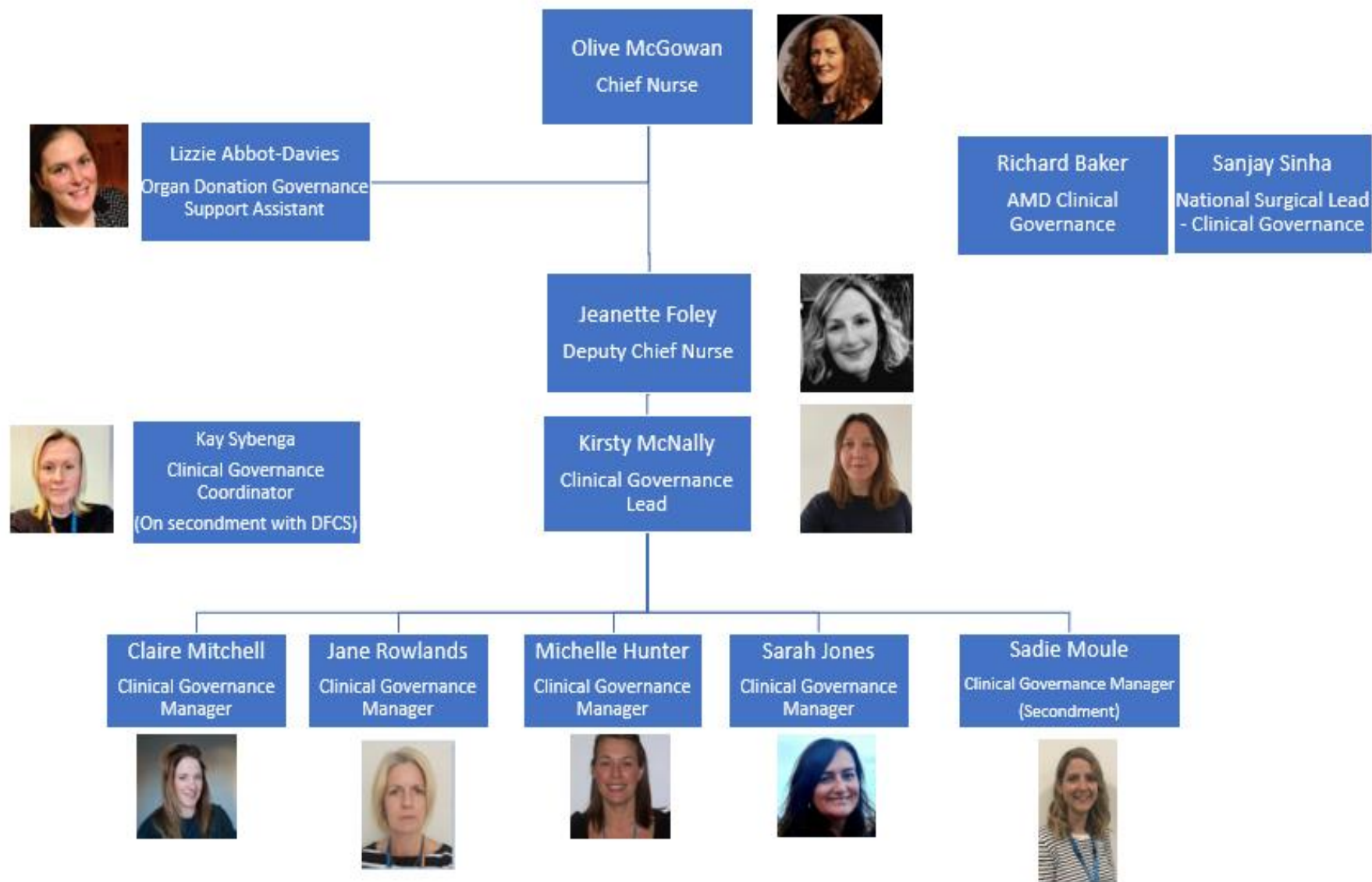


Patient Safety in Donation and Transplantation

ODT Clinical Governance



Patient Safety is what we do...

