The TP role in Belgium

Jana Vanden Broeck





TP2024 (15 May)



80%







TP: what & why?



Goal: to improve quality and safety of transfusion

&

support the implementation of PBM

Haemovigilance

Policy & procedure development/implementation

Audit/compliance monitoring

Education

Patient information

Reducing iatrogenic anemia Preop anaemia management

. . .





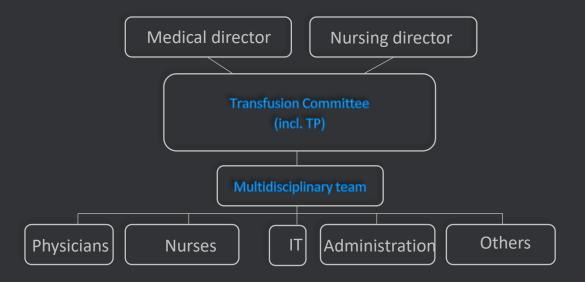


TP – national infrastructure: Belgium -

Continuous funding of hospitals to improve transfusion quality system since 2014: 4.340.000 EUR



⇒ For each hospital: lump sum 10.200 EUR + amount calculated based on blood consumption & hospital beds







Belgian "structure"

• Federal:









Communities:



Professional associations:



Belgian Society of Anesthesiology, Resuscitation, Perioperative medicine and Pain management



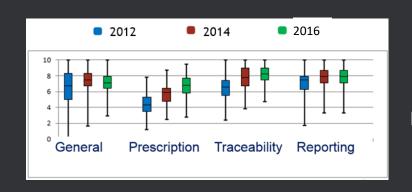
Scientific Association Transfusion Flanders

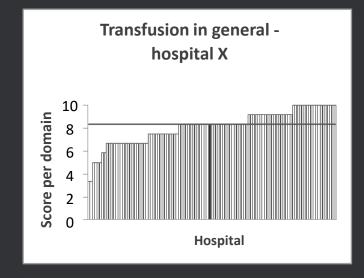




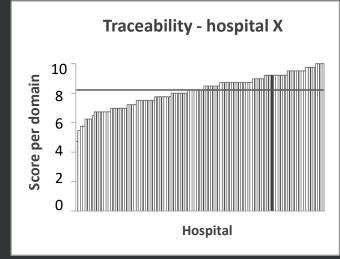
National survey 2012-16: quality of transfusion practice

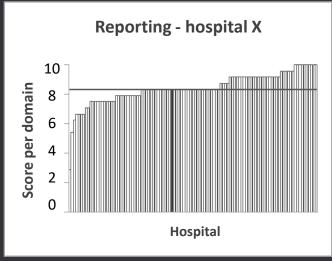
Benchmarking









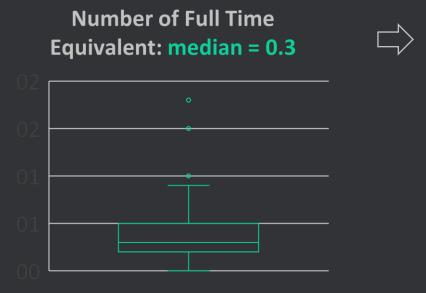




Number of TPs in Belgium

• 2020: In 88.5% of hospitals a TP is appointed (2016: 92.5%, 2014: 80%)

Hospitals with TP (n=85):



29 hospitals have a TP with allocated time to work on PBM



BeQuinT?









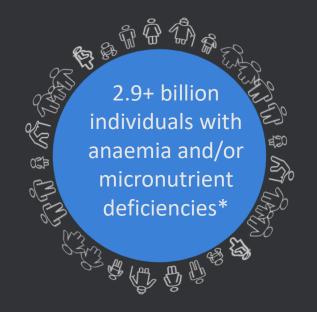








Blood is costly



Anaemia is a widespread problem



Guaranteeing blood supply





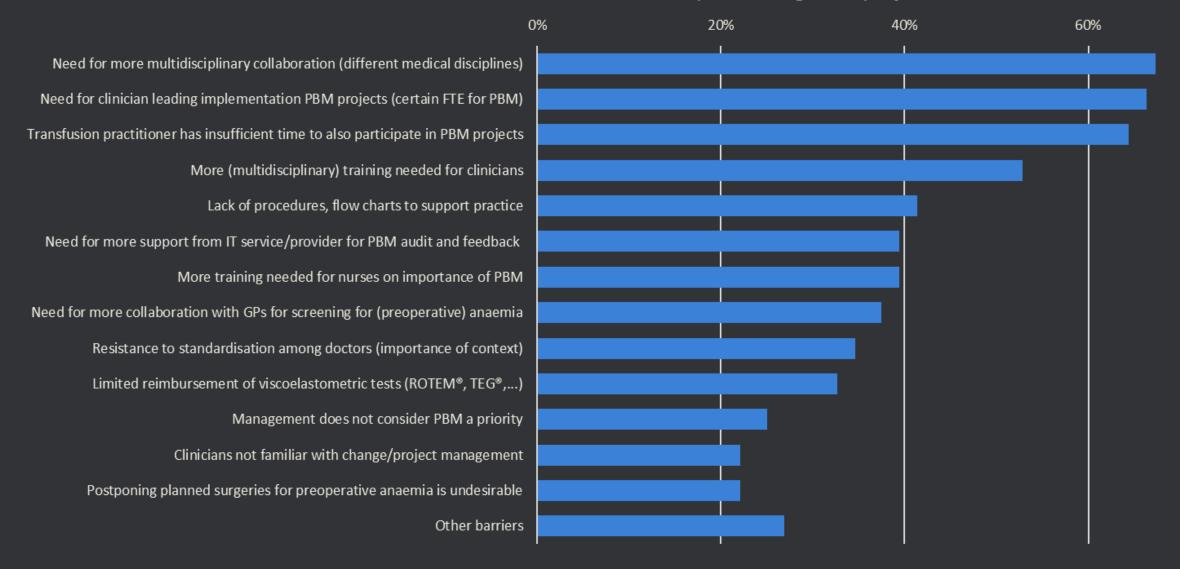
1. Improving quality of transfusion practice

2. Optimising blood use ⇒ implementing PBM





Encountered barriers to implementing PBM projects in 2020-2023







Strategies:

Structural: legislation & financial

Data collection, analysis & feedback

Education

Guidance







Data collection, analysis & feedback



Need for national PBM benchmarking



Haemovigilance benchmarking

[FAGG?]



National database antibodies







Guidance

O negative RBC

[Hoge Gezondheidsraad]

PBM in obstetrcs

PBM for GPs





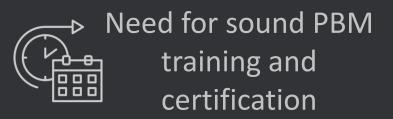
National blood shortage plan [+ stakeholders]







Education







Education

Webinars since pandemic

- 1. Results PBM survey BeQuinT + 2 PBM experts (UK)
- 2. Optimizing the perioperative management of anti-coagulants and anti-aggregants
 - 3. Single unit transfusion policy of RBC
 - 4. Visco-elastic testing as a PBM tool in today clinical practice
 - 5. How to manage preoperative anaemia?
 - 6. Blood conservation strategies
 - 7. Reduction of iatrogenic anaemia
 - 8. PBM in obstetrics





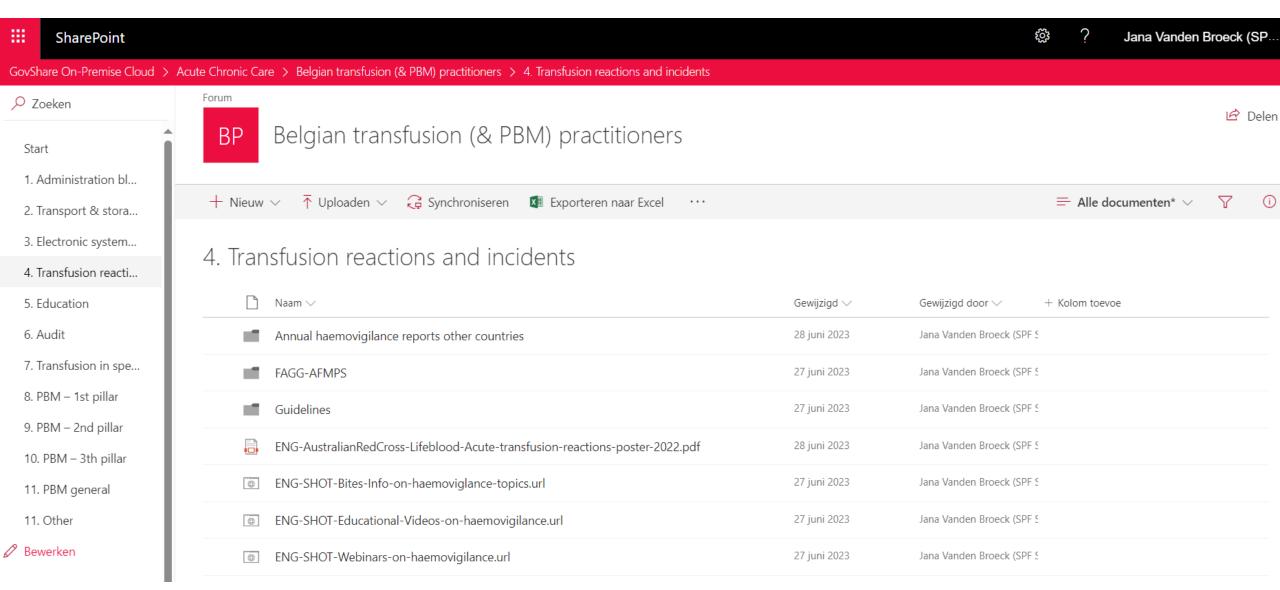
Interactive workshops for TPs

- 1. Retrospective analysis of transfusion incidents
- 2. (PBM) project management
- 3. Local transfusion audit





