



Resolution

CLYDE&CO

Kennedys

South Regional Safety and Learning Networking Forum

23 September 2025

12:00 – 14:00



@NHSResolution

Advise / Resolve / Learn

Housekeeping

- This event is being recorded.
- Please use the chat box to submit any questions.
- Feel free to share your innovations in the chat.
- An online feedback link will be shared towards the end of the meeting and emailed after the webinar.



Agenda



Resolution

Time	Item	Speaker
12:00	Introduction to the forum (5 mins)	Chair, Ingrid Henderson, Associate Safety & Learning Lead – London & South – NHS Resolution (NHSR)
12:05	Learning from claims data & consent (10 mins)	Caroline Latham-Parker, Safety & Learning Lead – South - NHSR
12:15	Consent & the law, how to avoid the pitfalls in clinical practice (20 mins)	- Rob Wilson, Partner - Clyde & Co LLP - Jennifer Harris, Legal Director - Capsticks LLP
12:35	Consent in general & colorectal surgery (15 mins)	Mark Coleman, General & Colorectal Surgeon - University Hospitals Plymouth NHS Trust
12:50	Responding to consent concerns: An advisory perspective (15mins)	Rinke Schram, Lead Assessment and Remediation Adviser, NHSR
13:05	Q&A (20 mins)	Ed Glasgow, Partner – Kennedys
13:25	Member led breakout groups facilitated by NHSR (30 mins)	- Caroline Latham-Parker - Ingrid Henderson - Kelly Birleson
13:55	Summary and Close (5 mins)	Caroline Latham-Parker

Failed to Warn-Informed Consent CNST claims 2020/21 to 2024/25

National & regional overview
(South East & South West)

Safety & Learning Team

Presentation Overview

1. Introduction
2. National - Volume and value of Failed to Warn-Informed Consent CNST claims per year.
3. National & Regional – Status of Failed to Warn-Informed Consent CNST claims 2020/21 to 2024/25.
4. National & Regional – Top specialities for Failed to Warn-Informed Consent CNST claims volume
5. National & Regional – Top 5 Cause codes for Failed to Warn-Informed Consent CNST claims volume.

Learning Objectives: To provide an overview of the national and regional themes seen in Failed to Warn-Informed Consent claims brought under the CNST scheme.

- How does these themes compare with what you see in your local claims via the Trust scorecard?
- Do they align with your current quality metrics, e.g. complaints and patient safety events data?
- How can this help you formulate priorities for future patient safety improvement?

Introduction

- Supporting decision making (consent) is a vital part of patient care.
- Consent claims often centre on the information, or insufficient information, provided to patients to make informed decisions about their healthcare.
- Supported decision making ensures that risks are not omitted (including the risk that treatment may not achieve its intended aims).
- Between April 2020 and March 2025, 2185, claims were reported with a cause code of 'Fail To Warn-Informed Consent'. The estimated value of these claims is £807.7 Million.

