Patient safety in oncology: quantifying the hospital-associated risks with a discrete-event simulation

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BACKGROUND

The WHO reports that the occurrence of adverse safety events due to unsafe care is likely to be one of the 10 leading causes of death and disability in the world, and that approximately 50% of patient safety events are considered preventable¹. There are safety risks at every stage of the oncology pathway and it is critical to identify them in order to act and improve patient outcomes.

OBJECTIVES

This research aimed at quantifying the patient safety risks in a hospital setting considering 2 potential oncology scenarios:

- 1: day-case patients getting antineoplastic preparations as part of their intravenous (IV) chemotherapy.
- 2: hospitalized patients undergoing surgery.

METHODS

The patient pathways of the scenarios were mapped according to the UK setting, and their applicability was tested through experts interviews in the EMEA region. The baseline risks associated with each step of the two patient journeys were quantified through desk literature research, including the additional length of stay associated with each potential adverse event. A discrete event simulation was developed to recreate the hospital environment and quantify the overall risks along the patient journeys if no preventive actions were put in place.

RESULTS

The patient journeys were validated in different EMEA hospitals and the following type of risks that could impact the patients, if not prevented, were identified: pre-treatment diagnostic errors leading to delay or incorrect treatment; medication management errors (including dispensing, preparation, and administration) and (including catheter-related hospital acquired infections bloodstream infections; surgical and urinary infections for the surgical pathway only). The simulation quantified that in Scenario 1, every 1'000 patients getting day-case IV chemotherapy, there is a risk of 22 possible complications: 11 in the diagnostic phase and 11 in the medication management process. In Scenario 2, 147 complications can occur, every 1'000 patients undergoing surgery, including 45 surgical site infections and 10 urinary tract infections, leading to more than 3'800 extra bed days in the wards and 396 extra bed days in the intensive care unit.

CONCLUSION

An oncology patient entering a hospital is facing many hazards that can put safety at risk and increase pressure on hospital capacity. The adoption of best practices and effective technologies to prevent such risks should be incentivized by hospital decision makers.

WHAT'S NEW

- 1.Discrete event simulations used to quantify the risks associated with in-hospital oncology patients treatment during their stay
- 2. Possibility to extend a similar approach to other areas of disease

