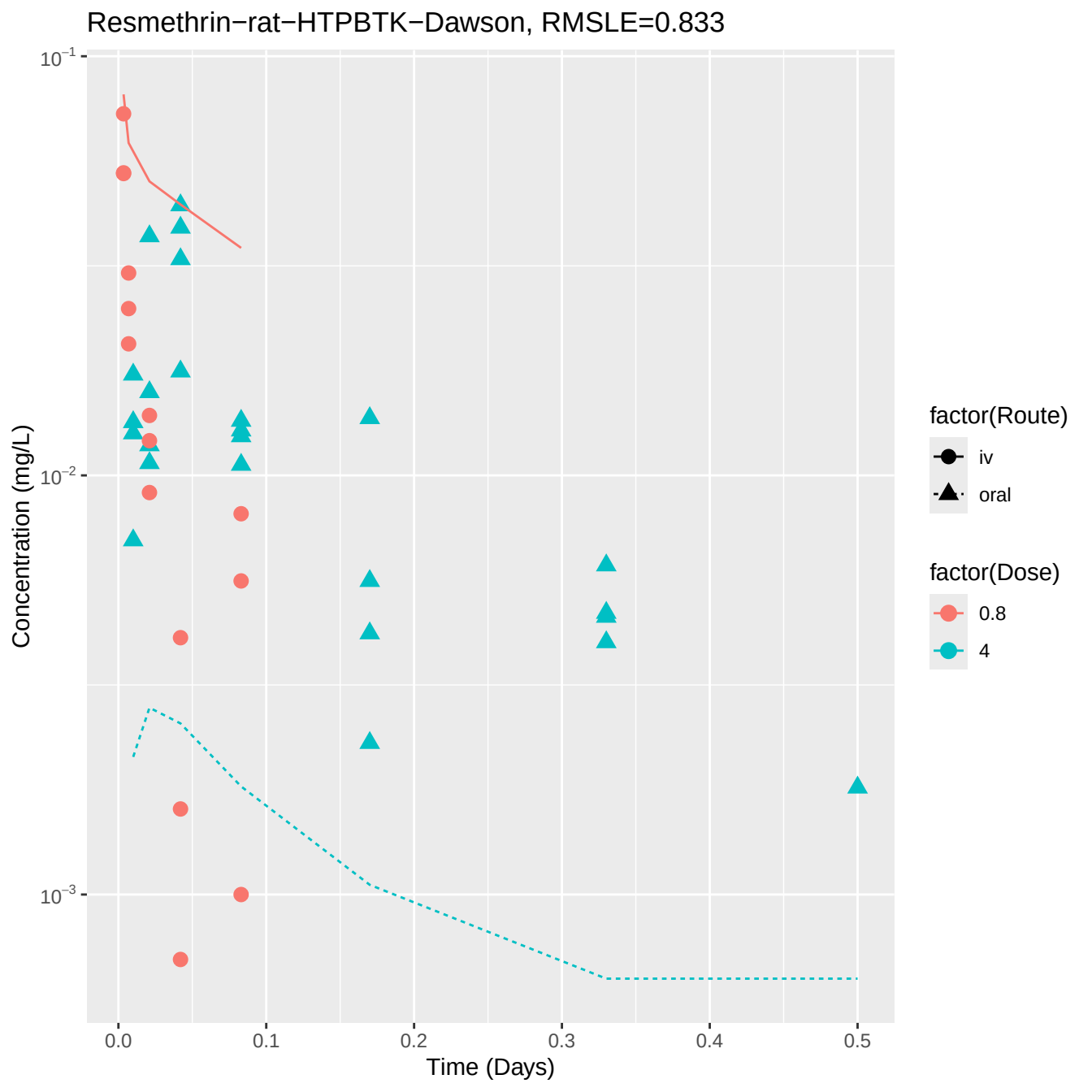

Ondansetron−rat−HTPBTK−Dawson, RMSLE=0.6
Concentration (mg/L)
10^2
10
1
10^{-1}
0.00
0.05
0.10
0.15
Time (Days)
factor(Route)
iv
oral
factor(Dose)
1
4
8
20

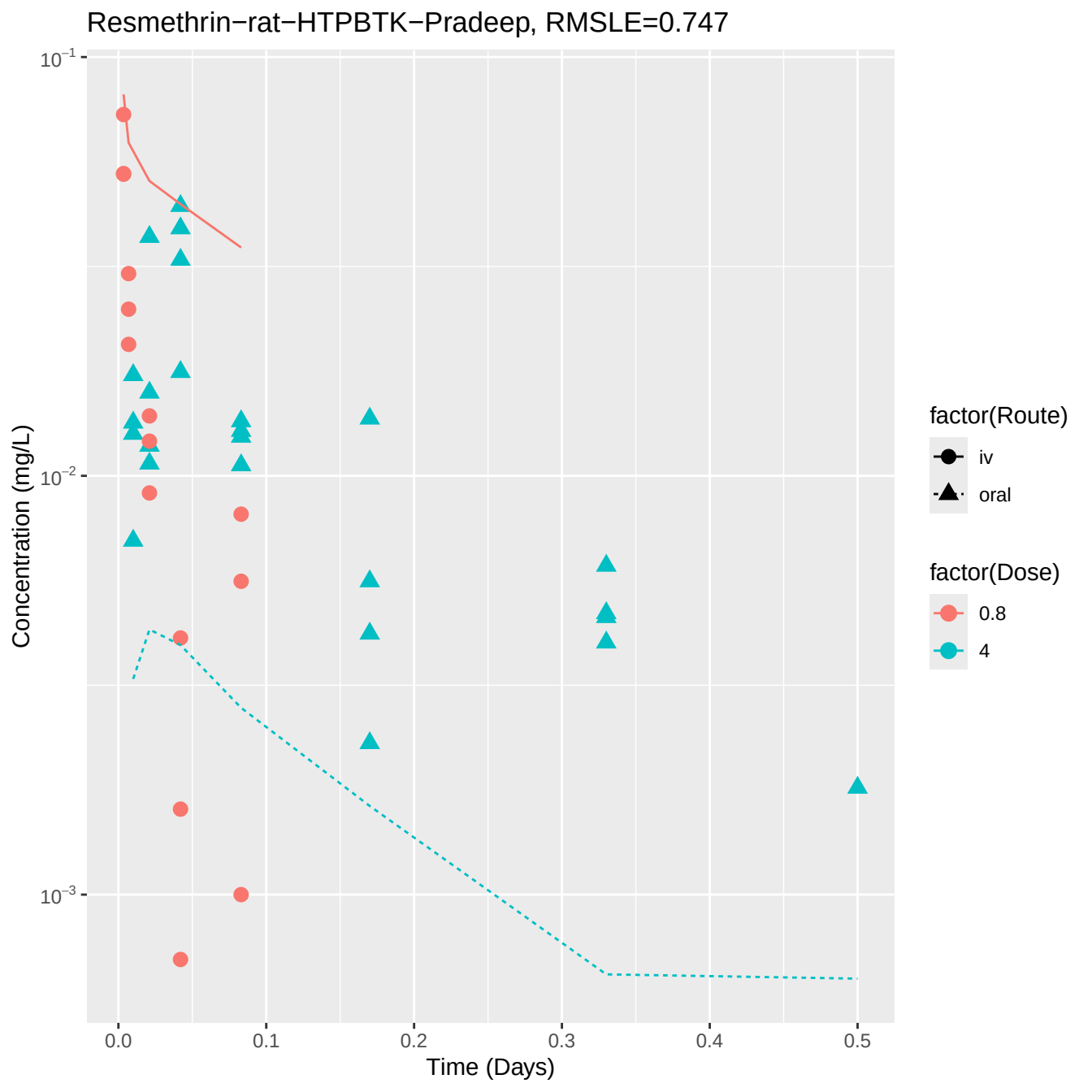

Ondansetron−rat−HTPBTK−Pradeep, RMSLE=0.822
Concentration (mg/L)
10^2
10
1
10^{-1}
0.00
0.05
0.10
0.15
Time (Days)
factor(Route)
iv
oral
factor(Dose)
1
4
8
20

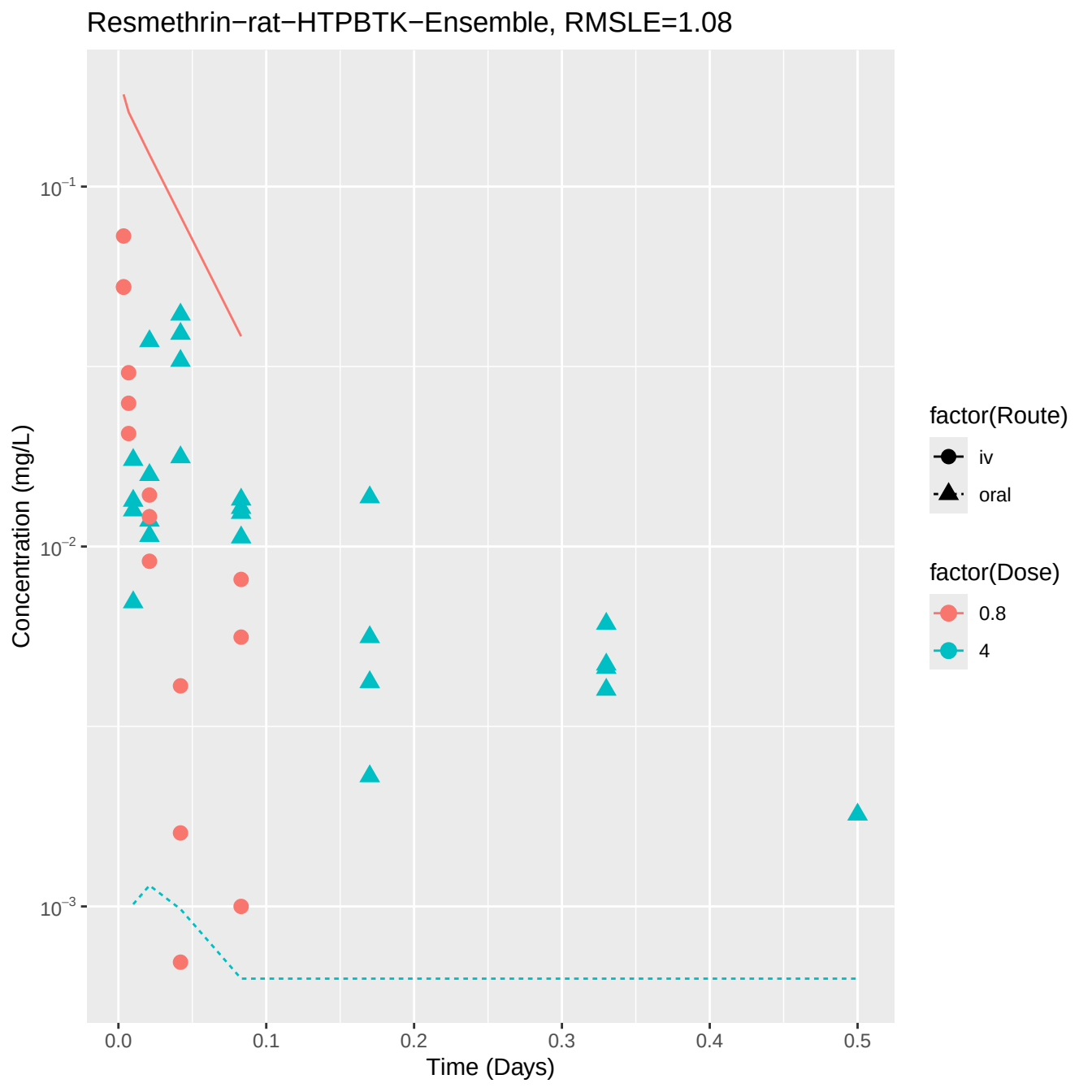

Ondansetron−rat−HTPBTK−Ensemble, RMSLE=0.608
Concentration (mg/L)
10^2
10
1
10^{-1}
0.00
0.05
0.10
0.15
Time (Days)
factor(Route)
iv
oral
factor(Dose)
1
4
8
20

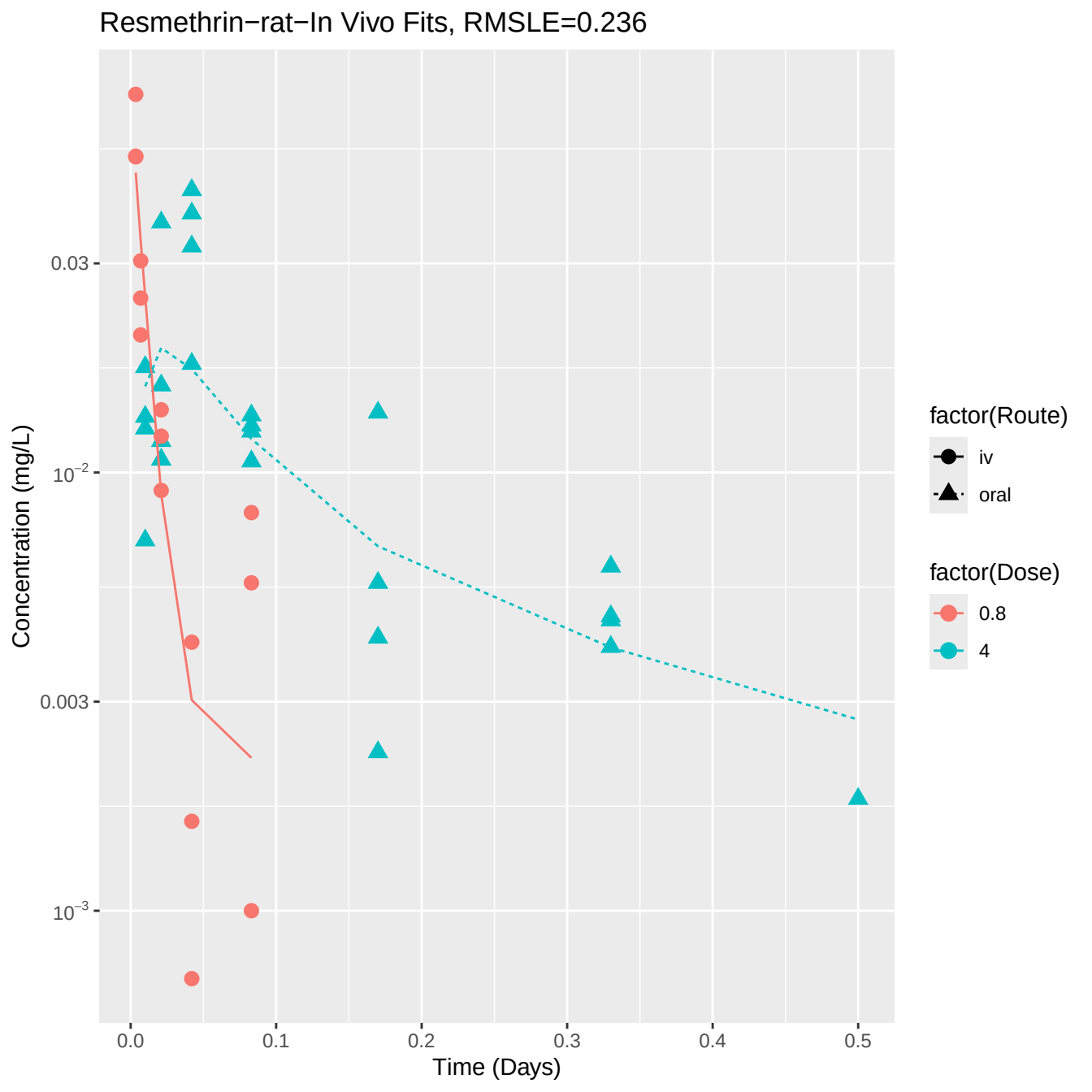

Ondansetron−rat−In Vivo Fits, RMSLE=0.212
Concentration (mg/L)
10^2
10
1
10^{-1}
0.00
0.05
0.10
0.15
Time (Days)
factor(Route)
iv
oral
factor(Dose)
1
4
8
20

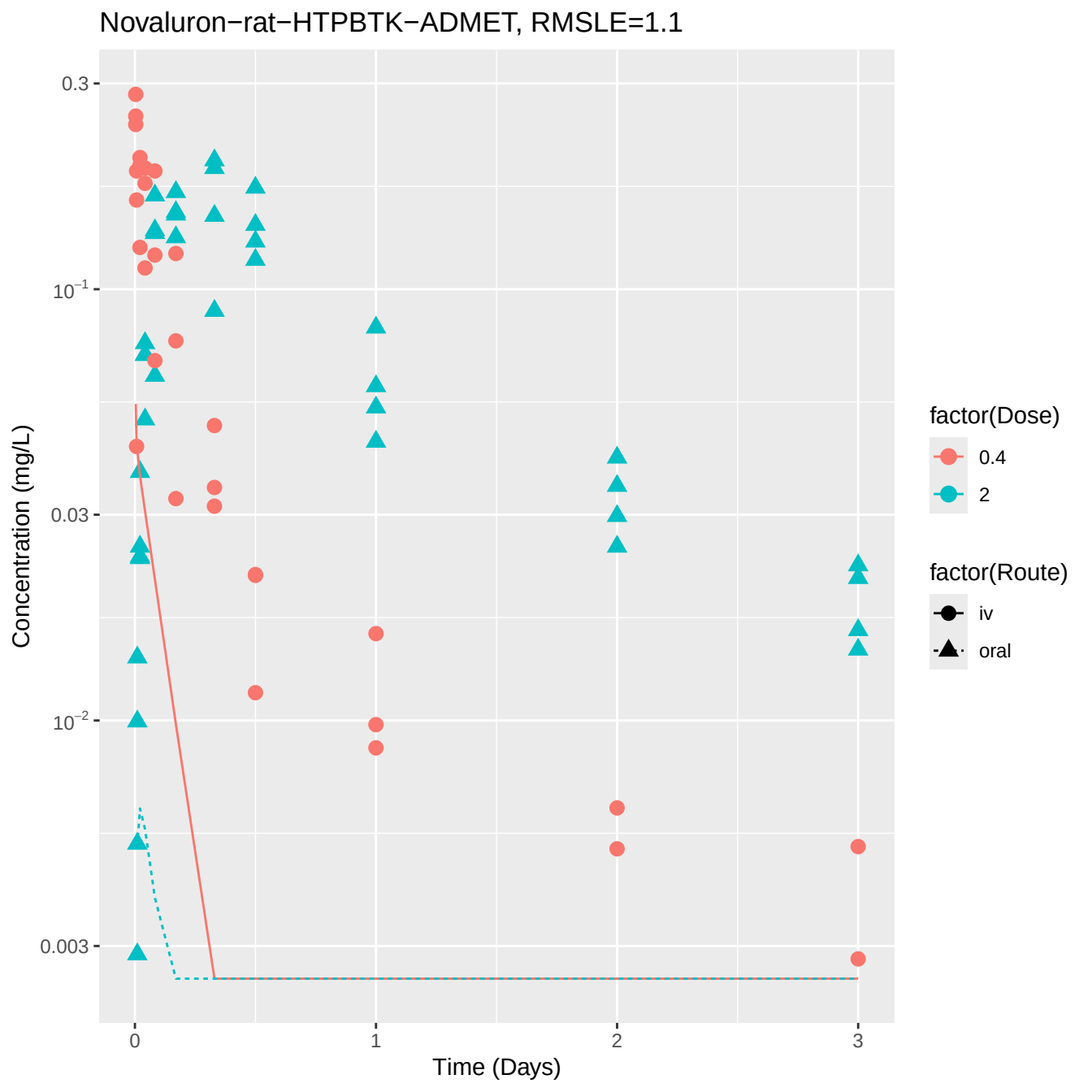

Resmethrin−rat−HTPBTK−ADMET, RMSLE=1
Concentration (mg/L)
10^{-1}
10^{-2}
10^{-3}
0.0
0.1
0.2
0.3
0.4
0.5
Time (Days)
factor(Route)
iv
oral
factor(Dose)
0.8
4

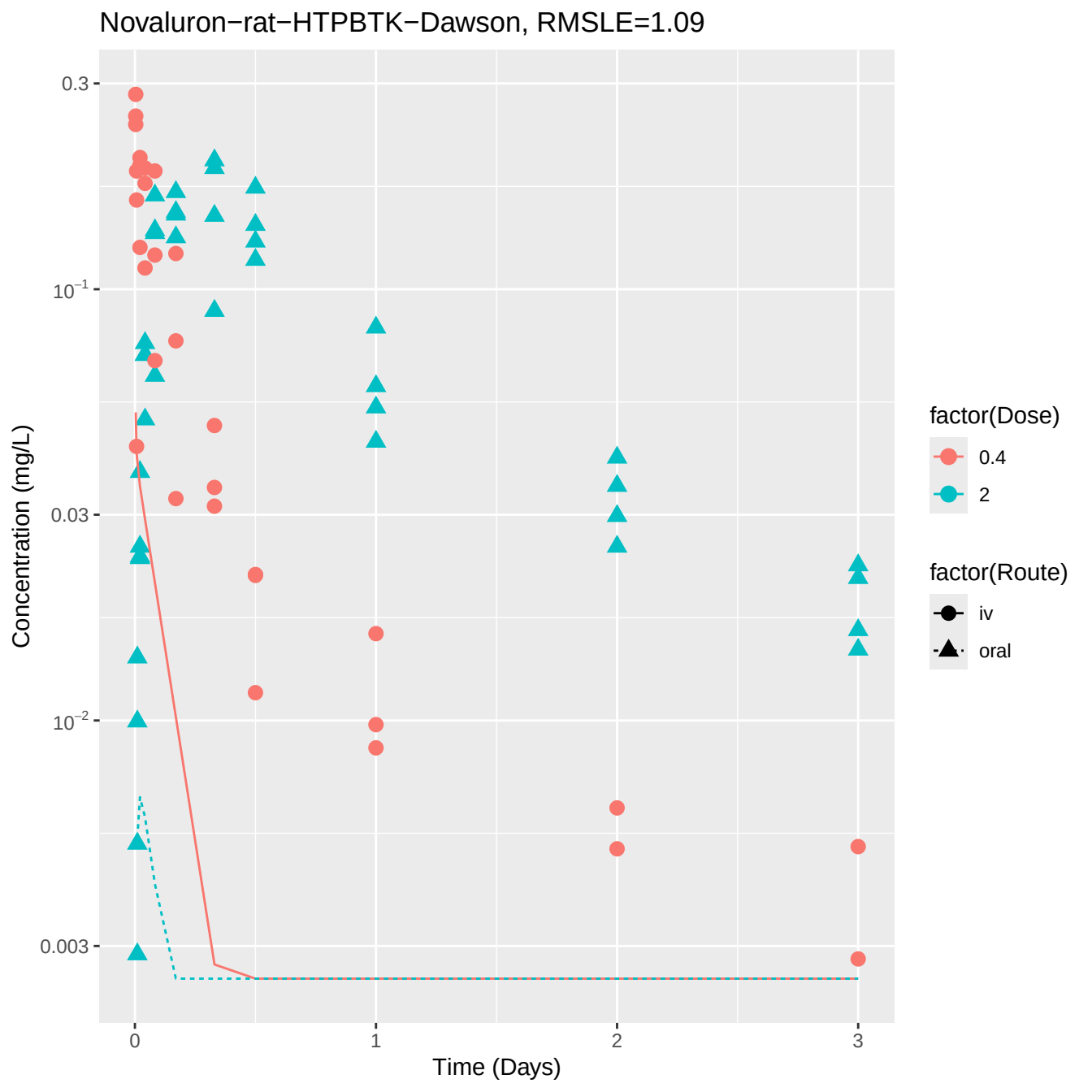

Resmethrin−rat−HTPBTK−Dawson, RMSLE=0.833
Concentration (mg/L)
10^{-1}
10^{-2}
10^{-3}
0.0
0.1
0.2
0.3
0.4
0.5
Time (Days)
factor(Route)
iv
oral
factor(Dose)
0.8
4

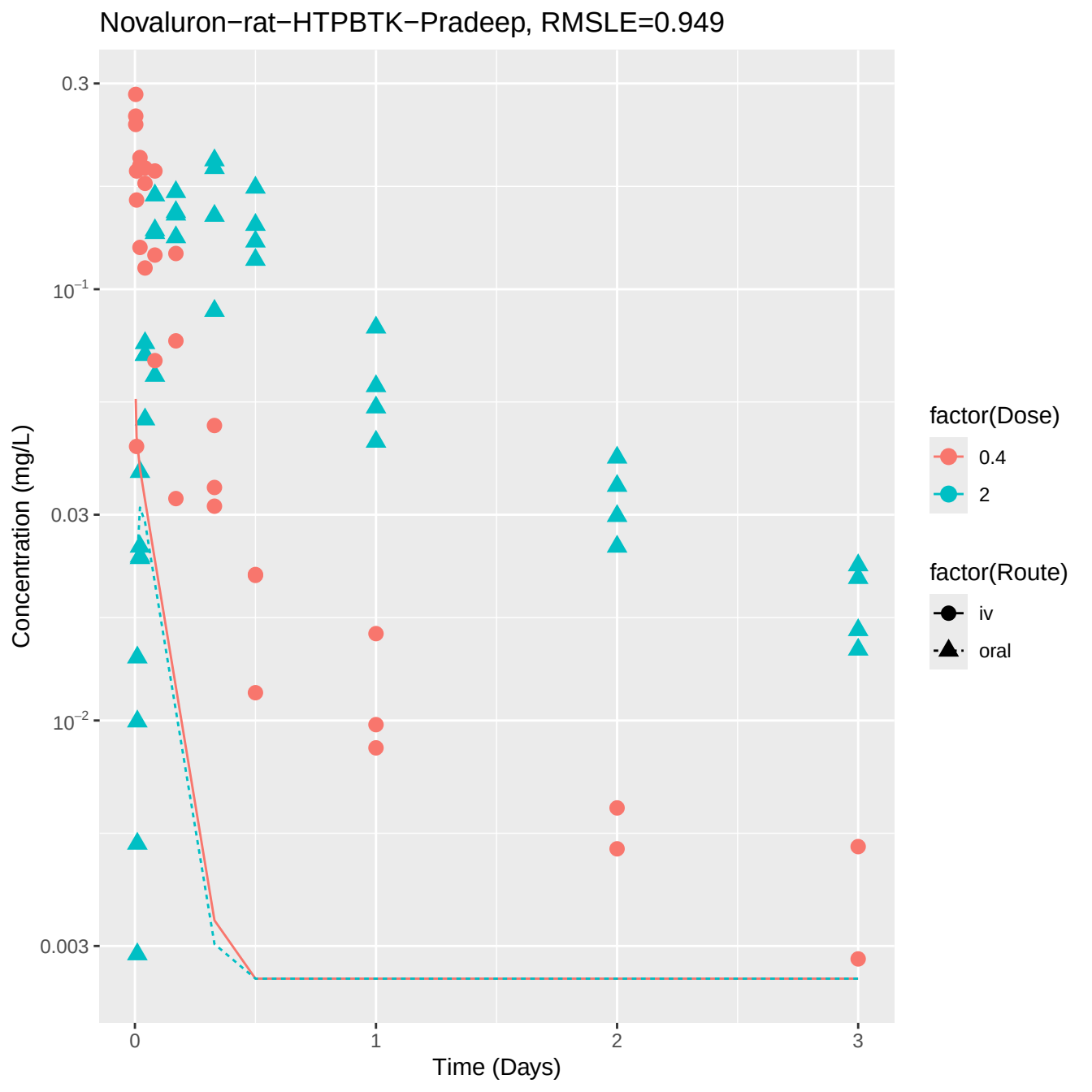

Resmethrin−rat−HTPBTK−Pradeep, RMSLE=0.747
Concentration (mg/L)
10^{-1}
10^{-2}
10^{-3}
0.0
0.1
0.2
0.3
0.4
0.5
Time (Days)
factor(Route)
iv
oral
factor(Dose)
0.8
4

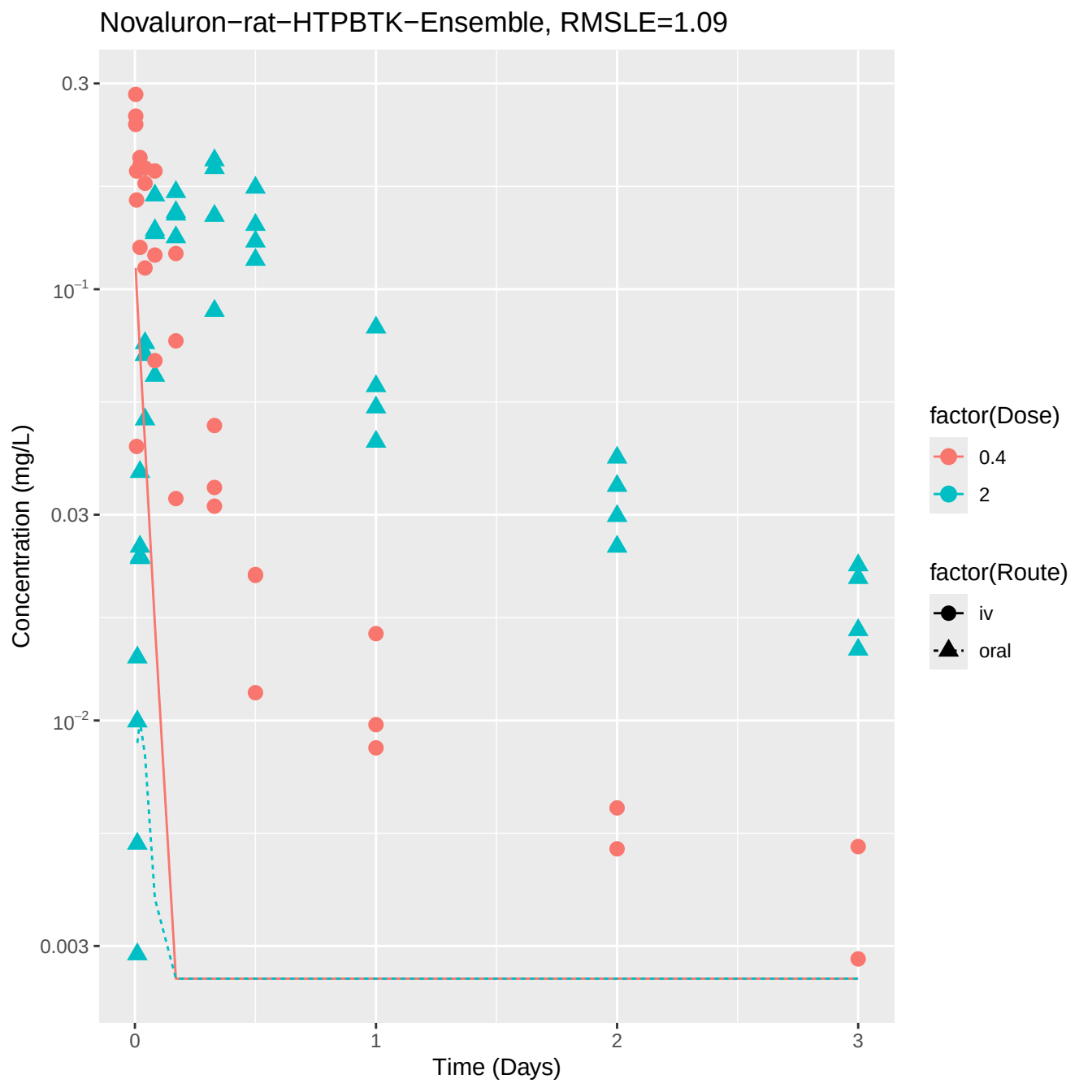

Resmethrin−rat−HTPBTK−Ensemble, RMSLE=1.08
Concentration (mg/L)
10^{-1}
10^{-2}
10^{-3}
0.0
0.1
0.2
0.3
0.4
0.5
Time (Days)
factor(Route)
iv
oral
factor(Dose)
0.8
4

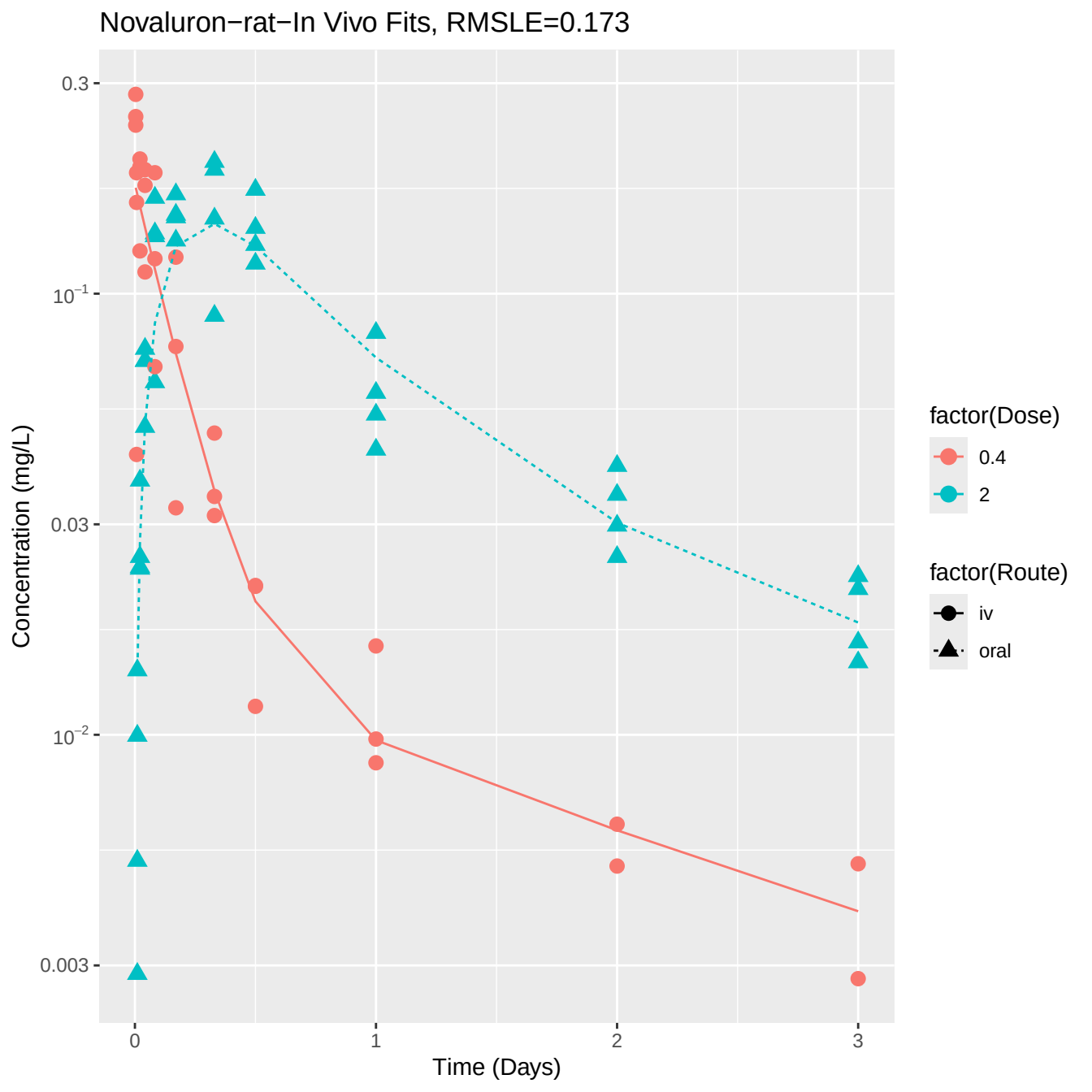

Resmethrin−rat−In Vivo Fits, RMSLE=0.236
Concentration (mg/L)
0.03
10^{-2}
0.003
10^{-3}
0.0
0.1
0.2
0.3
0.4
0.5
Time (Days)
factor(Route)
iv
oral
factor(Dose)
0.8
4

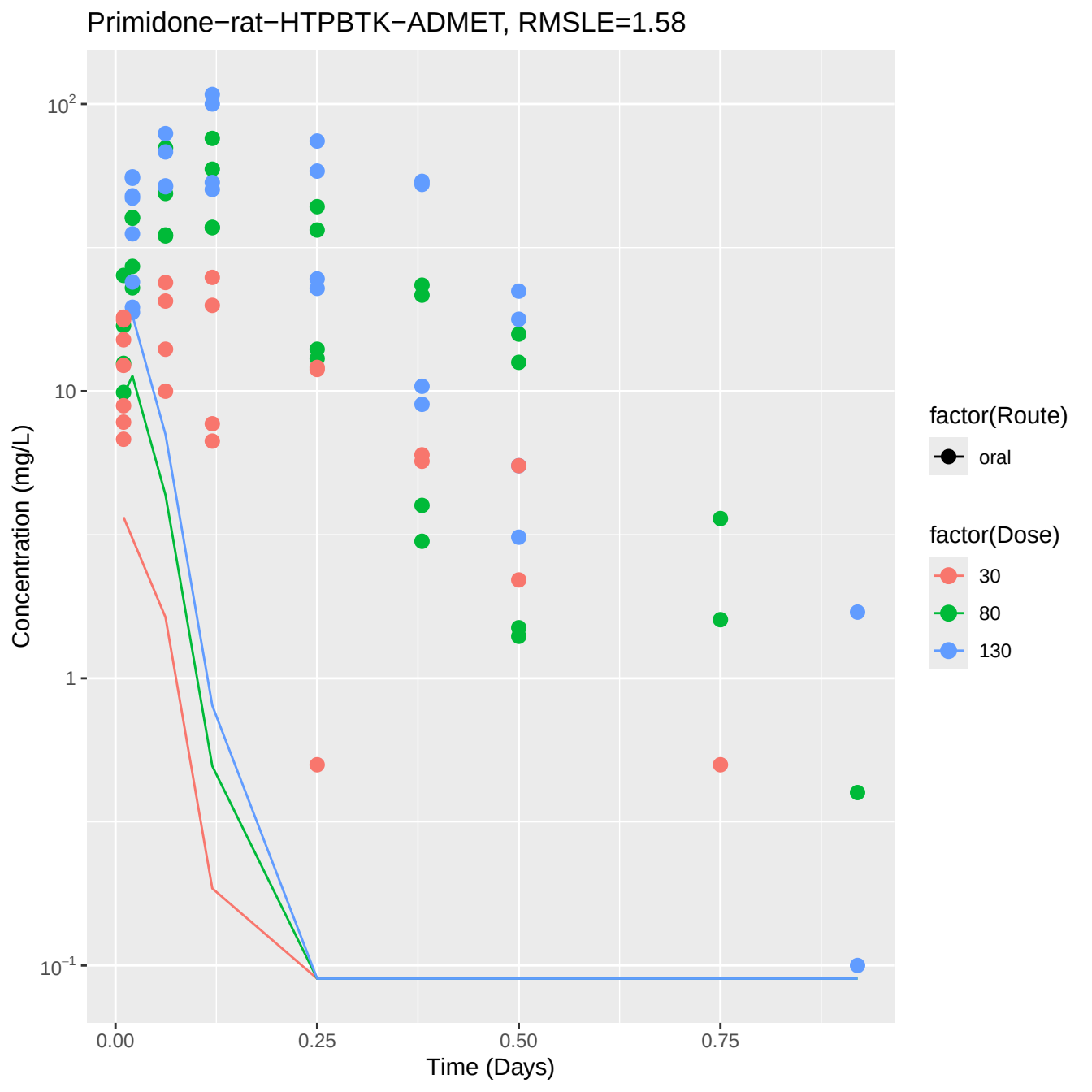

Novaluron−rat−HTPBTK−ADMET, RMSLE=1.1
Concentration (mg/L)
0.3
10^{-1}
0.03
10^{-2}
0.003
0
1
2
3
Time (Days)
factor(Dose)
0.4
2
factor(Route)
iv
oral

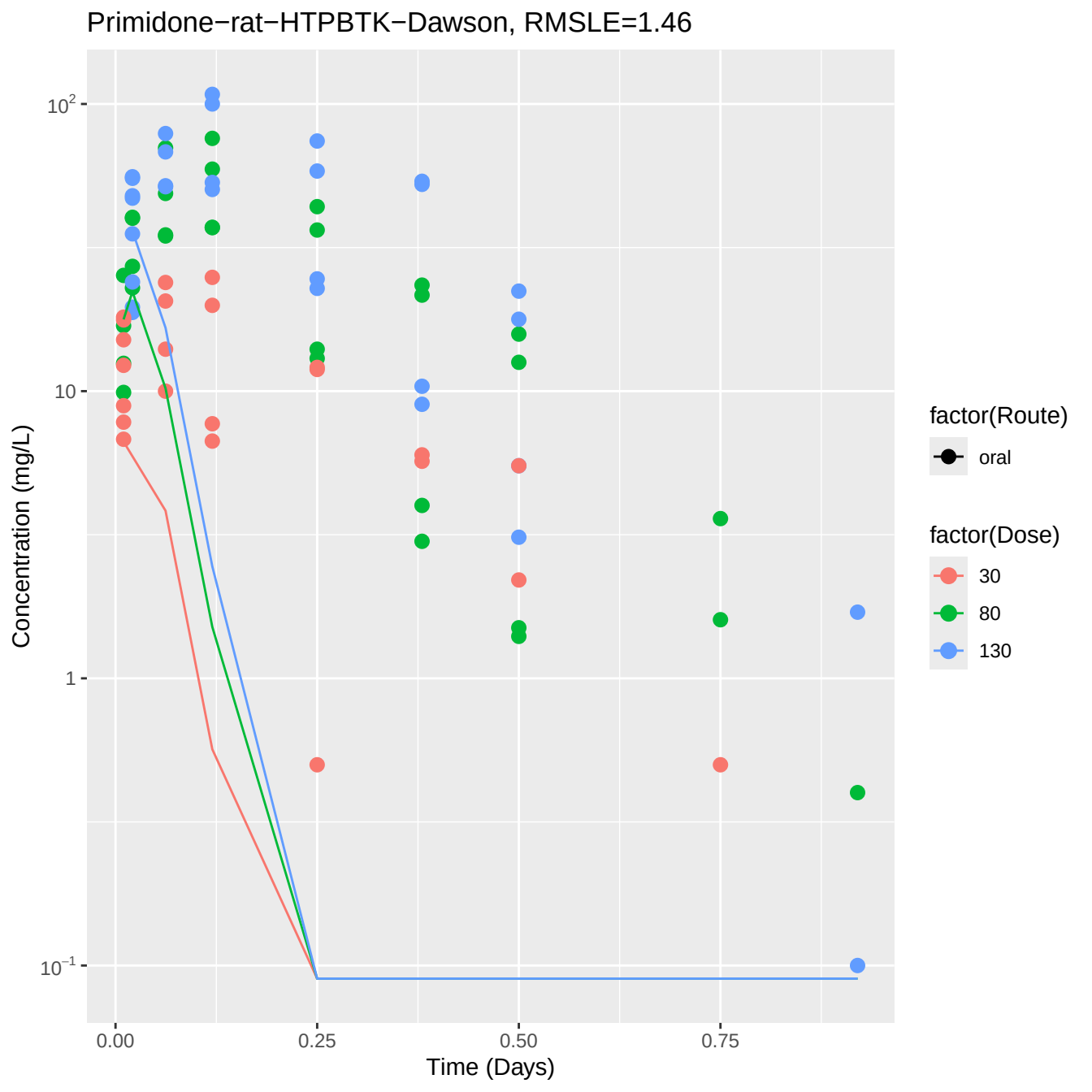

Novaluron−rat−HTPBTK−Dawson, RMSLE=1.09
0.3
10^{-1}
0.03
10^{-2}
0.003
Concentration (mg/L)
0
1
2
3
Time (Days)
factor(Dose)
0.4
2
factor(Route)
iv
oral

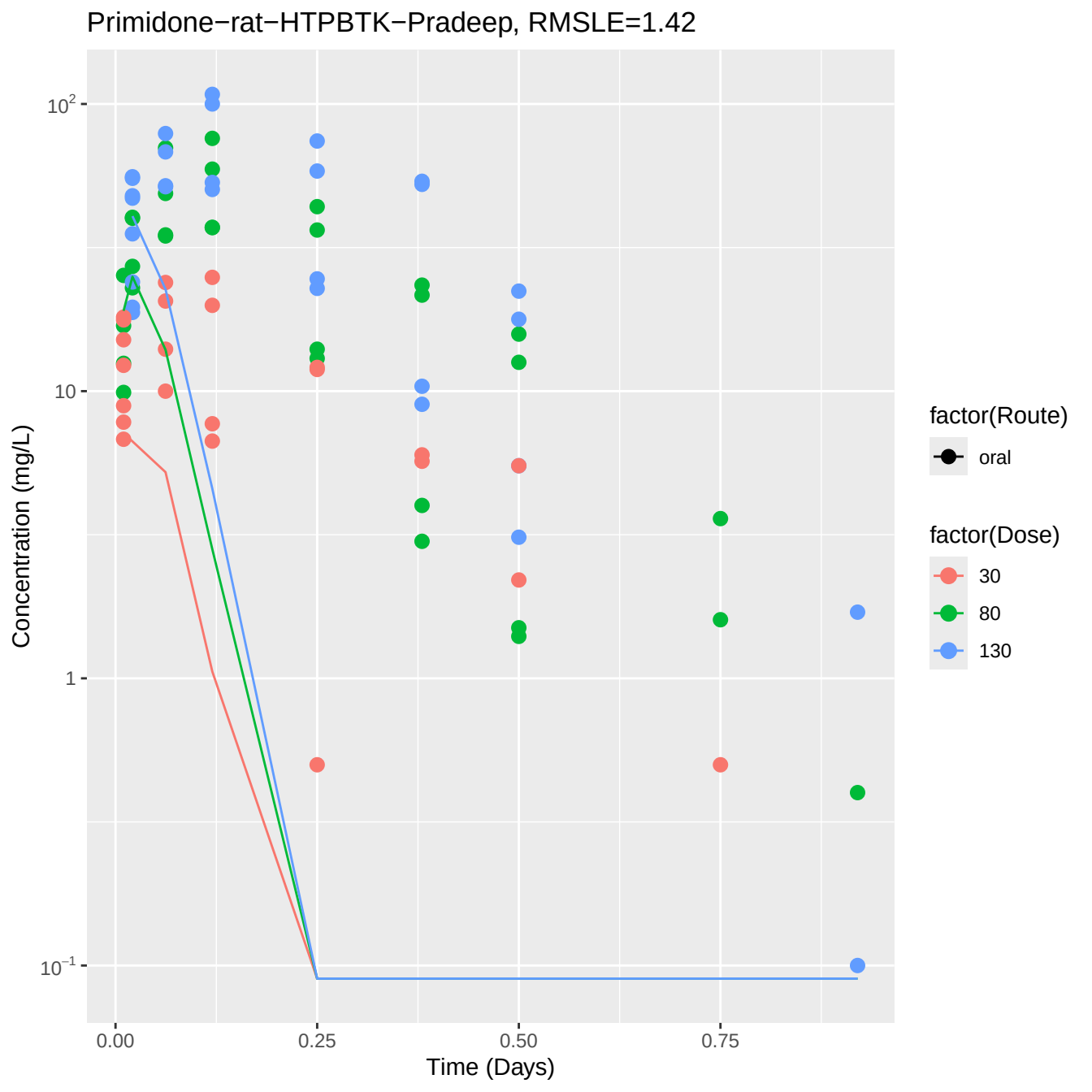

Novaluron−rat−HTPBTK−Pradeep, RMSLE=0.949
Concentration (mg/L)
0.3
10^{-1}
0.03
10^{-2}
0.003
0
1
2
3
Time (Days)
factor(Dose)
0.4
2
factor(Route)
iv
oral

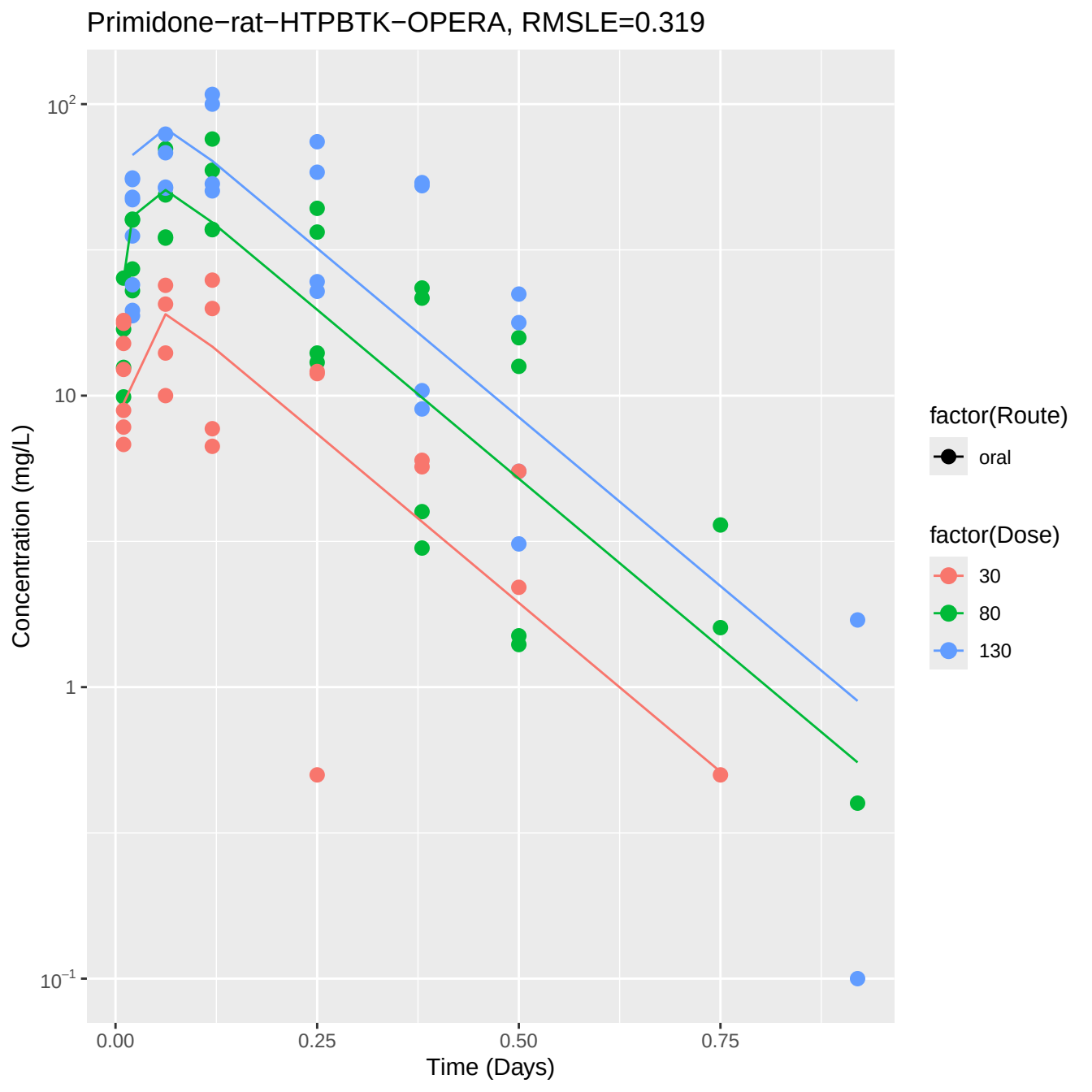

Novaluron−rat−HTPBTK−Ensemble, RMSLE=1.09
Concentration (mg/L)
0.3
10^{-1}
0.03
10^{-2}
0.003
0
1
2
3
Time (Days)
factor(Dose)
0.4
2
factor(Route)
iv
oral

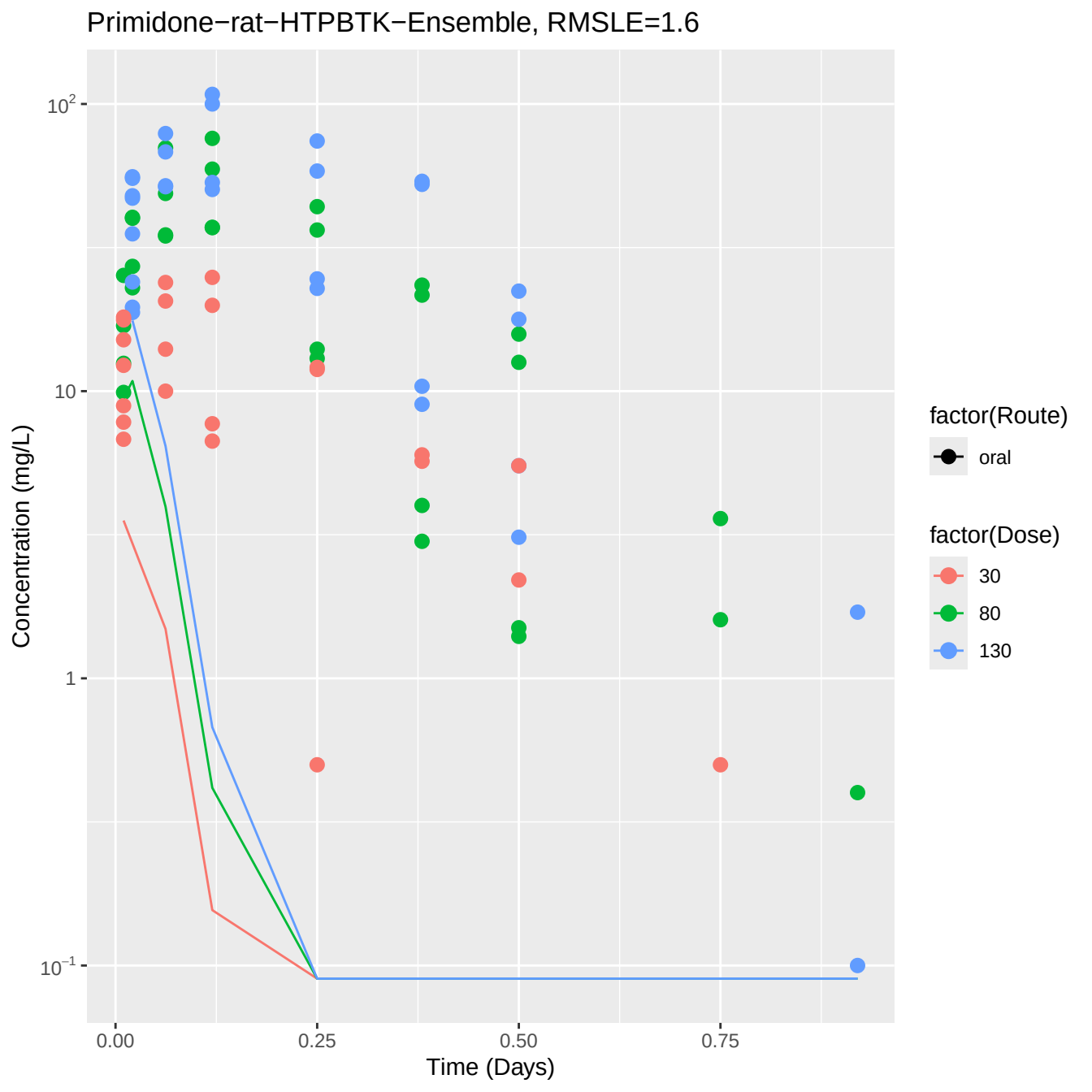

Novaluron−rat−In Vivo Fits, RMSLE=0.173
0.3
10^{-1}
0.03
10^{-2}
0.003
Concentration (mg/L)
0
1
2
3
Time (Days)
factor(Dose)
0.4
2
factor(Route)
iv
oral

