

DISCLAIMER

This file is to support the local research time. The lay out of this file can differ from REDCap, which is leading. The data needs to be filled in REDCap by the local investigator

www.redcap.heart-institute.nl

1.	Patient Identification Number	
2.	Date:	

Patient demographics

1	Age	у	ears
2	Sex	☐ female ☐ male	
3	Are there principal or religious reasons		
	for this patient to refuse blood products?	☐ Yes ☐ No ☐ Unknown	
4	Does the patient have a 'do not		
	resuscitate' (DNR) order at ICU		
	admittance?	☐ Yes ☐ No ☐ Unknown	
5	Does this patient have a 'do not		
	intubate' (DNI) order at ICU admittance?	☐ Yes ☐ No ☐ Unknown	
6	What is the referring specialism?	☐ Cardiology	
		Cardiothoracic surgery	
		☐ Gastro-enteral surgery	
		☐ Gynecology	
		☐ Internal medicine	
		☐ Neurology	
		□ Neurosurgery	
		☐ Orthopedic surgery	
		☐ Pulmonology	
		☐ Surgery	
		☐ Trauma surgery	
		☐ Urology	
		Other, please specify:	
7	Type of admission	☐ Elective ☐ Emergency	
8	Referred from	Operating theater	
		☐ Emergency department	
		☐ General ward	
		☐ Other hospital	
		Other, please specify:	

9	What is the main reason for ICU	Shock
	admission?	Respiratory insufficiency
		☐ Acute brain injury
		☐ Metabolic disturbances including intoxication,
		acute kidney injury and liver failure
		☐ Monitoring post-surgery
		☐ In/out of hospital arrest
		 □ Trauma
		☐ Other
10	Did this patient undergo surgery within	
	24 hours prior to admittance, or during	
	the first 24 hours of ICU admittance?	☐ Yes ☐ No ☐ Unknown
	If yes, please specify <i>type of surgical</i>	Cardiothoracic
	patient	 ☐ Gastro-intestinal
		 ☐ Gynecological
		☐ Neurosurgical
		 ∏ Trauma
		☐ Other, please specify
11	Presence of shock on day of admission	☐ Yes ☐ No ☐ Unknown
	If yes, please specify type of shock	☐ Anaphylactic
		☐ Cardiogenic
		☐ Hypovolemic
		☐ Neurogenic
		☐ Obstructive including pulmonary embolism
		☐ Septic
	In case of septic shock, please specify	Abdominal
	focus	Lungs/pneumosepsis
		☐ Urinary tract
		Unknown
		☐ Other, please specify:
12	APACHE IV score	
	In case of non- cardiothoracic surgery	points
	EURO-score	
	In case of cardiothoracic surgery	points

13	Was the patient supported with	
	mechanical ventilation at ICU	
	admission?	☐ Yes ☐ No ☐ Unknown
14	Did the patient receive supportive	☐ Renal replacement therapy
	therapy at the day of admission?	☐ VA-ECMO
		☐ VV-ECMO
		☐ Other mechanical cardiac support (e.g.
		LVAD, Impella IABP)
		☐ Other please specify
		☐ None
15	Relevant comorbidities	☐ Acute coronary syndrome
		☐ Benign hematological disease
		☐ Chronic kidney failure
		☐ Chronic obstructive pulmonary disease
		☐ Heart failure
		☐ Hematological malignancy
		Liver failure
		☐ Solid tumor
		☐ Organ transplant*
		☐ Bone marrow transplant
		☐ Other, please specify
		☐ None
	If Organ transplant	☐ Heart
		☐ Kidney
		☐ Lung(s)
		☐ Pancreas
		Other:

17	Hemoglobin level prior to ICU admission	mmol/L	OR
		g/dL	OR
		g/L	OR
		☐ Not measured within 24 hours prior to ICU add	mission
	OR hematocrit level prior to ICU	L/L	OR
	admission	☐ Not measured within 24 hours prior to ICU add	mission
18	Platelet count prior ICU admission	x 10 ⁹ /L	OR
		g/dL	OR
		☐ Not measured within 24 hours prior to ICU add	mission
19	INR or Prothrombin time prior to ICU	INR	OR
	admission	PT seconds	OR
	If both available please use INR	☐ Not measured within 24 hours prior to ICU add	mission
20	Activated partial thromboplastin time	seconds	
	prior to ICU admission	☐ Not measured within 24 hours prior to ICU add	mission
DA'	Y 28: OUTCOME		
1	Number of days admitted to the ICU	days	
2	Patient outcome at day 28	☐ Death ☐ Alive ☐ Unknowr	1
3	If death	☐ Died during ICU admittance	
		☐ Died after ICU admittance	
		☐ Died after ICU admittance, outside hos	spital
	If unknown	☐ Discharged from ICU, status after disc	harge
		unknown	
		☐ Transferred to other ICU, current statu	s
		unknown	