SharePoint for TPs



SharePoint for TPs



Start

- 1. Administration bloo...
- 2. Transport & storage
- 3. Electronic systems (...
- 4. Transfusion reaction...
- 5. Education
- 6. Audit
- 7. Transfusion in specif...
- 8. PBM 1st pillar
- 9. PBM 2nd pillar
- 10. PBM 3th pillar
- 11. PBM general
- 11. Other

Bewerken



Agenda

+ Gebeurtenis toevoegen

23 JUN.

ISBT Congres zo 23 jun., Hele dag Alles weergeven

Forum

Belgian transfusion (& PBM) practitioners

KOPPELINGEN BEWERKEN

Forum

• nieuwe discussie

Recent Mijn gesprekken Onbeantwoorde vragen ...

Beste, Zijn er ziekenhuizen waarbij ontdooid plasma wordt meegeven tijdens een MTP?M... Door Sonja Vanhaeren | 18 maart

IZ-191-1

Bonjour à tous,Je cherche un contact chez Dedalus pour le logiciel Kalilab.Quelqu'un utili... Door Carine Maggetto | 16 februari

Bloedproducten en TPN via een PICC katheter

Beste, in de richtlijnen van bloedtransfusie wordt beschreven dat men via een katheter m...

Door Sonja Vanhaeren | 15 november 2023

Bloedafhaling door zorgkundigen

o Goeiemorgen allemaal, bij ons in het ziekenhuis luidt het protocol voor bloedafhaling ... Door Heidi Cardinaels │ Nieuwste antwoord door Sonja Vanhaeren │ 15 november 2023

New documents

In deze discussie zal ik meedelen welke documenten ik heb opgeladen op deze Sharepoi...

Door Jana Vanden Broeck (SPF Santé Publique - FOD Volksgezondheid) | Nieuwste antwoord door

Jana Vanden Broeck (SPF Santé Publique - FOD Volksgezondheid) | 18 oktober 2023



Short survey (new) TPs when signing up for SharePoint

TPs could contact each other with questions on specific topics. On which topic(s) could fellow TPs contact you for advice?

Administration of blood comp.	Transport and storage of blood comp.	Electr.	Transf. reactions		_	_		Transf. in haemato oncology	with (solid) organ	PBM: screening and treatment of anaemia	PBM: manage ment of massive haemo- rrhage	PBM: cell	PBM:	Data analysis (transf./P BM) (indicators graphs, etc.)
Arrival Character														
and a second														
Maria Maria														
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Andrea Andrea														
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SOLE SALES														

Short survey (new) TPs when signing up for SharePoint

I would like to contribute the following (for Sharepoint and/or subsequent meetings)

- Protocol about:
 - transport & storage of blood components
 - scanning of blood components (at least before start transfusion)
 - transfusion reactions
 - informed consent transfusion
- **PPT** for training:
 - transfusion basics for nurses
 - scanning of blood components, electronic order/lab system
- Contribute to document 'Frequently Asked Questions' about:
 - blood scanning (Cybertrack, GLIMS, HiX,...)
 - Informed Consent for transfusion
 - practical aspects of implementing preoperative screening and treatment of anaemia
- Give a presentation on 1 of my projects to other TPs
- None of the above







Median of 0.3 FTE for TP role

Haemovigilance

Policy & procedure development/implementation

Audit/compliance monitoring

Education

Patient information

Reducing iatrogenic anemia Preop anaemia management

...

⇒ Project & time management!







Project charter

- Projects summary
- 2. SCOPE
- 2.1 Business value
- 2.2 Project goals
- 2.3 Initial project requirements
- 2.4 Iniatial project description
- 2.4.1 Project bounderies
- 2.4.2 Main deliverables
- TIME Initial milestones
- 4. COST Initial budget
- 5. PEOPLE Main stakeholders
- 6. RISKS General project risks

Project scope document

- 1. Description of the product scope
- 2. Project deliverables & acceptation criteria
- 3. Project exclusions
- ⇒ Example: latrogenic anemia







Initial project requirements

- 1) A baseline audit of iatrogenic blood loss is realised including at least the following parameters:
 - o daily phlebotomy loss per patient, cumulative total of blood sample volume, hemoglobin on admission, hemoglobin on discharge, nadir hemoglobin during hospitalization/audit, number of RBC units transfused per patient
- 2) The feasibility of at least the following strategies is evaluated:
 - rationalisation blood tests and collections,
 - o smaller blood tubes & small volume equipment.

Other strategies can be: return of void volumes (closed system sampling), removal of sampling lines, non-invasive technics and Point-Of-Care.

This feasibility study should include the budgetary impact (costs) of each strategy.

- 1) The feasibility report includes specific KPIs estimating the effect of the above strategies on the reduction of blood loss by phlebotomy:
 - Intermediate outcomes: evolution blood loss from phlebotomy (reduction?)
 - Health outcomes: evolution of decline in haemoglobin levels, iatrogenic anaemia and need for transfusion (reduction?)
- 2) The transfusion committee is regularly informed/updated about the project.

(According to local agreements, approval for the project is obtained)

3) The TP plans and monitors this project in collaboration with stakeholders of the laboratory, clinicians, nurses and phlebotomists.







prosource.be

Training (PBM) project management

Initial milestones

Milestone	Description	Date
M1 - Project start	Project charter approved	1 December 2021
M2 - Presentation of baseline audit	Presentation to the transfusion committee	15 June 2022
results		
M3 - Presentation of the feasibility	Presentation to the transfusion committee	15 October 2022
report (incl. KPIs)		
M4 - Selection of strategies to reduce	Selection based on the feasibility report and final approval	15 October 2022
iatrogenic blood loss	by chair of transfusion committee and head of the lab	
M5 - Publication of new/modified	Publication in the digital document management system	1 January 2022
procedures/policies	and communication to the involved stakeholders	
M6 - Start education of stakeholders who	First training session	1 January 2022
have to implement the new strategies to		
reduce iatrogenic blood loss		
M7 - Modified electronic lab tests order	Modifications to the Laboratory Information System	1 March 2023
(sets) if applicable/possible	to facilitate the rationalisation blood tests and collections	
M8 - Presentation of the post-	Presentation to the transfusion committee and other	30 September 2023
intervention(s) audit results	involved stakeholders	







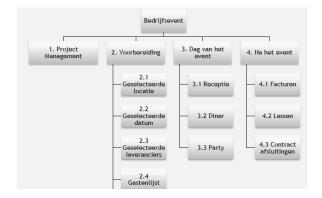
Deliverable	Acceptation criteria
Baseline audit report about iatrogenic blood loss and	Including patient demographics (sex, age, weight, etc.)
anaemia	Including at least the following parameters:
	daily phlebotomy loss per patient, cumulative total of blood sample volume,
	hemoglobin on admission, hemoglobin on discharge, nadir hemoglobin during
	hospitalization/audit, number of RBC units transfused per patient.
Feasibility report about the proposes strategies to	Including at least the feasibility of the following strategies:
reduce iatrogenic blood loss:	o rationalisation blood tests and collections (which tests can be realised on
	which blood tubes, minimum blood volume required for each test, which
	tests (+ frequency) are necessary?)
	o smaller blood tubes & small volume equipment,
	Other strategies can be:
	o return of void volumes (= closed system sampling),
	o removal of sampling lines,
	o Point-of-Care testing.
	• Including the budgetary impact (costs) of each strategy.
	• Including specific KPIs estimating the effect of the above strategies on the
	reduction of blood loss by phlebotomy.
New policy/modified protocols for selected strategies	Format according to hospitals' template
	Validated by some key persons and communicated to stakeholders
Training for involved clinicians, nurses, lab technicians and	• Including a formal assessment with minimum score 7.5/10.
phlebotomists	
Modified electronic lab tests order (sets) if applicable/	Modifications to the Laboratory Information System to facilitate the
possible	rationalisation blood tests and collections
Implementation of selected (feasible) strategies to reduce	Ensure policy & modified protocols are followed
iatrogenic blood loss	Ensure Lab. IS systems are implemented and validated
	Ensure Lab material is available for the Labs
Availability of (lab) equipment	Ensure necessary (Lab) equipment is ordered for start-up
Post-intervention(s) audit report about iatrogenic blood	Structure similar to baseline audit report
loss and anaemia	Including comparison with baseline audit results and targeted KPIs
	Including suggestions for improvement actions
	Including suggestions for improvement actions