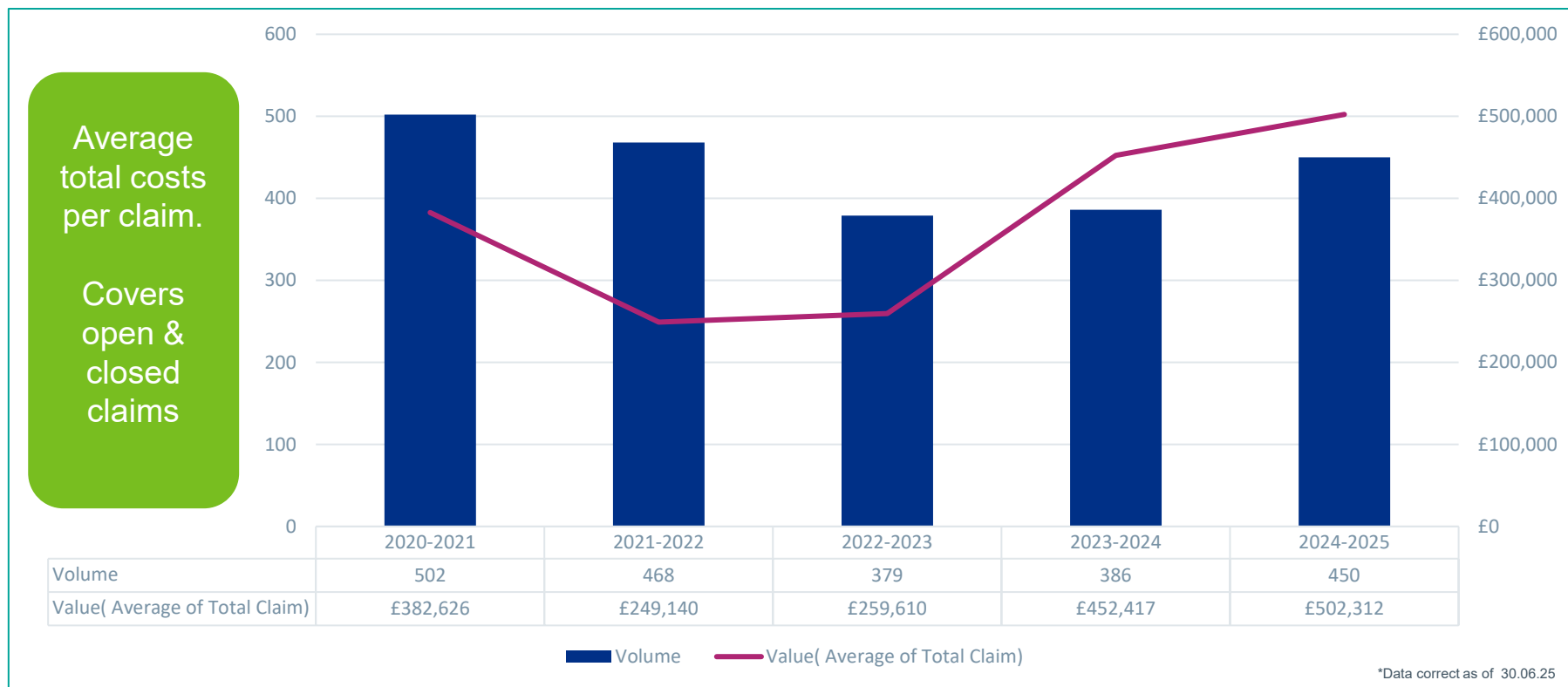


<https://resolution.nhs.uk/resources/the-benefits-of-supported-decision-making-consent/>
<https://resolution.nhs.uk/resource-foi-module/consent/>

National - Failed to Warn-Informed Consent CNST claims per year 2020/21 to 2024/25 via claims notification date



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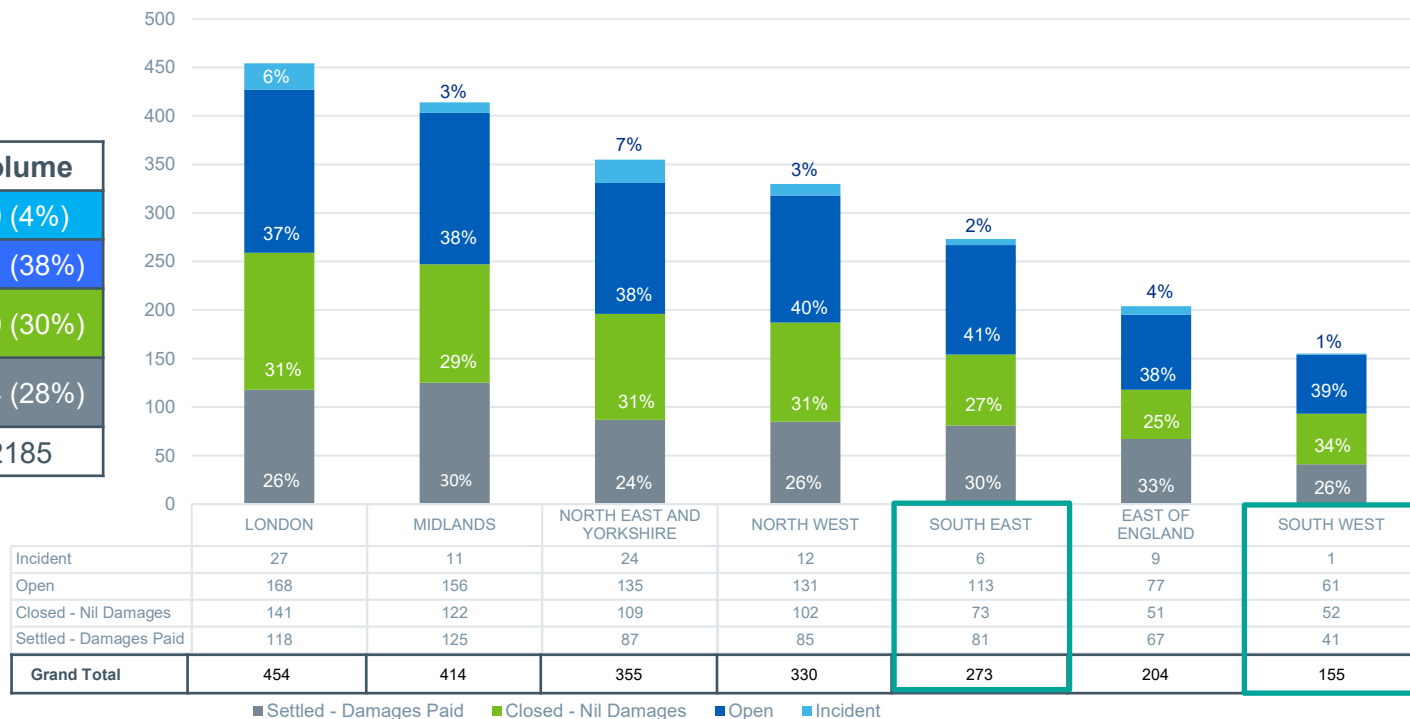


