

NATIONAL INSTITUTE OF BIOLOGICALS
(Ministry of Health & Family Welfare)

Biological Products Test Reports Released

Date: 18.08.2017

Blood Product Lab

S. No	Name of the Product	Manufacturer	Lot / Batch No.	Expiry Date	Date of receipt of Sample	Date of release of report	Report Dispatch No.	Status of Report	Remarks
1	Human Albumin 20%EP, 100ml	M/s Baxter AG, Austria	PAA1S083	31.03.2020	11.07.2017	12.08.2017	N.3-142/2017-18-SRRD/BP	CoA Released	Nil
2	Human Albumin 20%EP, 100ml	M/s Baxter AG, Austria	PAA1S084	03.2020	11.07.2017	12.08.2017	N.3-143/2017-18-SRRD/BP	CoA Released	Nil
3	Human Albumin 20%EP, 100ml	M/s Baxter AG, Austria	PAA1S092	31.03.2020	11.07.2017	12.08.2017	N.3-144/2017-18-SRRD/BP	CoA Released	Nil
4	RIXUBIS(coagulation Factor IX (Recombinant)250 IU	M/s Baxalta Bioscience, US Inc., USA	TNA16013AE	01.09.2019	30.06.2017	12.08.2017	N.3-128/2017-18-SRRD/BP	CoA Released	Nil
5	Hemofil-M, Anthihemophilic Factor(Human), Method M, Monoclonal Purified Nanofiltration	M/s Baxalta US Inc., USA	THA17214AA	15.09.2019	30.06.2017	12.08.2017	N.3-127/2017-18-SRRD/BP	CoA Released	Nil
6	Hemofil-M, Anthiemophilic Factor(Human), Method M, Monoclonal Purified Nanofiltration	M/s Baxalta US Inc., USA	THA17227AB	04.11.2019	06.07.2017	12.08.2017	N.3-138/2017-18-SRRD/BP	CoA Released	Nil

NATIONAL INSTITUTE OF BIOLOGICALS
(Ministry of Health & Family Welfare)

Biological Products Test Reports Released

S. No	Name of the Product	Manufacturer	Lot / Batch No.	Expiry Date	Date of receipt of Sample	Date of release of report	Report Dispatch No.	Status of Report	Remarks
7	Hemofil-M, Antihemophilic Factor(Human), Method M, Monoclonal Purified Nanofiltration	M/s Baxalta US Inc., USA	THA17215AA	21.09.2019	06.07.2017	12.08.2017	N.3-137/2017-18-SRRD/BP	CoA Released	Nil
8	Hemofil-M, Nanofiltered, Antihemophilic Factor (Human) Method M, Monoclonal Purified, F VIII	Baxalta US Inc., USA	THL16230AA	06.09.2018	16.12.2016	20.02.2017	N.3-572/2016-SRRD/BP	CoA Released	Nil
9	Hemofil-M, Nanofiltered, Antihemophilic Factor (Human) Method M, Monoclonal Purified, F VIII	Baxalta US Inc., USA	THL16228AA	03.09.2018	16.12.2016	20.02.2017	N.3-571/2016-SRRD/BP	CoA Released	Nil
10	IMMUNATE 250 IU, Purified Freeze-Dried Human Coagulation Factor VIII, Virus Inactivated	Baxter AG, Austria	VNC3R047	30.04.2018	20.01.2017	28.02.2017	N.3-628/2016-17-SRRD/BP	CoA Released	Nil

Recombinant Product Lab

S. No	Name of the Product	Manufacturer	Lot / Batch No.	Expiry Date	Date of receipt of Sample	Date of release of report	Report Dispatch No.	Status of Report	Remarks
1	Insulatard Flexpen, Isophane insulin injection IP, 100 IU/ml	Novo Nordisk Producao Farmaceutica do Brazil Ltda., Brazil	GR79302	7.2019	13.07.2017	16.08.2017	N.4-95/2017-18-SRRD/RP	CoA Released	Nil

NATIONAL INSTITUTE OF BIOLOGICALS
(Ministry of Health & Family Welfare)