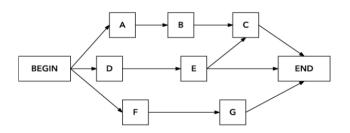


Exercises in groups of +-4 persons:

1) Project planning: Make a Work Breakdown Structure of the Project "latrogenic anaemia" (30 minutes)



2) Project planning: order the activities: Start to make a Precedence Diagram for the Project "latrogenic anaemia" (45 minutes)



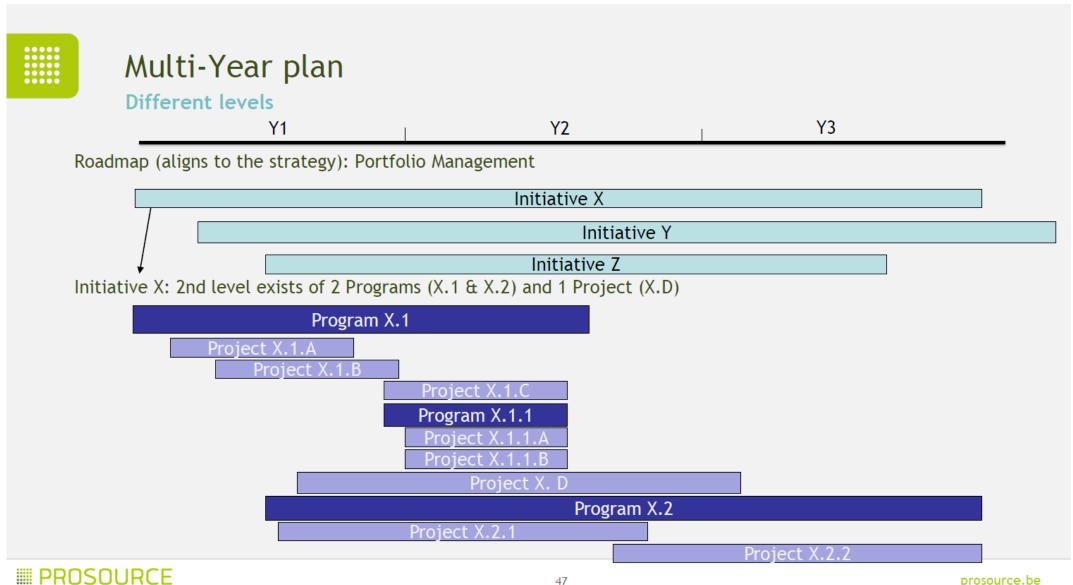
3) Make a Risk analysis for the Project "latrogenic anaemia" (30 minutes) Risk, Probability, Impact, PI-index, Effect/Impact of the Risk, Cost if the Risk occurs,

Risk Owner, Strategy, Concrete Mitigated action, Cost of the action,...





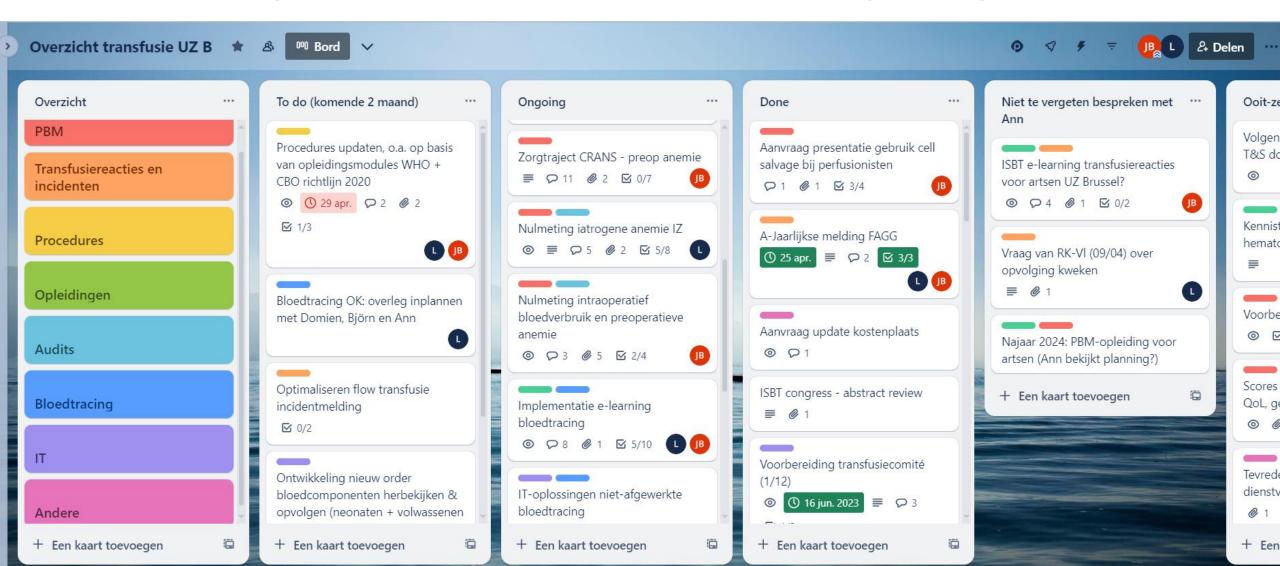






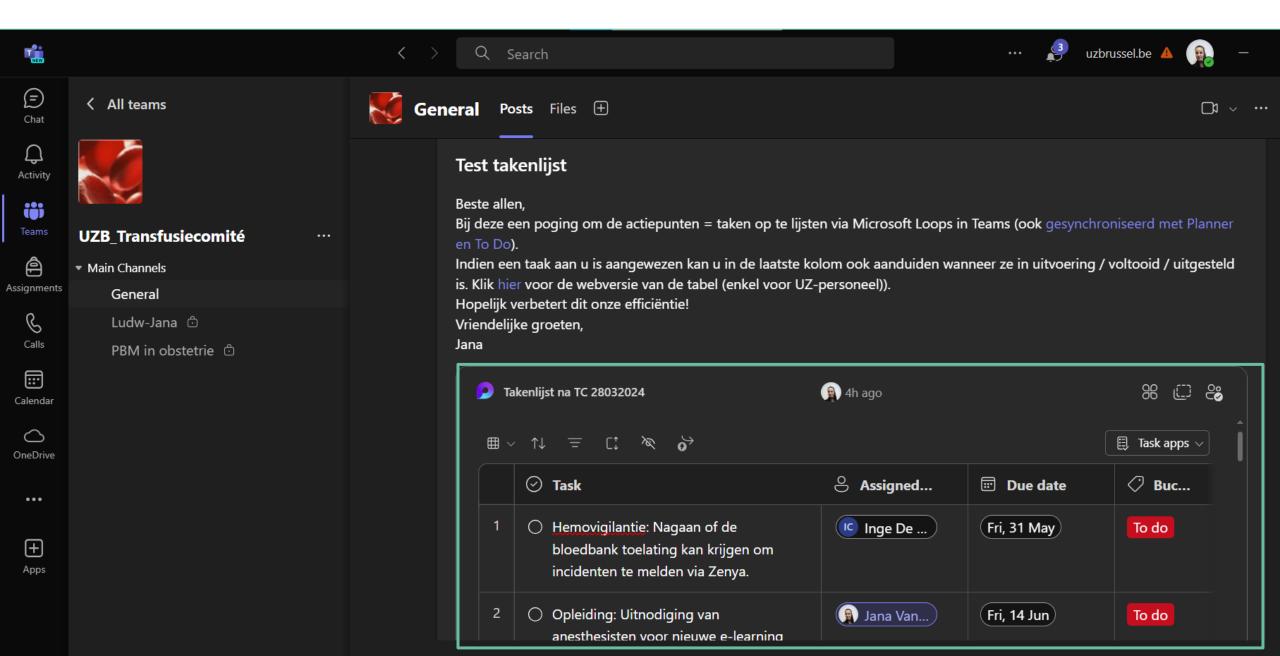


Some examples of the TP role in University Hospital Brussels



Time management: Microsoft Loop component in Teams group for actions by Transfusion Committee

YouTube: <u>How to use Loop components in Teams | Microsoft 365 TimeSavers</u>



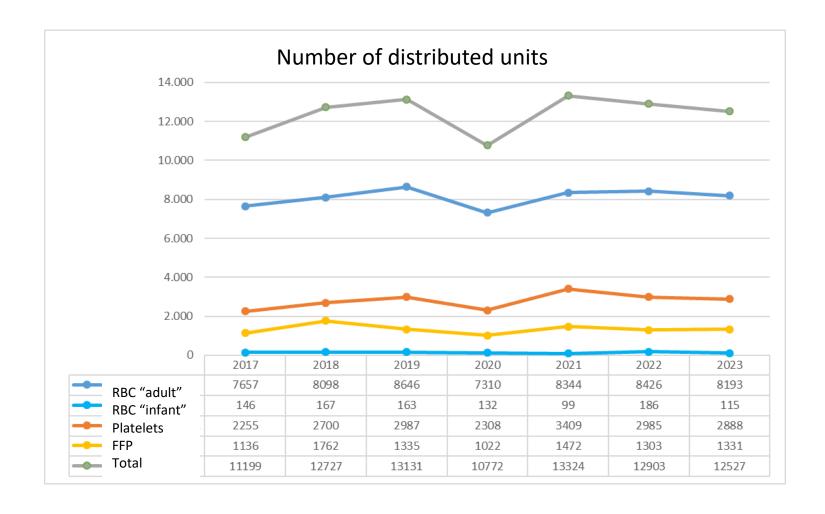
Some examples of the TP role in University Hospital Brussels

- 1) Haemovigilance charts: annual overview
- 2) Training: e-learning blood tracking/scanning
- 3) Communication on single-unit transfusion
- 4) Audit massive haemorrhage, flowchart MHP
- 5) Data cell salvage
- Data PBM (in obstetrics)
- 7) Collaborations as Subject Matter Expert with IT: mock-ups...