Status of National & Regional Failed to Warn-Informed Consent CNST claims 2020/21 to 2024/25 via claims notification date



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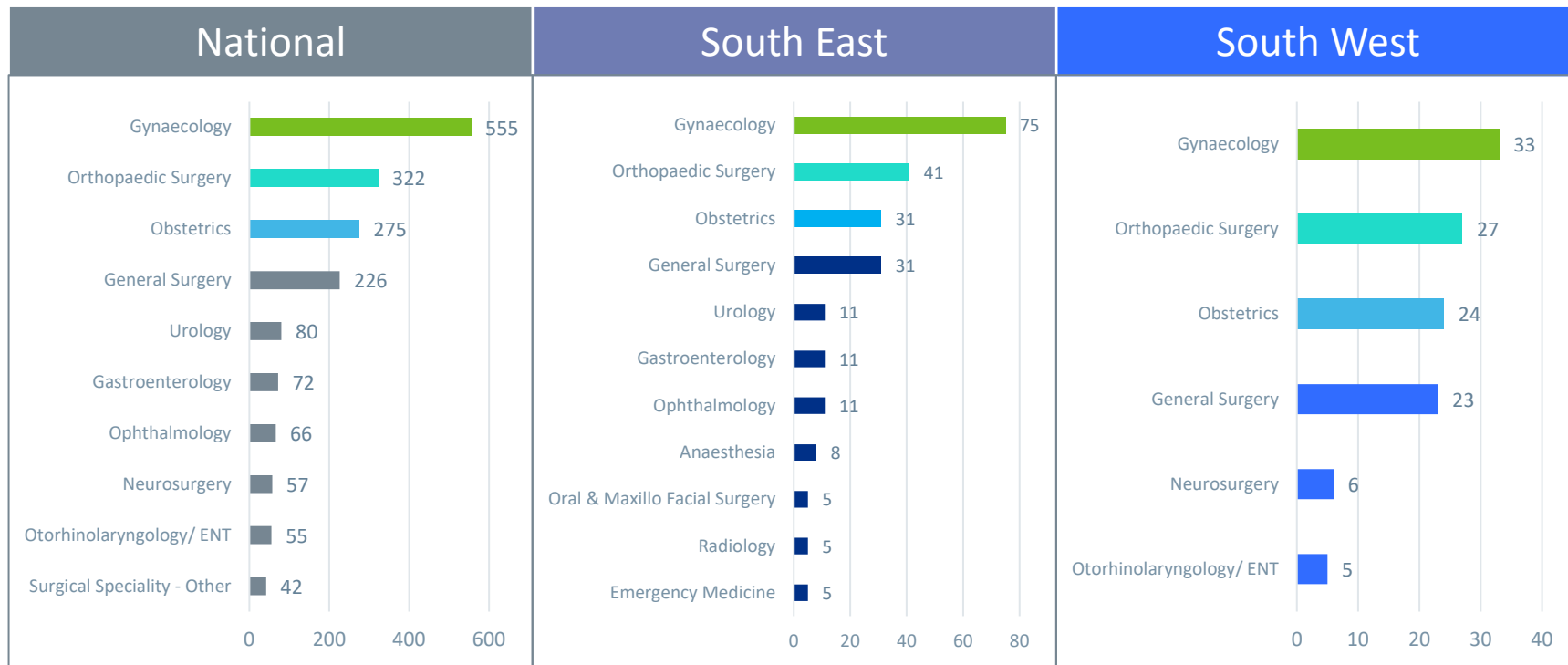
National	Volume
Incident	90 (4%)
Open	841 (38%)
Closed - Nil Damages	650 (30%)
Settled - Damages Paid	604 (28%)
Grand Total	2185



