



सत्यमेव जयते

National Institute of Biologicals, NOIDA
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Ministry of Health & Family Welfare
Government of India

HAEMOVIGILANCE NEWSLETTER

Haemovigilance Programme of India

Haemovigilance
Newsletter Vol. No. 8,
Issue 16, July-
December, 2020



HvPI Initiated State wise
CMEs in the month of
February- March, 2020

08
National Level
CME via Webinar

04-05
Trainings/CMEs
under HvPI

03
Haemovigilance
Programme
of India-Milestones

06
State Wise CMEs

10
Version-2 Adverse
Blood Donor Reaction
Reporting Form Launched
w.e.f. 1st Jan, 2020

"The aim of the newsletter is to disseminate information on Haemovigilance Programme of India so as to create awareness amongst healthcare professionals & other stakeholders on safe Blood Transfusion & Blood Products' Administration Practices"

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TABLE OF CONTENTS

Haemovigilance Programme of India-Milestones	3
Trainings/CMEs under HvPI.....	4
HvPI Initiated State wise CMEs in the month of Feb- March, 2020.....	6
National Level CME on Haemovigilance & Voluntary Blood Donation in COVID-19 Pandemic	8
Version-2 Adverse Blood Donor Reaction Reporting Form w.e.f 1st Jan, 2020 launched.....	10
Transfusion Reaction Reporting Form.....	12
New Members Enrolled under Haemovigilance Programme of India	14
How to Enroll your Centre under HvPI.....	15

HAEMOVIGILANCE PROGRAMME OF INDIA - MILESTONES

Haemovigilance Programme of India was launched on 10th December, 2012 at the National level in 90 medical institutions across the country by National Institute of Biologicals (NIB), NOIDA, Ministry of Health & Family Welfare, Government of India as the National Coordinating Centre (NCC). The objective of this programme is to track Adverse Reactions associated with Blood Transfusion and Blood Donation.

Haemovigilance is defined as 'a set of surveillance procedures covering the whole transfusion chain from the collection of blood and its components to the follow-up of its recipients i.e. from the vein of the donor to the vein of the recipient. It is intended to collect and assess information on unexpected or undesirable effects resulting from the therapeutic use of labile blood products and to prevent their occurrence and recurrence'. Haemovigilance is a tool to improve the quality of the blood transfusion chain, primarily focusing on safety.

1. The recipient's arm i.e. reporting of Adverse Reactions with respect to Blood Transfusion in the patient is being covered under **Haemovigilance Programme of India (HvPI)** with the launch of the programme on 10th December, 2012 in the country.
2. The donor's arm i.e. Reporting of Adverse Reactions associated with Blood Donations is being covered under **National Blood Donor Vigilance Programme (NBDVP)** which was launched on 14th June, 2015 on World Blood Donor Day at Science City Kolkata under the ambit of HvPI.
3. Reporting of Adverse Transfusion Reactions is done online via Haemo-Vigil software & reporting of Adverse Blood Donor Reactions is done via Donor-Vigil software available on NIB website www.nib.gov.in

Implementation and coordination of activities of Haemovigilance Programme of India became one of the Mandate's of NIB as per its bye-laws 3.4.1 of the Institute as approved in the Governing Body meeting of NIB held under chairpersonship of Secretary (Health & F.W.)/ Chairman, Governing Body of NIB on 12th Dec, 2014.

DCG (I) issued an office memorandum dated 4th December, 2015 w.r.t. enrolment of all licensed blood centres under HvPI. These licensed blood centres are required to obtain their user ID and password from NIB to uplink their adverse transfusion data to Haemo-Vigil software under HvPI.

National Accreditation Board for Hospitals and Healthcare Providers (NABH) in its third edition of accreditation standards on Blood Centres and transfusion services issued in year 2016 has included enrolment by Blood Centres under National Haemovigilance Program of India and monitor adverse donor reactions and adverse transfusion reactions as per the direction issued.

NCC-HvPI, NIB issues certificate of participation to the centres who are actively reporting under Haemovigilance Programme of India.

TRAININGS/CMEs UNDER HVPI

1. CME on Haemovigilance Programme of India was organized by National Institute of Biologicals (NIB), NOIDA in collaboration with SRCC TRUST'S SRCC CHILDREN'S HOSPITAL BLOOD BANK, Mumbai Maharashtra on 18th- 19th January, 2020 at Maharashtra.



2. CME on Blood Safety and Haemovigilance was organized by National Institute of Biologicals (NIB), NOIDA in collaboration with Blood Centre Technologists' Association Maharashtra (BCTAM), Maharashtra on 28th and 29th February, 2020 at Aurangabad, Maharashtra.



- About 200 participants attended the aforesaid CMEs which included Blood Bank Official(s), Medical Officer(s), Staff Nurse(s), Senior Supervisor(s), Technical Supervisor(s) and Technician(s).
- Presentation on various aspects of Haemovigilance followed by Hands on Training on Haemovigilance software(s) were imparted to the participants.

3. NIB- National Health Mission (NHM) Collaborative Six days Residential Training Programme for Strengthening of Blood Services for the state Maharashtra, Chhattisgarh & Uttar Pradesh.



Maharashtra on 20th – 25th
January, 2020



Chhattisgarh on 03rd– 08th
February, 2020



Uttar Pradesh on 02nd– 07th
March, 2020

4. National Skill Development and Hands - on Training Program on Quality Control of Biologicals for M.Sc. Biotechnology, Microbiology and Biochemistry students of Various Universities of the Country.



Jammu University, J & K and
Tezpur University, Assam on
06th– 17th January, 2020



Mizoram University, Mizoram &
Himachal Pradesh University,
Shimla, Himachal Pradesh on 27th
January– 07th February, 2020



Gauhati University, Assam &
Central Research Institute,
Kasauli, Himachal Pradesh on
17th–28th February, 2020

Head- HvPI invited as a faculty to deliver a presentation on:-

- (a) Haemovigilance Programme of India during CME on current trends & challenges in immunohematology & blood transfusion held on 20th February, 2020 at Deen Dayal Upadhyay Hospital, Hari Nagar Delhi.



- (b) Haemovigilance Programme of India on 03rd March, 2020 during 6th Asia Pacific Pharmacovigilance Training Course schedule from 24th February to 6th March 2020 organized by Indian Pharmacopoeia Commission, Ghaziabad.

