The Association Between Pressure Ulcer Development And Patient Comorbidities In Varied Care Settings: A Review Of The Literature

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Abstract

Pressure ulcers can be described as localized injury to the skin which can develop in the patients placed in various types of care setting. The comorbidities, or underlying diseases of the patients, are thought to be strong risk factors for the development of pressure ulcers. The purpose of this paper is to review the previous literature investigating the association between patient comorbidities and pressure ulcer development. The author classified the literature into five categories by the type of care setting, namely, the nursing home, hospital, perioperative, rehabilitation, and home settings, and summarized the study results for each. The author found a large dispersion in the results according to design and sample size, even within the same settings. Nonetheless, half or more of the studies found that diabetes mellitus had a close association with pressure ulcer development in all of the settings while cardiac, renal, and respiratory diseases were also found to be associated with many of the settings. The author surmises that recognition of such comorbidities could serve as an important step towards improving pressure ulcer prevention.

Keywords: Cardiac Disease; Comorbidity; Diabetes Mellitus; Pressure Ulcer; Renal Disease; Respiratory Disease; Underlying Disease

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Introduction

A pressure ulcer (PU) is defined as “localized injury to the skin and/ or underlying tissue, usually over a bony prominence, as a result of pressure or pressure in combination with shear” [1]. In addition to such mechanical forces, however, patient comorbidity or an underlying disease may also contribute to PU development. Several clinical guidelines for PU prevention refer to comorbidity as a risk factor for PU development, citing, for example, diabetes mellitus [1-4], circulatory disease [1], spinal cord injury [2,4], hip fracture [4], cerebrovascular disease [4], etc. In addition, risk assessment scales for PU prevention such as CBO [5], Spinal Cord Injury Pressure Ulcer Scale (SCIPUS) [6], SCIPUS-A [7] and Waterloo [8] include factors related to patient comorbidities such as diabetes mellitus, and pulmonary and cardiac diseases.

PUs can develop in patients placed in any types of care setting. The aim of this paper is to review the previous English-language literature investigating the association between patient comorbidities and PU development. The author classified the literature into five categories according to the type of care setting, namely the nursing home, hospital, perioperative, rehabilitation, and home settings, and further examined the kind of PU-related comorbidity in each of these settings.

Nursing Home Setting

Nursing homes constitute representative long-term care settings for the immobile or inactive elderly population. The previous literature dealing with the association between comorbidities and PU development in nursing homes is summarized in Table 1. To the author’s knowledge, seven studies have been published to date on this subject [9-15], three cohort studies [10,13,14] and four cross-sectional studies [9,11,12,15], all of which utilized large databases at multiple facilities. It should be noted that six of the seven studies cited [9,11,15] concluded that diabetes mellitus was associated with PU development. Moreover, in some studies, hip fracture [13], Parkinson’s disease [9], and peripheral vascular disease were also listed as risk factors, while other studies showed inverse results [9,11,12,14].

Hospital Setting

Inpatient hospital care ranges from acute or intensive care to chronic or geriatric care. Accordingly, hospitalized patients may present many types of comorbidities in association with PU development. As shown in Table 2, we found three cohort studies [18-20], three cross-sectional studies [16,21,22], one cross-sectional/cohort study [17], and one case-control study [23], or a total of eight studies dealing with PU-related comorbidities in the hospital setting. Three of these studies [21-23] utilized a large inpatient database derived from multiple facilities. On the other hand, six [16-20] utilized relatively small samples from single hospitals including two teaching hospitals [16,19], two chronic care hospitals [17,18], and one medical ICU in a public hospital [20]. In four of the studies [17,21-23], diabetes mellitus was described...
### Table 1. Nursing home setting