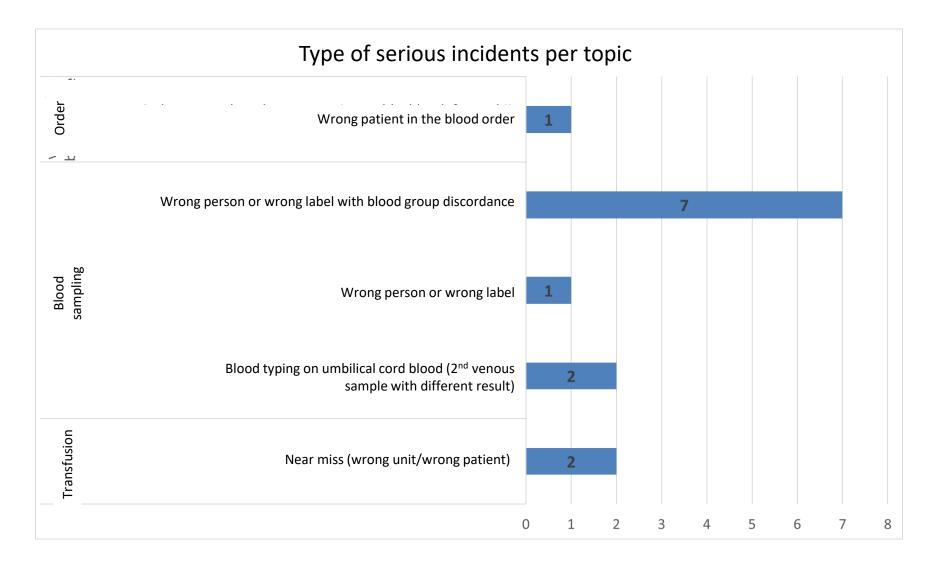
BLOOD USE BY UNIVERSITY HOSPITAL BRUSSELS







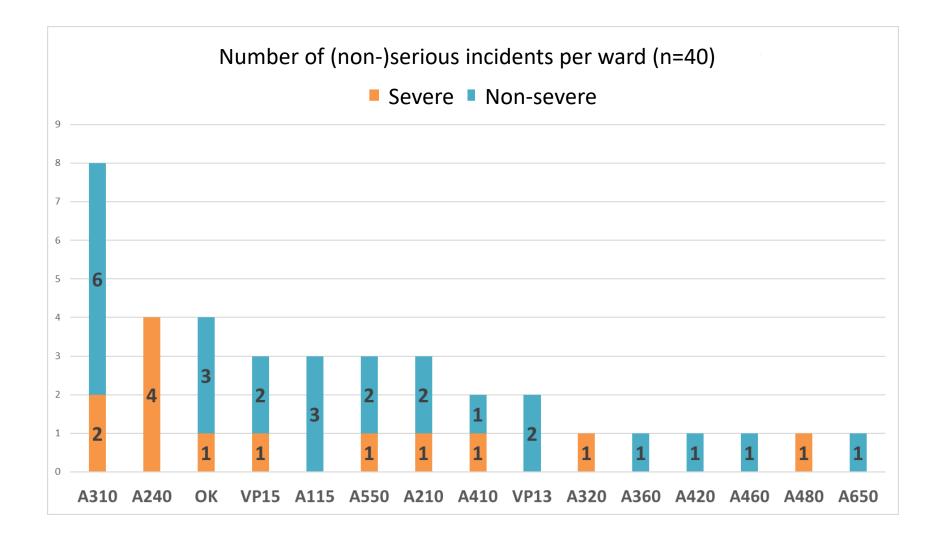
••• HAEMOVIGILANCE – OVERVIEW INCIDENTS 2023







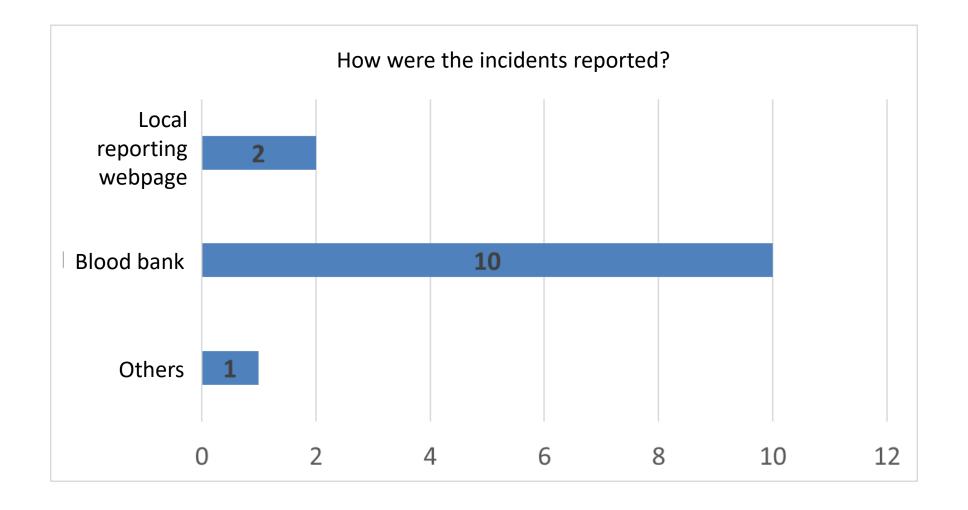
HAEMOVIGILANCE – OVERVIEW INCIDENTS 2023







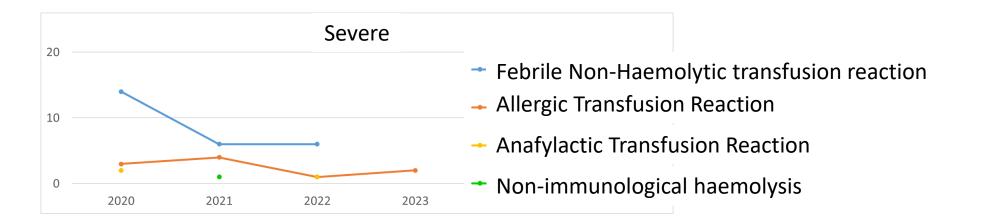
••• HAEMOVIGILANCE – OVERVIEW INCIDENTS 2023

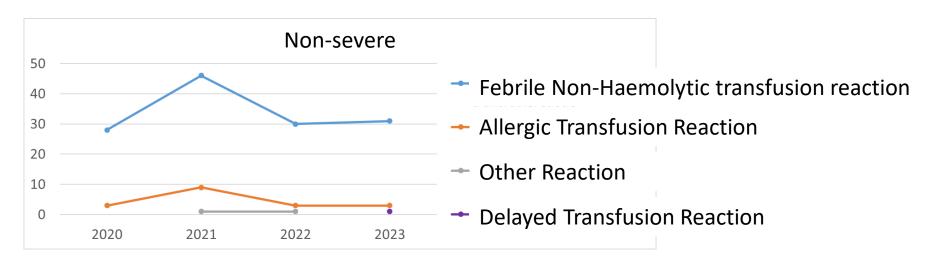






••• HV - OVERVIEW TRANSFUSION REACTIONS: EVOLUTION PER TYPE

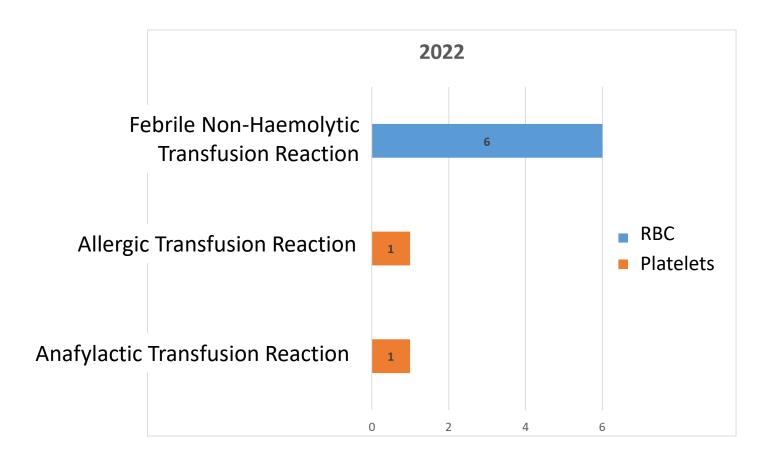


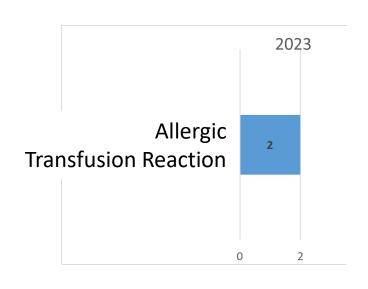






HV -TYPE OF SERIOUS TRANSFUSION REACTIONS PER BLOOD COMP.