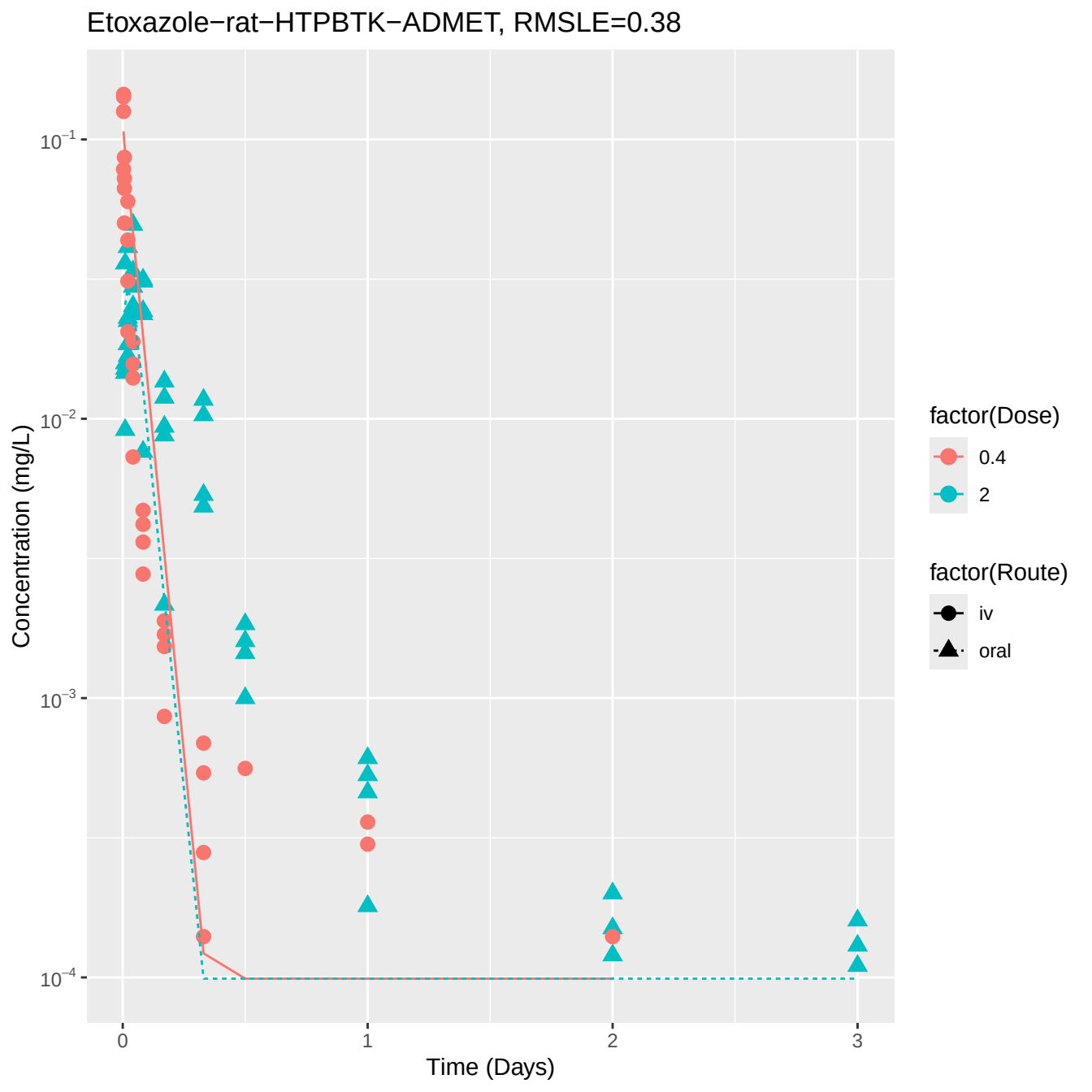

Primidone−rat−HTPBTK−ADMET, RMSLE=1.58
Concentration (mg/L)
10^2
10
1
10^{-1}
0.00
0.25
0.50
0.75
Time (Days)
factor(Route)
oral
factor(Dose)
30
80
130

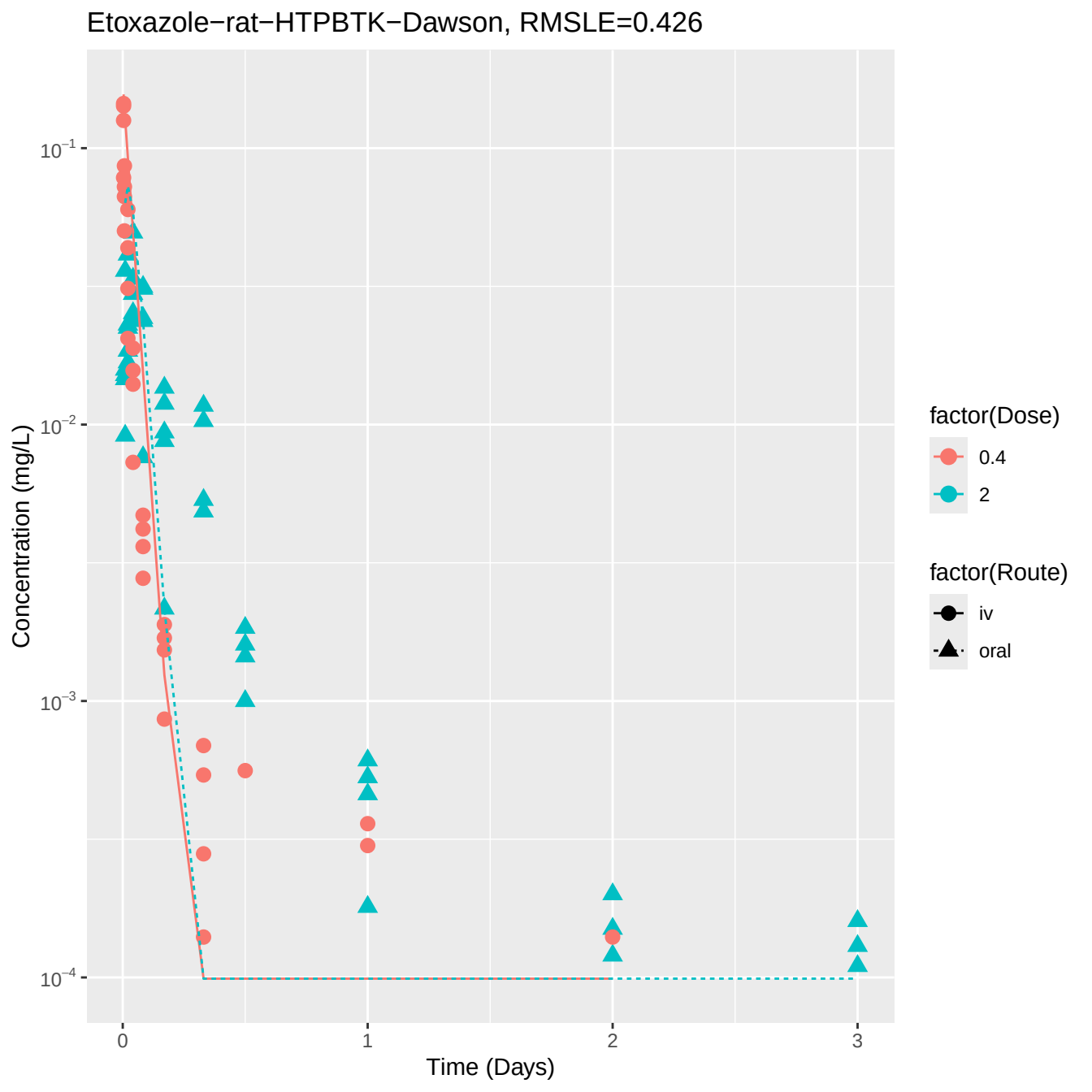

Primidone−rat−HTPBTK−Dawson, RMSLE=1.46
Concentration (mg/L)
10^2
10
1
10^{-1}
0.00
0.25
0.50
0.75
Time (Days)
factor(Route)
oral
factor(Dose)
30
80
130

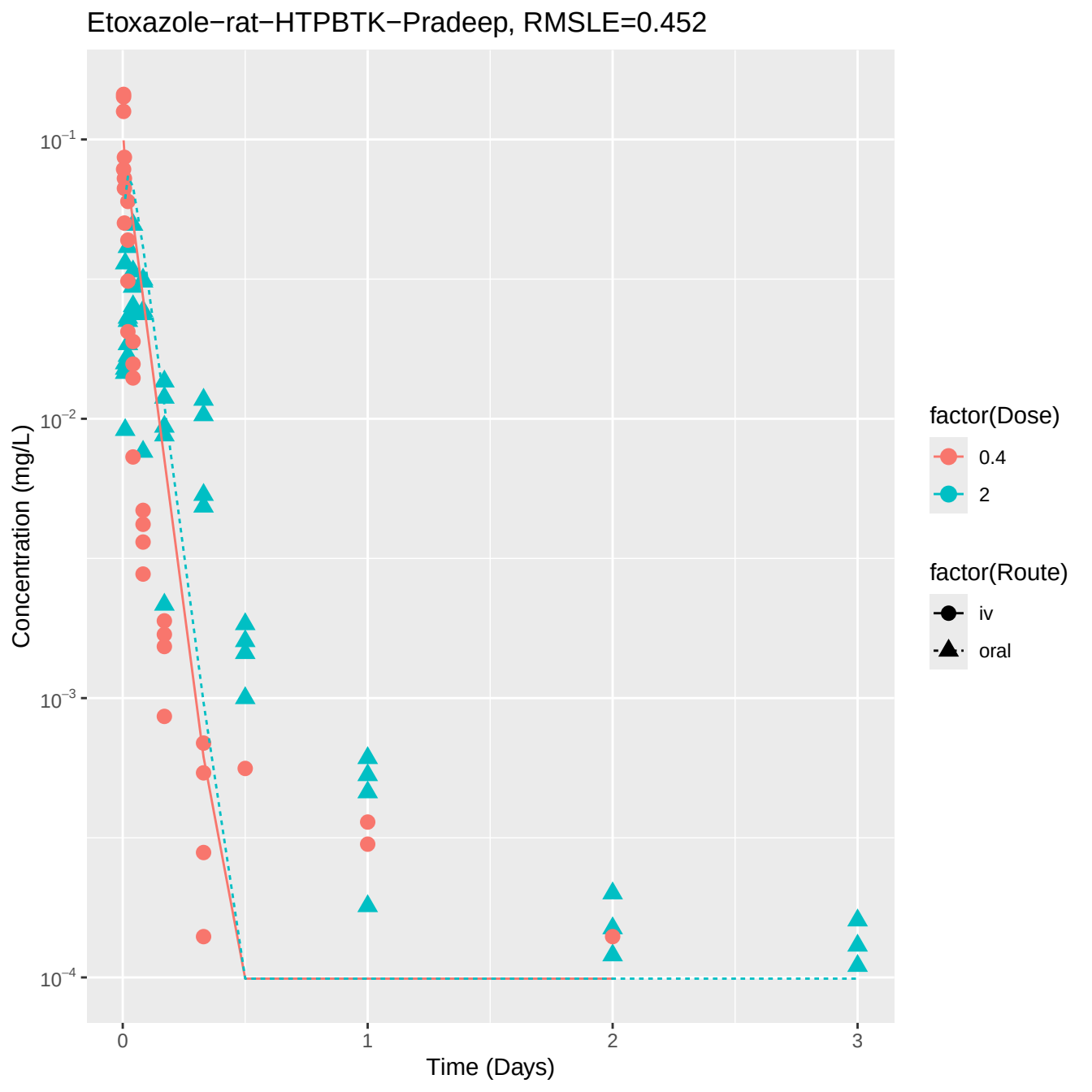

Primidone−rat−HTPBTK−Pradeep, RMSLE=1.42
Concentration (mg/L)
10^2
10
1
10^{-1}
0.00
0.25
0.50
0.75
Time (Days)
factor(Route)
oral
factor(Dose)
30
80
130

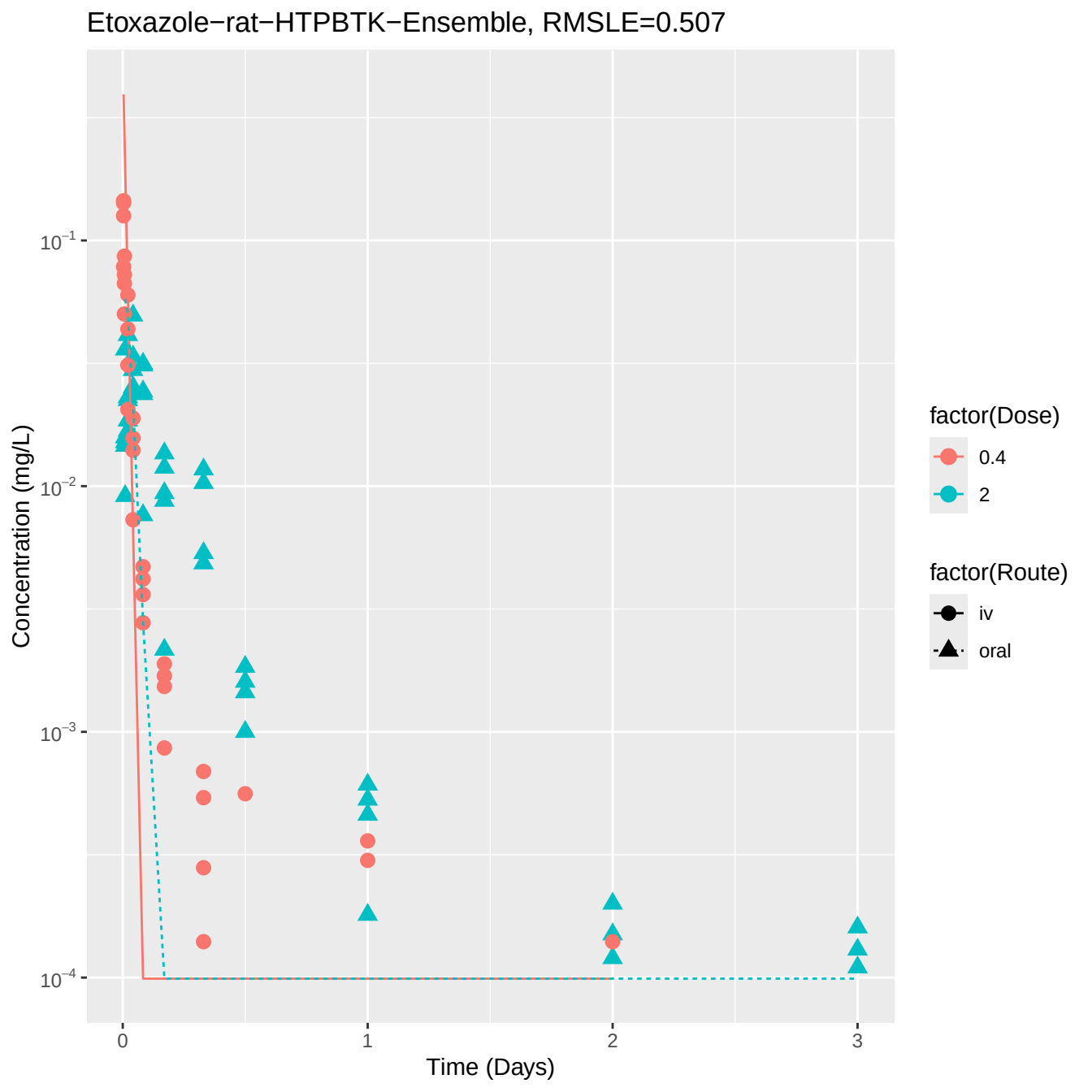

Primidone−rat−HTPBTK−OPERA, RMSLE=0.319
Concentration (mg/L)
10^2
10
1
10^{-1}
0.00
0.25
0.50
0.75
Time (Days)
factor(Route)
oral
factor(Dose)
30
80
130

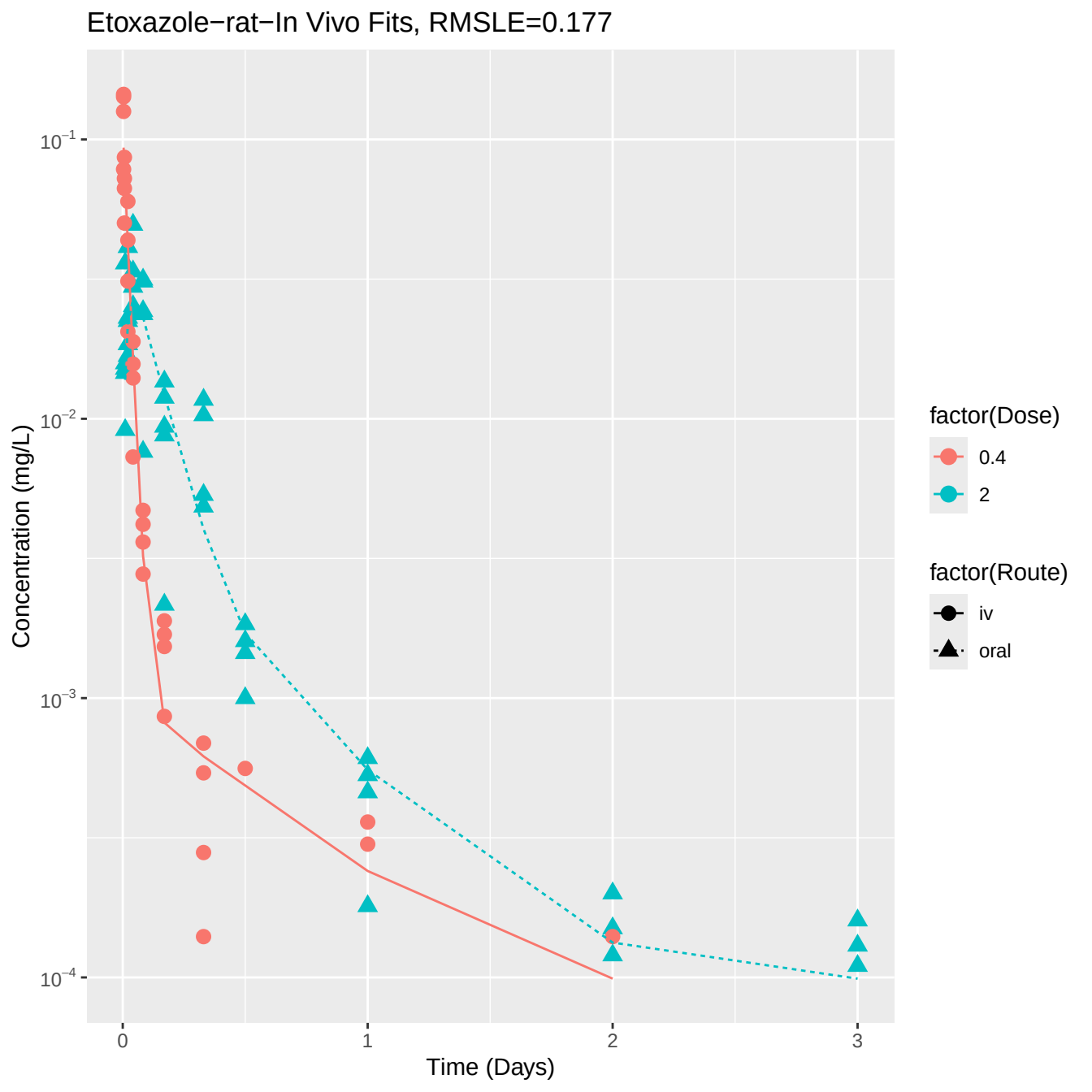

Primidone−rat−HTPBTK−Ensemble, RMSLE=1.6
Concentration (mg/L)
10^2
10
1
10^{-1}
0.00
0.25
0.50
0.75
Time (Days)
factor(Route)
oral
factor(Dose)
30
80
130

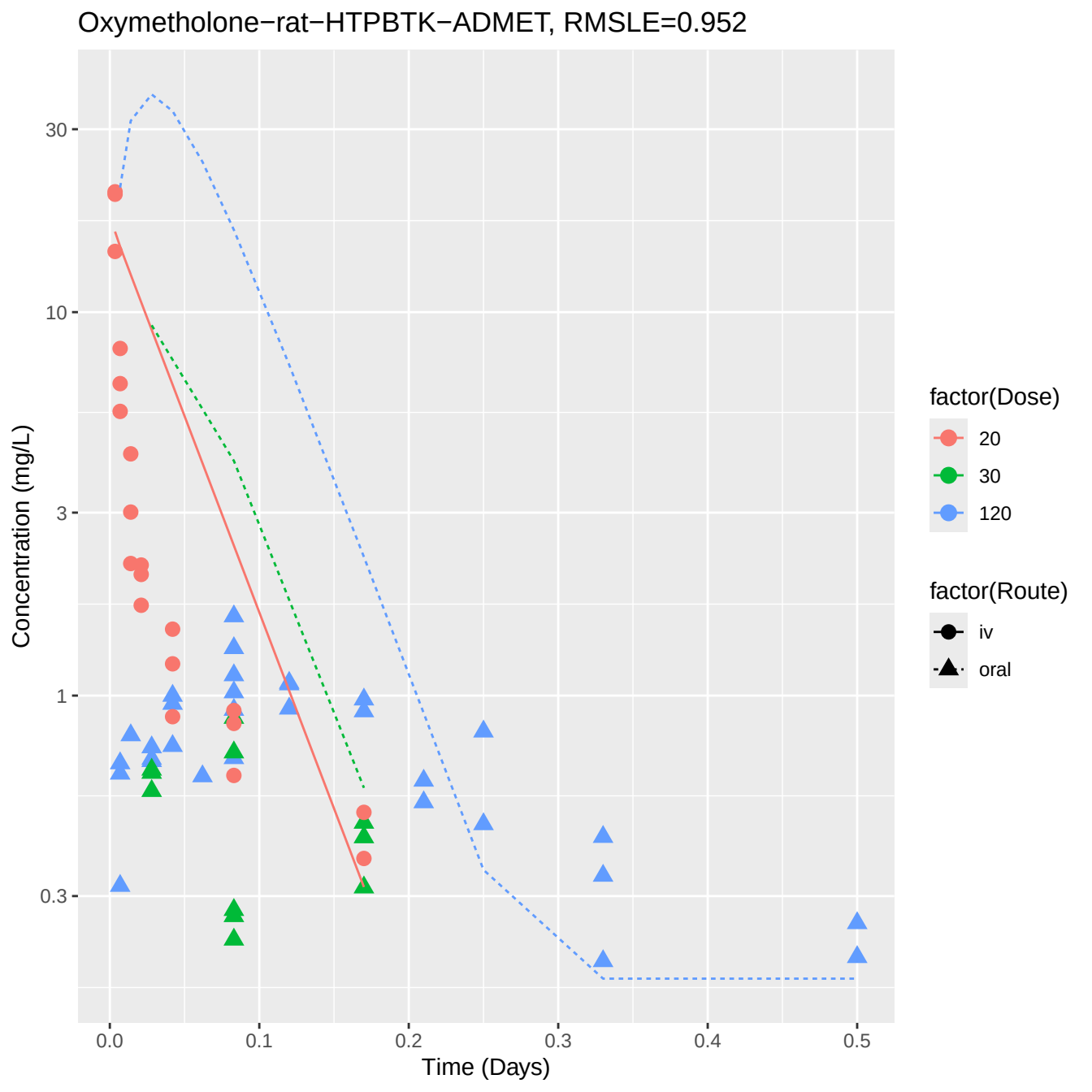

Etoxazole−rat−HTPBTK−ADMET, RMSLE=0.38
Concentration (mg/L)
10^{-1}
10^{-2}
10^{-3}
10^{-4}
0
1
2
3
Time (Days)
factor(Dose)
0.4
2
factor(Route)
iv
oral

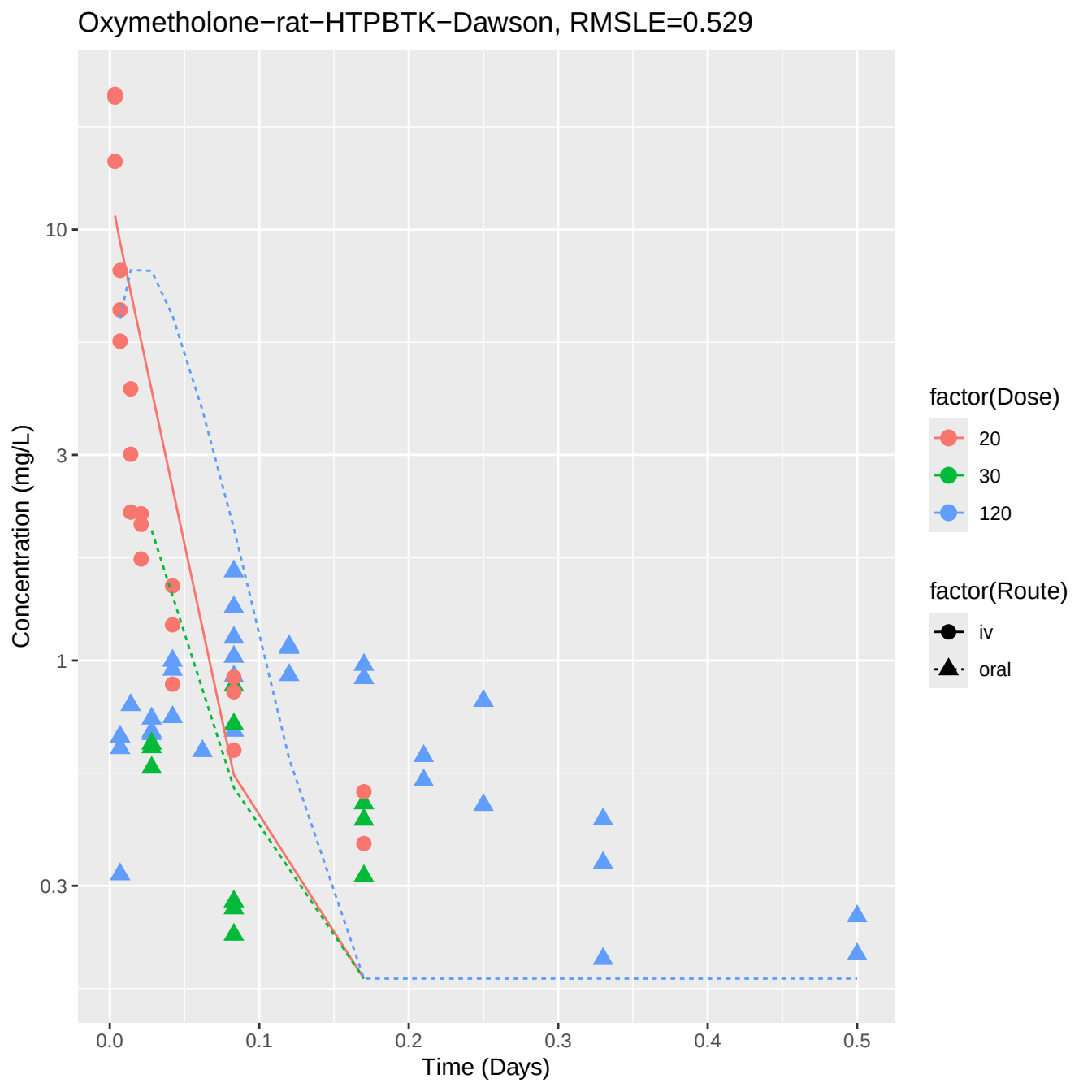

Etoxazole−rat−HTPBTK−Dawson, RMSLE=0.426
Concentration (mg/L)
10^{-1}
10^{-2}
10^{-3}
10^{-4}
0
1
2
3
Time (Days)
factor(Dose)
0.4
2
factor(Route)
iv
oral

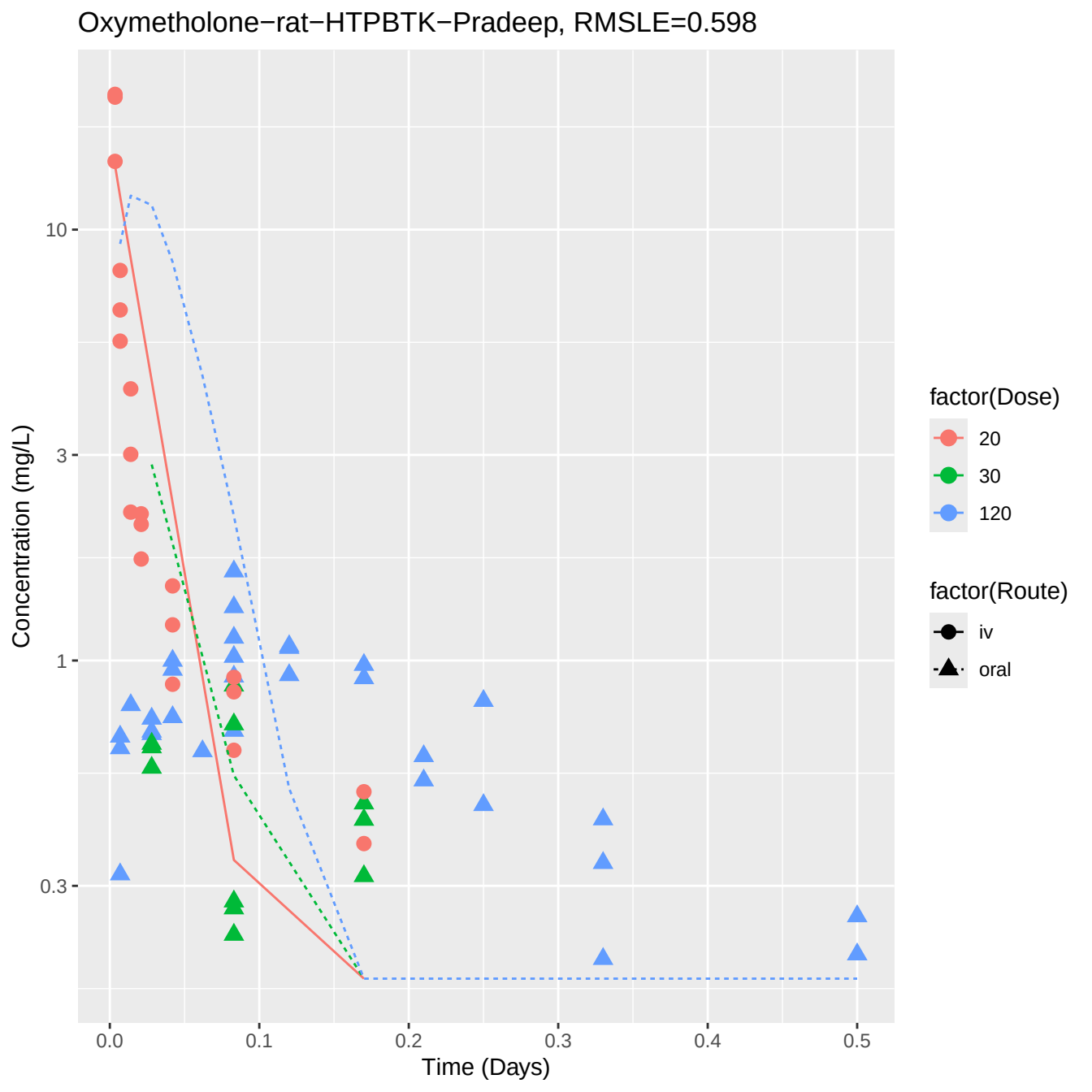

Etoxazole−rat−HTPBTK−Pradeep, RMSLE=0.452
Concentration (mg/L)
10^{-1}
10^{-2}
10^{-3}
10^{-4}
0
1
2
3
Time (Days)
factor(Dose)
0.4
2
factor(Route)
iv
oral

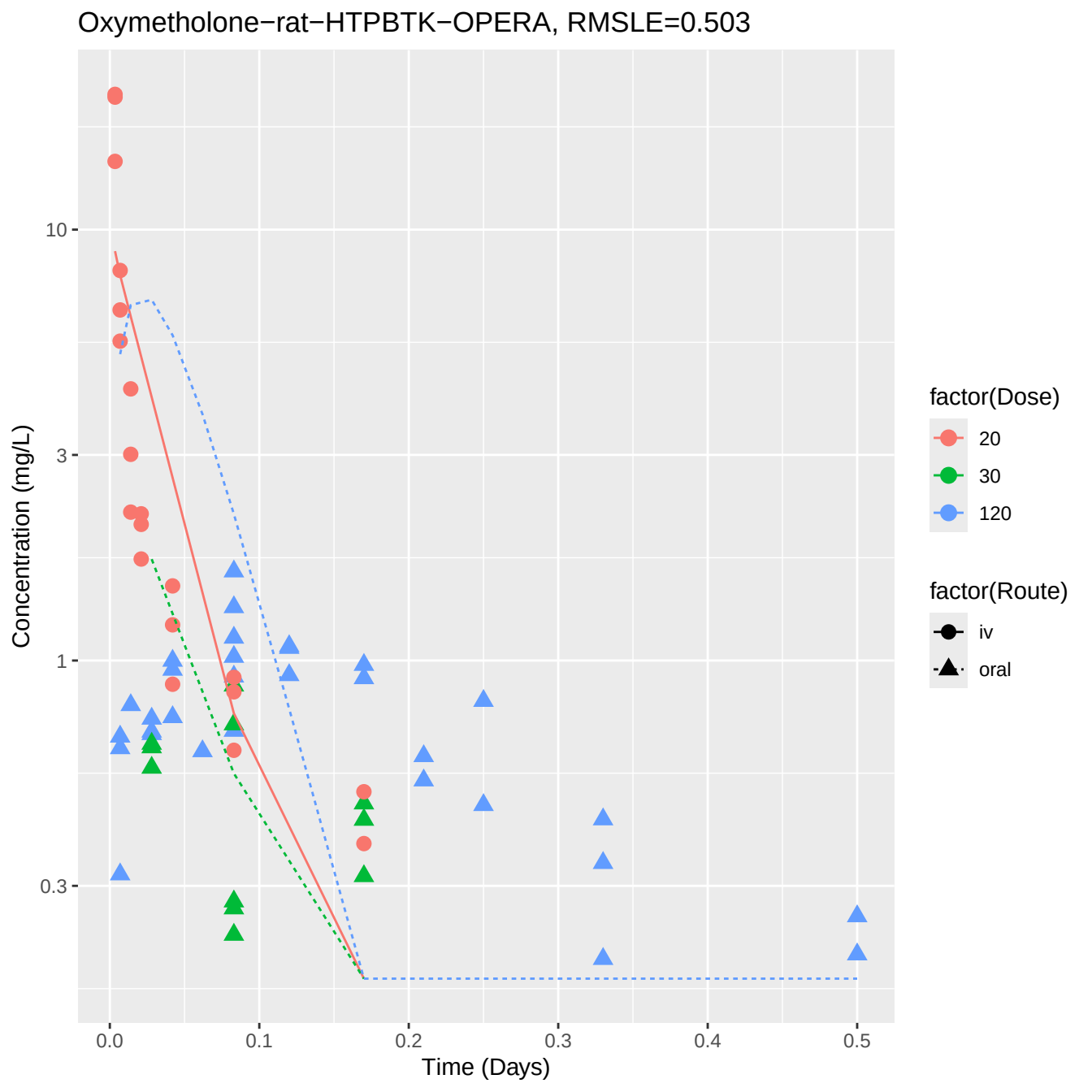

Etoxazole−rat−HTPBTK−Ensemble, RMSLE=0.507
Concentration (mg/L)
10^{-1}
10^{-2}
10^{-3}
10^{-4}
0
1
2
3
Time (Days)
factor(Dose)
0.4
2
factor(Route)
iv
oral

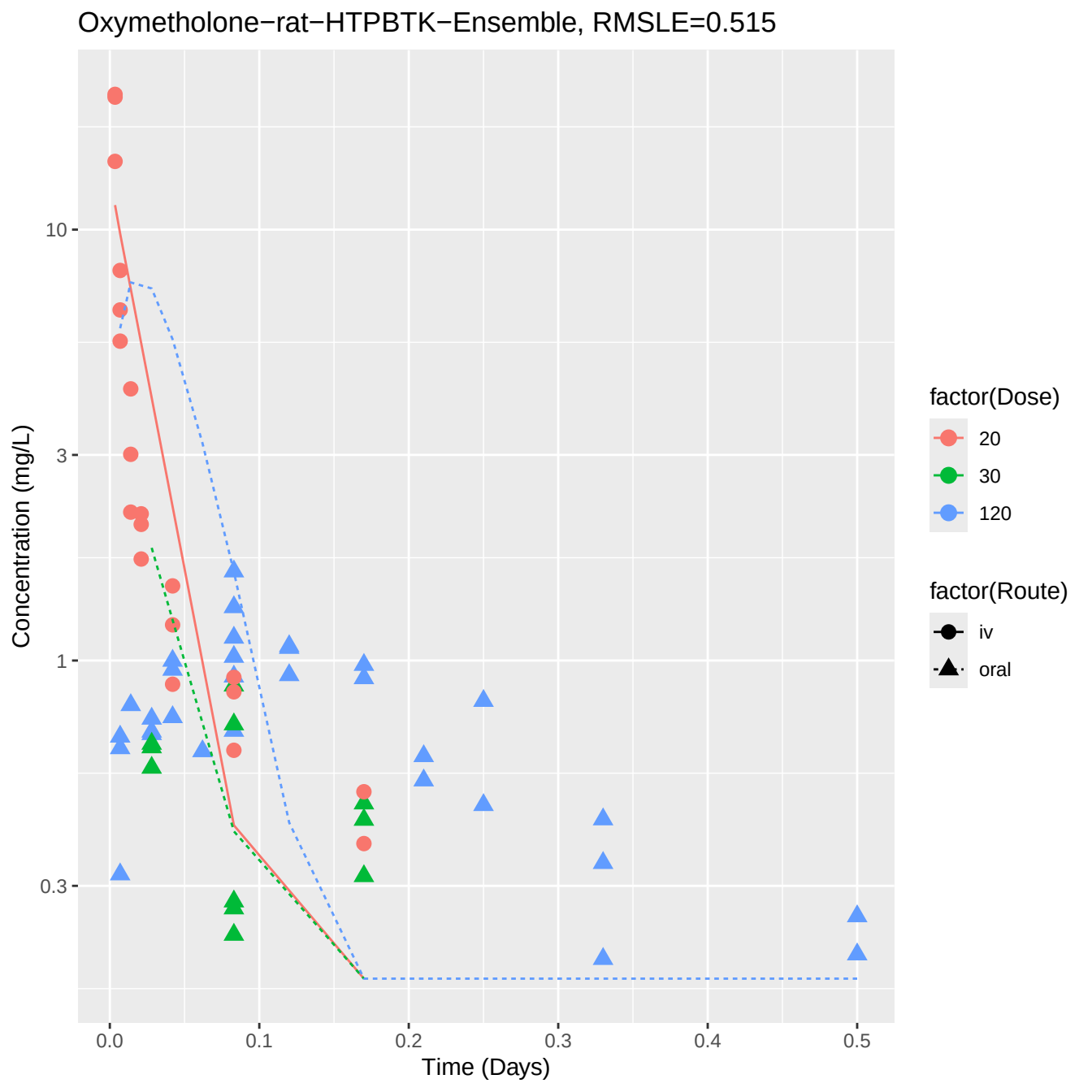

Etoxazole−rat−In Vivo Fits, RMSLE=0.177
Concentration (mg/L)
10^{-1}
10^{-2}
10^{-3}
10^{-4}
0
1
2
3
Time (Days)
factor(Dose)
0.4
2
factor(Route)
iv
oral

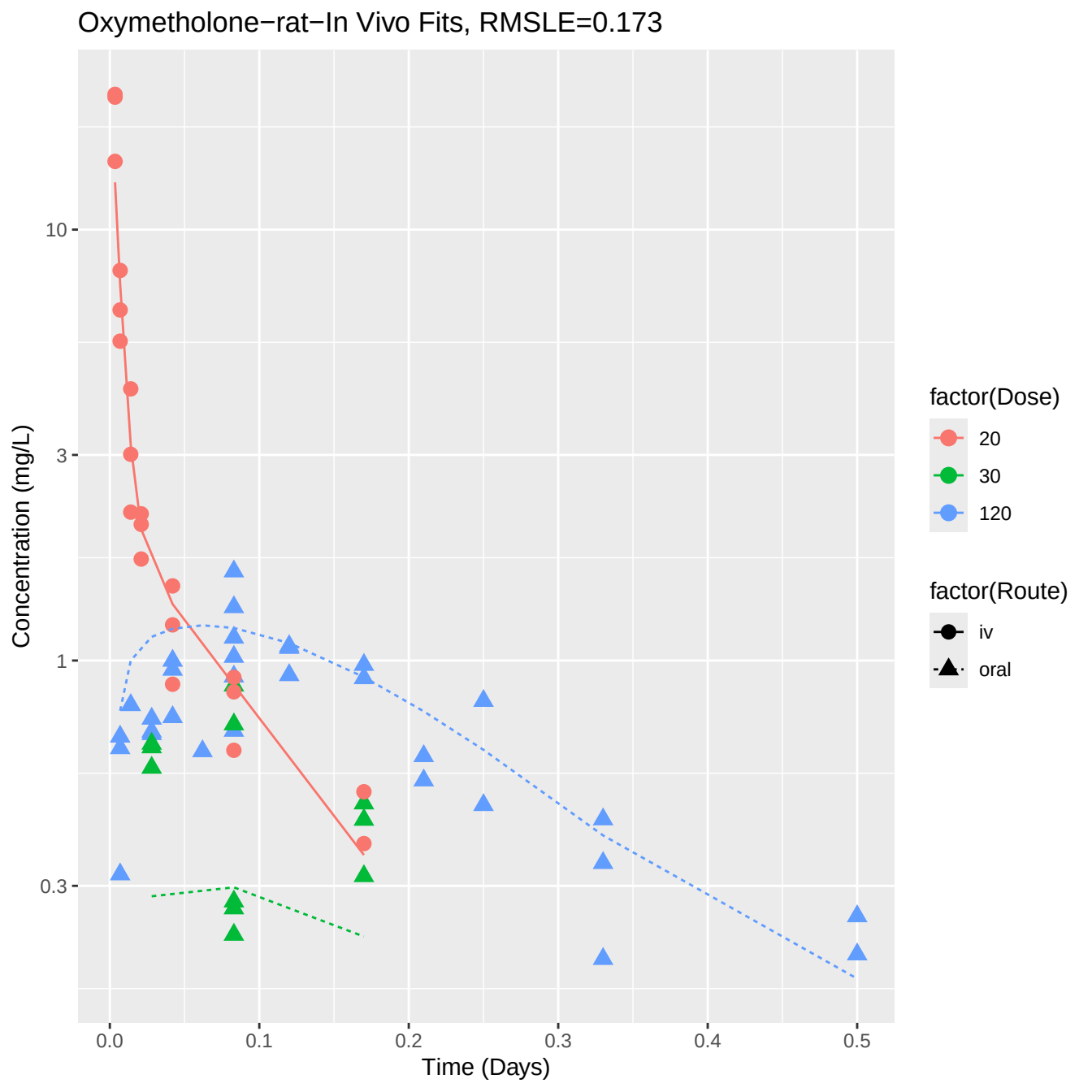

Oxymetholone−rat−HTPBTK−ADMET, RMSLE=0.952
Concentration (mg/L)
30
10
3
1
0.3
0.0
0.1
0.2
0.3
0.4
0.5
Time (Days)
factor(Dose)
20
30
120
factor(Route)
iv
oral

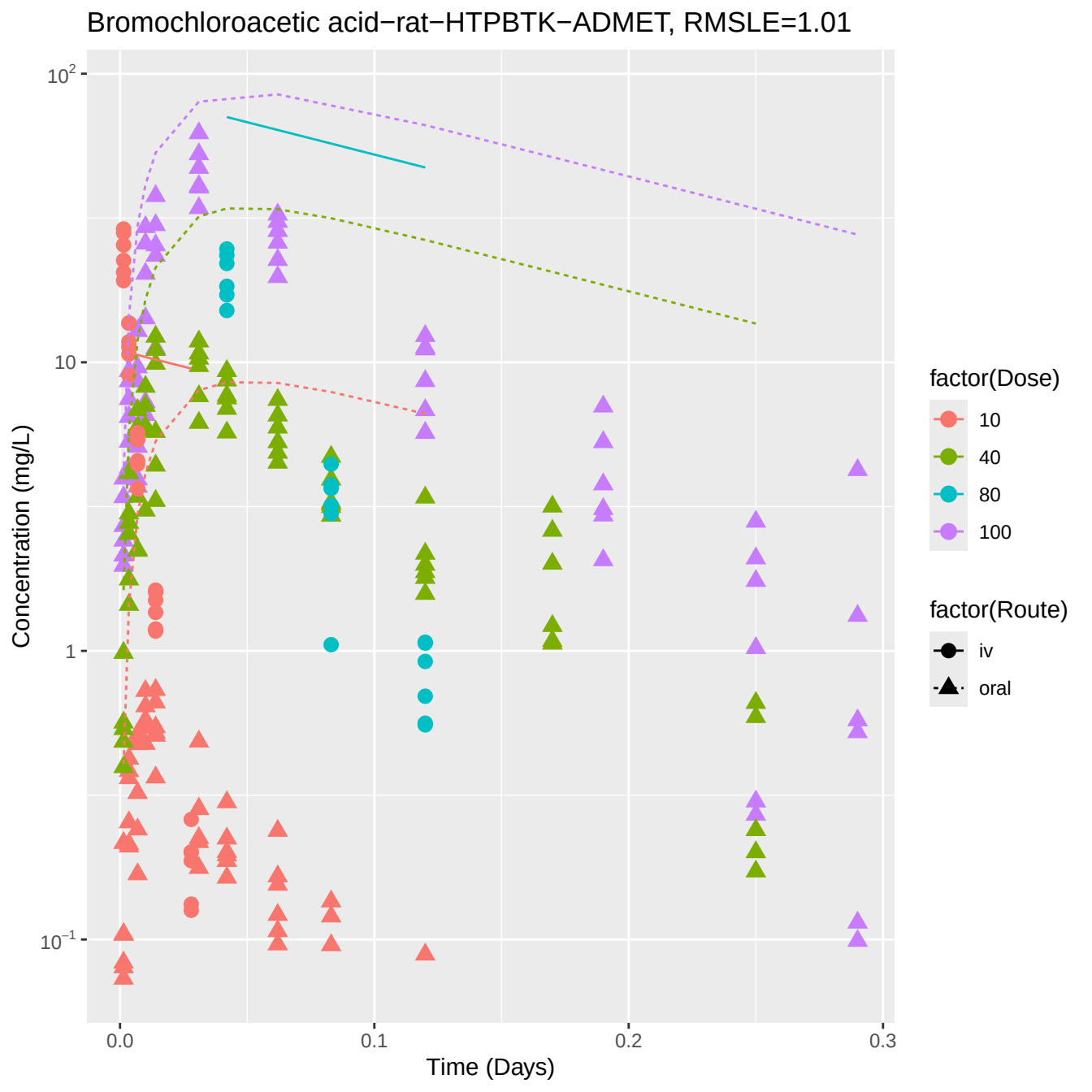

Oxymetholone−rat−HTPBTK−Dawson, RMSLE=0.529
Concentration (mg/L)
10
3
1
0.3
0.0
0.1
0.2
0.3
0.4
0.5
Time (Days)
factor(Dose)
20
30
120
factor(Route)
iv
oral

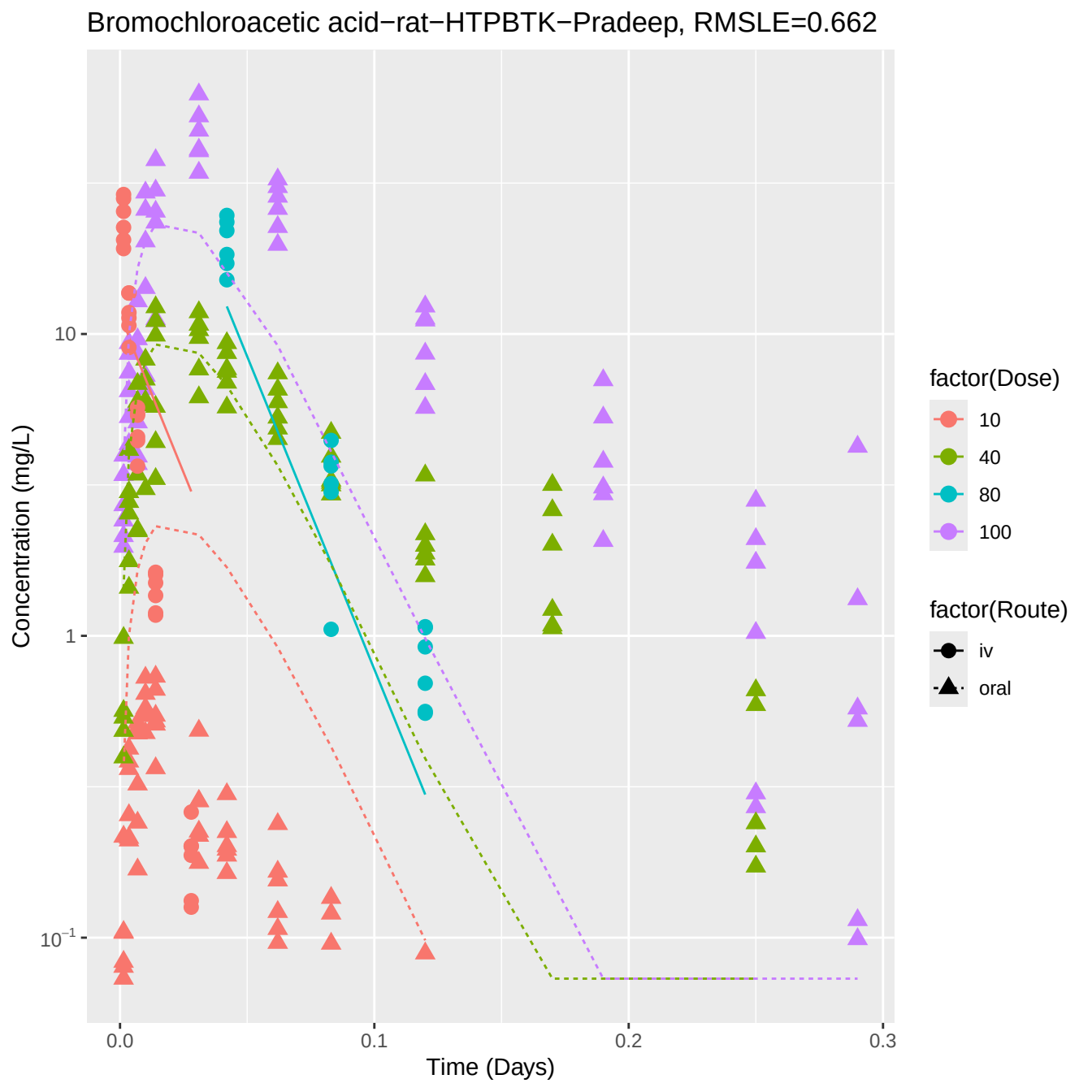

Oxymetholone−rat−HTPBTK−Pradeep, RMSLE=0.598
Concentration (mg/L)
10
3
1
0.3
0.0
0.1
0.2
0.3
0.4
0.5
Time (Days)
factor(Dose)
20
30
120
factor(Route)
iv
oral