* Data correct as of 30.06.25

Failed to Warn-Informed Consent CNST claims by top Specialities & **NHS** volume 2020/21 to 2024/25 via claims notification date

Resolution



Volume of Failed to Warn-Informed Consent CNST claims by injury code 2020/21 to 2024/25 via claims notification date



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Successful risk navigation



Consent and the law – how to avoid the pitfalls in clinical practice

Rob Wilson, Clyde & Co

Jennifer Harris, Capsticks

Liability

Breach of Duty – not “Bolam” in consent cases

- Failure to obtain informed consent. Not adequately explaining alternative options/risk etc.

Causation

- Had the Claimant been appropriately consented he/she would not have proceeded with the treatment

Montgomery/McCulloch

- Doctors must provide information about all material risks which a reasonable person in the patient's position would attach significance.
- Include any reasonable alternatives – as per “Bolam” – supported by a reasonable body of medical opinion.
- Doctor to make judgement about how best to explain to individual patient: a “material” risk assessment is sensitive to the patient's characteristics.
- GMC guidance



NMC
Nursing &
Midwifery
Council

The Code

Professional standards of practice and behaviour for nurses, midwives and nursing associates



CONSENT:
SUPPORTED
DECISION-MAKING



Royal College
of Surgeons

CONSENT: SUPPORTED DECISION-MAKING

A Guide to Good Practice

Guidance on professional standards and ethics for doctors

Decision making and consent

Working with doctors Working for patients

General
Medical
Council

G I R F T

GETTING IT RIGHT FIRST TIME



Royal College of
General Practitioners

MRCGP RCA guidance and proforma consent form

Patient consent for recording of telephone and video consultations in general practice settings

Updated 18 June 2020

BMA

Consent and refusal by adults with decision- making capacity

A toolkit for doctors



COMMUNICATION

Informed Consent

- **Who** is responsible for getting consent?
 - Responsibility of individual performing procedure.
 - Can delegate but only if you are satisfied they are suitably qualified.

- **How** do you have the discussion?
 - Face to Face / Over the Phone / Video Link
 - Avoid Jargon, bombarding with technical information
 - Consider language / cultural barriers

Mordel v Royal Berkshire NHS Foundation Trust [2019]



COMMUNICATION

Informed Consent

■ When should you get consent?

- Depends on the nature of the procedure.
- On the day / Days or Weeks in advance / Both
- Is it invasive? High Risk? Minor or assessment only?
- Do you need to allow time to consider the options?

Thefaut v Johnston [2017]

■ Without Pressure or Coercion

- Family / Religious leaders / clinicians

**It is a process
Not a single point in time**



CONTEXT IS KING

Informed Consent

What does this look like in real life?

Diamond v Royal Devon & Exeter NHSFT [2017/2019]

Failure to consider the impact of options on the future

Plant v El-Amir [2020]

Even if successful will you improve the situation sufficiently to warrant taking the risk?

Thefault v Johnston [2017]

Inaccurate advice on success rates and risks.

Remember:

- **Consider the future.**
- **Will you achieve the patient's aims?**
- **Get the stats right!**



CONTEXT IS KING

Informed Consent

What does this look like in real life?

A v East Kent Hospitals University NHS Foundation Trust [2015]

“a small but well-established risk of a serious adverse outcome” is significant for the purposes of consent - GMC Guidance.

