Organisation of the sympathetic skin response in spinal cord injury

P Cariga, M Catley, C J Mathias, G Savic, H L Frankel, P H Ellaway

The sympathetic skin response (SSR) is a technique that records changes in skin conductance after activation of sweat glands in skin areas rich in eccrine glands (commonly palmar and plantar sites) under the neural control of sympathetic cholinergic (sudomotor) fibres. Potentials generated by the SSR can be recorded in response to various stimuli; these include electrical peripheral nerve stimulation, acoustic stimuli, or magnetic stimulation of nerves or the brain, although magnetic stimulation lacks specificity in terms of sensory pathways involved (it comprises acoustic, magnetic, and tactile stimuli). The SSR is dependent on integrity of peripheral sympathetic cholinergic pathways, as it is preserved in selective sympathetic adrenergic failure (as in dopamine β-hydroxylase deficiency), and is absent in pure (peripheral) autonomic failure (with sympathetic adrenergic and cholinergic failure) and in pure cholinergic dysautonomia.

Although the SSR is an established technique for the study of peripheral sympathetic neuropathies, its clinical application in investigation of central sudomotor pathways is still debated. Although there is evidence in humans of arousal related cerebral centres that control the SSR, as reported in recent neuroimaging studies, previous reports suggested that the SSR can be generated within the isolated spinal cord.

In addition to limitations related to the use of SSR in general, such as habituation and response variability, the application of the technique in patients with spinal cord injury (SCI) needs to be considered in the light of two factors. The first is that with the wide range of neurological features of SCI, the segmental level and completeness of the lesion are of importance because of the thoracolumbar sympathetic outflow. Secondly, autonomic function is not taken into account in the International Standard for Neurological and Functional Classification of SCI.

The aim of our study therefore was to assess the capability of an isolated spinal cord to generate SSR and determine the required levels of integrity of spinal cord and supraspinal connections. Electrical peripheral nerve stimulation only was used, as such a method offers the advantage of providing an objective measure of stimulus intensity and it can be compared in nerves above and below the neurological level of SCI.

We used electrically evoked SSR in healthy volunteers and patients with SCI clinically classified at different levels of lesion. Stimulation of peripheral nerves above and below the lesion was employed with simultaneous, bilateral recording from palmar and plantar sites.

SUBJECTS AND METHODS

Subjects

Twenty-nine patients with SCI (21 male and eight female), mean age (SD) 37 (16) years, and 10 age matched healthy control subjects (six male and four female), mean age (SD) 37 (12) years participated in the study after giving informed consent. Ethical approval was obtained from the Aylesbury Vale local research ethics committee. None of the participants was taking any medication with known autonomic effects. The cause of SCI was traumatic in 22 patients, surgical in four, and infective in three. The time from injury ranged from 3 to over 1000 weeks, with an average of 176 weeks from injury. Subjects with peripheral neuropathy, diabetes, carpal tunnel syndrome (evaluated with nerve conduction studies), traumatic brain injury (excluded by imaging and detailed neurological examination), or any neurological condition other than SCI were excluded from the study. Only subjects free from pain, pressure sores and clinical depression, and fit enough to be tested in a sitting position, were recruited. Subjects with clinical syndromes involving spinal cord hemisection (Brown-Séquard) or peripheral nerve damage (cauda equina syndrome) were also excluded from the study. A detailed clinical examination was performed and each subject was clinically classified at different levels of SCI.

Methods

Palmar and plantar SSR to peripheral nerve electrical stimulation (median or supraorbital nerve above the lesion, and peroneal nerve below the lesion) were recorded in 29 patients with SCI at various neurological levels and in 10 healthy control subjects.

Results

In complete SCI at any neurological level, SSR was absent below the lesion. Palmar SSR to median nerve stimuli was absent in complete SCI with level of lesion above T6. Plantar SSR was absent in all patients with complete SCI at the cervical and thoracic level. In incomplete SCI, the occurrence of SSR was dependent on the preservation of supraspinal connections. For all stimulated nerves, there was no difference between recording from ipsilateral and contralateral limbs.