Learning Sharing Strengthening



Human Factors in
Incident Investigation



NHS
Resolution

Duty of candour

An illustration of two people, a man and a woman, sitting at a desk with a computer. The man is on the left, wearing a blue shirt, and the woman is on the right, wearing a blue dress. They are both looking at the computer screen. The text "Duty of candour" is written in a blue, cursive font below the illustration. In the top right corner, the NHS Resolution logo is visible.

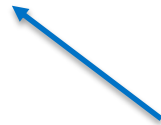
What is an ODT incident?

Any event in the organ donation and/or transplantation process which can or does affect the donor, recipient, safety or the quality of the organs for transplantation

May have national or wider learning

NHSBT has an assisted function with the HTA - Legal requirement to report under HTA regulations

May relate to organs being sent/received from overseas



NHS

Blood and Transplant



Serious Adverse Event (SAE) -
any event that occurs that impacts
or has the potential to impact on a
patient

Serious Adverse Reaction (SAR)
- An unintended reaction that
impacts on the patient

Never Event- Preventable
incidents and events that should
never occur, dictated
by NHSE and NHSI and
reportable.

Examples of reportable SAEARs

Damage to an organ during retrieval.

An organ was inadvertently damaged during retrieval. The damage associated with retrieval resulted in an otherwise transplantable organ rendered unsuitable for transplantation.



SAE

Transmission of a communicable disease.

A cytomegalovirus (CMV) test result was reported incorrectly as negative when the actual test result was positive. At the time of implantation in a CMV negative recipient the transplant surgeon was unaware that the donor test was CMV positive.



SAE

The recipient will be subject to monitoring. **If CMV related disease develops** (transmission of disease), this will be reported as a serious adverse reaction.



SAR

An unnecessary procedure performed on organ recipient.

An organ recipient was anaesthetised in preparation for an organ transplant. On inspection of the organ, the surgeon following a risk and benefit analysis found the organ was unsuitable for transplantation and the procedure was aborted.



SAR

Donor malignancy discovered during post mortem.

The organ donor's medical history did not indicate potential malignancy and lesions were not evident at retrieval.



SAE

The recipient will be subject to monitoring. **If a related malignancy develops** (transmission of disease), this will be reported as a serious adverse reaction.



SAR

Surgical removal of transplanted organ.

An organ damaged during retrieval was repaired and implanted. Following implantation, the recipient developed complications associated with the transplant and the organ had to be removed.



SAR

Extended cold ischaemic time (CIT).

An unlabelled organ box was not accepted for transport resulting in extended CIT. Following delay, the organ was unsuitable for transplantation.



SAE

Donor had previous history of malignancy which was known at the time

<https://www.odt.nhs.uk/odt-structures-and-standards/governance-and-quality/tell-us-about-an-incident/>



Blood and Transplant

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Incident Reporting

Urgent incidents

Call ODT Hub Operations on 0117 975 7580 if the incident is urgent and may affect the quality and safety of an organ for transplantation or the treatment of recipients or potential recipients.

This call should be followed by completing this [online form](#)

Tell us about an incident

Tell us about an incident by completing this [online form](#)

Positive transport fluid results

Tell us about positive transport fluid results by downloading and completing the [Rapid Alert – Positive transport fluid results form](#) and emailing it to odthub.operations@nhsbt.nhs.uk

In this section

[Shared Learning](#)

