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Spinal cord injury associated with cervical spinal canal stenosis: Outcomes and prognostic factors

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ABSTRACT

Objectives: To specify outcomes and identify prognostic factors of neurologic and functional recovery in patients with an acute traumatic spinal cord injury (SCI) associated with cervical spinal canal stenosis (SCS), without spinal instability.

Methods: A retrospective study was conducted using data from a Regional Department for SCI rehabilitation in France. A description of the population characteristics, clinical data and neurological and functional outcomes of all patients treated for acute SCI due to cervical trauma associated with SCS was performed. A statistical analysis provided insights into the prognostic factors associated with the outcomes.

Results: Sixty-three patients (mean age 60.1 years) were hospitalized for traumatic SCI with SCS and without instability between January 2000 and December 2012. Falls were the most frequent cause of trauma (77.8%). At admission, most patients had an American Spinal Injury Association Impairment Scale (AIS) grade of C (43.3%) or D (41.7%) and the most frequent neurological levels of injury were C4 (35.7%) and C5 (28.6%). Clinical syndromes were frequently identified (78.6%), with the most frequent being the Brown-Sequard plus syndrome (BSPS) (30.9%), followed by central cord syndrome (CCS, 23.8%). Almost 80% of survivors returned to the community, 60% were able to walk and 75% recovered complete voluntary control of bladder function. Identified prognostic factors of favourable functional outcomes were higher AIS at admission, age under 60 years and presence of BSPS or CCS.

Conclusion: Traumatic SCI, associated with SCS results mostly in incomplete injuries, can cause various syndromes and is associated with favourable functional outcomes.

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1. Introduction

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Cervical spondylosis affects up to 50% of persons over 40 years of age [1]. It often causes spinal canal stenosis (SCS), leading to neck pain, cervical radiculopathy and cervical myelopathy [1]. Subjects with cervical spondylosis have a higher risk of spinal cord lesions since the cord cannot move freely within the spinal canal, thus mild cervical trauma can cause devastating spinal cord

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http://dx.doi.org/10.1016/j.rehab.2017.09.003 1877-0657/© 2017 Elsevier Masson SAS. All rights reserved. injuries [2]. The anatomical features of such injuries usually 20 include haematoma, oedema and myelomalacia, mostly affecting 21 22 the central part of the spinal cord [3]. Schneider and Cherry provided the classical description of the symptoms that occur 23 following cervical spinal cord injury (SCI) on an already 24 compressed spinal cord [3]: weakness mostly affecting the upper 25 limbs and various impairments of the lower limbs, including loss of 26 bladder and bowel function. This clinical condition is today 27 referred to as 'traumatic central cord syndrome' (CCS) and criteria 28 for its diagnosis have recently been precisely defined [4]. Despite 29 the generally older age of patients with CCS [5], this syndrome is 30 associated with more favourable functional outcomes than other 31 SCI-related syndromes [5–7]. 32

CCS and traumatic SCI with CCS are frequently confused, 33 probably because of the initial statement by Schneider and Cherry 34

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Abbreviations: ACS, anterior cord syndrome; AIS, ASIA Impairment Scale; ASIA, American Spinal Injury Association; BSPS, Brown-Sequard plus syndrome; CCS, central cord syndrome; MRI, magnetic resonance imaging; NLI, neurological level of injury; SCI, spinal cord injury; SCS, spinal canal stenosis; TDM, tomodensitometry.

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35 that CCS affects older patients with cervical spondylosis [3]. How-36 ever, various situations may arise in subjects with acute SCI and 37 concomitant cervical spondylosis [8]. Although this condition was 38 first described more than 50 years ago, data relating to 39 sociodemographic characteristics, clinical signs and symptoms 40 and outcomes are lacking [8,9]. Such knowledge would help to 41 determine appropriate management for this condition. One 42 retrospective study investigated the clinical features and prognos-43 tic factors of CCS, however this was in a subset of patients who underwent surgical decompression [10]. 44

45 The aims of this study were therefore:

47 • to describe the clinical features at the time of admission in 48 patients with acute traumatic SCI associated with cervical CCS, 49 without spinal instability:

50 to identify outcomes and prognostic factors of neurologic and 51 functional recovery in these patients.

