

LETTER TO THE EDITOR

Pressure ulcer stage IV caused by cervical collar in patients with multiple trauma in intensive care unit

Dear Editors,

Cervical collars are necessary to immobilise the cervical spine of patients with multiple trauma (1,2). This device remains in situ until the radiographic cervical spine clearance is completed (2). A 29-year-old male with multiple trauma and subdural haemorrhage admitted to the intensive care unit (ICU) was selected for this case study. The patient had been taken to hospital by emergency medical service (EMS) following an accident. He was fitted with a Philadelphia rigid collar at the site of the accident by EMS technicians. After admission to the hospital and assessment by emergency physicians, neurosurgeons and orthopaedic surgeons, he was transferred to the ICU. Five days later, the spinal clearance process was completed. On day 6 after ICU admission, physicians ordered removal of the cervical collar. After removal of the collar by critical care nurses, two stage IV pressure ulcers were observed. One ulcer developed in his occiput and the other on his chin (Figures 1 and 2).



Figure 1 Ulcer in the occiput.



Figure 2 Ulcer on the chin.

Physicians assessed the ulcers and confirmed that they were caused by the pressure of the cervical collar.

Risk of pressure ulcer development increased with the use of medical devices (3–6). This risk is higher among critically ill patients because this group of patients requires more medical devices for monitoring and therapeutic purposes (4,7). Coyer *et al.* reported that the prevalence of medical device-related pressure ulcer was 3.1% in ICU patients (8). The present case shows that skin may be affected by cervical collar, and as a result pressure ulcer may develop. Ackland *et al.* determined four risk factors for pressure ulcer development during cervical collar use, which include ICU admission, mechanical ventilation, the necessity for cervical magnetic resonance imaging and the time to cervical spine clearance and collar removal (2). For the prevention of medical device-related pressure ulcers, the National Pressure Ulcer Advisory Panel (NPUAP) has identified seven recommendations that include: (i) choose the correct size of medical devices to fit the individual; (ii) cushion and protect the skin with dressings

in high-risk areas; (iii) remove or move the device daily to assess skin under device; (iv) avoid placement of device over area of prior or existing pressure ulceration; (v) educate staff on correct use of devices and prevention of skin breakdown; (vi) be aware of oedema under device(s) and potential for skin breakdown; and (vii) confirm that devices are not placed directly under an individual who is bedridden or immobile (9). These recommendations should be considered by health care members, especially nurses who care for critically ill patients.

Abbas Abdoli Tafti, MD¹, Sanaz Sajadi, MD² & Hossein Rafiei, MSc³

¹Orthopedic Department
Kashani Hospital, Shahrekord University of Medical Sciences
Shahrekord, Iran

²Emergency Department
Kashani Hospital, Shahrekord University of Medical Sciences
Shahrekord, Iran

³Department of Intensive and Critical Care
School of Nursing and Midwifery, Shahrekord University of Medical Sciences
Shahrekord, Iran
hosseinrafiei21@yahoo.com

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