Bayley v George Eliot Hospital NHS Trust [2017]

Duty to notify of all “reasonable alternatives”.

Has to be known not experimental.

Mills v Oxford University Hospitals NHS Trust [2019]

Need to highlight if a procedure is “new” and address the “standard” option too.

Remember:

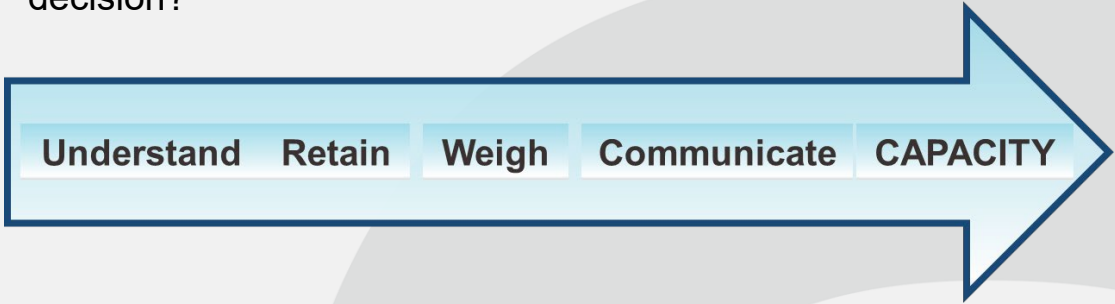
- **Real not theoretical risk**
- **Reasonable alternatives**
- **Highlight innovations**

Informed Consent

CAPACITY – *Mental Capacity Act 2005*

2 Stage Test

1. Do they have an impairment of their mind or brain?
(can be an illness or external factor such as alcohol or drugs)
2. Does the impairment mean they cannot make a specific decision?



Understand Retain Weigh Communicate CAPACITY

*Remember capacity can fluctuate with time
& may not apply to all types of decisions*

East Lancashire Hospitals NHS Trust v GH [2021]

Informed Consent

Mental Capacity Act 2005 - 5 Principles

1. A presumption of capacity

Every adult (16 yrs plus) has the right to make their own decisions and must be assumed to have capacity unless proven otherwise.

2. Individual supported to make their own decision

Do all you can to help the person make a decision.

3. Unwise decision

Unwise does not equal unable to make a decision.

Which includes refusal of life prolonging care.

4. Best Interests

If they lack capacity you must act in their best interests.

5. Least restrictive option



Kings College Hospital NHS Foundation Trust vs C [2015]

Re MB (Medical Treatment) [1997]

St George's Healthcare NHS Trust v S

Documentation

- 3 legged stool:
 - Information booklets
 - Patient-centred dialogue
 - Consent forms
- 3 exceptions:
 - Emergency
 - Lack of capacity
 - Impractical/impossible (voluntarily waived)

Thank you.

Rob Wilson, Clyde & Co
Jennifer Harris, Capsticks

Surgical Consent

Mark Coleman

- Consultant Surgeon University Hospitals Plymouth (UHP) NHS Trust (25 years)
- Chair of the National Joint Advisory Group for GI Endoscopy 2020-2023 (thejag.org.uk)
- Executive Council of the Association of Coloproctology GB&I 2011-2016 (acpgbi.org.uk)
- Service Line Director Colorectal Unit and Theatres UHP
- Co-Director Lapco International Consultancy

mcoleman1@nhs.net

+44 7789 873190



Disclosures

Introduction

- Surgical consent is a vital part of patient care
- Ensures patients are informed and voluntarily agree
- Protects patient rights and legal interests

“Decisions are more important than incisions”

“a fool with a tool is still a fool”

“Patients want to know that you care before they care what you know”



Is there a problem?