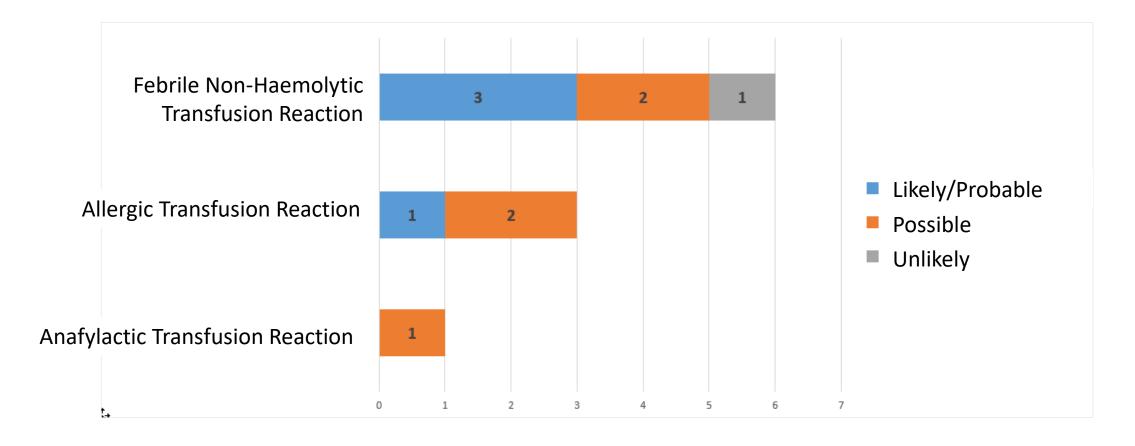






••• HV -IMPUTABILITY PER TYPE OF TRANSFUSION REACTION

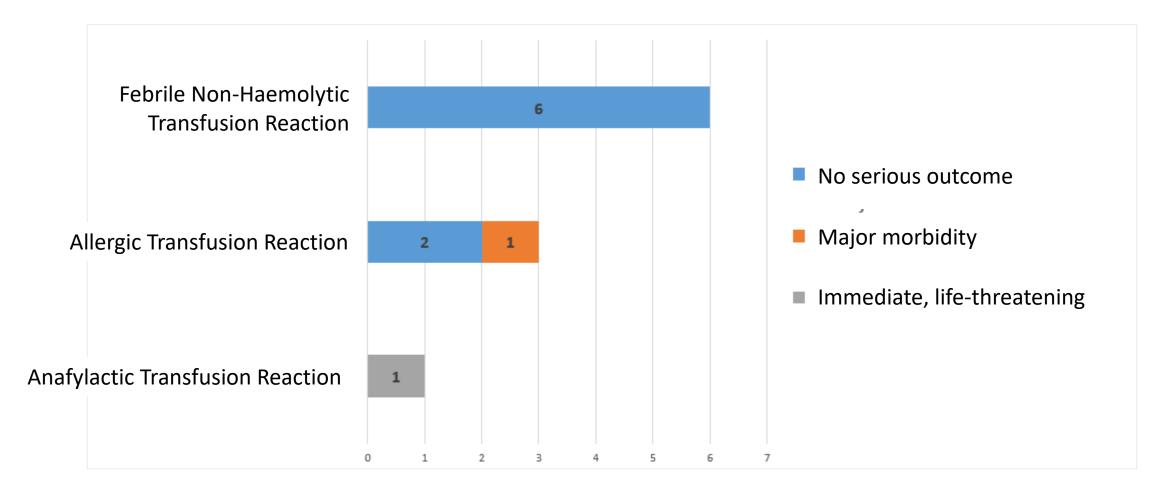
(2022-23)





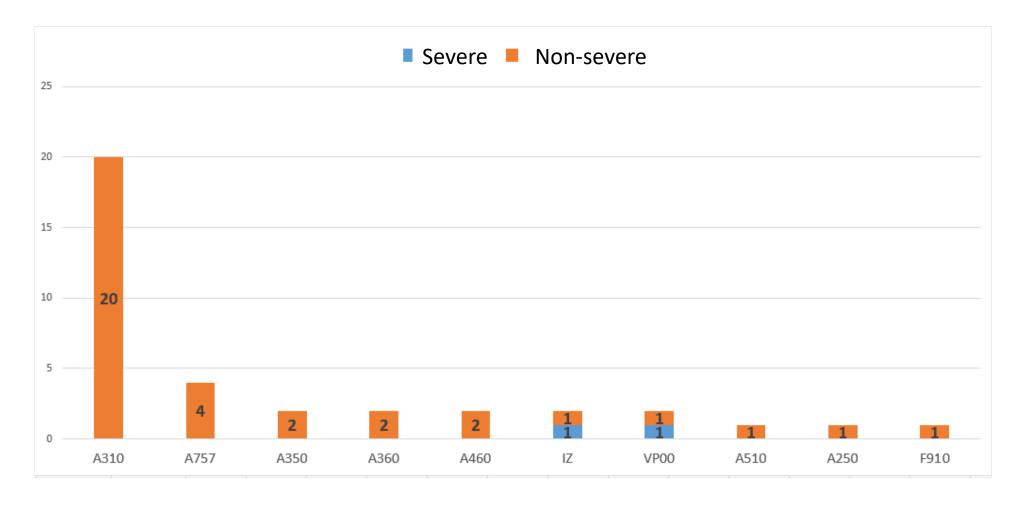
•• HV – OVERVIEW TR: SEVERITY PER TYPE OF SERIOUS REACTION

(2022-23)





HV –NUMBER OF (NON) SERIOUS TRANSF. REACTIONS PER WARD







TRAINING: E-LEARNING BLOOD TRACKING/SCANNING

- 1) Blood is costly
- 2) Correct storage and transport of blood components
- 3) Blood tracking
 - What is blood tracking?
 - Blood tracking at the University Hospital Brussels
 - What to do in case of a technical problem
 - When to record which parameters during transfusion
 - What if blood components have to be returned to the blood bank?







UZ Brussel -Bloedtracing verpleegkundige n en vroedvrouwen

 \equiv

0

13% COMPLETI

▼ BLOED IS KOSTBAAR

Bloed is kostbaar

Correcte bewaring en transport van bloedcomponenten

▼ BLOEDTRACING

Wat is bloedtracing?

Welke controles? Wat bij technisch probleem scanner?

Wanneer welke parameters opnemen tijdens transfusie?

Teruggave van bloedcomponenten

Welke controles? Wat bij technisch probleem scanner?

d-teach online training

Herinner je je Thomas? Het is bijna ochtend en zijn shift zit er bijna op. Het is druk op de afdeling. Patiënt Mika moet bloed krijgen en er is net een nieuwe patiënt, Louise, bijgekomen die eveneens bloed moet ontvangen. Logistiek medewerkster Anne gaat op vraag van Thomas naar de bloedbank om de bestelde bloedcomponenten te gaan halen.



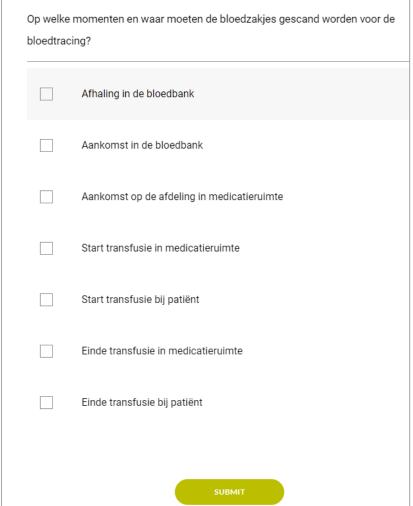
Anne scant de unit voor "Afhaling" en "Aankomst" en verwittigt Thomas dat het bloed in de koelkast



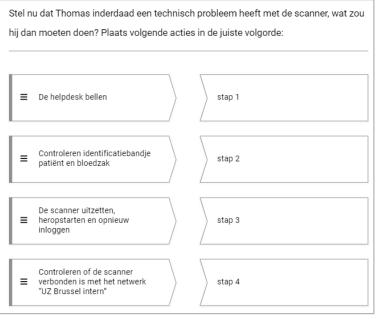




TRAINING: INTERACTIVE E-LEARNING

















ge-unit distribution of RBC



stable
normovolemic
adult patients
without clin. sign. bleeding
>1 day hospitalisation



