

Client: Example Client ABC123 123 Test Drive Salt Lake City, UT 84108

UNITED STATES

Physician: Doctor, Example

Patient: Patient, Example

DOB 5/29/1948

Gender: Male

Patient Identifiers: 01234567890ABCD, 012345

Visit Number (FIN): 01234567890ABCD **Collection Date:** 00/00/0000 00:00

Hypoglycemia Panel (Sulfonylureas), Serum or Plasma

ARUP test code 3005636

Rosiglitazone

None Det ng/mL Serum or Plasma

Reporting Limit: 40 ng/mL

Synonym(s): Avandia(R); Avandaryl(R); Avandamet(R) Peak plasma concentrations of approximately 70-430 ng/mL and 240-830 ng/mL were achieved 1 hour after administration of 4 mg and 8 mg

daily doses, respectively.
Analysis by High Performance Liquid Chromatography/
Tandem Mass Spectrometry (LC-MS/MS)

Chlorpropamide

None Det mcg/mL

Serum or Plasma

Reporting Limit: 0.10 mcg/mL

Synonym(s): Diabinese(R)

Peak plasma concentrations of approximately 75-360 mcg/mL were achieved 2 hours following chronic daily doses of 250-1000 mg.

The blood to plasma ratio of Chlorpropamide is not known.

Analysis by High Performance Liquid Chromatography/ Tandem Mass Spectrometry (LC-MS/MS)

Glimepiride

None Det ng/mL Serum or Plasma

Reporting Limit: 25 ng/mL

Synonym(s): Duetact(R); Avandaryl(R); Amaryl(R) Peak plasma concentrations of approximately 60-340 ng/mL were achieved 2-3 hours after administration of 4 mg of glimepiride. The blood to plasma ratio of Glimepiride

is not known.

Analysis by High Performance Liquid Chromatography/ Tandem Mass Spectrometry (LC-MS/MS)

Glipizide

330 ng/mL

H=High, L=Low, *=Abnormal, C=Critical

4848



Serum or Plasma

Reporting Limit: 40 ng/mL

Synonym(s): Glynase; Glucotrol(R); Glibenese Peak plasma concentrations of approximately 310-610 ng/mL were achieved after administration of a single 5 mg dose of both immediate and extended release formulations. Maximum concentrations were reached in approximately 1.5-4.5 and 3.5-7 hours after immediate and extended release dosing respectively.

The blood to plasma ratio of Glipizide is not known. Analysis by High Performance Liquid Chromatography/Tandem Mass Spectrometry (LC-MS/MS)

Pioglitazone

130 ng/mL

Serum or Plasma

Reporting Limit: 40 ng/mL

Synonym(s): Duetact(R); ActoPlus Met(R); Actos(R); Oseni(R) Peak plasma concentrations of approximately 530-2600 ng/mL were achieved 1-4 hour after administration of 45 mg of pioglitazone.

Analysis by High Performance Liquid Chromatography/ Tandem Mass Spectrometry (LC-MS/MS)

Glyburide

None Det ng/mL

Serum or Plasma

Reporting Limit: 40 ng/mL

Synonym(s): PresTab(R); Micronase(R); Glibenclamide; Glynase(R)

Glynase(R)
Peak plasma concentrations of approximately
130-200 ng/mL following a single 5 mg dose have
been reported. A group of ten diabetic patients given
daily oral 2.5 mg doses for 6 weeks attained peak
plasma glyburide concentrations averaging 140 ng/mL
at 3 hours after the first dose and 240 ng/mL
at 2.4 hours after the last dose.
Analysis by High Performance Liquid Chromatography/
Tandem Mass Spectrometry (LC-MS/MS)

Nateglinide

None Det mcg/mL

Serum or Plasma

Reporting Limit: 0.10 mcg/mL

Synonym(s): Starlix(R)

Peak plasma concentrations of approximately
1.3-7.5 mcg/mL were achieved 0.5 hours following

a single 60 mg dose.

Analysis by High Performance Liquid Chromatography/

Tandem Mass Spectrometry (LC-MS/MS)

Tolazamide

None Det mcg/mL

H=High, L=Low, *=Abnormal, C=Critical

4848



Serum or Plasma

Reporting Limit: 0.10 mcg/mL

Synonym(s): Tolinase(R)

No plasma concentrations have been

reported in the literature Analysis by High Performance Liquid Chromatography/ Tandem Mass Spectrometry (LC-MS/MS)

Tolbutamide

None Det mcg/mL

Serum or Plasma

Reporting Limit: 0.10 mcg/mL

Synonym(s): Orinase(R)

Peak plasma concentrations of approximately 50-100 mcg/mL were achieved 3-5 hours following chronic daily doses.

Analysis by High Performance Liquid Chromatography/ Tandem Mass Spectrometry (LC-MS/MS)

Repaglinide

None Det ng/mL

Serum or Plasma

Reporting Limit: 10 ng/mL

Synonym(s): Prandin(R); PrandiMet(R)

Peak plasma concentrations of approximately <10-180 ng/mL were achieved 1 hour after administration of 4 mg of repaglinide.

Analysis by High Performance Liquid Chromatography/
Tandem Mass Spectrometry (LC-MS/MS)
This test was developed and its performance characteristics determined by NMS Labs. It has not been cleared or approved by the US Food and Drug

Administration.

Digital data review may have taken place remotely by qualified NMS staff utilizing a secure VPN connection for some or all of the reported results. This is in accordance with and follows CLIA regulations.

Testing performed at NMS Labs, Inc. 200 Welsh Road
Horsham, PA 19044-2208
CLIA 39D0197898

H=High, L=Low, *=Abnormal, C=Critical

4848



VERIFIED/REPORTED DATES				
Procedure	Accession	Collected	Received	Verified/Reported
Rosiglitazone	24-155-110257	00/00/0000 00:00	00/00/0000 00:00	00/00/0000 00:00
Chlorpropamide	24-155-110257	00/00/0000 00:00	00/00/0000 00:00	00/00/0000 00:00
Glimepiride	24-155-110257	00/00/0000 00:00	00/00/0000 00:00	00/00/0000 00:00
Glipizide	24-155-110257	00/00/0000 00:00	00/00/0000 00:00	00/00/0000 00:00
Pioglitazone	24-155-110257	00/00/0000 00:00	00/00/0000 00:00	00/00/0000 00:00
Glyburide	24-155-110257	00/00/0000 00:00	00/00/0000 00:00	00/00/0000 00:00
Nateglinide	24-155-110257	00/00/0000 00:00	00/00/0000 00:00	00/00/0000 00:00
Tolazamide	24-155-110257	00/00/0000 00:00	00/00/0000 00:00	00/00/0000 00:00
Tolbutamide	24-155-110257	00/00/0000 00:00	00/00/0000 00:00	00/00/0000 00:00
Repaglinide	24-155-110257	00/00/0000 00:00	00/00/0000 00:00	00/00/0000 00:00

END OF CHART

H=High, L=Low, *=Abnormal, C=Critical