

PROTOCOL

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Risk factors for pressure injury development in critically ill patients in the intensive care unit: a systematic review protocol

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Abstract

Background: Pressure injuries (PIs) create a significant burden in the health care system. Up to 49% of critically ill patients develop PIs. Identifying and understanding potential risk factors is essential to the provision of effective targeted prevention strategies to mitigate risk. The objectives of this review are to identify patient-centred clinical factors that may be associated with PI development in the adult intensive care environment and to determine the effect size of the relationship between identified factors and PI development in this unique population.

Method/design: The review will follow the PRISMA reporting guidelines for systematic reviews. Electronic databases (Cochrane; PubMed/MEDLINE; CINAHL (EBSCOhost); Embase; Scopus; PsycINFO; Proquest; Networked Digital Library of Theses and Dissertations; Australian Digital Theses Program, Grey literature, Google scholar, and Clinical Trial Registries) will be systematically searched. A suite of search terms will identify articles that have examined the patient-centred risk factors for PI development in adult intensive care units. The search strategy will be designed to retrieve studies published since inception to 2016 in English language. Quality of the studies will be assessed by using an assessment framework designed to appraise quality in prognostic studies and methodological considerations in the analysis and publication of observational studies. Screening, study selection process, and data extraction will be undertaken by two independent reviewers. Disagreement will be resolved by discussion and, if required, a third independent reviewer. Clinical and methodological heterogeneity across studies will be assessed and, if possible, meta-analyses will be performed.

Discussion: The evidence synthesis arising from this review will identify person-centred risk factors that are associated with PI development among critically ill patients in intensive care. Findings from this review will demonstrate potential patient risk factors that may influence practice and research priorities to prevent PI development and improve the quality of care provided.

Systematic review registration: PROSPERO CRD42016037690

Keywords: Intensive/critical care, Pressure injury, Pressure ulcer, Predictors, Risk factors

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Protocol

Background

Pressure injuries (PIs) represent a common but potentially preventable condition seen most often in high-risk populations such as elderly persons, those with physical impairments, and the critically ill [1–3]. Pressure injuries, also known as pressure ulcers, are defined as a damage or lesion to the skin and underlying soft tissue, resulting from unrelieved pressure, shear, friction, moisture, or a combination of these, usually over a bony prominence or an anatomical area related to medical devices [4]. Pressure injuries differ in size and in the severity of affected tissue layers, with the latter ranging from skin erythema to muscle and underlying bone damage [4]. Moreover, PIs have significant negative impacts related to patients, society, and health systems (e.g. pain, increased infection rates, morbidity and mortality, increased length of stay in hospital, and raised financial costs) [5–7]. Preventing PI development to reduce the burden of PIs for patients and health care systems is considered a core aim of healthcare organizations.

Evidence suggests that PIs can be prevented with the implementation of PI prevention guidelines or care bundles, which target known risk factors associated with PI development [8–10]. Therefore, the identification and understanding of risk factors is required in order to provide appropriate prevention interventions and better utilize resources in practice.

Initial searching identified a systematic review published on the risk factors associated with PI development in adult hospitalized patients [11]. Findings identified that the three main factors which contributed to PI development were reduced mobility/activity, perfusion alterations (e.g. diabetes, vascular disease, poor circulation, blood pressure changes, smoking, oedema), and skin or PI status (e.g. a history of a previous PI occurrence). Further, the review concluded that PI occurrences cannot be explained by a single factor. This significant systematic review reported a total of 54 studies of which 13 studies were conducted in the intensive care environment [11]. However, the review reported only aggregate data from the 54 studies and no analyses were reported on specific subpopulations such as critically ill intensive care unit (ICU) patients. In comparison to general adult acute care in-patients, critically ill patients are more susceptible to risk factors for PI development as the majority of critically ill patients are ventilated and sedated and, therefore, unable to care for themselves, move, or change position. Further, the patient's critical illness may involve hemodynamic instability and oxygenation disorders, which potentially may complicate and accelerate the effects of prolonged immobility such as PI development. Extensive exposure

to pressure, from lying or sitting, on a specific part of the body renders patients at greater risk of skin breakdown. The highest PI in-hospital prevalence and incidence rates are noted in critically ill patients, thus confirming that, when compared to the total hospital population, the critically ill are at greatest risk for PI development [12]. It is estimated that up to 40% of patients develop PIs during their admission to ICUs [13].