Non applicable

- RBC for OR, ambulatory ward haemato-oncol. and gastro-enterology
- acute bleeding
- apheresis/exchange transf.
- infants or children



Order 1 unit (exceptionally more units allowed if necessary)



1 unit per patient released by blood bank



immediate* transfusion of 1 unit RBC



Re-assess patient: laboratory and/or clinical assessment

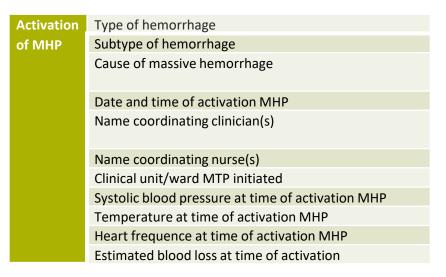


Other units max. 24h stand-by in blood bank

*ICU: 1 RBC max. 24h in blood fridge (+2 à +6°C)

••• AUDIT MASSIVE HAEMORRHAGE TEMPLATE

Demographics, patient	Medical record number					
characteristics	Admission number					
	Admission date : start & end					
	Age					
	Sex					
	ВМІ					
	Ethnicity					
	Blood type					
	Antibody screen					
	Diagnosis					
	Name of surgery					
	Duration of surgery					
	Injury Severity Score (ISS) ?					
	Abbreviated Injury Scale (AIS)?					
	Admission Glasgow Coma Scale (GCS) score					
	Did the patient have a pre-existing bleeding risk? =					
	Preadmission anticoagulation (= Active prescription					
	for an anticoagulant or anti-PLT agent at the time of					
	hospital admission)					
	Trauma: mechanism of injury: Blunt (Yes,					
	type:/No), Penetrating (Yes, type:/No)					





AUDIT MASSIVE HAEMORRHAGE TEMPLATE

Transfusion	Number of transfused RBC from emergency stocks						
	Number of transfused RBC from emergency stock						
	blood bank (uncrossmatched)						
	Number of transfused RBC (gekruist)						
	Number of transfused plasma						
	Number of transfused Platelet Concentrates						
	Number of returned RBC						
	Number of wasted RBC (non-conform storage, etc.)						
	Number of wasted plasma						
	Number of wasted Platelet Concentrates						
	Plasma: RBC transfusion ratio						
	Platelets : RBC transfusion ratio Time to first RBC transfusion						
	Time to first plasma transfusion						
	Location of transfusion during MHP: emergency						
	department, theatre, intensive care unit, delivery						
	ward						
	Use of blood warmer						
	> 10 units RBC transfused in 24h?						
	> 4 units RBC transfused in 4h?						

Conventional	Hb (g / dL) [Baseline, lowest, post-resuscitation]						
Coagulation	Platelet count (109 / L) [Baseline, lowest, post-resuscitation]						
Assays:							
haematologic	INR [Baseline, lowest, post-resuscitation]						
parameters	Prothrombin time (PT) [Baseline, lowest, post-resuscitation]						
	Activated Partial Thromboplasin Time (APTT) [Baseline, lowest, post-resuscitation]						
	Fibrinogen mg / dL [Baseline, lowest, post-resuscitation]						
	pH [Baseline, lowest, post-resuscitation]						
	Base deficit [Baseline, lowest, post-resuscitation]						
	Maintainance stage: hourly monitoring of haematologic parameters?						
ROTEM	ROTEM used to guide hemostatic resuscitation?						



AUDIT MASSIVE HAEMORRHAGE TEMPLATE

Coagualation testing versus transfusion

Patient had a Hb result within 24 hr of MHP activation

Patient had an INR result within 24 hr of MTP activation

Patient had a PLT count result within 24 hr of MTP activation

Patient had a fibrinogen level result within 24 hr of MTP activation

1 Patient with INR > 1.5 received at least 1 unit of FFP within an hour after the result

Patient with a PLT count < $50,000/\mu l$ of blood received 1 unit of PLT within an hour after the result

Patient with fibrinogen level < 200 mg/dl or 2 g/l of blood received 2 g of fibrinogen concentrate within an hour after the result

Patient with fibrinogen level < 150 mg/dl or 1.5 g/l of blood received 4 g of fibrinogen concentrate within an hour after the result

Patient had a ROTEM EXTEM assay result within 24 hr of MTP activation

Patient had a ROTEM FIBTEM assay result within 24 hr of MHP activation

Patient with ROTEM EXTEM A10 <45 mm and ROTEM FIBTEM A10 >13 mm received 1 unit of PLTs within an hour after the result

Patient with ROTEM EXTEM A10 < 45 mm and ROTEM FIBTEM A10 < 13 mm received 2g of fibrinogen concentrate within an hour after the result

Patient with CT > 80 s received at least 1 unit of FFP within an hour after the result

Cell salvage	Cell salvage volume reinfused (mL)			
Hemostatic agents	Anti-fibrinolytica: tranexaminezuur [Exacyl®] en aprotinine [Trasylol®] > administration of tranexamic acid within 24 hr of MTP activation			
	Stollingsfactoren: fibrinogeen [Riastap®], protrombinecomplex-concentraat [Cofact®, Confidex®], geactiveerde stollingsfactoren [Novoseven®], Factor VIII + factor von Willebrand [Haemate P®], Factor VIII [Advate®]			
	Antidota anticoagulantia: Idarucizumab [Praxbind®], Protaminesulfaat [Protamine Sulfaat Leo Pharma®]			
Vasopressine-analogen	Desmopressine [Minirin®]			
Calcium				
Crystalloids vs Colloids	Volume of intraoperative crystalloid (mL) Volume of intraoperative colloid (mL)			
Additional measures	What was implemented to stop bleeding (Surgical management, angiography, other)?			
Temperatuur	Temperatuur patiënt Wat gebruikt om hypothermie te vermijden?			



Sanderson 2020 - How well does your massive transfusion protocol perform? A scoping review of quality indicators Margolin 2023 - Massive Transfusion Protocol Adherence: Relationship to Trauma Patient Outcomes Shields 2019 - Obstetrical hemorrhage reporting and systems learning Broxton 2018 - Implementation of a Massive Transfusion Protocol: Evaluation of Its Use and Efficacy

References:

AUDIT MASSIVE HAEMORRHAGE TEMPLATE

Intensive care: length of stay (days) Hospital length of stay (days) Number of ventilator-days Peripartum hysterectomy (in case of obstetric haemorrhage) Hospital mortality Survival after the initial 24 hours after MHP activation Survival at discharge Survival at 30 days Administration Cessation of MHP: date and time Debrief documented within 72 hours? Time from MHP activation to arrival of first red blood cell from the blood bank Was the patient's identity known at the time of activation? Feedback about multidisciplinary collaboration and communication [orally and retrospectively collected by the Transfusion Practitioner]: Were the blood bank, emergency lab, perfusionist notified of massive blood loss? Were the blood bank, emergency lab notified of MHP	Outcome	Trauma-induced coagulopathy				
Number of ventilator-days Peripartum hysterectomy (in case of obstetric haemorrhage) Hospital mortality Survival after the initial 24 hours after MHP activation Survival at discharge Survival at 30 days Administration Cessation of MHP: date and time Debrief documented within 72 hours? Time from MHP activation to arrival of first red blood cell from the blood bank Was the patient's identity known at the time of activation? Communication Feedback about multidisciplinary collaboration and communication [orally and retrospectively collected by the Transfusion Practitioner]: Were the blood bank, emergency lab, perfusionist notified of massive blood loss? Were the blood bank, emergency lab notified of MHP		Intensive care: length of stay (days)				
Peripartum hysterectomy (in case of obstetric haemorrhage) Hospital mortality Survival after the initial 24 hours after MHP activation Survival at discharge Survival at 30 days Administration Cessation of MHP: date and time Debrief documented within 72 hours? Time from MHP activation to arrival of first red blood cell from the blood bank Was the patient's identity known at the time of activation? Communication Feedback about multidisciplinary collaboration and communication [orally and retrospectively collected by the Transfusion Practitioner]: Were the blood bank, emergency lab, perfusionist notified of massive blood loss? Were the blood bank, emergency lab notified of MHP		Hospital length of stay (days)				
haemorrhage) Hospital mortality Survival after the initial 24 hours after MHP activation Survival at discharge Survival at 30 days Administration Cessation of MHP: date and time Debrief documented within 72 hours? Time from MHP activation to arrival of first red blood cell from the blood bank Was the patient's identity known at the time of activation? Communication Feedback about multidisciplinary collaboration and communication [orally and retrospectively collected by the Transfusion Practitioner]: Were the blood bank, emergency lab, perfusionist notified of massive blood loss? Were the blood bank, emergency lab notified of MHP		Number of ventilator-days				
Hospital mortality Survival after the initial 24 hours after MHP activation Survival at discharge Survival at 30 days Administration Cessation of MHP: date and time Debrief documented within 72 hours? Time from MHP activation to arrival of first red blood cell from the blood bank Was the patient's identity known at the time of activation? Communication Feedback about multidisciplinary collaboration and communication [orally and retrospectively collected by the Transfusion Practitioner]: Were the blood bank, emergency lab, perfusionist notified of massive blood loss? Were the blood bank, emergency lab notified of MHP		Peripartum hysterectomy (in case of obstetric				
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Survival at discharge Survival at 30 days Administration Cessation of MHP: date and time Debrief documented within 72 hours? Time from MHP activation to arrival of first red blood cell from the blood bank Was the patient's identity known at the time of activation? Communication Feedback about multidisciplinary collaboration and communication [orally and retrospectively collected by the Transfusion Practitioner]: Were the blood bank, emergency lab, perfusionist notified of massive blood loss? Were the blood bank, emergency lab notified of MHP		Hospital mortality				
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the Transfusion Practitioner]: Were the blood bank, emergency lab, perfusionist notified of massive blood loss? Were the blood bank, emergency lab notified of MHP	Communication	Feedback about multidisciplinary collaboration and				
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Were the blood bank, emergency lab notified of MHP		Were the blood bank, emergency lab, perfusionist				
		notified of massive blood loss?				
		Were the blood bank, emergency lab notified of MHP				
CESSALIOH!		cessation?				



