SHORT REPORT

Callosal disconnection syndrome and knowledge of the body: a case of left hand isolation from the body schema with names

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Abstract

A patient is described who presented with a disturbance of body cognition confined to the left side of the body. She showed difficulties in naming the left fingers and in moving the named left fingers. She also showed great difficulty in pointing to named parts of the body with her left hand. Earlier in the course of the disease she showed a personification phenomenon of the left hand. Brain MRI showed involvement of the entire corpus callosum, probably due to occlusion of a branch of the anterior cerebral artery. It is speculated that this syndrome is caused by disconnection of the right hemisphere from the left hemispheric body schema.

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We encountered a patient who showed an array of disconnection syndromes after an infarction of the corpus callosum. We have already reported the left sided avoiding reaction of this patient and discussed the symptom in terms of possible disconnection of the asymmetrically organised manual spatial function. Other intriguing symptoms of this patient were a left sided alien hand sign in the early course of the disease and persistent difficulty in identifying body parts with the left hand. To our knowledge, reports of unilateral autotopagnosia associated with the callosal disconnection syndrome have not appeared in the medical literature.

We studied the symptoms in detail and speculate on an underlying mechanism in terms of lateralisation of a body recognising system—that is, a body schema.

Case presentation

The details of this 56 year old right handed housewife have already been reported. In March 1988 she had a cerebral infarction, most likely a thrombosis of a branch of the left anterior cerebral artery. Neuroimaging studies including CT and MRI confirmed an ischaemic lesion confined to the left half of the corpus callosum from the genu through the entire length of the body (fig 1).

Subsequently, she showed various typical disconnection syndromes including alien left hand sign, diagnostic apraxia, self grasping by the left hand, left hand tactile anomia, left hand agraphia, left ideomotor apraxia, left sided motor initiation difficulty, left sided crossed avoiding, and left unilateral neglect when drafting with the right hand. This left unilateral spatial neglect was never seen with tasks involving the left hand. General behavioural neglect toward the left was not seen in daily activities.

On 20 December 1988 she was alert and cooperative. Cranial nerves were normal including the visual fields. Movements of the left upper and lower limbs were clumsy in finger to nose and heel to shin tests, but no clear weakness or ataxia was present. Muscle tone was normal. Deep tendon reflexes were increased bilaterally but no pathological reflexes were detected. Superficial sensation was normal. Joint position sense was impaired in the fingers of the left hand. Vibration was slightly decreased bilaterally. For stereognosis, the right hand was normal and the left hand
was correct for 80% in non-verbal response tasks.

No appreciable change was detected in the neurological signs at a re-examination three years after the onset. In this examination we tested for the first time left-right transfer of tactile localisation and finger pattern. Transfer from the right to the left often elicited stiffness of the left hand. Even when the left-right transfer was feasible, it ended in errors.

Brain CT taken four years after the onset showed an advance of the cortical atrophy in the left frontal area. The anterior horn of the left hemisphere was enlarged and the damage of the corpus callosum at this level was clear.

**Studies on body identification**

**PERSONIFICATION**

For three months after the onset she showed a pronounced personification phenomena of the left, non-paralysed, hand. She often behaved as if the left hand were a real person. The personification usually occurred when she could not control her left hand voluntarily. Here is an example:

When she was being tested for apraxia, the left hand grasped the right hand performing a task. She tried to let the left hand go to no avail. Then she started talking towards the left hand, "Grandma, would you please let me go. You are warm, but my hand is sweating and uneasy."

**IDENTIFICATION OF THE PATIENT'S OWN BODY**

Based on the method described by Benton, extensive tests of body identification were performed over a two year period one year after the onset. During this period no personification phenomena were seen. The table summarises the test results.

For naming of fingers three methods were employed, visual naming and two non-verbal methods of naming a finger touched by an examiner with the patient blindfolded and with the hand behind the patient's back. For comprehension of finger names, two methods were used: pointing to a named finger with the contralateral hand and moving a named finger. For non-verbal identification of fingers the patient was asked to move a touched finger. For identification of body parts the patient was asked to name a touched part, point to a named part, and to point to a touched part.

The patient's performance was perfect for visual naming of the fingers and the body. Non-verbal naming, and naming of the body parts and the right fingers were perfect; however, the patient performed very poorly for non-verbal naming of the left fingers. The patient performed accurately for pointing to a named left finger with the right hand with the eyes open or closed, but was poor in pointing to the right fingers with the left hand, being much worse when blindfolded. Pointing to a named body part with the right hand was perfect (Fig 2), but it was poor with the left hand both with the eyes closed and open.

Identification of a named finger by moving it was perfect for the right hand with the eyes open, the eyes closed, and the hand behind the body. It was remarkably poor for the left hand, being much worse when blindfolded. With the non-verbal identification task of moving a touched finger, the left fingers responded correctly in both visual and non-visual conditions. Body part identification by pointing to a touched part was fairly good with the left hand, even when non-verbal.

**IDENTIFICATION OF THE EXAMINER'S BODY**

Visual naming of the examiner's body parts including the fingers was perfect. Pointing to the named parts of the examiner's body was also without error if performed with the right hand. Correct pointing rate with the left hand, however, fell to 79%.

**POINTING TO OBJECTS**

Finally we tested pointing ability of the left hand with objects. Five objects were placed on a desk, the patient pointed to a named object when able to see and when blindfolded. Under the visual condition, the patient's response was good. With 10 trials (50 pointings) she was 97% correct. Under the non-visual condition, the response became slow and poor, but the performance improved after she memorised the location of objects. With 10 trials (50 pointings) she was 67% correct. We tested visual pointing capacity of her left hand for six objects, placed on her forehead, ear, shoulder, breast, navel, and knee. With the total of five sessions (30 pointings) she was 80% correct. When the task was switched to pointing to a named body part she was correct in only 37% of 30 pointings. For both tests the performance of the right hand was normal.

