

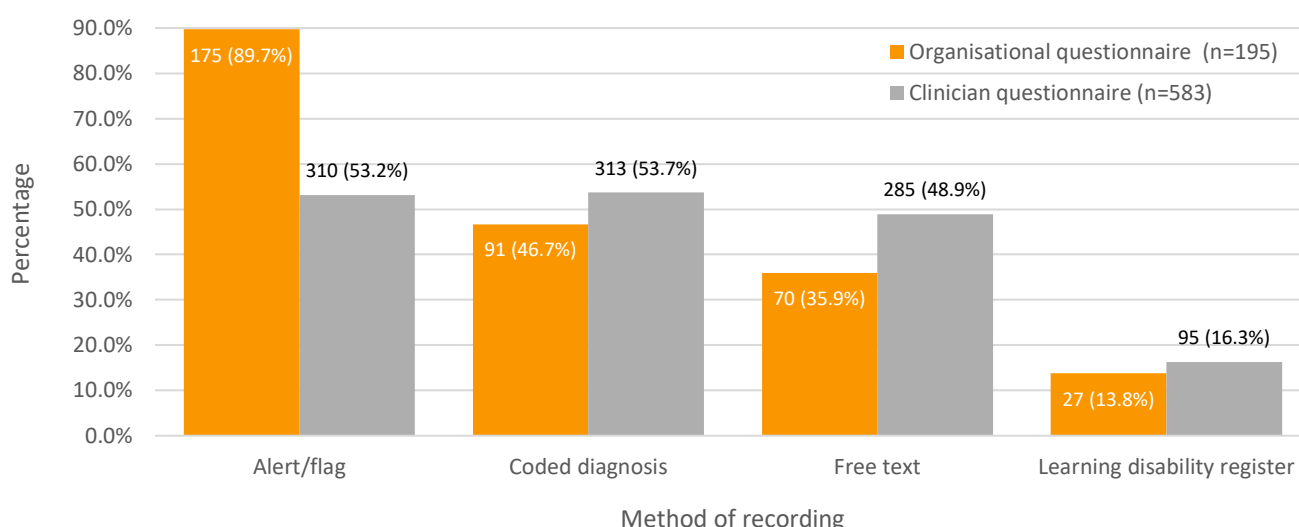
### 3 IDENTIFICATION AND RECORDING OF PATIENTS WITH A LEARNING DISABILITY

#### Hospital records systems

The Equality Act 2010 requires public bodies to consider all individuals when delivering services so that people with a disability are not disadvantaged when accessing healthcare.<sup>[12]</sup> It is recommended that a learning disability should be recorded in hospital case notes and electronic health records to enable healthcare professionals to identify if someone has a learning disability when they arrive at hospital.<sup>[13]</sup>

It was possible to identify a patient with a learning disability on the patient record system in 195/199 (98.0%) acute hospitals.

The way in which patients with a learning disability were identified and recorded varied by organisation. A total of 175/196 (89.7%) organisations reported using alerts or flags on electronic patient records, while clinicians reported only 310/583 (53.2%) patients had such alerts (F3.1).



#### F3.1 How a learning disability is recorded by data source

*Organisational and clinician questionnaire data. Answers may be multiple*

Review of the case notes showed that 119/366 (32.5%) patients were described as having a learning difficulty rather than a learning disability and that the two terms were often recorded interchangeably throughout the patients' notes. If patients with a learning disability were miscoded as having a learning difficulty throughout an admission they would not have been identified as part of the study population, and more importantly this could impact their care while in hospital.<sup>[14]</sup>

Reviewers gave numerous examples of the confusion between a learning disability and a learning difficulty. One reviewer noted "shortened to 'LD' so not sure what they mean" and another noted that the terms were "used interchangeably in the same assessment by the same medic".

Healthcare professionals in acute settings and the community also reported that it was not easy to identify patients with a learning disability from electronic patient records or hospital systems, noting that

it was slightly harder for healthcare professionals in acute care settings than for those in the community (178/415; 42.9% vs 171/301; 56.8%) (T3.1).

