

How to use artificial intelligence tools to predict and prevent care-related harm

Context

In the Australian setting, risk prediction aligns with the goals of the National Safety and Quality Health Service Standards (especially Clinical Governance and Preventing and Controlling Healthcare-Associated Infection Standards), as well as the aims of the RACP Evolve initiative, which seeks to reduce low-value care and promote evidence-based practice.

Low-value care tests, treatments, or procedures that provide little or no clinical benefit remains a significant challenge in the Australian health system. Research suggests that up to one in three healthcare services delivered may be of low or uncertain value, contributing to avoidable patient harm, inefficiencies, and escalating costs.¹

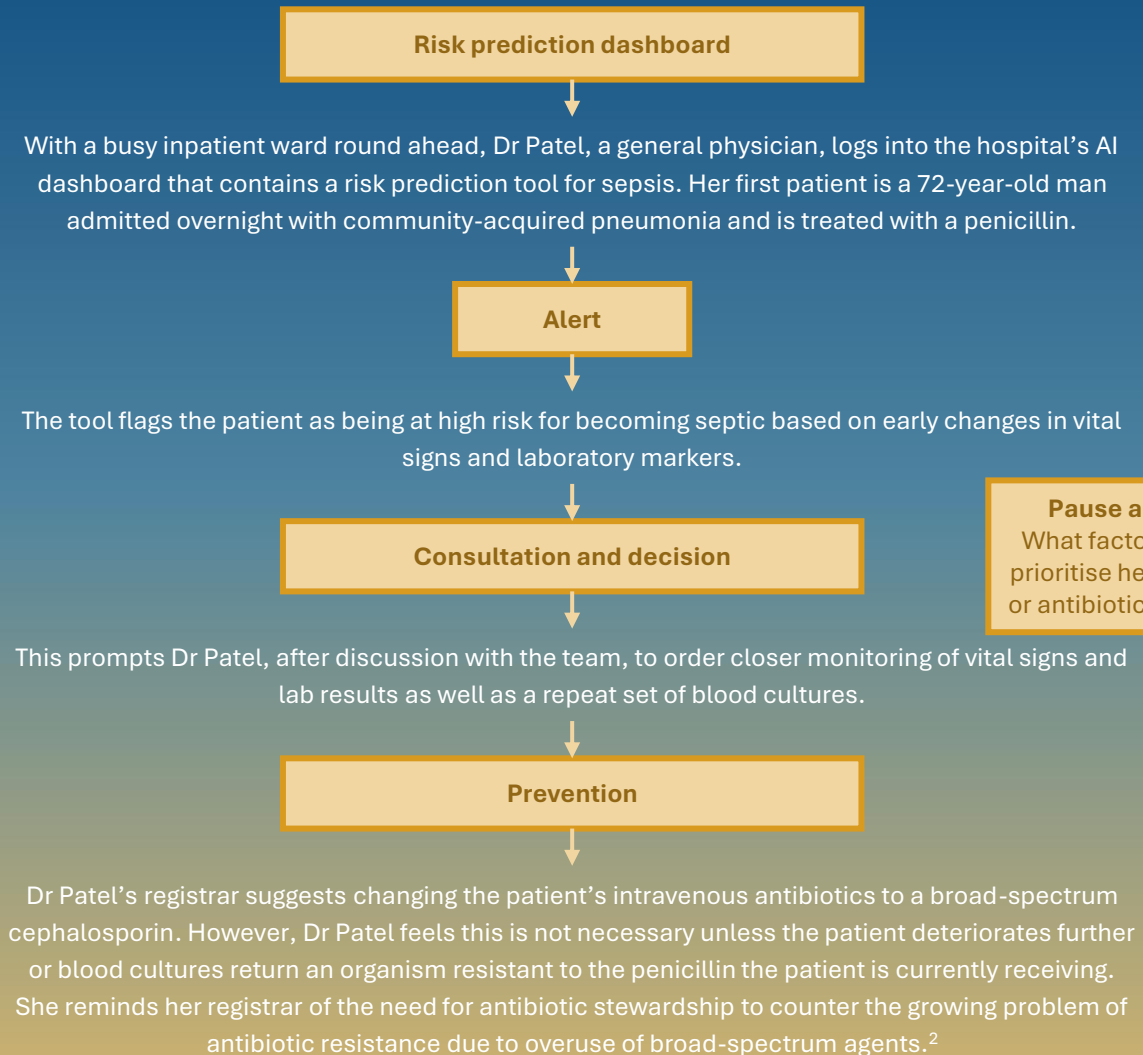
Artificial intelligence (AI)-driven risk prediction tools are being explored across healthcare to identify patients at risk of harm before it occurs – such as deterioration, sepsis, falls, or unplanned readmissions – and to target such individuals with evidence-based preventive strategies and avoid low-value practices that provide limited clinical benefit.

Ensuring physicians evaluate and effectively use AI tools is essential to addressing low-value care and improving patient outcomes and system sustainability.

