



# Staff Survey: Patient Experience ERAS UK 2013

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## The NHS Outcomes Framework 2013

### NHS Patient Experience Framework

In October 2011 the NHS National Quality Board (NQB) adopted a definition of patient experience to guide the measurement of patient experience across the NHS. This framework outlines those elements which define patients' experience of NHS Services.

- **Respect for patient-centred values, and respect for patients' expressed needs**, including: cultural issues; the independence of patients and service users; an awareness of patients' issues; and shared decision making;
- **Coordination and integration of care** across the care system;
- **Information, communication, and education**, including: progress, prognosis, and processes of care in order to support care and health promotion;
- **Physical comfort** including pain management, health and safety, living, and clean and comfortable surroundings;
- **Emotional support** and alleviation of fear and anxiety, including: as clinical status, prognosis, and the impact of illness on patients and their finances;
- **Welcoming the involvement of family and carers**, including: patients and service users rely, in decision-making, on the awareness and accommodation of their needs as care-givers;
- **Transition and continuity** as regards informal carers, including: care for themselves away from a clinical setting, and support to ease transitions;
- **Access to care** with attention for example, to the time between admission or time between admission and placement in a care setting, and waiting time for an appointment or visit in a care or social care setting.

This framework is based on a modified version of the Picker Institute's Principles of Patient-Centred Care, an evidence based definition of a good patient experience. When using this framework the NHS is required under the Equality Act 2010 to take account of its Public Sector Equality Duty including eliminating discrimination, harassment and victimisation, promoting equality and fostering good relations between people.