IHN Teleconference:-

Head- HvPI & Secretary of International Haemovigilance Network (IHN) attended teleconference of IHN Board on 29th April, 2020 at 07:00 pm of Indian Standard Time.

STATE WISE TARGETED CONTINUING MEDICAL EDUCATION (CMEs) ON HAEMOVIGILANCE PROGRAMME OF INDIA

Initiative of organizing State wise targeted CMEs on Haemovigilance Programme of India by National Institute of Biologicals, NOIDA. The objective of the CMEs are to enroll & encourage reporting by the blood centres under HvPI in the respective states/ UTs. The following CMEs were organized in this regard during January - June, 2020.

S. No.	Name of the Medical College/ Organization/ Institute Involved	Date & Place of CME
1.	Drugs Control Organization, Rajasthan & R.N.T. Medical College, Udaipur	7th February, 2020 at Udaipur
2.	Drugs Control Organization, Rajasthan & All India Institute of Medical Sciences, Jodhpur	10th February, 2020 at Jodhpur
3.	Gujarat State AIDS Control Society (GSAC) & Gujarat State Council for Blood Transfusion (GSCBT) and Surat Raktdan Kendra	15th February, 2020 at Surat
4.	Gujarat State AIDS Control Society (GSAC) & Gujarat State Council for Blood Transfusion (GSCBT) and Indian Red Cross Society, Ahmedabad	17th February, 2020 at Ahmedabad
5.	Gujarat State AIDS Control Society (GSAC) & Gujarat State Council for Blood Transfusion (GSCBT) and Red Cross Blood Bank, Rajkot	24th February, 2020 at Rajkot
6.	Department of Drugs Control Administration, Tamil Nadu & Ganga Medical Center & Hospitals (P) Ltd., Coimbatore	7th March, 2020 at Coimbatore
7.	Department of Drugs Control Administration, Tamil Nadu, Chennai	14th March, 2020 at Chennai



Udaipur (07th February, 2020)



Jodhpur (10th February, 2020)



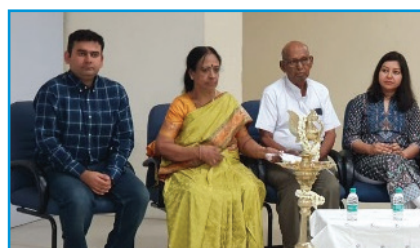
Surat (15th February, 2020)



Ahmedabad (17th February, 2020)



Rajkot (24th February, 2020)



Coimbatore (07th March, 2020)



Chennai (14th March, 2020)

Salient Features:

- About 300 participants which included Blood Centre officials, clinicians, technicians & nurses were trained during the aforesaid CMEs.
- Presentations on Haemovigilance Programme of India, Recipient Haemovigilance, Blood Donor Vigilance, Regulation of Blood Centres followed by Case Studies pertaining to Recipient Haemovigilance & Blood Donor Vigilance.
- Haemovigilance Software(s) Demonstration & Hands on Training imparted to the participants.
- On the spot enrolment for the Blood Banks under HvPI.
- Panel Discussion was held on action plan for reporting reactions under HvPI.

National Level CME on Haemovigilance & Voluntary Blood Donation in COVID-19 Pandemic on 14th–15th June, 2020.

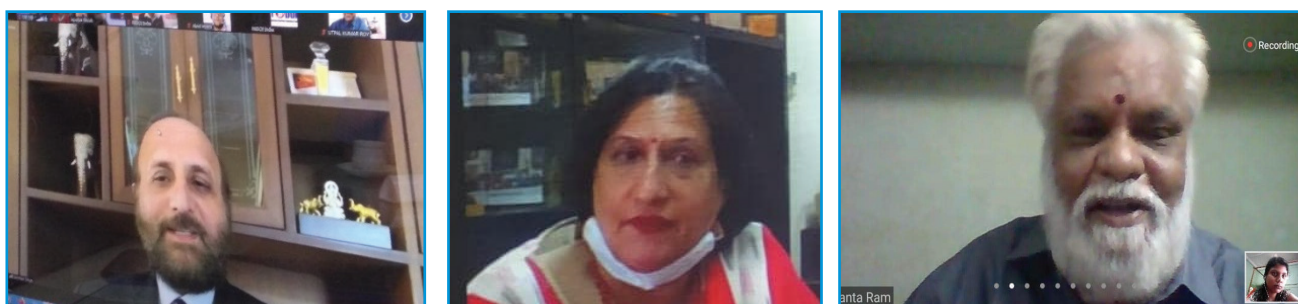
The National Level CME on Haemovigilance and Voluntary Blood Donation in COVID-19 Pandemic organized by National Institute of Biologicals (NIB), NOIDA, Ministry of Health and Family Welfare, Government of India and Federation of Blood Donor Organizations of India and West Bengal Voluntary Blood Donors Forum was held on 14th and 15th June, 2020. The two-day webinar was supported by Mani Trust, Kalimpong and Animesh Gosh, Cancer Awareness Center, West Bengal.

In webinar, about 119 eminent personalities from 21 States all cross the country participated.

This webinar focused on importance of blood donations during the present pandemic situation but also brain stormed on ways and means to continue blood donations post the COVID-19 period.

The webinar was inaugurated by Dr. Surinder Singh, Vice Chancellor, JSS, AHER, Mysuru, Dr. Reba Chhabra, Director i/c, NIB, NOIDA, and Shri L. Santaram, Secretary AVBD, Chennai

Inauguration:-



From Left to Right Dr. Surinder Singh, Vice Chancellor, JSS, AHER, Mysuru , Dr. Reba Chhabra, Director i/c, NIB & Shri L. Santaram, Secretary AVBD, Chennai

Presentations & Lectures:-

First Day (14.06.2020)



1. Dr. Reba Chhabra, Director, In-charge, NIB, NOIDA : Gave presentation on Haemovigilance which focusses on National Blood Donor Vigilance Programme (NBDVP) under HvPI.
2. Shri L. Santaram, Secretary AVBD, Tamil Nadu, Chennai : Discussed about the national data of blood units collected in the past year and the target for the year 2020- 2021.