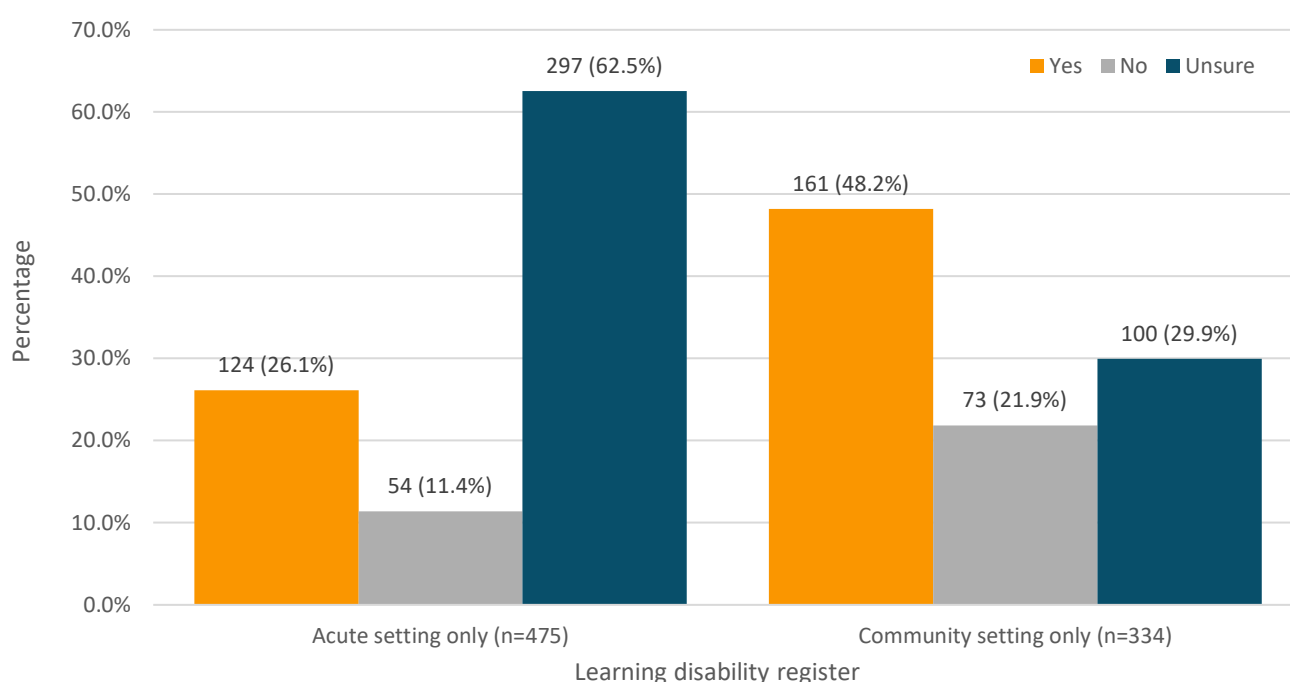
T3.1 The ease of identification of patients with a learning disability on patient record systems	Acute (physical health) setting		Community/primary care setting	
	Number of responses	%	Number of responses	%
Yes	178	42.9	171	56.8
No	237	57.1	130	43.2
<b>Subtotal</b>	<b>415</b>		<b>301</b>	
Unsure	54		33	
Not applicable	15		14	
<b>Total</b>	<b>484</b>		<b>348</b>	

Health and social care professional survey

This lack of clarity was compounded by unclear diagnoses and the use of a variety of terms to describe a learning disability. Often a patient would be described as having ‘Down Syndrome’ due to its strong association with learning disabilities and outdated terms such as ‘mental retardation’ were also still sometimes used.

A register of people with a learning disability could support a hospital response to the delivery of care. However, it was reported that only 27/195 (13.8%) acute hospitals had a register.

More widely there were 285/809 (35.2%) health and social care survey respondents who reported that their organisation had a learning disability register, but this was much more common in community settings than in acute settings (F3.2). In addition, 297/475 (62.5%) survey respondents working in acute settings were unsure whether a register existed in their organisation. Therefore, if a register is developed, teams will need to be aware that it is available.



### F3.2 Learning disability registers within organisations

Health and social care professional survey