It has been argued that PI development is a complex phenomenon that is enhanced by the presence of multiple, rather than single, risk factors in the individual [14]. Two literature reviews have been retrieved that have addressed the risk factors of PI development in the intensive care context [15, 16]. Findings from the first review [15] revealed that the potential risk factors for PI development were the same among hospitalized patients, despite critically ill patients having more than one factor. These findings were supported with a second review [16], which additionally explored different risk factors that accelerate the development of PI in ICU contexts and have an influence on the level and extent of tissue necrosis. These factors were identified and conceptualized in two categories: intrinsic (inherent factors of critical illness) and extrinsic (related to external forces) factors. A total of 28 factors were identified as the main risk factors for PI development in ICU settings in two or more studies. The intrinsic factors identified in two or more studies were older age, increased length of stay in the ICU, and history of cardiovascular disease. The extrinsic factors identified in two or more studies were the administration of norepinephrine and patient repositioning (turning). However, these were literature reviews [15, 16] in which no assessment of the methodological quality of the studies reviewed was undertaken thus making the interpretation of findings more susceptible to bias. In addition, Coleman and colleagues [11] suggest in their systematic review that multivariable statistical modelling, the identification of potential factors associated with PI development, should be independent to other risk factor variables included. An independent risk factor does not infer causality. Rather, it is a risk factor that 'retains statistical association with the outcome when other established risk factors are included in the statistical model' [17]. Thus, truly independent predictive intrinsic, patient-centred risk factors for critically ill patients in the ICU have yet to be conclusively established.

A systematic review is required to synthesize comprehensive current evidence in order to identify potential independent patient-centred clinical factors that are associated with PI development among critically ill patients in intensive care. Currently, after searching across PubMed, CINAHL, the Cochrane Library, the Joanna Briggs Institute Database of Systematic Reviews

and Implementation Reports, and Google Scholar, no current systematic review addressing risk factors of PIs in this context has been identified. Identifying the potential independent person or patient-centred factors will facilitate decision-makers such as researchers, clinicians, and policy-makers to provide appropriate interventions to alleviate the pressing problem of PI development.

Objectives

The objectives of this review are to (1) identify patient-centred clinical factors that may be associated with PI development in adult intensive care and (2) determine the effect size of the association between identified factors and PI development. The systematic review will synthesize existing knowledge and make recommendations for future research.

Methods/design

This review will replicate the methodology used by Coleman and colleagues who identified PI risk factors in adult hospitalized patients [11]. However, this review will focus on adult intensive care patients.

This protocol adheres to the Preferred Reporting Items for Systematic Reviews and Meta-Analyses Protocol (PRISMA-P) statement [18], and this is provided as an Additional file 1. This review is registered with PROSPERO—registration number CRD42016037690.

Criteria for considering studies for this review

Types of studies

Eligible quantitative studies will be included in this review. The systematic review will consist of studies using either epidemiological designs such as retrospective or prospective cohort studies, or experimental designs such as randomized controlled trials or quasi-experimental trials, where analyses for potential patient-centred clinical factors of PI development in intensive care have been reported. Cross-sectional, qualitative, case studies and abstract-only reports, for which full text is not available, will be excluded from this review.

Type of clinical settings

Studies conducted in intensive/critical care settings will be included.

Type of participants

This review will consider studies that included adult critically ill patients aged 18 or above, who have not been diagnosed with a PI that developed before the patient was admitted to the ICU.

Type of exposure

No restriction will be imposed for risk factors that are significantly associated with PI development in intensive

care. Patient risk factors are defined as those factors related to individual patient descriptors including age, gender, diagnosis, comorbidities, body mass index, severity of illness, treatment modalities related to critical illness, complications of treatment, clinical pathology, length of ICU stay, and ICU outcome (discharge, transfer, or death) [16]. In addition, studies that examine the factors of PI development for adults across hospitals but have separate statistical analyses reported for PI development in adult intensive care will also be included.

Type of outcome measure

This review will consider studies that include all stages of PI or equivalent as the outcome measure. According to the National Pressure Ulcer Advisory Panel, European Pressure Ulcer Advisory Panel, and Pan Pacific Pressure Injury Alliance Clinical practice guidelines, the PI stages reflect the level of skin tissue damage [4, 19].

