Assessment of autonomic dysreflexia in patients with spinal cord injury

Armin Curt, Boris Nitsche, Biljana Rodic, Brigitte Schurch, Volker Dietz

Abstract

Objectives and methods—To assess the impairment of supraspinal control over spinal sympathetic centres and the occurrence of autonomic dysreflexia in patients with spinal cord injury. Autonomic dysreflexia is caused by the disconnection of spinal sympathetic centres from supraspinal control and is characterised by paroxysmal hypertensive episodes caused by non-specific stimuli below the level of the lesion. Therefore, patients with spinal cord injury were examined clinically and by different techniques to assess the occurrence of autonomic dysreflexia and to relate disturbances of the sympathetic nervous system to episodes of autonomic dysreflexia.

Results—None of the paraplegic patients, but 59% (13/22) of tetraplegic patients (91% of the complete, 27% of the incomplete patients) presented signs of autonomic dysreflexia during urodynamic examination. Only 62% of the tetraplegic patients complained about symptoms of autonomic dysreflexia. Pathological sympathetic skin responses (SSRs) of the hands were related to signs of autonomic dysreflexia in 93% of cases. No patient with preserved SSR potentials of the hands and feet showed signs of autonomic dysreflexia, either clinically or during urodynamic examination. Ambulatory blood pressure measurements (ABPMs) indicated a loss of circadian blood pressure rhythm (sympathetic control) but preserved heart rate rhythm (parasympathetic regulation) only in patients with complete tetraplegia. Pathological ABPM recordings were seen in 70% of patients with symptoms of autonomic dysreflexia.

Conclusions—The urodynamic examination was more sensitive in indicating signs of autonomic dysreflexia in patients with spinal cord injury, whereas SSR allowed the assessment of the degree of disconnection of the sympathetic spinal centres from supraspinal control. Using ABPM recordings the occurrence of episodes of autonomic dysreflexia over 24 hours and the effectiveness of therapeutical treatment can be assessed.

Keywords: spinal cord injury; autonomic dysreflexia; urodynamic examination; 24 hour ambulant blood pressure measurements; sympathetic skin response

In patients with spinal cord injury both the spinal somatic (motor-sensory) and autonomic (sympathetic) nervous system can be affected. Therefore, patients can have sympathetic failure (for example, orthostasis) due to a disconnection of the spinal sympathetic centres from supraspinal control (hypothalamus, brainstem). By contrast with the spinal sympathetic pathway, parasympathetic nerve fibres innervating the cardiovascular system are conveyed by the vagus nerve and run outside the spinal cord. Therefore, patients with spinal cord injury usually have autonomic failure due to the disconnection of the spinal sympathetic system whereas cardiac parasympathetic pathways remain unaffected.

Autonomic failure can lead to autonomic dysreflexia characterised by paroxysmal hypertensive episodes combined with headache, profuse sweating, and concurrent bradycardia. Due to the loss of supraspinal sympathetic control, non-specific stimuli below the level of injury (for example, bladder distension, rectal manipulation, painful stimuli to the skin) can induce excessive sympathetic spinal outflow below the level of lesion, which causes the cardiovascular changes. This sympathetic hyperactivity is balanced by activation of the vagus nerve via the baroreceptor reflex, which leads to the concurrent bradycardia. An adequate diagnosis and management of autonomic dysreflexia is critical for the rehabilitation programme of patients with spinal cord injury, as these episodes are potentially fatal. Those patients with a level of injury higher than T6 are particularly prone to autonomic dysreflexia. Therefore, this study focused on improving the diagnostic assessment of the spinal sympathetic nervous system and autonomic dysreflexia in patients with traumatic tetraplegia.

The aim of this study was to evaluate the effect of impaired function of the descending sympathetic spinal tract on (1) the sympathetic skin responses (SSRs); (2) the results of 24 hour ambulant blood pressure measurements (ABPMs); and (3) the frequency of occurrence of symptoms of autonomic dysreflexia during urodynamic examination. These variables would be expected to provide information related to the damage of the spinal
sympathetic tract in patients with spinal cord injury and also to indicate the best diagnostic approach for the assessment of autonomic dysreflexia even if it is not clinically evident.

Patients and methods

Patients with chronic (>6 months post-trauma) paraplegia and tetraplegia due to traumatic spinal cord injury who were admitted to the paraplegic centre between 1993 and 1995 were studied. All patients were younger than 65 and none were receiving cardiac or hypertensive treatment. Furthermore, patients were excluded if they had a history of cardiovascular diseases (coronary heart disease, arrhythmias, hypertension, or orthostatic dysregulation). All patients underwent neurological examination according to the international American Spinal Injury Association (ASIA) protocol, allowing the classification of the lesion as either complete or incomplete (any sensory or motor function in the sacral dermatomes). Furthermore, the ASIA motor (maximum 100 points) and sensory scores (maximum for both anesthesia and algesia 112 points) indicated the severity of the spinal cord lesion.