••• AUDIT MASSIVE HAEMORRHAGE – REPORT: RECOMMENDATIONS

Recommendations	Objectives	Improvement action (SMART)	Planning (who, when)	Barriers and challenges	Outcome	Follow-up
3. Advice of haematologist should be requested especially in case of lifethreatening coagulopathy	Optimal treatment of coagulopathy (final aim: reducing and managing the blood loss)	a) Adding some haematological aspects to the massive haemorrhage protocol + clearer instruction to contact haematologist. b) Including the consulting a haematologist as an indicator in annual overview of massive haemorrhages	a) A. De Becker: Q1 2024 b) J. Vanden Broeck: rest 2024	More experience in this topic needed for haematologists of our hospital	Advice haematologist should be asked <u>at</u> <u>least</u> for every activation of MHPs ≥ 4 hours or if >3 ROTEMs	A. De Becker, J. Vanden Broeck





••• CHALLENGES FLOWCHART MASSIVE HAEMORRHAGE PROTOCOL

(1) Indicatie Massale Bloedingsprotocol: Bij een actieve of recente bloeding met één van VOLDOET AAN INDICATIE MASSALE BLOEDINGSPROTOCOL(1) olgende voorwaarden: Universitair 1) Tekens van hemorragische shock Ziekenhuis 2) Bij wie transfusie van meer dan 4 bloedcomponenten verwacht wordt het eerstvolgende uur (2) Bij opstart MBP steeds aan Bloedbank Supervisie CC Spoedgevallendienst Activatie van MBP 2) Zorgcoördinator Spoedgevallendienst (3903) Naam, voornaam en dossiernummer patiënt 3) Bloedbank (6720) en Urgentielabo (5044) (2) Geslacht Dienst waar de patient zich bevindt Naam en DECT van Coördinerend Arts GELIJKTIJDIG GELIJKTIJDIG GELIJKTIJDIG **HEMOSTASE - STOP THE BLEEDING DETECTIE COAGULOPATHIE** Standaard cABCDE Benadering EXTERNE HEMOSTASE: Voorzie uitgebreide bloedafname met kruisproef 1) Lokale Druk en hoogstand en bloedgroepbepaling Voorzie 2x large Bore IV Access (2x 2,7ml Oranje, 4x 1,6ml Rood, 1x3 ml Groen)) Drukverband Overweeg onmiddellijke plaatsing Rapid STALEN AF TE NEMEN VOOR START Overhechten/stapler Infusion Catheter 4) Tourniquet --> Noteer de tijd! TRANSFUSIE Minimaliseer Cristalloïd / Colloïd gebruik *Bekkensling bij Bekkentrauma. Installeer Ranger en Rapid infuser - prime preferentieel met Bloed INTERNE HEMOSTASE: NEEM 1X EXTRA ROTEM TUBE AF Ifv bloedingsfocus activeer de specifieke (1x 3ml Groen - VOLLEDIG STAAL VULLEN discipline: TOT REFERENTIELIJN) Sta Permissieve Hypotensie toe - interventioneel radioloog VPK analyseert stalen onmiddellijk op ROTEM TOT CONTROLE BLOEDING (3) (2820 > 6048 > 5373 > 5100) · Vermijd Hypothermie - voorzie in externe toestel OK. Senior Chirurg (2686) warmtestraler / Bairhugger zo mogelijk Orthopedist (2628) Vermijd Acidose: overweeg verhogen Sa na of de patiënt gekend is met minuutvolume tot laag normale capnie Gastro-Enterologg (3305) 1) Aangeboren / verworven Coagulopathie Pneumoloog (17110) 2) Anticoagulatie / anti-agregantia inname KNO (3166) tiidstip laatste inname?) Gynaecoloog niet zwanger (6504) Gynaecoloog zwanger (6788) EERSTE TRANSFUSIESTRATEGIE Ifv mogelijk interventies buiten de = RATIO BASED Spoedgevallendienst, verwittig vroegtijdig BEHANDELING COAGULOPATHIE Anesthesie (3124/3123) Transfusieratio: 1U ECL / 1U FFP / 1U PLT DIEN TXA ASAP TOE ZO GEÏNDICEERD Bestel bloedproducten via order 'Masaal Bii ongekende Focus: (TRAUMA/POSTPARTUM) Bloedverlies' per pakket in PRIMUZ. Bolus TXA: 1g / 10 min L) VOORZIE IN POCUS E-FAST / RUSH gevolgd door Spp: 1g / 8 uur Keuze tussen - 4U ECL / 4U FFP / 0U PLT - 4U ECL / 4U FFP / 1U PLT 2) OVERWEEG CT-GRAFISCHE Voorzie ASAP Reversal van Anti-coagulatie BEELDVORMING TIJDENS RESUSCITATIE - #U ECL / #U FFP / #U PLT (ifv ROTEM) (cfr MBP Zenya - specifieke protocol) Plaatjestransfusie van een pool plaatjes pas na OVERWEEG ONMIDDELLIJKE TRANSFER Overweeg stollingsfactoren / vitamine K bij NAAR OK VOOR DAMAGE CONTROL 8U ECL en 8U FFP in totaal verworven coagulopathie SURGERY / INTERVENTIONELE RADIOLOGIE TIJDENS RESUSCITATIE Houdt Calcium en Fibrinogeen boven de transfusiedoelen (4) Startpakket 4U ECL en 4U FFP terug te vinden Vuistregel 1g CaCl in NaCl 0.9% 50cc over 10 in noodvoorraad Spoedgevallendienst min na elke - 4U ECL O ongekruist in koelkast VANAS - 2U ECL of FFP - 4U FFP in Diepvries naast Plasmatherm - na elke 1 pool PLT

ROTEM GEKEND?

Controleer geiöniseerd Calcium regelmatig,

ninstens na ieder pakket.