Biological Products Test Reports Released

S. No	Name of the Product	Manufacturer	Lot / Batch No.	Expiry Date	Date of receipt of Sample	Date of release of report	Report Dispatch No.	Status of Report	Remarks
2	Human Mixtard, Biphasic Isophane insulin injection IP, 100 IU/ml	Novo Nordisk A/S, Denmark	FS61R54	3.2019	12.01.2017	15.02.2017	N.4-544/2016-17-SRRD/RP	CoA Released	Nil
3	NovoMix 30 Flexpen, I.P Mixture of insulin aspart & protamine crystallized insulin aspart, 100 U/ml	Novo Nordisk A/S, Denmark	FP53174	8.2018	27.12.2016	13.02.2017	N.4-519/2016-SRRD/RP	CoA Released	Nil
4	NovoMix 50 Flexpen, Biphasic Insulin aspart I.P., 100 U/ml	Novo Nordisk A/S, Denmark	FP53209	9.2018	27.12.2016	13.02.2017	N.4-520/2016-SRRD/RP	CoA Released	Nil
5	NovoMix 50 Penfill, Biphasic Insulin aspart I.P., 100 U/ml	Novo Nordisk A/S, Denmark	FS61H86	7.2018	27.12.2016	13.02.2017	N.4-517/2016-SRRD/RP	CoA Released	Nil
6	NovoMix 50 Penfill, Biphasic Insulin aspart I.P., 100 U/ml	Novo Nordisk A/S, Denmark	FS61J43	7.2018	27.12.2016	13.02.2017	N.4-518/2016-SRRD/RP	CoA Released	Nil

NATIONAL INSTITUTE OF BIOLOGICALS
(Ministry of Health & Family Welfare)

Biological Products Test Reports Released

Infection Diagnostic Lab

S. No	Name of the Product	Manufacturer	Lot / Batch No.	Expiry Date	Date of receipt of Sample	Date of release of report	Report Dispatch No.	Status of Report	Remarks
1	HIV1 & 2 ELISA KIT	Trivitron Healthcare Pvt. Ltd., Chennai	HIVT E61109	1.2018	13.12.2016	09.01.2017	N.1-358/2016-SRRD/IDKL	CoA Released	Nil
2	HIV1 & 2 ELISA KIT	Trivitron Healthcare Pvt. Ltd., Chennai	HIVT E61107	1.2018	13.12.2016	09.01.2017	N.1-356/2016-SRRD/IDKL	CoA Released	Nil
3	HIV Ag-Ab ELISA KIT	Trivitron Healthcare Pvt. Ltd., Chennai	HIVF E61110	1.2018	13.12.2016	11.01.2017	N.1-359/2016-SRRD/IDKL	CoA Released	Nil
4	STANDARD Q HCV Ab Rapid Test	SD Biosensor Inc., Republic of Korea	QHC1016002-2S	24.10.2018	28.12.2016	01.02.2017	N.1-374/2016-SRRD/IDKL	CoA Released	Nil
5	HCV ELISA KIT	Trivitron Healthcare Pvt. Ltd., Chennai	HCV E61106	1.2018	13.12.2016	11.01.2017	N.1-355/2016-SRRD/IDKL	CoA Released	Nil
6	HCV ELISA KIT	Trivitron Healthcare Pvt. Ltd., Chennai	HCV E61105	1.2018	13.12.2016	11.01.2017	N.1-354/2016-SRRD/IDKL	CoA Released	Nil

NATIONAL INSTITUTE OF BIOLOGICALS
(Ministry of Health & Family Welfare)

Biological Products Test Reports Released

S. No	Name of the Product	Manufacturer	Lot / Batch No.	Expiry Date	Date of receipt of Sample	Date of release of report	Report Dispatch No.	Status of Report	Remarks
7	STANDARD Q HCV Ab Rapid Test	SD Biosensor Inc., Republic of Korea	QHC1016002-3S	24.10.2018	28.12.2016	01.02.2017	N.1-375/2016-SRRD/IDKL	CoA Released	Nil