3. Dr. Debasish Gupta, Founder President, FBDI & Expert BTS : Highlighted the importance of blood requirement not only during this pandemic but also in the future.
4. Dr. Akanksha Bisht: Scientist II & Head, Haemovigilance, NIB, NOIDA : Gave insight about NBDVP and online reporting system of donor reactions.
5. Dr. Amit Dutt Dwuary, Oncologist, Apollo Cancer Hospital, Kolkata : Presented on the blood requirements and components for a Cancer patient.
6. Dr. Kajal Krishna Banik, Expert Public Health, Department of Health & Family Welfare Government of West Bengal; Former General Secretary, Indian Medical Association, West Bengal Chapter : Presented on the importance of "blood donors", SOPs on safety measures and safe blood donation.
7. Dr. Manisha Srivastava, Expert BTS & Superintendent, AIIMS Hospital, Bhopal : Gave update about the COVID-19 spread and donor awareness.

Second Day (15.06.2020)



1. Dr. Naresh Kumar Bhatia, President FBDI & Expert on Blood Banking and BTS : presented the guidelines on how an organization can conduct a blood donation camp during the COVID- 19 outbreak.
2. Shree Dhanaji Raane, Chairman, FBDI Mumbai, Maharashtra : gave presentation on motivating the blood donors and on how to organize a blood donation camp (outdoors) to maintain safety.
3. Dr. Shyamai Bran Mukherjee, Secretary, IRCS : apprised about the sanitization procedure pre and post blood donation camp.
4. Mr. Deepak Rai, CC VBDAS, Sikkim : described the data of the state where 56 blood donation camps held during the lockdown in which 672 units of blood was collected.
5. Mr. Ashu Paul, Vice President FBDI & President TYGA : explained the significance of future development of blood donation camps and how the rural areas can also be involved with the camps.

Webinar concluded with the vote of thanks addressed by Dr. Surinder Singh, Vice Chancellor, JSS, AHER, Mysuru.



National Institute of Biologicals
Ministry of Health & Family Welfare, Govt. of India
NATIONAL BLOOD DONOR VIGILANCE PROGRAMME
(Haemovigilance Programme of India)



Adverse Blood Donor Reaction Reporting Form

Version-2

A) Donor Information	
Donor Id *: _____	Type of Donation* (a) Whole Blood (b) Apheresis__ (Platelets/Plasma/ Plasma + Platelets/RBC/Granulocyte/ Peripheral Blood Stem Cells)
Sex * _____ (Male/Female/Other)	Donor Type* (a) Voluntary (b) Replacement (c) Family Donor (d) Autologous (First Time/Repeat)
Weight of Donor (kg) * _____ Height of Donor (cm)* _____	Site of Donation* _____ (Blood Centre/Camp)
Age/ Date of Birth * Yrs: _____ Month: _____ Days: _____ OR _____	Date of Donation * _____
Pre-Donation Vitals* Pulse: _____ per min BP (Systolic): _____ mmHg BP (Diastolic): _____ mmHg	Time of Donation Hr _____ Min _____
B) Whole blood Details of Blood Collected/Apheresis Details of Blood Collected	
(a) Whole Blood Lot No. of Blood Bag* _____ Volume Collected (ml)* _____ Manufacturer of Blood Bag* _____ (Terumo Penpol Limited/Mitra Industries Pvt. Ltd/ HLL Lifecare Ltd/Fresenius Kabi AG/Fenwal Inc/Polymed/Other) Expiry Date of Blood Bag* _____	
(b) Apheresis Lot No. Kit* _____ Volume Collected (ml)* _____ Expiry Date of Kit* _____	
C) Adverse Reaction Details	
Date and Time of reaction* _____ Hr _____ Min _____	Type of Reaction* _____ (Localised/Generalized/Both)
Vitals at the time of Reaction Pulse: _____ per min BP (Systolic): _____ mmHg BP (Diastolic): _____ mmHg	Data Captured* _____ (Onsite/Call back by donor/ Call back by Blood Centre)
Reaction Time* _____	(Pre-Donation/During Donation/After Donation)
Venipuncture Site* _____ (Left/Right)	Injury* _____ (Yes/No)
Venipuncture* _____ (1/2/>2)	Site of Reaction* _____ (At Donation Site/ Outside Donation Site)
	Donation Completed* _____ (Yes/No)
D) Type of Complications:*	
Localised Complications <input type="checkbox"/> A1-Complications mainly characterized by the occurrence of blood outside the vessels (a) <input type="checkbox"/> Haematoma (bruise) (b) <input type="checkbox"/> Arterial puncture (c) <input type="checkbox"/> Delayed(bleeding/Re-bleeding) <input type="checkbox"/> (Within 30 minutes of Donation/After 30 minutes of Donation) <input type="checkbox"/> A2-Complications mainly characterized by pain (a) <input type="checkbox"/> Nerve injury/irritation (b) <input type="checkbox"/> Other Painful arm <input type="checkbox"/> A3-Localised infection/inflammation along the course of a vein (a) <input type="checkbox"/> Thrombophlebitis (b) <input type="checkbox"/> Cellulitis <input type="checkbox"/> A4- Allergy (local): Itching and redness at the <input type="checkbox"/> (Venipuncture Site/Medical Adhesive Medicated Tape/Skin Disinfection Area) <input type="checkbox"/> A5-Other major blood vessel injury -Serious conditions needing specialist medical diagnosis and attention (a) <input type="checkbox"/> Deep venous thrombosis (DVT) (b) <input type="checkbox"/> Arteriovenous fistula (c) <input type="checkbox"/> Compartment syndrome (d) <input type="checkbox"/> Brachial artery pseudoaneurysm	

Haemovigilance Programme of India launched the second version of Adverse Blood Donor Reaction Reporting Form (ABDRRF Version-2) w.e.f. 1st January, 2020.