Logistieke hulp of nachtvervoer haalt

de Bloedbank

ondertussen 2e pakket Bloedproducten op bij

⇔ easy and quick

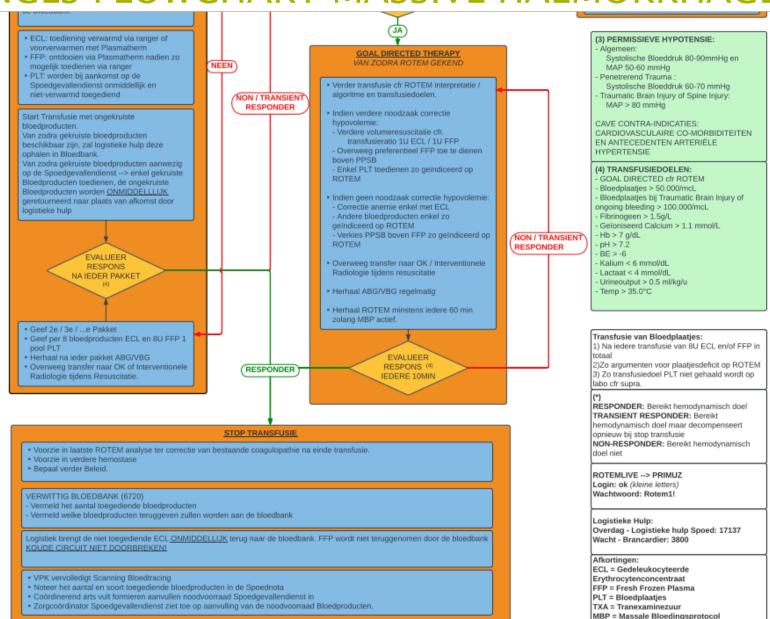
to understand in

emergencies



••• CHALLENGES FLOWCHART MASSIVE HAEMORRHAGE PROTOCOL

⇔ easy and quick to understand in emergencies

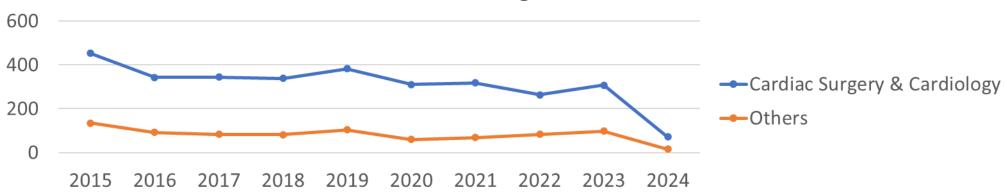


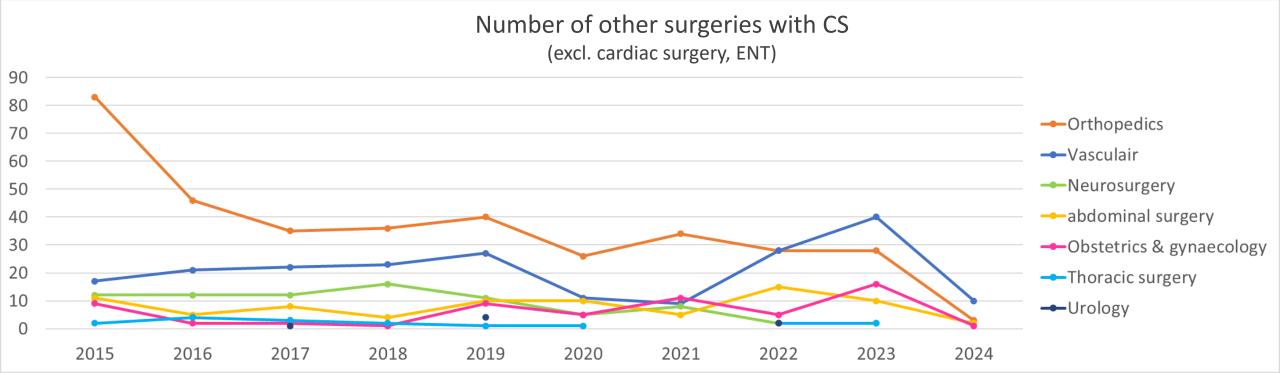




••• USE OF CELL SALVAGE



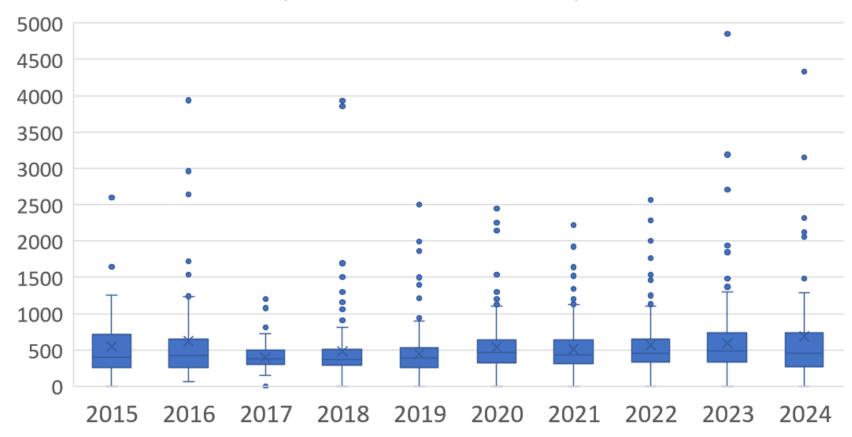




••• USE OF CELL SALVAGE

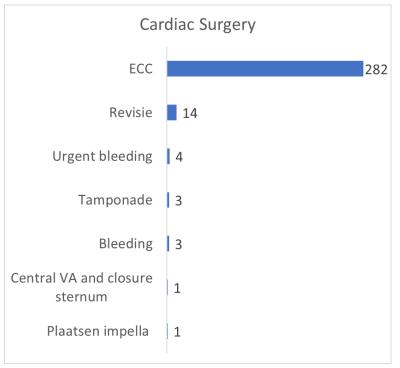
Cardiac surgery: range of returned volume

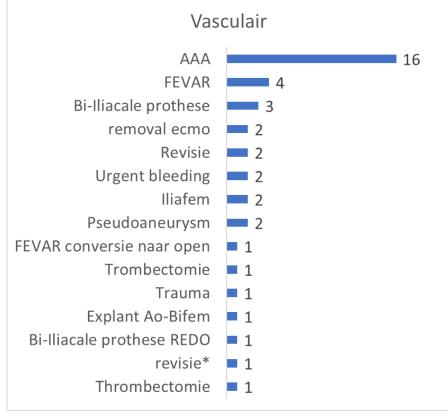
(extra outlier in 2022: 8.000 mL)

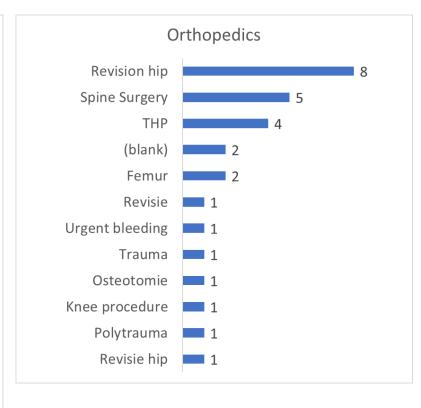


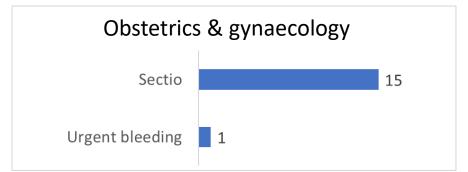


INDICATION CELL SALVAGE PER MEDICAL DISCIPLINE (2023)













••• INCIDENT REPORTING ON CELL SALVAGE?

TERM	DEFINITION	WHAT TO REPORT
Cell Salvage	Events and reactions in relation to the use of intraoperative and postoperative cell salvage.	 This category currently includes: Adverse events due to operator error, where the event has the potential to impact on patient care Adverse events due to machine or disposable failure where the event has the potential to impact on patient care Adverse events related to the availability of trained staff which impact on the patient Adverse clinical events during the cell salvage process, such as hypotensive events Pathological reactions to <i>reinfused</i> blood

Recommendations:

Review suitability of cell salvage documentation (paper or electronic) and its appropriate use. Ensure
the record of cell salvage is accessible and complete, particularly in relation to communicating
pertinent details at handover

Action: Cell salvage leads, theatre teams, hospital transfusion teams

• Establish clear responsibilities and lines of reporting for cell salvage incidents. Review pathways and structure for governance and communicate these processes to all stakeholders

Action: Cell salvage leads, theatre leads, HTC, clinical governance leads







DATA EXPORT: PBM (IN OBSTETRICS) + RH D PROPHYLAXIS

IV iron, fibrinogen concentrate, tranexamic acid, anti-D immunoglobulins

- Record number, date and time of administration, name/number of prescribing clinician
- Date of birth, weight (before pregnancy), BMI, etnicity
- Gravidity, parity, expected delivery date, delivery date and time, multiple pregnancy, delivery: vaginal (normal/instrumental), caesarean section (elective/emergency)
- Oral supplements in pregnancy: iron, multivitamins, others, unknown
- Antepartum/peripartum bleeding, amniocentesis, chorionic villus sampling, external cephalic version, abdominal trauma, extra-uterine pregnancy, miscarriage/abortus/still birth, pregnancy-related hypertensive disease: pre-eclampsia
- Lab results (+ date)
 - ABO & Rhesus D blood type
 - Screening and identification anti-D antibodies & titer anti-D antibodies if positive
 - Kleihauwer Betke test
 - Flow cytometry
- Record number baby + ABO blood type & Rhesus D, birth weight and length

(to compare with all O Rh negative pregnant/delivered women who didn't receive any of these medications?)



Order for Desmet, Arthur (XXXXXXX)

Prescriber				Destin	ation				
Prescribing ward:	Prescribing ward: A210 Neonatology		Transfusing w	rard: A210	Neonatology				
Prescriber :	Dr. CO	OOLS, Fili	р	Tel. :	6763		s after this order ate, crossm.	22/07/2022 13:00	
					Transfusi	☐ Urgent,	uncrossmatched		
	Pat	tient					Blood com	ponent(s)	
Gestational age:	34	weeks	2	days		RBC	Number:	Specific requirements:	
Postnatal age:	0	week	21	days		⇒ Indication ———————————————————————————————————	Postnatal blood	naturity (top-up transfusion) loss (bv. intraventricular, pulmeto catheter insertion)	nonary, gastro-intestinal
Extra information:	(a)·					⇒ Indication	Perinatal blood I umbilical cord ru Surgery	oss (foetomaternal transfusion pture or abruptio placentae)	on, placenta/vasa praevia,
☐ Other:	(6).					Platelets	Exchange transfu Other	usion	_
						⇒ Indication Reconstit. whole bloo	0		<u> </u>

Non-complete transfusion record in the OR: Mock-up pop-up for anaesthesiologists

Were these blood components transfused during the surgery?							
Unit number	Transfused	Timing					
XXX		10:15					
YYY							
ZZZ							

Save



Thank you & let's continue the good work as TPs