NCGC National Clinical Guideline Centre

## Patient experience in adult NHS services: improving the experience of care for people using adult NHS services

Patient experience in generic terms

*Clinical Guidance*

*Methods, evidence and recommendations*

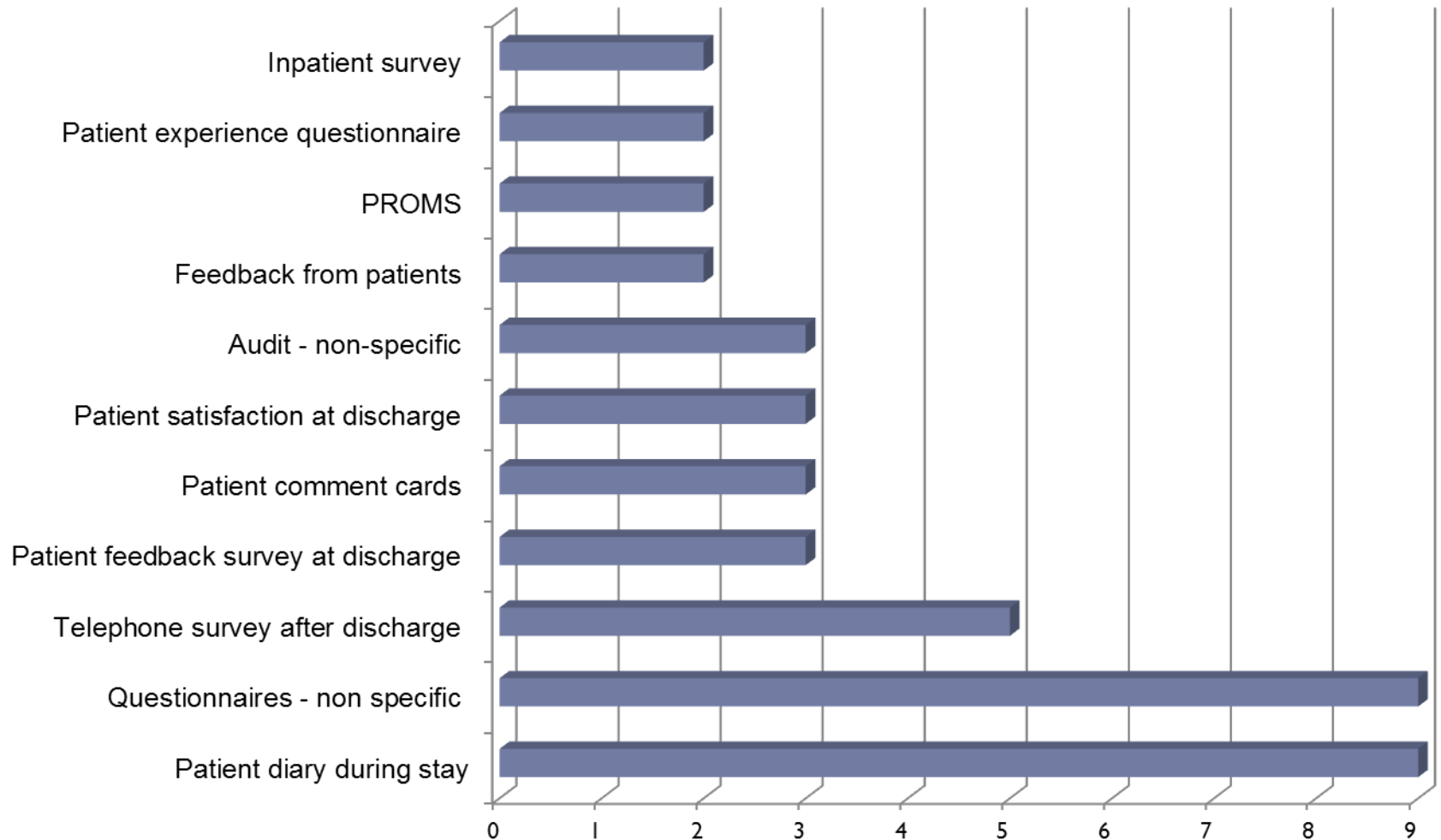
*February 2012*

*Commissioned by the National Institute for Health and Clinical Excellence*