Understanding of Consent	56%–76%	Procedure-specific forms publishing.rcseng
Time Taken for Consent	~50% <10min, often rushed	RCS/RCSEng guidance publishing.rcseng
Errors in Consent Forms	>50% (paper forms)	Concentric Health/BJs study concentric



Definition of Surgical Consent



A legal and ethical process



Permission given by a patient
after understanding the
procedure, risks, and alternatives



Voluntary acceptance without
coercion

Importance of Surgical Consent



RESPECTS PATIENT
AUTONOMY



REDUCES LEGAL RISKS
FOR HEALTHCARE
PROVIDERS



ENHANCES PATIENT
TRUST AND
SATISFACTION



STATUTORY
REQUIREMENT IN
HEALTHCARE

Types of Consent



Expressed Consent: Oral or written



Implied Consent: Actions indicating agreement (e.g., extending arm for injections)

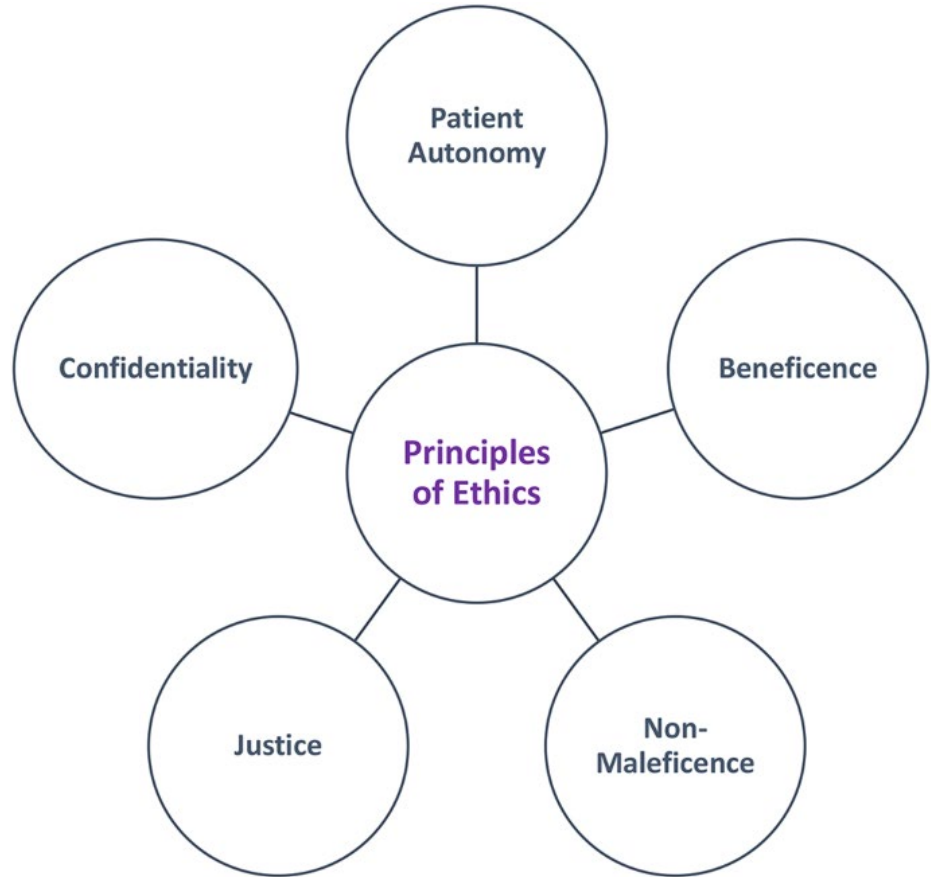


Informed Consent: Full disclosure of information

Components of Informed Consent

- Explanation of the procedure
 - Risks and benefits
 - Alternatives available
 - Questions and clarifications
 - Voluntary decision
- **BRAN?:**
 - **B**enefits
 - **R**isks
 - **A**lternatives
 - **N**othing
 - **?Q**uestions

Legal and Ethical Principles



When is Consent Not Necessary?

- Emergency situations when patient is unconscious or unable to consent
- Implied consent for life-saving procedures
- Minors with parental or guardian consent

[< Back to episode](#)

Capacity assessment

i You must provide a reason for conducting an assessment, details of support being provided, and answer 'yes' or 'no' to all sections.

Reason for assessment

What is the reason for conducting this capacity assessment?

Where an adult's capacity to consent to treatment is being assessed, document the nature of the impairment or disturbance and how it is presenting itself.

Please fill in this field.

Support being provided

What support is being provided to support the patient to make the decision for themselves?

Before concluding that a patient lacks capacity all practicable help must be provided to support the patient to make the decision for themselves. If it is impractical to provide support, explain why.