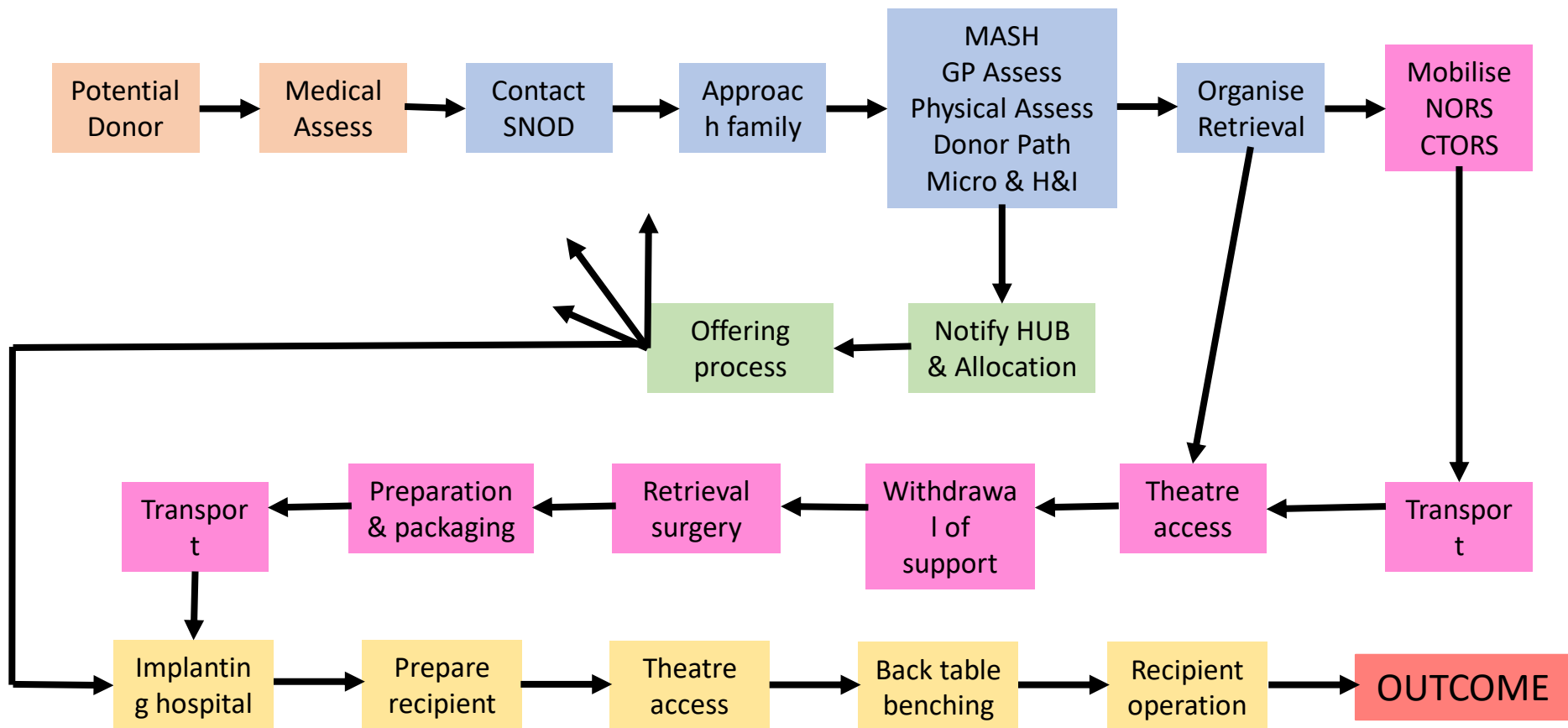
[Incident Reporting](#)

[Learning from excellence](#)

Healthcare is a complex socio-technical system...multiply that by 10 for us!

Highly variable, uncertain and dynamic; **Ultra-adaptive**

Patient safety 'emerges' from interactions and **not from a single component, such as the actions of people**



Thinking about patient safety....

- Systems thinking – “human factors” **
- Psychological safety in teams **
- Restorative just culture
- Shared learning *
- Learning from Excellence – what works ?
- Incident investigation – how do you involve people? **



Systems thinking



- Alarm overload
- Usability
- Accessibility
- Availability
- confusing tool, equipment or work procedure
- inadequately designed interfaces

- Shift work/work schedules
- Supervision
- Leadership
- Organisational culture
- Staffing levels

Individual characteristics:

- Knowledge, competence
- cognitive factors (frustration, stress)

Collective characteristics

- Team cohesion

- Excessive physical or cognitive demands
- High or low workloads
- Complexity
- Monotonous or repetitive

Characteristics of physical environment such as:

- smell, noise, heat, cold, cramped conditions, lighting, ventilation, ease of access
- work space design

Targets, regulatory or policy obligations, accreditation standards, global events outside organisation control