Conclusions

No evidence was found to support the hypothesis that the spinal cord isolated from the brain stem could generate an SSR. The results indicate that supraspinal connections are necessary for the SSR, together with integrity of central sympathetic pathways of the upper thoracic segments for palmar SSR, and possibly all thoracic segments for plantar SSR.
neurological examination was performed on all subjects, after which they were divided into the following groups, based on the American Spinal Injury Association (ASIA) impairment scale modified from Frankel et al:

**Group A**
Seventeen patients with SCI at various neurological levels from C3 to L1, with ASIA grade A (complete lesion: no motor function preserved below the lesion and no sensory function in the sacral segments S4-S5).

**Group B**
Four patients with SCI at neurological level T1 or above, with ASIA grade B (incomplete lesion, sensory preservation including the sacral segments S4-S5 but no motor preservation below the neurological level).

**Group C**
Four patients with SCI at neurological level T1 or above, with ASIA grade C. This includes incomplete lesion with some degree of motor preservation; more than half of key muscles below the neurological level have a muscle grade less than 3 on a scale of 0–5 (0=no contraction; 5=normal strength), according to Medical Research Council guidelines.

**Group D**
Four patients with SCI at neurological level T1 or above, with ASIA grade D (incomplete lesion with some degree of motor preservation; more than half of key muscles below the neurological level have a muscle grade greater than or equal to 3).

**Group N**
Ten healthy control subjects.

The essential individual characteristics of all patients with SCI, including neurological level, ASIA motor and sensory scores, and time from onset of paralysis are shown in table 1.

## Investigations
The SSR was recorded from palmar and plantar surfaces, both left and right, using neonatal ECG Ag/AgCl surface electrodes (Arbo) with differential recording. One electrode was placed on the volar aspects of the hands or feet, on the skin above the third metacarpal or metatarsal bones (at 3 cm from the distal end). The second electrode was placed on the corresponding area of the dorsal aspect of the relevant hand or foot.

Optically isolated amplifiers were used with gain of 100 and filters at –3 dB below 0.1 and above 100 Hz, connected to a computer through an analog/digital interface. The signals were sampled at a rate of 200 Hz. The traces were recorded from 0.5 seconds before to 8 seconds after the trigger stimulus. To minimise interference from spontaneous potentials, the operator monitored the baseline on an oscilloscope before releasing the electric stimulus, which was delivered at a varying time (30 to 35 seconds) after the previous stimulus.

During the test the subject sat in a chair or wheelchair and was relaxed but not asleep. The room temperature was maintained at 24±0.5°C. The skin temperature of the upper and lower limbs was monitored and stayed constant from the beginning to the end of the test (intrasubject variability <0.2°C). This protocol was followed as temperature is known to affect conduction velocity of unmyelinated sympathetic fibres.

Each stimulus consisted of a single electric pulse (width 0.5 ms). All subjects were stimulated at the median nerve at the wrist and peroneal nerve at the ankle. The stimulus intensity was set to 1.5 times the motor threshold of the stimulated nerve. In patients with complete SCI who did not have SSR to

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Table 1 Presence or absence of palmar and plantar SSR to stimuli above or below the lesion in patients with SCI

- No SSR; + SSR, amplitude <50% of normal mean; ++, SSR amplitude between 50% and 75% of normal mean; +++, SSR amplitude >75% of normal mean.
peroneal stimulation, the stimulation intensity was progressively increased up to levels equal to five times the motor threshold of normal subjects, unless the electric stimulus caused a muscle response diffuse enough to be noticed by the patient. The intensity of this stimulation would have been painful in healthy normal subjects. In tetraplegics patients, with a level of lesion higher than the root supply of the median nerve, the supraorbital nerve was also stimulated at the forehead, a technique used in patients with SCI to study the SSR with stimuli above the lesion level. A minimum of four events was recorded for each nerve and intensity level.