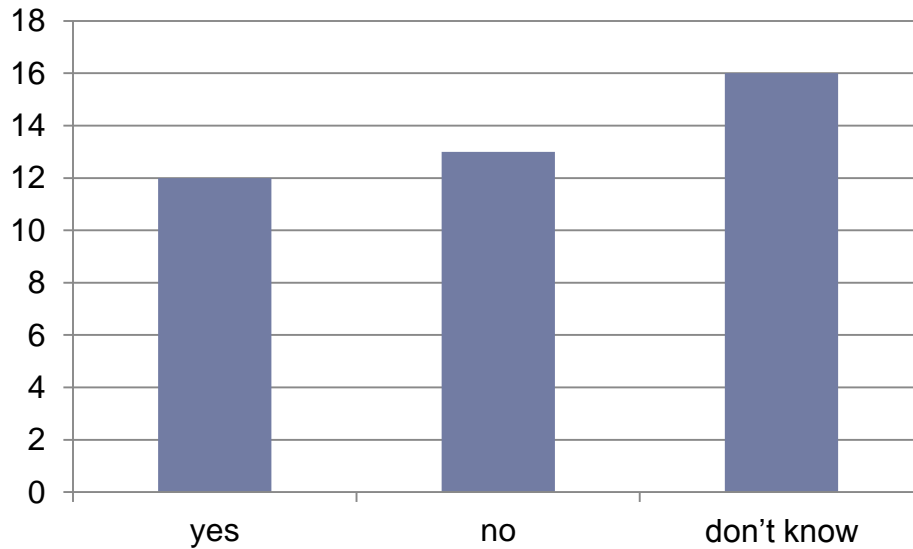




# Methods of measurement: 2012



# Detecting patient experience: 2012



Yes - the questionnaires are anonymous (4)

No - We do not know what is important for a good or bad experience (2)

I don't know – Not enough data to judge this (9)

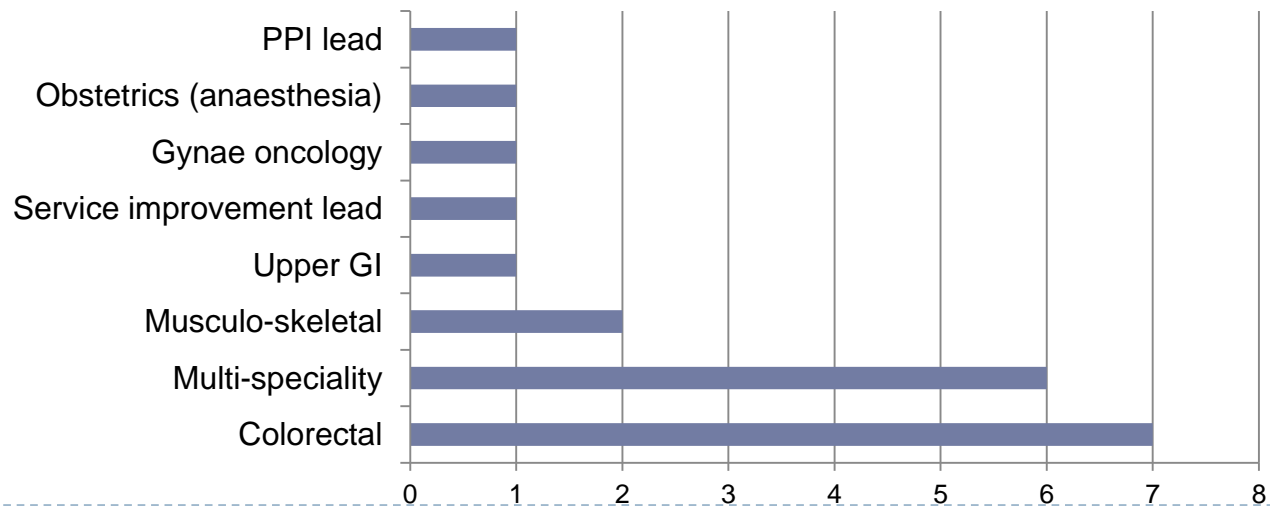
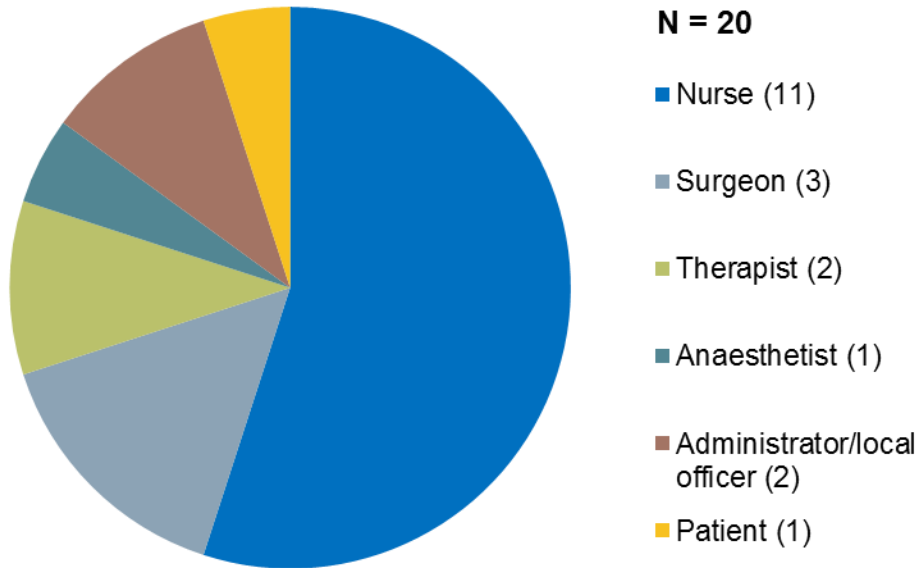
Survey, questionnaires and diaries often not completed (4)