Guidance

Support may come in the form of assistance with communication or other practical help. Communication assistance may include use of simple language and repetition, communication devices, and help from a family member. Practical help may include 'Easy Read' information or drawings, and assessment in a place where they are at ease. Examples of attempts made to reverse any temporary causes of incapacity may include treating hypoxia, pain and infection, or allowing any sedative medication to clear.

Documenting Consent

- Written consent form signed by patient
- Details of the procedure, risks, and patient understanding
- Witness signatures if required

Patient agreement to investigation or treatment CONSENT FORM 1

Patient details (or pre-printed label)

Patient's surname/family name	_____
Patient's first names	_____
Date of birth	_____
NHS number (or other identifier)	_____
☐ Male	☐ Female

Responsible health professional _____

Job title _____

Special requirements
(eg other language/other communication method) _____

Name of proposed procedure or course of treatment

(include brief explanation if medical term not clear) _____

Statement of health professional (to be filled in by health professional with appropriate knowledge of proposed procedure, as specified in consent policy)

I have explained the procedure to the patient. In particular, I have explained:

The intended benefits _____

Serious or frequently occurring risks _____

Any extra procedures which may become necessary during the procedure

☐ blood transfusion ☐ other procedure (please specify) _____

I have also discussed what the procedure is likely to involve, the benefits and risks of any available alternative treatments (including no treatment) and any particular concerns of this patient.

☐ The following leaflet/tape has been provided: _____

This procedure will involve:

☐ general and/or regional anaesthesia ☐ local anaesthesia ☐ sedation

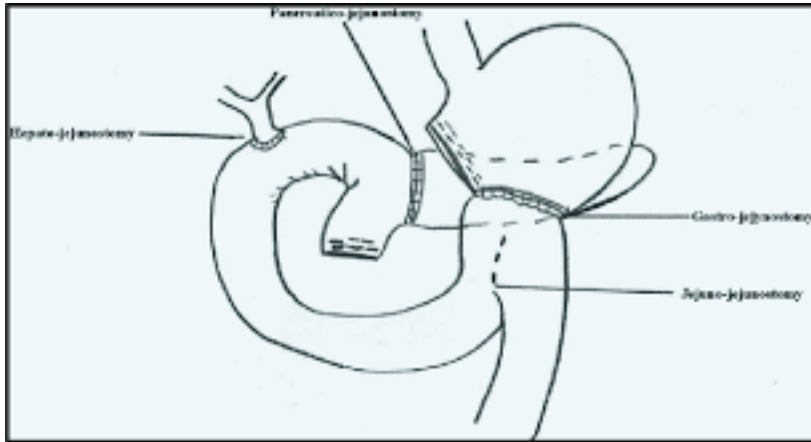
Signed _____ Date _____

Challenges & Common Issues

- Language barriers
- Cognitive impairments
- Medical jargon
- Coercion or undue influence



Improving Consent Process



- Use of plain language
- Visual aids and diagrams
- Confirm understanding
- Use interpreter services if needed

Conclusion

Consent is a
cornerstone of
ethical surgical
care

Must be
voluntary,
informed, and
documented

Continuous
communication
improves
outcomes



THANK YOU

Questions?



Case study: Lessons on consent and communication in clinical examinations

Addressing consent-related concerns about doctors: An advisory perspective.

Rineke Schram

Lead Assessment and Remediation Adviser



Case summary

Setting: GP
practice –
resident doctor
under supervision

- Policy: Chaperone required only for intimate examinations
- Event:
- Young female patient, chest (heart) examination clinically indicated
- Allegation: Doctor touched patient's breast (over bra) during examination

Outcome:

- Immediate exclusion pending investigation
- Police investigation – no further action due to insufficient evidence
- Internal investigation discontinued

Key issues raised

Consent:

- Was explicit, informed consent obtained before examination?
- Did the patient fully understand what the examination involved?

Chaperone policy:

- Does the 'intimate examination only' policy give sufficient protection?
- Should the chaperones be offered more widely, especially for cross gender examinations?

Communication:

- Were examination steps explained in real time?
- Was the patient given an opportunity to ask questions or refuse?



Potential preventive actions (Doctor)

Clear Pre-Examination Explanation:

Explain why the exam is needed, what areas will be touched, and duration

Explicit Consent:

Ask directly: 'Is it OK if I proceed?'