☐ Discharged from hospital, status after

discharge unknown

☐ Discharged from hospital

☐ ICU readmission

General ward

☐ ICU

If alive: current location at day 28

Daily questionnaire

1	Date	
		dd mm yyyy
2	Day since admission	
	day of admission = day 0	
3	Estimated blood loss	mL
4	SOFA score	points
5	Patient is suffering from:	Acute coronary syndrome
		Acute respiratory distress syndrome
		☐ Acute kidney injury
		☐ Bone marrow failure
		☐ Sepsis
		Liver failure
		☐ Failure to wean
		☐ Ischemic cerebrovascular accident
		Hemorrhagic cerebrovascular accident
		☐ Gastro-intestinal bleeding
		☐ Retinal bleeding
		☐ None of the above
6	Does the patient currently receive	Renal replacement therapy
	supportive therapy?	☐ VA-ECMO
		☐ VV-ECMO
		Other mechanical cardiac support
		(e.g. LVAD, Impella IABP)
		Invasive mechanical ventilation
		☐ Non-invasive mechanical ventilation
		☐ Other please specify
		☐ None
7	Lowest hemoglobin or hematocrit value	mmol/L OR
	this day	g/dL OR
		g/L OR
		L/L OR
		☐ Not measured

8	Lowest platelet count this day	x 10 ⁹ /L OR
		g/dL OR
		☐ Not measured
	If platelet count < 150 x 10 ⁹ cells/L	☐ Disseminated intravascular coagulation (DIC
	Is this patient diagnosed with:	☐ Heparin induced thrombocytopenia (HIT)
		☐ Idiopathic thrombocytopenic purpura (ITP)
		☐ Thrombotic thrombocytopenic purpura (TTP)
		☐ None of the above
9	Highest INR/PT this day	INROR
	If both available please use INR	PT seconds OR
		☐ Not measured
10	Highest aPTT value this day	seconds
		☐ Not measured
11	Was point of care visco-elastic tests	Yes, if yes specify results:
	(e.g. ROTEM or TEG) used this day?	☐ Normal coagulation status
		☐ Fibrinogen deficiency
		☐ Platelet deficiency
		☐ Clotting factor deficiency
		☐ Hyperfibrinolysis
		□ No
12	Was iron administered today?	☐ Yes:mg
		□ No
		Unknown
13	Was EPO administered today	
		☐ Yes ☐ No ☐ Unknown
14	Transfused blood products or	☐ None
	administration coagulation factors or	☐ Red blood cells *
	antifibrinolytic agents?	☐ Platelets *
		☐ Plasma *
	* Please fill in transfusion questionnaire	☐ Coagulation factors or antifibrinolytic agents
	for each transfusion event.	(e.g. tranexamic acid, fibrinogen, prothrombin
		complex concentrate, vitamin K, cryoprecipitate
		Other please specify
		☐ None of the above

Red blood cell transfusion

1	Date	
		dd mm yyyy
2	Time of administration of blood product	☐ During office hours (7:30 am- 6:00 pm)
		☐ Outside office hours (6:00 pm- 06:30 am)
3a	Certification level of transfusion	☐ Intensivist*
	requestor	☐ Specialist non-intensivist practicing ICU*
		☐ Resident, specialist in training*
		☐ Student
		☐ Nurse
		Other, please specify:
3b	* Please specify medical specialism:	☐ Anesthesiology
		☐ Cardiology
		☐ Internal medicine
		☐ Neurology
		☐ Pulmonology
		☐ Surgery
		☐ Other, please specify:
4	Location of transfusion	☐ ICU
		☐ Operating theater
5	Number of transfused RBC units this	units
	transfusion episode	
6	Reason(s) for red blood cell transfusion	Low hemoglobin level
		☐ Active bleeding
		☐ Age patient
		☐ Coronary ischemia
		☐ Hemodynamic instability
		☐ Improvement of general state
		☐ Improvement peripheral O2 perfusion
		☐ Improve weaning
		☐ Before procedure/surgery
		☐ Other, please specify:

7	Hemoglobin or hematocrit value prior to		mmol/L	OR
	transfusion (<24 hours prior to		g/dL	OR
	transfusion, if not available leave open,		g/L	OR
	if both Hb and Ht available, note Hb		L/L	
	level)			
8	Hemoglobin or hematocrit threshold for		mmol/L	OR
	this patient		g/dL	OR
			g/L	OR
			L/L	
9	Hemoglobin or hematocrit value after		mmol/L	OR
	transfusion		g/dL	OR
			g/L	OR
			L/L	
10	Was there a physiological trigger other than Hb or Ht to transfuse? (multiple answers possible)	☐ Tachycardia ☐ Hypotension ☐ Arrhythmia ☐ Significant ECG changes ☐ SvO2 (mixed venous saturation of ☐ ScO2 (mixed central saturation of ☐ Lactate > 2mmol ☐ Acidosis ☐ Other, please specify ☐ None		
11	Characteristic of blood product (multiple answers applicable).	☐ Autologous☐ Cell salvaged☐ Irradiated☐ Washed☐ None of the above		

Platelet transfusion

1	Date	
		dd mm yyyy
2	Time of administration of blood product	☐ During office hours (7:30 am- 6:00 pm)
		☐ Outside office hours (6:00 pm- 06:30 am)
3	Certification level of transfusion	☐ Intensivist*
	requestor	☐ Specialist non-intensivist practicing ICU*
		☐ Resident, specialist in training*
		☐ Student
		☐ Nurse
		Other, please specify:
3b	* Please specify medical specialism:	☐ Anesthesiology
		☐ Cardiology
		☐ Internal medicine
		☐ Neurology
		☐ Pulmonology
		☐ Surgery
		☐ Other, please specify:
4	Location of transfusion	☐ ICU
		☐ Operating theater
5	Number of platelet units transfused this	
	Number of platelet units transfused this episode	Operating theater units
		units
5	episode	units
5	episode	units Active bleeding Prophylactic
5	episode	units Active bleeding Prophylactic (in the absence of an upcoming procedure)
5	episode	units Active bleeding Prophylactic (in the absence of an upcoming procedure) Upcoming procedure, please specify:
5	episode	□ Active bleeding □ Prophylactic (in the absence of an upcoming procedure) □ Upcoming procedure, please specify: □ As part of a clinical trial
5	episode	
5	episode Reason(s) for plasma transfusion	□ Active bleeding □ Prophylactic (in the absence of an upcoming procedure) □ Upcoming procedure, please specify: □ As part of a clinical trial
5	episode	
5	episode Reason(s) for plasma transfusion	□ Active bleeding □ Prophylactic (in the absence of an upcoming procedure) □ Upcoming procedure, please specify: □ As part of a clinical trial □ Results viscoelastic testing (ROTEM, TEG) □ Other, please specify:
5	episode Reason(s) for plasma transfusion	units Active bleeding Prophylactic (in the absence of an upcoming procedure) Upcoming procedure, please specify: As part of a clinical trial Results viscoelastic testing (ROTEM, TEG) Other, please specify: Abdominal drain placement

		☐ General surgery		
		Lumbar puncture		
		☐ Neurosurgery		
		☐ Organ biopsy (liver, kidney)		
		☐ Thorax drain placement		
		☐ Tracheostomy		
		Other, please specify:		
7	Platelet count prior to transfusion		x 10 ⁹ /L	OR
			g/dL	OR
		☐ Not measured		
8	Platelet count target for this patient		x 10 ⁹ /L	OR
			g/dL	OR
		☐ Not measured		
9	Platelet count after transfusion		x 10 ⁹ /L	OR
			g/dL	OR
		☐ Not measured		
10	Did the patient use antiplatelet drugs the	□ No		
	past 7 days?	☐ Yes, please specify		
		Abciximab		
		☐ Acetylsalicylic acid		
		☐ Carbasalatecalcium		
		Clopidogrel		
		Dipyradimol		
		☐ Ptifibatide		
		☐ Prasugrel☐ Ticagrelor		
		Tirofiban		
		Other, please specify:		
	If yes: was therapeutic anticoagulant			
	usage a trigger to transfuse plasma?	Yes No		
11	Was the decision to transfuse guided by viscoelastic test (ROTEM, TEG)?	☐ Yes ☐ No		