Tools & Technology

- Some LIMS systems will not allow blood product issue without a BG entered
- Human can override the system to enter O
- Can LIMS be overridden without entering a flag/note on system? (refer to Never Event FFP for benefit of comments only). What does LIMS say/do if overridden?
- Result visible, comments not always visible, dependant on system and user
- LIMS systems significantly different across UK – John Radcliffe really advanced, others from 1970's
- LIMS doesn't have ability to flag if patient an OD if known
- Analyser may 'release' a result if a previous result has been entered (even if reverse and forward groups differ)
- If reverse and forward groups differ when analysed, and no prior group confirmed, a human is required to interpret
- SNODs reviewing hospital systems that unfamiliar with
- DonorPath only asks 'any transfusions in the last week' – why? Rationale of timescale?
- Haemodilution does not mention blood group
- What is 'mass transfusion' – calculation? Impact of incorrect weight

Internal environment

- RCI support likely to be remote working – viewing results via scanned copies
- Conversations often in noisy environments with conflicting 'mental models'

External environment

- Lack of national guidelines in blood group confirmation following mass transfusion
- SHOT reportable events in relation to incompatible ABO FFP transfusion – info not shared with OTDT
- No mention of organ donation or transplantation in any haematology guidelines

Person

SNOD

- Assumption that there are 'special' tests in RCI labs to confirm BG in mass transfusion (not the case)
- Haemodilution is felt to be for microbiology/tissues only not BG
- Time pressured to get result. May inadvertently pressurise BMS to confirm a result
- Weight of organ donors often 'not taken seriously'
- SNOD assumes BMS understands why need a BG and impact on organ donation

BMS

- Stressed due to competing demands if on call
- Likely never involved in organ donation – being told patient is an OD irrelevant to decision making
- Mind set is focused around transfusion and "If in doubt state O" – O is safest
- Risk adverse
- Mental model impacts on decision – think fast and state O
- Experience may impact on whether discrepancies escalated
- There is a lack of confidence to interpret BG when mass transfusions occurred
- BMS may assume SNOD just needs 'a BG' and not understand the implications

Tasks

- Escalation by the of who and when to escalate will vary
- Handover from BMS to senior BMS may 'lose' key info around a patient being an organ donor
- Huge variation from SNODs around sending of blood to confirm blood group
 - Repeat on referral
 - Repeat if 'old' (how old?)
 - Sending samples to RCI as 'special' test to confirm BG in mass transfusion – not the case
 - Concerns around accuracy of when 'historic' samples sent – so not able to correlate with transfusions
- Who and when to SNODs escalate to?
Not all hospitals send 2 blood samples to confirm blood group – 'safetrack' use means only 1 sample sent
O often entered 'as safest for the patient' – however this is not always the case (FFP Never Event)
Discussions between BMS and SNOD could be 'cross wired'

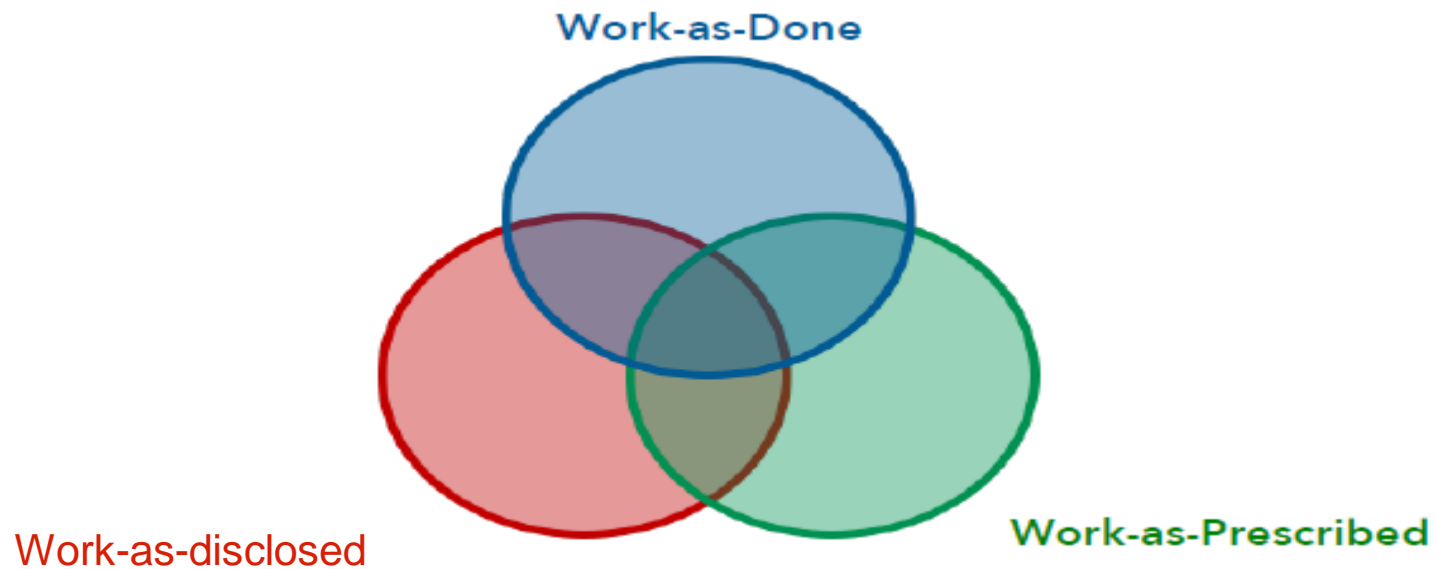
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Incident Investigation