| Study               | Study design | Participants | Study setting and database | Country          | Comorbidities in associated with PU development | Comorbidities not associated with PU development |
|---------------------|--------------|--------------|-----------------------------|------------------|------------------------------------------------|------------------------------------------------|
| Spector, 1994 [9]   | Cross-sectional | 2803         | 699 nursing homes in NMES IPC 1987 | United States    | Diabetes mellitus, Parkinson’s disease          | Hip fracture, stroke                             |
| Brandies et al, 1994 [10] | Cohort       | 4232         | 78 nursing homes belonging to National Health Corp | United States    | None                                            | Diabetes mellitus                                |
| Brandies et al, 1995 [11] | Cross-sectional | 2011         | 270 nursing homes in MDS 1990 | United States    | Diabetes mellitus                               | Alzheimer’s disease, hip fracture, peripheral vascular disease |
| Spector and Förtin-sky, 1998 [12] | Cross-sectional | 15121       | 843 nursing homes in MDS+ 1994 | United States    | Diabetes mellitus                               | Parkinson’s disease, hip fracture, stroke         |
| Berkowitz et al, 2001 [13], [14] | Cohort       | 14607        | 109 nursing homes in MDS 1997 | United States    | Diabetes mellitus, hip fracture, peripheral vascular disease | None                                           |
| Berkowitz et al, 2001 [15] | Cohort       | 13457        | 108 nursing homes in MDS 1998 | United States    | Diabetes mellitus                               | End-stage disease, hip fracture, peripheral vascular disease |
| Casimro et al, 2002 [16] | Cross-sectional | 827          | 50 geriatric facilities     | United Kingdom   | Diabetes mellitus                               | None                                           |

NMES IPC (Institutional Population Component of the National Medical Expenditure Survey): A nationally representative sample of residents in nursing homes.

MDS+ (Minimum Data Set Plus): The expanded version of a national resident assessment instrument specifically for Multistate Nursing Home Case Mix and Quality Demonstration Project.

MDS (Minimum Data Set): A valuable tool for assessing the quality of nursing home care containing clinical information describing the health status of all nursing home residents.

### Table 2. Hospital settings

| Study               | Study design | Participants | Study setting and database | Country          | Comorbidities in associated with PU development | Comorbidities not associated with PU development |
|---------------------|--------------|--------------|-----------------------------|------------------|------------------------------------------------|------------------------------------------------|
| Allman et al, 1996 [16] | Cross-sectional | 634          | Single teaching hospital    | United States    | Fracture                                       | Anemia, dementia, diabetes mellitus, heart failure, malignancy, pneumonia, renal failure, spinal cord injury, stroke, urinary tract infection |
| Berkowitz and Wilking, 1989 [17] | Cross-sectional and cohort | 301          | Single chronic care hospital | United States    | Diabetes mellitus, cerebrovascular accident     | Dementia, diabetes mellitus, heart failure, fracture, malignancy |
| Bianchetti et al, 1993 [18] | Cohort       | 148          | Single psychogeriatric hospital | Italy            | None                                          | Diabetes mellitus, heart failure, peripheral vascular disease, pneumonia, stroke |
| Allman et al, 1995 [19] | Cohort       | 266          | Single teaching hospital    | United States    | None                                          | Chronic heart failure, diabetes mellitus, fracture, paraplegia |
| Jiricka et al, 1995 [20] | Cohort       | 85           | Medical / surgical ICU in a single public hospital | United States    | None                                          | Diabetes mellitus                                 |
| Scott et al, 2006 [21] | Cross-sectional | 100000       | CHARS 1987-2000              | United States    | Diabetes mellitus, spinal cord injury           | Injury, infection                                 |
| Fogerty, 2008 [22] | Cross-sectional | 94758        | NIS 2003                     | United States    | Anemia, cerebrovascular disease, chronic heart failure, diabetes mellitus, osteomyelitis, pneumonia, renal failure, respiratory failure, sepsis, urinary tract infection | Not described                                    |
| Leder et al, 2012 [23] | Case-control | 51842        | MPSMS 2006-2007              | United States    | Cerebrovascular disease, chronic heart failure, chronic obstructive pulmonary disease, diabetes mellitus | None                                           |

CHARS (Comprehensive Hospital Abstract Reporting System): The annual data from all hospital admissions in the Washington state.