Analysis

The peak to peak amplitude and latency of SSR were measured. Only repeatable responses, consistent in latency, and temporal relation to the electric stimulus, were considered as electrically evoked SSRs. Potentials obtained in the presence of muscle spasms, passive or voluntary limb movement, deep inspiration, or coughing were excluded from the analysis. The latency (in ms) and amplitude (in mV) of SSRs were measured for the first four trials to each nerve or, when the stimulus intensity was increased above 1.5 motor threshold, from the first trial producing a response. Values are expressed as mean (SD). Statistically significant levels reported compare a given group with controls, unless otherwise specified. A $\chi^2$ (or Fisher’s exact test where appropriate) was used to compare the rate of responders/non-responders (where responders refer to subjects presenting SSR), and analysis of variance (ANOVA) was used to compare latency and amplitude of responses in different groups.

RESULTS

The results reported refer to right palmar and plantar SSR to stimulation of the right median or supraorbital nerve above, or the peroneal nerve below the lesion level. Responses were also obtained from left nerve stimulation and left palmar/plantar recording. There was no significant difference between sides for response rate, amplitude, and latency.

Control subjects

All normal subjects had palmar and plantar SSR to median and peroneal nerve stimulation. For palmar SSR to median nerve stimulation, latency was 1476 (162) ms and amplitude 2.98 (1.5) mV. Plantar SSR to the same stimuli had longer latency (2231 (305) ms) and smaller amplitude (1.77 (1.11) mV) than palmar SSR (fig 1).

Figure 1 Superimposed SSRs to four consecutive right median nerve stimuli in a normal subject. Stimulus was applied at time 0 (artefact in right hand trace).

For palmar SSR to peroneal nerve stimulation, latency was 1568 (SD 156) ms and amplitude 2.32 (SD 1.82) mV. Similarly to median nerve stimulation, plantar SSR to peroneal nerve stimulation had longer latency (2205 (215) ms) and smaller amplitude (2.29 (1.44) mV) than palmar SSR to the same stimuli.

All normal subjects had palmar and plantar SSR to median nerve stimulation, latency was 1476 (162) ms and amplitude 2.98 (1.5) mV. Plantar SSR to the same stimuli had longer latency (2231 (305) ms) and smaller amplitude (1.77 (1.11) mV) than palmar SSR (fig 1).

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Palmar SSR to supralesion stimuli

In group A (complete SCI), the presence of a palmar SSR to supralesion stimuli (whether median or supraorbital nerve) was related to the level of the lesion (table 1). In only one of 11 patients with complete SCI above level T6 (patient 7, level T2) was there a palmar SSR obtained (latency 1358 (124) ms; amplitude 2.18 (0.25) mV; fig 2). One of the two patients with complete SCI at level T6 also had a response (patient 12: latency 1540 (140) ms; amplitude 1.67 (0.86) mV). All patients with complete SCI below T6 had a palmar SSR (latency 1495 (91) ms; amplitude 4.42 (3.61); see fig 3). There were no significant differences in latency or amplitude between the SSR obtained in group A and in the control group (p>0.5).

Figure 2 Superimposed SSRs to four consecutive right median nerve stimuli in a patient with SCI grade A, level T2 (patient 7). There are repeatable, bilateral palmar responses of consistent latency whereas no plantar responses are recorded. Stimulus was applied at time 0 (artefact in right hand trace). An occasional artefact (arrow), inconsistent with the properties of an evoked response [non-repeatable, inappropriate latency to the stimulus] is present in the left foot trace.

Figure 3 Superimposed SSRs to four consecutive right median nerve stimuli in a patient with complete, low thoracic SCI (patient 15: level T11, ASIA grade A). Stimulus was applied at time 0 (artefact in right hand trace). Repeatable palmar SSRs of normal amplitude are present in both sides. No plantar responses are present.