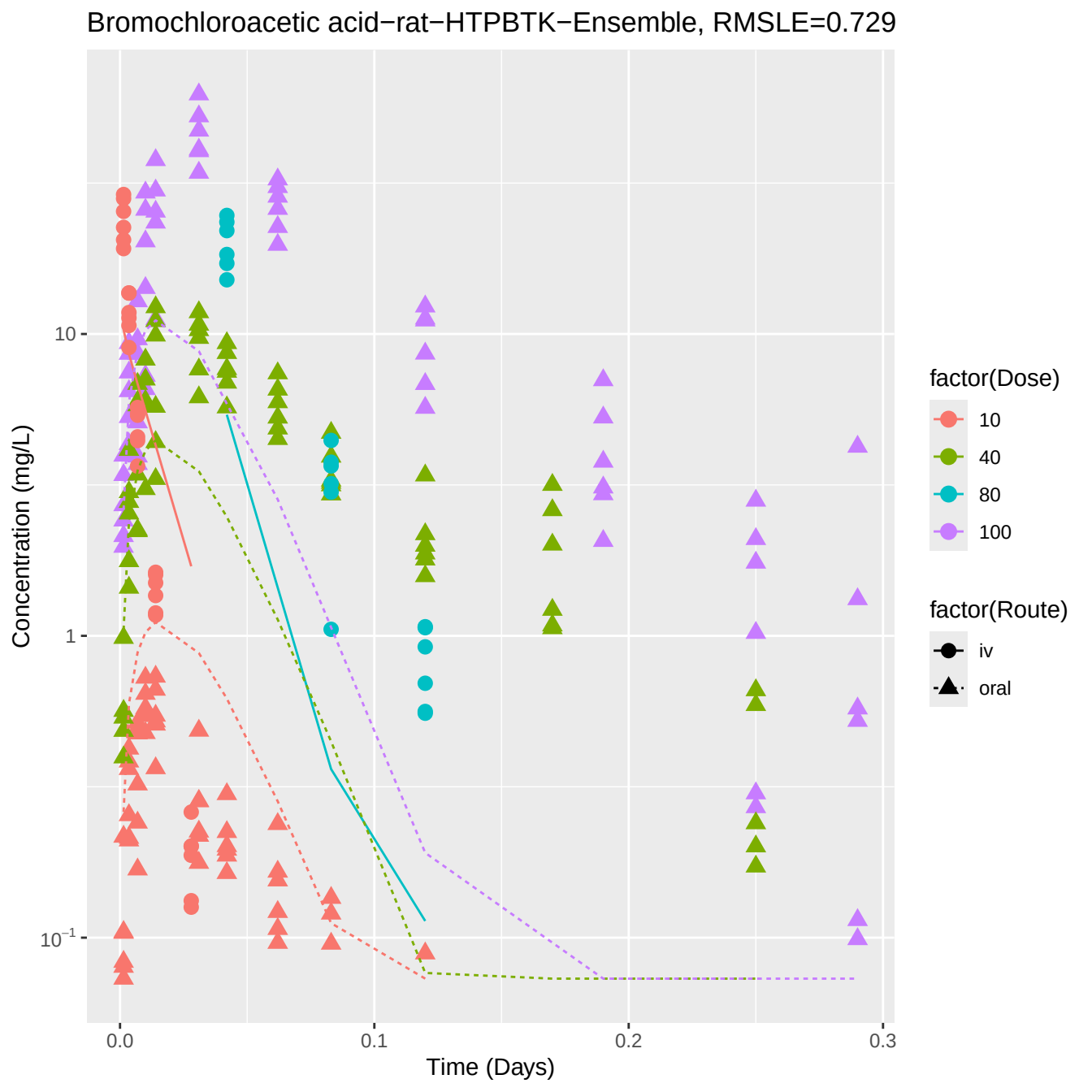

Oxymetholone−rat−HTPBTK−OPERA, RMSLE=0.503
Concentration (mg/L)
10
3
1
0.3
0.0
0.1
0.2
0.3
0.4
0.5
Time (Days)
factor(Dose)
20
30
120
factor(Route)
iv
oral

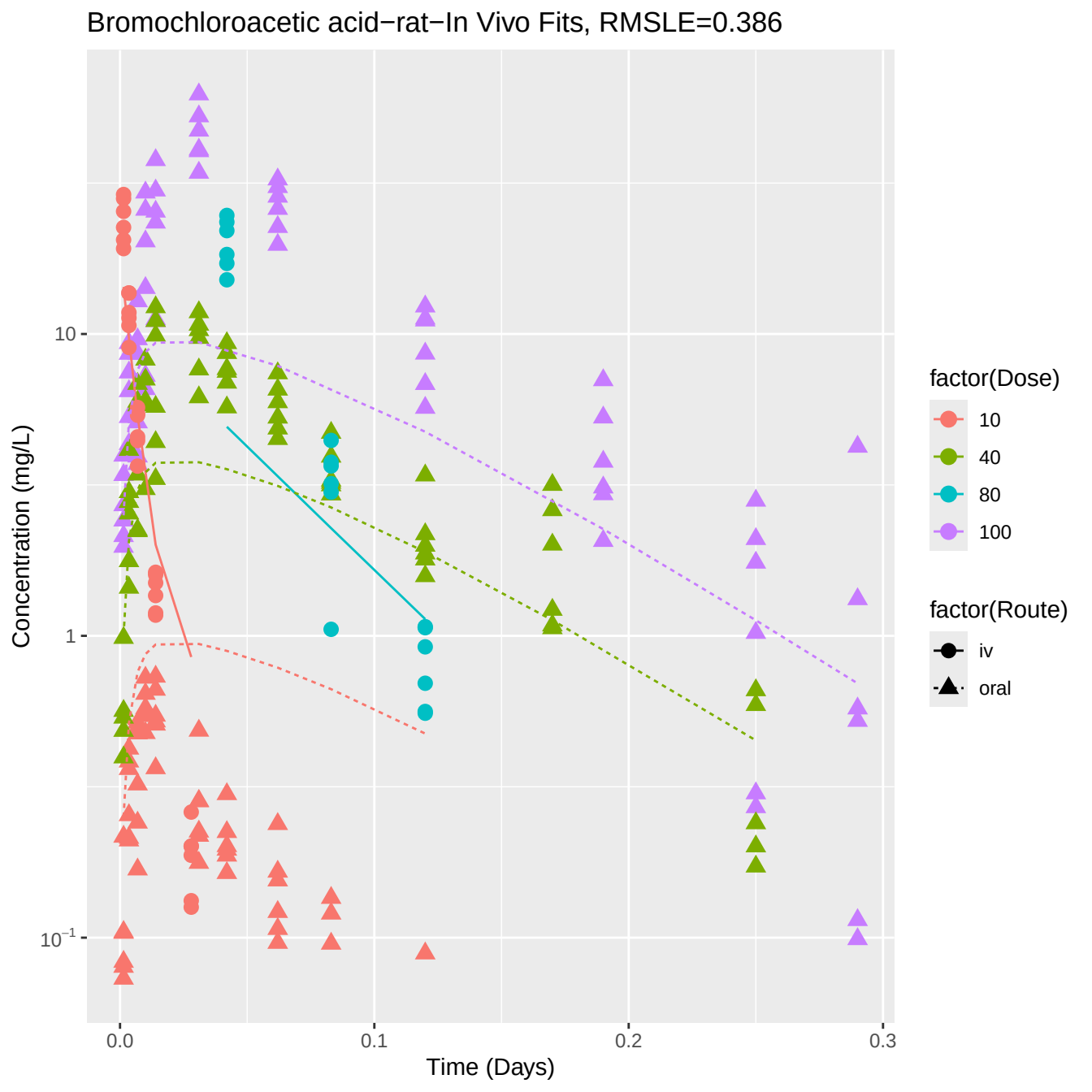

Oxymetholone−rat−HTPBTK−Ensemble, RMSLE=0.515
Concentration (mg/L)
10
3
1
0.3
0.0
0.1
0.2
0.3
0.4
0.5
Time (Days)
factor(Dose)
20
30
120
factor(Route)
iv
oral

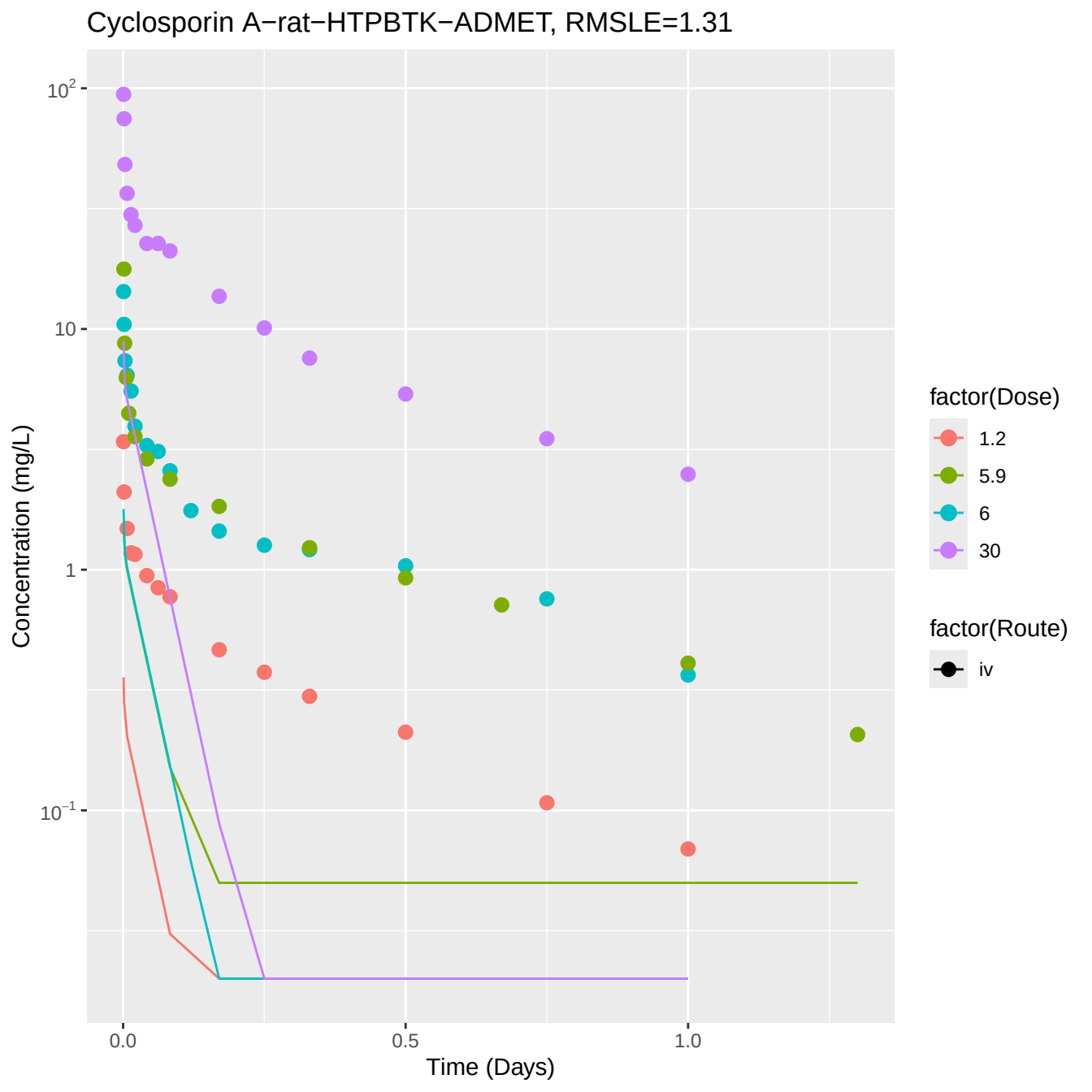

Oxymetholone−rat−In Vivo Fits, RMSLE=0.173
Concentration (mg/L)
10
3
1
0.3
0.0
0.1
0.2
0.3
0.4
0.5
Time (Days)
factor(Dose)
20
30
120
factor(Route)
iv
oral

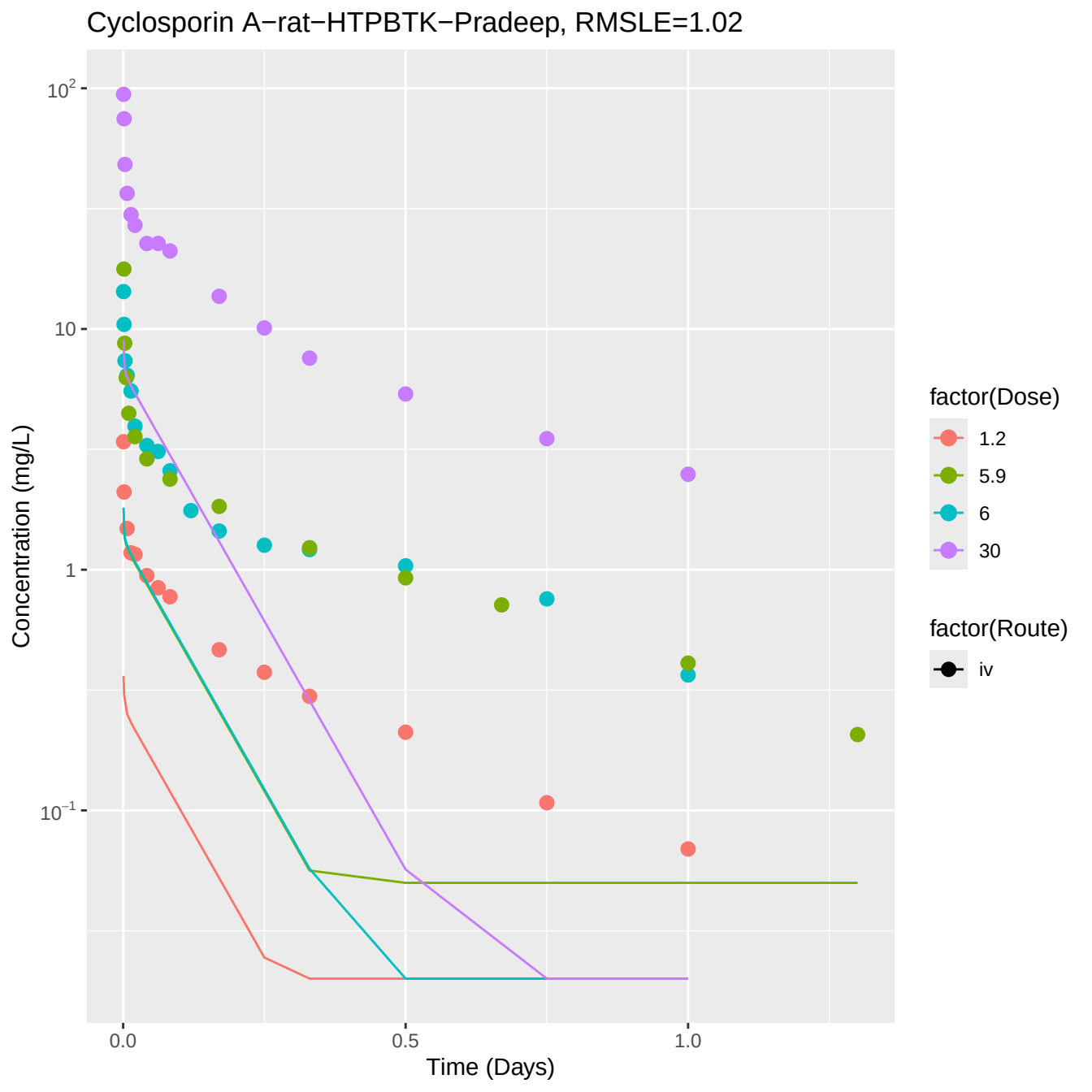

Bromochloroacetic acid−rat−HTPBTK−ADMET, RMSLE=1.01
Concentration (mg/L)
10^2
10
1
10^{-1}
0.0
0.1
0.2
0.3
Time (Days)
factor(Dose)
10
40
80
100
factor(Route)
iv
oral

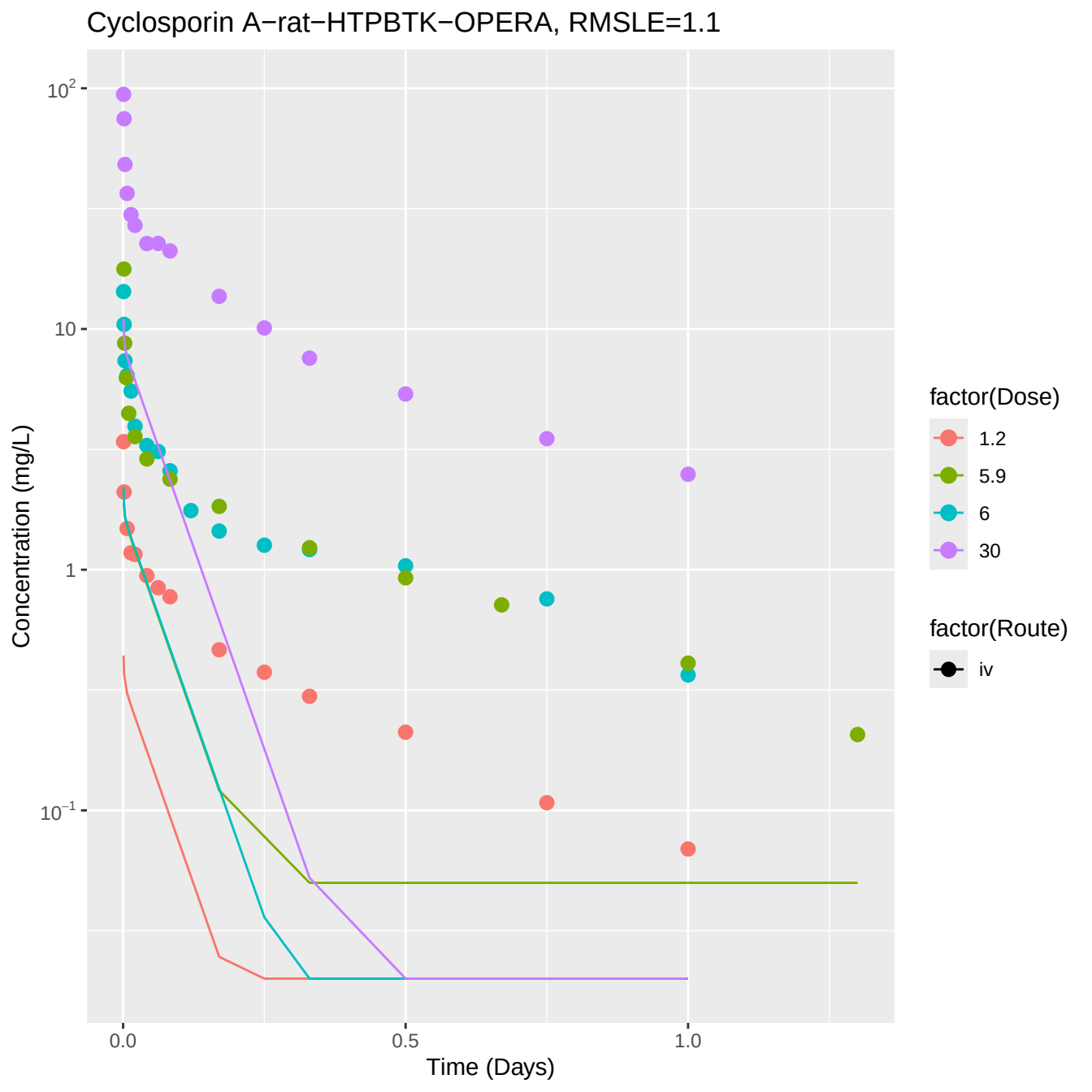

Bromochloroacetic acid−rat−HTPBTK−Pradeep, RMSLE=0.662
Concentration (mg/L)
10
1
10^{-1}
0.0
0.1
0.2
0.3
Time (Days)
factor(Dose)
10
40
80
100
factor(Route)
iv
oral

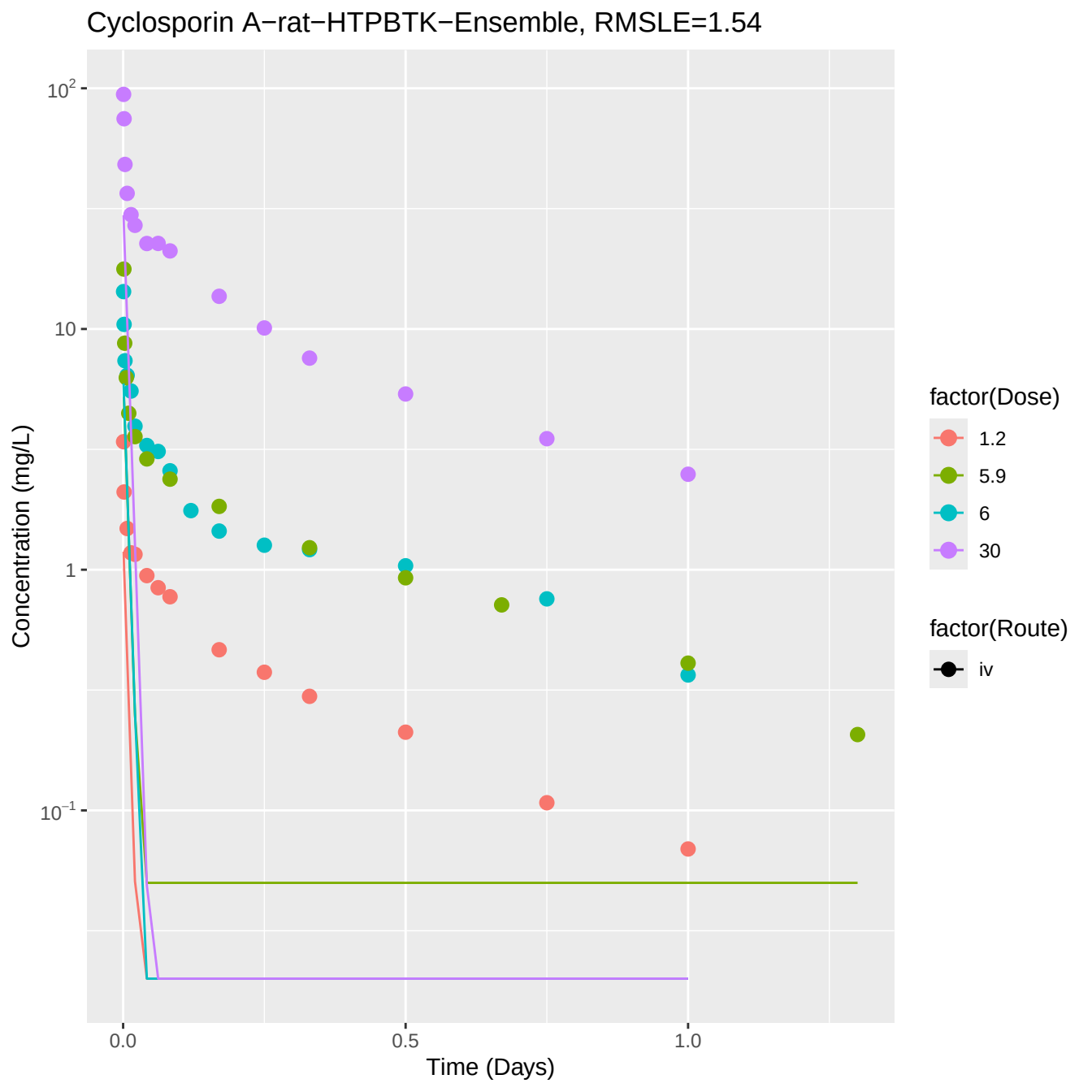

Bromochloroacetic acid−rat−HTPBTK−Ensemble, RMSLE=0.729
Concentration (mg/L)
10
1
10^{-1}
0.0
0.1
0.2
0.3
Time (Days)
factor(Dose)
10
40
80
100
factor(Route)
iv
oral

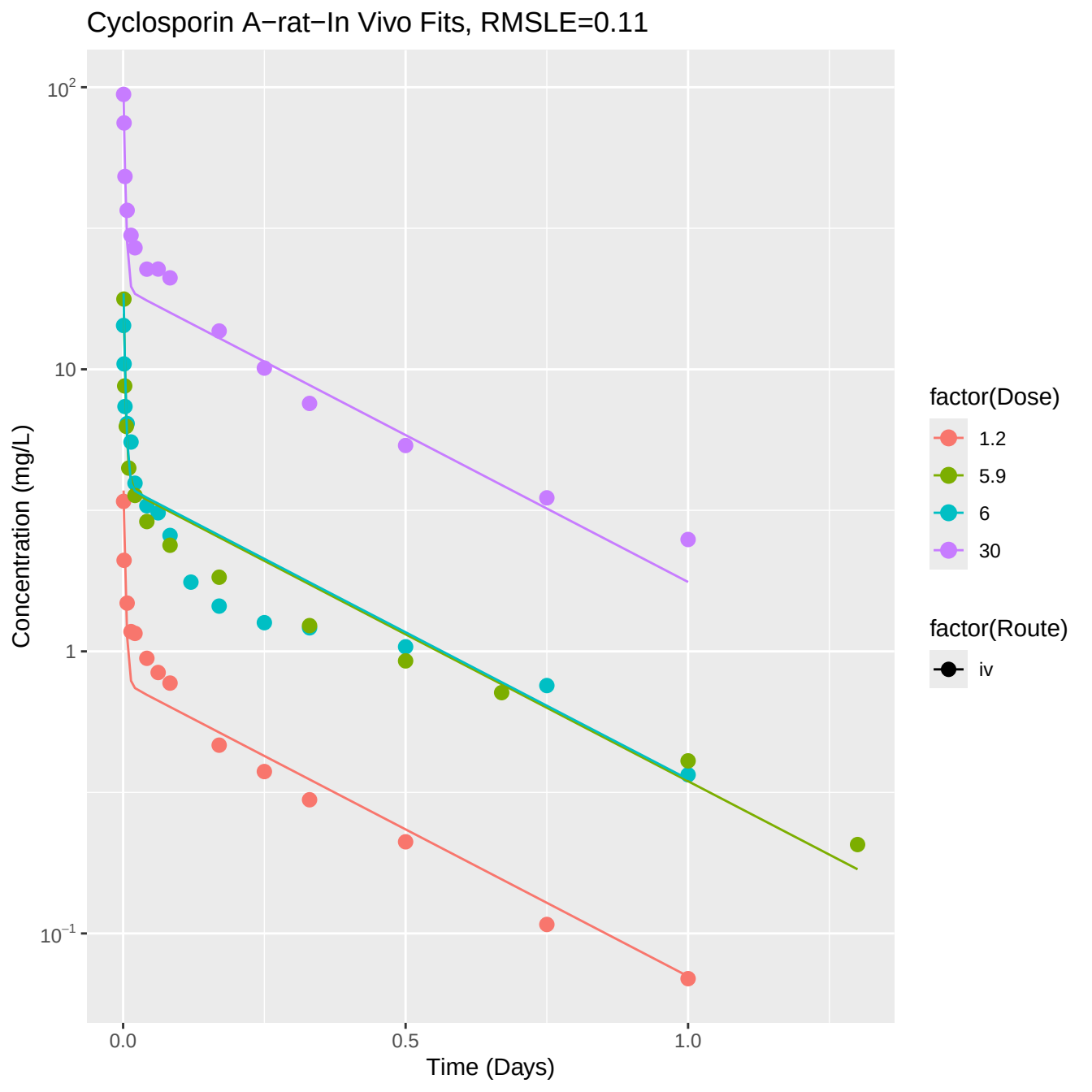

Bromochloroacetic acid−rat−In Vivo Fits, RMSLE=0.386
Concentration (mg/L)
10
1
10^{-1}
0.0
0.1
0.2
0.3
Time (Days)
factor(Dose)
10
40
80
100
factor(Route)
iv
oral

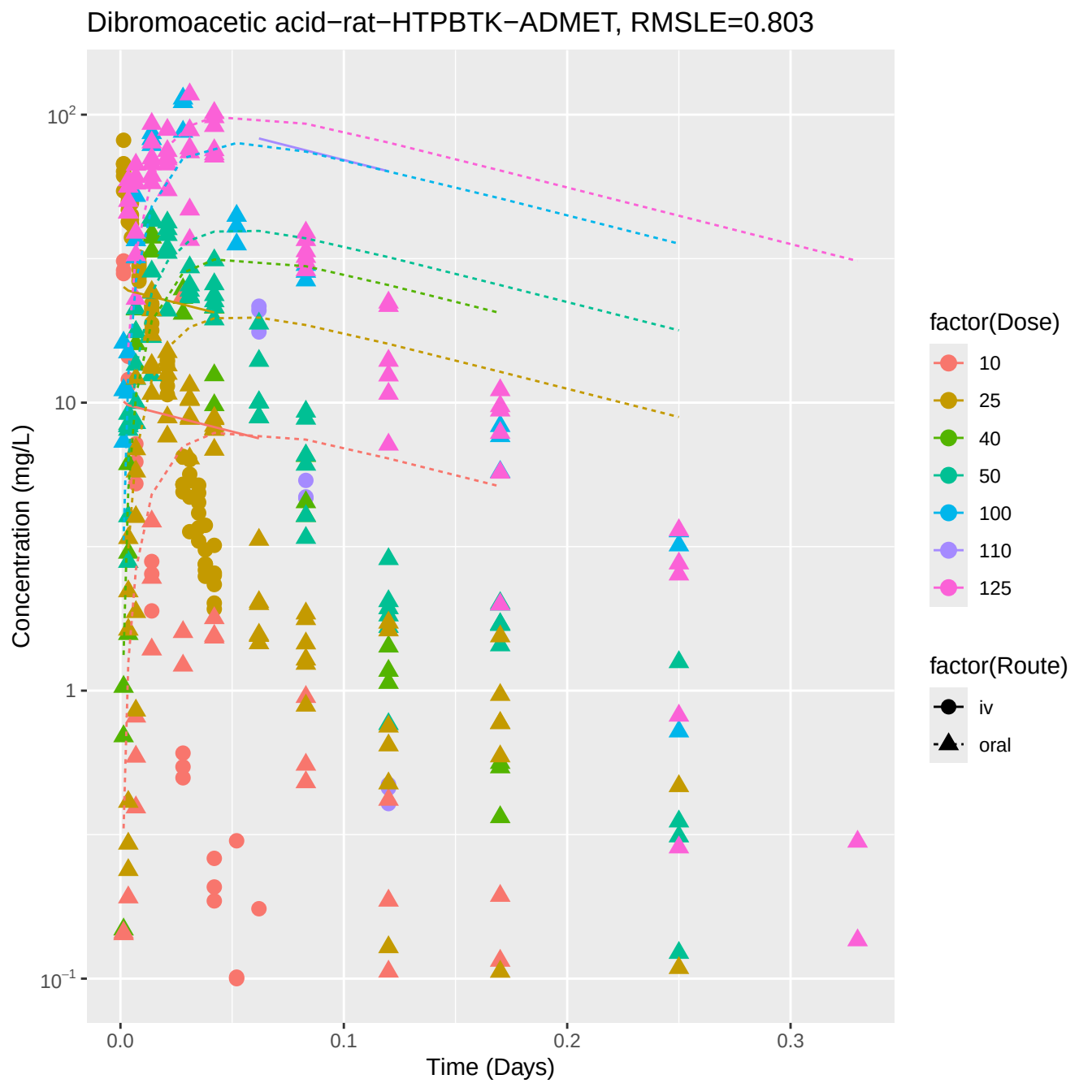

Cyclosporin A−rat−HTPBTK−ADMET, RMSLE=1.31
Concentration (mg/L)
10^2
10
1
10^{-1}
0.0
0.5
1.0
Time (Days)
factor(Dose)
1.2
5.9
6
30
factor(Route)
iv

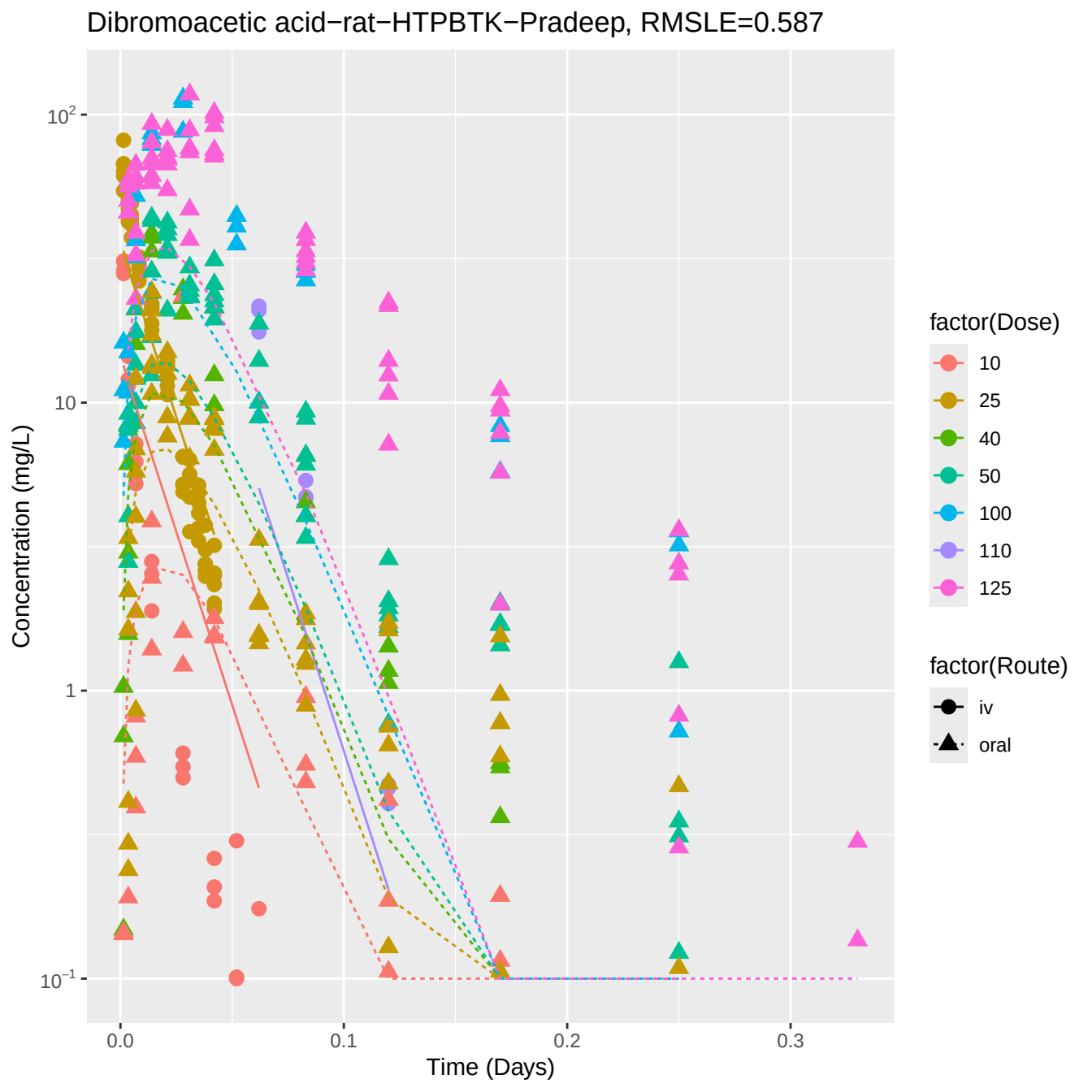

Cyclosporin A−rat−HTPBTK−Pradeep, RMSLE=1.02
Concentration (mg/L)
10^2
10
1
10^{-1}
0.0
0.5
1.0
Time (Days)
factor(Dose)
1.2
5.9
6
30
factor(Route)
iv

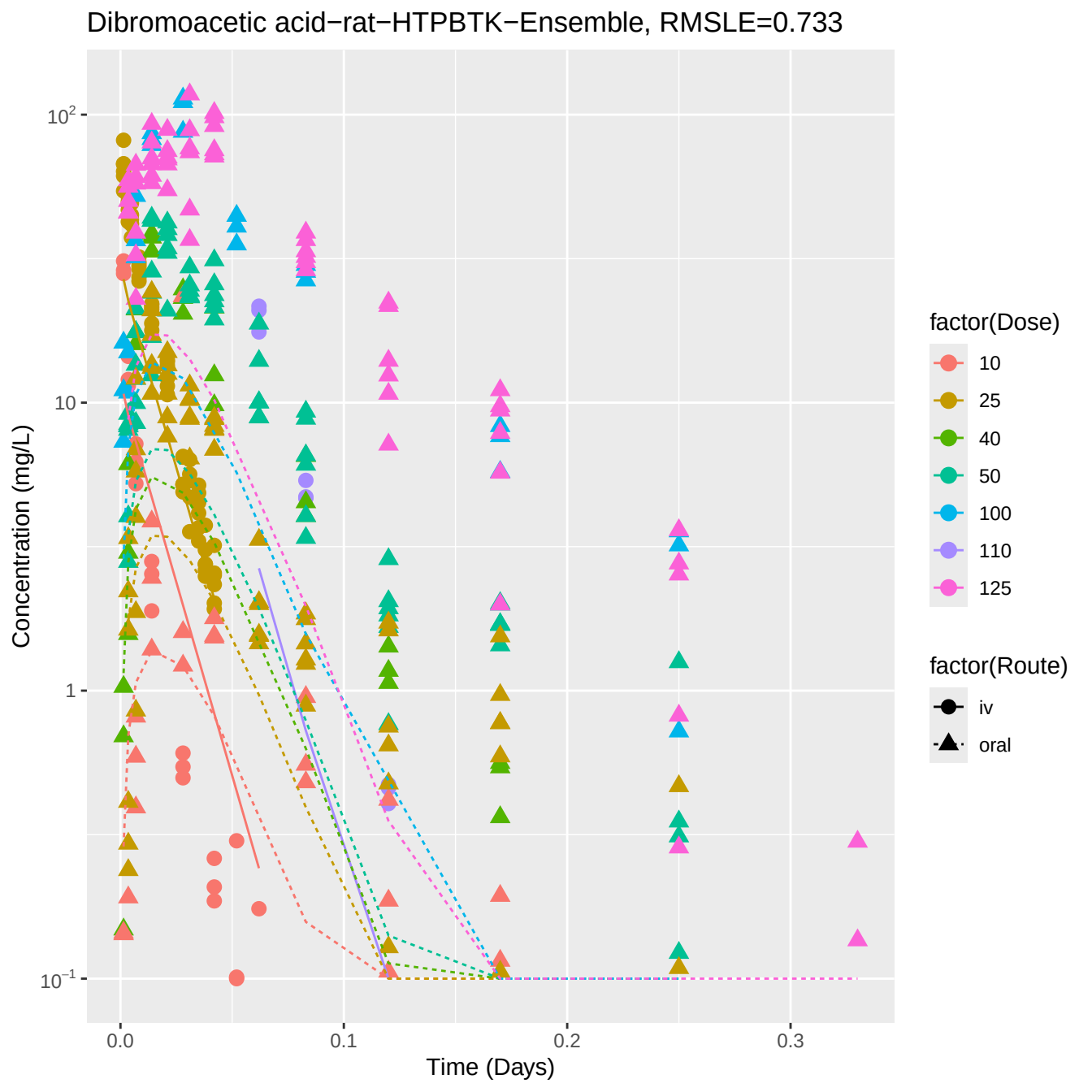

Cyclosporin A−rat−HTPBTK−OPERA, RMSLE=1.1
Concentration (mg/L)
10^2
10
1
10^{-1}
0.0
0.5
1.0
Time (Days)
factor(Dose)
1.2
5.9
6
30
factor(Route)
iv

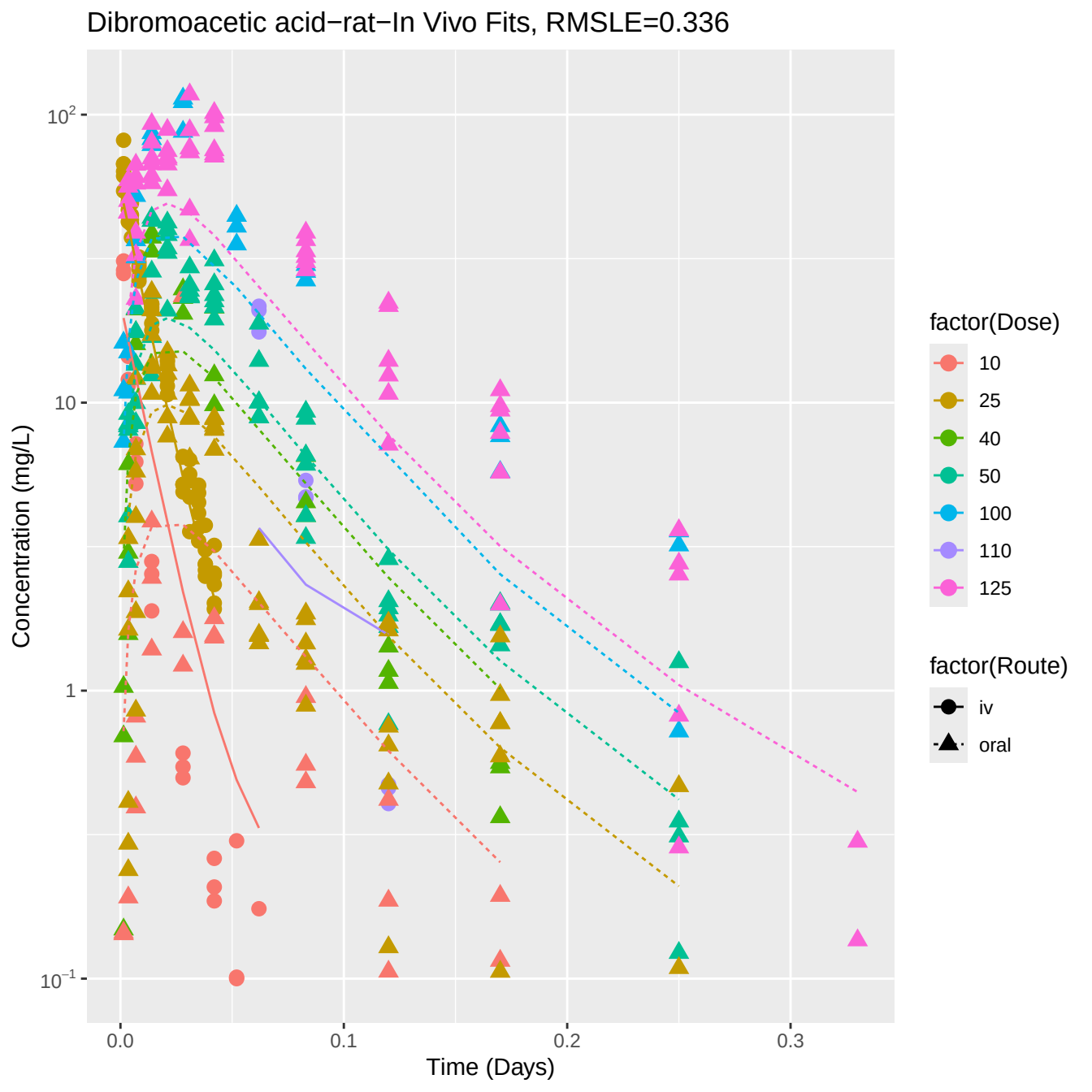

Cyclosporin A−rat−HTPBTK−Ensemble, RMSLE=1.54
Concentration (mg/L)
10^2
10
1
10^{-1}
0.0
0.5
1.0
Time (Days)
factor(Dose)
1.2
5.9
6
30
factor(Route)
iv

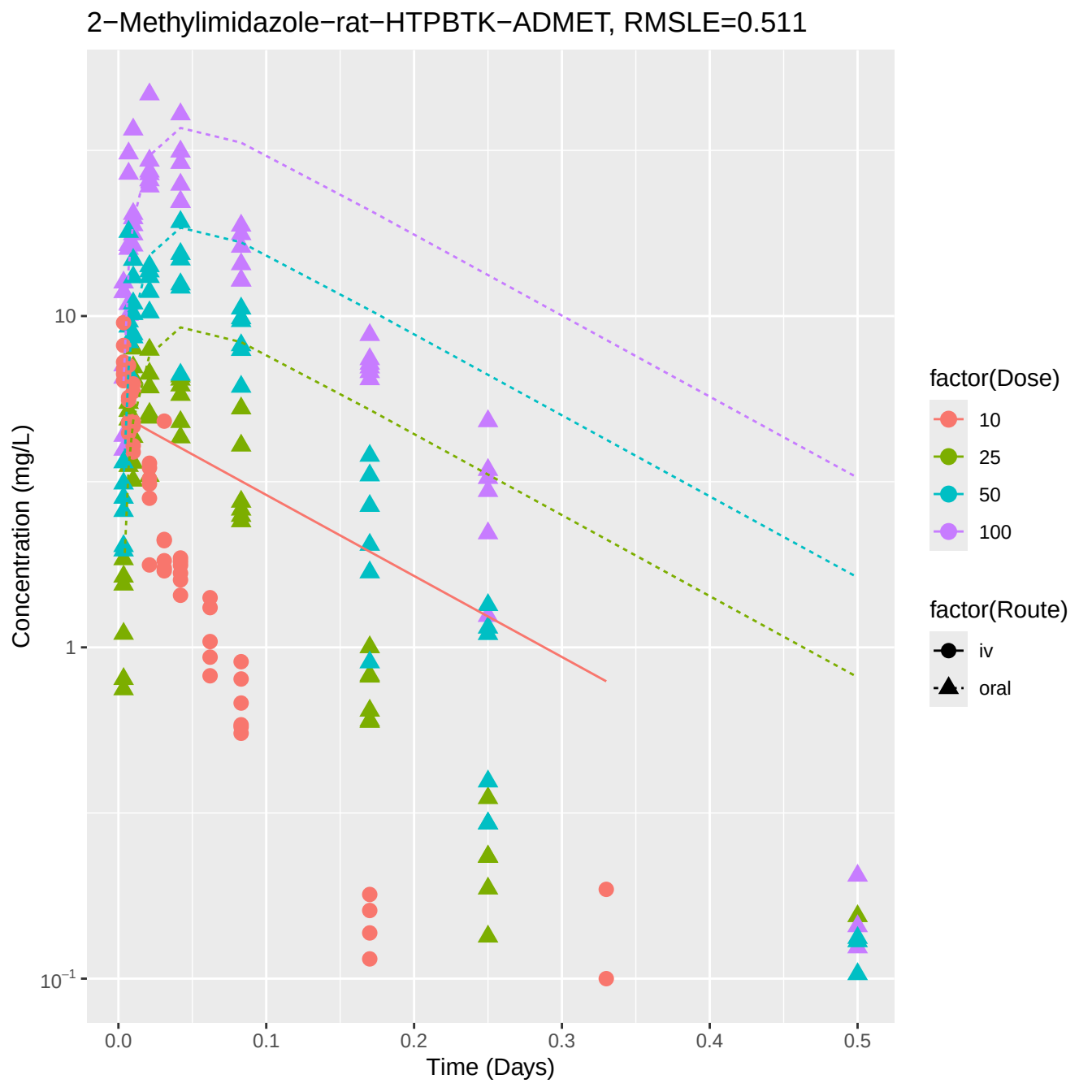

Cyclosporin A−rat−In Vivo Fits, RMSLE=0.11
Concentration (mg/L)
10^2
10
1
10^{-1}
0.0
0.5
1.0
Time (Days)
factor(Dose)
1.2
5.9
6
30
factor(Route)
iv

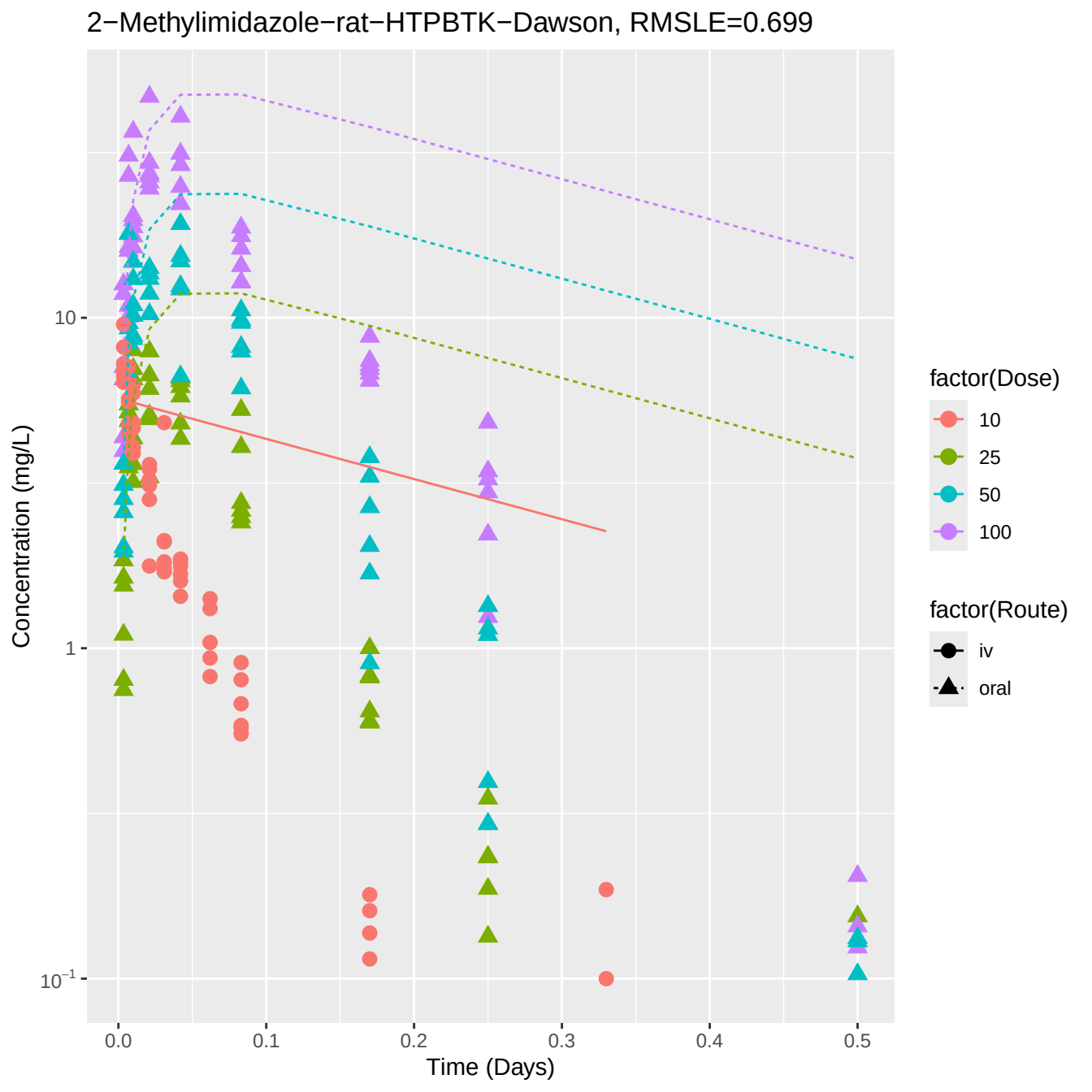

Dibromoacetic acid−rat−HTPBTK−ADMET, RMSLE=0.803
Concentration (mg/L)
10^2
10
1
10^{-1}
0.0
0.1
0.2
0.3
Time (Days)
factor(Dose)
10
25
40
50
100
110
125
factor(Route)
iv
oral

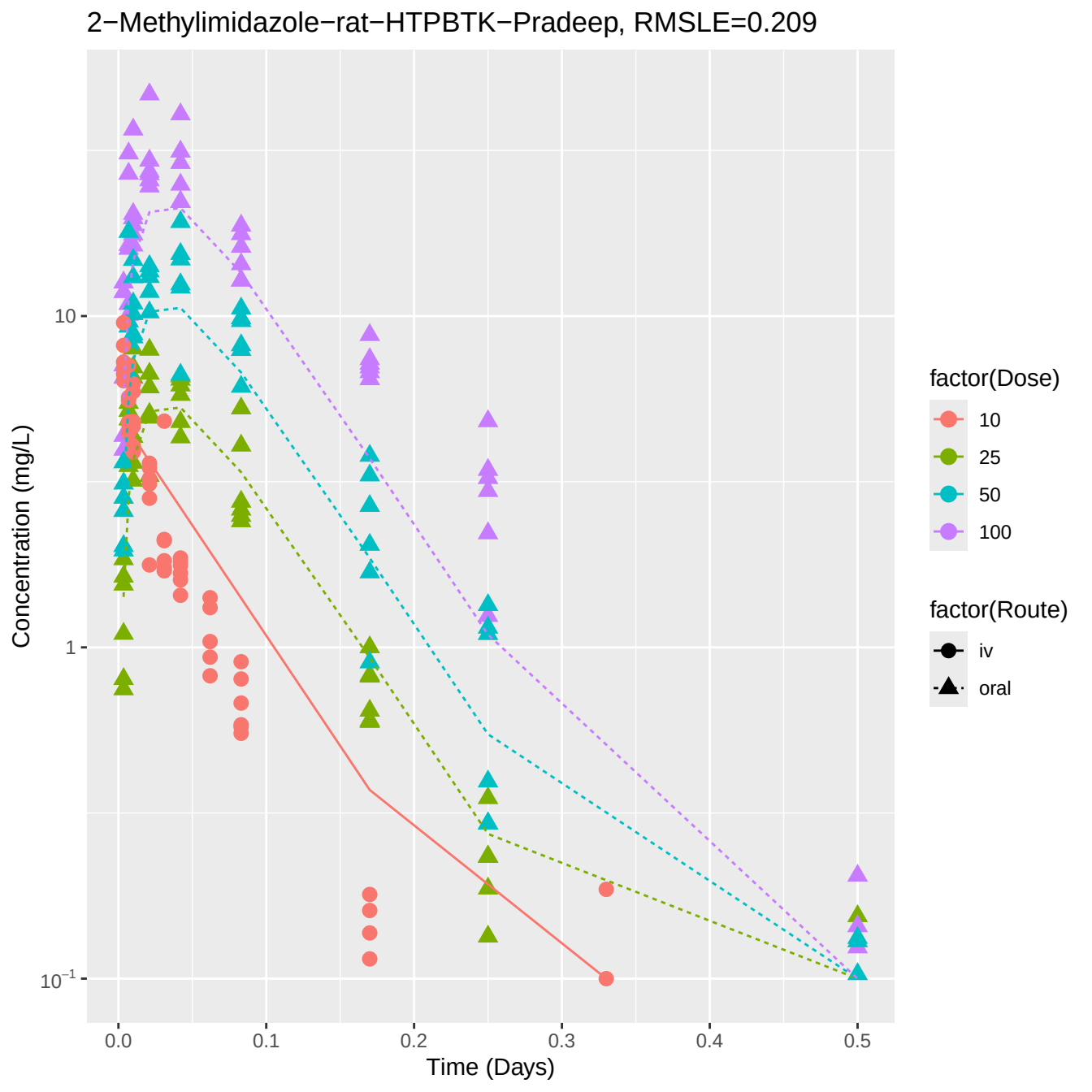

Dibromoacetic acid−rat−HTPBTK−Pradeep, RMSLE=0.587
Concentration (mg/L)
10^2
10
1
10^{-1}
0.0
0.1
0.2
0.3
Time (Days)
factor(Dose)
10
25
40
50
100
110
125
factor(Route)
iv
oral