SYMPATHETIC SKIN RESPONSE

For the evaluation of the function of the descending spinal sympathetic tract the SSR of the right hand and foot were examined in accordance with techniques described elsewhere. The SSR was recorded using conventional EMG apparatus (Amplaid). Surface EMG disc electrodes (gold cup, Dantec) were attached to the ventral and dorsal surface of the right hand and foot. The SSR was recorded over 10 seconds and sampled at a band pass of 0.3 Hz to 3 kHz. Electrical stimulation was applied using single, square pulses (0.2 ms duration, 10–20 mA) to the contralateral median nerve at the wrist. To avoid pathological SSR recordings due to lesions of the afferent nerve fibres (for example, in traumatic tetraplegic patients) a second method was performed in parallel. Single, transcortical magnetic stimuli were applied (Mag Pro, Dantec, Denmark) using a flat round coil (MC 125 Coil Dantec, 1.4 Tesla) localised over Cz (according to the international 10–20 system). The stimulus duration was 0.1 ms and the intensity 80–100 A/μs. To prevent habituation the interstimulus interval was set at 60 seconds for both stimulation techniques. In accordance with other studies the SSR potentials were considered to be normal when they could reliably be reproduced three times by both stimulation techniques. Those potentials which were either not sufficiently reproducible or absent were classified as pathological.

TWENTY FOUR HOUR AMBULATORY BLOOD PRESSURE MEASUREMENT

The ABPM was performed with a SpaceLabs 90207 Ambulatory Blood Pressure System (SpaceLabs Medical GmbH, Kaarst) as previously described. Systolic blood pressure, diastolic blood pressure, and heart rate were measured and recorded automatically. For statistical analysis, measurements recorded every 20 minutes between 0800 and 1200 represented the day period, whereas hourly recordings from midnight to 0400 represented the night period. This definition was made for the following reasons. During defined day periods patients were involved in an intensive rehabilitation programme, usually mobilised in a wheelchair. The period of bed rest (night period) usually started at midnight and ended, depending on the state of rehabilitation, at 0400 with the next bladder emptying. Therefore, the definition of the day/night period was adapted to the particular needs of the patients in the rehabilitation programme.

URODYNAMIC EXAMINATION

Patients were examined urodynamically while in a supine position with 15° pelvic obliquity. Two microtransducer catheters (No 5 and 3 radio-opaque, FR) were used for the urethrovaginal and anorectal pressure recordings respectively. Proper location of the transducers was controlled by image intensifier. Slow bladder filling (5 ml/min) was performed with a 24% contrast medium (3-acetylaminomethyl-5-acetylamino-2,4,6-tribenzoic acid). Drugs that could interfere with the urethrovaginal function were discontinued 24 hours before examination. In all patients blood pressure and heart rate were continuously monitored throughout the examination (Ohmeda 2300 Finnarex). As described elsewhere, the patients were expected to exhibit signs of autonomic dysreflexia, when they showed paroxysmal dystonic and diastolic hypertensive episodes with concurrent bradycardia due to manipulations of the bladder (bladder distension, voiding, catheter positioning).

DATA ANALYSIS

The mean (SD) of the ABPM, systolic blood pressure, diastolic blood pressure, and heart rate examinations were calculated and the significance of any differences was determined with a two tailed, paired t test. The level of significance was set at P < 0.01.

Results

PATIENTS

Parallel studies were performed on 22 patients with chronic tetraplegia (11 complete, 11 incomplete; 18 men, four women) and nine patients with complete paraplegia (level of lesion <T6, seven men, two women). The second group served as controls, as they were not likely to develop autonomic dysreflexia. Table 1 shows the demographic and clinical data (ASIA scores).

SYMPATHETIC SKIN RESPONSE

No patient with complete tetraplegia displayed normal SSR potentials in the hands or feet using either of the stimulation techniques. In all patients with complete paraplegia SSR potentials of the hands could be recorded, but only 60% (six of 11) gave responses of the feet. In 73% (eight of 11) of the patients with
incomplete tetraplegia (as defined by the ASIA protocol which focuses on somatic nerve function) normal SSR potentials could be recorded in both the hands and the feet (fig 1). Therefore, the clinical ASIA classification failed to accurately assess the extent of disconnection of the spinal sympathetic system in 27% (three of 11) of patients within the group with incomplete spinal cord injury.

No patient with preserved SSR potentials of the hands and feet showed symptoms of autonomic dysreflexia either clinically or during urodynamic examination. Conversely, all patients with symptoms of autonomic dysreflexia showed pathological SSR potentials of the hands and feet. Loss of SSR potentials of the hands was associated with symptoms of autonomic dysreflexia in 93% (12 of 13) of the patients.