National Institute of Biologicals
Ministry of Health & Family Welfare, Govt. of India
NATIONAL BLOOD DONOR VIGILANCE PROGRAMME
(Haemovigilance Programme of India)





Adverse Blood Donor Reaction Reporting Form

Version-2

Generalized Complications			
<input type="checkbox"/> B1-Vasovagal reactions			
(a) <input type="checkbox"/> Generalized Weakness	(b) <input type="checkbox"/> Anxiety	(c) <input type="checkbox"/> Dizziness	(d) <input type="checkbox"/> Nausea
(e) <input type="checkbox"/> Vomiting	(f) <input type="checkbox"/> Pallor(skin and lips)	(g) <input type="checkbox"/> Rapid Pulse	(h) <input type="checkbox"/> Convulsions
(i) <input type="checkbox"/> Cold extremities	(j) <input type="checkbox"/> Hyperventilation	(k) <input type="checkbox"/> Hypotension	(l) <input type="checkbox"/> Low Vol Pulse
(m) <input type="checkbox"/> Feeling of warmth	(n) <input type="checkbox"/> Tetany	(o) <input type="checkbox"/> Loss of bowel or bladder control	(p) <input type="checkbox"/> Cyanosis
(q) <input type="checkbox"/> Loss of Consciousness(LOC) <input type="text"/> (<60 Sec/>60 Sec)			
<input type="checkbox"/> B2-Allergic reactions (Generalized)			
(a) <input type="checkbox"/> Cyanosis	(b) <input type="checkbox"/> Wheezing	(c) <input type="checkbox"/> Flushing,swelling of eyes,lips or tongue	
(d) <input type="checkbox"/> Chest tightness	(e) <input type="checkbox"/> Cardiac arrest		
<input type="checkbox"/> B3-Other serious complications related to blood donation			
(a) <input type="checkbox"/> Acute cardiac symptoms(other than myocardial infarction or cardiac arrest)		(b) <input type="checkbox"/> Myocardial infarction(MI)	
(c) <input type="checkbox"/> Cardiac arrest		(d) <input type="checkbox"/> Transient Ischemic attack (TIA)	
		(e) <input type="checkbox"/> Death	
Apheresis Complication			
<input type="checkbox"/> C-Complications related to apheresis			
(a) <input type="checkbox"/> Citrate reaction			
<input type="checkbox"/> tingling/vibrations-lips,fingers	<input type="checkbox"/> light-headedness	<input type="checkbox"/> Metallic taste	<input type="checkbox"/> Muscle twitching
<input type="checkbox"/> Carpopedal spasm	<input type="checkbox"/> Shock	<input type="checkbox"/> Cardiac arrest	<input type="checkbox"/> Tetany
<input type="checkbox"/> Prophylactic Calcium given before reaction <input type="text"/> (Yes/No)			
(b) <input type="checkbox"/> Haemolysis during procedure			
(c) <input type="checkbox"/> Air embolism			
(d) <input type="checkbox"/> Unable to return red cell(>200ml)			
Other Complication			
<input type="checkbox"/> D-Other Reactions Please Specify <input style="width: 150px;" type="text"/>			
Outcome*			
<input type="checkbox"/> Resolved on donation site		<input type="checkbox"/> Resolved on follow up	<input type="checkbox"/> Recovered with Sequelae
<input type="checkbox"/> Permanently disabled		<input type="checkbox"/> Death following the adverse reactions	<input type="checkbox"/> Unknown
Imputability*			
<input type="checkbox"/> Definite (Certain)		<input type="checkbox"/> Probable (Likely)	<input type="checkbox"/> Possible
<input type="checkbox"/> Unlikely (Doubtful)		<input type="checkbox"/> Excluded	
Any Other Information: <input style="width: 100px;" type="text"/>			
Reporter		Date of Report	
Denominator Data about All Donor			
Total Donation in the month (of reporting)			
<input type="checkbox"/> Whole blood <input style="width: 50px;" type="text"/>			
Volume of donation (Total)*	No. of 350 ml bags <input style="width: 50px;" type="text"/>	No. of 450 ml bags <input style="width: 50px;" type="text"/>	
<input type="checkbox"/> Apheresis if apheresis <input style="width: 50px;" type="text"/>	RBC <input style="width: 50px;" type="text"/>	Platelets <input style="width: 50px;" type="text"/>	Plasma <input style="width: 50px;" type="text"/>
	Plasma+Platelets <input style="width: 50px;" type="text"/>	Granulocyte <input style="width: 50px;" type="text"/>	Peripheral Blood Stem Cells <input style="width: 50px;" type="text"/>
Gender of Donor(Total)*	Male <input style="width: 50px;" type="text"/>	Female <input style="width: 50px;" type="text"/>	Other <input style="width: 50px;" type="text"/>
Type of Donation(Total)*	Voluntary <input style="width: 50px;" type="text"/>	Replacement <input style="width: 50px;" type="text"/>	Family Donor <input style="width: 50px;" type="text"/>
			Autologous <input style="width: 50px;" type="text"/>
Donor Types(Total)*	First-Time Donors <input style="width: 50px;" type="text"/>	Repeat Donors <input style="width: 50px;" type="text"/>	
Site of Donation(Total)*	Blood Centre <input style="width: 50px;" type="text"/>	Camp <input style="width: 50px;" type="text"/>	

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Transfusion Reaction Reporting Form (TRRF Version-2)