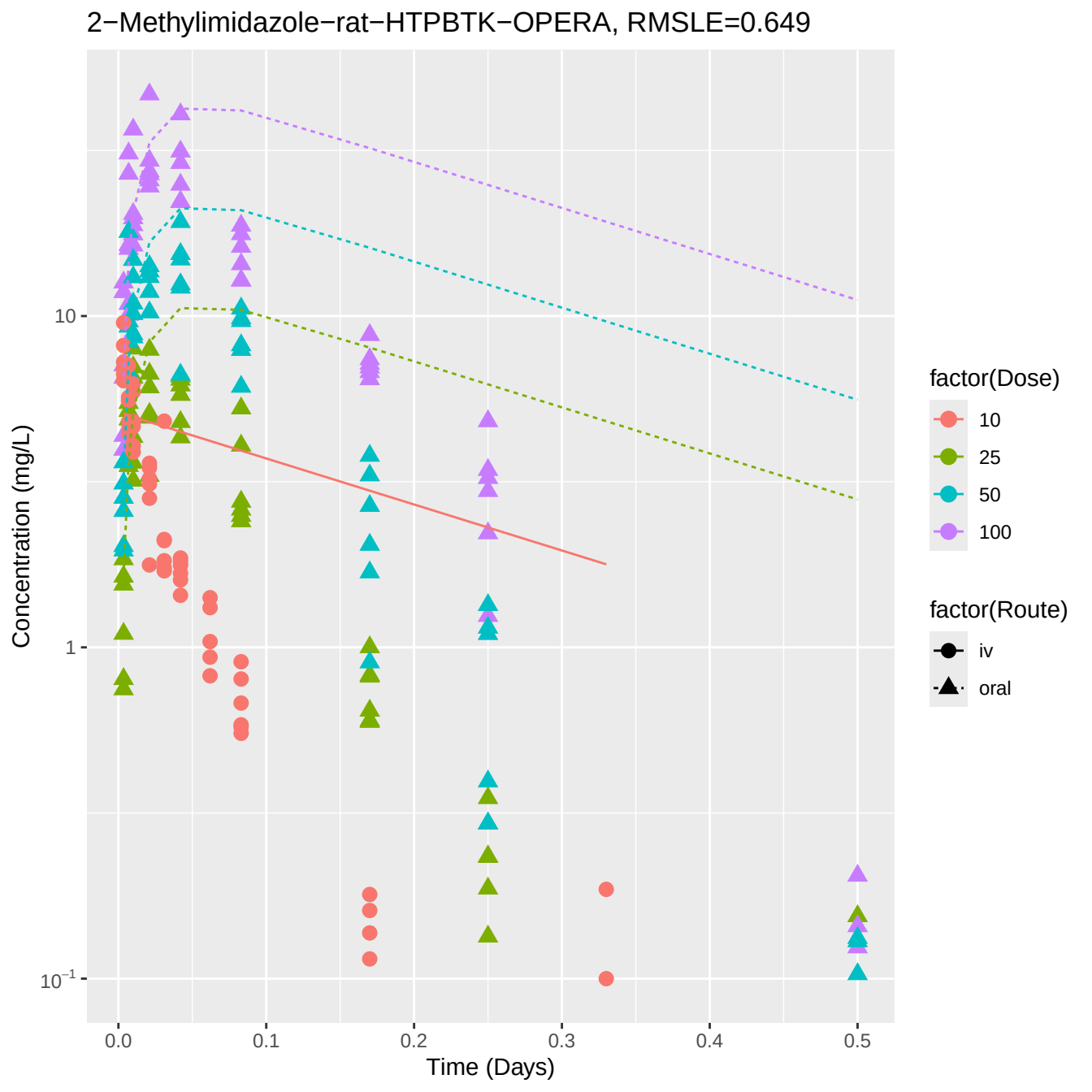

Dibromoacetic acid−rat−HTPBTK−Ensemble, RMSLE=0.733
Concentration (mg/L)
10^2
10
1
10^{-1}
0.0
0.1
0.2
0.3
Time (Days)
factor(Dose)
10
25
40
50
100
110
125
factor(Route)
iv
oral

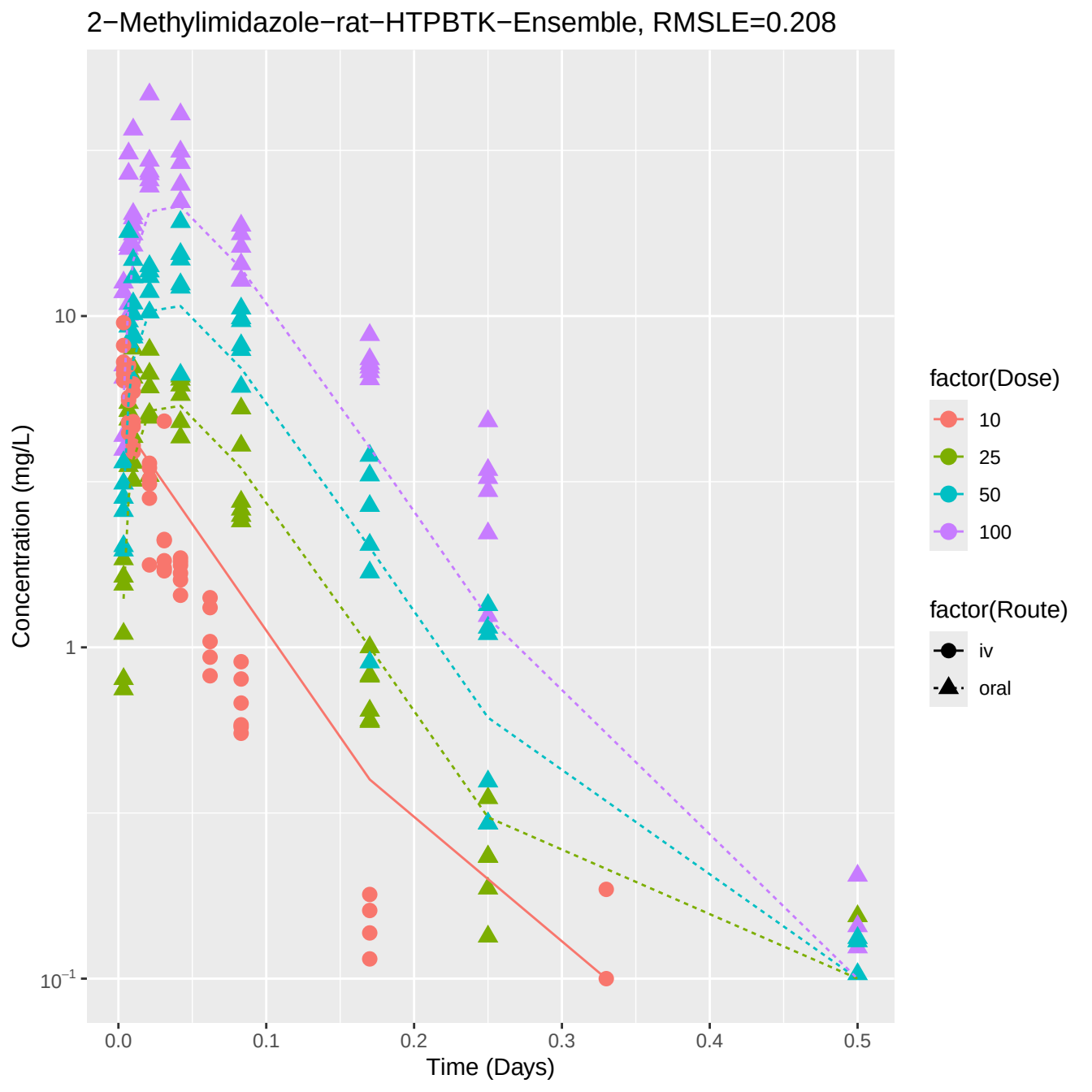

Dibromoacetic acid−rat−In Vivo Fits, RMSLE=0.336
Concentration (mg/L)
10^2
10
1
10^{-1}
0.0
0.1
0.2
0.3
Time (Days)
factor(Dose)
10
25
40
50
100
110
125
factor(Route)
iv
oral

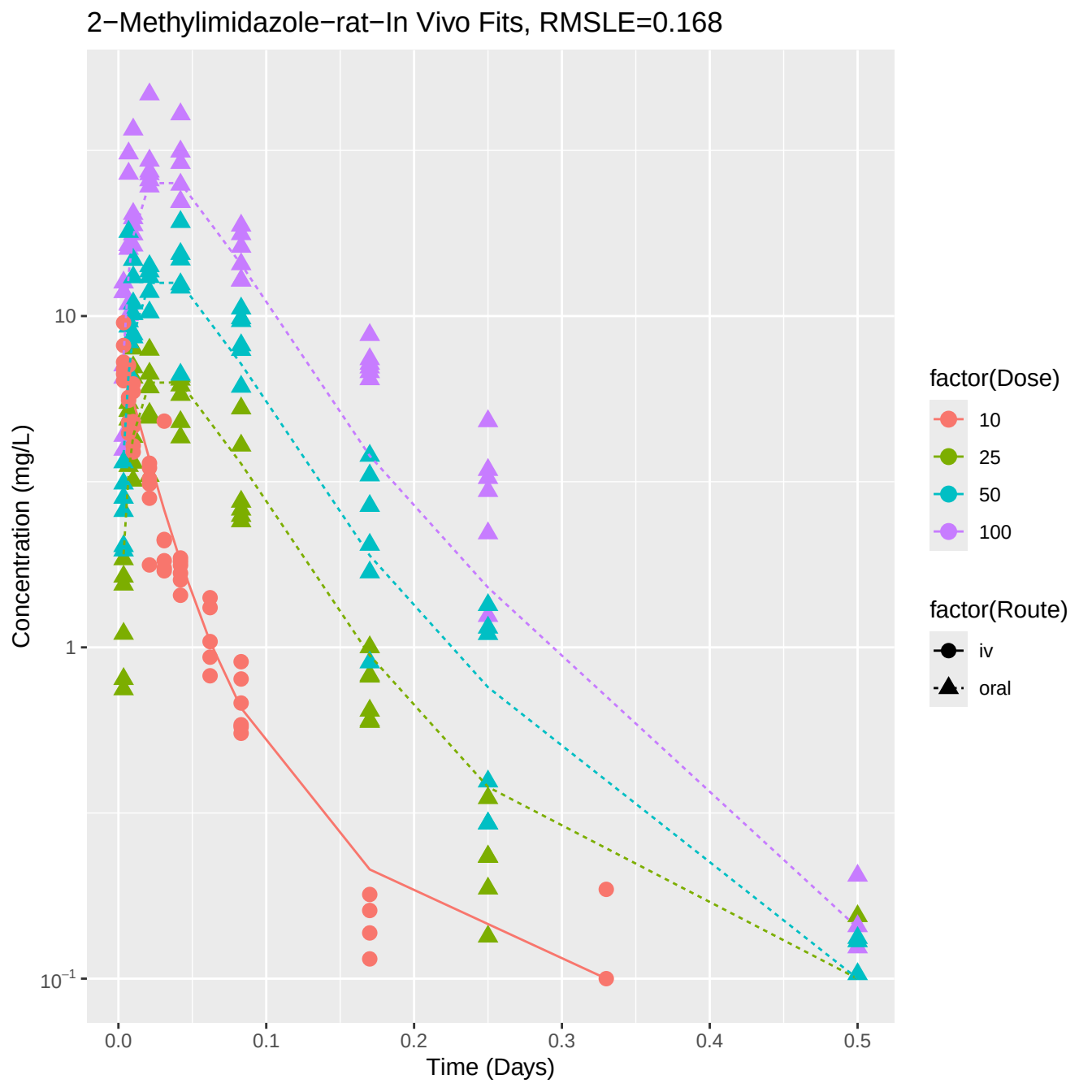

2−Methylimidazole−rat−HTPBTK−ADMET, RMSLE=0.511
Concentration (mg/L)
10
1
10^{-1}
0.0
0.1
0.2
0.3
0.4
0.5
Time (Days)
factor(Dose)
10
25
50
100
factor(Route)
iv
oral

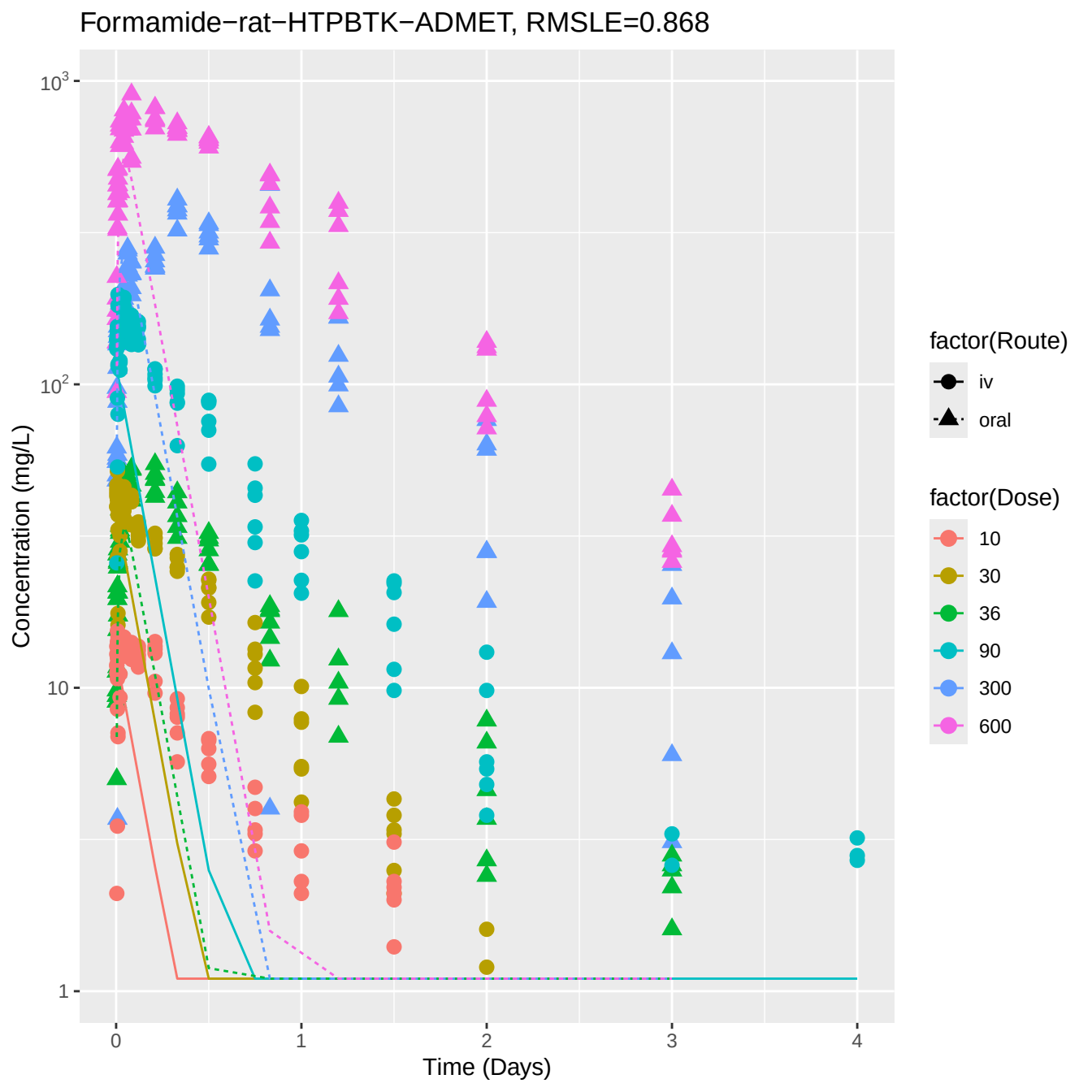

2−Methylimidazole−rat−HTPBTK−Dawson, RMSLE=0.699
Concentration (mg/L)
10
1
10^{-1}
0.0
0.1
0.2
0.3
0.4
0.5
Time (Days)
factor(Dose)
10
25
50
100
factor(Route)
iv
oral

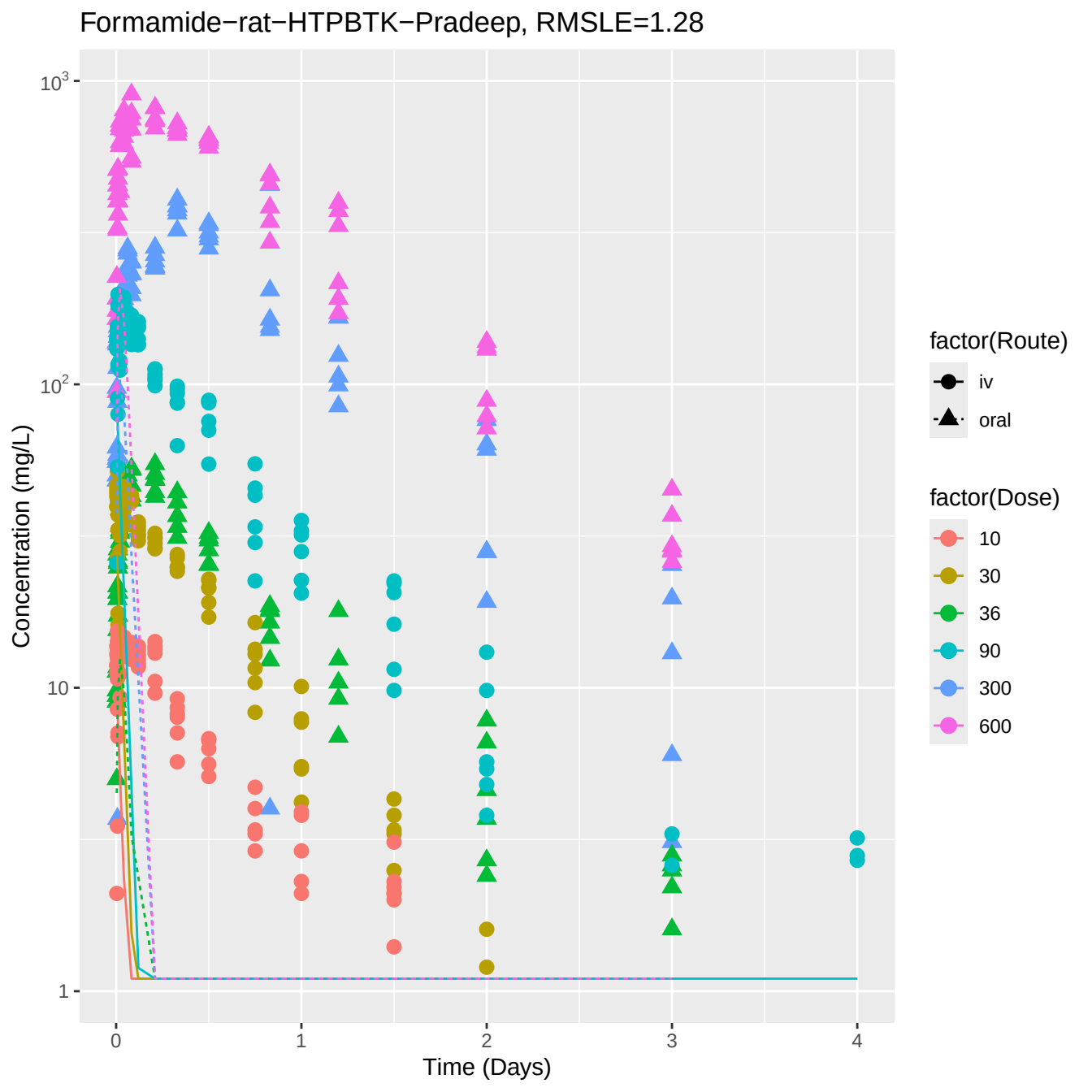

2−Methylimidazole−rat−HTPBTK−Pradeep, RMSLE=0.209
Concentration (mg/L)
10
1
10^{-1}
0.0
0.1
0.2
0.3
0.4
0.5
Time (Days)
factor(Dose)
10
25
50
100
factor(Route)
iv
oral

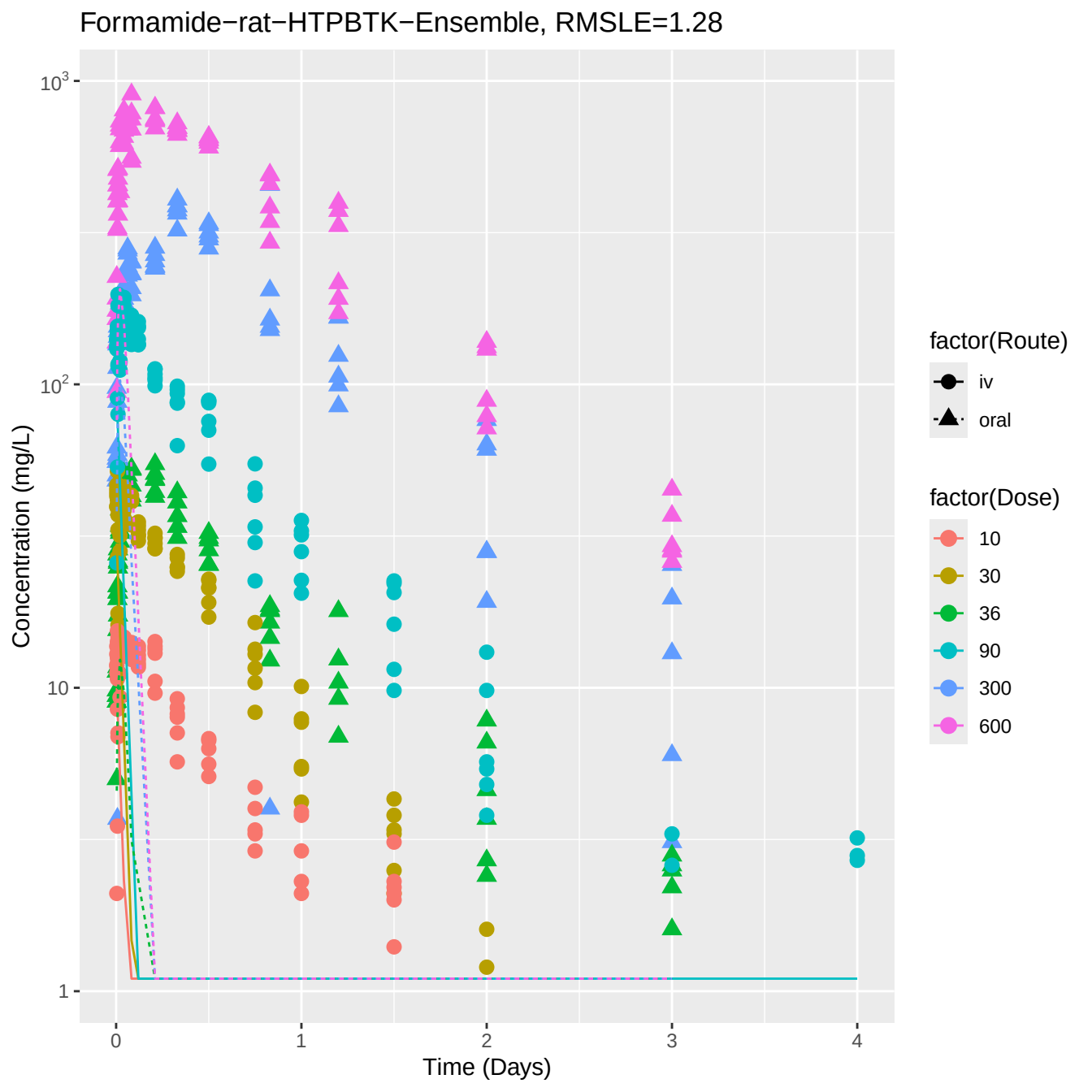

2−Methylimidazole−rat−HTPBTK−OPERA, RMSLE=0.649
Concentration (mg/L)
10
1
10^{-1}
0.0
0.1
0.2
0.3
0.4
0.5
Time (Days)
factor(Dose)
10
25
50
100
factor(Route)
iv
oral

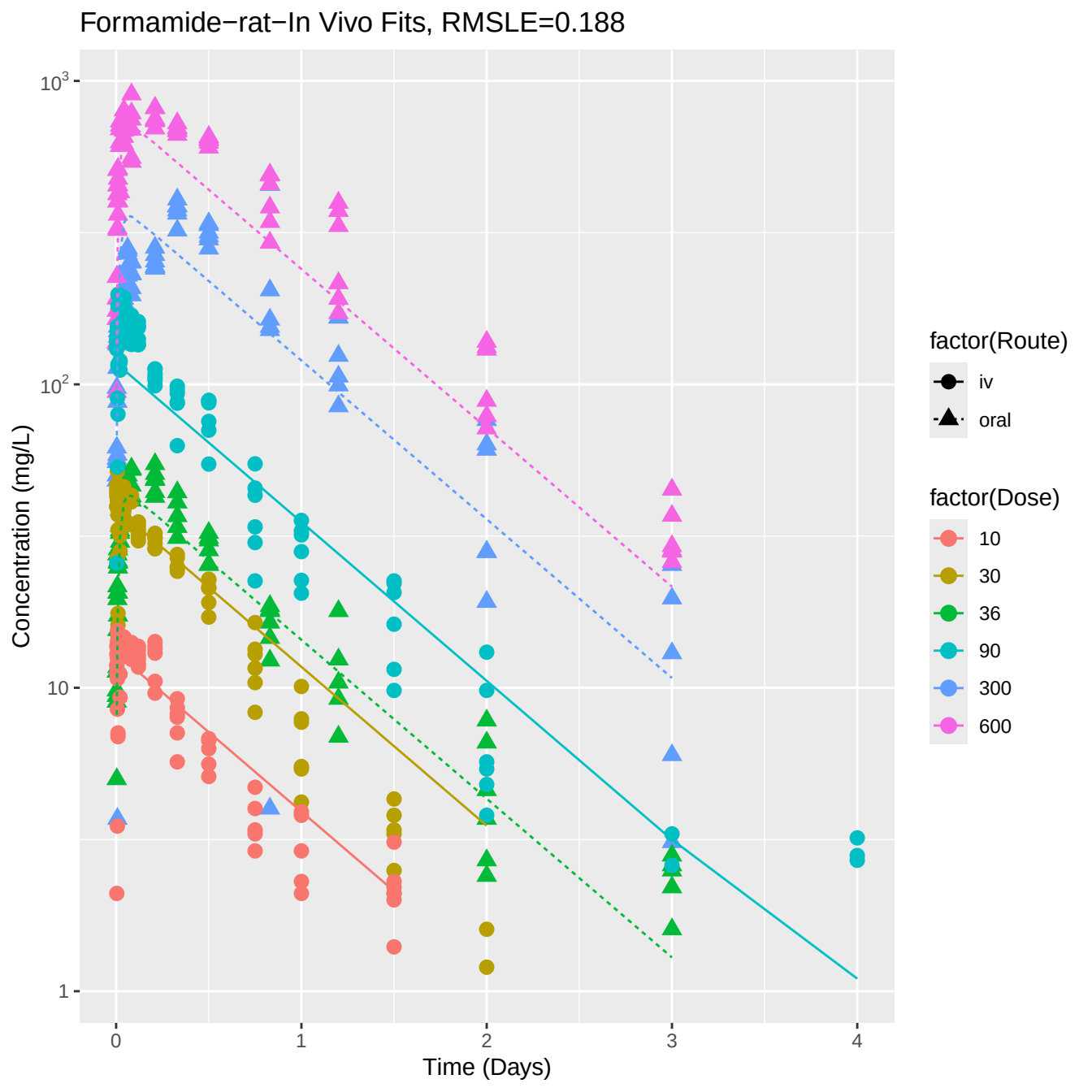

2−Methylimidazole−rat−HTPBTK−Ensemble, RMSLE=0.208
Concentration (mg/L)
10
1
10^{-1}
0.0
0.1
0.2
0.3
0.4
0.5
Time (Days)
factor(Dose)
10
25
50
100
factor(Route)
iv
oral

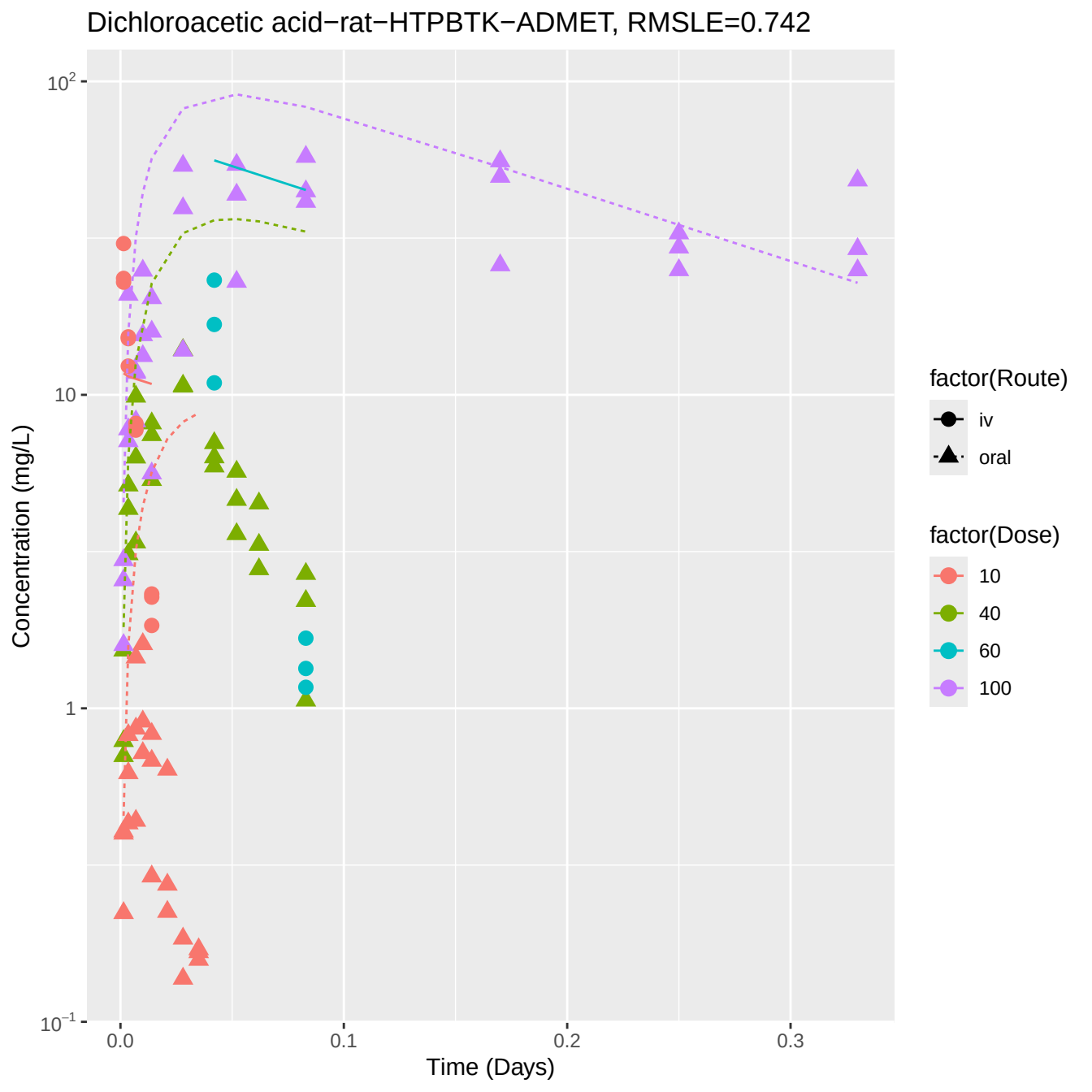

2−Methylimidazole−rat−In Vivo Fits, RMSLE=0.168
Concentration (mg/L)
10
1
10^{-1}
0.0
0.1
0.2
0.3
0.4
0.5
Time (Days)
factor(Dose)
10
25
50
100
factor(Route)
iv
oral

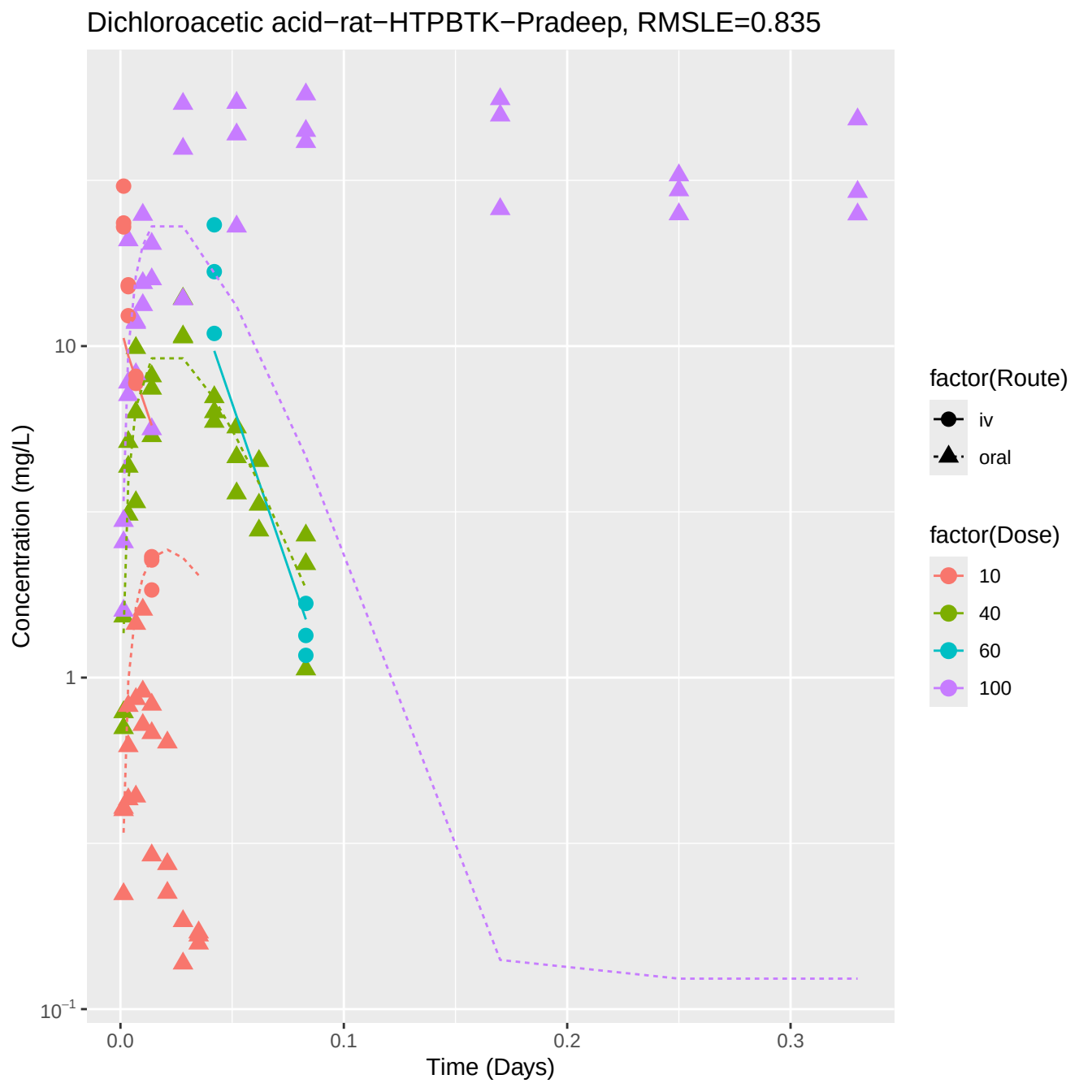

Formamide−rat−HTPBTK−ADMET, RMSLE=0.868
Concentration (mg/L)
10^3
10^2
10
1
0
1
2
3
4
Time (Days)
factor(Route)
iv
oral
factor(Dose)
10
30
36
90
300
600

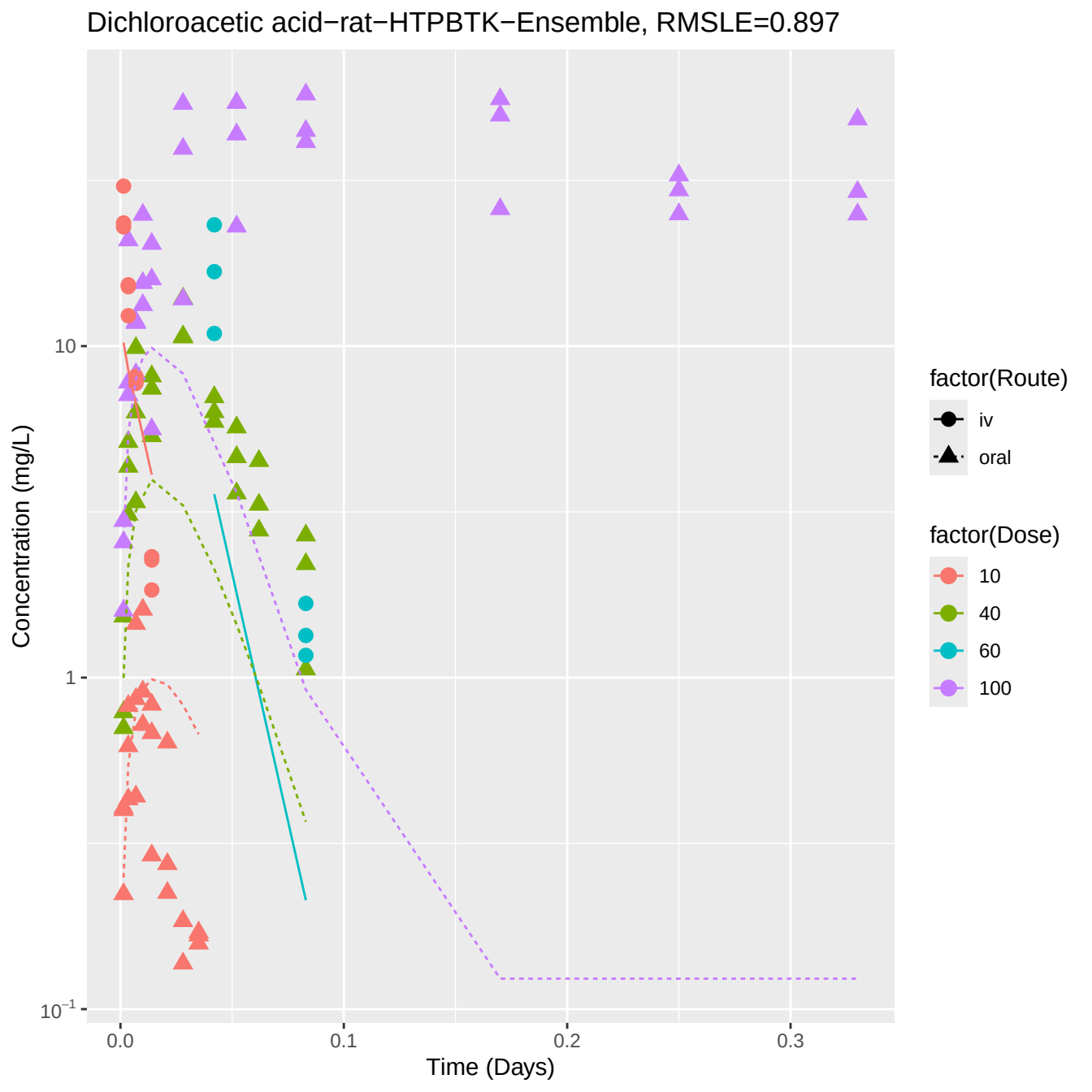

Formamide−rat−HTPBTK−Pradeep, RMSLE=1.28
Concentration (mg/L)
10^3
10^2
10
1
0
1
2
3
4
Time (Days)
factor(Route)
iv
oral
factor(Dose)
10
30
36
90
300
600

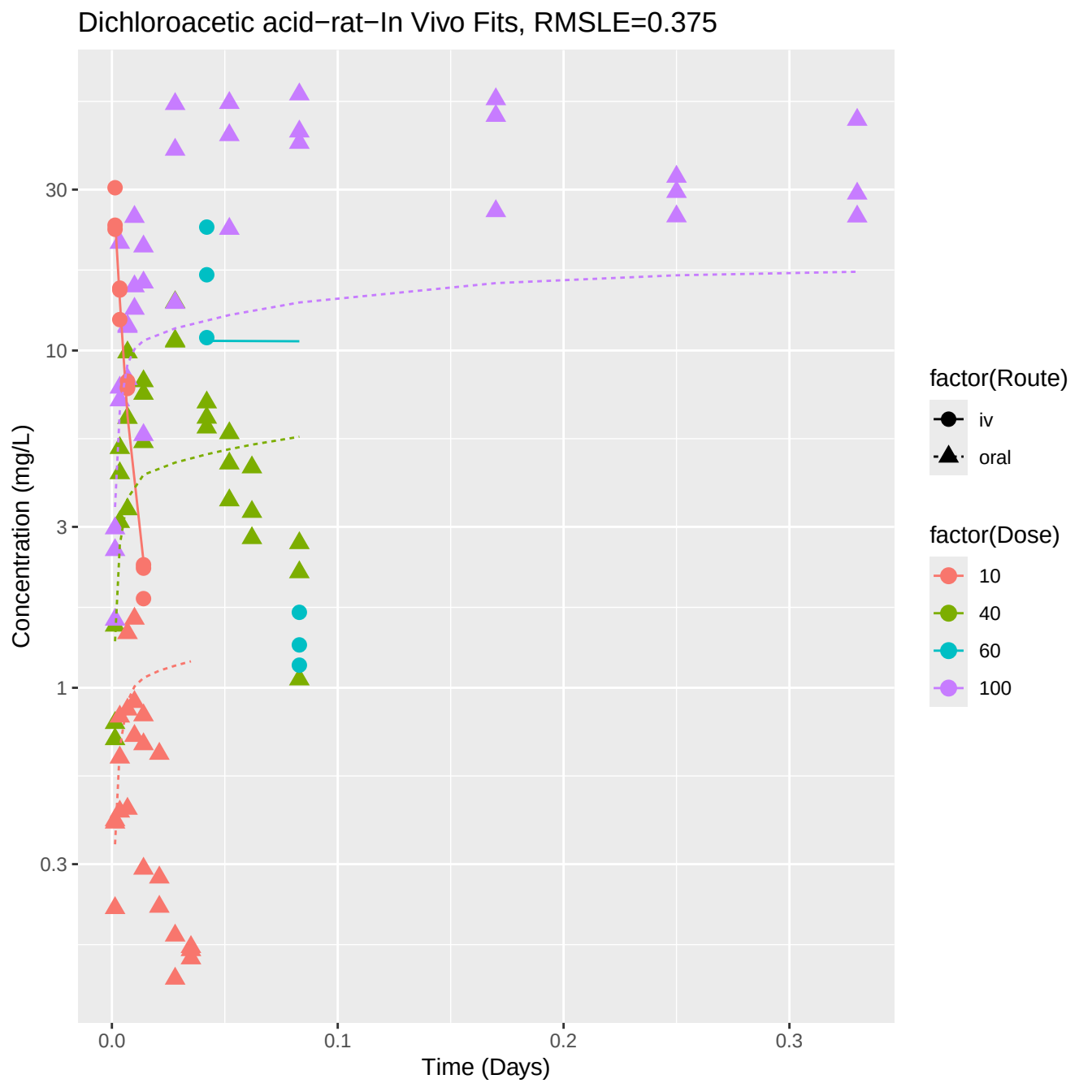

Formamide−rat−HTPBTK−Ensemble, RMSLE=1.28
Concentration (mg/L)
10^3
10^2
10
1
0
1
2
3
4
Time (Days)
factor(Route)
iv
oral
factor(Dose)
10
30
36
90
300
600

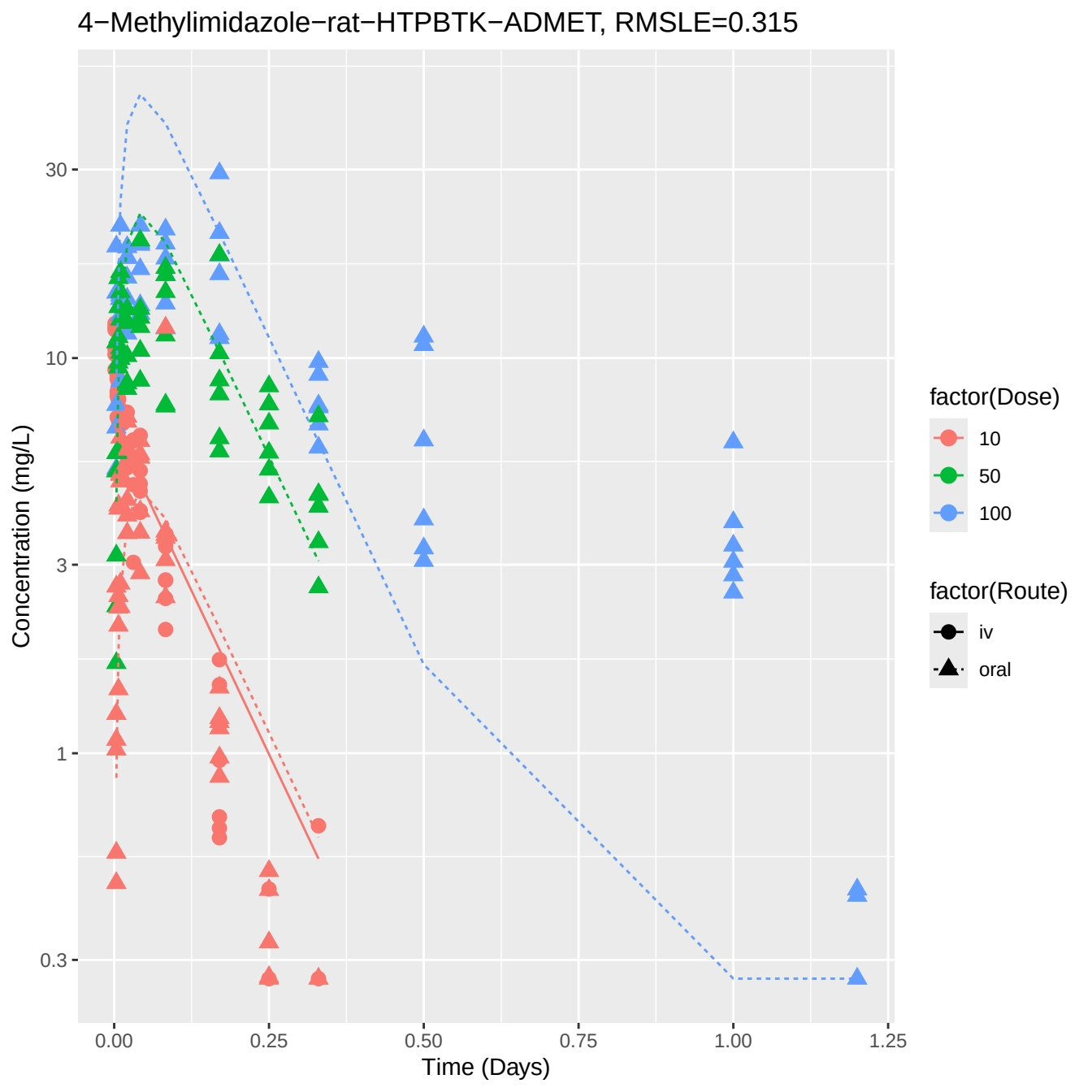

Formamide−rat−In Vivo Fits, RMSLE=0.188
Concentration (mg/L)
10^3
10^2
10
1
0
1
2
3
4
Time (Days)
factor(Route)
iv
oral
factor(Dose)
10
30
36
90
300
600

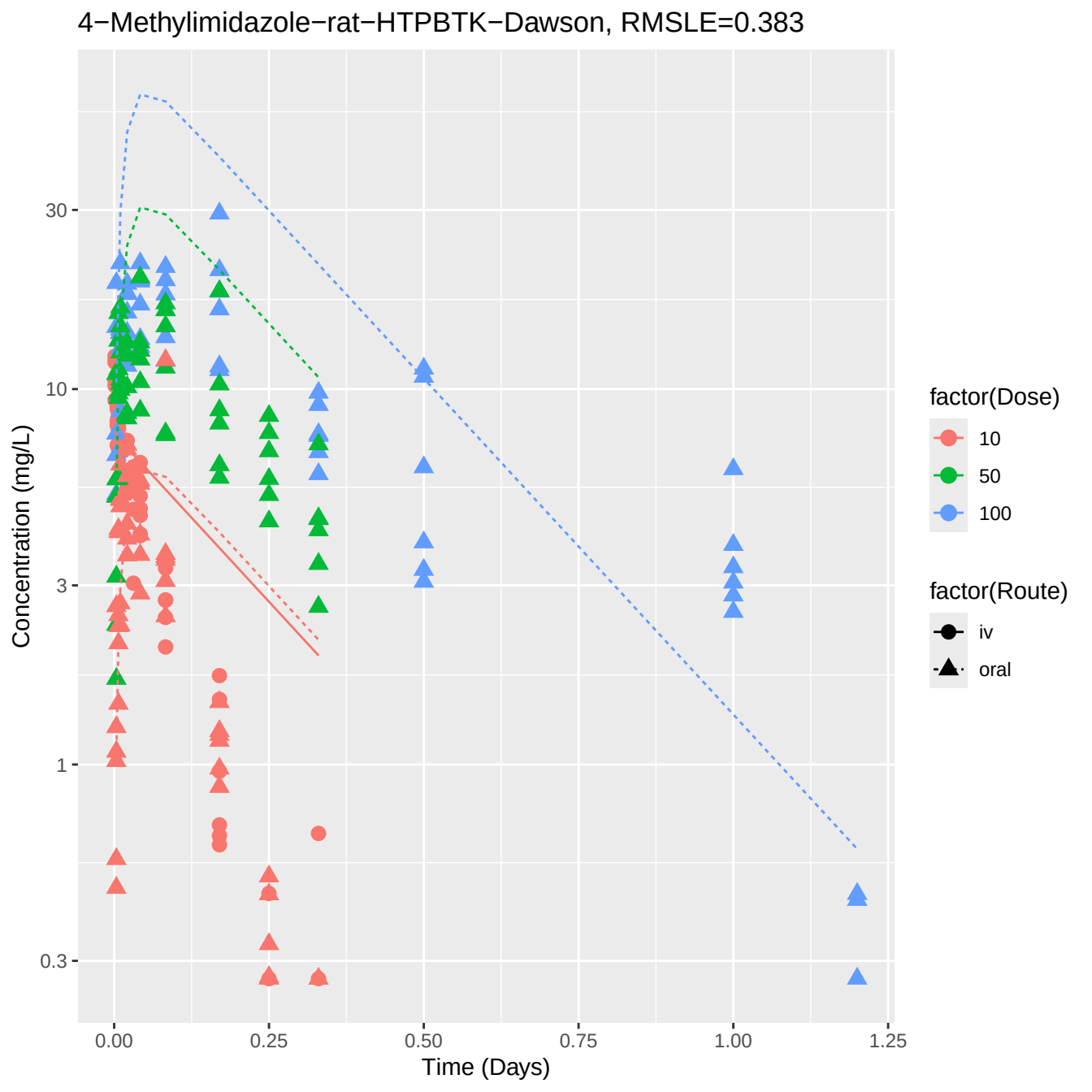

Dichloroacetic acid−rat−HTPBTK−ADMET, RMSLE=0.742
Concentration (mg/L)
10^2
10
1
10^{-1}
0.0
0.1
0.2
0.3
Time (Days)
factor(Route)
iv
oral
factor(Dose)
10
40
60
100

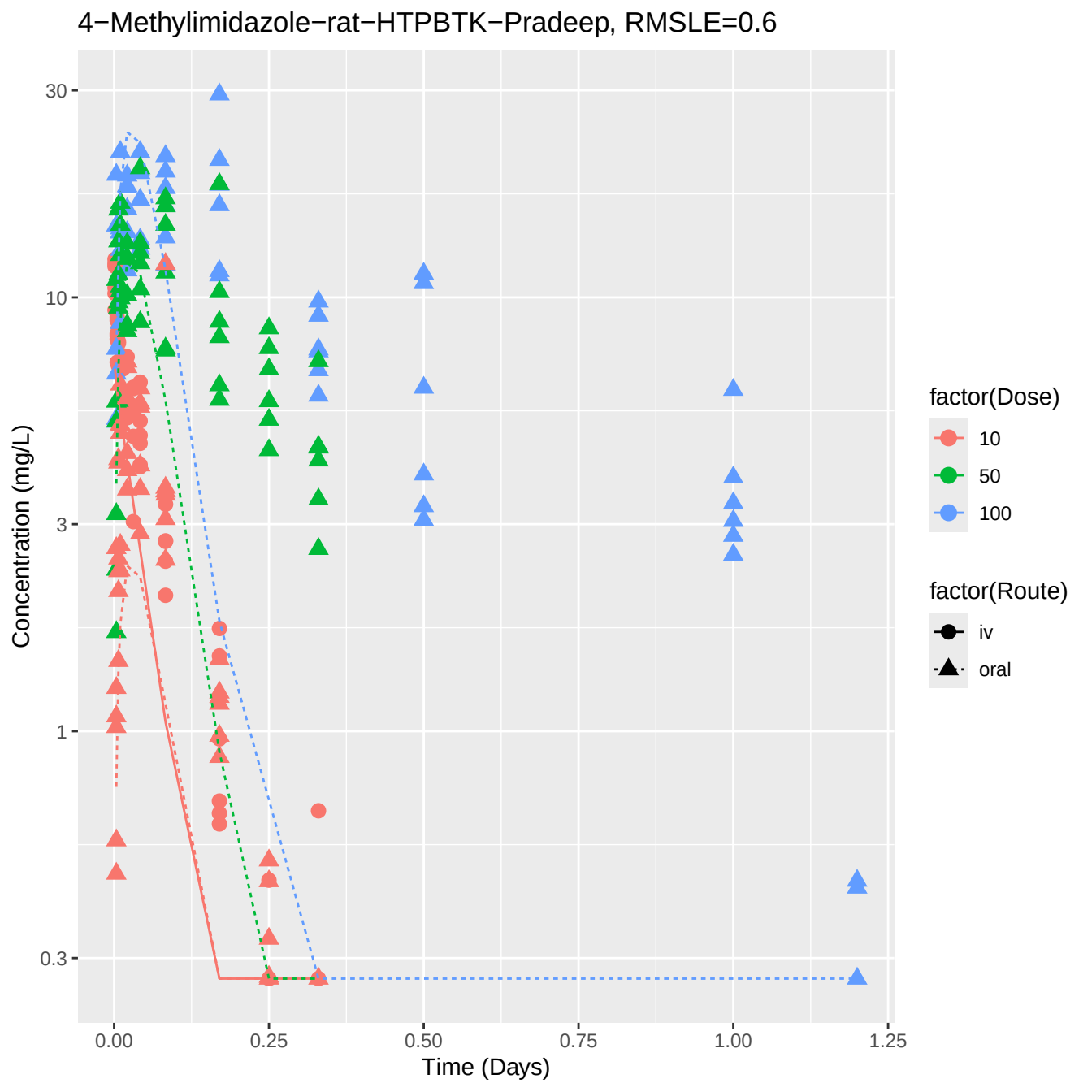

Dichloroacetic acid−rat−HTPBTK−Pradeep, RMSLE=0.835
Concentration (mg/L)
10
1
10^{-1}
0.0
0.1
0.2
0.3
Time (Days)
factor(Route)
iv
oral
factor(Dose)
10
40
60
100