**Table 1** Demographic and clinical data of the tetraplegic and paraplegic patients studied

<table>
<thead>
<tr>
<th>Patients</th>
<th>Level of injury</th>
<th>Age (mean range)</th>
<th>ASIA scores (mean SD)</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td></td>
<td></td>
<td>motor (max 100)</td>
</tr>
<tr>
<td>Complete tetraplegia</td>
<td>C4-C7</td>
<td>29-3</td>
<td>17</td>
</tr>
<tr>
<td>(n = 11)</td>
<td></td>
<td>(17-56)</td>
<td>(9-2)</td>
</tr>
<tr>
<td>Incomplete tetraplegia</td>
<td>C2-C8</td>
<td>40-1</td>
<td>80</td>
</tr>
<tr>
<td>(n = 11)</td>
<td></td>
<td>(15-64)</td>
<td>(24)</td>
</tr>
<tr>
<td>Complete paraplegia</td>
<td>T6-L4</td>
<td>33-4</td>
<td>52</td>
</tr>
<tr>
<td>(n = 9)</td>
<td></td>
<td>(19-50)</td>
<td>(3-9)</td>
</tr>
</tbody>
</table>

blood pressure with reduction from 72.5 (9.7) mm Hg to 62.9 (6.5) mm Hg (P < 0.0001). In 91% (10 of 11) of patients with complete tetraplegia (n = 32 recordings) the physiological rhythmicity of circadian blood pressure was absent (fig 2B, C). There was no significant difference between the values of the day period (systolic blood pressure 109-7 (8.5) mm Hg; diastolic blood pressure 62-6 (7.5) mm Hg) and those of the night period (systolic blood pressure 113-6 (9.4) mm Hg; diastolic blood pressure 63-1 (8.1) mm Hg).

Two patients with complete tetraplegia with symptoms of autonomic dysreflexia and episodes of severe cardiovascular dysregulation (paroxysmal hypertensive episodes up to 224 mm Hg with concurrent bradycardia up to 46 bpm) disclosed a pathological ABPM profile reflecting these episodes (fig 2B). After effective treatment (antihypertensive drugs, sphincterotomy) the ABPM profile showed the characteristics of patients with complete tetraplegia (loss of circadian rhythmicity) without the episodes of severe blood pressure dysregulation (fig 2C).

By contrast with the blood pressure recordings, the circadian rhythmicity of the heart rate was preserved in all tetraplegic and paraplegic patients. The patients with complete tetraplegia (heart rate 71.1 (17.7) to 60.4 (11.4) bpm) as well as the patients with incomplete tetraplegia (heart rate 83.2 (13.2) to 68.1 (10.6) bpm) showed a significant (P < 0.001) reduction in heart rate during the night period. Therefore, the results of ABPM showed severe dysfunction of the spinal sympathetic system in patients with complete tetraplegia, whereas the parasympathetic regulation was normal (because the circadian rhythmicity in heart rate is controlled by the extraspinal vagal nerve).

Of those patients presenting symptoms of autonomic dysreflexia 70% (nine of 13) exhibited pathological ABPM recordings. The loss of circadian rhythmicity of blood pressure was associated with the occurrence of autonomic dysreflexia in 91% (12 of 13) of patients. Of those patients with pathological SSR potentials 27% (three of 11) still showed normal ABPM measurements.

**URODYNAMIC EXAMINATION**

During urodynamic examination, which was combined with continuous monitoring of the blood pressure and heart rate, no paraplegic patient (level of spinal cord injury < T6), but 59% (13 of 22) of the tetraplegic patients displayed signs of autonomic dysreflexia. Of the patients with clinically complete tetraplegia...
patients showed episodes with significant (P < 0.001) increases of systolic and diastolic blood pressures of over 30% (table 2) and a concurrent significant reduction of the heart rate of about 15% (P < 0.01). Signs of autonomic dysreflexia were induced by bladder distension and were related mainly to hyperreflexive bladder contractions and detrusor-sphincter dysynergia. Only 62% (eight of 13) of the patients had clinical symptoms of autonomic dysreflexia.

**Discussion**

Tetraplegic patients and patients with a high level of thoracic spinal cord injury (> T 6) are prone to develop autonomic dysreflexia, which is an autonomic failure due to the loss of supraspinal control of spinal sympathetic centres below the level of lesion. After recovering from spinal shock the remaining sympathetic spinal centres can become hyperreflexive such that non-specific stimuli below the level of lesion can provoke uncontrolled, excessive sympathetic outflow with subsequent cardiovascular effects. The most effective stimulus to provoke such hyperreflexive sympathetic activity below the level of lesion seems to be bladder distension. In patients with spinal cord injury the enhanced sympathetic activity due to bladder stimulation could be shown by recording SSR potentials during urodynamic examination.