	National Institute of Biologicals Ministry of Health & Family Welfare, Govt. of India (National Coordinating Center) HAEMOVIGILANCE PROGRAMME OF INDIA																																																							
Transfusion Reaction Reporting Form (TRRF) For Blood & Blood Components & Plasma Products																																																								
* Mandatory Field																																																								
(A) Patient Information																																																								
Hospital Code No.:																																																								
Patient Initials*: Gender*: Blood Group*:																																																								
Hospital Admission No. *: Age/Date of Birth*:YrsMonthDaysHrsMins																																																								
Primary Diagnosis*:																																																								
Medical History:																																																								
(B) Transfusion Reaction Details*																																																								
Was the patient under anaesthesia during transfusion: Yes/No if Yes type : GA/Spinal/LA																																																								
Pre-transfusion Vitals: Temp: Pulse: BP: RR: SPO2:																																																								
Vitals at the time of reaction: Temp: Pulse: BP: RR: SPO2:																																																								
Please tick mark the relevant signs and symptoms listed below																																																								
<table border="1" style="width: 100%; border-collapse: collapse;"> <thead> <tr> <th colspan="2">Generalised</th> <th>Pain</th> <th>Respiratory</th> <th>Renal</th> <th>Circulatory</th> </tr> </thead> <tbody> <tr> <td><input type="checkbox"/> Fever</td> <td><input type="checkbox"/> Anxiety</td> <td><input type="checkbox"/> Chest Pain</td> <td><input type="checkbox"/> Dyspnoea</td> <td><input type="checkbox"/> Haematuria</td> <td><input type="checkbox"/> Tachycardia</td> </tr> <tr> <td><input type="checkbox"/> Chills</td> <td><input type="checkbox"/> Itching (Pruritus)</td> <td><input type="checkbox"/> Abdominal</td> <td><input type="checkbox"/> Wheeze</td> <td><input type="checkbox"/> Haemoglobinuria</td> <td><input type="checkbox"/> Hypertension</td> </tr> <tr> <td><input type="checkbox"/> Rigors</td> <td><input type="checkbox"/> Edema (Site) _____</td> <td><input type="checkbox"/> Back/Flank Pain</td> <td><input type="checkbox"/> Cough</td> <td><input type="checkbox"/> Oliguria</td> <td><input type="checkbox"/> Hypotension</td> </tr> <tr> <td><input type="checkbox"/> Nausea</td> <td><input type="checkbox"/> Jaundice</td> <td><input type="checkbox"/> Infusion Site Pain</td> <td><input type="checkbox"/> Hypoxemia</td> <td><input type="checkbox"/> Other _____</td> <td><input type="checkbox"/> Raised JVP</td> </tr> <tr> <td><input type="checkbox"/> Urticaria</td> <td><input type="checkbox"/> Other _____</td> <td><input type="checkbox"/> Other _____</td> <td><input type="checkbox"/></td> <td></td> <td><input type="checkbox"/> Arrhythmias</td> </tr> <tr> <td><input type="checkbox"/> Flushing</td> <td></td> <td></td> <td><input type="checkbox"/> Bilateral Infiltrates on</td> <td></td> <td><input type="checkbox"/> Other _____</td> </tr> <tr> <td><input type="checkbox"/> Restlessness</td> <td></td> <td></td> <td><input type="checkbox"/> Chest X-ray</td> <td></td> <td></td> </tr> <tr> <td><input type="checkbox"/> Vomiting</td> <td></td> <td></td> <td><input type="checkbox"/> Other _____</td> <td></td> <td></td> </tr> </tbody> </table>			Generalised		Pain	Respiratory	Renal	Circulatory	<input type="checkbox"/> Fever	<input type="checkbox"/> Anxiety	<input type="checkbox"/> Chest Pain	<input type="checkbox"/> Dyspnoea	<input type="checkbox"/> Haematuria	<input type="checkbox"/> Tachycardia	<input type="checkbox"/> Chills	<input type="checkbox"/> Itching (Pruritus)	<input type="checkbox"/> Abdominal	<input type="checkbox"/> Wheeze	<input type="checkbox"/> Haemoglobinuria	<input type="checkbox"/> Hypertension	<input type="checkbox"/> Rigors	<input type="checkbox"/> Edema (Site) _____	<input type="checkbox"/> Back/Flank Pain	<input type="checkbox"/> Cough	<input type="checkbox"/> Oliguria	<input type="checkbox"/> Hypotension	<input type="checkbox"/> Nausea	<input type="checkbox"/> Jaundice	<input type="checkbox"/> Infusion Site Pain	<input type="checkbox"/> Hypoxemia	<input type="checkbox"/> Other _____	<input type="checkbox"/> Raised JVP	<input type="checkbox"/> Urticaria	<input type="checkbox"/> Other _____	<input type="checkbox"/> Other _____	<input type="checkbox"/>		<input type="checkbox"/> Arrhythmias	<input type="checkbox"/> Flushing			<input type="checkbox"/> Bilateral Infiltrates on		<input type="checkbox"/> Other _____	<input type="checkbox"/> Restlessness			<input type="checkbox"/> Chest X-ray			<input type="checkbox"/> Vomiting			<input type="checkbox"/> Other _____		
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<input type="checkbox"/> Rigors	<input type="checkbox"/> Edema (Site) _____	<input type="checkbox"/> Back/Flank Pain	<input type="checkbox"/> Cough	<input type="checkbox"/> Oliguria	<input type="checkbox"/> Hypotension																																																			
<input type="checkbox"/> Nausea	<input type="checkbox"/> Jaundice	<input type="checkbox"/> Infusion Site Pain	<input type="checkbox"/> Hypoxemia	<input type="checkbox"/> Other _____	<input type="checkbox"/> Raised JVP																																																			
<input type="checkbox"/> Urticaria	<input type="checkbox"/> Other _____	<input type="checkbox"/> Other _____	<input type="checkbox"/>		<input type="checkbox"/> Arrhythmias																																																			
<input type="checkbox"/> Flushing			<input type="checkbox"/> Bilateral Infiltrates on		<input type="checkbox"/> Other _____																																																			
<input type="checkbox"/> Restlessness			<input type="checkbox"/> Chest X-ray																																																					
<input type="checkbox"/> Vomiting			<input type="checkbox"/> Other _____																																																					
Any Other(Specify) :																																																								
(C) Transfusion Product(s) Details*																																																								
Select*	Select Component	Select Indication	Date & Time of Issue of Blood Component	Date & Time of onset Transfusion	Unit Id (Transfused)	Blood Group	Volume Transfused (ml)	Expiry date of Blood Component	Manufacturer of Blood Bag	Batch / Lot No. of the Blood Bag	1st time/ repeat Transfusion																																													
<input type="checkbox"/>	Whole blood										<input type="checkbox"/>																																													
<input type="checkbox"/>	Packed Red blood cells (PRBC)										1st Time																																													
<input type="checkbox"/>	Buffy coat depleted PRBC																																																							
<input type="checkbox"/>	Leucofiltered PRBC																																																							
<input type="checkbox"/>	Random Donor platelets/ pooled										<input type="checkbox"/>																																													
<input type="checkbox"/>	Apheresis Platelets										Repeat 1 to 10																																													
<input type="checkbox"/>	Fresh Frozen Plasma																																																							
<input type="checkbox"/>	Cryoprecipitate										<input type="checkbox"/>																																													
<input type="checkbox"/>	Any Other _____										Repeat > 10																																													
Add New Plasma Product																																																								
Select	Plasma Product	Indication	Date of Administration	Manufacturer	Expiry Date of the Plasma Product	Batch No. / Lot No.	1st Time / Repeat																																																	
							<input type="checkbox"/> 1st Time																																																	
							<input type="checkbox"/> Repeat 1 to 10																																																	
							<input type="checkbox"/> Repeat > 10																																																	