NIS (Nationwide Inpatient Sample): A national sample of inpatient discharge data.

MPSMS (Medical Patient Safety Monitoring System): A nationwide surveillance system within the hospitalized fee-for-service Medicare population.
as a risk factor for PU development, although the remaining four studies denied this association [16,18-20]. Additionally, anemia [20], cerebrovascular disease [17,22,23], chronic heart failure [22,23], chronic obstructive pulmonary disease [23], fracture [16], osteomyelitis [22], pneumonia [22], renal failure [22], respiratory failure [22], sepsis [22], spinal cord injury [21], and urinary tract infection [22] were mentioned as risk factors for PU development.

**Perioperative Setting**

Patients who undergo a lengthy operation are potentially at high risk for PU development because repositioning may not be possible for several hours during the intraoperative and postoperative periods. The author has found 10 studies to date concerning PU-related comorbidities in the perioperative setting [24-33], including nine cohort studies [24-31,33] and one meta-analysis [32](Table 3). Eight of these studies were based on data from a

| Study                          | Participants | Study setting and database | Country                  | Comorbidities in associated with PU development | Comorbidities not associated with PU development |
|-------------------------------|--------------|----------------------------|--------------------------|-------------------------------------------------|--------------------------------------------------|
| Papantonio et al, 1994 [24]   | Cohort 136   | Single teaching hospital   | United States            | Diabetes mellitus, respiratory disease          | Hypertension, peripheral vascular disease, renal disease |
| Lewicki et al, 1997 [25]      | Cohort 337   | Single academic medical center | United States          | Diabetes mellitus                               | None                                             |
| Schultz et al, 1999 [26]      | Cohort 413   | Single tertiary care center | United States            | Diabetes mellitus                               | None                                             |
| Pokorny et al, 2003 [27]      | Cohort 351   | Single medical center      | United States            | Heart failure                                    | None                                             |
| Frankel et al, 2007 [28]      | Cohort 820   | Surgical ICU in a single teaching hospital | United States        | Diabetes mellitus, spinal cord injury           | None                                             |
| Haleem et al, 2008 [29]       | Cohort 4546  | Single hospital            | United Kingdom           | Diabetes mellitus                               | Malignant disease, Rheumatoid arthritis          |
| Lindholm et al, 2008 [30]     | Cohort 635   | Accident and Emergency Departments in six countries | Sweden, Finland, United Kingdom, Spain, Italy, Portugal | Cardiovascular disease, diabetes mellitus, pulmonary disease | Gastrointestinal disease, malignancy, urological disease |
| Slowikowski and Funk, 2010 [31] | Cohort 369   | Surgical ICU in a single hospital | United States          | Diabetes mellitus                               | Renal disease, vascular disease                  |
| Liu et al, 2012 [32]          | Meta-analysis | 2453 Six observational studies | United States (5 studies) and Belgium (one study) | Diabetes mellitus                               | None                                             |
| O’ Brien et al, 2013 [33]     | Cohort 2695  | Surgical ICUs in a single teaching hospital | United States          | Congestive heart failure, renal failure         | Hypertension, liver disease                      |

**Table 3. Perioperative settings**

| Study                          | Study design | Participants | Study setting and database | Country                  | Comorbidities in associated with PU development | Comorbidities not associated with PU development |
|-------------------------------|--------------|--------------|----------------------------|--------------------------|-------------------------------------------------|--------------------------------------------------|
| Salzberg et al, 1996 [6]      | Case-control | 219          | Single spinal cord injury unit in veterans affair medical center | United States            | Cardio disease, diabetes mellitus, pulmonary disease, renal disease | Urinary tract infection, sepsis                     |
| Cakmak et al, 2009 [34]       | Case-control | 64           | Single physical therapy and rehabilitation hospital | Turkey                   | None                                            | Diabetes mellitus, hypertension                    |
| Verschueren et al, 2011 [35]  | Cohort       | 193          | Eight rehabilitation hospitals with SCI units | The Netherlands          | Pulmonary disease                                | Cardiovascular disease, spine fracture, urinary tract infection |
| Wang et al, 2013 [36]         | Case-control | 5804         | UDSMR 2009-2011            | United States            | Dementia, diabetes mellitus, peripheral vascular disease | Amputation, arthritis, cardiac disorders, pulmonary disorders, spinal cord injury, stroke, |