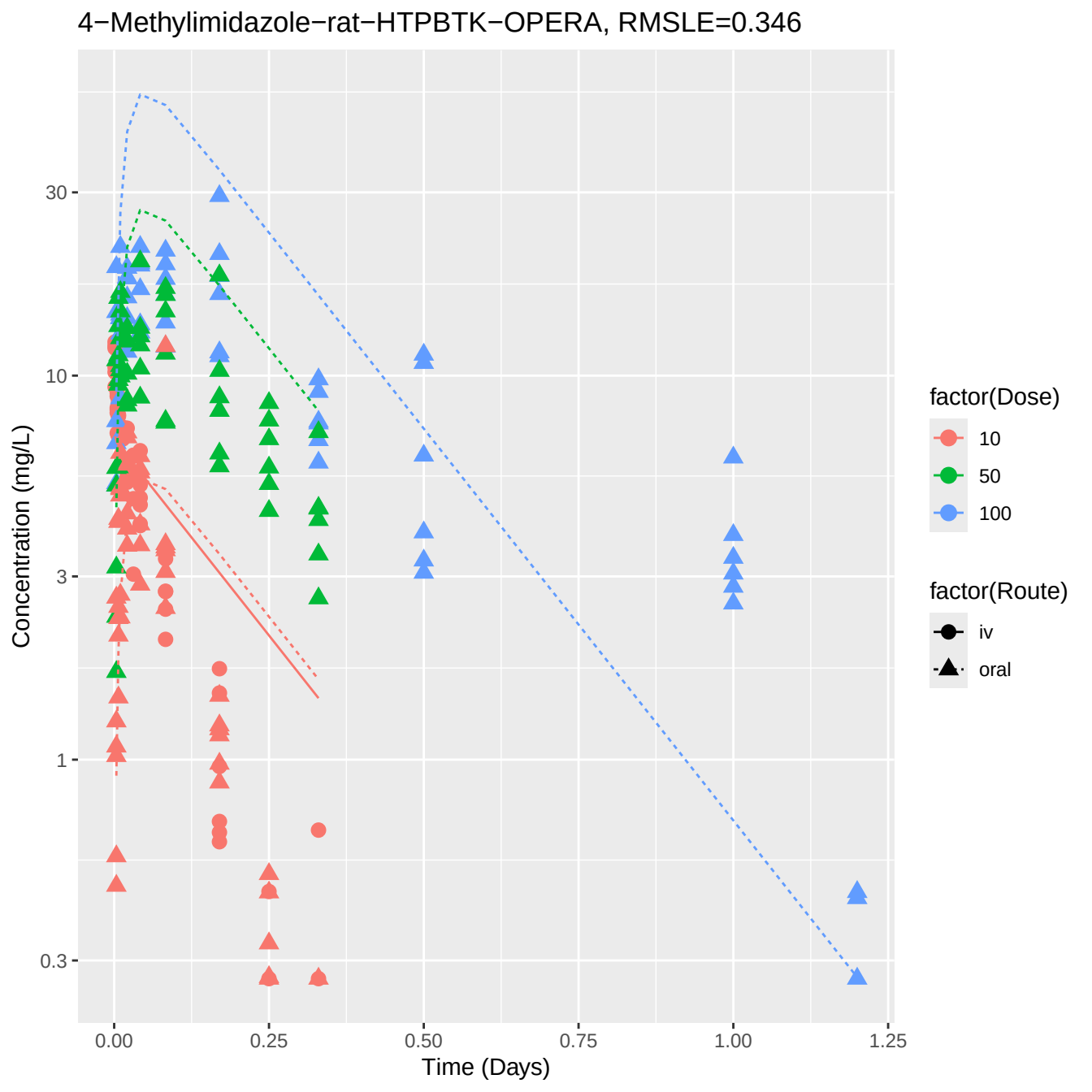

Dichloroacetic acid−rat−HTPBTK−Ensemble, RMSLE=0.897
Concentration (mg/L)
10
1
10^{-1}
0.0
0.1
0.2
0.3
Time (Days)
factor(Route)
iv
oral
factor(Dose)
10
40
60
100

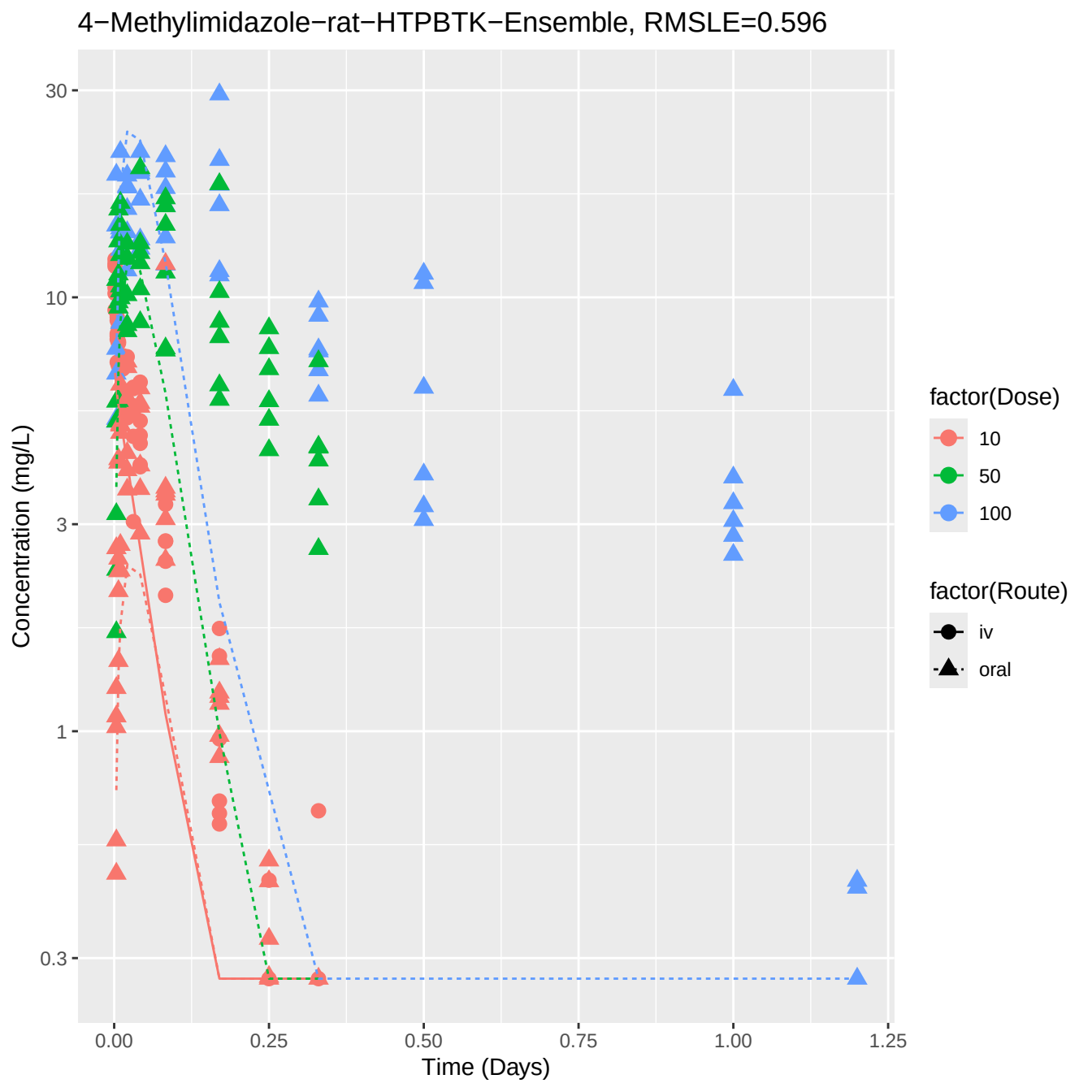

Dichloroacetic acid−rat−In Vivo Fits, RMSLE=0.375
Concentration (mg/L)
30
10
3
1
0.3
0.0
0.1
0.2
0.3
Time (Days)
factor(Route)
iv
oral
factor(Dose)
10
40
60
100

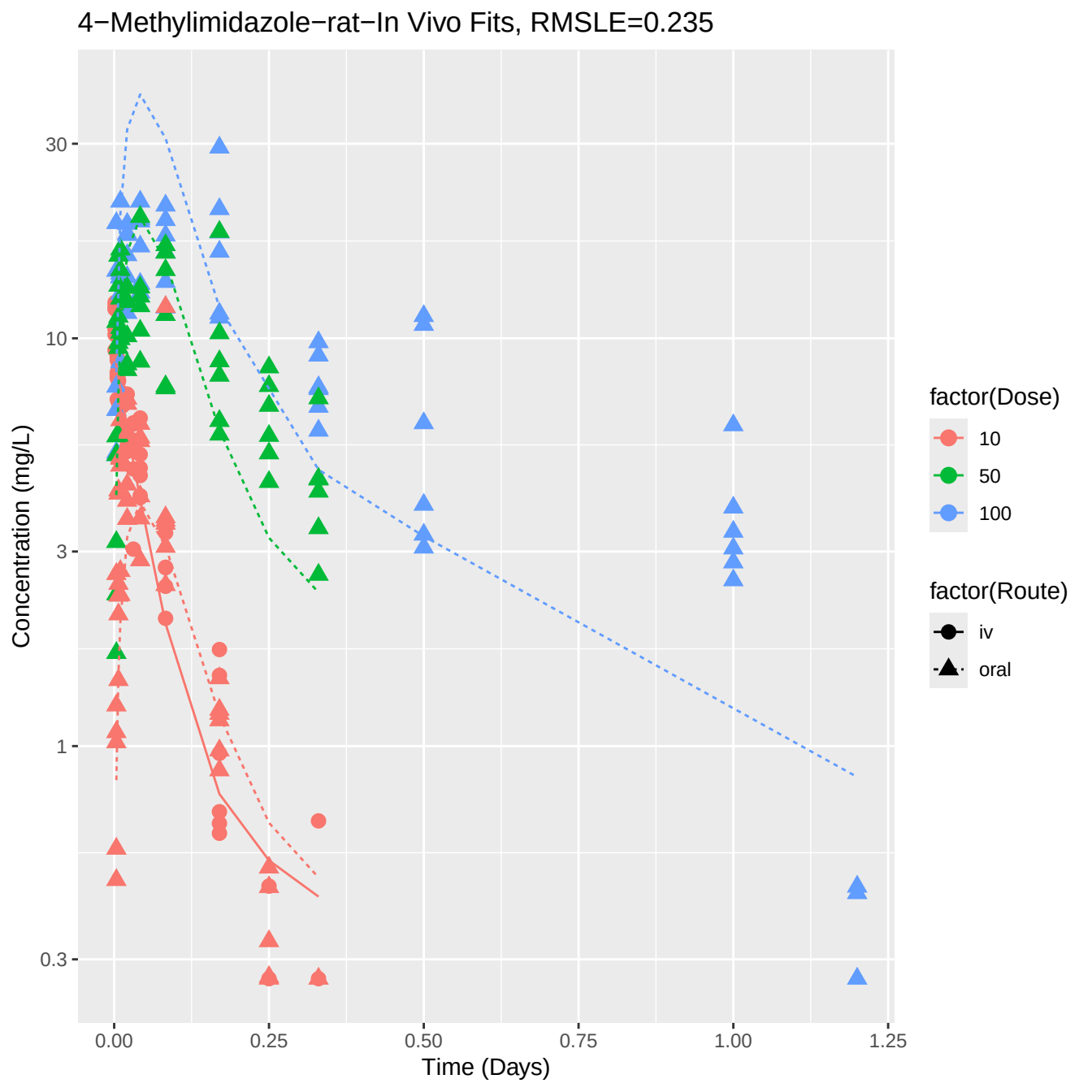

4−Methylimidazole−rat−HTPBTK−ADMET, RMSLE=0.315
Concentration (mg/L)
30
10
3
1
0.3
0.00
0.25
0.50
0.75
1.00
1.25
Time (Days)
factor(Dose)
10
50
100
factor(Route)
iv
oral

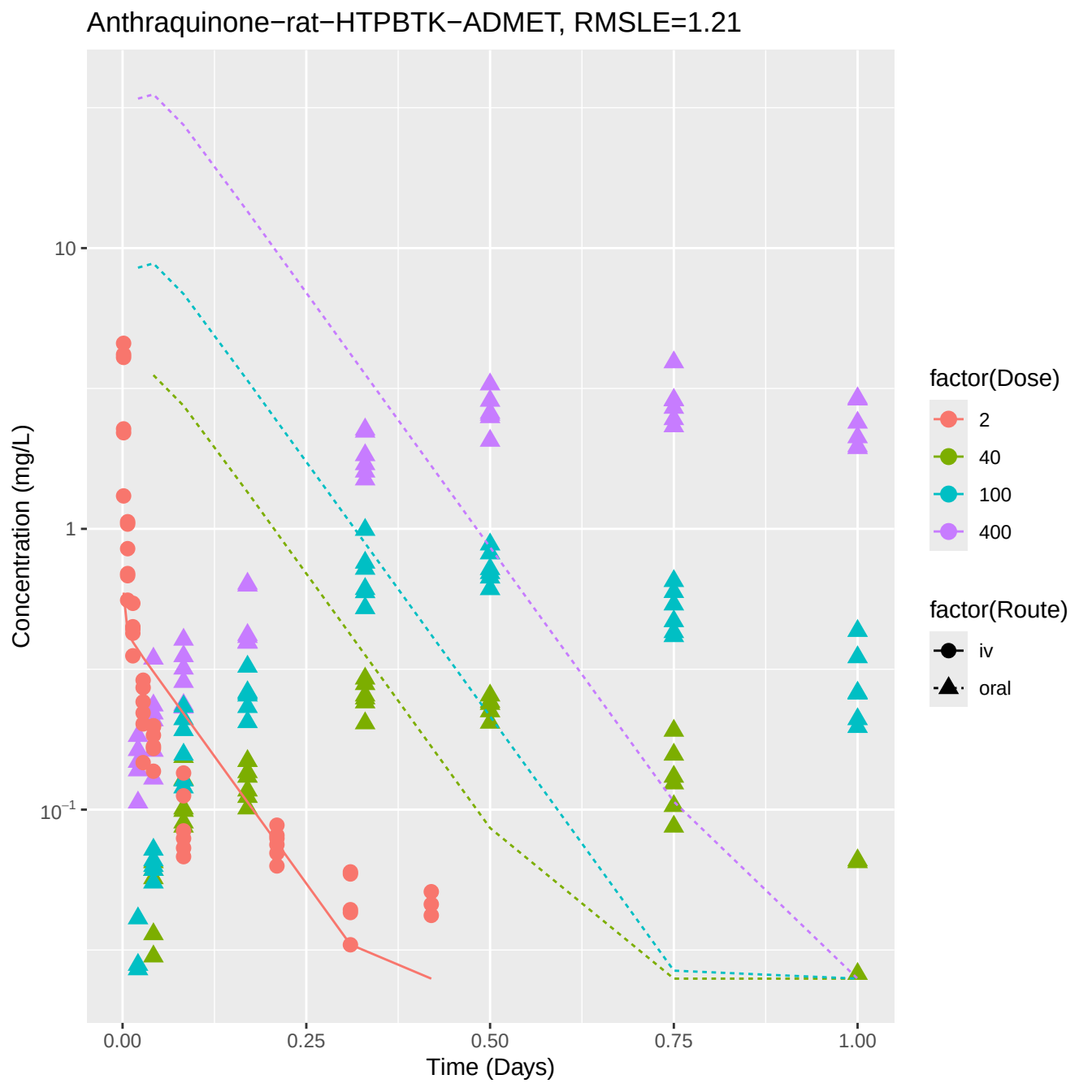

4−Methylimidazole−rat−HTPBTK−Dawson, RMSLE=0.383
Concentration (mg/L)
30
10
3
1
0.3
0.00
0.25
0.50
0.75
1.00
1.25
Time (Days)
factor(Dose)
10
50
100
factor(Route)
iv
oral

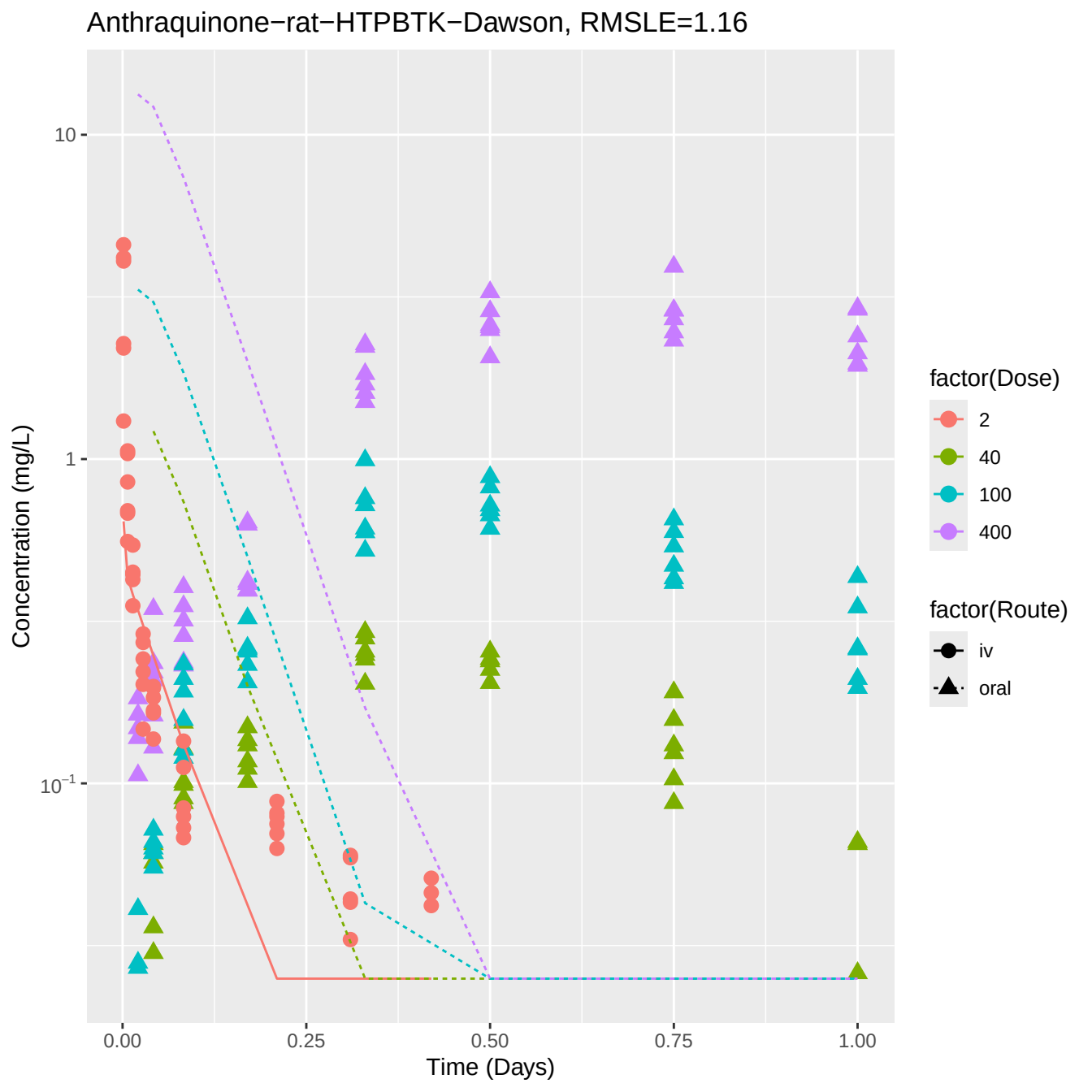

4−Methylimidazole−rat−HTPBTK−Pradeep, RMSLE=0.6
Concentration (mg/L)
30
10
3
1
0.3
0.00
0.25
0.50
0.75
1.00
1.25
Time (Days)
factor(Dose)
10
50
100
factor(Route)
iv
oral

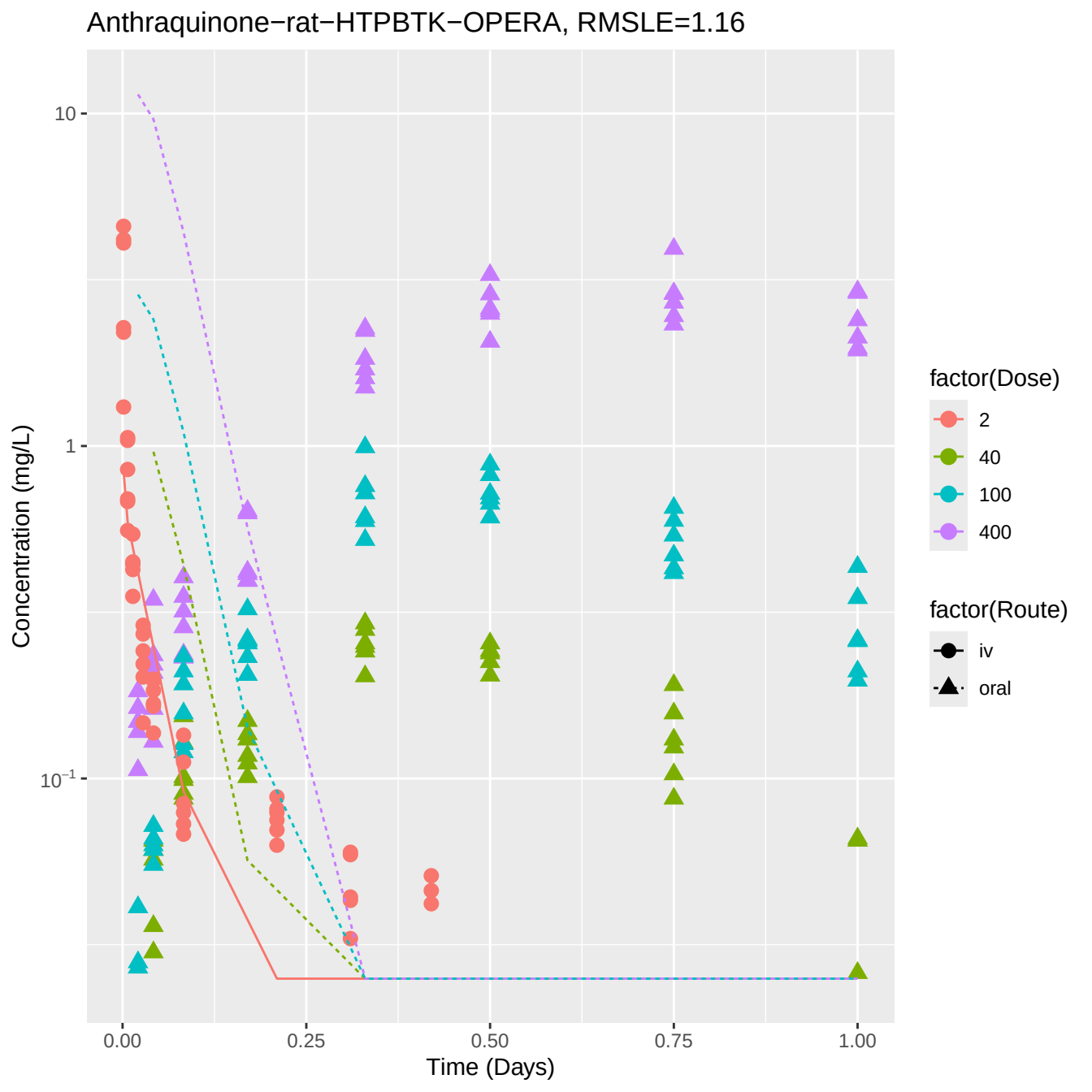

4−Methylimidazole−rat−HTPBTK−OPERA, RMSLE=0.346
Concentration (mg/L)
30
10
3
1
0.3
0.00
0.25
0.50
0.75
1.00
1.25
Time (Days)
factor(Dose)
10
50
100
factor(Route)
iv
oral

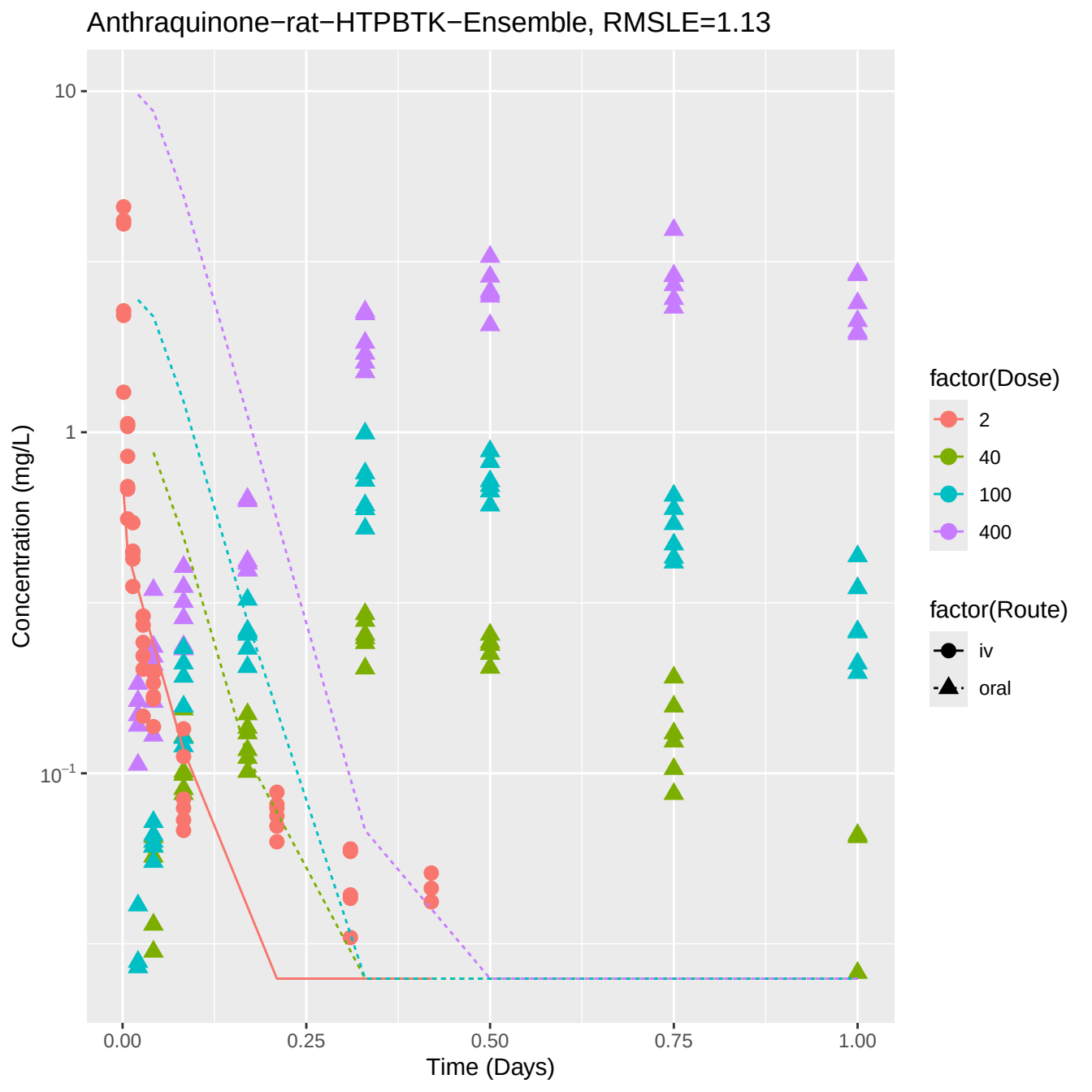

4−Methylimidazole−rat−HTPBTK−Ensemble, RMSLE=0.596
Concentration (mg/L)
30
10
3
1
0.3
0.00
0.25
0.50
0.75
1.00
1.25
Time (Days)
factor(Dose)
10
50
100
factor(Route)
iv
oral

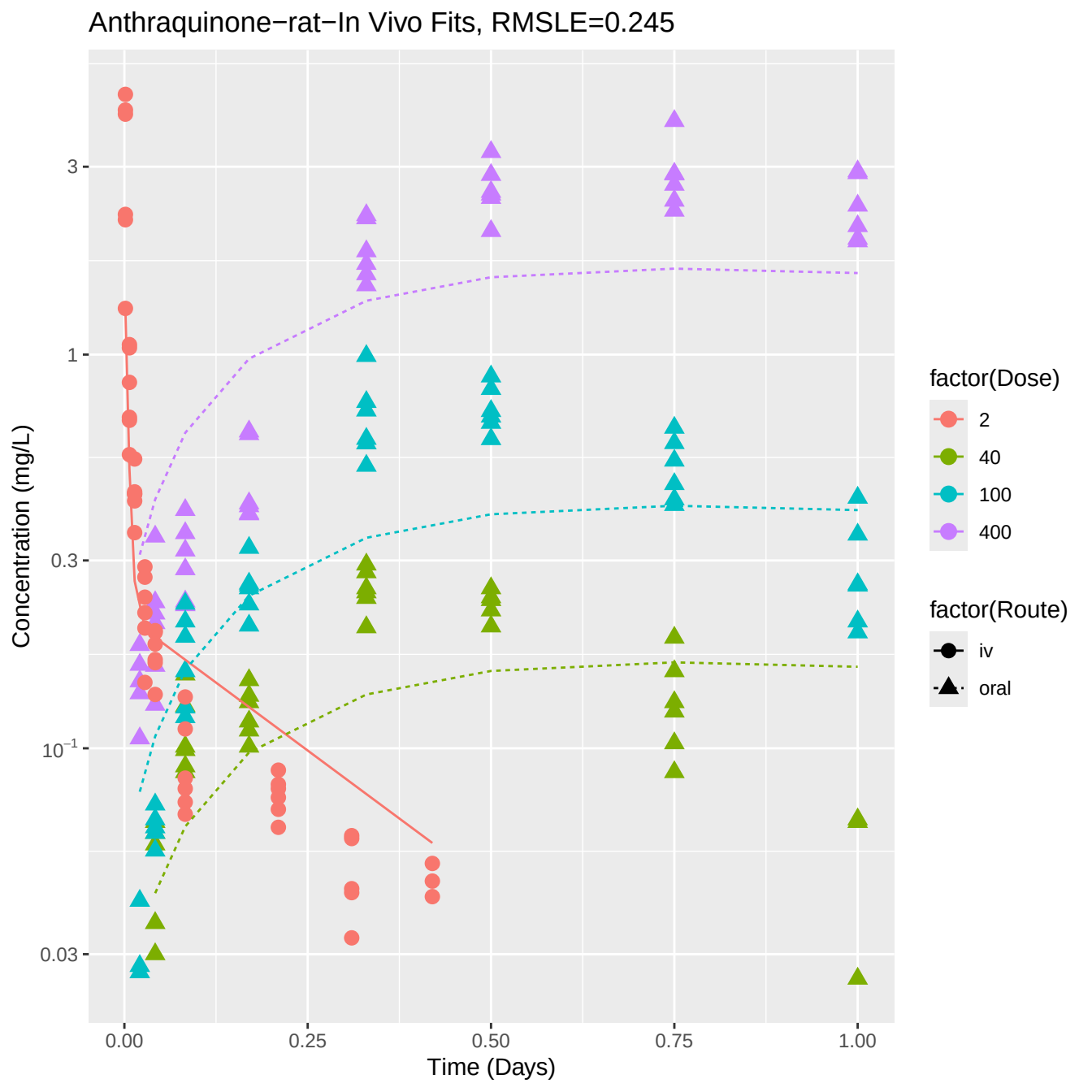

4−Methylimidazole−rat−In Vivo Fits, RMSLE=0.235
Concentration (mg/L)
30
10
3
1
0.3
0.00
0.25
0.50
0.75
1.00
1.25
Time (Days)
factor(Dose)
10
50
100
factor(Route)
iv
oral

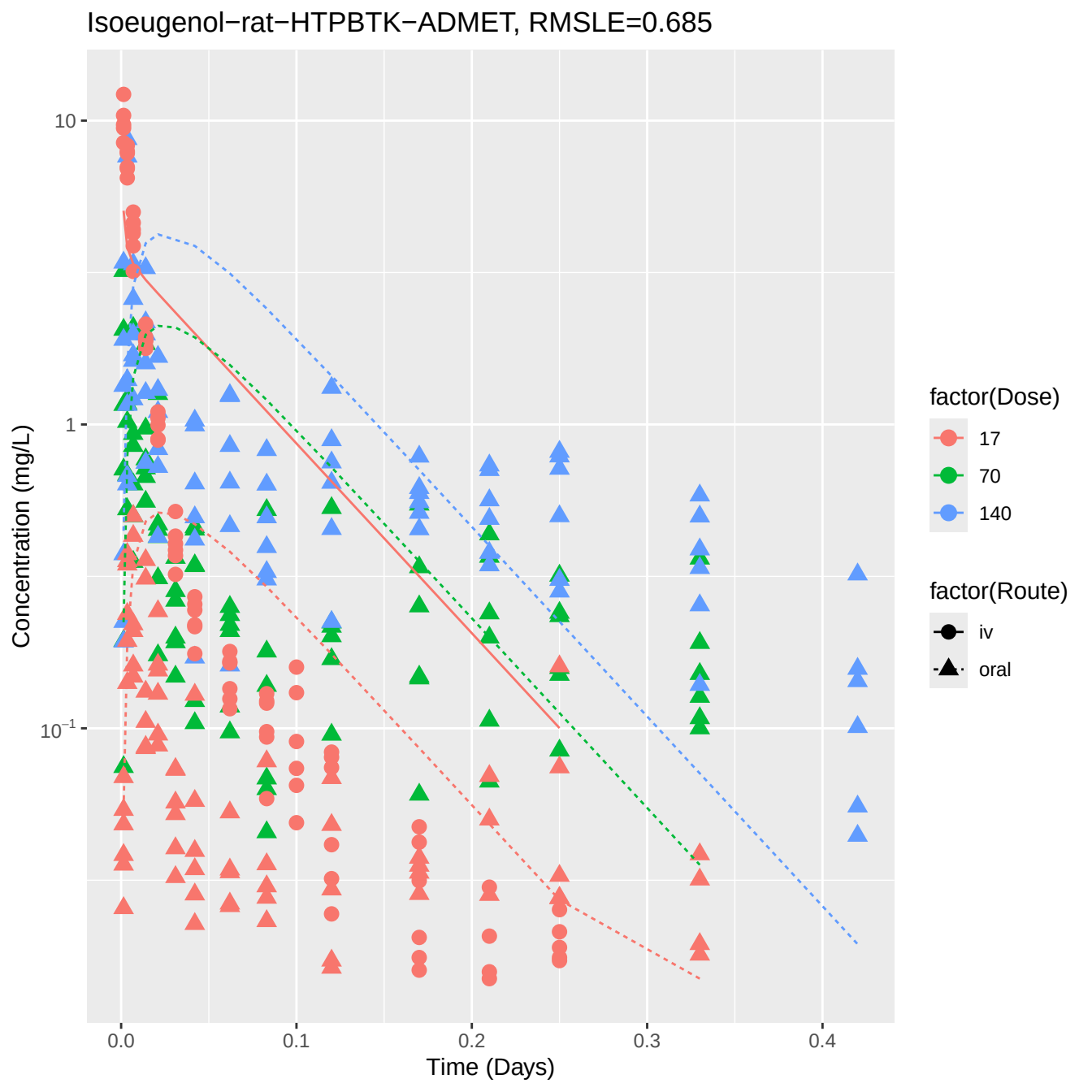

Anthraquinone−rat−HTPBTK−ADMET, RMSLE=1.21
Concentration (mg/L)
10
1
10^{-1}
0.00
0.25
0.50
0.75
1.00
Time (Days)
factor(Dose)
2
40
100
400
factor(Route)
iv
oral

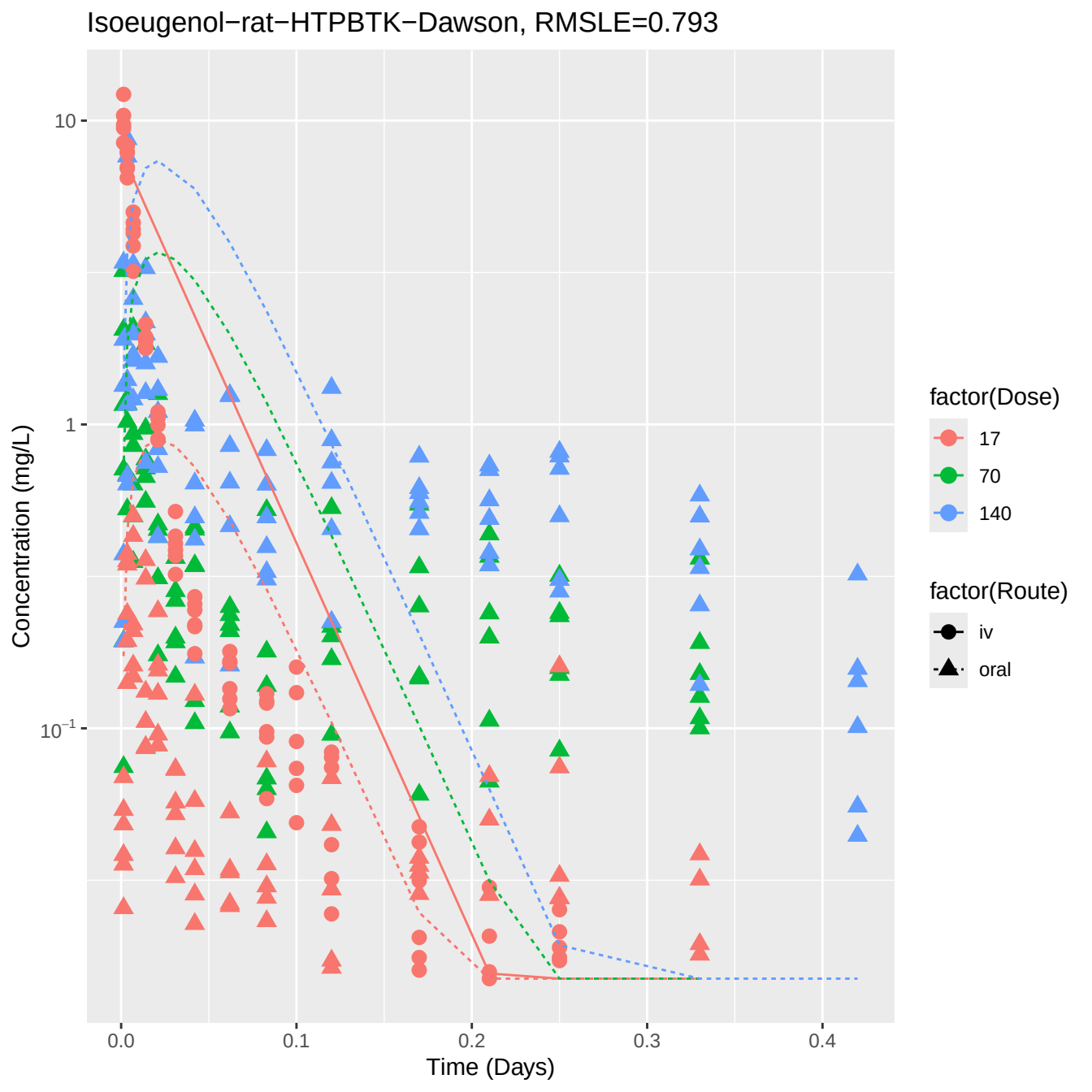

Anthraquinone−rat−HTPBTK−Dawson, RMSLE=1.16
10
1
10^{-1}
Concentration (mg/L)
0.00
0.25
0.50
0.75
1.00
Time (Days)
factor(Dose)
2
40
100
400
factor(Route)
iv
oral

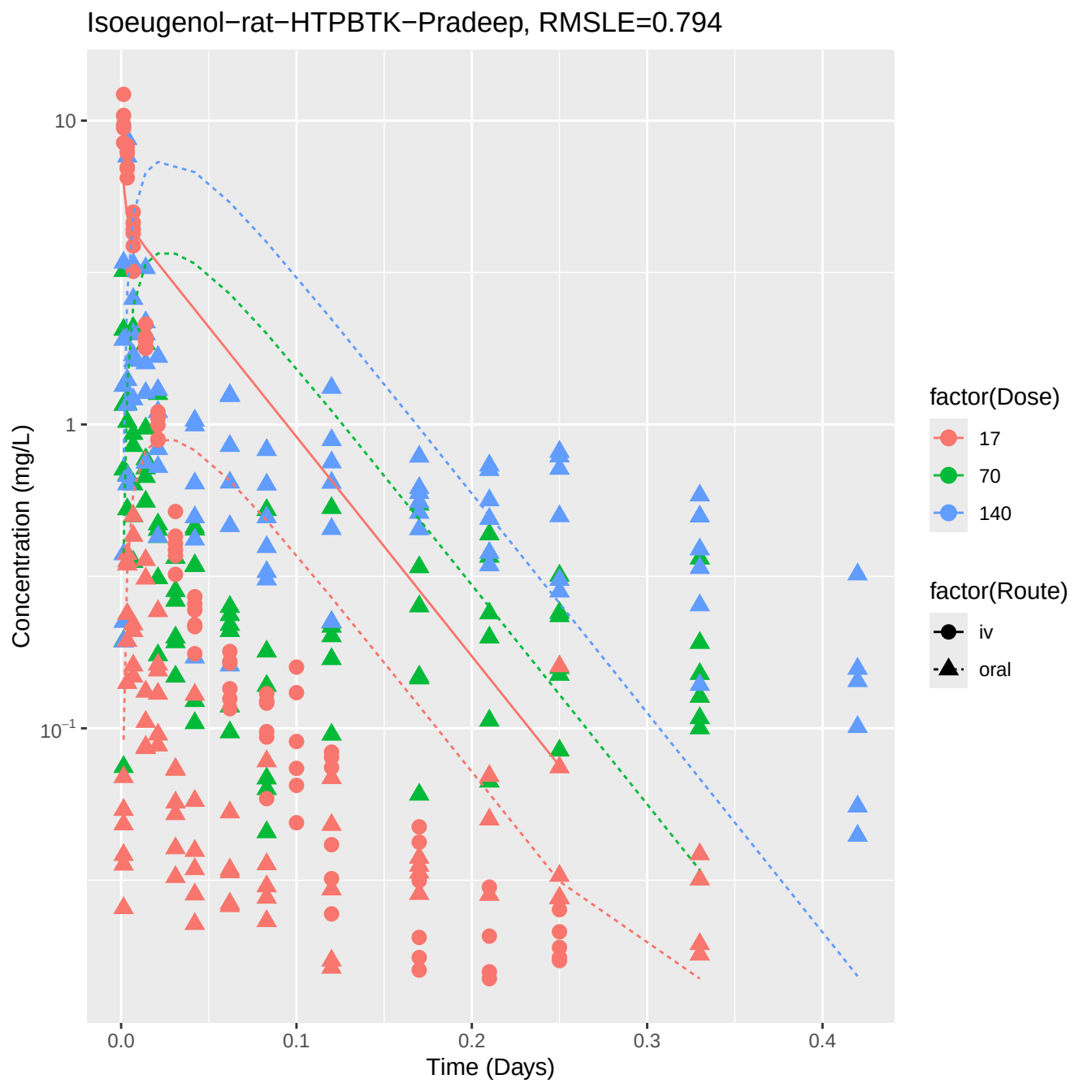

Anthraquinone−rat−HTPBTK−OPERA, RMSLE=1.16
Concentration (mg/L)
10
1
10^{-1}
0.00
0.25
0.50
0.75
1.00
Time (Days)
factor(Dose)
2
40
100
400
factor(Route)
iv
oral

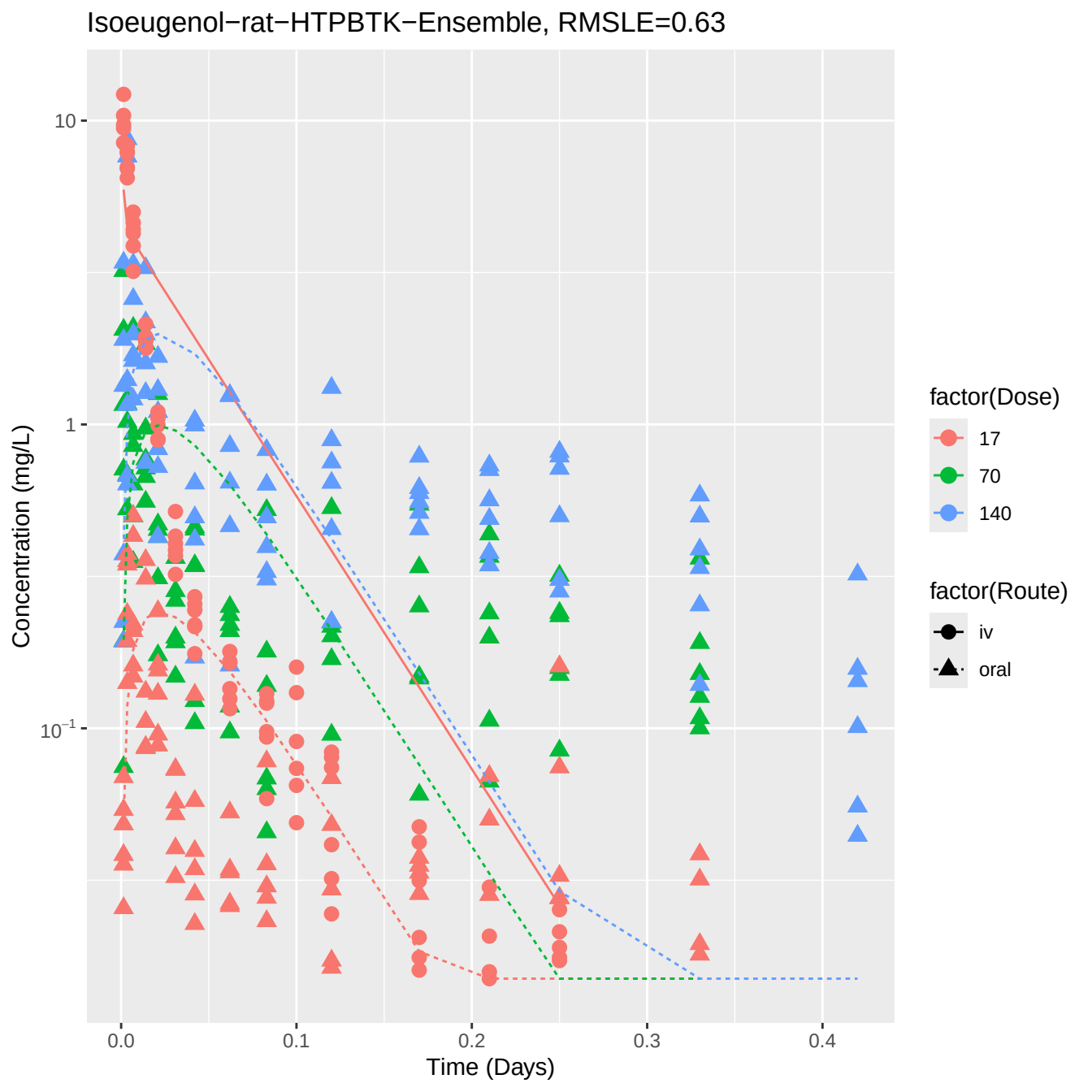

Anthraquinone−rat−HTPBTK−Ensemble, RMSLE=1.13
Concentration (mg/L)
10
1
10^{-1}
0.00
0.25
0.50
0.75
1.00
Time (Days)
factor(Dose)
2
40
100
400
factor(Route)
iv
oral

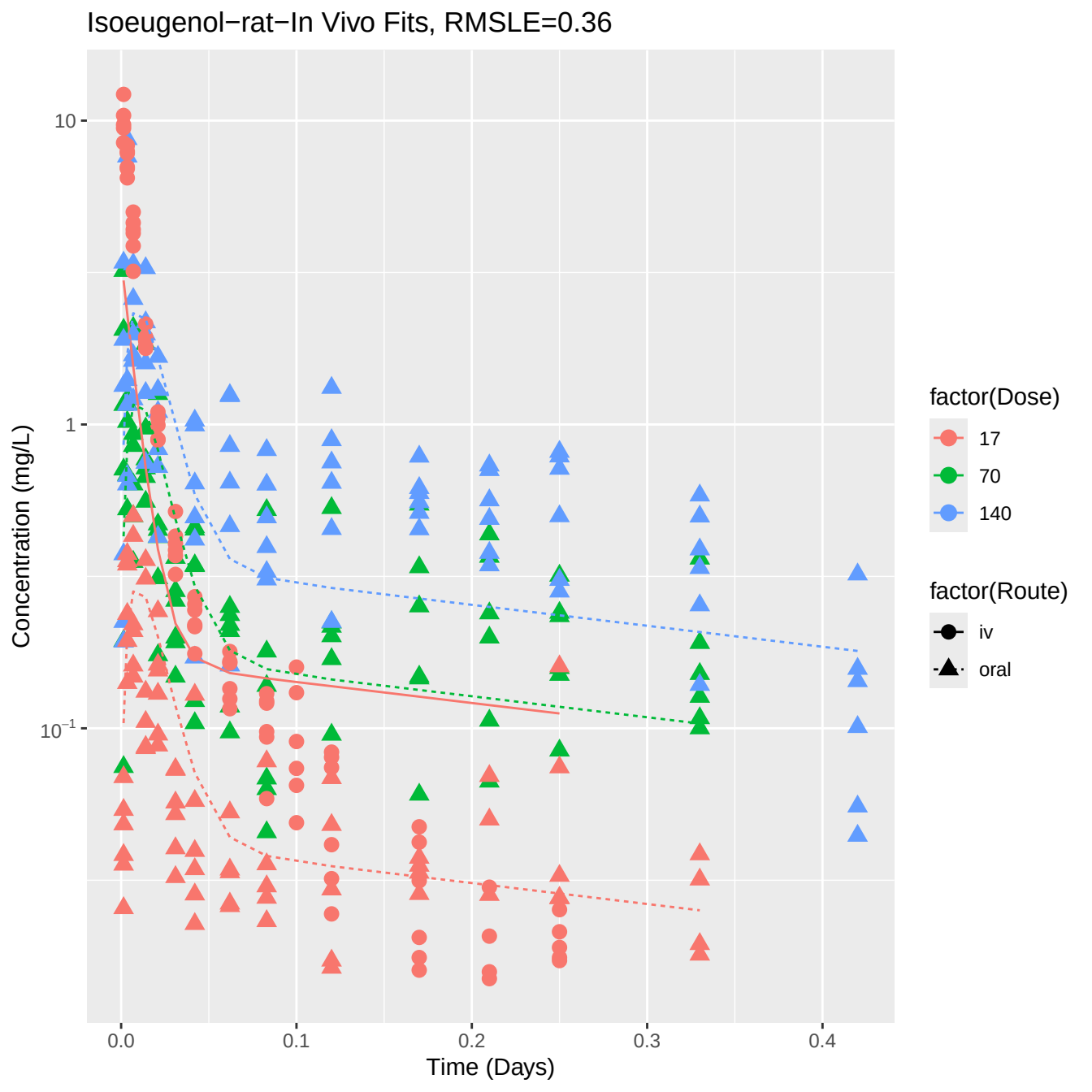

Anthraquinone−rat−In Vivo Fits, RMSLE=0.245
Concentration (mg/L)
3
1
0.3
10−1
0.03
0.00
0.25
0.50
0.75
1.00
Time (Days)
factor(Dose)
2
40
100
400
factor(Route)
iv
oral