In the present study SSR potentials were recorded in patients with spinal cord injury to test the function of the descending spinal sympathetic tract. In accordance with studies performed in other spinal cord diseases the technique used here allows the assessment of the degree to which supraspinal control of the spinal sympathetic system is impaired. The findings show that a loss of or impaired spinal sympathetic control is related to the development of autonomic dysreflexia. No patient with preserved SSR potentials showed autonomic dysreflexia symptoms either clinically or during urodynamic examination. By contrast, all patients with autonomic dysreflexia symptoms had pathological SSR. This fits in with the finding that in three tetraplegic patients with incomplete lesions of the somatic nervous system (motor and sensory functions), but complete disconnection of the autonomic spinal centres (abolished SSR potentials after supraspinal stimulation) displayed typical signs of autonomic dysreflexia during urodynamic examination. Therefore, autonomic failure can be assessed by the recording of SSR potentials, even when they are not clinically evident.

The influence of the descending spinal sympathetic tract on the cardiovascular system can also be assessed using ABPM recordings. By contrast with healthy subjects and both paraplegic and patients with incomplete tetraplegia, almost all patients with complete tetraplegia showed a loss of physiological regulation of circadian blood pressure. This is consistent with findings from patients with pure autonomic failure, who also show a loss of cir-

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**Figure 2** (A) Normal ABPM measurements with a preserved physiological drop of blood pressure during the night period (midnight to 0400). (B) Recordings from a patient with complete tetraplegia and multiple episodes of autonomic dysreflexia. (C) Recordings from the same patient after effective treatment of autonomic dysreflexia with the antihypertensive drug nifedipine. Note that there is still a loss of the circadian regulation of blood pressure.

**Table 2** Changes in systolic (SBP) and diastolic (DBP) blood pressures and heart rate (HR) in patients with signs of autonomic dysreflexia during urodynamic examination

<table>
<thead>
<tr>
<th>Cardiovascular values</th>
<th>Basic mean (SD)</th>
<th>Dysreflexia mean (SD)</th>
<th>Difference (%)</th>
<th>Absolute difference</th>
</tr>
</thead>
<tbody>
<tr>
<td>SBP (mm Hg)</td>
<td>115 (17)</td>
<td>155 (32)</td>
<td>35 ± 5</td>
<td>40 ± 25</td>
</tr>
<tr>
<td>DBP (mm Hg)</td>
<td>64 (13)</td>
<td>82 (16)</td>
<td>31 ± 2</td>
<td>17 ± 16</td>
</tr>
<tr>
<td>HR (bpm)</td>
<td>59 (6)</td>
<td>53 (10)</td>
<td>-14 ± 3</td>
<td>8 ± 6</td>
</tr>
</tbody>
</table>

P < 0.01.
cadian rhythmity of blood pressure.  

By contrast with the impaired cadian rhythmity of blood pressure in patients with complete tetraplegia, the regulation of the cadian heart rate rhythmity was preserved. This is due to the fact that the cadian regulation is under the control of the vagus nerve.  

Nevertheless one third of patients lacking SSR had a preserved rhythmity of physiological cadian blood pressure and only about two thirds of patients with typical signs of autonomic dysreflexia during urodynamic examination showed a loss of the physiological cadian blood pressure regulation. These findings indicate that blood pressure regulation is more resistant to impairment after disconnection of the spinal sympathetic centres. However, the ABPM recordings are useful in recording the frequency and severity of episodes of autonomic dysreflexia during daily living. Furthermore, the effectiveness of medical (antihypertensive drugs) and surgical (sphincterotomy) treatment of these patients can be monitored using ABPM recordings.

Urodynamic examination can be regarded as an effective and standardised diagnostic procedure for evoking signs of autonomic dysreflexia. In this study it was clear that urodynamic examination is a very sensitive tool in the diagnosis of autonomic dysreflexia, as only half of the patients who showed signs of autonomic dysreflexia during urodynamic examination presented clinical symptoms of autonomic dysreflexia. As described earlier episodes of autonomic dysreflexia during urodynamic examination were characterised by increases of both systolic and diastolic blood pressures by about 30%, while the heart rate decreased by about 15%. Therefore, the recording of cardiovascular variables during urodynamic examinations is important to assess signs of autonomic dysreflexia.

Conclusion

In patients with spinal cord injury autonomic failure due to the impaired function of the spinal sympathetic tract can be assessed by different techniques. The SSR recordings are the most sensitive at indicating the degree of damage of the descending spinal sympathetic system. The combination of urodynamic examinations and ABPM recordings allows the prediction of the occurrence and severity of autonomic failure (sympathetic disconnection). Therefore, early diagnosis and treatment of autonomic dysreflexia, one of the most serious complications in patients with a high level (> T6) of spinal cord injury, becomes possible.

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