Bias: patients know the staff (3)

Are patients honest about us when they are in hospital? (3)



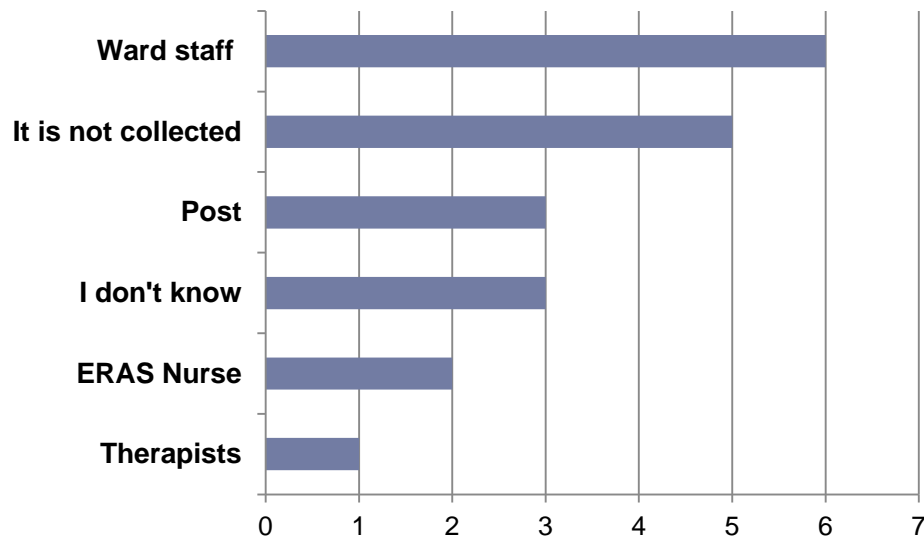
# ERAS UK Staff Survey 2013



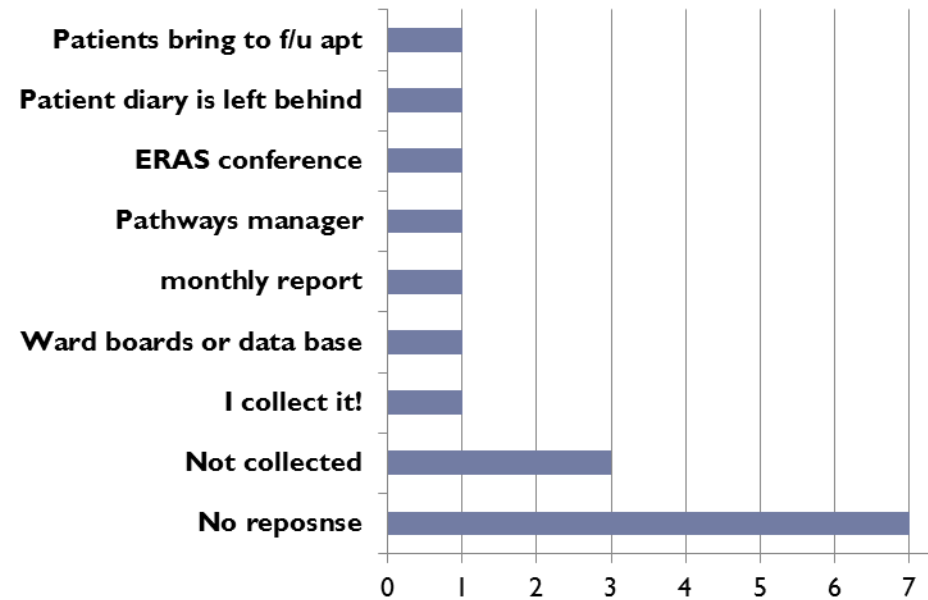
# Capture of PROMs in ERAS

“Any report of the status of a patient’s health condition that comes directly from the patient themselves, without interpretation by a clinician or anyone else”