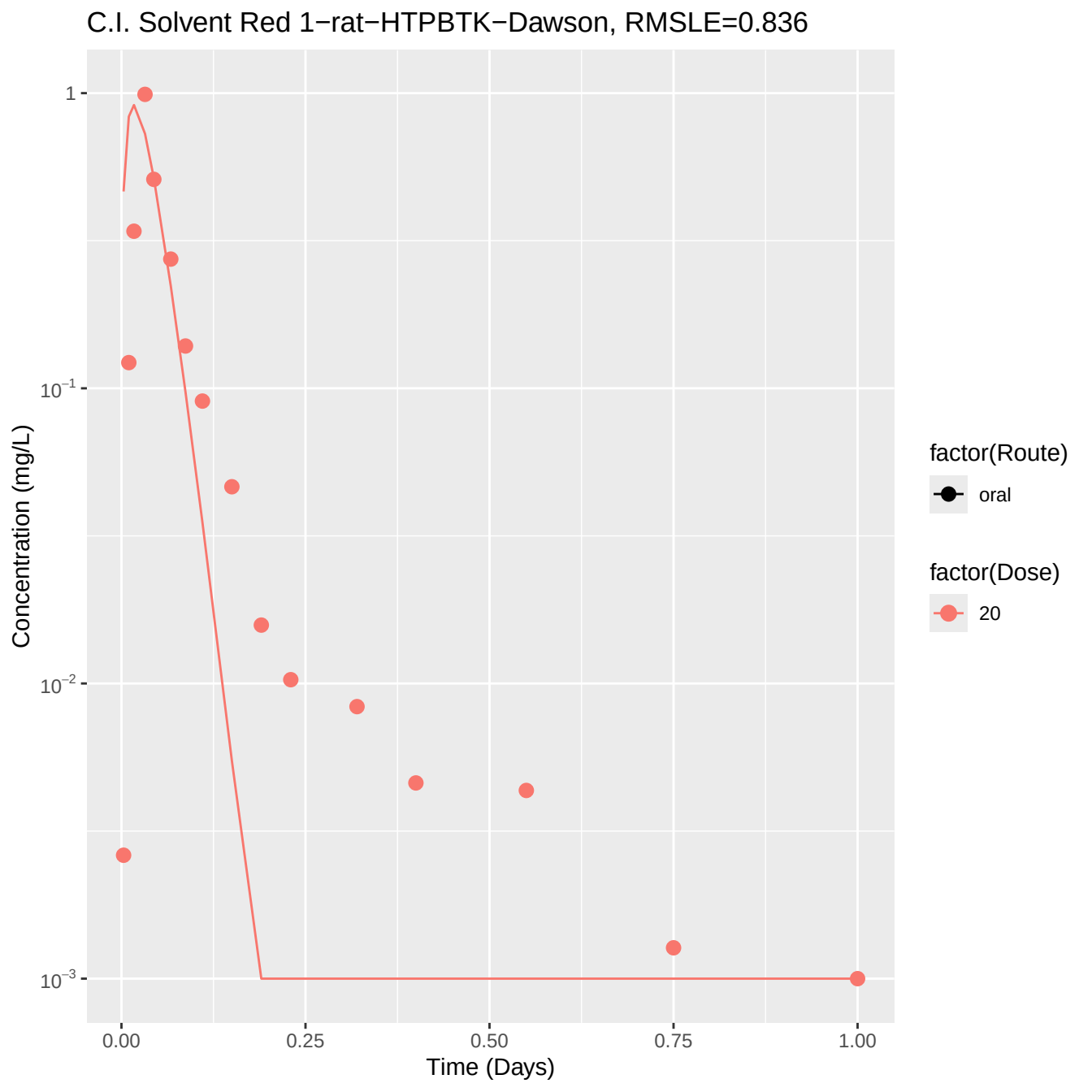

Isoeugenol−rat−HTPBTK−ADMET, RMSLE=0.685
10
1
10^{-1}
Concentration (mg/L)
0.0
0.1
0.2
0.3
0.4
Time (Days)
factor(Dose)
17
70
140
factor(Route)
iv
oral

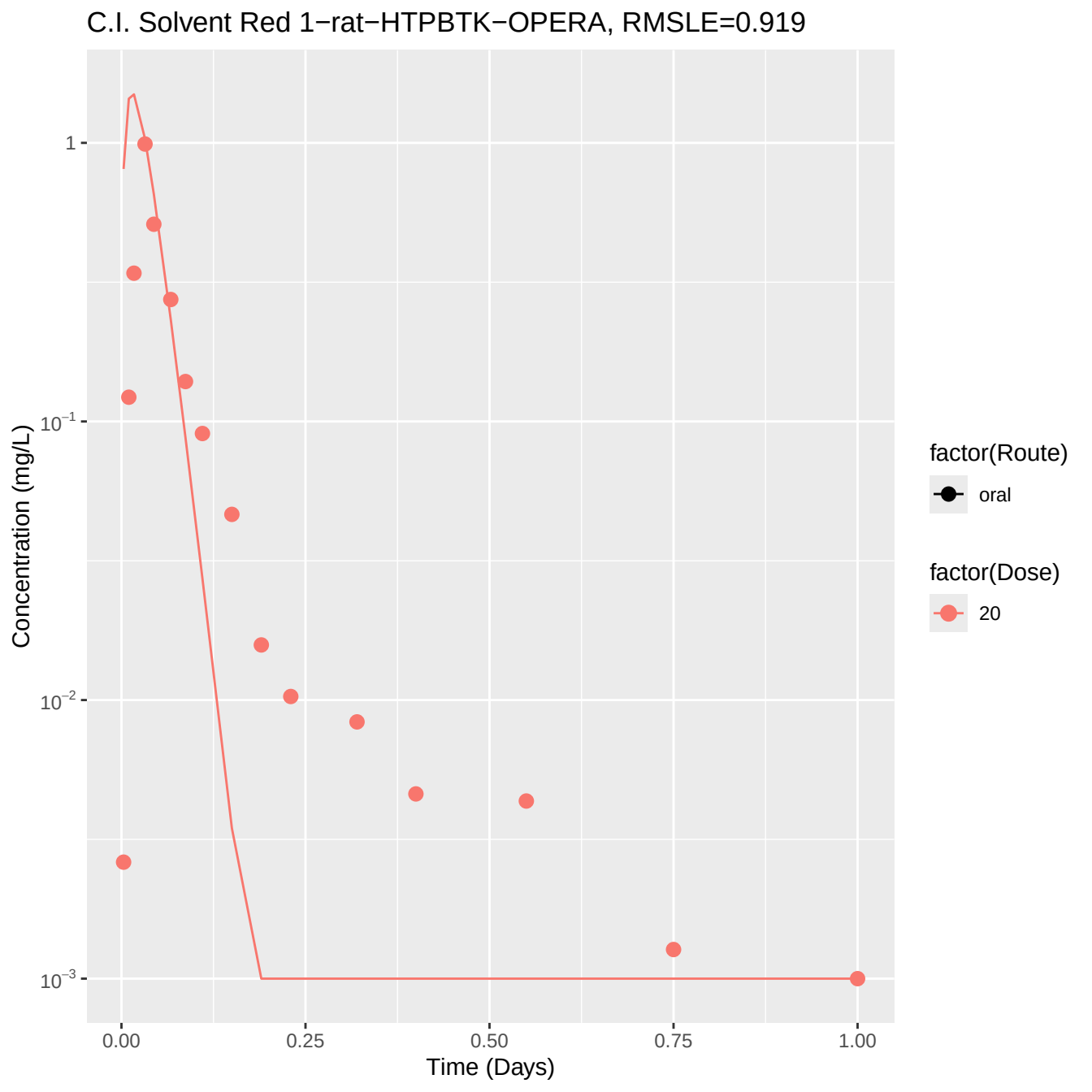

Isoeugenol−rat−HTPBTK−Dawson, RMSLE=0.793
Concentration (mg/L)
10
1
10^{-1}
0.0
0.1
0.2
0.3
0.4
Time (Days)
factor(Dose)
17
70
140
factor(Route)
iv
oral

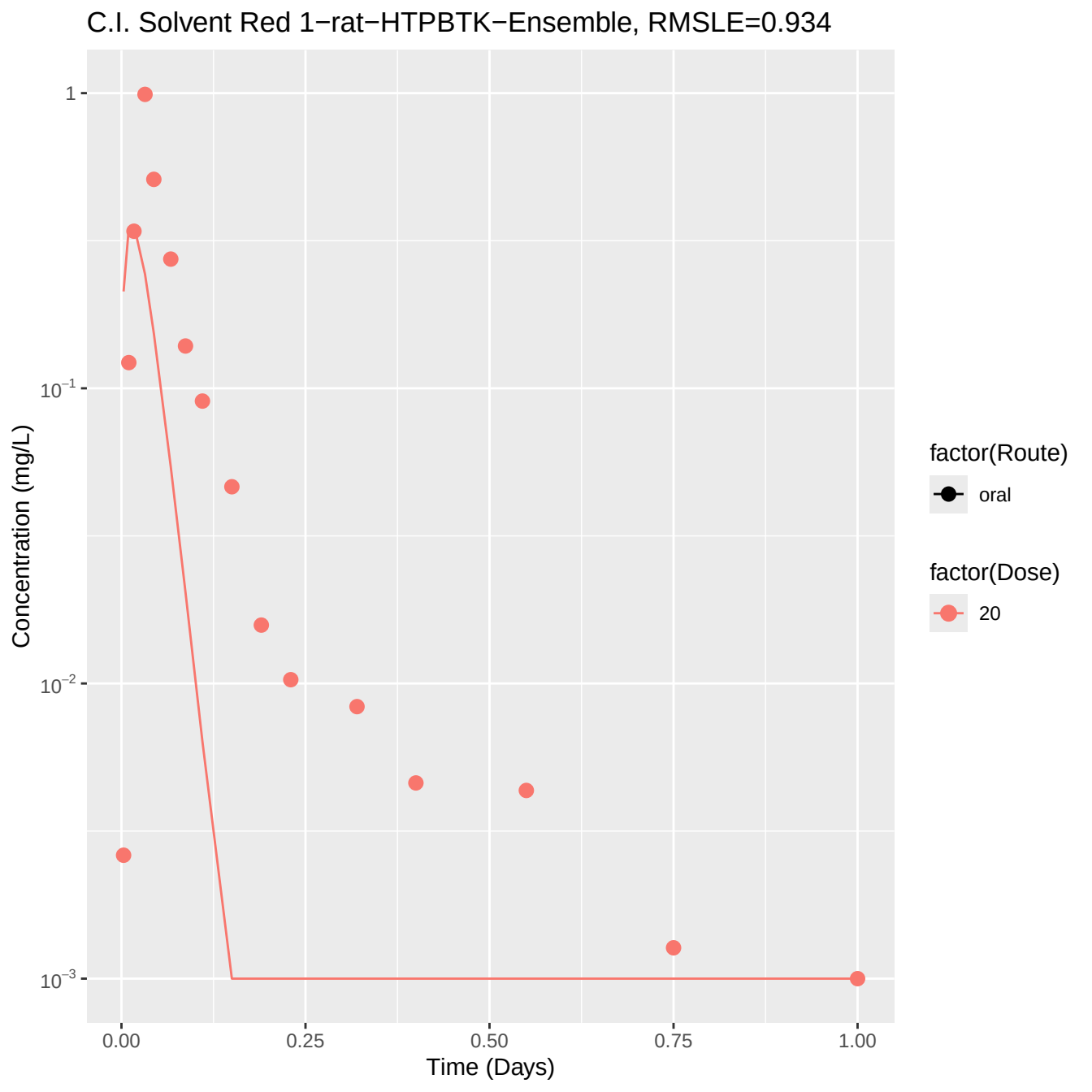

Isoeugenol−rat−HTPBTK−Pradeep, RMSLE=0.794
Concentration (mg/L)
10
1
10^{-1}
0.0
0.1
0.2
0.3
0.4
Time (Days)
factor(Dose)
17
70
140
factor(Route)
iv
oral

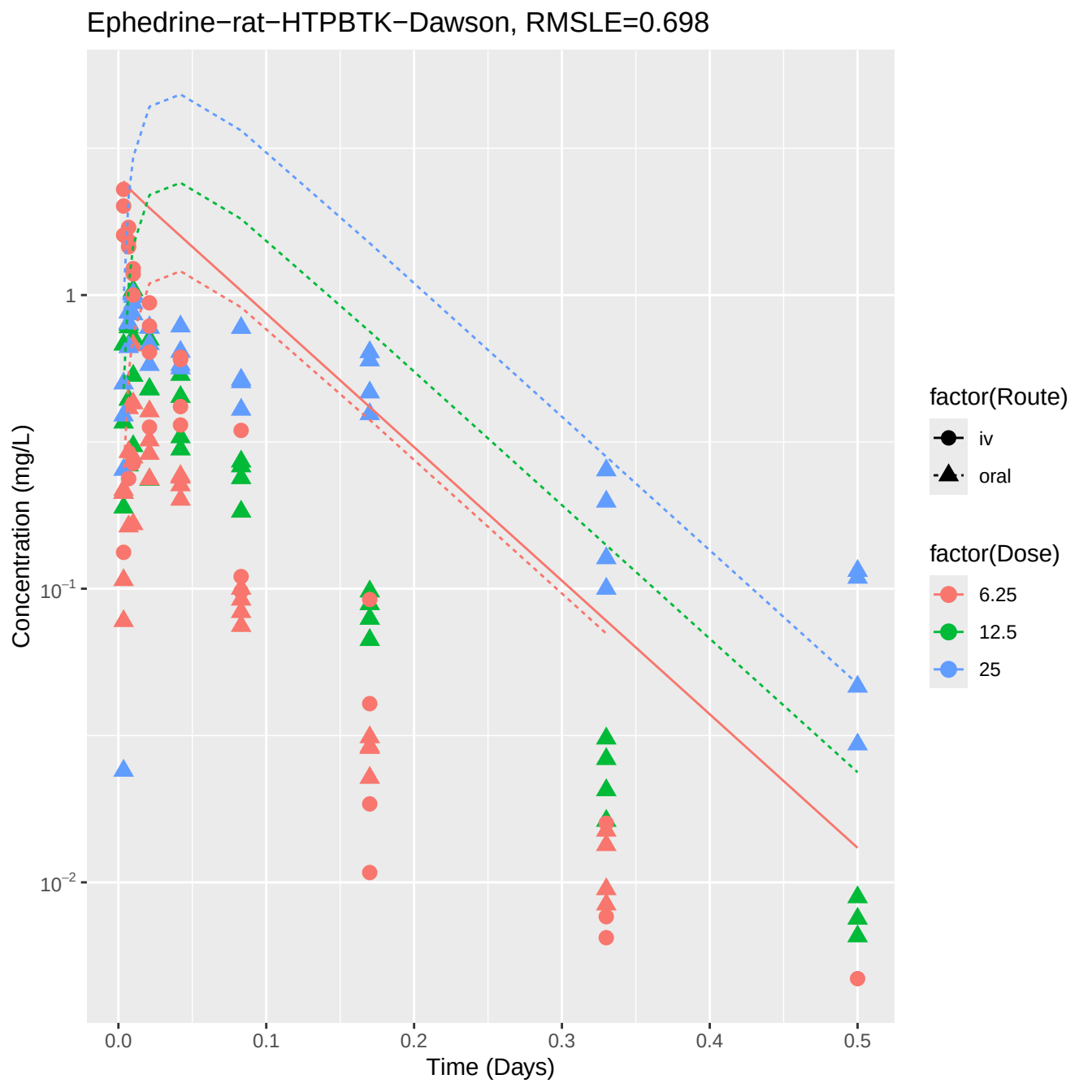

Isoeugenol−rat−HTPBTK−Ensemble, RMSLE=0.63
Concentration (mg/L)
10
1
10^{-1}
0.0
0.1
0.2
0.3
0.4
Time (Days)
factor(Dose)
17
70
140
factor(Route)
iv
oral

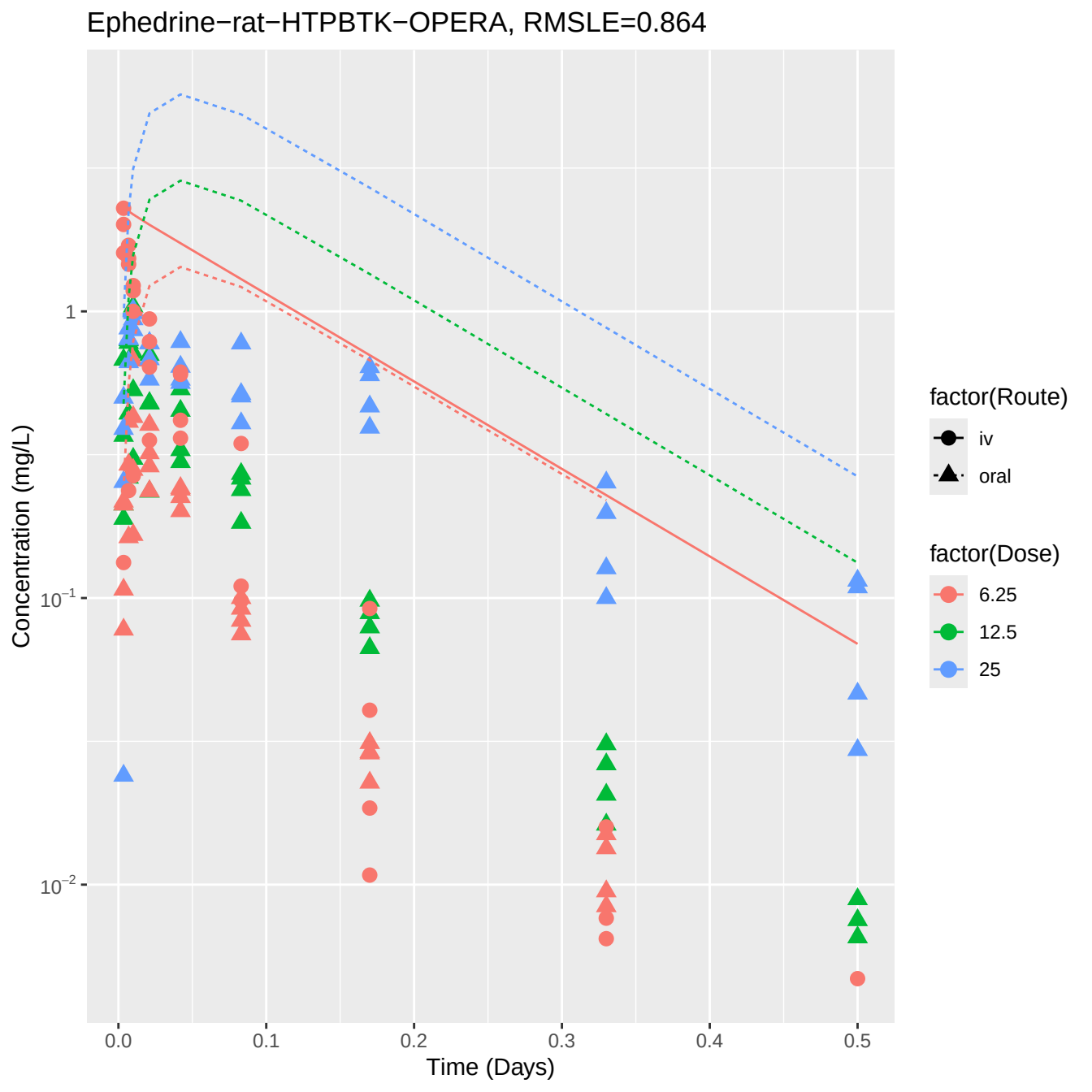

Isoeugenol−rat−In Vivo Fits, RMSLE=0.36
Concentration (mg/L)
10
1
10^{-1}
0.0
0.1
0.2
0.3
0.4
Time (Days)
factor(Dose)
17
70
140
factor(Route)
iv
oral

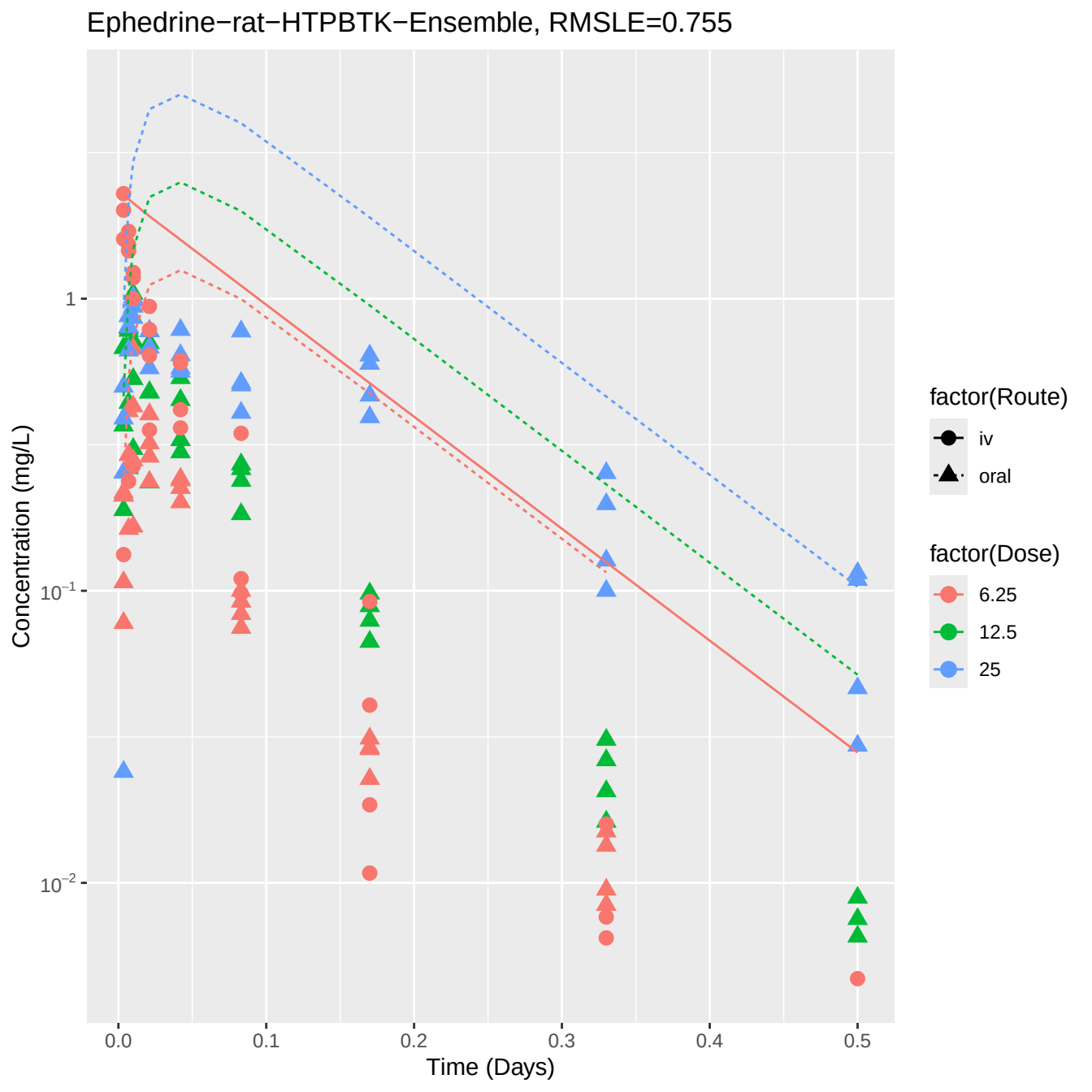

C.I. Solvent Red 1−rat−HTPBTK−Dawson, RMSLE=0.836
1
10^{-1}
10^{-2}
10^{-3}
Concentration (mg/L)
0.00
0.25
0.50
0.75
1.00
Time (Days)
factor(Route)
oral
factor(Dose)
20

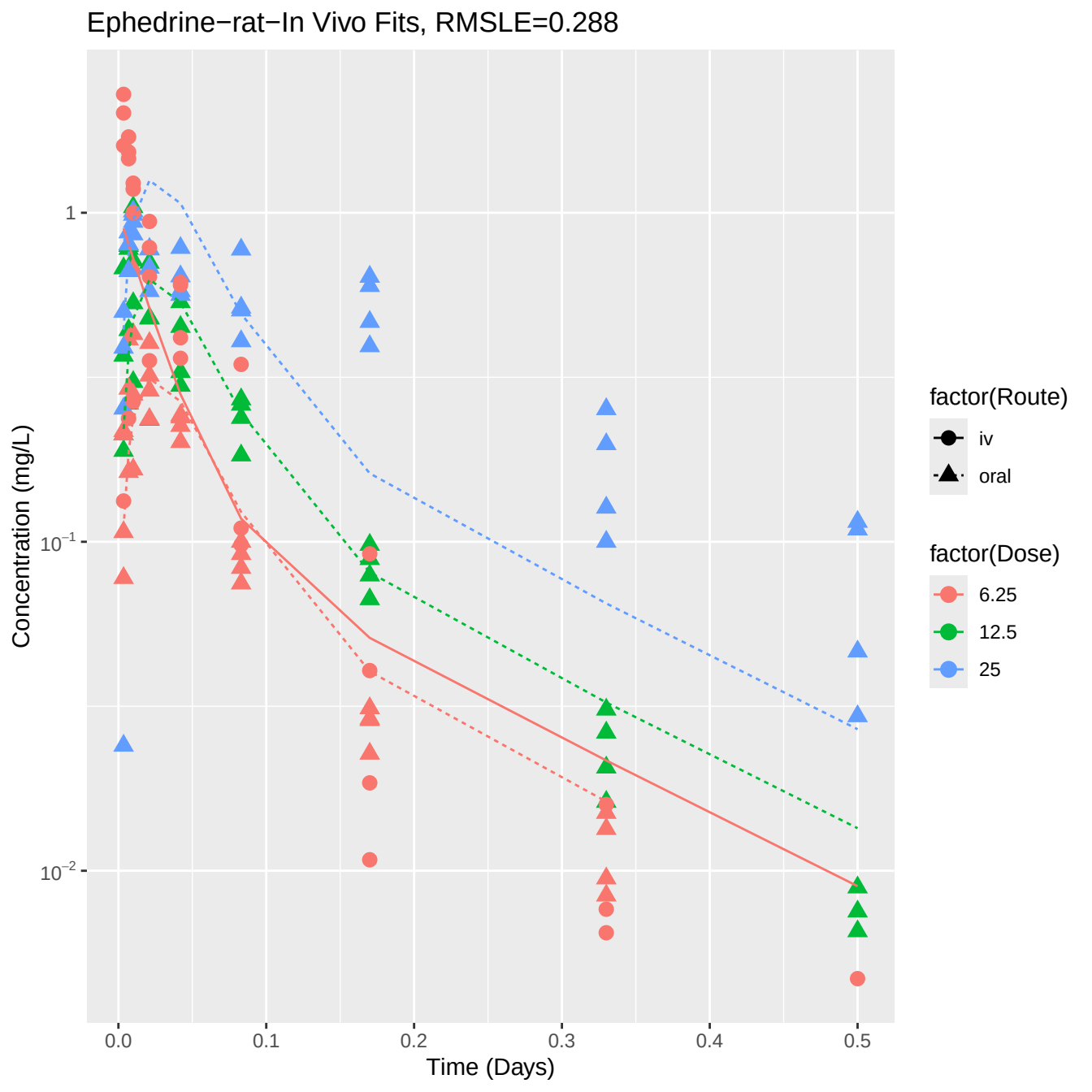

C.I. Solvent Red 1−rat−HTPBTK−OPERA, RMSLE=0.919
1
10^{-1}
10^{-2}
10^{-3}
Concentration (mg/L)
0.00
0.25
0.50
0.75
1.00
Time (Days)
factor(Route)
oral
factor(Dose)
20

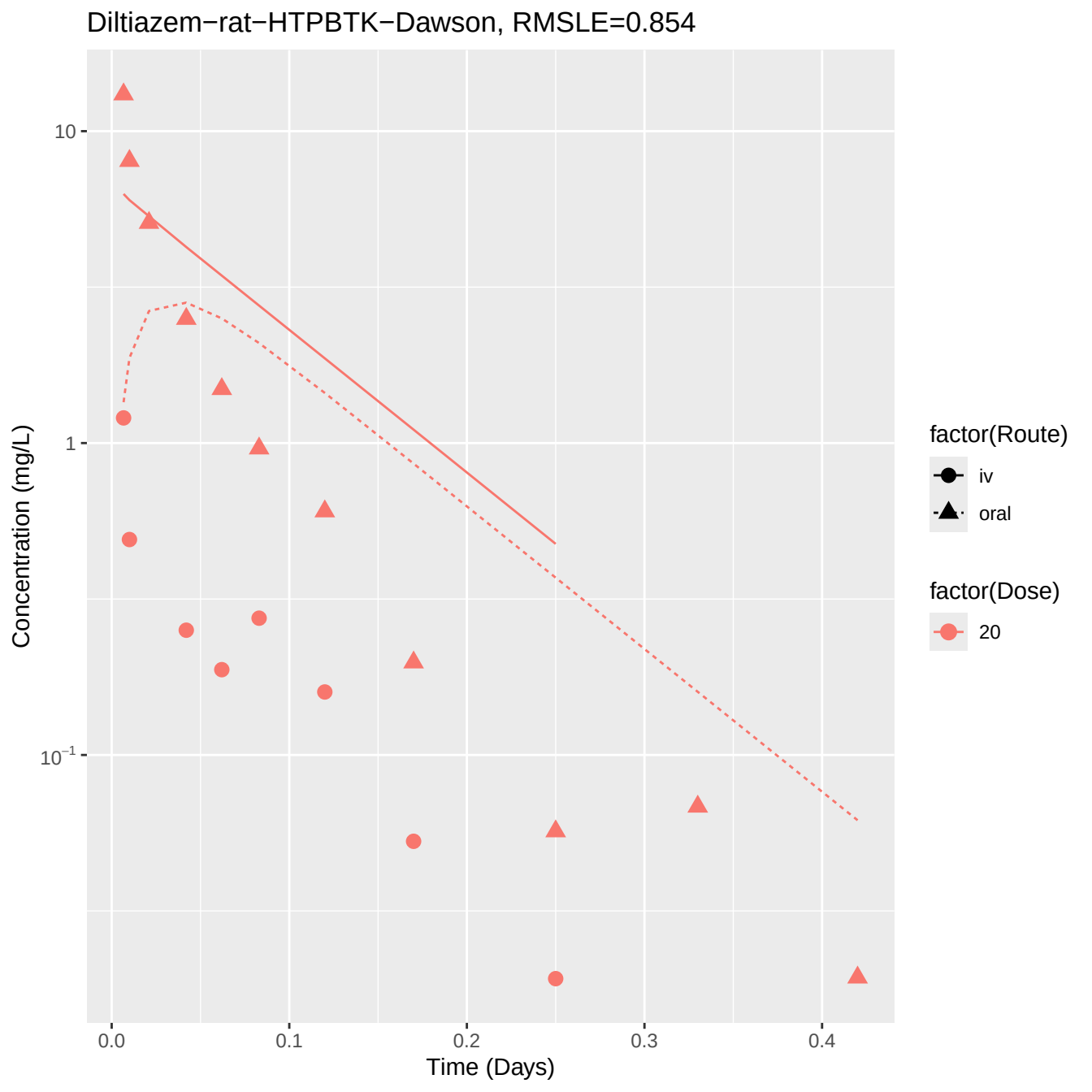

C.I. Solvent Red 1−rat−HTPBTK−Ensemble, RMSLE=0.934
1
10^{-1}
10^{-2}
10^{-3}
Concentration (mg/L)
0.00
0.25
0.50
0.75
1.00
Time (Days)
factor(Route)
oral
factor(Dose)
20

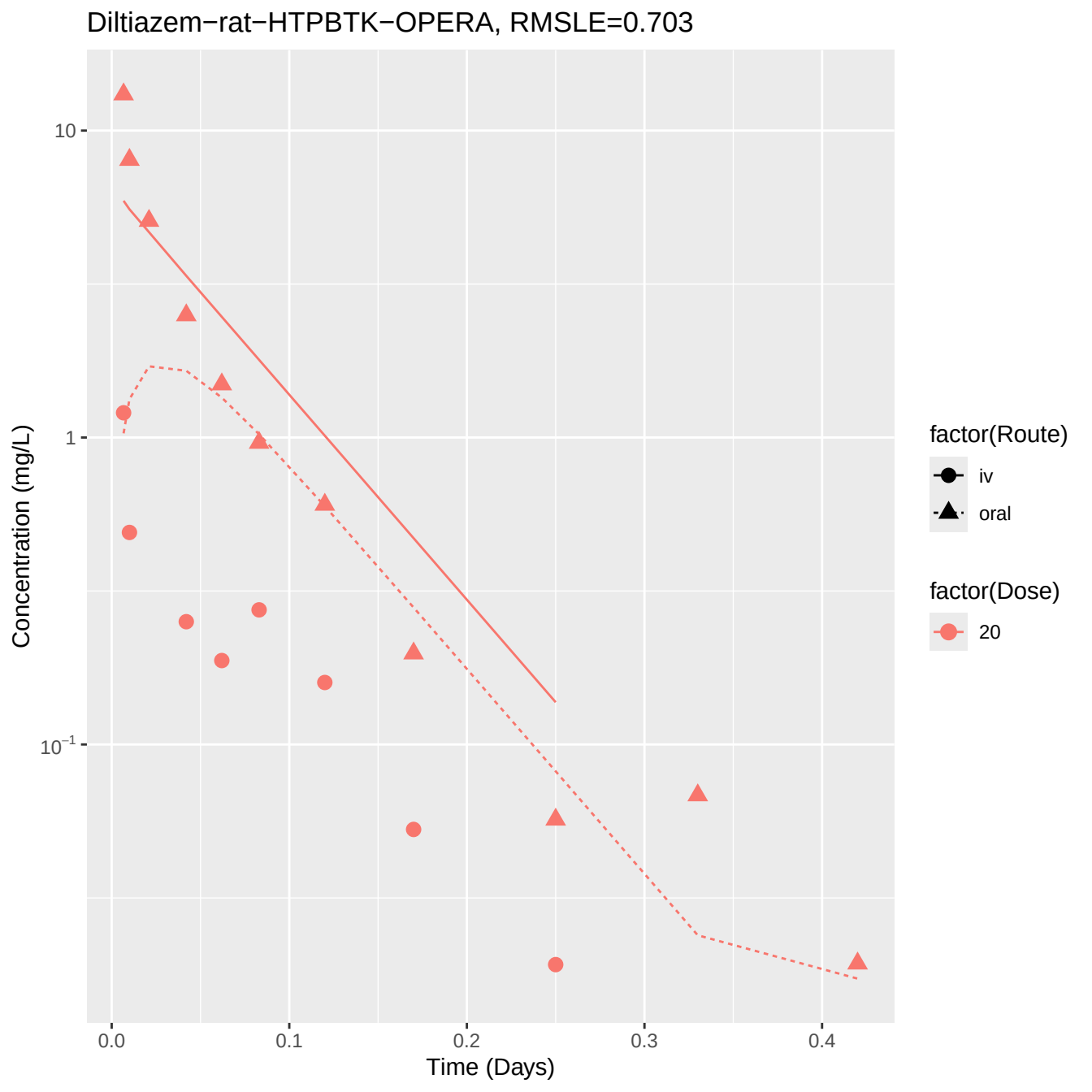

Ephedrine−rat−HTPBTK−Dawson, RMSLE=0.698
Concentration (mg/L)
1
10^{-1}
10^{-2}
0.0
0.1
0.2
0.3
0.4
0.5
Time (Days)
factor(Route)
iv
oral
factor(Dose)
6.25
12.5
25

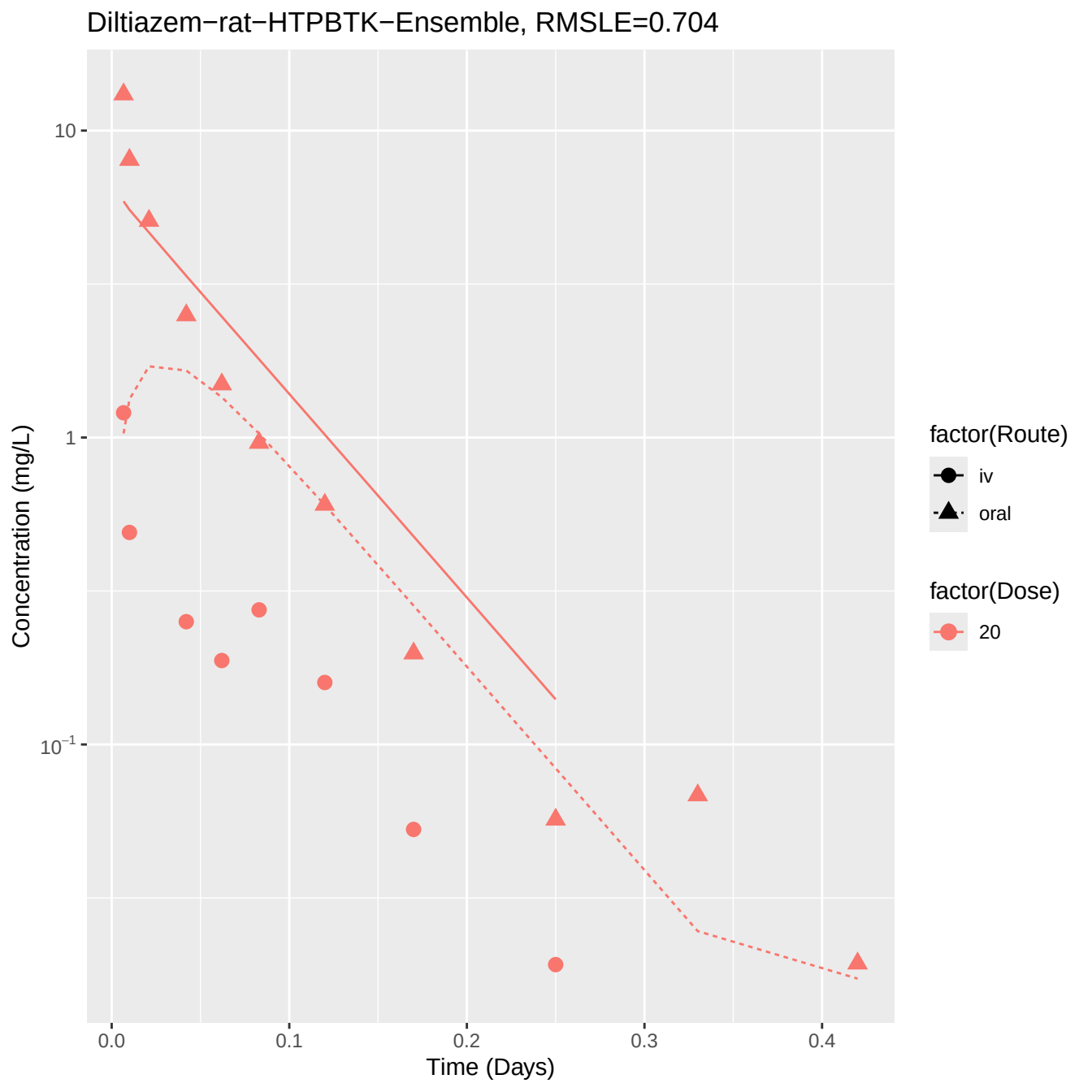

Ephedrine−rat−HTPBTK−OPERA, RMSLE=0.864
Concentration (mg/L)
1
10^{-1}
10^{-2}
0.0
0.1
0.2
0.3
0.4
0.5
Time (Days)
factor(Route)
iv
oral
factor(Dose)
6.25
12.5
25

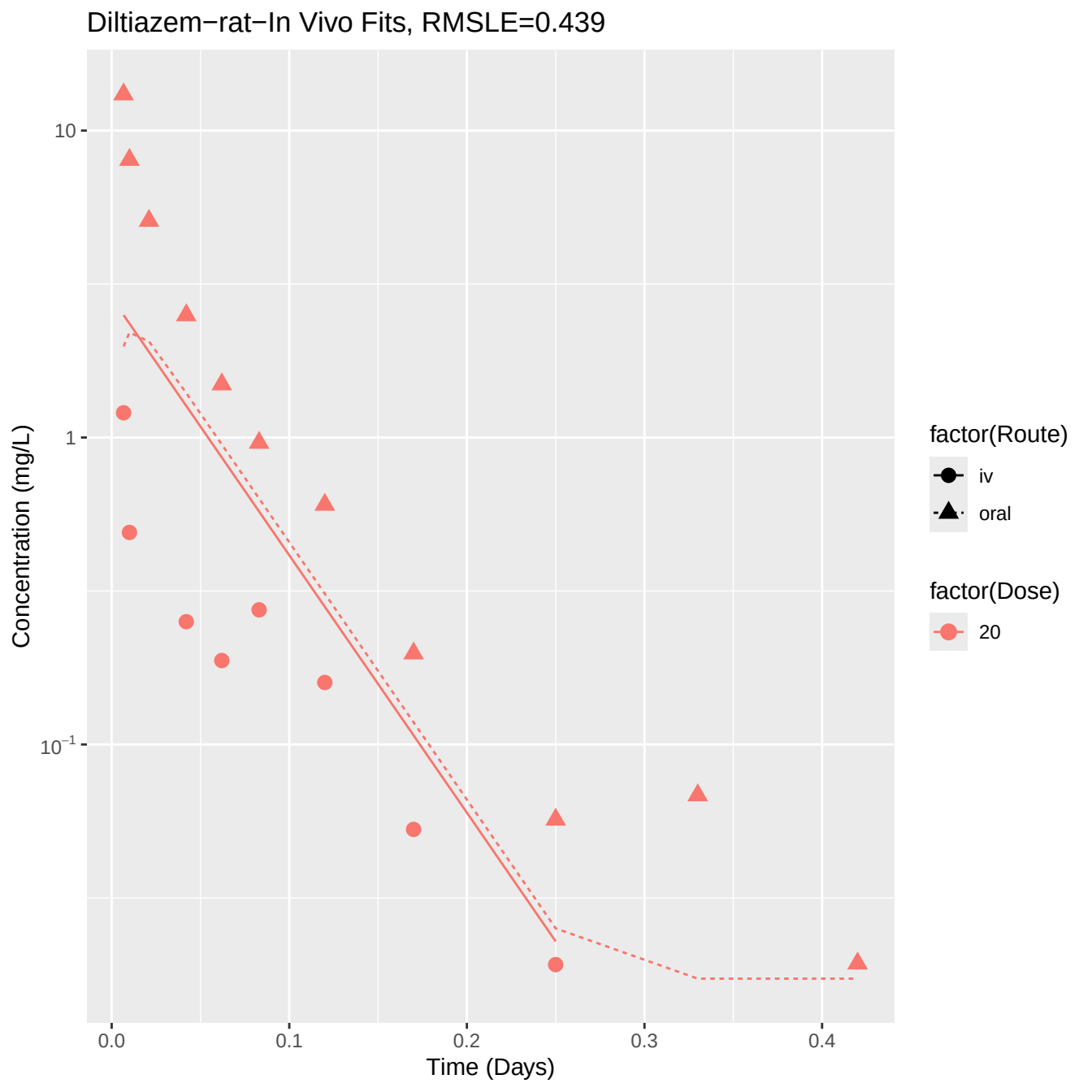

Ephedrine−rat−HTPBTK−Ensemble, RMSLE=0.755
Concentration (mg/L)
1
10^{-1}
10^{-2}
0.0
0.1
0.2
0.3
0.4
0.5
Time (Days)
factor(Route)
iv
oral
factor(Dose)
6.25
12.5
25

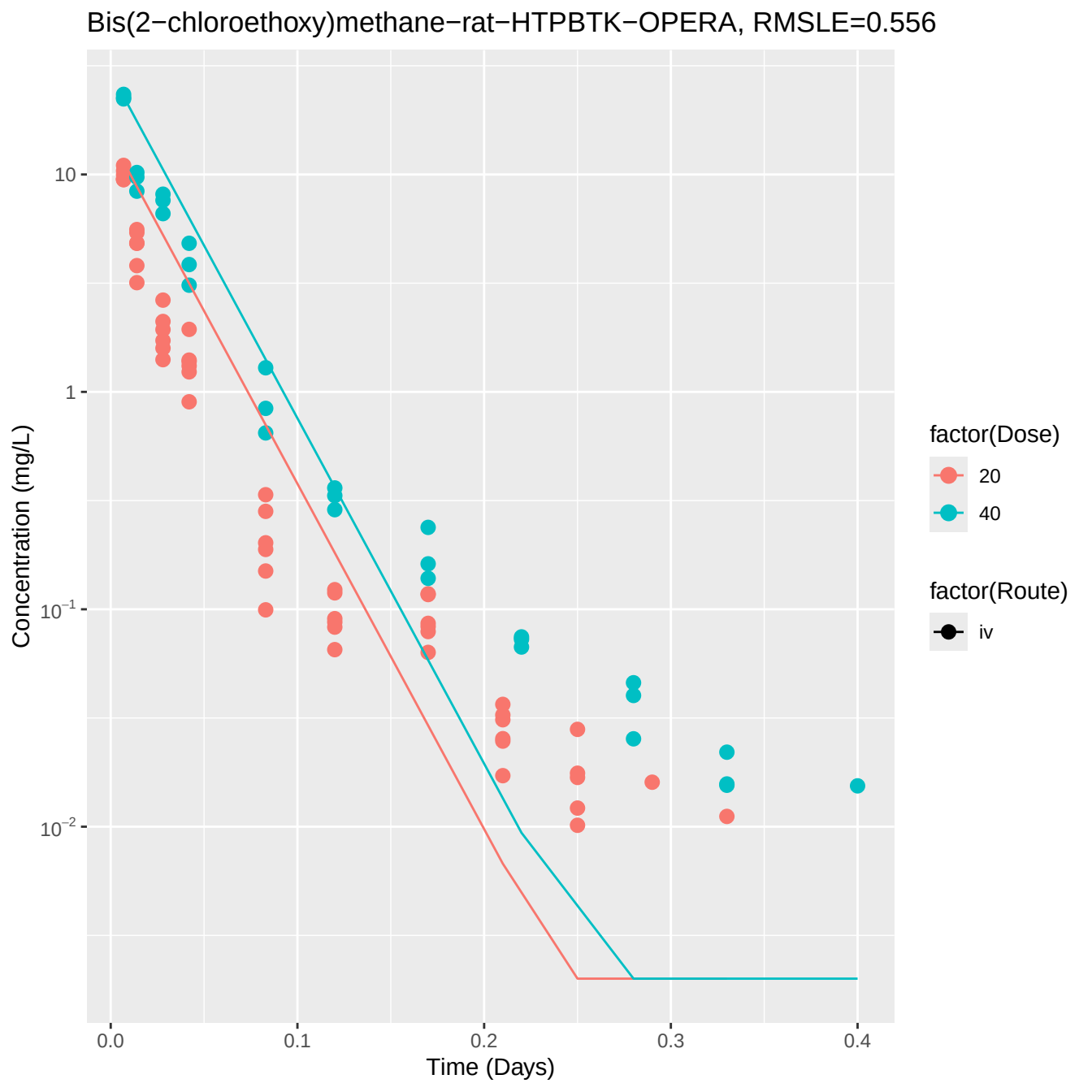

Ephedrine−rat−In Vivo Fits, RMSLE=0.288
Concentration (mg/L)
1
10^{-1}
10^{-2}
0.0
0.1
0.2
0.3
0.4
0.5
Time (Days)
factor(Route)
iv
oral
factor(Dose)
6.25
12.5
25

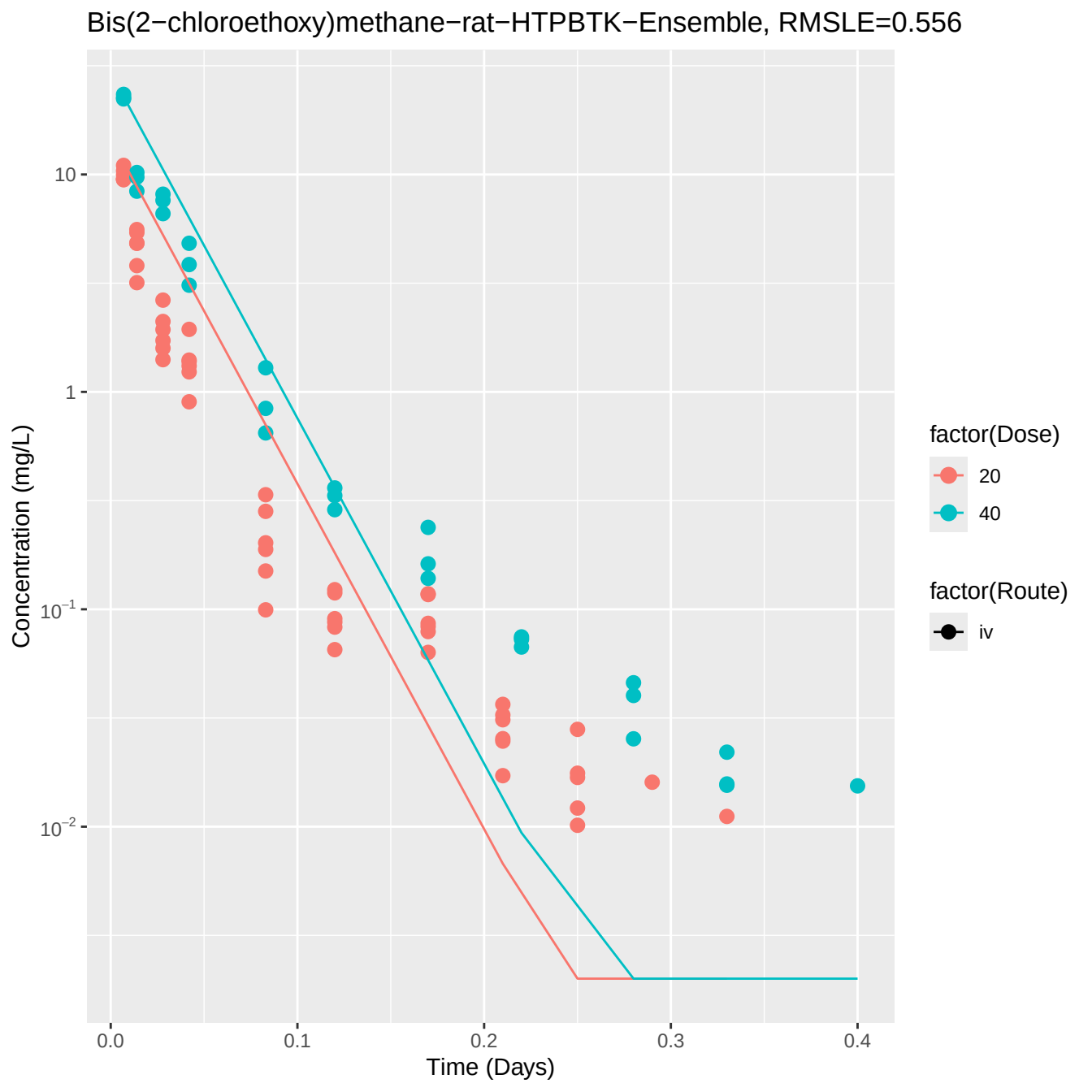

Diltiazem−rat−HTPBTK−Dawson, RMSLE=0.854
Concentration (mg/L)
10
1
10^{-1}
0.0
0.1
0.2
0.3
0.4
Time (Days)
factor(Route)
iv
oral
factor(Dose)
20

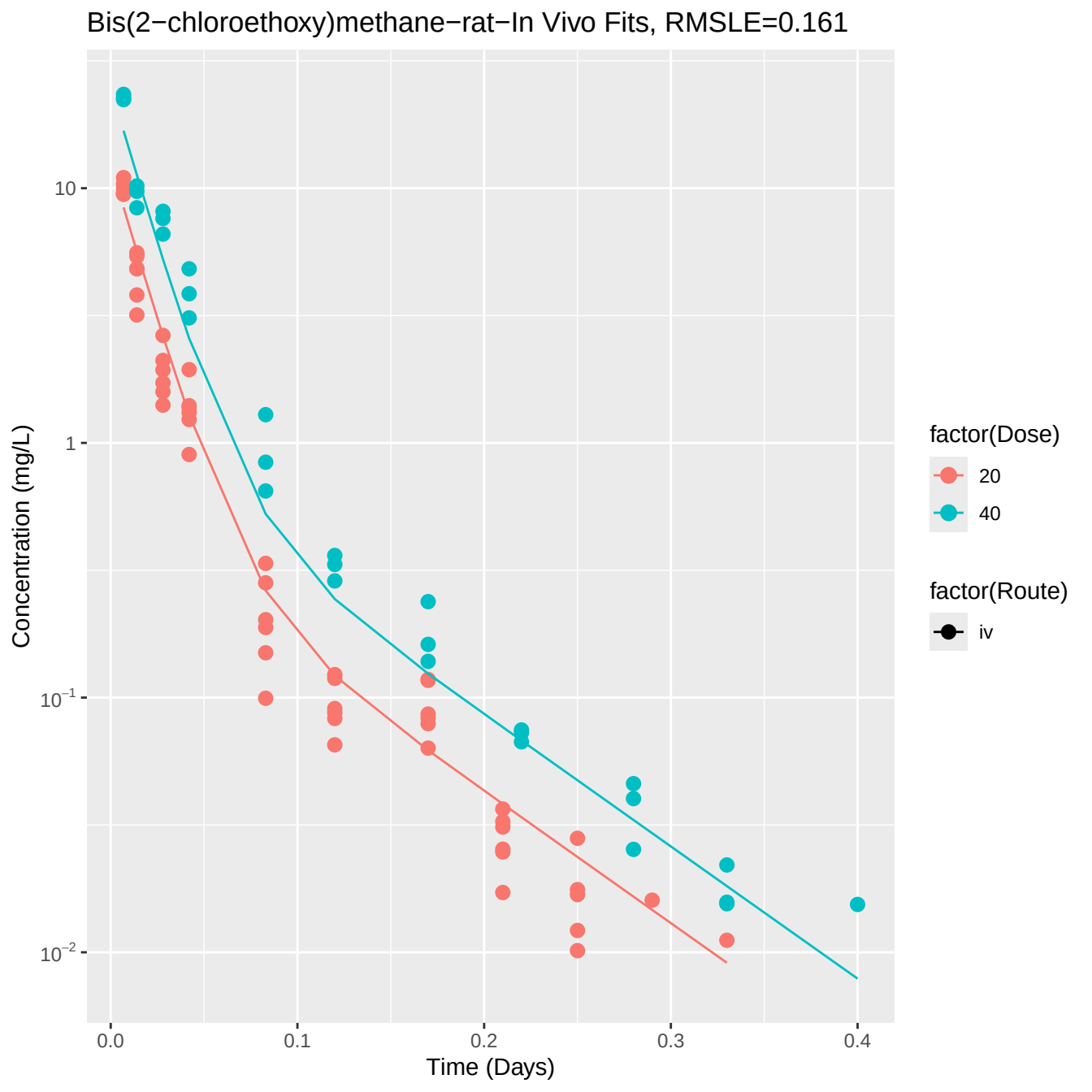

Diltiazem−rat−HTPBTK−OPERA, RMSLE=0.703
Concentration (mg/L)
10
1
10^{-1}
0.0
0.1
0.2
0.3
0.4
Time (Days)
factor(Route)
iv
oral
factor(Dose)
20