Important to involve those directly involved in the incident – you are often the people who do that, including donor families

Look at ‘work as prescribed’ versus ‘work as done’

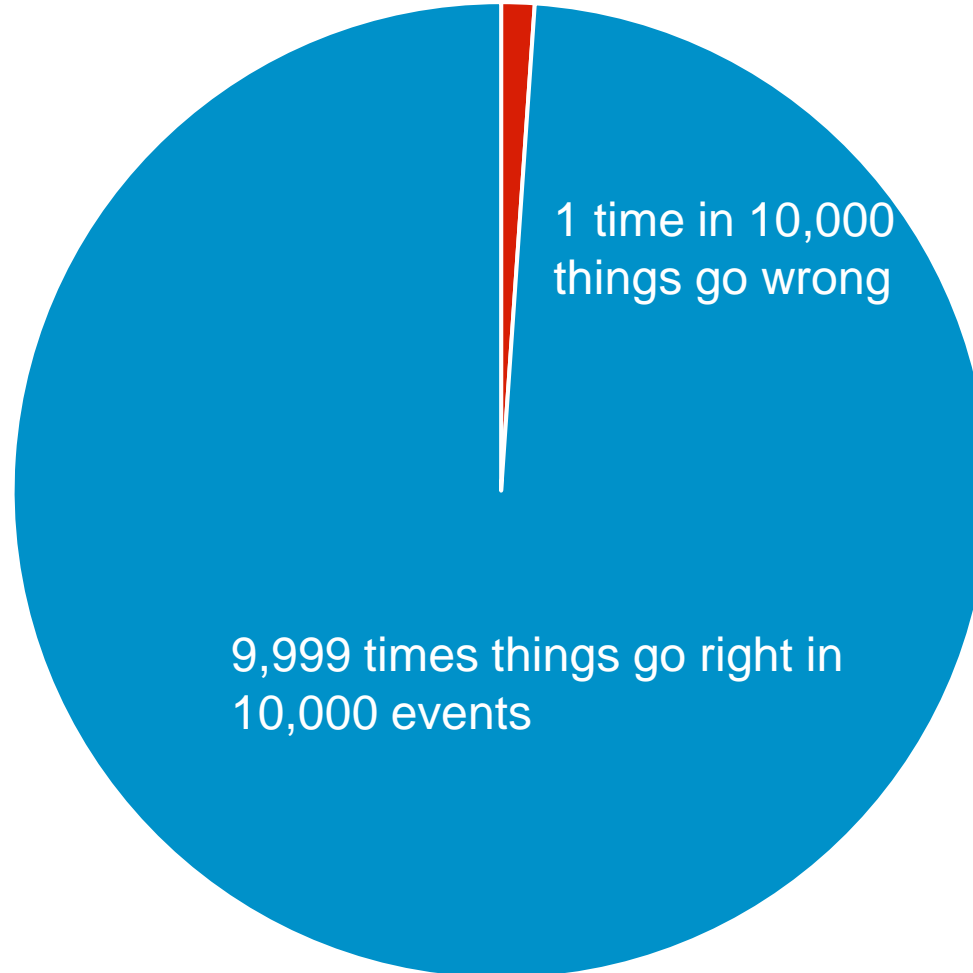


Psychological Safety in teams

<https://www.youtube.com/watch?v=eP6guvRt0U0>



Safety I and Safety II





In this section

Shared learning

Incident Reporting