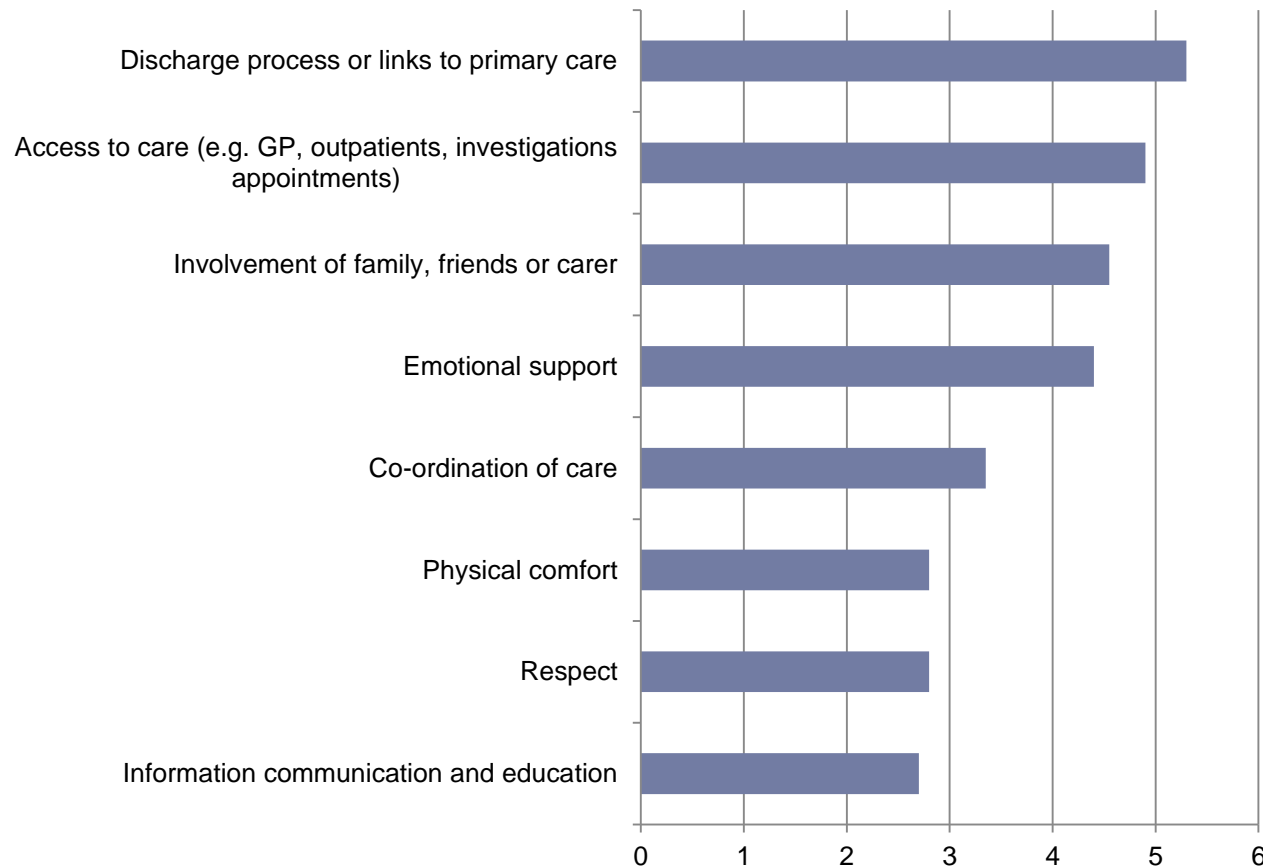
## Collection



## Feed back



# Ranked dimensions of experience (1 = most important)



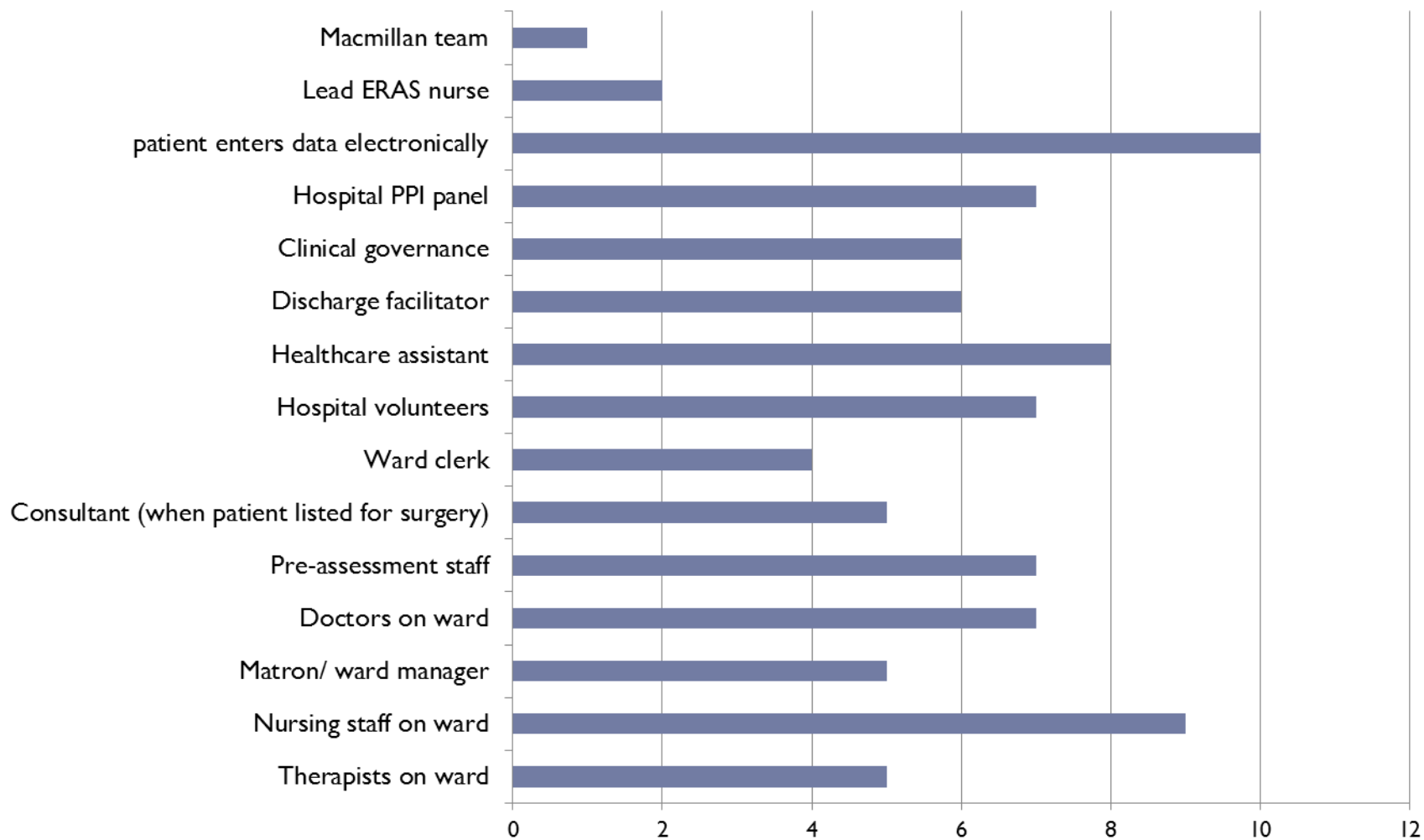