(D) Investigations				
<input type="checkbox"/> Clerical Checks		Specify Error Found if any: _____		
Investigation	Pre-transfusion sample	Post-transfusion sample		
<input type="checkbox"/> Visual Check				
* <input type="checkbox"/> Repeat Blood Grouping	O+ /A+ /B+ /AB+ /O- /A- /B- /AB- /Others/Not Done	O+ /A+ /B+ /AB+ /O- /A- /B- /AB- /Others/Not Done		
* <input type="checkbox"/> Repeat Crossmatch	<input type="checkbox"/> Compatible <input type="checkbox"/> InCompatible <input type="checkbox"/> Not Done	<input type="checkbox"/> Compatible <input type="checkbox"/> InCompatible <input type="checkbox"/> Not Done		
* <input type="checkbox"/> Repeat Antibody screen	<input type="checkbox"/> Negative <input type="checkbox"/> Positive <input type="checkbox"/> Not Done	<input type="checkbox"/> Negative <input type="checkbox"/> Positive <input type="checkbox"/> Not Done		
<input type="checkbox"/> Antibody Identification				
* <input type="checkbox"/> Direct antiglobulin test	<input type="checkbox"/> Negative <input type="checkbox"/> Positive <input type="checkbox"/> Not Done	<input type="checkbox"/> Negative <input type="checkbox"/> Positive <input type="checkbox"/> Not Done		
<input type="checkbox"/> Hemoglobin				
<input type="checkbox"/> Plasma Hemoglobin				
<input type="checkbox"/> Urine hemoglobin				
<input type="checkbox"/> Bilirubin (Total/conjugated)				
<input type="checkbox"/> Platelet count				
<input type="checkbox"/> PT/INR				
* <input type="checkbox"/> Blood culture of Blood Bag	<input type="checkbox"/> Negative <input type="checkbox"/> Positive <input type="checkbox"/> Not Done	Specify Organism if positive _____		
* <input type="checkbox"/> Blood culture of Patient	<input type="checkbox"/> Negative <input type="checkbox"/> Positive <input type="checkbox"/> Not Done	<input type="checkbox"/> Negative <input type="checkbox"/> Positive <input type="checkbox"/> Not Done		
<input type="checkbox"/> Chest X-ray of the patient in case of suspected TRALI				
In case of Non-immune hemolysis (which of the following was the case?)				
<input type="checkbox"/> Hemolysis due to freezing of PRBC Units				
<input type="checkbox"/> Hemolysis due to inappropriate warming of PRBC Units				
<input type="checkbox"/> Hemolysis due to infusion of any other fluid through same BT set. Specify Fluid: _____				
<input type="checkbox"/> Mechanical damage				
In Case of ABO Mismatch (which of the following was the case?)				
<input type="checkbox"/> Wrong blood in tube				
<input type="checkbox"/> Grouping error				
<input type="checkbox"/> Labelling error				
<input type="checkbox"/> Wrong unit transfused				
(E) Nature of Adverse Reaction(s)*				
Select	Reaction	Date & Time of Onset of Reaction	Date & Time of Recovery	Outcome
<input type="checkbox"/>	Febrile Non Haemolytic Reactions (FNHTR) 1° C rise in temperature <input type="checkbox"/> 2° C rise in temperature <input type="checkbox"/> Only Chills & Rigors <input type="checkbox"/>			<input type="checkbox"/> 1. Death following the Adverse Reaction(s)
<input type="checkbox"/>	Allergic reaction			<input type="checkbox"/> 2. Recovered
<input type="checkbox"/>	Anaphylaxis			
<input type="checkbox"/>	Immunological Haemolysis due to ABO Incompatibility			<input type="checkbox"/> 3. Recovered with Sequelae
<input type="checkbox"/>	Immunological Haemolysis due to other Allo-Antibodies			
<input type="checkbox"/>	Non Immunological Haemolysis			<input type="checkbox"/> 4. Unknown
<input type="checkbox"/>	Hypotensive Transfusion Reaction			
<input type="checkbox"/>	Transfusion Related Acute Lung Injury (TRALI) Definite <input type="checkbox"/> Possible <input type="checkbox"/>			
<input type="checkbox"/>	Transfusion Associated Dyspnoea (TAD)			
<input type="checkbox"/>	Transfusion Associated Circulatory Overload (TACO)			
<input type="checkbox"/>	Transfusion Transmitted Bacterial Infection			
<input type="checkbox"/>	Transfusion Transmitted Parasitic Infection (Malaria)			
<input type="checkbox"/>	Post Transfusion Purpura			
<input type="checkbox"/>	Transfusion Associated Graft versus Host Disease (TAGvHD)			
<input type="checkbox"/>	Other Reaction (s) _____ Add New _____			
IMPUTABILITY ASSESSMENT				
(F) Imputability Assessment*				
S. No.	Reaction Term	Transfusion Product/ Component	*Imputability Assessment (Please mention from the below list)	
*Imputability: 1. Definite (Certain), 2. Probable (Likely), 3. Possible, 4. Unlikely (Doubtful), 5. Excluded, 6. Not Assessed				
Monthly Denominator Reporting Form *				
Hospital Code :		Month/Year:		
Blood Component		No. of Units Issued		
1) Fresh Frozen Plasma				
2) Whole Blood				
3) Packed Red Blood Cells (PRBC)				
4) Buffy Coat Depleted PRBC				
5) Leucofiltered PRBC				
6) Random Donor Platelets/ Pooled				
7) Apheresis Platelets				
8) Cryoprecipitate				
9) Any Other _____				

New Members Enrolled under Haemovigilance Programme of India (108)