Plasma transfusion

1	Date	
		dd mm yyyy
2	Time of administration of blood product	☐ During office hours (7:30 am- 6:00 pm)
		☐ Outside office hours (6:00 pm- 06:30 am)
3	Certification level of transfusion	☐ Intensivist*
	requestor	☐ Specialist non-intensivist practicing ICU*
		☐ Resident, specialist in training*
		☐ Student
		☐ Nurse
		Other, please specify:
3b	* Please specify medical specialism:	☐ Anesthesiology
		☐ Cardiology
		☐ Internal medicine
		☐ Neurology
		☐ Pulmonology
		☐ Surgery
		☐ Other, please specify:
4	Location of transfusion	☐ ICU
		☐ Operating theater
5	Number of plasma units transfused this	units
	episode	
6	Reason(s) for plasma transfusion	☐ Active bleeding
		☐ Prophylactic
		(in the absence of an upcoming procedure)
		☐ Upcoming procedure, please specify:
		<u> </u>
		☐ As part of a clinical trial
		
		<u> </u>
	In case of upcoming procedure	☐ Results viscoelastic tests (ROTEM, TEG) ☐ Other, please specify:
	In case of upcoming procedure	☐ Results viscoelastic tests (ROTEM, TEG) ☐ Other, please specify:
	In case of upcoming procedure	☐ Results viscoelastic tests (ROTEM, TEG) ☐ Other, please specify:
	In case of upcoming procedure	☐ Results viscoelastic tests (ROTEM, TEG) ☐ Other, please specify:
	In case of upcoming procedure	☐ Results viscoelastic tests (ROTEM, TEG) ☐ Other, please specify:

		Lumbar puncture
		☐ Neurosurgery
		☐ Organ biopsy (liver, kidney)
		☐ Thorax drain placement
		☐ Tracheostomy
		☐ Other, please specify:
7	INR or PT prior to transfusion	INR OR
		PT seconds OR
		☐ Not measured
8	INR or PT target for this patient	INR OR
		PT seconds OR
		☐ Not measured
9	INR or PT after transfusion	INR OR
		PT seconds OR
		☐ Not measured
10	Did the patient use therapeutic	
	anticoagulant drugs the past 7 days	□ No
		Yes, please specify
		☐ DOAC (e.g. dabigatran, apixaban)
		Acenocoumarol
		☐ Fenprocoumon ☐ LMWH
		Heparin
		☐ Warfarin
		Other, please specify:
	If yes: was therapeutic anticoagulant	☐ Yes ☐ No
	usage a trigger to transfuse plasma?	
11	Was the decision to transfuse guided by viscoelastic test (ROTEM, TEG)?	☐ Yes ☐ No

Administration of fibrinogen or cryoprecipitate

1	Date				
		dd mm yyyy			
2	Time of administration of blood product	☐ During office hours (7:30 am- 6:00 pm)			
		Outside office hours (6:00 pm- 06:30 am)			
3a	Certification level of transfusion	☐ Intensivist*			
	requestor	☐ Specialist non-intensivist practicing ICU*			
		☐ Resident, specialist in training*			
		☐ Student			
		Nurse			
		Other, please specify:			
3b	* Please specify medical specialism:	Anesthesiology			
		☐ Cardiology			
		☐ Internal medicine			
		☐ Neurology			
		☐ Pulmonology			
		☐ Surgery			
		☐ Other, please specify:			
4	Location of transfusion	☐ ICU			
		☐ Operating theater			
5	Dosage administered this episode	gram (fibrinogen) OR			
		units (cryoprecipitate)			
6	Reason(s) for transfusion	☐ Active bleeding			
		☐ Prophylactic (in the absence of an upcoming procedure)			
		Upcoming procedure, please specify:			
		☐ As part of a clinical trial			
		☐ Results viscoelastic testing (ROTEM, TEG)			
		☐ Other, please specify:			
	In case of upcoming procedure				
		Abdominal drain placement			
		☐ Bone marrow biopsy			
		☐ Cardiothoracic surgery			
		☐ Central venous catheter			
		☐ General surgery			

		Lumbar puncture
		☐ Neurosurgery
		☐ Organ biopsy (liver, kidney)
		☐ Thorax drain placement
		☐ Tracheostomy
		☐ Other, please specify:
7	Fibrinogen level prior to transfusion	g/L OR
		☐ Not measured
8	Fibrinogen level target for this patient	g/L OR
		☐ Not measured
9	Fibrinogen level after transfusion	g/L OR
		☐ Not measured
10	Was the decision to transfuse guided	☐ Yes ☐ No
	by viscoelastic test (ROTEM, TEG)?	