Search methods of identification of studies

The search strategy aims to find both published and unpublished studies exploring the risk factors that are associated with PI development in intensive care. The literature will be searched using Medical Subject Headings (MeSH) and combinations of key terms. The search is limited to English language publications with no restriction on the year of publication. The electronic databases searches will include the Cochrane database (1991 (established) to present); PubMed/MEDLINE (1969–present); CINAHL (EBSCOhost) (1989 to present); Embase (1966–present); Scopus (1975–present); PsycINFO (2009 to present); Proquest (1959 to present); and Google scholar. In addition, the reference lists of articles meeting the criteria for this review will be searched for further relevant studies. The search for unpublished studies will include Networked Digital Library of Theses and Dissertations; Australian Digital Theses Program; Open Grey (greynet.org); science.gov; clinicaltrials.gov; International Standards for Randomised Controlled Trials (IRISCTN) Registry; and the Australian and New Zealand Clinical Trials Registry (ANZCTR).

The search syntax will be designed and guided by the Cochrane Handbook [20] to retrieve the relevant results. Search strings will be focused on the terms pressure ulcer, pressure injury, risk factors, predictors, adult intensive care, and their synonyms.

Search terms

Medical Subject Headings (MeSH) and free text word terms will be developed specifically for each database and will be subsequently pilot tested. For example, for MeSH, terms include (pressure ulcer* OR pressure injur* OR bed sore* OR decubitus ulcer* OR pressure sore*)

AND (risk factor* OR predictor* OR predictor factor*)
AND (intensive care* OR ICU OR critical care) AND
(adult OR age ≥ 18).

Data collection and analysis

Selection of studies

EndNote (Thomson Reuters) software will be used to manage records for retrieval of articles and inclusion/exclusion decisions. Two independent reviewers (both authors) will perform the initial screening of titles and abstracts to identify eligible studies (Additional file 2). Full copies of all studies that meet the criteria as outlined below will be obtained for further assessment. The inclusion criteria are (i) length of patient follow-up that is at least 48 h (this has been identified as an accepted time frame for PI development) [21], (ii) multivariate analyses undertaken to identify factors affecting PI outcome, and (iii) not specific to geographical areas. The exclusion criteria are (i) observational studies that will be excluded if >20% of the study sample were excluded from analysis for reasons including withdrawal, death, loss to follow-up, and missing records and (ii) control trial studies will be excluded unless all of the following minimum criteria are applied: parts of the study were prospectively designed and intention to treat analyses are reported [20].

Data extraction and management

Data will be extracted independently by the two reviewers using the data extraction form. This form will include the study characteristics (for example, study population, recruitment type used, PI definition, and analysis method) and risk factors investigated in the multivariate models, including those found to be significant. Non-significant factors through using a stepwise regression will be included if it reported as independent correlated variables to PI development. Disagreement will be reconciled by discussion leading to mutual agreement or, if this is not possible, consultation with a third independent reviewer. In addition, authors will be contacted if papers are unobtainable or the methods reported need clarification.

Assessment of risk of bias in included studies

Both authors will undertake the risk of bias assessment of the included studies independently, as guided by the assessment framework for assessing quality in prognostic studies and methodological considerations in the analysis and publication of observational studies [11] as described below.

Quality assessment

The methodological quality of eligible studies will be assessed and critically appraised by two independent

reviewers (both authors) using the assessment framework that is designed to appraise quality in prognostic studies and methodological considerations in the analysis and publication of observational studies [11]. Any disagreements that arise between the reviewers will be resolved through discussion or, if agreement is not reached, consultation with a third, independent reviewer.

The quality assessment tool will rate the relevant methodological parameters of studies across seven areas [11]: selection bias (selection of target population), confounders (whether confounders were controlled using appropriate adjustment), data collection methods (validity and reliability by using a clear definition or description of the risk factor), reporting of continuous variables (cut point), measurement of the range of potential risk factors, withdrawals and dropouts (dropout rates and completion of study rates), and no selective reporting of results.

In addition, this tool has four considerations to appraise domains quality:

1. Is there a sufficient number of events (rule of thumb, 10 events per risk factor)?
2. Is there sufficient presentation of data to assess the adequacy of method and analysis?
3. Is the strategy for model building (i.e. inclusion of variables) appropriate and based upon a conceptual framework?
4. Is the selected model adequate for the design?

Each criteria will be assessed as being met (yes/no/partial/unsure). Consequently, assessment of domain criteria will assist the determination of overall study quality [11].