Chhattisgarh

1. Jawaharlal Nehru Hospital and Research Centre, Bhilai, Durg

Gujarat

1. Blood Bank S.S.G. Hospital, Baroda, Vadodara
2. Indian Red Cross Society Voluntary Blood Bank, Dahod
3. Mehsana Jaycees Charitable Trust Sanchalit Voluntary Blood Bank, Mehsana
4. Shri Kutchi Leva Patel Education & Medical Trust Sanchalit MMPJ Hospital Blood Bank, Bhuj-Kutch
5. Blood Bank, Narayana Hrudayalaya Ltd., Narayana Multispeciality Hospital, Ahmedabad
6. Dr. Jivraj Mehta Smarak Health Foundation Blood Bank and Bakeri Medical Research Centre, Ahmedabad
7. Indian Red Cross Society C.R. Parikh Blood Bank, Kapadwanj, Kheda
8. Indian Red Cross Society, Branch Blood Bank, Anand
9. Jivan Jyot Blood Bank, Bhuj-Kutch
10. Nathani Voluntary Blood Bank, Rajkot
11. Blood Bank General Hospital Jam-Khanbhalia, Dwarka
12. Shree Jivanprakash Foundation Voluntary Blood Bank, Junagadh
13. Navkar Charitable Trust Blood Bank, Mahuva
14. Samved Voluntary Blood Bank and Research Center, Jamnagar
15. Late N. H. Ramani Memorial Voluntary Blood Bank, Aravli
16. GCS Medical College, Hospital and Research Centre, Ahmedabad
17. Blood Bank, C. U. Shah Medical College and Hospital, Surendranagar
18. General Hospital, Godhra Blood Bank, Godhra
19. Bhailal Amin General Hospital Blood Bank, Vadodara
20. Omkar Paramedical & Charitable Trust Blood Bank, Botad
21. M/s. "Cadila Healthcare Limited" Zydus Medical College and Hospital, Blood Bank, Dahod
22. Smt. Chandaben Mohanbhai Patel Blood Bank, Charusat Hospital, Anand
23. Parul Sevashram Hospital Blood Bank, Vadodara
24. Atma Vallabh Hospital Voluntary Blood Bank IDAR, Sabarkantha
25. Shantabaa Medical College & General Hospital Blood Bank, Amreli
26. Saurashtra Voluntary Blood Bank & Research Centre, Rajkot
27. GMERS Medical College & Hospital, Valsad
28. Navdeep Voluntary Blood Bank, Junagarh
29. Navjeevan Blood Bank & Clinical Laboratory, Junagarh
30. S.M.S. Multispeciality Hospital & Dr. M.K. Shah Medical College & Research Centre, Ahmedabad
31. Sanskar Blood Bank, Morbi
32. Smt. Shardaben Hospital, Ahmedabad
33. Savior Voluntary Blood Bank & Research Centre, Surat
34. Aasha Blood Bank & Component Centre, Porbandar
35. Dr. N.D. Desai Faculty of Medical Science & Research, Nadiad
36. Lions Club Bilimora Charitable Foundation Managed, Nanubhai Mavljbhai Patel Blood Bank & Dr. A.C. Mapara Blood Component Centre, Navsari
37. Hi-Tech Vol. Blood Bank, Aravalli

Haryana

1. Panchkula Welfare Trust (Regd) Charitable Diagnostic Center, Panchkula
2. Blood Bank, Columbia Asia Hospital, Palam Vihar, Gurugram

Himachal Pradesh

1. Maharishi Markandeshwar Medical College & Hospital, Solan

Jammu & Kashmir

1. Shri Mata Vaishno Devi Superspeciality Hospital, Reasi

Karnataka

1. HKE Society, Mahadevappa Rampure Medical College, Kalaburagi

Kerala

1. M.U.M. Hospital Blood Bank, Kottayam
2. Kozhikode District Co-operative Hospital Pvt Ltd., Kozhikode
3. Blood Bank, Government Medical College, Kannur
4. Malabar Institute of Medical Sciences Ltd. Kottakkal, Malappuram
5. St Thomas Hospital Blood Bank, Kottayam

Madhya Pradesh

1. All is Well Multispeciality Hospital, Burhanpur

Maharashtra

1. Ashtavinayak Blood Bank, Ahmednagar
2. Rotary D G Goenka Blood Bank, Andheri West Mumbai
3. Triumph Blood Bank, Thane
4. K. J. Somaiya Medical College and Hospital, Blood Bank, Mumbai
5. Masina Hospital Blood Bank, Mumbai
6. Pallavi Blood Bank, Govandi East, Mumbai

7. Sant Nirankari Blood Bank, Vileparle East, Mumbai
8. Kohinoor Hospital Blood Bank, Kurla West, Mumbai
9. Bombay Hospital Blood Bank, Mumbai
10. Jaslok Hospital & Research Center Blood Bank, Mumbai
11. Smt. S. R. Mehta & Sir K. P. Cardiac Institute Blood Bank, Mumbai
12. Datta Meghe Medical College, Dattatraya Blood Bank, Nagpur
13. Sarvodaya Hospital Samarpan Blood Bank, Ghatkopar West, Mumbai
14. Jagjivan Ram Hospital (Western Railway) Blood Bank, Mumbai
15. Blood Bank Government Medical College and Hospital, Chandrapur
16. Balasaheb Thackeray Blood Bank, Jogeshwari (E) Mumbai
17. Civil Surgeon Blood Bank, Alibag-Raigad
18. Bloodline Blood Bank, Thane (West)
19. Sanjeevani Blood Bank, Pune
20. Latur Blood Bank and Components, Latur
21. NKP Salve Institute of Medical Sciences & Research Centre & Lata Mangeshkar Hospital Blood Bank, Nagarpur
22. Adarsh Blood Bank, Aurangabad
23. Vande Mataram Samajik Sansthan's Jeevandhara Blood Bank, Kolhapur
24. MIMSR Medical College and Y.C.R. Hospital, Smt. Saraswati Karad Blood Bank, Latur

Meghalaya

1. Regional Blood Bank, Pasteur Institute, Shillong

New Delhi

1. Blood Bank Dr. Hedgewar Arogya Sansthan Govt. of N.C.T. of Delhi, Karkardooma
2. Regional Blood Transfusion Centre, Hindu Rao Hospital, Malka Ganj
3. Lal Bahadur Shastri Hospital, Govt. of National Territory of Delhi (GNCTD), Khichripur, Delhi

Orissa

1. Christian Hospital Blood Bank, Rayagada

Punjab

1. Preet Multispecialty Hospital & Surgical Centre, Ludhiana

Rajasthan

1. MP Birla Hospital & Research Center, MPBHRC, Chittorgarh
2. RAPS Hospital Blood Bank, Chittorgarh
3. Blood Bank Arihant Hospital and Research Sansthan, Bhilwara
4. American International Institute of Medical Sciences Blood Bank, Udaipur
5. Government General Hospital, Sirohi
6. Bhagwan Mahavir Hospital Component Blood Bank, Pali
7. Geetanjali Medical College and Hospital, Udaipur
8. Purohit Charitable Laboratory Sansthan, Sriganganagar
9. Life Line Blood Bank, Hanumangarh Jun.
10. M/s Sanjeevani Jan Sewa Samiti, Bikaner
11. Dr. S. S. Tania Medica College & Research Centre, Tania University, Sri Ganganagar
12. M/S Jeevan Jyoti Blood Bank, Bikaner
13. D. H. Blood Bank, Jalore