Administration of tranexamic acid

1	Date					
		dd mm yyyy				
2	Time of administration of blood product	☐ During office hours (7:30 am- 6:00 pm)				
		☐ Outside office hours (6:00 pm- 06:30 am)				
3	Certification level of transfusion	☐ Intensivist*				
	requestor	☐ Specialist non-intensivist practicing ICU*				
		☐ Resident, specialist in training*				
		☐ Student				
		☐ Nurse				
		☐ Other, please specify:				
3b	* Please specify medical specialism:	☐ Anesthesiology				
		☐ Cardiology				
		☐ Internal medicine				
		☐ Neurology				
		☐ Pulmonology				
		☐ Surgery				
		☐ Other, please specify:				
4	Location of transfusion	☐ ICU ☐ Operating theater				
4 5	Location of transfusion Dosage administered this episode	☐ ICU ☐ Operating theater milligram				
		milligram				
5	Dosage administered this episode	milligram				
5	Dosage administered this episode	☐ Active bleeding ☐ Prophylactic				
5	Dosage administered this episode	☐ Active bleeding ☐ Prophylactic (in the absence of an upcoming procedure)				
5	Dosage administered this episode	☐ Active bleeding ☐ Prophylactic				
5	Dosage administered this episode	milligram ☐ Active bleeding ☐ Prophylactic ☐ (in the absence of an upcoming procedure) ☐ Upcoming procedure, please specify: ☐ As part of a clinical trial				
5	Dosage administered this episode	milligram Active bleeding Prophylactic (in the absence of an upcoming procedure) Upcoming procedure, please specify: As part of a clinical trial Results viscoelastic testing (ROTEM, TEG)				
5	Dosage administered this episode Reason(s) for transfusion	milligram ☐ Active bleeding ☐ Prophylactic ☐ (in the absence of an upcoming procedure) ☐ Upcoming procedure, please specify: ☐ As part of a clinical trial				
5	Dosage administered this episode	milligram Active bleeding Prophylactic (in the absence of an upcoming procedure) Upcoming procedure, please specify: As part of a clinical trial Results viscoelastic testing (ROTEM, TEG)				
5	Dosage administered this episode Reason(s) for transfusion	milligram Active bleeding Prophylactic (in the absence of an upcoming procedure) Upcoming procedure, please specify: As part of a clinical trial Results viscoelastic testing (ROTEM, TEG) Other, please specify:				
5	Dosage administered this episode Reason(s) for transfusion	milligram Active bleeding Prophylactic (in the absence of an upcoming procedure) Upcoming procedure, please specify: As part of a clinical trial Results viscoelastic testing (ROTEM, TEG) Other, please specify: Abdominal drain placement				
5	Dosage administered this episode Reason(s) for transfusion	milligram Active bleeding Prophylactic (in the absence of an upcoming procedure) Upcoming procedure, please specify: As part of a clinical trial Results viscoelastic testing (ROTEM, TEG) Other, please specify: Abdominal drain placement Bone marrow biopsy				
5	Dosage administered this episode Reason(s) for transfusion	milligram Active bleeding Prophylactic (in the absence of an upcoming procedure) Upcoming procedure, please specify: As part of a clinical trial Results viscoelastic testing (ROTEM, TEG) Other, please specify: Abdominal drain placement Bone marrow biopsy Cardiothoracic surgery				
5	Dosage administered this episode Reason(s) for transfusion	milligram Active bleeding Prophylactic (in the absence of an upcoming procedure) Upcoming procedure, please specify: As part of a clinical trial Results viscoelastic testing (ROTEM, TEG) Other, please specify: Abdominal drain placement Bone marrow biopsy Cardiothoracic surgery Central venous catheter				

		☐ Neurosurgery
		☐ Organ biopsy (liver, kidney)
		☐ Thorax drain placement
		☐ Tracheostomy
		Other, please specify:
10	Was the decision to transfuse guided by viscoelastic test (ROTEM, TEG)?	☐ Yes ☐ No

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Administration of vitamin K or prothrombin complex concentrate