# Measurement of experience dimensions

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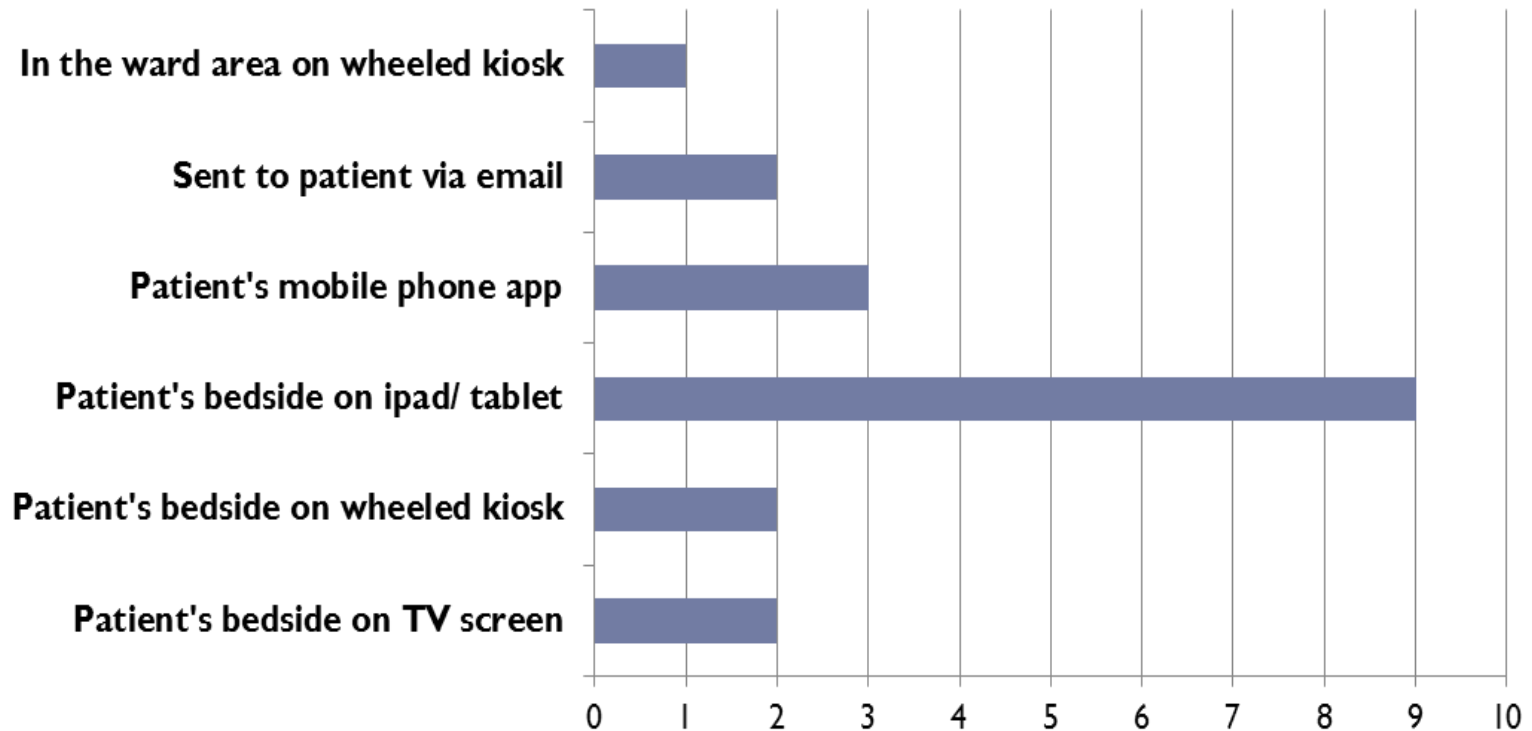