2. Methods 52

2.1. Patients 53

54 A retrospective analysis of all patients referred to our SCI centre 55 was conducted. This centre includes an acute care Neurotrauma 56 unit and a Physical and Rehabilitation Medicine (PRM) depart-57 ment. All cases, admitted between the 1st of January 2000 and the 58 31st of December 2012, were reviewed. The inclusion criteria 59 were:

- trauma to the cervical spinal cord with neurological impairment 61 62 in the acute phase;
- 63 • no unstable spinal lesion, verified by computed tomography (CT) 64 and magnetic resonance images (MRI) using the criteria defined 65 by White and Panjabi [11] (all patients systematically under-66 went CT and MRI). Any patients with fracture, dislocation, disc 67 lesion or vertebral sprain that was at risk of displacement or 68 compromising the spinal canal were excluded. This choice is in 69 accordance with many previous studies [9,10,12]: patients with 70 radiological features of spinal instability were excluded because 71 in many centres, including ours, these patients undergo 72 immediate surgery to treat the cause of instability and reduce 73 cord compression. Besides, it is not possible to differentiate 74 between neurological involvement due to spinal stenosis and 75 cord oedema and that due to acute cord compression by bony or 76 disc elements:
- 77 presence of cervical spinal canal stenosis. Cervical spinal canal 78 stenosis was confirmed by the 'Pavlov' ratio on CT images [13]. A 79 ratio < 0.8 at any level between C3 and C7 was considered as 80 stenosis, with or without evidence of spondylosis.

81 All patients who met these three criteria were considered, 82 whether they underwent surgery during the initial stages or not 83 and regardless of corticosteroid administration.

84 2.2. Data collection

85 All applicable institutional and governmental regulations 86 concerning the ethical use of human volunteers were followed 87 during the course of this research. This study conformed to the 88 Helsinki Declaration of 1975, revised in 2013 and was approved by 89 our institutional ethics board. Data were collected retrospectively, 90 during the year 2015. The two-year delay between the end of the 91 inclusion period and the beginning of data collection was 92 necessary in order to obtain discharge data for all patients.

93 Two time points were considered:

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- admission to the acute care unit;
- discharge from the PRM department.

97 The endpoint evaluation was performed at discharge since data at other time points (for instance, 6 months post-injury) were not 98 available for all patients. Moreover, it is common practice to report 99 data at discharge in epidemiological studies of patients with SCI 100 [6,14,15]. 101 102

All patients underwent standard neurological assessment. The neurological examination was performed according to the International Standards for Neurological Classification of Spinal Cord Injury [16]. The American Spinal Injury Association (ASIA) Impairment Scale (AIS), the neurological level of injury (NLI) and the presence of a clinical syndrome were evaluated. Clinical syndromes included central cord syndrome (CCS), Brown-Sequard plus syndrome (BSPS), transverse syndrome, anterior cord syndrome (ACS) and posterior cord syndrome. CCS was defined according to the criteria by the EM-SCI study group [4]: disproportionately more motor impairment of the upper than lower extremities, with a difference of at least 10 motor score points between the upper and lower extremities. BSPS was defined according to the description by Roth et al. [17]: asymmetric paresis with more marked hypoalgesia on the less paretic side. Transverse syndrome was considered as complete SCI. ACS was defined as loss of motor function and pain/temperature sensation at and below the injury level, with preservation of light touch and joint position sense [16]. PCS was defined as isolated loss of proprioception and vibration sense below the level of injury [6].

Data regarding function, including ambulatory status and lower 122 urinary tract function on discharge and discharge mode were 123 collected from the medical files. Level of ambulation was defined as 124 the mode of locomotion for distances over 10 m (without gait aid, 125 with gait aid, manual or electric wheelchair). Lower urinary tract 126 function included bladder-emptying method (voluntary control, 127 self-intermittent catheterization, hetero-catheterization, indwell-128 129 ing catheter or reflex micturition). Discharge mode included return 130 home with or without any personal assistance for activities of daily living or hospital/institution care. 131

2.3. Data analysis and prognostic factors

For qualitative variables, data are provided as numbers and 133 percentages. For quantitative variables, data are presented as means 134 and/or medians and standard deviations. To search for explanatory 135 factors relating to progression or stability of the AIS grade between 136 admission and discharge (neurological prognosis), Fisher exact tests 137 were performed with improvements in grade (yes/no) on one side 138 and potential explanatory factors of improvement on the other side. 139 The potential explanatory factors were: 140

- age (≥ 60 or < 60); 142
- cause of SCI; 143 144
- surgery (yes/no);
- the type of clinical syndrome caused by the SCI.

Three categories of clinical syndrome were considered to 146 determine prognostic factors (BSPS, CCS and no, or other, identified 147 syndrome) in order to have approximately similar numbers in each 148 149 group.

Then, in order to search for explanatory factors of functional 150 outcome, Fisher exact tests were performed with patients with or 151 without favourable outcomes on one side and potential explana-152 tory factors of outcomes on the other side. Three outcomes were 153 154 assessed:

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