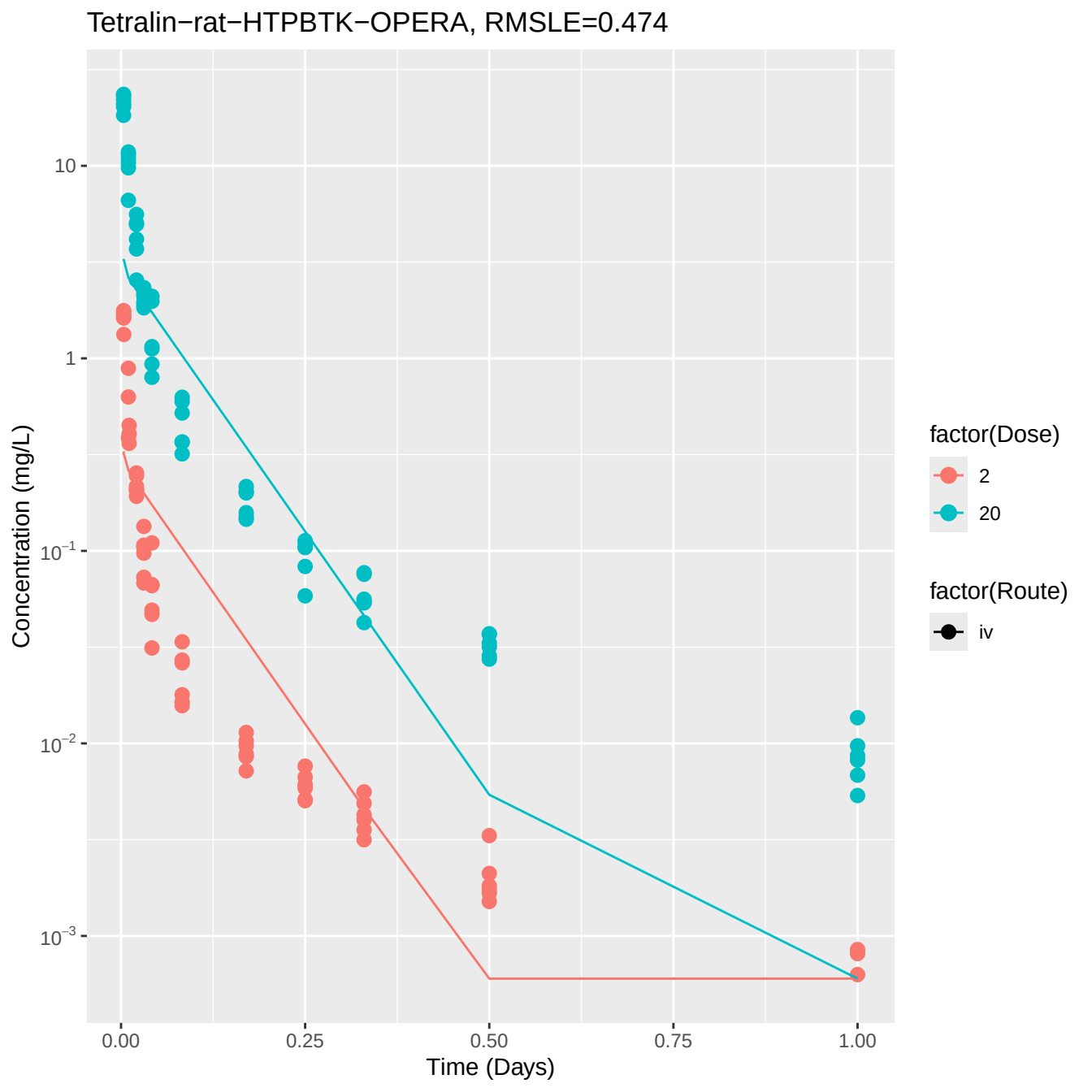

Diltiazem−rat−HTPBTK−Ensemble, RMSLE=0.704
Concentration (mg/L)
10
1
10^{-1}
0.0
0.1
0.2
0.3
0.4
Time (Days)
factor(Route)
iv
oral
factor(Dose)
20

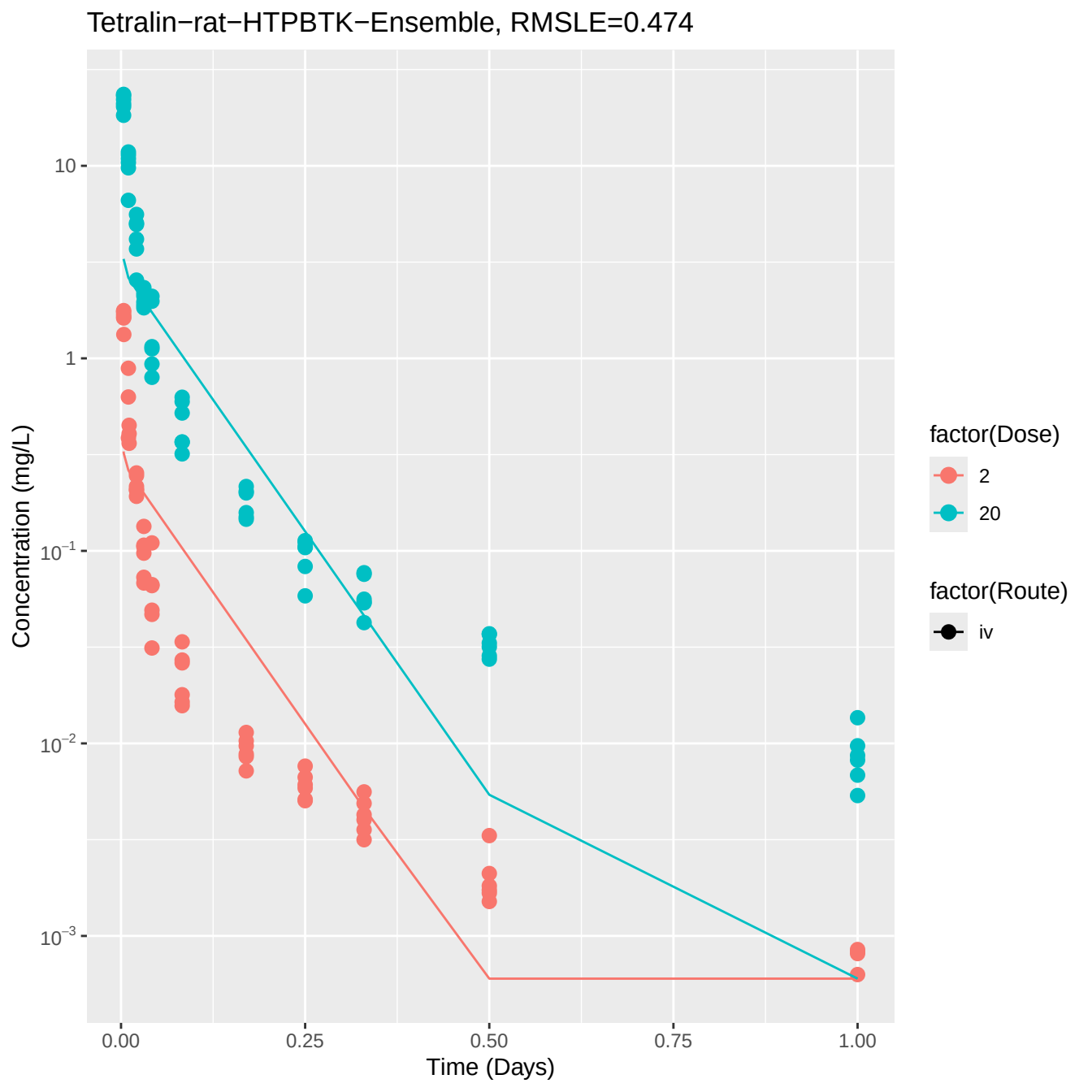

Diltiazem−rat−In Vivo Fits, RMSLE=0.439
10
1
10^{-1}
Concentration (mg/L)
0.0
0.1
0.2
0.3
0.4
Time (Days)
factor(Route)
iv
oral
factor(Dose)
20

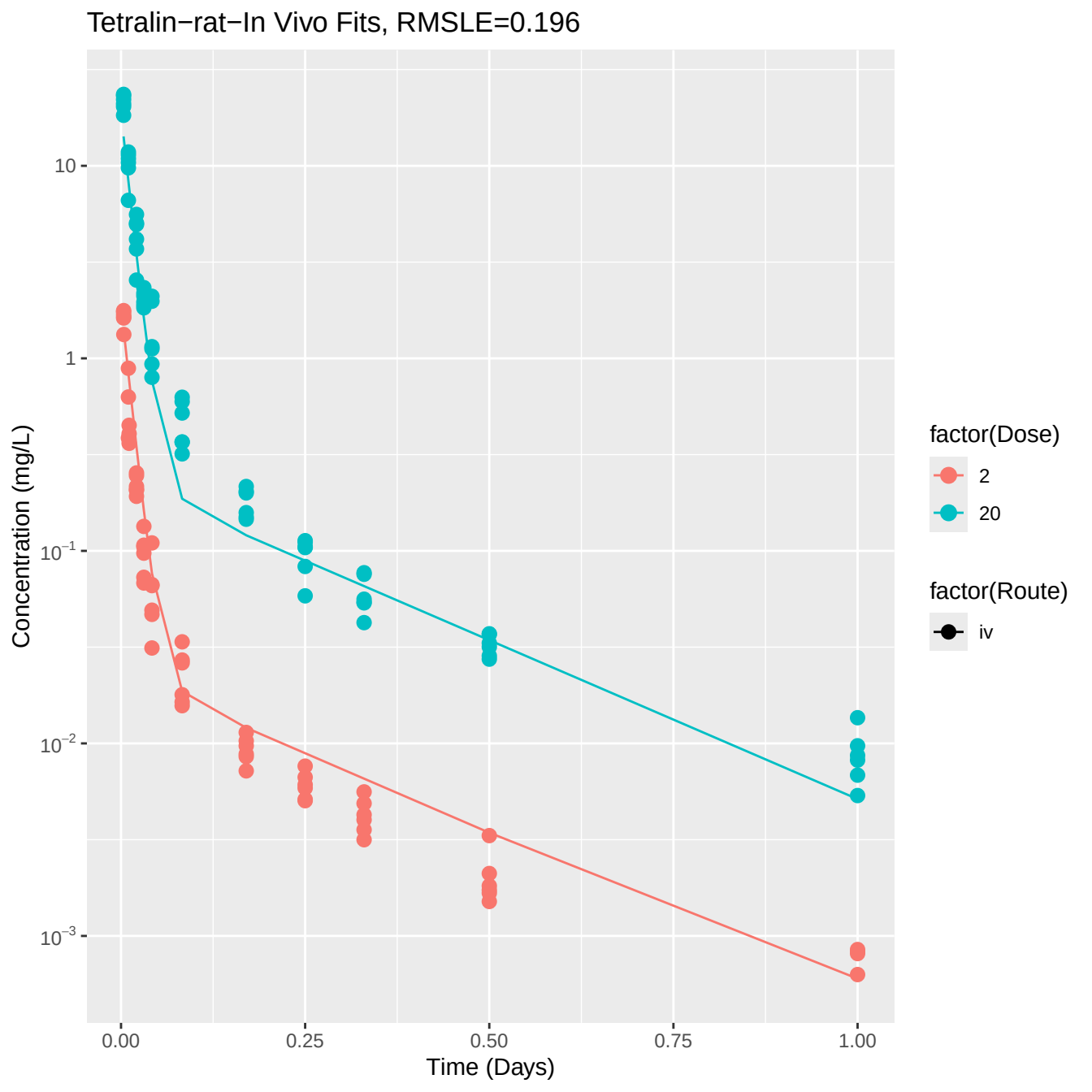

Bis(2−chloroethoxy)methane−rat−HTPBTK−OPERA, RMSLE=0.556
Concentration (mg/L)
10
1
10^{-1}
10^{-2}
0.0
0.1
0.2
0.3
0.4
Time (Days)
factor(Dose)
20
40
factor(Route)
iv

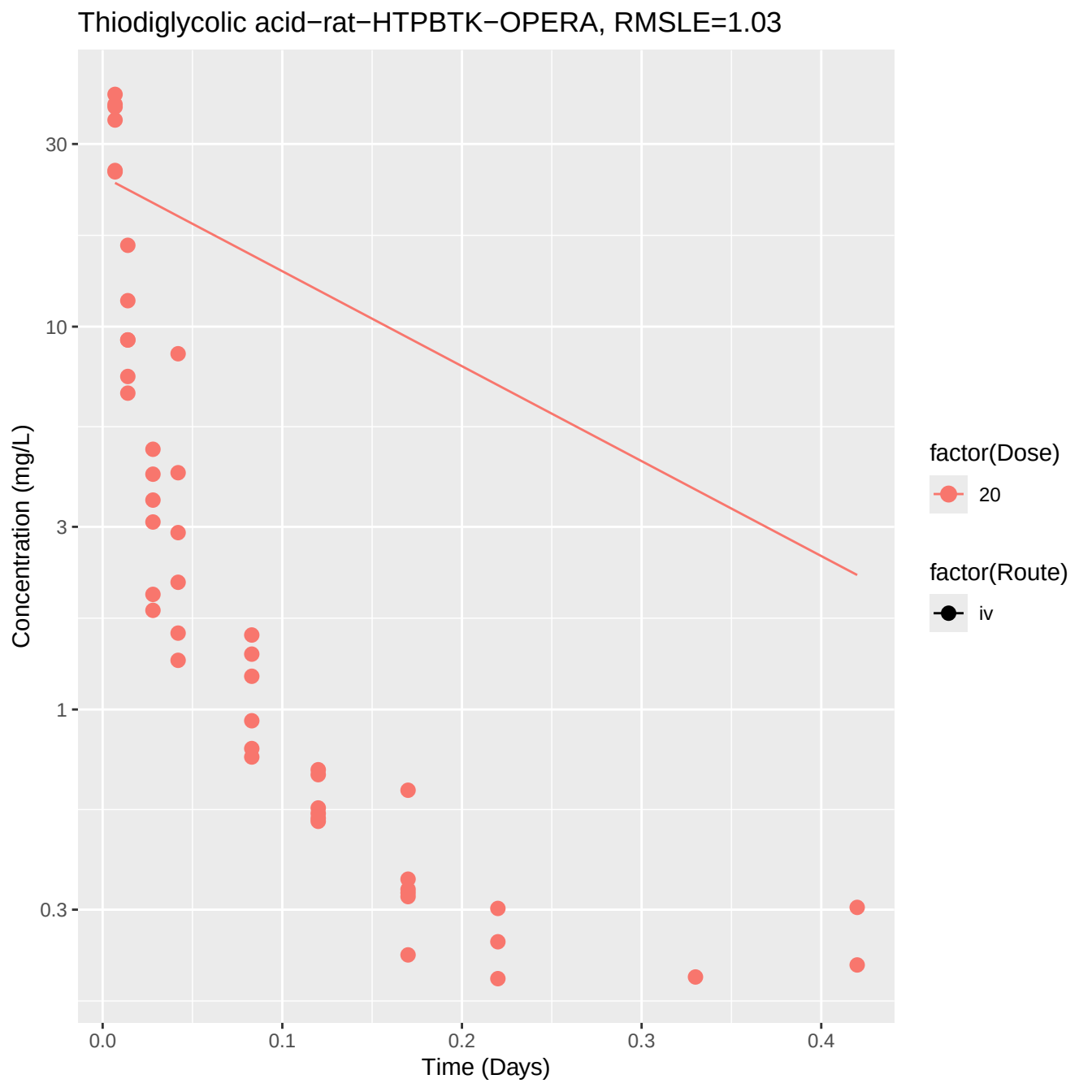

Bis(2−chloroethoxy)methane−rat−HTPBTK−Ensemble, RMSLE=0.556
Concentration (mg/L)
10
1
10^{-1}
10^{-2}
0.0
0.1
0.2
0.3
0.4
Time (Days)
factor(Dose)
20
40
factor(Route)
iv

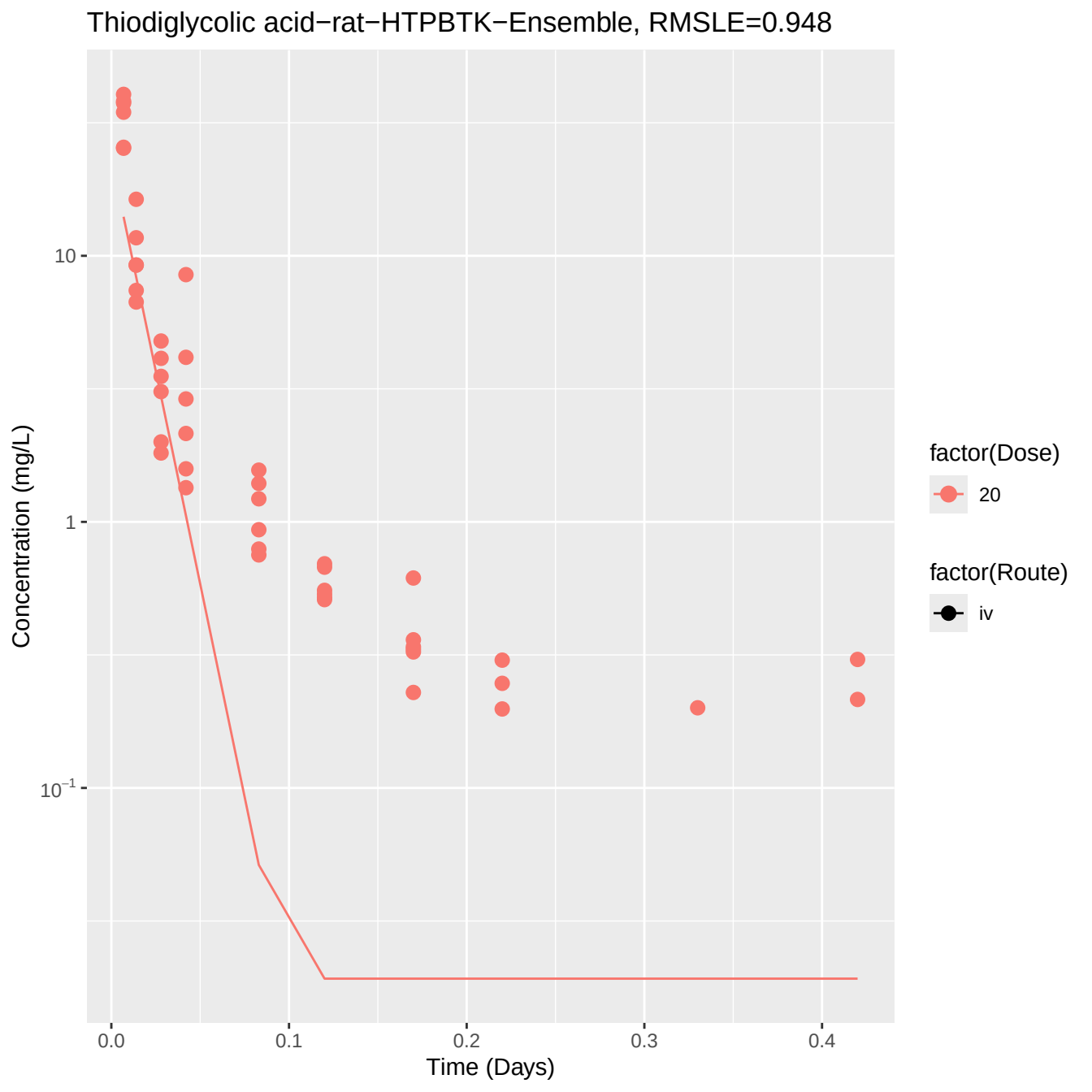

Bis(2−chloroethoxy)methane−rat−In Vivo Fits, RMSLE=0.161
Concentration (mg/L)
10
1
10^{-1}
10^{-2}
0.0
0.1
0.2
0.3
0.4
Time (Days)
factor(Dose)
20
40
factor(Route)
iv

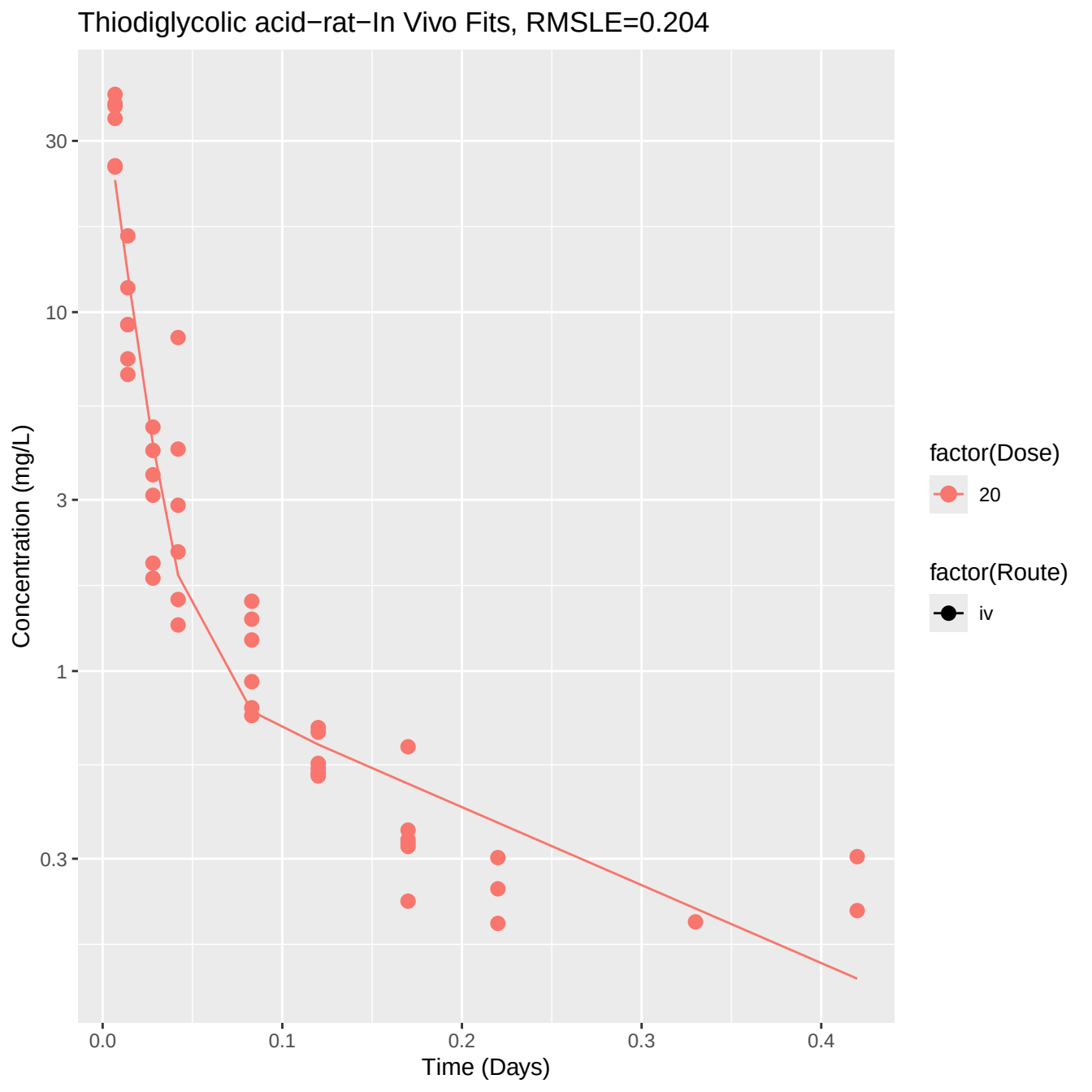

Tetralin−rat−HTPBTK−OPERA, RMSLE=0.474
Concentration (mg/L)
10
1
10^{-1}
10^{-2}
10^{-3}
0.00
0.25
0.50
0.75
1.00
Time (Days)
factor(Dose)
2
20
factor(Route)
iv

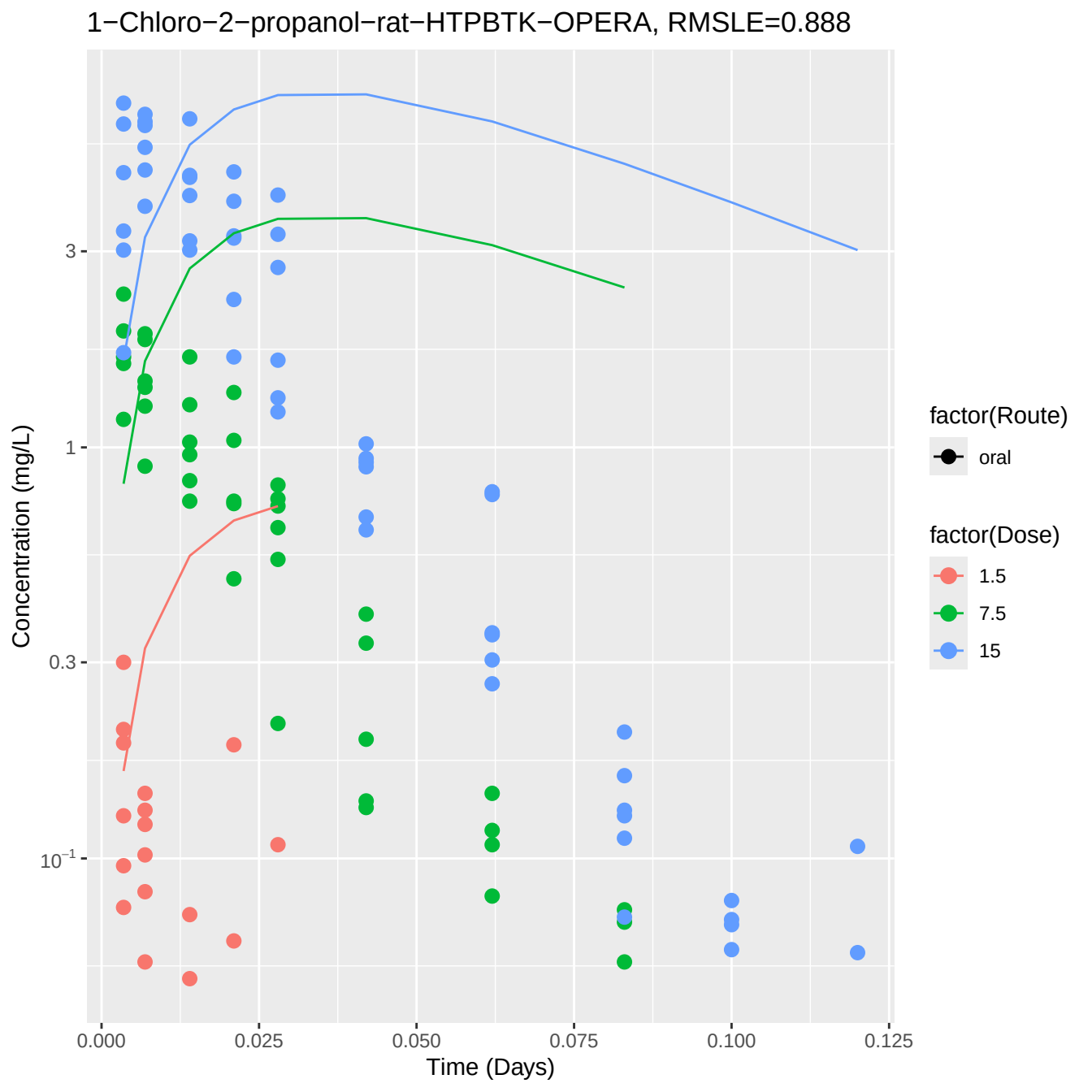

Tetralin−rat−HTPBTK−Ensemble, RMSLE=0.474
Concentration (mg/L)
10
1
10^{-1}
10^{-2}
10^{-3}
0.00
0.25
0.50
0.75
1.00
Time (Days)
factor(Dose)
2
20
factor(Route)
iv

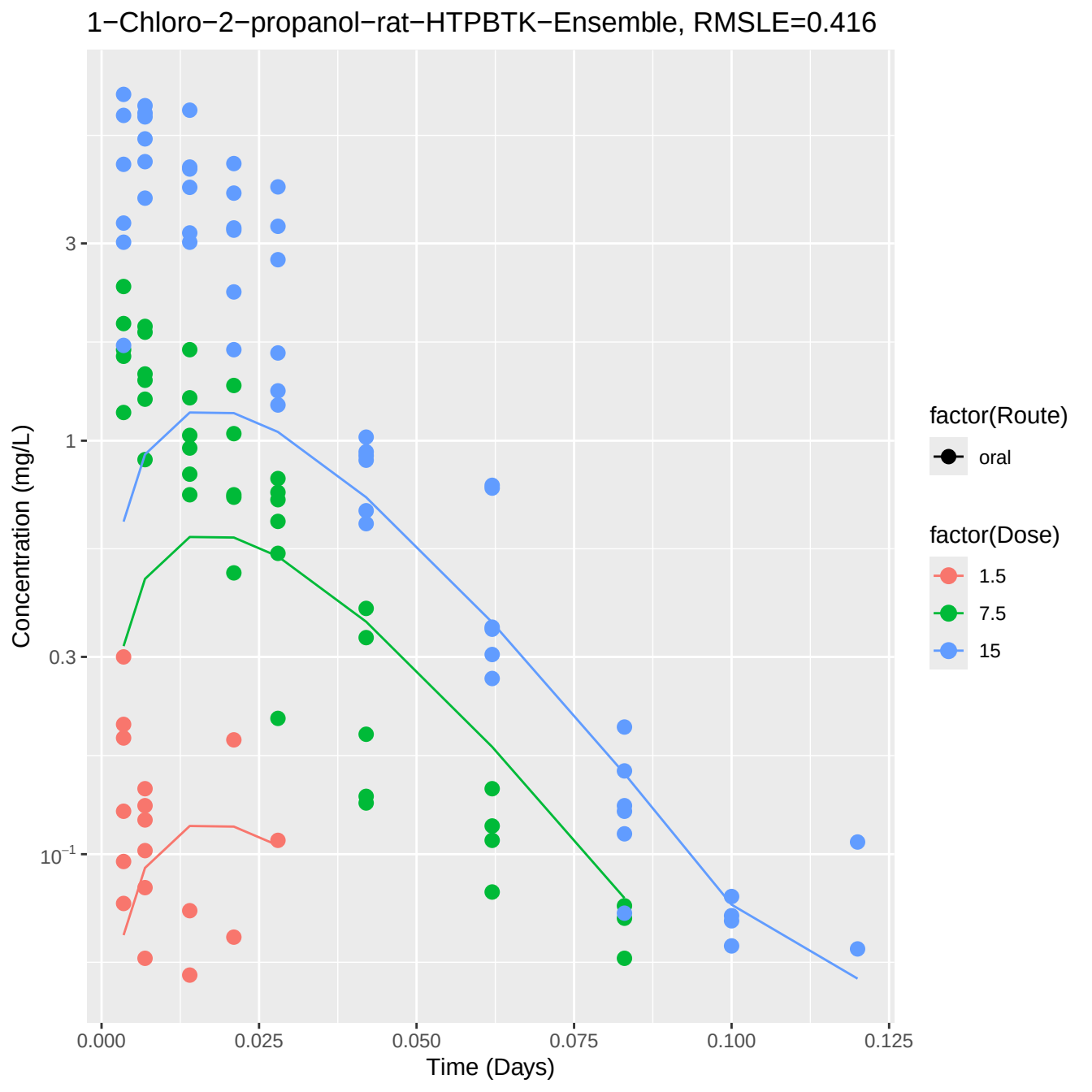

Tetralin−rat−In Vivo Fits, RMSLE=0.196
Concentration (mg/L)
10
1
10^{-1}
10^{-2}
10^{-3}
0.00
0.25
0.50
0.75
1.00
Time (Days)
factor(Dose)
2
20
factor(Route)
iv

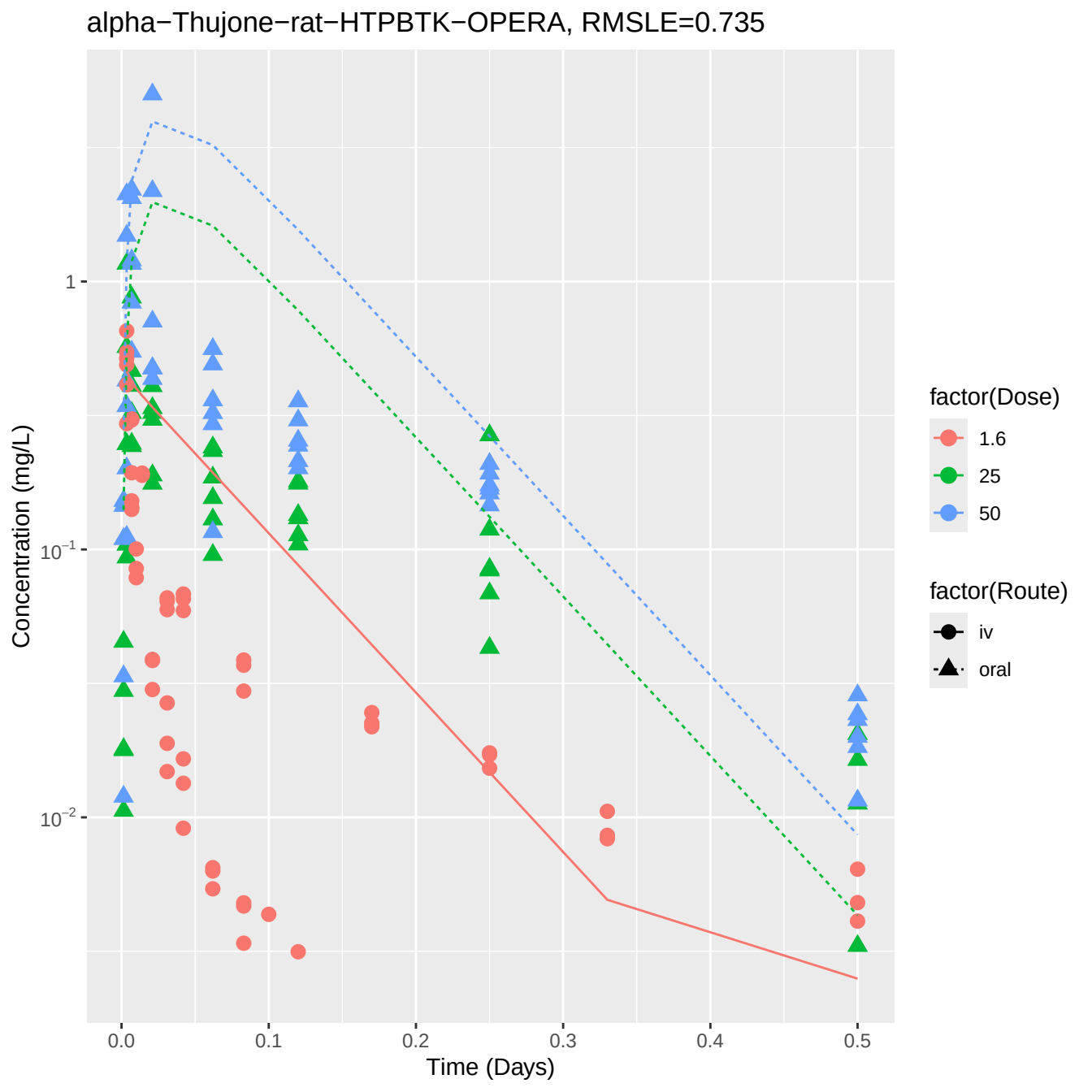

Thiodiglycolic acid−rat−HTPBTK−OPERA, RMSLE=1.03
Concentration (mg/L)
30
10
3
1
0.3
0.0
0.1
0.2
0.3
0.4
Time (Days)
factor(Dose)
20
factor(Route)
iv

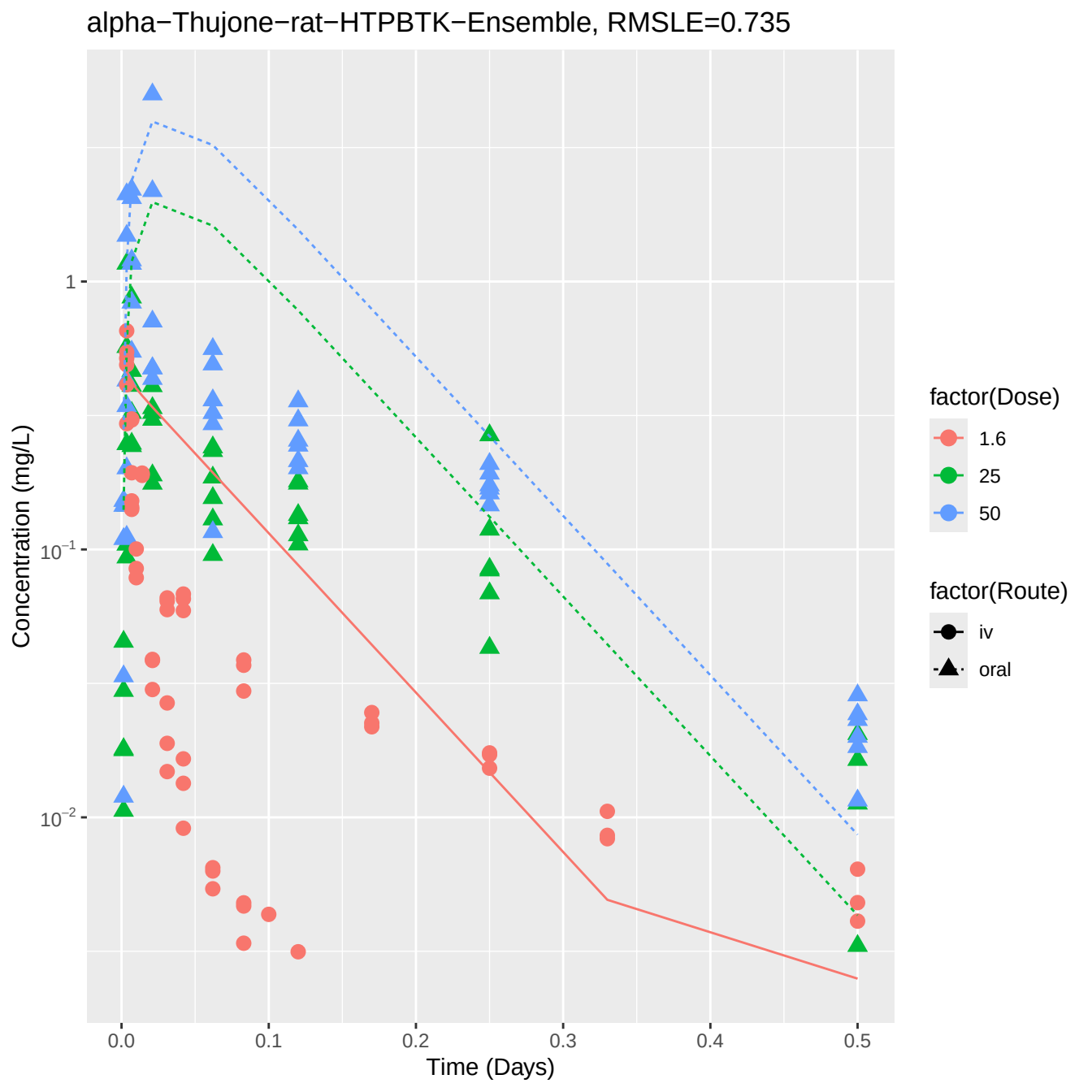

Thiodiglycolic acid−rat−HTPBTK−Ensemble, RMSLE=0.948
Concentration (mg/L)
10
1
10^{-1}
0.0
0.1
0.2
0.3
0.4
Time (Days)
factor(Dose)
20
factor(Route)
iv

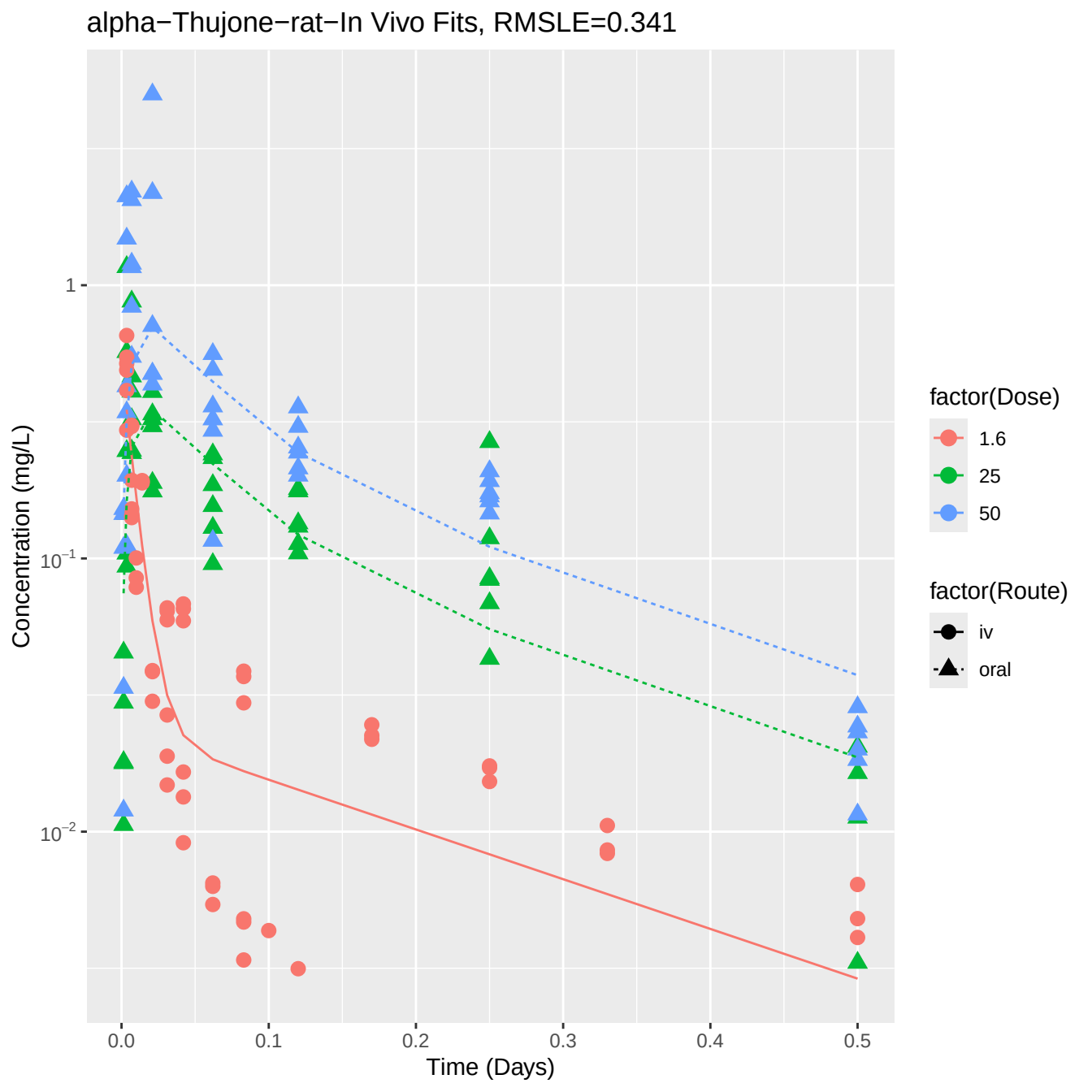

Thiodiglycolic acid−rat−In Vivo Fits, RMSLE=0.204
Concentration (mg/L)
30
10
3
1
0.3
0.0
0.1
0.2
0.3
0.4
Time (Days)
factor(Dose)
20
factor(Route)
iv

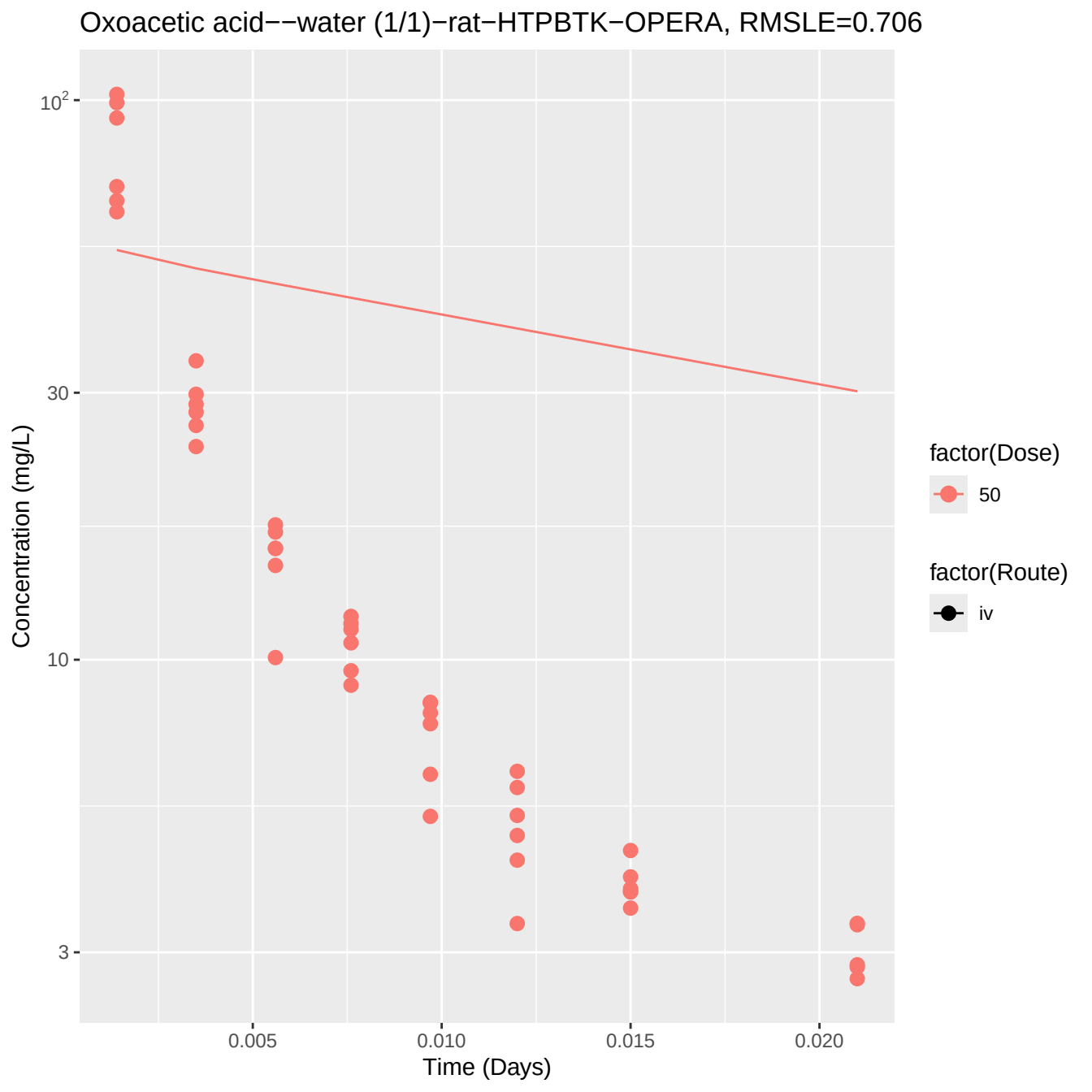

1−Chloro−2−propanol−rat−HTPBTK−OPERA, RMSLE=0.888
Concentration (mg/L)
3
1
0.3
10^{-1}
0.000
0.025
0.050
0.075
0.100
0.125
Time (Days)
factor(Route)
oral
factor(Dose)
1.5
7.5
15

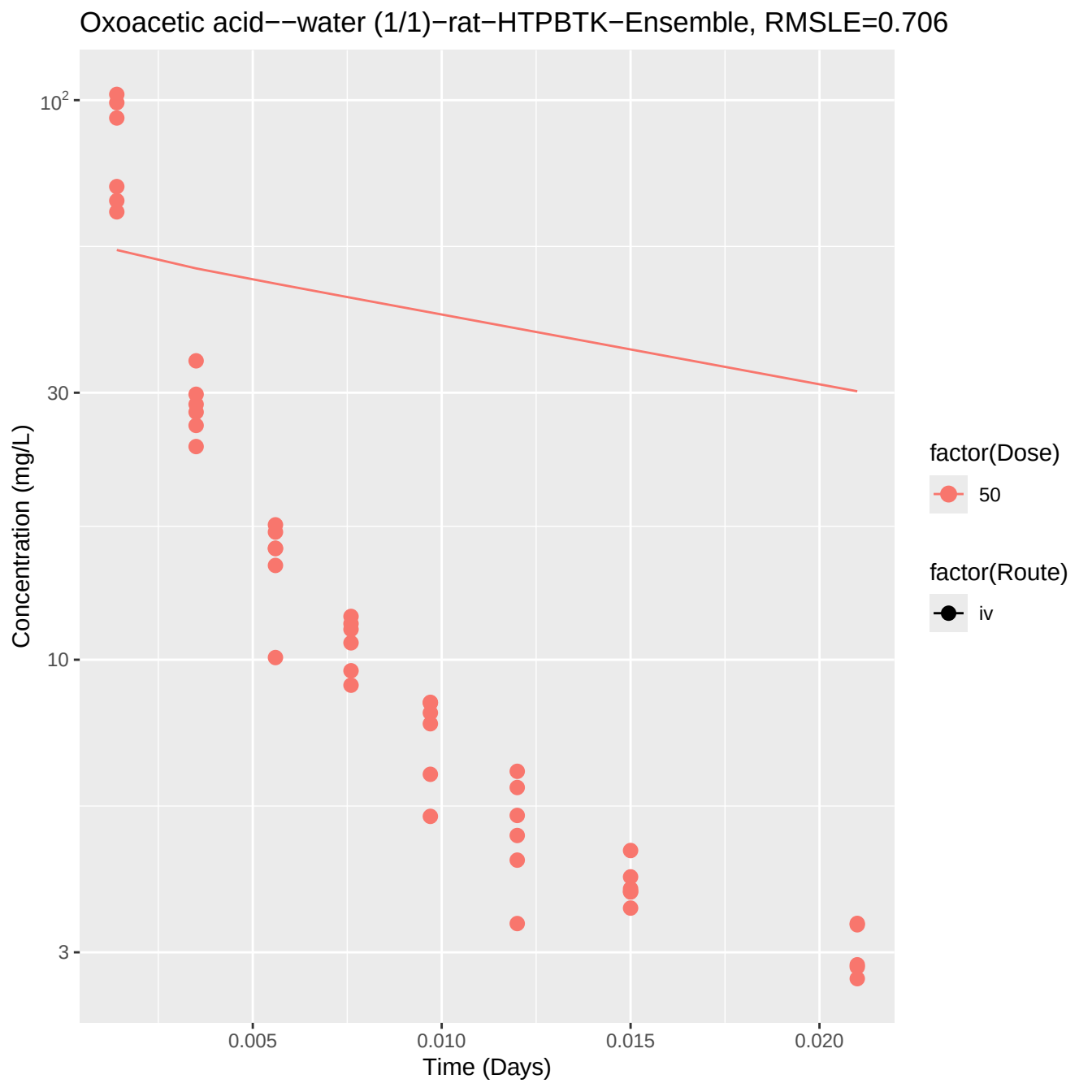

1−Chloro−2−propanol−rat−HTPBTK−Ensemble, RMSLE=0.416
Concentration (mg/L)
3
1
0.3
10−1
0.000
0.025
0.050
0.075
0.100
0.125
Time (Days)
factor(Route)
oral
factor(Dose)
1.5
7.5
15

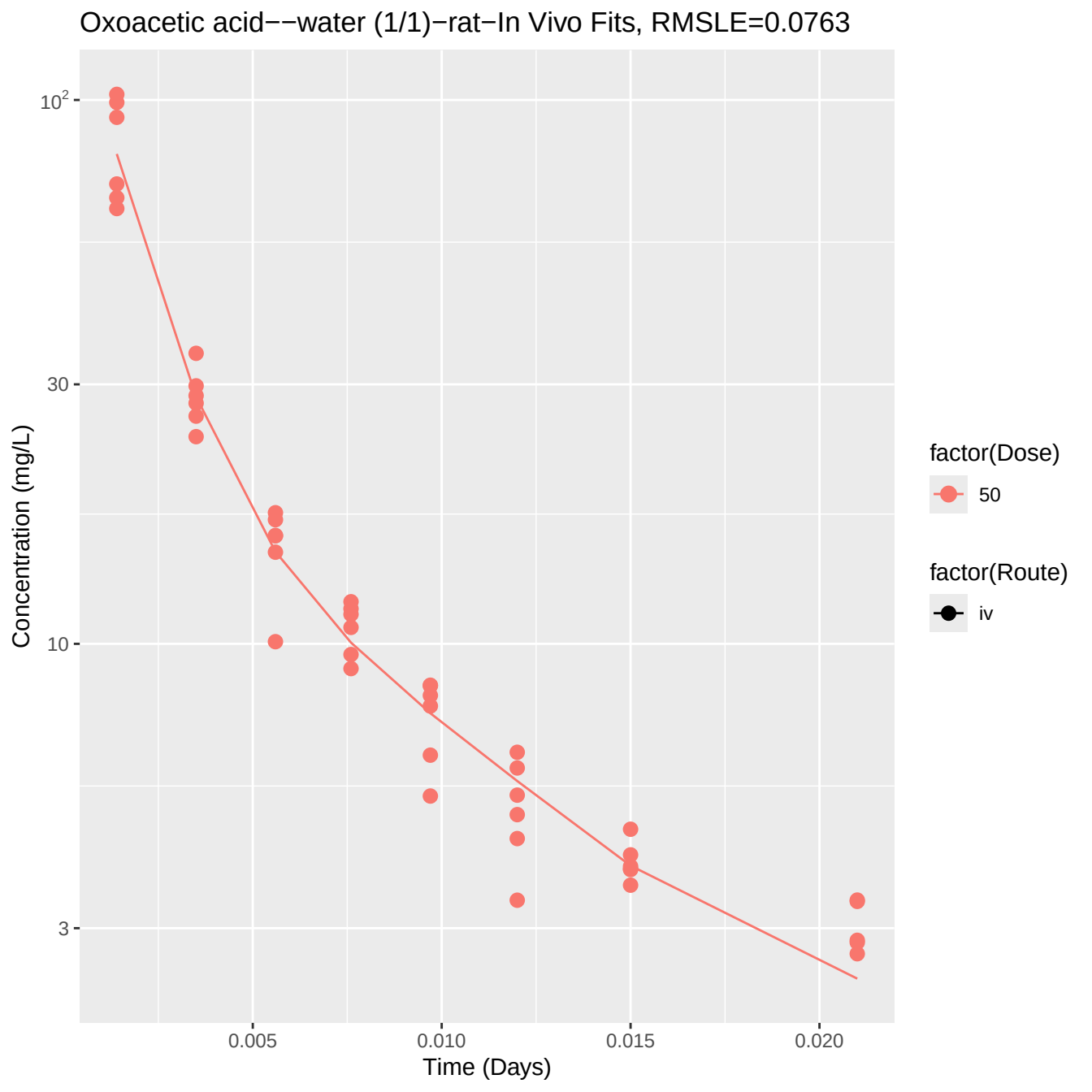

alpha−Thujone−rat−HTPBTK−OPERA, RMSLE=0.735
Concentration (mg/L)
1
10^{-1}
10^{-2}
0.0
0.1
0.2
0.3
0.4
0.5
Time (Days)
factor(Dose)
1.6
25
50
factor(Route)
iv
oral