Classification of study quality

The quality of the studies will be classified according to the assessment framework for assessing quality and methodological consideration in the analysis, meta-analysis, and publication of observational studies [11] as high, moderate, low, and very low quality based on the following criteria:

High: yes for all four criteria

Moderate: yes for criteria 1 and at least 2 for other criteria

Low: no for criteria 1 and no or partial for 2 other criteria

Very low: no for criteria 1 and no or partial for all 3 other criteria

Unit/scale of analysis issues

The unit of analysis will be based on the individual patient.

Assessment of heterogeneity

The studies will be assessed by considering variability in the clinical and methodological heterogeneity to ensure sufficient homogeneity and rigour of the studies to conduct the meta-analysis procedure. Clinical heterogeneity of the studies will be assessed by considering their population, risk factor description, and outcome. Methodological heterogeneity will be assessed using the data extracted on study design, procedure, and analysis reported. In the absence of clinical and methodological heterogeneity statistical measures, I^2 and H_M^2 statistic measures [22, 23] will also be used to determine the level of consistency between studies. I^2 and H_M^2 values will be used to measure the impact of heterogeneity between and within the study variance. An $I^2 \geq 50\%$, which corresponds to $H_M^2 > 0$, will be considered a significant level of heterogeneity.

Assessment of publication bias

The risk of publication bias will be minimized by comprehensively searching databases and obtaining data from unpublished work to reduce the risk of reporting bias [24, 25]. Moreover, if more than 10 studies are reviewed that fit the criteria, a funnel plot will be used to determine the relationship between study size and effect power of cohort clinical trial studies by signs of asymmetry.

Data synthesis

A meta-analysis may not be able to be conducted due to the clinical or statistical heterogeneity of the eligible studies. However, if included studies are sufficiently homogenous and rigorous, a meta-analysis using a random-effects model will be conducted. A relative risk will be the appropriate indicator of effect for potential correlates in this review with a 95% confidence interval (CI) calculated for binary outcomes (PI development). However, if there is considerable clinical and design heterogeneity in the included studies, the findings will be presented in narrative form including tables to aid in data presentation where appropriate.

Summary of findings

Patient-centred clinical risk factors for PI development will be categorized by domains and subdomains according to Coleman and colleagues [11]. Coding contributory factors into different domains will be conducted by the two authors independently. The identified domain and sub-domain factor will be summarized and presented in tables. Each sub-domain factor will be summarized including a number of significant studies using a multi-variable analysis and the total number and quality of the studies entering the variable. Disagreements will be resolved by discussion to achieve consensus or, if this is not possible, by a third independent reviewer.

Additional files

Additional file 1: PRISMA-P (Preferred Reporting Items for Systematic Review and Meta-Analysis Protocols) 2015 checklist. (PDF 278 kb)

Additional file 2: Figure S1. Literature searches, screening and selection of articles for inclusion. (PDF 102 kb)

Abbreviations

CI: Confidence interval; ICU: Intensive care unit; PI: Pressure injury

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Availability of data and materials

The dataset(s) supporting the conclusions of this systematic review will be included within the final review article (and its additional files).

Authors' contributions

FC is the guarantor of the review and the principal investigator. FC and NT contributed equally to designing the systematic review and drafting the protocol. NT developed the search strategy. Both authors have read and approved the final manuscript, agreed to be accountable for all the aspects of the work, and acknowledged that all those entitled to authorship are listed as authors.

Authors' information

Dr Fiona Coyer holds a joint appointment as Professor of Nursing with the School of Nursing, Queensland University of Technology and Metro-North Hospital Health Services. She is based at Intensive Care Services Department, Royal Brisbane and Women's Hospital. Fiona has extensive experience as a leader in academic and research programs in intensive care nursing, with research interests including nursing care of the critically ill patient and particularly skin integrity in critically ill patients. Nahla Tayyib is a Lecturer, School of Nursing, Umm Al-Qura University, Saudi Arabia. Nahla completed her doctoral degree at Queensland University of Technology in 2016. Nahla's doctoral work explores the use of an interventional pressure ulcer bundle to reduce pressure ulcer incidence in intensive care patients in Saudi Arabia. Thus far, she has published seven articles from her doctoral work.

Competing interests

NT declares no competing interests. FC declares educational consultancy interests with 3M and Teleflex companies.

Consent for publication

Not applicable. This protocol manuscript does not, nor will the final systematic review, contain any individual person's data in any form.

Ethics approval and consent to participate

Not applicable. This review will retrieve only published material.

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