UDSMR (Uniform Data System for Medical Rehabilitation): A large nongovernment registry for standardized medical rehabilitation information.
single hospital [24-29,31,33]. Some studies targeted patients who received cardiac surgery [24,25,27] or hip fracture surgery [29,30]. In seven of nine cohort studies [24-26,28-32], diabetes mellitus was identified as a risk factor for PU development. This result was strongly supported by one recent meta-analysis [32] involving six observational studies with a total 2453 patients; when compared to patients with normal glucose tolerance, patients with diabetes mellitus were more likely to develop PUs (odds ratio = 2.15, 95% confidence interval: 1.62-2.84). In addition, other comorbidities including cardiac disease [27,30,33], renal failure [33], respiratory disease [24,30,33], and spinal cord injury [28] were also described as risk factors.

Rehabilitation Setting

Because of restrictions on mobility, patients undergoing rehabilitation are thought to be at extremely high risk for PU development. Table 4 lists four studies, including three case-control studies [6,34,36] and one cohort study [35], investigating PU-related comorbidities at rehabilitation facilities [6,35-37]. Among these, three studies utilized data obtained from a single hospital [6,34] or small number of hospitals [35], while one utilized a large inpatient database [36]. Two studies targeting patients admitted to spinal cord injury units [6,35]. Diabetes mellitus [6,36] and pulmonary disease [6,35] were each found to be risk factors for pressure ulcer development in two studies. Additionally, cardiac disease [6], dementia [36], peripheral vascular disease [36], and renal disease [6] were also listed as risk factors.

Discussion And Summary

PU’s can develop in various care settings. In this paper, we categorized the relevant literature published to date on the subject according to five types of care setting. Importantly, patient background, study design, sample size, and risk factor variables varied widely in each study. Moreover, a large dispersion of the results was seen between the studies, even in the same setting. For this reason the author concluded that the outcomes from each study mentioned above are in some instances not amenable to comparison. Nonetheless, among the numerous comorbidities referred to in these studies, diabetes mellitus seemed to present the strongest association, with half or more of the studies in all of the care settings and one meta-analysis pointing to this condition as a risk factor for PU development. With respect to other comorbidities, cardiac, renal, and respiratory disorders were commonly found to be risk factors in four of the five care settings: hospital, perioperative, rehabilitation, and home. However, the author could not find a clear difference in the kinds of PU-related comorbidities between settings.

In summary, the findings of previous studies investigating risk factors for PU development underscore the importance of recognizing patient comorbidities in order to prevent PUs. Despite the large dispersion seen between the study results, diabetes mellitus, cardiac disease, renal disease, and respiratory disease were found to be risk factors across a variety of settings.

Table 5. Home settings

| Study                | Study design | Participants | Study setting and database | Country            | Comorbidities in associated with PU development | Comorbidities not associated with PU development |
|----------------------|-------------|--------------|----------------------------|--------------------|-----------------------------------------------|-----------------------------------------------|
| Margolis et al, 2003 | Cohort      | 75,168       | GPRD 1988-1996             | United Kingdom     | Alzheimer's disease, diabetes mellitus, chronic heart failure, chronic obstructive pulmonary disease, cerebral vascular accident, deep venous thrombosis, hip fracture, limb paralysis, lower limb edema, malignancy, osteoporosis, Parkinson's disease, rheumatoid arthritis, urinary tract infection | Angina, hypertension, pneumonia |

GPRD (General Practice Research Database): a large outpatient record database from the United Kingdom.

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