Group A

Palmar SSR to supralesion stimuli

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Plantar SSR to supralesion stimuli

A plantar SSR to supralesion stimuli (latency 2183 ms, amplitude 1.1 mV; fig 4) was recorded in only one patient from group A (table 1, subject 17). That patient had SCI at level L1, the lowest in the group.

Peroneal nerve stimuli

No patients from group A had either palmar or plantar SSRs to peroneal nerve stimuli (p<0.001), even at five times normal motor threshold stimulation (fig 5).

Group B

No patients from group B (all with SCI at cervical level) had palmar or plantar SSRs to either median or peroneal nerve stimuli (p<0.001), similarly to all patients with complete SCI with a comparable level of lesion.
and sensory cortex, as they could feel the activating stimuli). Whom the sensory stimulus was conducted to the brain stem level or stimulation intensity. An SSR to stimuli below the trical stimuli. In none of our patients with complete SCI (ASIA DISCUSSION

stimuli (latency 2205 (215) ms; amplitude 2.55 (0.52)). Two of four patients had plantar SSRs to peroneal nerve stimuli below the lesion (latency 1661 (99) ms; amplitude 1.53 (0.63) mV). Three of four patients had palmar SSRs to peroneal nerve stimuli (latency 2330 (229) ms, amplitude 1.77 (1.11) mV). No plantar SSRs to peroneal nerve stimuli were recorded in this group (p<0.001).

Group D

All patients from group D had palmar SSRs to median nerve stimuli (latency 1343 (212) ms, amplitude 2.33 (2.02) mV), and three of four patients had plantar SSRs to the same stimuli (latency 2288 ms; amplitude 0.25 mV), and one patient had palmar SSRs to peroneal nerve stimuli (patient 22: latency 1598 ms; amplitude 0.73 mV). No plantar SSRs to peroneal nerve stimuli were recorded in this group (p<0.001).

DISCUSSION

In our study of complete SCI at different levels there was no evidence of isolated spinal cord sudomotor responses to electrical stimuli. In none of our patients with complete SCI (ASIA grade A) was a palmar or plantar SSR obtained in response to electrical stimuli applied below the lesion, regardless of lesion level or stimulation intensity. An SSR to stimuli below the lesion was also absent in patients with SCI of grade B (in whom the sensory stimulus was conducted to the brain stem and sensory cortex, as they could feel the activating stimuli). Conversely, some patients with a degree of preservation of both motor and sensory function (ASIA grades C and D) had palmar and plantar SSRs to stimuli below the lesion. The mean amplitude of these responses was smaller than normal, although the difference was not statistically significant. The latency of the SSRs was within the normal range, suggesting that the responses were conducted through the same neurological pathways used in normal subjects.

A plantar sudomotor response to lower extremity noxious stimuli has been reported in patients with functionally complete cord transection at cervical and thoracic levels. Such responses, however, were not recorded consistently, and the duration of electrical stimulation (pulse trains of 0.5 s) did not favour a precise analysis of the latency. Also, in those two studies, patients with SCI were only broadly classified by a single neurological level, and the partial preservation of sensory and motor pathways in segments below the lesion level was not documented using ASIA groups or quantified using ASIA motor and sensory scores. Therefore, the possibility of some degree of supraspinal connection and partial preservation of pathways below the lesion was not excluded in those patients.

Our results in healthy control subjects demonstrated no significant differences between median and peroneal nerve stimulation in SSR amplitude or response rate. In patients with SCI, however, differences in response rate and SSR amplitude could be seen according to the degree of preservation of the stimulated sensory pathway. In general, supra lesion stimuli produced closer to normal responses than did peroneal nerve stimuli. A similar effect was shown by groups with a different degree of motor preservation (group D had closer to normal responses than group C). Habitation of the SSR, which has been suggested to have a non-neurological, peripheral component at sweat gland level, might contribute to these differences. Patients with SCI often have trophic skin disturbances including severe skin dehydration, especially in the lower limbs. However, we did not find a significant reduction in amplitude over the first four responses either in patients with SCI or in normal subjects, indicating that habituation did not affect our results. We suggest therefore that a quantitative effect of damage to both sensory and motor pathways is responsible for the differences in SSR amplitude between the different ASIA groups.