Chaperone Offer:

Offer regardless of policy when patient may feel vulnerable

During the Examination:

Describe actions before and as they occur ('I'm now going to place my hand on...')

Post-Examination Check:

Ask if the patient has any concerns or questions

Potential preventive actions (Practice)

Chaperone Policy Review:

Consider making chaperone offer standard for examinations of sensitive areas

Staff Training:

Communication skills for explaining examinations
Handling and documenting consent

Documentation:

Record that consent was obtained, chaperone offered/declined, and patient's response

Patient Information:

Posters or leaflets explaining chaperone availability and patients' rights

Learning points



Even clinically appropriate examinations can lead to complaints if communication and consent are unclear



Offering a chaperone can protect both patient and clinician



Documentation of consent and chaperone discussions is critical



A proactive consent and chaperone culture supports trust and safety

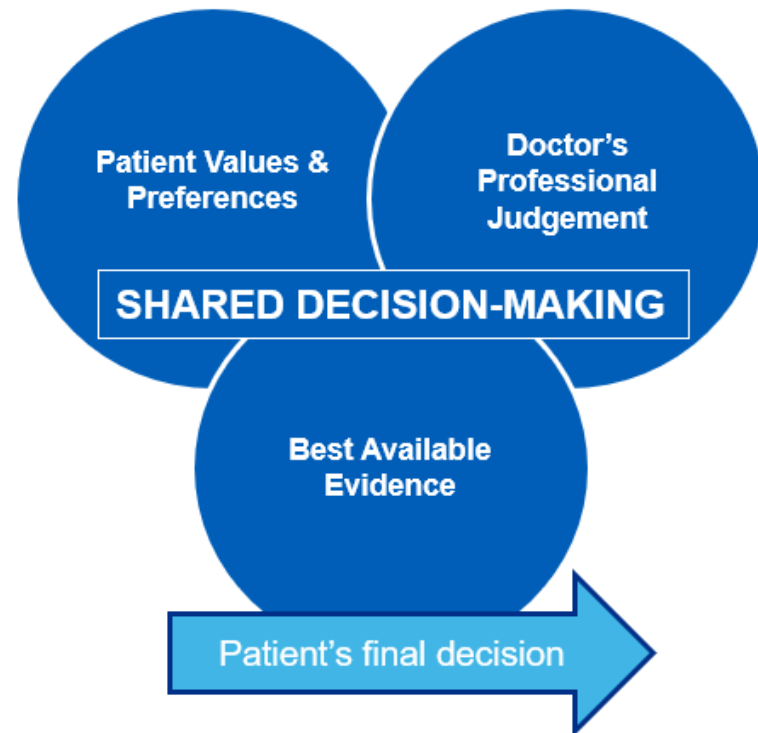
Consent in Practice – Balancing Values, Judgement, and Evidence

Core elements in seeking consent

- Patient – values, beliefs, lifestyle, preferences; priorities for outcomes and risks
- Doctor – professional knowledge, clinical judgement; duty to explain material risks and alternatives (Montgomery ruling)
- Evidence – clinical data, guidelines, treatment/procedure outcomes; benefits, harms, uncertainties

Shared decision-making framework

- Dialogue – open, honest exchange of information
- Understanding – patient comprehends benefits, risks, alternatives
- Respect – patient's perspective guides options discussed



Consent in Practice – Balancing Values, Judgement, and Evidence



Key principle

- Montgomery standard: Material risks are those a reasonable patient would want to know, and those the clinician should know are significant to that patient.

Final decision

- The patient is the decision-maker – the clinician advises, informs, and supports. Consent must be informed, voluntary, and given by a patient with capacity.

Contact Practitioner Performance Advice



Resolution



020 7811 2600



nhsr.advice@nhs.net



NHS Resolution,
8th Floor
10 South Colonnade
Canary Wharf
London
E14 4PU

Events team:

nhsr.adviceeducation@nhs.net



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<https://resolution.nhs.uk>

Q&A led by Ed Glasgow, Partner – Kennedys

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SCAN ME

<https://forms.office.com/e/Q9CsgWnHZs>

Breakout groups

We are now going to split into three breakout groups:

1. Acute
2. Mental Health
3. Community Trust & Other

Please choose the room that reflects your organisation

Summary & Close

We hope you have found this event useful,
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Thank you for your time

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