# Who should collect patient experience data?



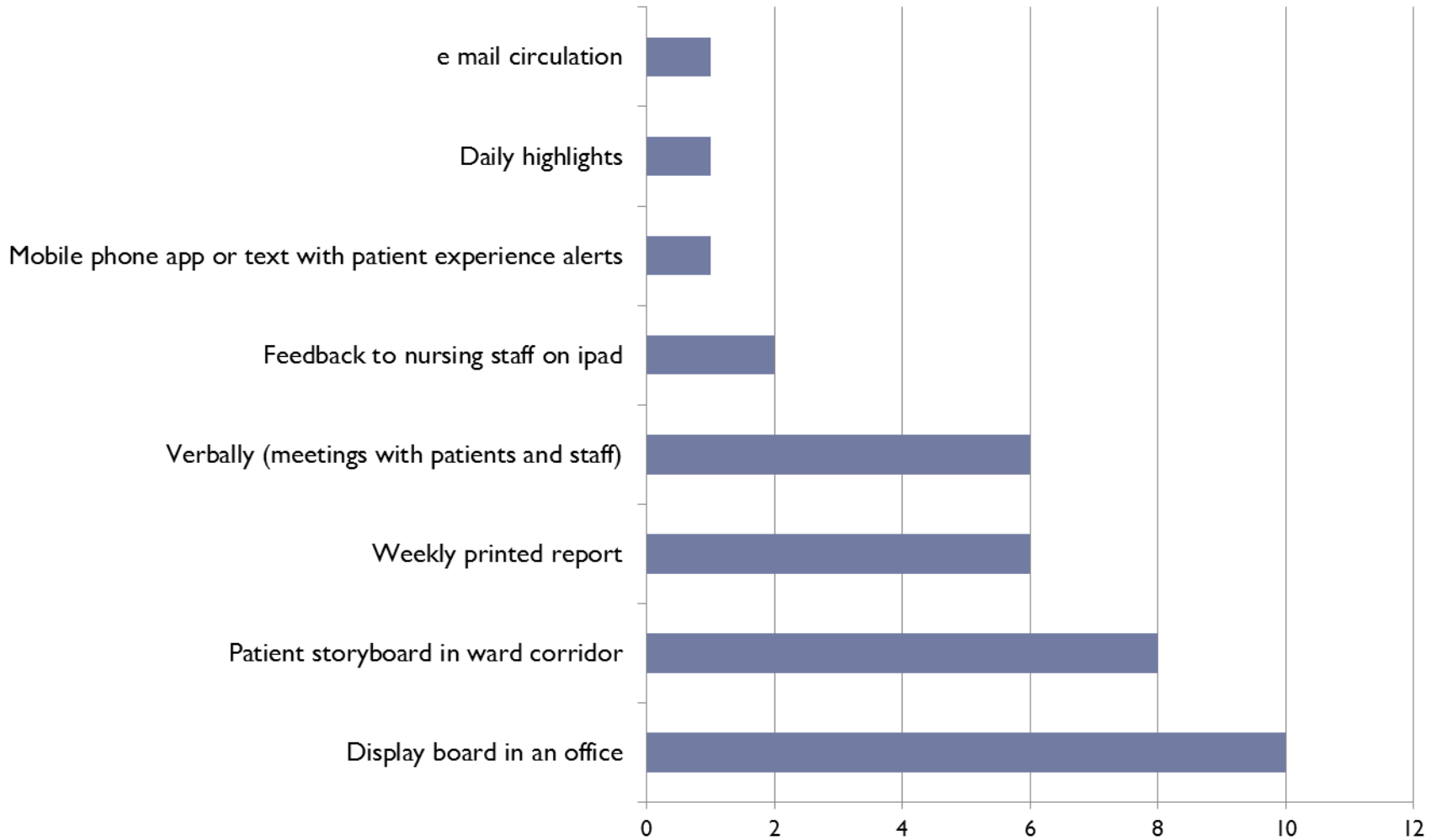
# Electronic data collection

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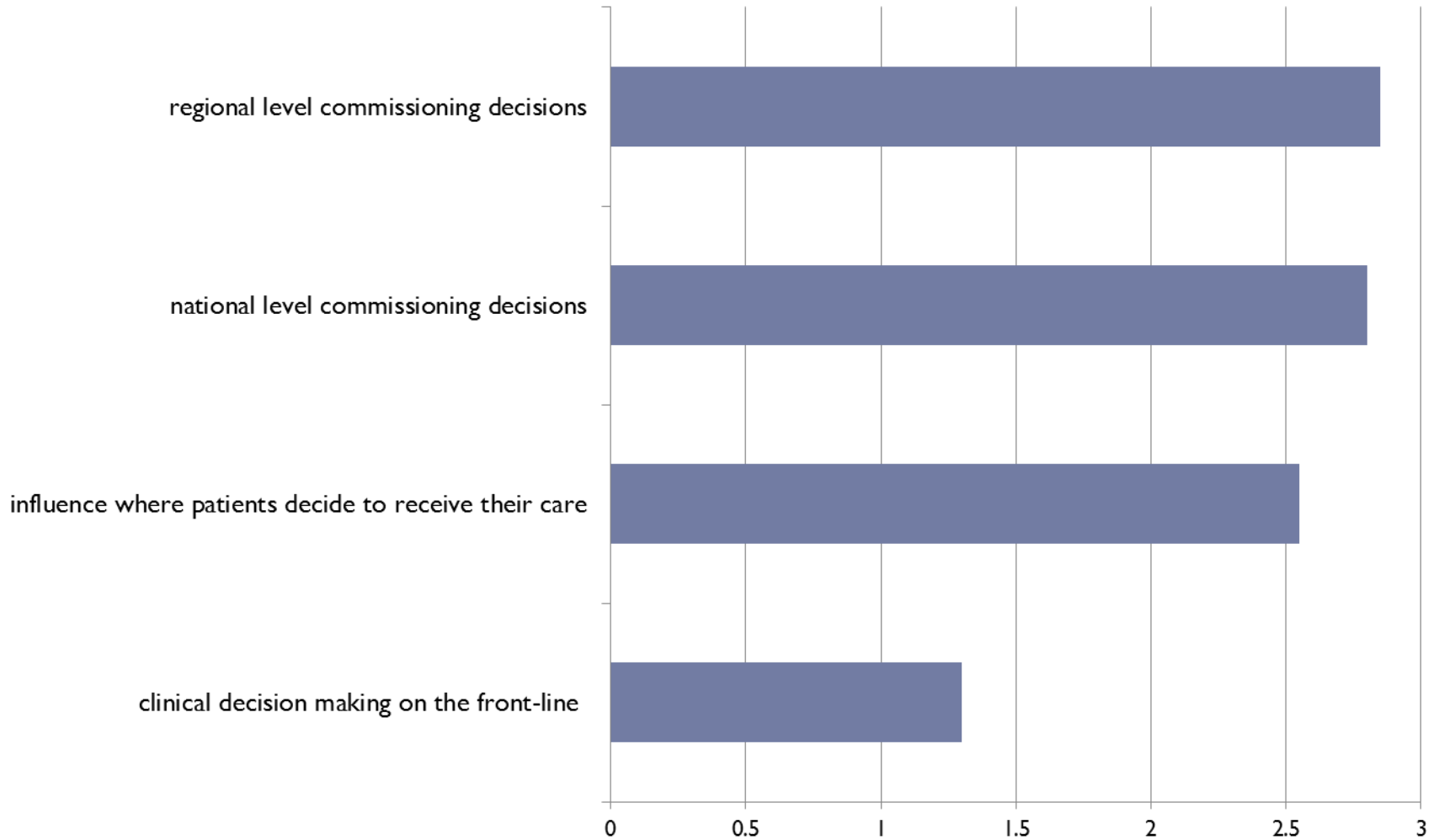
# Feed back of patient experience data