1	Date					
		dd mm yyyy				
2	Time of administration of blood product	☐ During office hours (7:30 am- 6:00 pm)				
		☐ Outside office hours (6:00 pm- 06:30 am)				
3	Certification level of transfusion	☐ Intensivist*				
	requestor	☐ Specialist non-intensivist practicing ICU*				
		☐ Resident, specialist in training*				
		☐ Student				
		☐ Nurse				
		Other, please specify:				
3b	* Please specify medical specialism:	☐ Anesthesiology				
		☐ Cardiology				
		☐ Internal medicine				
		☐ Neurology				
		☐ Pulmonology				
		☐ Surgery				
		☐ Other, please specify:				
4	Location of transfusion	☐ ICU ☐ Operating theater				
5	Dosage administered this episode	IE OR				
		milligrams				
6	Reason(s) for transfusion	☐ Active blooding				
		☐ Active bleeding				
		Prophylactic				
		(in the absence of an upcoming procedure)				
		Upcoming procedure, please specify:				
		☐ As part of a clinical trial				
		Results viscoelastic testing (ROTEM, TEG)				
	In case of uncoming procedure	Other, please specify:				
	In case of upcoming procedure	☐ Abdominal drain placement				
		☐ Bone marrow biopsy				
		☐ Cardiothoracic surgery				
		☐ Central venous catheter				
		☐ General surgery				
		Lumbar puncture				
		<u> </u>				

			☐ Neurosurgery
			☐ Organ biopsy (liver, kidney)
			☐ Thorax drain placement
			☐ Tracheostomy
			☐ Other, please specify:
7	INR or PT prior to transfusion	INR	OR
		PT	seconds OR
		│ □ No	t measured
8	INR or PT target for this patient	INR	OR
	G I	PT	seconds OR
		│ □ No	t measured
9	INR or PT after transfusion	INR	OR
	That Carlo and a carlo action	PT	seconds OR
		l	t measured
10	Did the patient use therapeutic	□No	
10	·		s, please specify
	anticoagulant drugs the past 7 days		o, product opening
			☐ DOAC (e.g. dabigatran, apixaban)
			Acenocoumarol
			☐ Fenprocoumon☐ LMWH
			Heparin
			☐ Warfarin
			Other, please specify:
	If yes: was therapeutic anticoagulant	 	es 🗆 No
	usage a trigger to transfuse plasma?		
11	Was the decision to transfuse guided by viscoelastic test (ROTEM, TEG)?	☐ Ye	es 🗌 No

Massive transfusion protocol

1	Date				
		dd mm yyyy			
2	Time of administration of blood product	☐ During office hours (7:30 am- 6:00 pm)			
		☐ Outside office hours (6:00 pm- 06:30 am)			
3a	Certification level of transfusion	☐ Intensivist*			
	requestor	☐ Specialist non-intensivist practicing ICU*			
		☐ Resident, specialist in training*			
		Student			
		☐ Nurse			
		Other, please specify:			
3b	* Please specify medical specialism:	☐ Anesthesiology			
		☐ Cardiology			
		☐ Internal medicine			
		☐ Neurology			
		☐ Pulmonology			
		☐ Surgery			
		☐ Other, please specify:			
4	Location of transfusion	☐ ICU ☐ Operating theater	☐ ICU ☐ Operating theater		
5	Number of transfused RBC units this				
	transfusion episode	units			
6	Number of transfused platelet units this	units			
	transfusion episode	units			
7	Number of transfused plasma units this	units			
	transfusion episode	units			
8	Hemoglobin or hematocrit value prior to	mmol/L OR)		
	transfusion (<24 hours prior to	g/dL OR	2		
	transfusion, if not available leave open)	g/L OF	₹		
		L/L			
9	Hemoglobin or hematocrit threshold for	mmol/L OR)		
	this patient	g/dL OR	?		
		g/L OF	₹		
		L/L			

10	Hemoglobin or hematocrit value after		mmol/L	OR
	transfusion		g/dL	OR
			g/L	OR
			L/L	
11	Platelet count prior to transfusion		x 10 ⁹ /L	OR
			g/dL	OR
		☐ Not measured		
12	Platelet count target for this patient		x 10 ⁹ /L	OR
			g/dL	OR
		☐ Not measured		
13	Platelet count after transfusion		x 10 ⁹ /L	OR
			g/dL	OR
		☐ Not measured		
14	INR or PT prior to transfusion	INR		OR
		PT	seconds	OR
		☐ Not measured		
16	INR or PT target for this patient	INR		OR
		PT	seconds	OR
		☐ Not measured		
17	INR or PT after transfusion	INR		OR
		PT	seconds	OR
		☐ Not measured		
18	Was the decision to transfuse guided by viscoelastic test (ROTEM, TEG)?	☐ Yes ☐ No		
19	Following factors given (multiple	□ None		
	answers possible)	☐ Aprotinin		
		☐ Cryoprecipitate		
		☐ Factor VIIa		
		☐ Factor XIII		
		☐ Fibrinogen		
		☐ Novoseven (eptacog alfa)		
		☐ Prothrombin complex concentrate		
		☐ Tranexamic acid		