# Challenges of conducting an audit of pressure ulcer monitoring systems in NHS England



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#### Background

A number of initiatives have been introduced throughout the NHS to reduce avoidable pressure ulcer harm, including monitoring systems such as:

- Safety Thermometer<sup>1</sup> (STh)
- Incident Reporting Systems (IRS)<sup>2</sup> (e.g. Datix/Ulysses)
- Strategic Executive Information System (STEIS) for the reporting of Serious Incidents Requiring Investigation<sup>3</sup> (SIRI).

Concerns about inconsistencies in the local implementation of these systems and over-interpretation of data, prompted the Tissue Viability Society (TVS) to fund a project, supported by NHS England, to assess the accuracy of pressure ulcer monitoring in England and to inform the interpretation and further development of pressure ulcer monitoring.

The results of this project were presented at EPUAP 2015 and are now published<sup>4,5</sup>.

This poster focuses on the challenges of conducting the audit of pressure ulcer monitoring systems in NHS England.

### Summary of results

Reporting System	Sensitivity (95% CI)	Specificity (95% CI)
STh (weighted estimates)	48.2% (35.4%-56.7%)	99.0% (99.0%-99.0%)
IRS (unweighted estimates)	53.4% (46.3% to 60.4%)	98.3% (97.7% to 98.8%)

Table 2: overall accuracy of current monitoring systems

Under-reporting of pressure ulcers observed across monitoring systems

Correct classification when skin damage is identified across systems

- PUs not reported as IADs
- IADs not reported as PUs
- Other wounds not reported as PUs

When both the PUWA and monitoring system report a pressure ulcer on the same skin site:

- Good levels of accuracy classification
- Good levels of accuracy origin of pressure ulcer

## Learning points

- Important to understand the data and working definitions of the systems you are assessing including the way these are implemented in practice.
- Important to interpret the results in combination with qualitative survey results to add **Context** to the results
- To evaluate the accuracy of tools used for clinical management of pressure ulcers, robust data collection and analysis methods are required.
- A good team and co-ordinated approach is required

#### Methods and Data Collection

The project aimed to compare and contrast current data sources including in-patient STh prevalence data, IRS and STEIS incident data against a 'gold standard' Pressure Ulcer/Wound Audit (PUWA) and if appropriate develop proposals for a standardised approach to pressure

To facilitate comparison of different monitoring systems it was important to understand the working definitions associated with each system and these are detailed in table 1. This demonstrates variation across monitoring systems and centres.

ulcer monitoring.

As per local Trust policy Origin 4. U or DTI at the time of admission within 72 hours of admission to your organisation. systemα A pressure ulcer that developed 72 hours or Not POA more after the patient was admitted to your A pressure ulcer which was not POA Acquired (HA) organisation. <sup>µ</sup> Pressure **Ulcer Data** Recorded 'Current category' recorded as the category of PU Current observed at the PUWA skin assessment All data recorded for a pressure For New and Old the classification of the patient during their 'Worst category' recorded as the worst category† ulcers worst pressure ulcer should be reported. current admission should reported in the patient's clinical records during this be recorded, including a classifications of ulcers 'Worst category' recorded as the worst category<sup>†</sup> Healed from all reports listed. N/A reported in the patient's clinical records during this

**Table 1: Definitions** 

\*This is the definition for a 'Old' ulcer on the Safety Thermometer; <sup>µ</sup> This is the definition for a 'New' ulcer on the Safety Thermometer; <sup>α</sup> Questionnaire responses indicated that only 10 use on admission definition; 12 use within 72 hours of admission; † Severity of classification from worst to best is 4, 3, Unstageable, DTI, 2; § Defined as complete re-epithelialisation in the absence of a scab including normal or erythematous skin.

This was an audit and used anonymised data, ethical approval was not required. However, verbal consent for skin inspection was obtained in line with usual clinical procedures and care. To allow comparison of monitoring systems all data was recorded in a single booklet by 3 nurses (figure 1).

ulcers

Comprehensive **training** for the *audit process* was required. We liaised with participating Trusts audit lead/staff to ensure that that everyone was clear about their role within the audit process prior