Ideal future

AI risk prediction tools are being used in the hospital environment to flag a patient as being at higher risk of disease by analysing subtle trends in vital signs, lab results, and clinical notes contained in electronic medical records (EMRs). The tools are typically trained on large datasets obtained from EMRs containing clinical, demographic, and administrative data to identify patterns associated with risk of future harm and enable early detection.

Clinical scenario



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Benefits

Earlier intervention and prevention: Clinicians are alerted to patients at risk of certain harms who warrant early initiation of preventive measures before harms occur.

Efficiency in resource allocation: High-risk patients can be identified, enabling interventions to be targeted to them while avoiding potentially unnecessary and wasteful care directed at those with lower risk.

Scalability and automation: AI tools in their training and deployment can process vast amounts of data, enabling rapid and more accurate risk prediction than current rule-based algorithms which often require manual data entry and human calculations.

Supporting clinical decision-making: AI tools can provide an additional insight to complement clinical judgement.

Quality improvement: Analyses of aggregated data from risk prediction tools can inform policy, guideline development, and education.



Risks

Data quality and bias: AI tools trained on flawed or incomplete data may misclassify risks or perpetuate inequities (e.g. among Indigenous or rural patients), negatively affecting the AI tool's external validity.

Alert fatigue: Excessive or non-specific alerts may reduce clinician engagement or even lead to dismissal of critical warnings.

Over-reliance: Clinicians may defer too readily to the tool rather than apply clinical reasoning, particularly in high-pressure settings.

False sense of objectivity: AI outputs may appear neutral and evidence-based, but underlying biases in training data and design such as encoding historical racial or socioeconomic disparities, underrepresenting rare conditions, or favouring billing-driven care patterns can perpetuate existing inefficiencies or low-value practices.

To find out more about the use of AI in medicine, see the *Evolve AI in Healthcare* webinar series on the Evolve website:

<https://www.racp.edu.au/evolve/home>

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Key considerations for physicians using AI tools to predict and prevent care-related harm



Understanding development and evaluation:

- Check whether the tool has undergone external validation in comparable Australian settings and patient groups. Use frameworks such as SALIENT³ and DECIDE-AI⁴ for early-phase clinical evaluation.
- Assess tool performance using metrics such as AUROC, AUPRC, sensitivity, and specificity, and question whether use of a seemingly high performing tool translates into meaningful improvement in patient outcomes.
- Determine whether the tool's predictive accuracy changes across different patient subgroups or in different clinical settings.



Balancing effectiveness, efficiency and utility⁶:

- Evaluate whether the tool integrates seamlessly into existing workflows without creating extra administrative burden. Use frameworks such as SALIENT³ to assess end-to-end implementation.
- Consider trade-offs in selecting the risk threshold for generating alerts: a tool optimised for sensitivity (i.e. lower risk threshold) may capture more at-risk patients however creates inefficiencies through generating more false positives in patients who have no or little risk, thus worsening alert fatigue. Conversely, a tool prioritising specificity (i.e. higher risk threshold) will reduce false alerts but will miss more at-risk patients.
- Determine if risk prediction tools actually alert clinicians to patients at risk of harm at an earlier time than clinicians would have recognised based on clinical judgement alone.
- Examine whether it supports stewardship of care by not causing overuse of care in many patients at low risk as the tool is too sensitive, while simultaneously maintaining patient safety by not missing many patients at high risk as the tool is too specific.



Maintain clinical accountability:

- AI tools should augment, not replace clinical decision-making.
- Physicians remain accountable and must weigh AI outputs against patient context and preferences.
- Discuss the reasons for considering AI-generated risk predictions with patients and carers to ensure trust and acceptability.



Governance and monitoring:

- Ensure compliance of AI tool use with the Privacy Act 1988⁵ and local data governance frameworks.
- Ensure compliance with regulatory standards that apply to AI tools which influence clinical decision-making.
- Advocate for continuous monitoring and recalibration of tools to maintain safety and equity.

Glossary

AUROC:

Area under the receiver operator characteristic curve

AUPRC:

Area under the precision-recall curve.

³ Van der Vegt, Scott IA, Dermawan K, et al. Implementation frameworks for end-to-end clinical AI: derivation of the SALIENT framework. J Am Med Inform Assoc 2023; 30(9):1503-1515.

⁴ Vasey B, Nagendran M, Campbell B, et al.; DECIDE-AI expert group. Reporting guideline for the early stage clinical evaluation of decision support systems driven by artificial intelligence: DECIDE-AI. BMJ 2022;377:e070904.

⁵ Office of the Australian Information Commissioner. The Privacy Act. Australian Government, <https://www.oaic.gov.au/privacy/privacy-legislation/the-privacy-act>

⁶ Van der Vegt AH, Campbell VK, Webb R, et al. A novel, standardised approach to balancing effectiveness, efficiency and utility of surveillance AI prediction models for hospitalised patients using sepsis prediction as an exemplar. J Am Med Inform Assoc 2025 Nov 11:ocaf192. doi: 10.1093/jamia/ocaf192. Epub ahead of print.