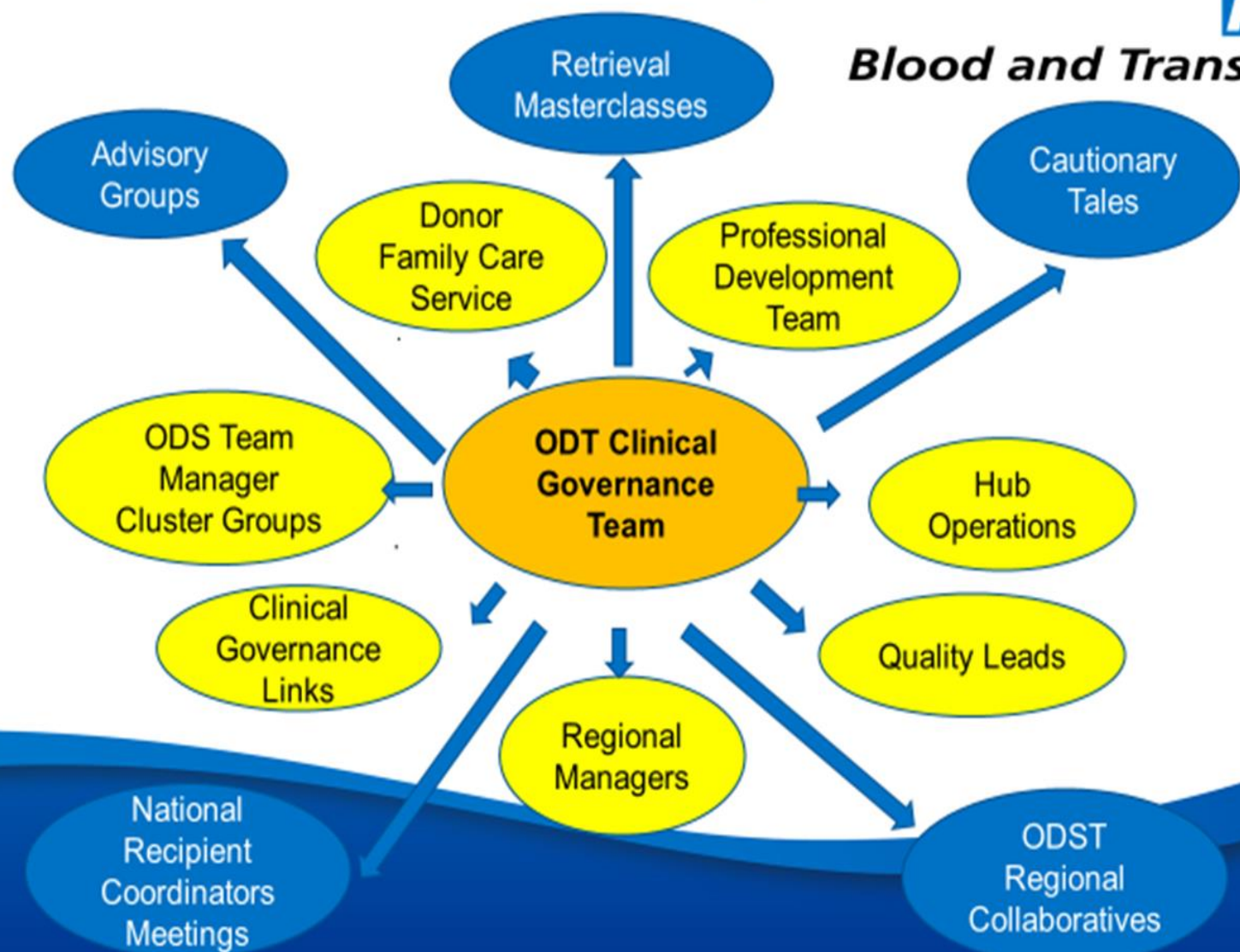
Learning from excellence

Complaints and compliments

Shared Learning

NHS

Blood and Transplant





NHS Patient Safety Syllabus

Safer culture, safer systems, safer patients



This programme is in partnership with...



<https://www.e-lfh.org.uk/programmes/patient-safety-syllabus-training/>



What questions do you have?