**OTHER TESTS RELATED TO THE BODY IMAGE**

We asked 14 verbal questions describing a relation of the various body parts. For
Naming and identification of body parts

<table>
<thead>
<tr>
<th>Right fingers</th>
<th>Left fingers</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>Naming of fingers (% correct):</strong></td>
<td><strong>Appearance</strong></td>
</tr>
<tr>
<td>Visual naming</td>
<td>100</td>
</tr>
<tr>
<td>Name a touched finger With eyes closed</td>
<td>100</td>
</tr>
<tr>
<td>With hand behind body</td>
<td>100</td>
</tr>
</tbody>
</table>

Identification of fingers:

| Point to a named finger | 100 | 100 |
| With the right hand With eyes open | 97 | 97 |
| With the left hand | 97 | 97 |

Move a named finger

| With eyes open | 100 | 100 |
| With eyes closed | 100 | 100 |
| With hand behind body | 100 | 100 |

Move a touched finger

| With eyes open | 100 | 100 |
| With eyes closed | 100 | 100 |

Eyes open

| Eyes closed | 98 |

Naming of body parts (% correct):

<table>
<thead>
<tr>
<th>Identification of body parts</th>
<th>Right</th>
<th>Left</th>
</tr>
</thead>
<tbody>
<tr>
<td>Name a touched part With the right hand</td>
<td>100</td>
<td>100</td>
</tr>
<tr>
<td>With the left hand</td>
<td>100</td>
<td>100</td>
</tr>
</tbody>
</table>

Discussion

The present case presented a unique combination of signs confined to the left hand. Essentially, there were two core symptoms. One was the difficulty of left finger identification. She could neither name the left fingers nor move the named left fingers when vision was excluded. Under visual conditions, these tasks were perfectly conducted. The other was the difficulty of pointing to the named body parts with the left hand. The difficulty covered the whole body including the right fingers. Again, the right hand showed no difficulty.

Autotopagnosia was first described by Pick in 1909 and defined as patients' inability to localise and name the parts of their own body correctly.10 Finger agnosia, described by Gerstmann in 1924,11 belongs to this category of autotopagnosia. Typical patients would not be able to name or move the parts of their own bodies touched by an examiner, or move or indicate the parts named by the examiner. The disorder was always bilateral. Drawing of a human figure may be abnormal.

The present case can be placed in this broad category of autotopagnosia but with exceptional features. Thus, we have a case of tactokinesthetic autotopagnosia confined to the left fingers associated with left hand disorientation for named body parts.

Tactokinesthetic left finger agnosia can be explained in terms of modality specific as well as category specific two way anomaia. Two way anomaia was first described by Geschwind and Fusillo in the colour domain.12 Their patient had difficulty naming colours and pointing to them but preserved the ability to sort colours according to hues, match colours, and recall colour names in response to verbal questions. They explained the symptoms as a disconnection of the linguistic system from the right hemispheric colour system because of combined damage to the left occipital visual system and the corpus callosum, through which the colour information of the right hemisphere passes to the left language area. The identical argument is feasible in the present case. Suppose a reference system for the location and the names of the body—call it the body image with names—is lateralised in the left hemisphere. If this body image with names was separated from the right hemisphere by an appropriately placed callosal lesion, anomaia as well as comprehension difficulty for the names of parts of left side of the body would inevitably result. Tactospatial information from the left finger area must be first processed in the contralateral right hemisphere and subsequently have to pass to the left hemisphere for reference for a body part identification and a name. When a name of the left fingers was provided, the left hemispheric body image with names was activated, but could not transmit this exact localising information to an appropriate area of the right motor area. Axial body information may reach the left hemisphere ipsilaterally resulting in no difficulty in identification. As the right somesthetic processing area is essentially intact, tactile localisation of a touched finger by moving the same finger should be possible.

How can we reconcile the peculiar left hand difficulty of pointing to the named body parts bilaterally with this two way anomaia theory? When given a name, the left hand showed disorientation and could not arrive at a correct destination. But the performance was better when visual than when blindfolded. When given tactile information (touch), the same hand localised the point fairly well under both conditions. This left hand performance could not be due to apraxia or visuomotor ataxia, as the spatial reaching at an object outside the patient's own body was good as shown by her ability in pointing to the body parts of the examiner and pointing to objects even on her own body. The left hand, under right hemispheric control, was unable to relate its spatiomotor information with the left hemispheric body image with names through the corpus callosum.

This theory is consistent with previous ones. Gerstmann,13 for instance, argued that the body schema should be lateralised to the left hemisphere. From our study of “gestures” autotopagnosia in which verbal and non-verbal identification of body parts is impaired, Semennza and Goodglass postulated the exis-
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Callosal disconnection syndrome but toward the interpret behaviourally. Having this hand could be left own body, resulting is disconnected from body. The disconnected right movement of right hemisphere would be thrown autonomy somesthetic information, an argument be activated.

Some might argue that the phenomenon could be interpreted as a partial symptom of the left sided neglect. It was true the patient showed left unilateral neglect with the right hand, but not with the left hand. No spatial inattention toward the left side was seen behaviourally. If there was a problem, it was toward the right. Thus it is difficult to interpret the symptoms in terms of left side neglect. Even the personification phenomenon seen in the present case was different from the similar phenomenon often seen in association with anosognosia. The first is seen in response to a moving left hand, whereas the second is seen in association with a paralysed left hand.

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