Figure 1: Patient pathways

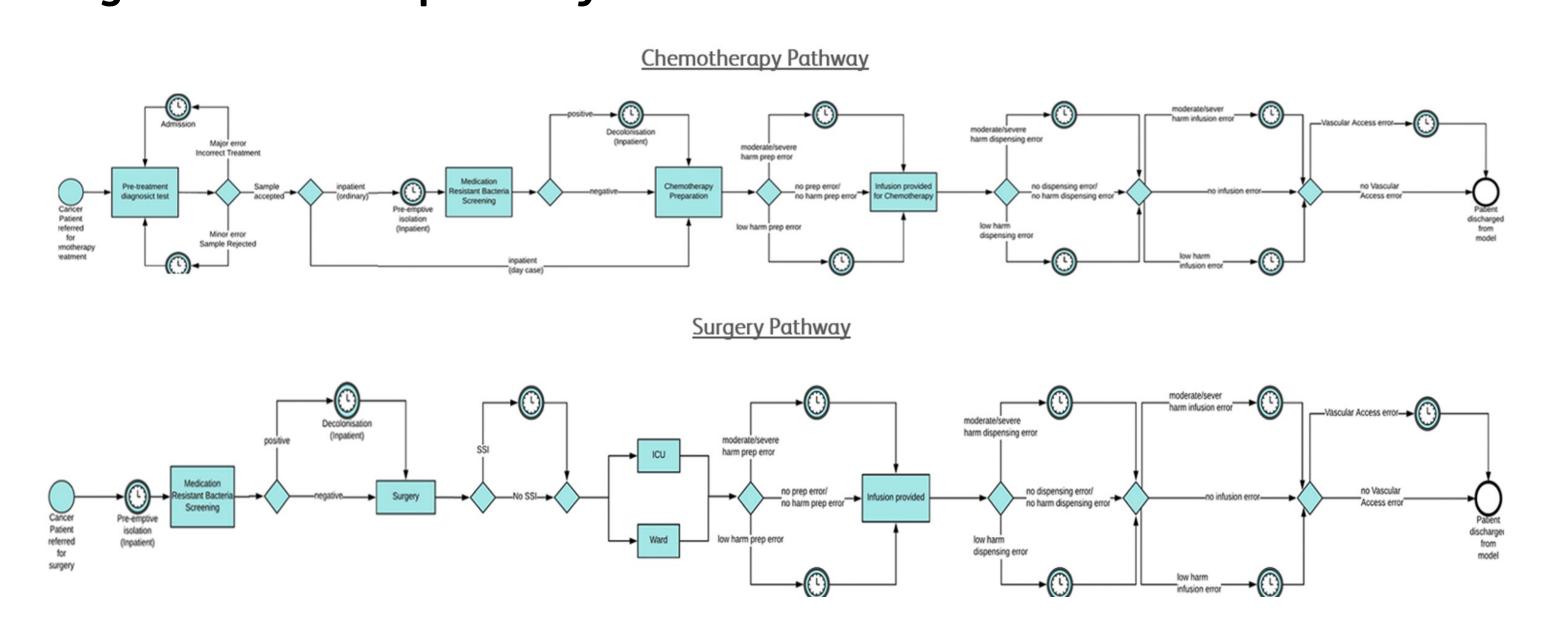


Figure 2: Schematic representation of the model

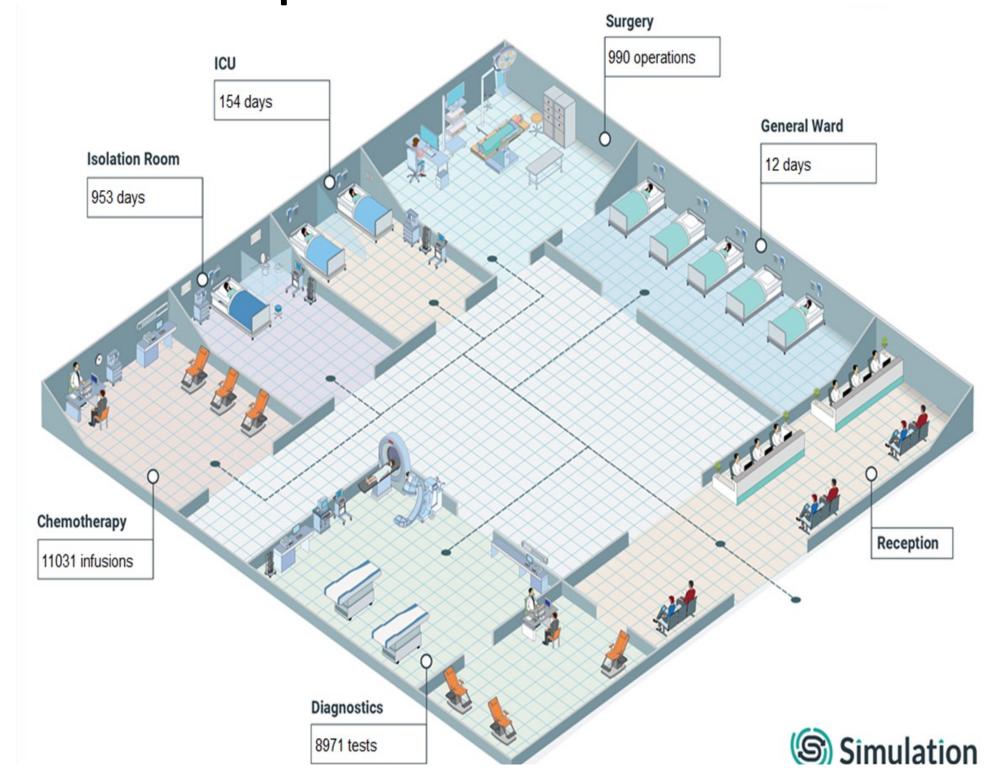


Table 1: Model's parameters

Parameter	Model's input
Risk of diagnostic errors per sample	1.67% ²
Risk of IV chemotherapy preparation errors (>10% deviation)	2.25%3
Risk of drug infusion administration errors	0.54%4
Risk of surgical site infection	4.67%5
Risk of urinary tract infection	2.1%6
Additional lenght of stay: surgical site infection	8 days ⁷
Additional lenght of stay: healthcare-associated urinary tract infection	4 days ⁸

REFERENCES

- 1. https://www.who.int/news-room/fact-sheets/detail/patient-safety
- 2. Carraro P, Zago T, Plebani M. Exploring the Initial Steps of the Testing Process: Frequency and Nature of Pre-Preanalytic Errors. Clinical Chemistry 58:3638–642 (2012)
- 3. Terkola R, Czekka M, Beribe J. Evaluaton of real-time data obtained from gravimateric preparation of antineoplastic agents shows mediction errors with possible therapeutic impact: Results of a large scale, multi centre, multinational, retrospective study. Journal of Clinical Pharmacy and Therapeutics. 2017:42(4);446-453
- 4. Kastrup M, Balzer F, Volk T, Spies C. Analysis of event logs from syringe pumps: a retrospective pilot study to assess possible effects of syringe pumps on safety in a university hospital critical care unit in Germany. Drug Safety 2012;35(7):563-74
- 5. Hannan et al. 2015. The Combined Impact of Surgical Team Education and Chlorhexidine 2% Alcohol on the Reduction of Surgical Site Infection following Cardiac Surgery. Surgical Infections. 2015;16(6): 799-805.
- 6. Thakker A, Briggs N, Maeda A, et al. Reducing the rate of post-surgical urinarytract infections in orthopaedic patients.BMJ Open Quality 2018;7:e000177
- 7. Jenks PJ, Laurent M, McQuarry S, Watkins R. Clinical and economic burden of surgical site infection (SSI) and predicted financial consequences of elimination of SSI from an English hospital. Journal of Hospital Infection. 2014 Jan;86(1):24-33
- 8. Mitchell BG, Ferguson JK, Anderson M, Sear J, Barnett A. Length of stay and mortality associated with healthcare-associated urinary tract infections: a multi-state model. J Hosp Infect. 2016 May;93(1):92-9