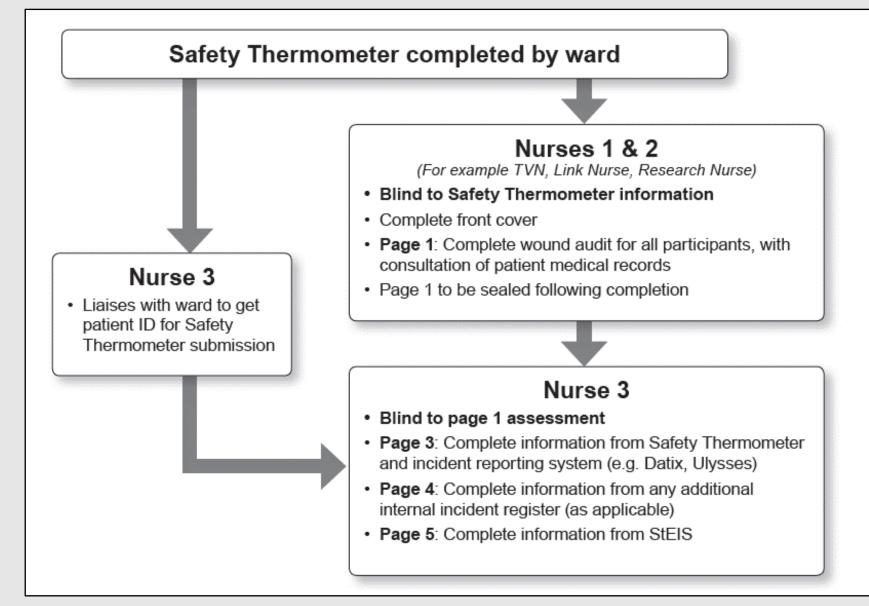
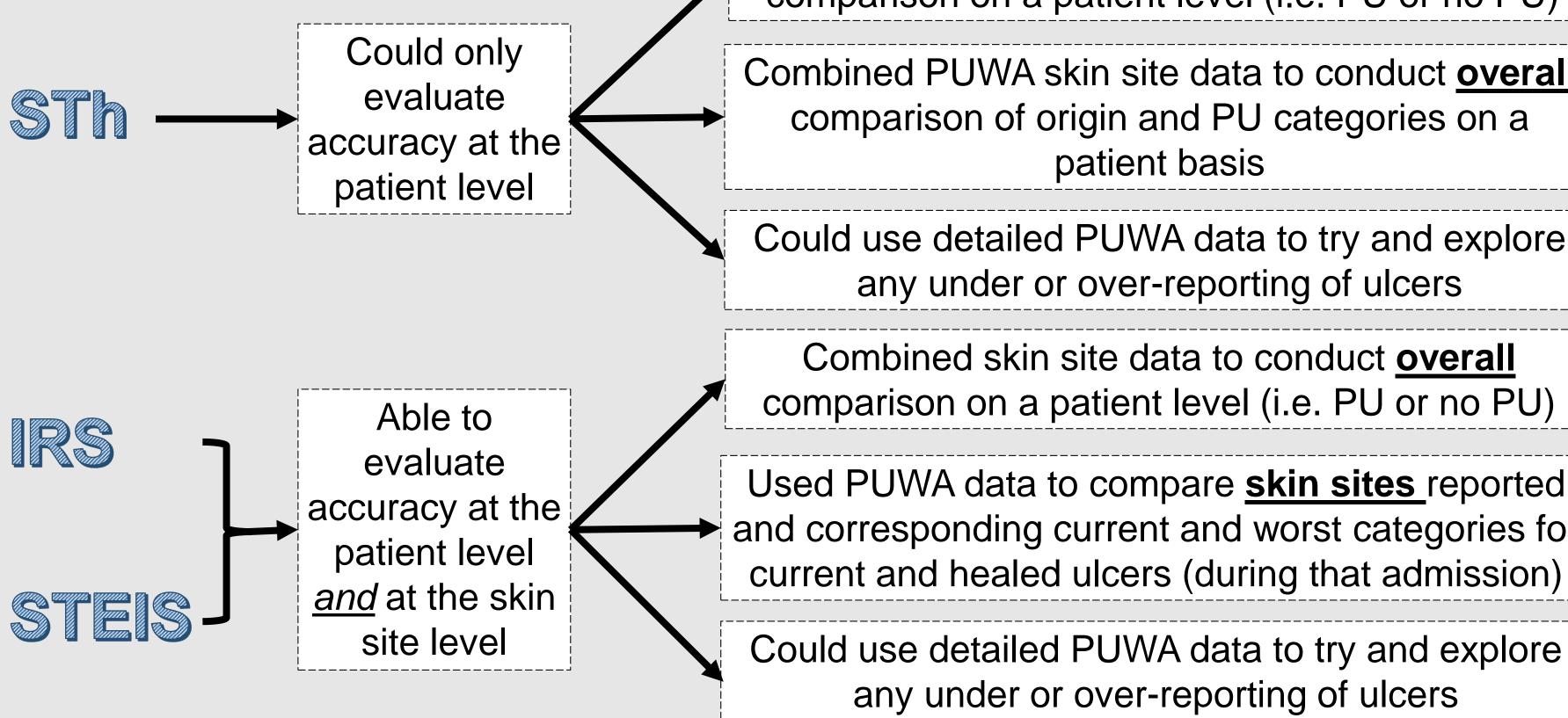


Figure 1: Audit process

to data collection on the October STh census. The assessors, were members of the Tissue Viability Team or ward based expert nurses and experienced in undertaking skin assessment and were given no additional training prior to the audit taking place

### Analysis

Data obtained from the PUWA required restructuring and combining depending on each system:



Combined PUWA skin site data to conduct **overall** comparison on a patient level (i.e. PU or no PU) Combined PUWA skin site data to conduct **overall** 

Could use detailed PUWA data to try and explore any under or over-reporting of ulcers

patient basis

Used PUWA data to compare **skin sites** reported and corresponding current and worst categories for current and healed ulcers (during that admission)

Could use detailed PUWA data to try and explore any under or over-reporting of ulcers

Other considerations:

- Weighting accuracy measures according to the sampling of Trusts
- Timelines for producing study report
- Preparation for analysis and reporting is key

#### **Acknowledgements**

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