Tamil Nadu

1. Maruthi Blood Bank with Component Centre, Dharmapuri
2. Department of Transfusion Medicine Panimalar Medical College Hospital and Research Institute, Chennai
3. Cancer Institute (W/A), Chennai
4. Gudalur Adivasi Hospital Blood Bank, The Nilgiris
5. Government Hospital, Gudalur
6. The Madras Medical Mission Hospital, Chennai
7. Vijaya Hospital Blood Centre, Chennai
8. Maga Blood Bank, Salem
9. SRM Institute for Medical Science Blood Bank, Chennai
10. MGM Health Care Pvt. Ltd. MGM Blood Centre, Chennai

Telangana

1. Indian Red Cross Society, Blood Bank, Mancheril
2. Dr. Patnam Mahender Reddy Institute of Medical Sciences & Hospital, Rangareddy Dist.
3. Thalassemia and Sickle Cell Society Vuppala Venkaiah Memorial Blood Bank, Rangareddy Dist.

Uttar Pradesh

1. Noida Charitable Blood Bank, Sector-22, Noida
2. Blood Centre, Sahara Hospital, Lucknow

West Bengal

1. North 24 Parganas District Hospital Blood Bank, Kolkata

How to Enroll your Centre under HvPI

Who can enrol?

Head/ In-charge of Transfusion Medicine Department / Blood centres

How to enrol?

- 1) Head / Incharge of Transfusion Medicine Department / Blood Centre provides the necessary details to the National Coordinating Centre (NCC) - Haemovigilance Programme of India (HvPI) by sending the duly filled **Enrolment Form** either to NCC at National Institute of Biologicals, Ministry of Health & Family Welfare, Plot No. A-32, Sector-62, Institutional Area, NOIDA - 201 309 (U.P.) or via E-mail to NCC at haemovigilance@nib.gov.in
- 2) NCC verifies the details provided by the centre.
- 3) After verification, NCC issues the User Id and Password to the Head / Incharge of Transfusion Medicine Department / Blood Centre to access the (a) Haemo - Vigil Software (b) Donor-vigil Software for onward Submission of Transfusion Reactions Reports and Adverse Blood Donor Reaction Reports to NCC.

Download Enrolment Form from the site:-<http://nib.gov.in/Annexure7.pdf>

How to Report?

Reporting of Adverse Transfusion Reactions via Haemo-Vigil Software & Adverse Blood Donor Reactions in donation via Donor-Vigil Software.

- a) Centres enrolled under HvPI receives unique User Id & Password from NCC-HvPI, NIB.
- b) User Id & Password is same for both the Softwares i.e. Haemo-Vigil (to report adverse transfusion reactions) & Donor-Vigil (to report adverse donor reactions).
- c) Software(s) link is available at NIB website i.e. www.nib.gov.in under the tab of Haemovigilance Programme of India.
- d) The adverse reaction reports can be uplinked and submitted online via the above mentioned software(s) to NCC-HvPI, NIB.

www.nib.gov.in

WhatsApp Web web....

भारत सरकार | Government of India
Hindi Language

National Institute of Biologicals
(राष्ट्रीय जैविक संस्थान)
Ministry of Health & Family Welfare, Government of India
स्वास्थ्य एवं परिवार कल्याण मंत्रालय, भारत सरकार

Site Updated on : 12/06/2020

Home About Us Services Careers Directory Tech. Expertise Contact Us Sitemap Help

Director's Profile
Governance
Right to Information
Infrastructure
Participation
Laboratories
CDL Notification
CHDTL Notification
Govt. Analyst
Medical Device Testing Officer
Monographs
Guidance Document
Inventory Module
Tenders
Notice to Bidders on E-Tendering
Collaboration
Proficiency Testing
Haemovigilance Programme of India
Annual Report

Parliamentary Committee Meeting held on 22nd January, 2020 at Ashoka Hotel, New Delhi

National Institute of Biologicals (NIB) had been set up in 1992 as an apex autonomous Institute under the administrative control of the Ministry of Health & Family Welfare, Government of India for promoting and protecting human health through various activities assigned to it.

The mandate of the Institute includes ensuring provision of quality biological drugs i.e. Invitro diagnostics, Vaccines and Biotherapeutics, including therapeutic monoclonal antibodies used by patients suffering from cancer and autoimmune diseases by undertaking high end science based testing with R&D interface for application of science. One of the main functions of the Institute, as per by-laws 3.5.8, is to undertake research, establish linkages and exchange personnel with different institutions in India and abroad for furtherance of its mandate. Other subjects forming part of its mandate are

(i) developing and validating standards for quality control testing;
(ii) developing linkages with other National/ International institutions and keep abreast with worldwide scientific research and technological developments;
(iii) providing training facilities in quality control of biologicals;
(iv) assessing from time to time the availability of qualified manpower; and
(v) implementing and co-ordinating activities of Haemovigilance Programme of India.

Sample Receipt & Report Dispatch Unit	National Reference Standards	Sera Panel
Sample Receipt & Report Dispatch Unit (SRRDU) Procedure. SRRDU has been set up as an independent unit in 2008. The sample receipt unit being the...	National Reference Standards	Sera Panel

Notifications !!

Revised conditions for purchase of :
1. Sera Panels



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National Institute of Biologicals

Ministry of Health and Family Welfare,
Government of India

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NIB website: <http://nib.gov.in/>

Tel: 0120-2400072, 0120-2593612 Fax: 0120-2403014,

Toll free No. **1800-180-2588** [Mon to Fri (9:00 a.m. to 5:30 p.m.)]
query related to Haemovigilance Programme of India.

**For any other
Information/ Suggestions/Query related to
Haemovigilance Programme of India kindly contact:**
Dr. Akanksha Bisht, Scientist Grade-II &
Head-Haemovigilance Programme of India,
NIB, NOIDA at: haemovigilance@nib.gov.in