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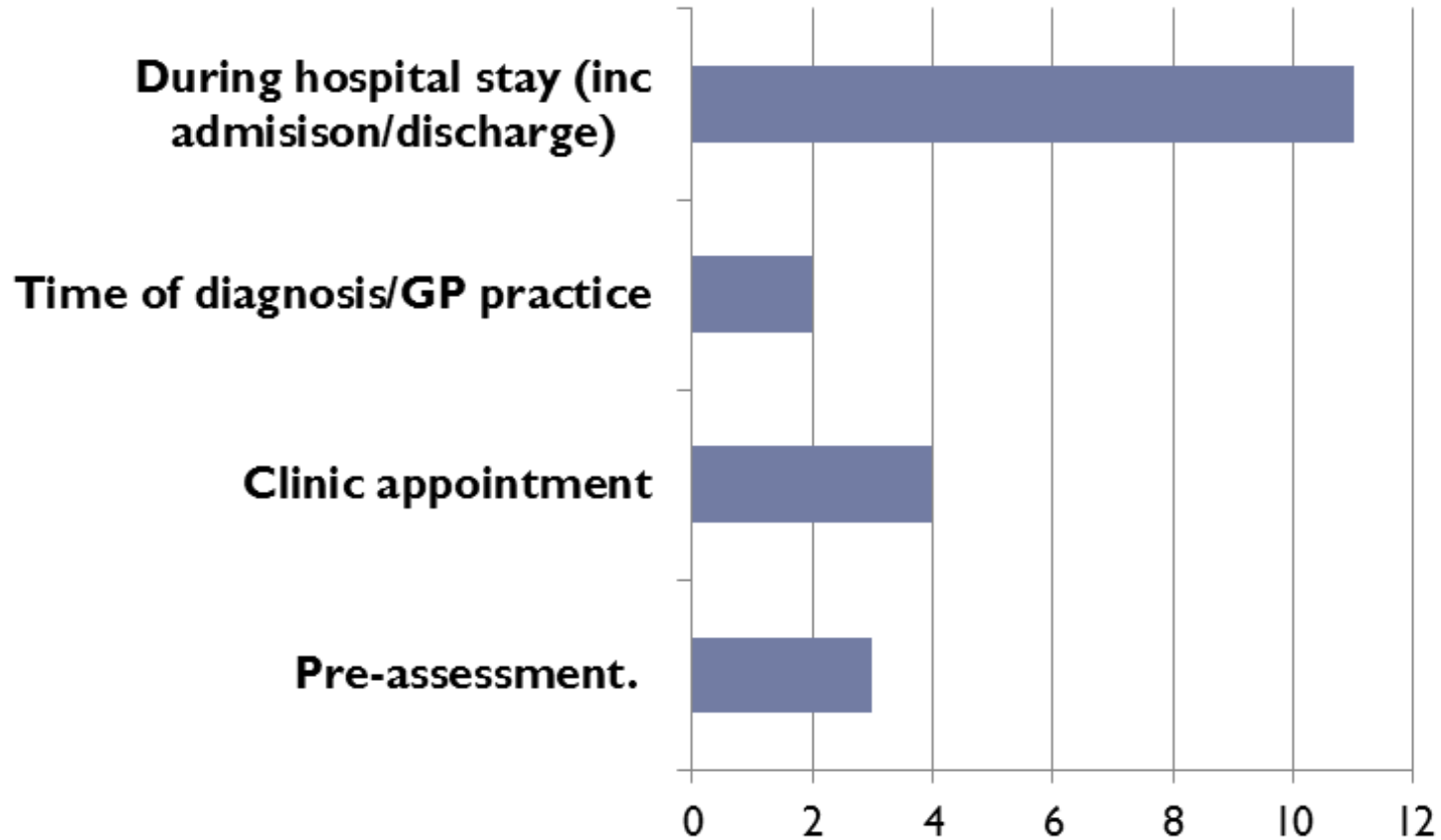
# Ranked utility of patient experience data (1 = most important)

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# Timing of patient experience data collection

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# Conclusions

- ▶ PROMs data feed-back is sub-optimal
- ▶ Staff perceive 'Information communication and education' as important
- ▶ Verbal and electronic data collection could be important across the patients whole journey

# The way forwards

- ▶ Why is this?
- ▶ Does this correlate with what patients think?

| FMC Streams       |     | Wait to See |   |     | Treat |     | Wait For Bed |     |     | Total Time in ED |     |    |  |   | EECU |  | Tot |
|-------------------|-----|-------------|---|-----|-------|-----|--------------|-----|-----|------------------|-----|----|--|---|------|--|-----|
| Stream            | Tot | WOT         | * | Tot | Tot   | WOT | *            | 0-2 | 2-3 | 3-4              | 4-6 | 6+ |  |   |      |  |     |
| LIKELY TO ADMIT   | 1   | 0           |   | 8   | 8     | 2   | ●            | 7   | 0   | 1                | 2   | 7  |  | 2 | 19   |  |     |
| LIKELY TO DISCHAR | 3   | 0           |   | 20  | 1     | 0   |              | 15  | 2   | 2                | 3   | 2  |  | 0 | 24   |  |     |
| RESUSCITATION     | 0   | 0           |   | 0   | 1     | 0   |              | 0   | 0   | 0                | 0   | 1  |  | 0 | 1    |  |     |
| UNCLASSIFIED      | 0   | 0           |   | 0   | 0     | 0   |              | 0   | 0   | 0                | 0   | 0  |  | 3 | 3    |  |     |
| Total             | 4   | 0           |   | 28  | 10    | 2   |              | 22  | 2   | 3                | 5   | 10 |  | 5 | 47   |  |     |

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“Clinicians’ views about outcome may not always be in the best interest of the patient”

“What patients want may not be what we perceive they want”

**Patient Experience**

“The method is not important, what matters is that patients have the opportunity to feed back”

“Data should be collected all the time: if our patients are not happy we should solve it there and then”

“Patient Experience is probably the only thing that matters”





# Implementing patient experience data: workshop

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- ▶ 1) How should we collect patient experience – dimensions, and methodology
- ▶ 2) Involving patients, carers and service users in the collection, and actioning of patient experience
- ▶ 3) Pragmatism in real time patient experience data collection –engaging frontline staff in developing responsive services