Overall, our findings indicate that connections with supraspinal structures are essential for evoking SSRs. The association of preserved somatomotor function and preservation of SSRs might be explained by the anatomical proximity, in the spinal cord, of the sympathetic preganglionic nuclei (intermediolateral columns from level T1 to L2) to the anterior horn nuclei.

Our data show that the SSR, which is an indicator of sympathetic cholinergic function, is very often abolished or impaired (both in palmar and plantar sites) in SCI above level T6 (depending on the completeness of SCI and ASIA level). On the other hand the sympathetic adrenergic function at these levels of lesion is released from supraspinal control and is typically hyperreactive to stimuli coming from below the lesion (skeletal muscle spasms, bladder or rectal activity, labour, noxious stimuli such as anal fissures). The condition leads to autonomic dysreflexia, to which patients with SCI above T6 (especially complete) typically are known to be prone. Dissociation between autonomic dysreflexia and presence of palmar SSR was also reported in previous studies by Curt et al. This might reflect differences in the sympathetic neural control of the cardiovascular and the sudomotor system.

Our data from patients with complete SCI indicate that supraspinal connections to the upper thoracic cord segments are required for the palmar SSR, possibly with borderline levels around T6. This does not mirror the levels required for the somatic innervation of the upper limbs via dorsal and ventral
roots, and is more probably related to distribution of preganglionic sympathetic neurons to the sympathetic ganglionic chain. These findings are consistent with the findings of Yokota et al.\(^3\) that the SSR probably reflects the integrity of central preganglionic pathways as well as peripheral postganglionic activity.

It is possible that supraspinal connections down to L1 might be essential for plantar SSR, but we had too few patients with relevant injuries to be certain. The high prevalence of nerve root damage (cauda equina syndromes) in lumbar SCI limits the availability of patients with purely central lesions at lumbar level.

An apparently exceptional finding was seen in patient 7, who had a palmar SSR to supraspinal stimuli despite having an ASIA grade A lesion at level T2. These responses, which had smaller amplitude than normal, were bilateral, repeatable, clearly related to the stimulus, and consistent in latency and waveform shape with the SSR recorded in normal volunteers. The neurological examination indicated a sensory level at T2 bilaterally, involving all types of sensation, with partial preservation at level T3. No motor or sensory function was detected below this level. The patient had no documented episodes of autonomic dysreflexia. A provocative test was performed using suprapubic bladder stimulation and blood pressure recording,\(^7\) which was negative. The injury was in its chronic phase (67 weeks) and therefore, if a predisposition to develop autonomic dysreflexia were present, it would have been manifest by the time of the study. This might be consistent with the possibility that the lesion was not complete with regard to the sympathetic pathways. It must be emphasised that ASIA classification does not take into account autonomic pathways, and partially preserved connection of some segments below the highest damaged level is not unusual, even in SCI classified as complete. It is possible that our subject had partially preserved sympathetic pathways for a few segments below the lesion, sufficient to produce palmar SSR of decreased amplitude. However we cannot exclude the possibility that, in a minority of cases, partial palmar responses might be present even when the sympathetic pathways are totally interrupted at a higher level than T6.

In conclusion, our study has defined the inability of the isolated spinal cord to generate SSR. It emphasises the potential of electrically evoked SSR in investigation of the neural control of sweating and central sudomotor pathways in SCI, particularly at a thoracic level. Furthermore, it might provide a useful technique in assessing the completeness of lesion in SCI and improve the current classification methods for neurological level and completeness of SCI, which are based on the somatic nervous system only.

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