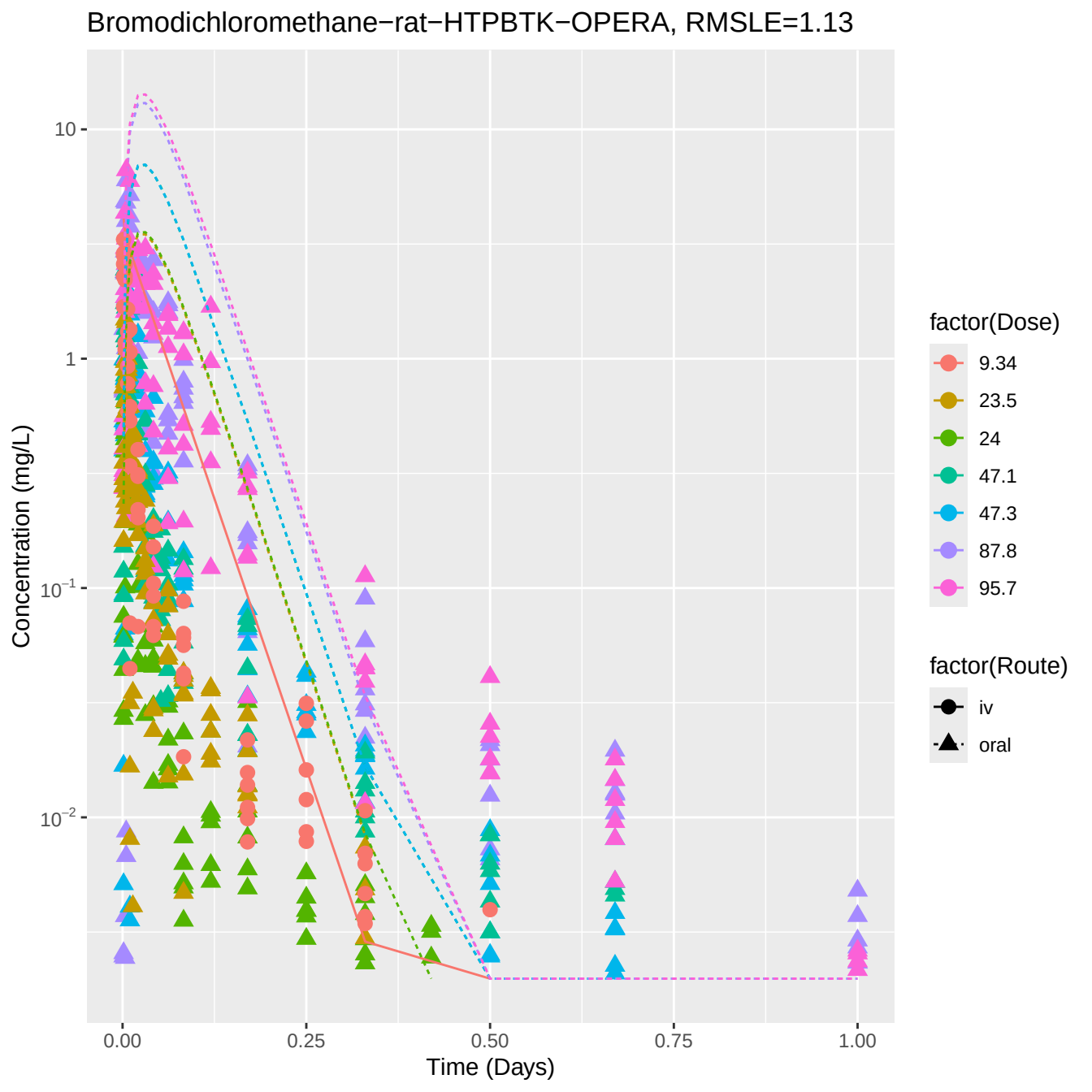

alpha−Thujone−rat−HTPBTK−Ensemble, RMSLE=0.735
Concentration (mg/L)
1
10^{-1}
10^{-2}
0.0
0.1
0.2
0.3
0.4
0.5
Time (Days)
factor(Dose)
1.6
25
50
factor(Route)
iv
oral

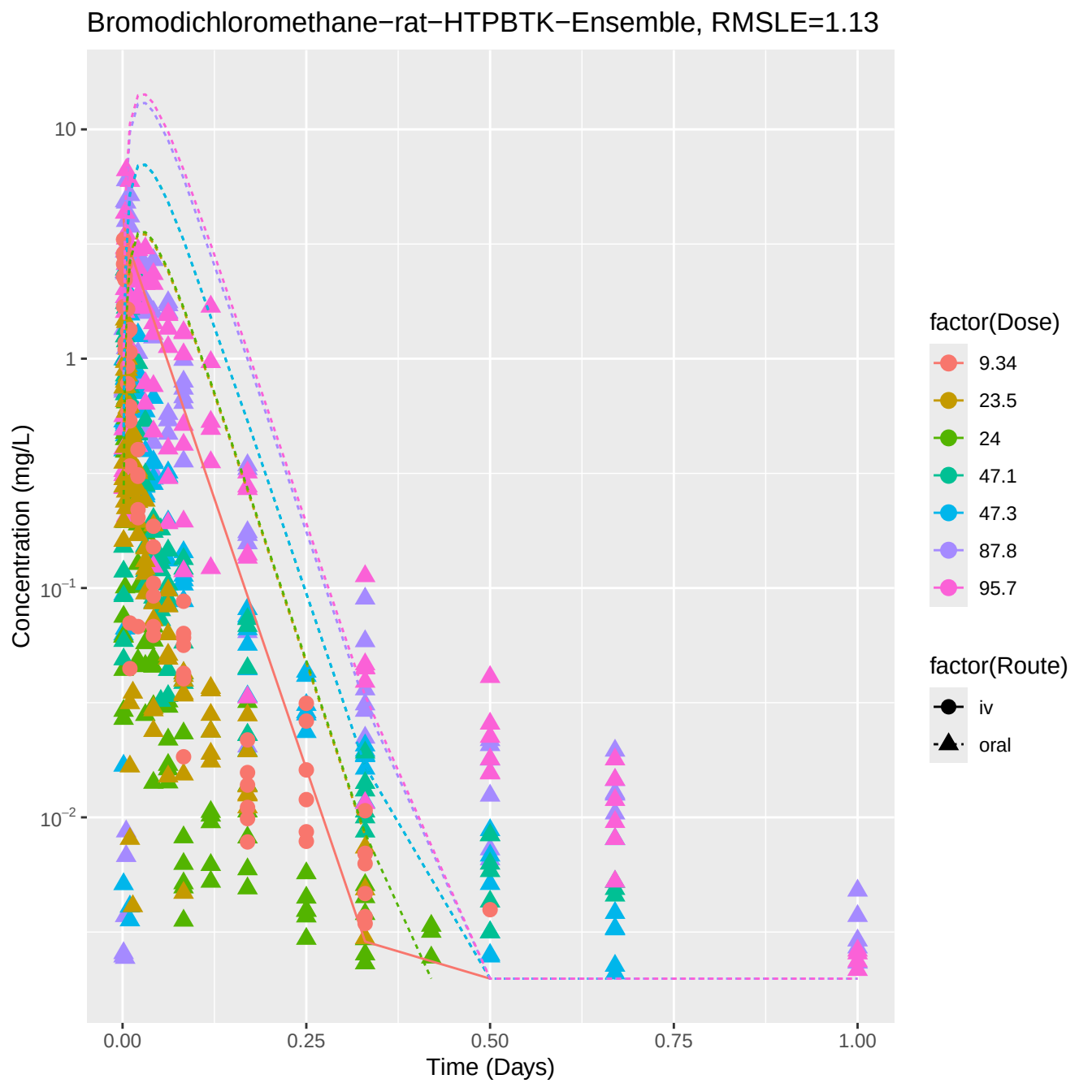

alpha−Thujone−rat−In Vivo Fits, RMSLE=0.341
Concentration (mg/L)
1
10^{-1}
10^{-2}
0.0
0.1
0.2
0.3
0.4
0.5
Time (Days)
factor(Dose)
1.6
25
50
factor(Route)
iv
oral

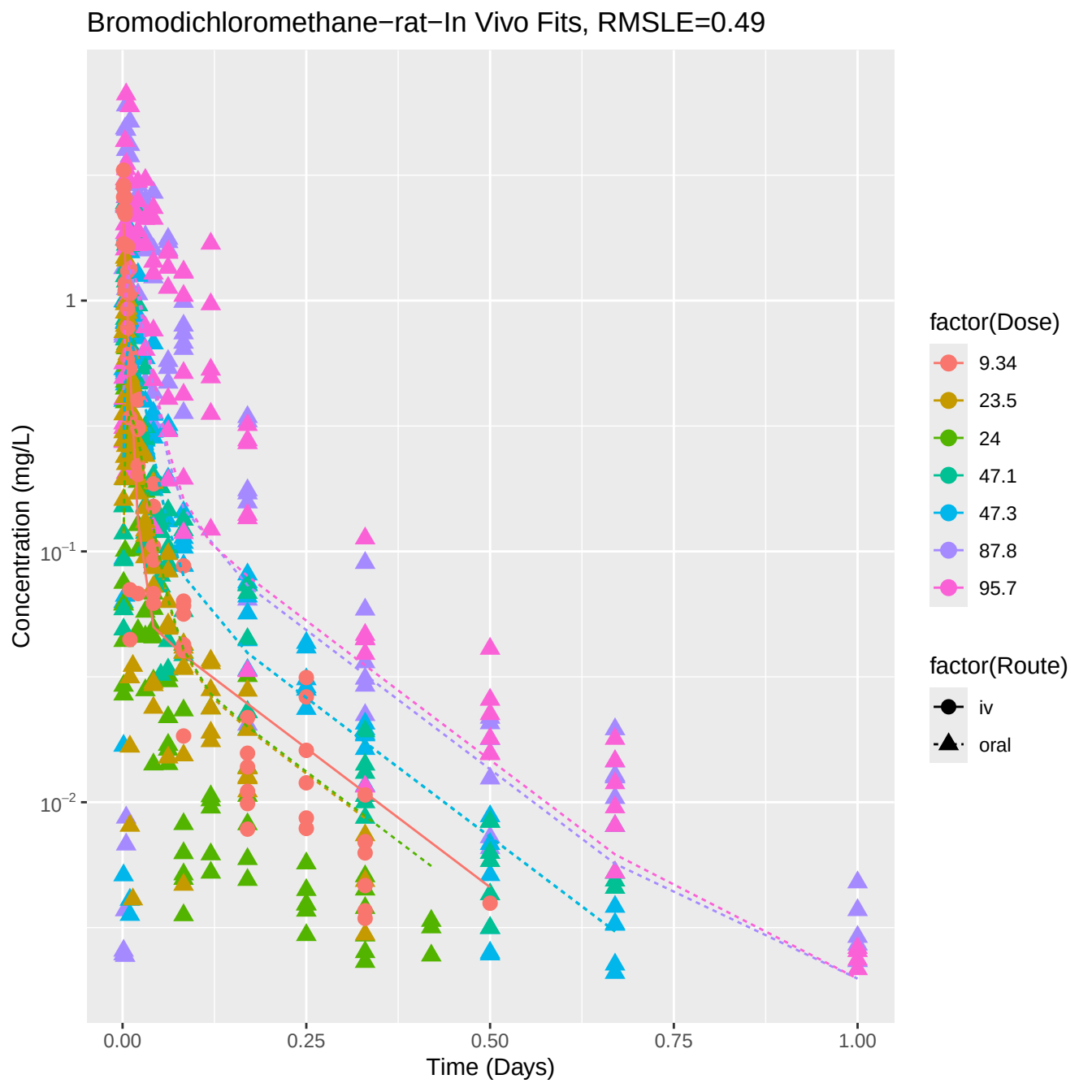

Oxoacetic acid−−water (1/1)−rat−HTPBTK−OPERA, RMSLE=0.706
Concentration (mg/L)
10^2
30
10
3
0.005
0.010
0.015
0.020
Time (Days)
factor(Dose)
50
factor(Route)
iv

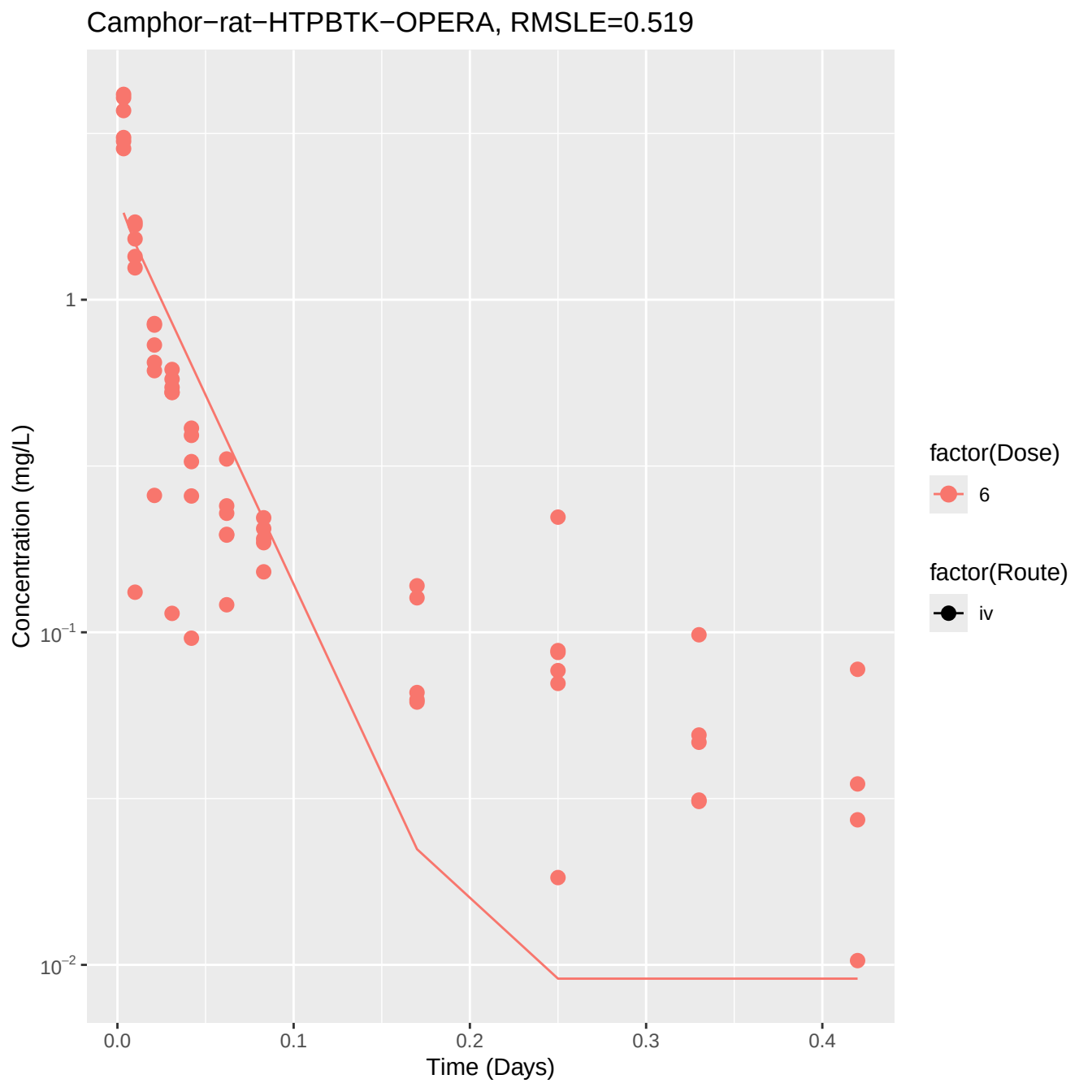

Oxoacetic acid−−water (1/1)−rat−HTPBTK−Ensemble, RMSLE=0.706
Concentration (mg/L)
10^2
30
10
3
0.005
0.010
0.015
0.020
Time (Days)
factor(Dose)
50
factor(Route)
iv

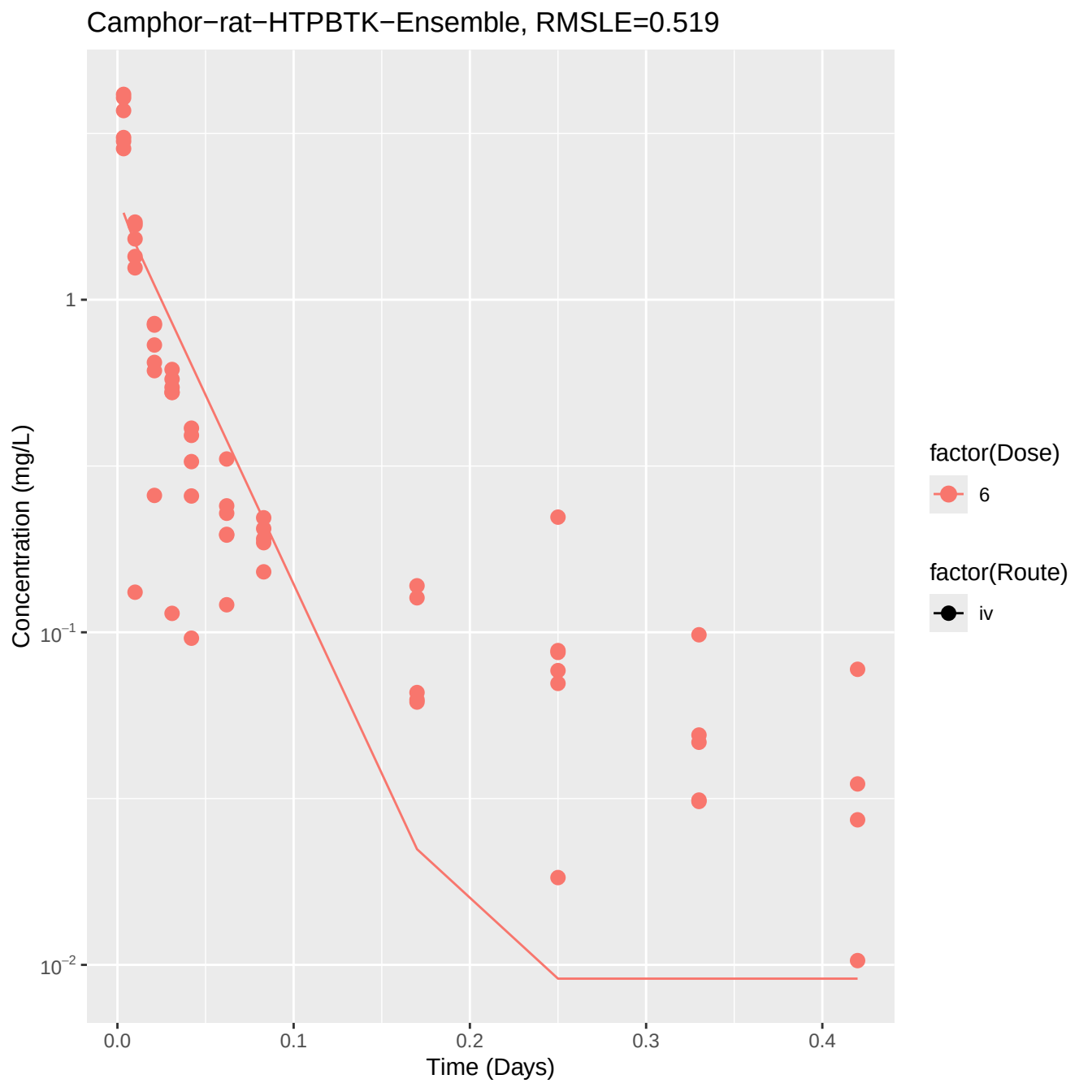

Oxoacetic acid−−water (1/1)−rat−In Vivo Fits, RMSLE=0.0763
Concentration (mg/L)
10^2
30
10
3
0.005
0.010
0.015
0.020
Time (Days)
factor(Dose)
50
factor(Route)
iv

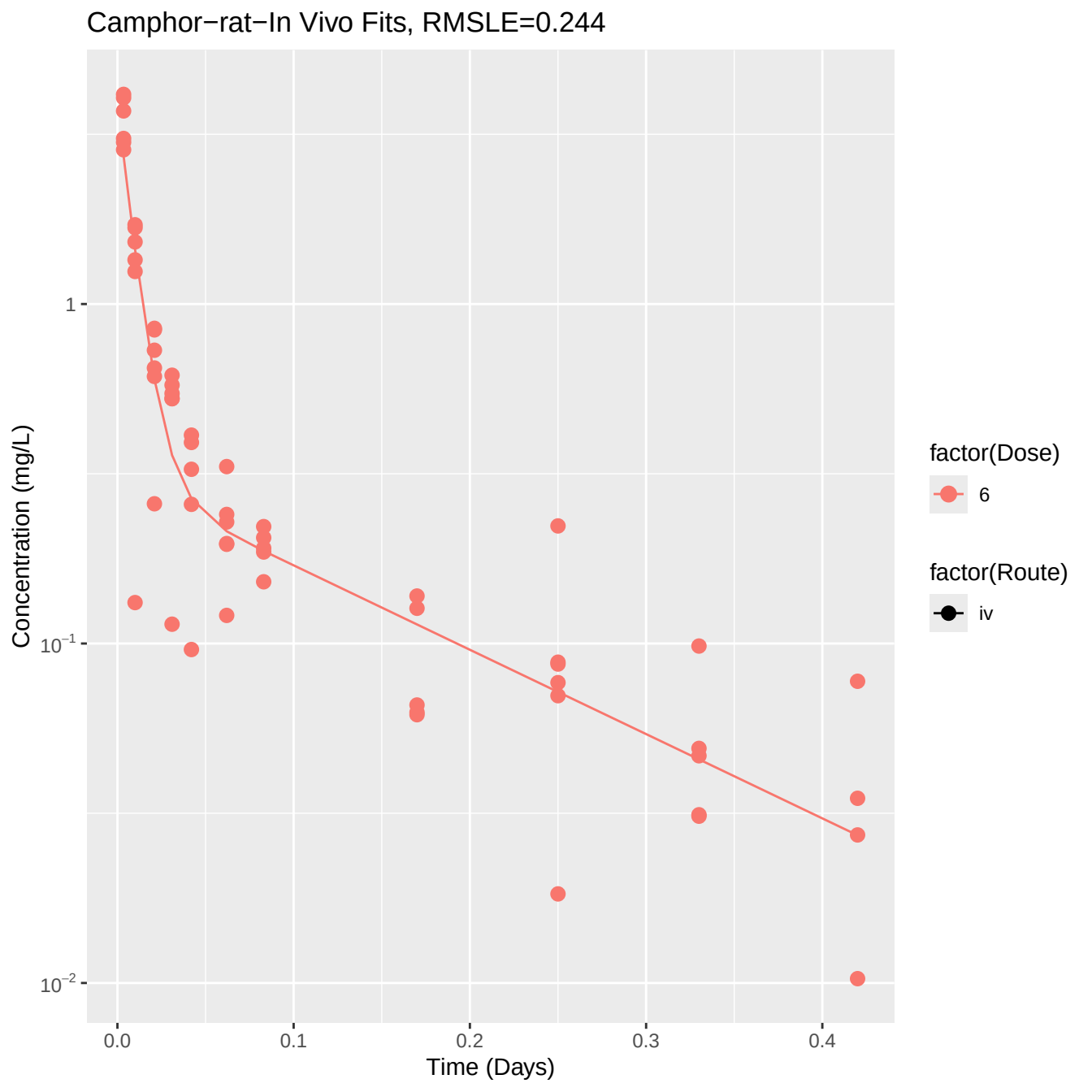

Bromodichloromethane−rat−HTPBTK−OPERA, RMSLE=1.13
Concentration (mg/L)
10
1
10^{-1}
10^{-2}
0.00
0.25
0.50
0.75
1.00
Time (Days)
factor(Dose)
9.34
23.5
24
47.1
47.3
87.8
95.7
factor(Route)
iv
oral

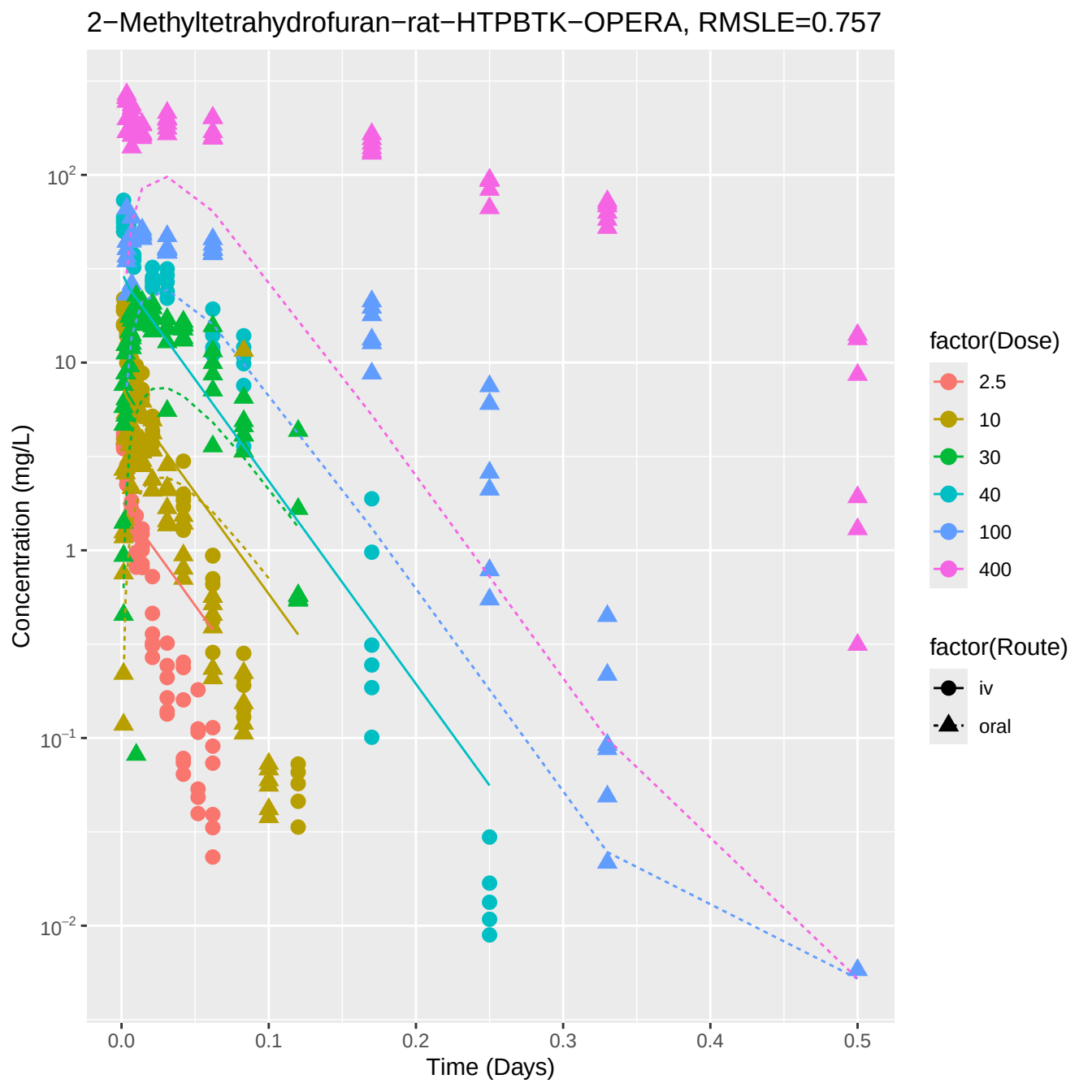

Bromodichloromethane−rat−HTPBTK−Ensemble, RMSLE=1.13
Concentration (mg/L)
10
1
10^{-1}
10^{-2}
0.00
0.25
0.50
0.75
1.00
Time (Days)
factor(Dose)
9.34
23.5
24
47.1
47.3
87.8
95.7
factor(Route)
iv
oral

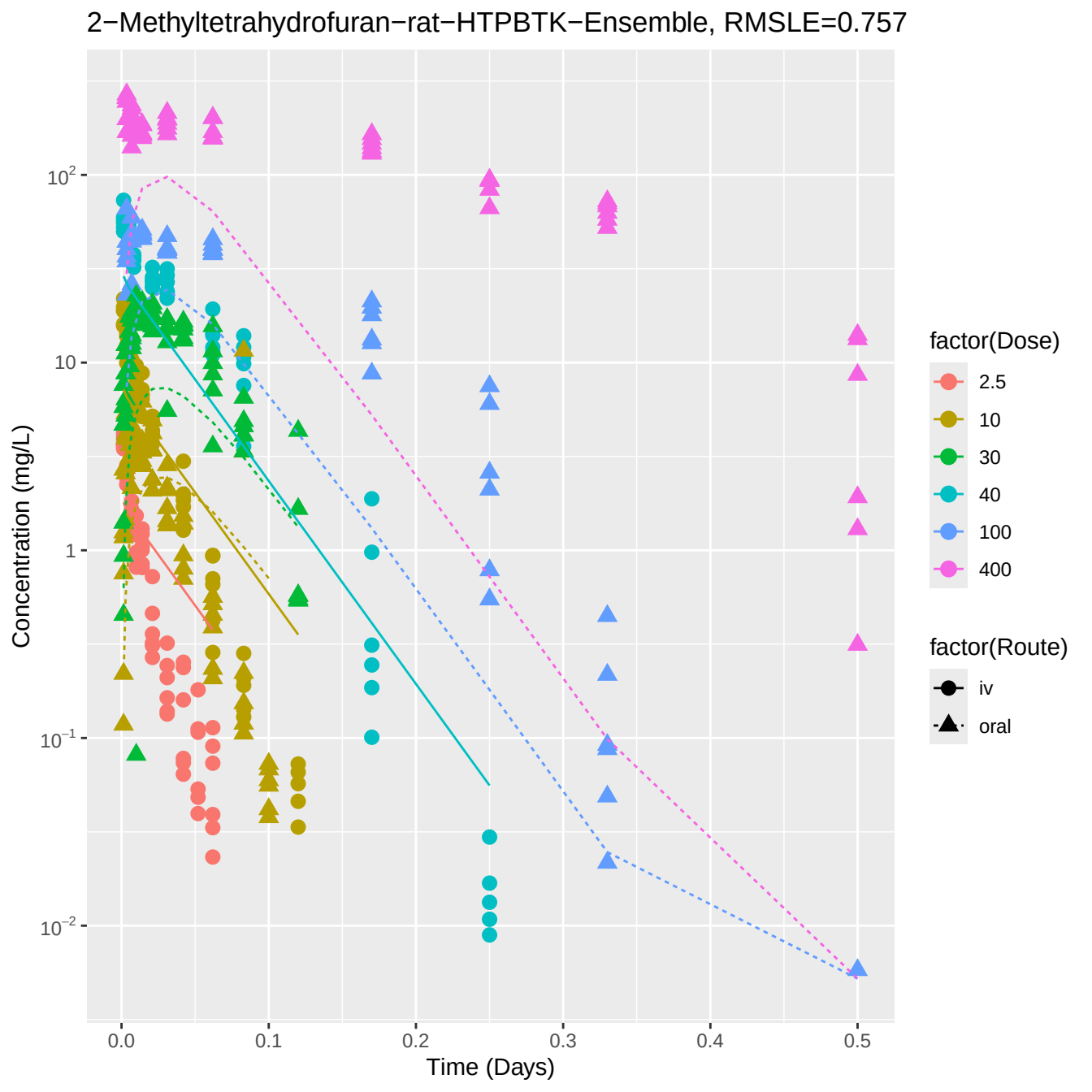

Bromodichloromethane−rat−In Vivo Fits, RMSLE=0.49
Concentration (mg/L)
1
10^{-1}
10^{-2}
0.00
0.25
0.50
0.75
1.00
Time (Days)
factor(Dose)
9.34
23.5
24
47.1
47.3
87.8
95.7
factor(Route)
iv
oral

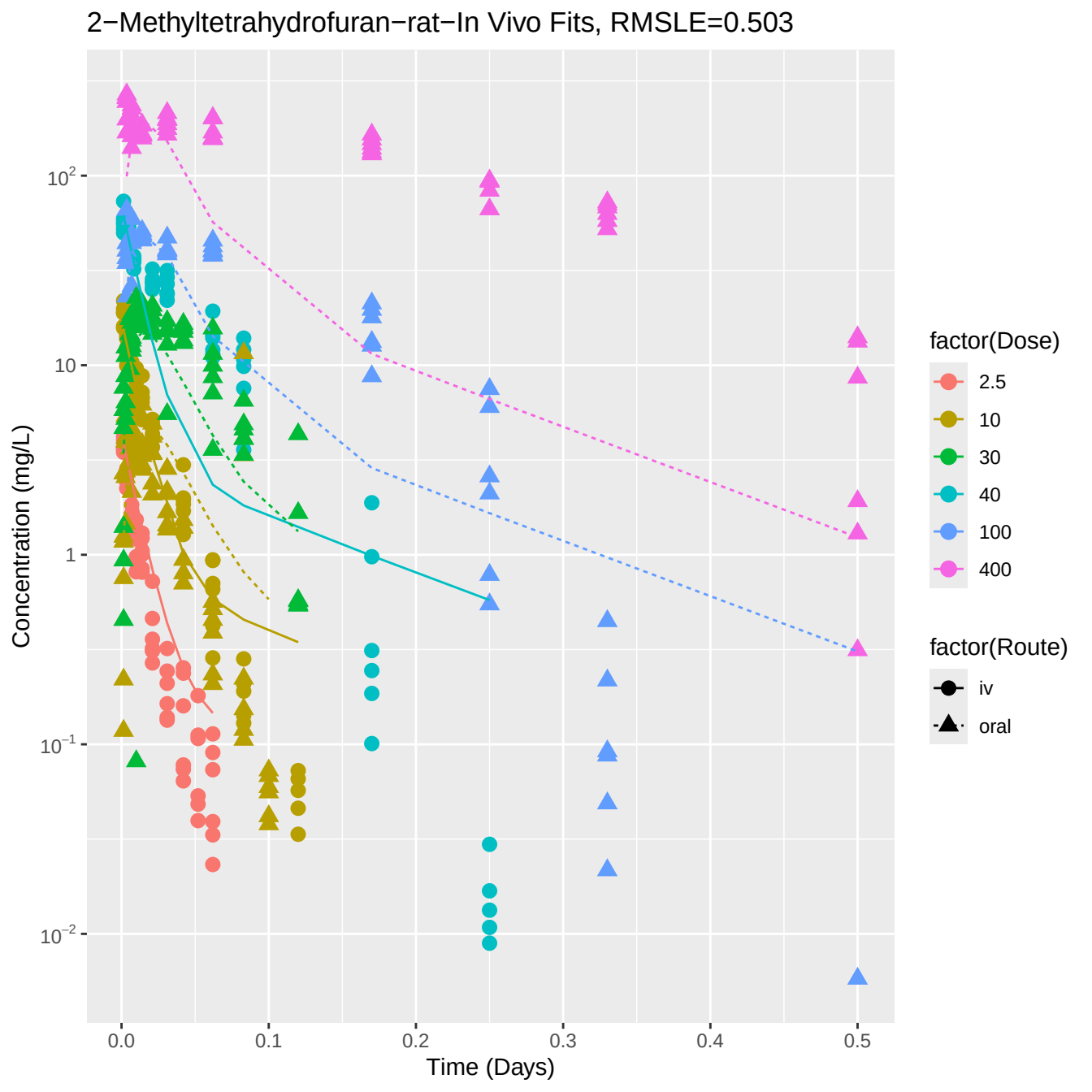

Camphor−rat−HTPBTK−OPERA, RMSLE=0.519
Concentration (mg/L)
1
10^{-1}
10^{-2}
0.0
0.1
0.2
0.3
0.4
Time (Days)
factor(Dose)
6
factor(Route)
iv

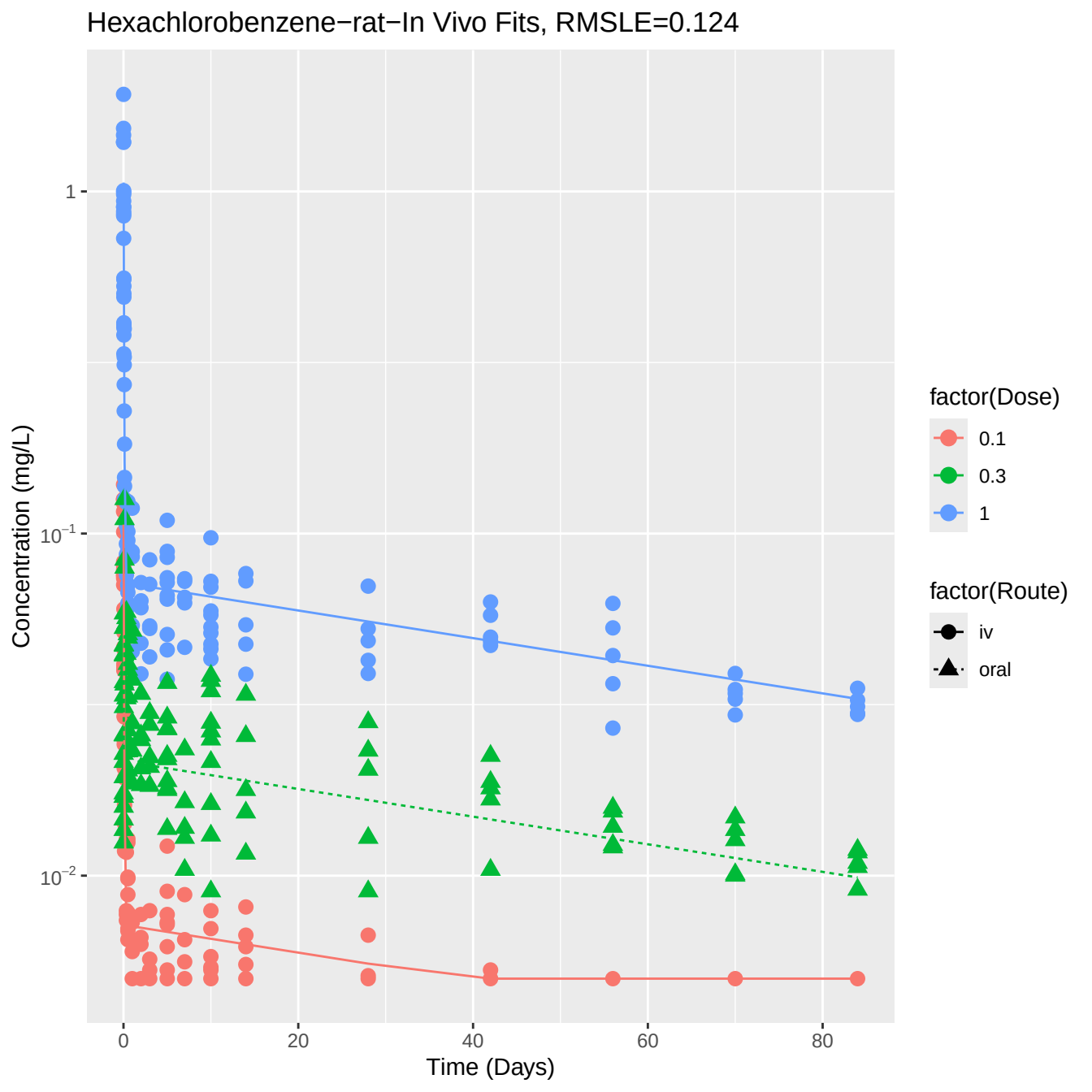

Camphor−rat−HTPBTK−Ensemble, RMSLE=0.519
Concentration (mg/L)
1
10^{-1}
10^{-2}
0.0
0.1
0.2
0.3
0.4
Time (Days)
factor(Dose)
6
factor(Route)
iv

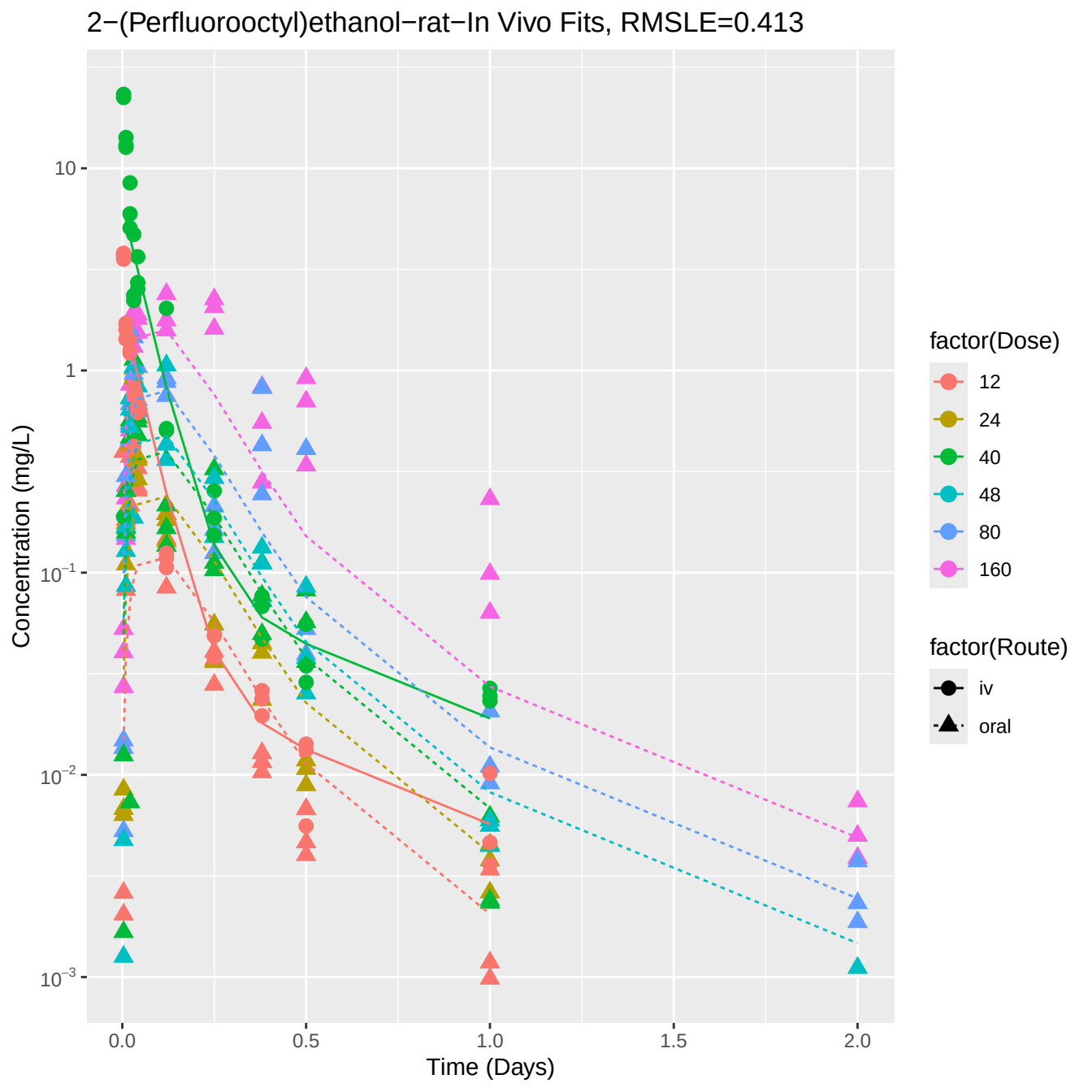

Camphor−rat−In Vivo Fits, RMSLE=0.244
Concentration (mg/L)
1
10^{-1}
10^{-2}
0.0
0.1
0.2
0.3
0.4
Time (Days)
factor(Dose)
6
factor(Route)
iv

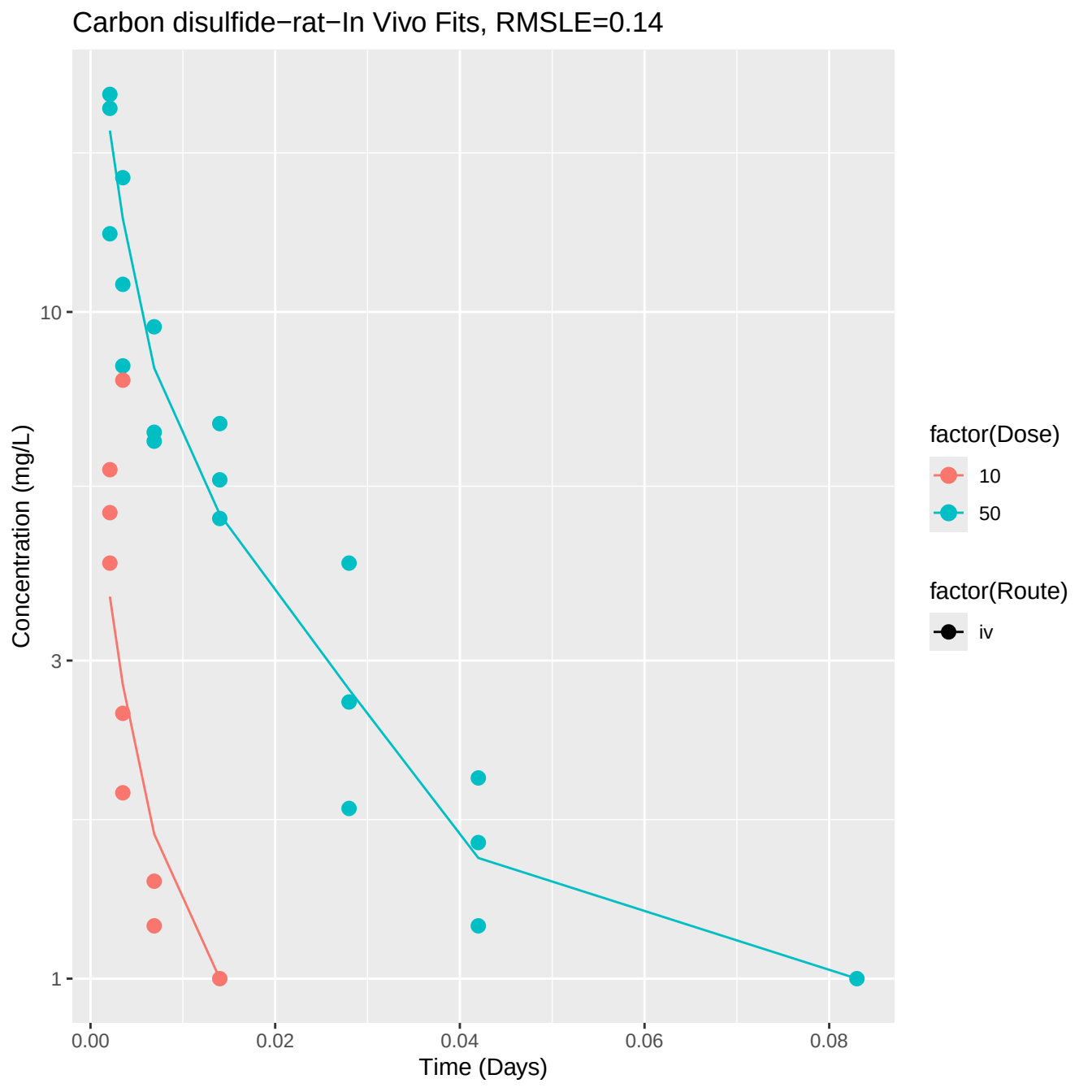

2−Methyltetrahydrofuran−rat−HTPBTK−OPERA, RMSLE=0.757
Concentration (mg/L)
10^2
10
1
10^{-1}
10^{-2}
0.0
0.1
0.2
0.3
0.4
0.5
Time (Days)
factor(Dose)
2.5
10
30
40
100
400
factor(Route)
iv
oral

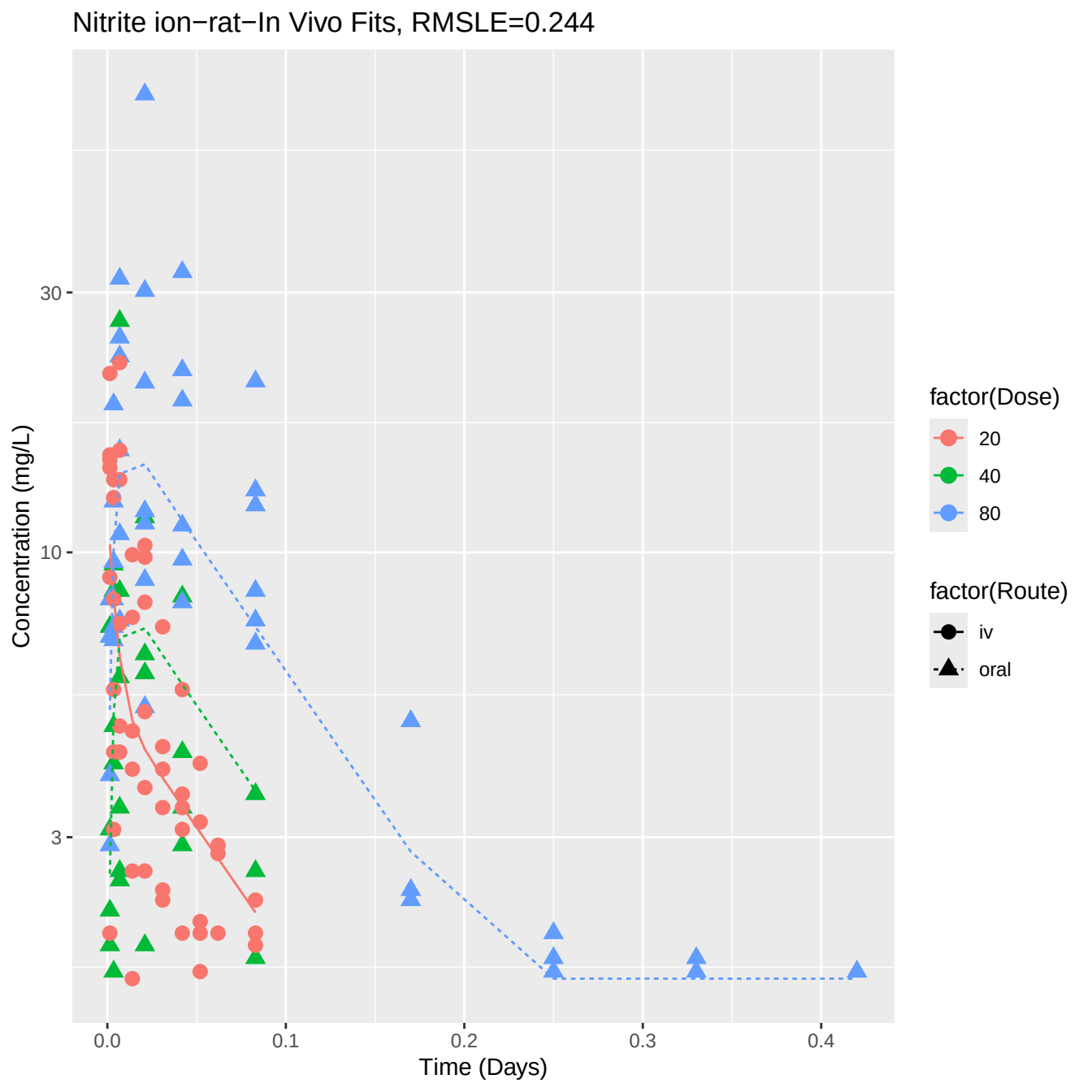

2−Methyltetrahydrofuran−rat−HTPBTK−Ensemble, RMSLE=0.757
Concentration (mg/L)
10^2
10
1
10^{-1}
10^{-2}
0.0
0.1
0.2
0.3
0.4
0.5
Time (Days)
factor(Dose)
2.5
10
30
40
100
400
factor(Route)
iv
oral


2−Methyltetrahydrofuran−rat−In Vivo Fits, RMSLE=0.503
Concentration (mg/L)
10^2
10
1
10^{-1}
10^{-2}
0.0
0.1
0.2
0.3
0.4
0.5
Time (Days)
factor(Dose)
2.5
10
30
40
100
400
factor(Route)
iv
oral


Hexachlorobenzene−rat−In Vivo Fits, RMSLE=0.124
Concentration (mg/L)
1
10^{-1}
10^{-2}
0
20
40
60
80
Time (Days)
factor(Dose)
0.1
0.3
1
factor(Route)
iv
oral


2−(Perfluorooctyl)ethanol−rat−In Vivo Fits, RMSLE=0.413
Concentration (mg/L)
10
1
10^{-1}
10^{-2}
10^{-3}
0.0
0.5
1.0
1.5
2.0
Time (Days)
factor(Dose)
12
24
40
48
80
160
factor(Route)
iv
oral


Carbon disulfide−rat−In Vivo Fits, RMSLE=0.14
Concentration (mg/L)
10
3
1
0.00
0.02
0.04
0.06
0.08
Time (Days)
factor(Dose)
10
50
factor(Route)
iv


Nitrite ion−rat−In Vivo Fits, RMSLE=0.244
Concentration (mg/L)
30
10
3
0.0
0.1
0.2
0.3
0.4
Time (Days)
factor(Dose)
20
40
80
factor(Route)
iv
oral