In Vivo Evidence of Structural Brain Asymmetry in Musicians

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of a cytosolic protein 60 kD in size over a distance of 150 to 200 μm will be about 3 to 5 min. Simple diffusional signaling between the synapse and cell body could thus account for the delay in the potentiation observed in the present study. Active retrograde transport is known to take place at a rate of about 50 to 180 μm/min (R. D. Allen, J. Metzual, I. Terasaki, S. T. Brady, S. P. Gilbert, Science 218, 1127 (1982)), which would allow much more rapid signaling than diffusion.

17. The increase in cytosolic Ca²⁺ was shown by direct Ca²⁺ imaging with the fluorescence dye FluO-3 in separate cultures, though use of methods previously described (Y. Dan and M. M. Poo, Nature 359, 733 (1992)).

18. The transfection was performed in normal culture medium to facilitate releasing of the plasma membrane. The medium was replaced with Ca²⁺-free solution [115 mM NaCl, 2 mM MgCl₂, 2.5 mM KCl, 10 mM Hepes, 3 mM EGTA, and 0.1% BSA (pH 7.3)] 20 min after the neurotoxin transfection.


24. Addition of actinomycin D at 2.5 μg/ml to Xenopus embryonic cell cultures was shown to inhibit >95% of [3H]uridine incorporation into the trichloroacetic acid-precipitable cell fraction from these cultures (D. K. O'Dowd, Nature 303, 619 (1983)). For blocking of RNA synthesis in Xenopus embryos, actinomycin D (25 μg/ml) has been used (S. Grafi and G. Gafi, FEBS Lett. 236, 403 (1988)). Incubation with cycloheximide at 0.6 μM (or 0.17 μg/ml) for 2 hours was found to achieve about 90% inhibition of [3H]uridine incorporation into the acid-precipitable fraction of these Xenopus cultures [L. A. G. Blair, J. Neurosci. 3, 1430 (1983)]. Higher concentrations of these drugs were used in the present study to ensure more complete inhibition.


26. We thank Regeneron Inc. for providing CNTF and BDNF. L.-G. Lu for technical assistance, and B. Lu, R. Giordano, and J. Alder for helpful comments on the manuscript. Support by grants from the NSF (IBN 22108) and the Office of Naval Research (N00014-80-J1803).

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In Vivo Evidence of Structural Brain Asymmetry in Musicians
Gottfried Schlaug,*† Lutz Jäncke, Yanxiong Huang, Helmuth Steinmetz*

Certain human talents, such as musical ability, have been associated with left-right differences in brain structure and function. In vivo magnetic resonance morphometry of the brain in musicians was used to measure the anatomical asymmetry of the planum temporale, a brain area containing auditory association cortex and previously shown to be a marker of structural and functional asymmetry. Musicians with perfect pitch revealed stronger leftward planum temporale asymmetry than nonmusicians or musicians without perfect pitch. The results indicate that outstanding musical ability is associated with increased leftward asymmetry of cortex subserving music-related functions.

A number of studies have demonstrated that the left hemisphere of the brain is dominant in the production and comprehension of language in the vast majority of persons (1). Similar attempts to localize musical functions have yielded conflicting data, mainly because studies of amusia—that is, impairment of musical skills as a result of cerebral lesions—have failed to reveal structural-functional maps similar to those of language organization (2). This situation has now changed with the introduction of positron emission tomography (PET) to measure regional cerebral blood flow and metabolism during the processing of verbal and nonverbal stimuli. Whereas left hemispheric activation sites are seen during phonological, lexical, or semantic language task performance (3), right hemispheric preponderances are found for melodic and pitch perception, at least in musically naive subjects (4). However, process-

Table 1. Means (±SD) for age, degree of anatomical planum temporale asymmetry (PTA), and size of left and right PT determined with in vivo magnetic resonance morphometry in healthy, right-handed musicians and nonmusicians.

<table>
<thead>
<tr>
<th>Subjects</th>
<th>Age</th>
<th>&amp;PTA</th>
<th>PT size (mm²)</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td></td>
<td></td>
<td>Left</td>
</tr>
<tr>
<td>Musicians (n = 30)</td>
<td>26 (4)</td>
<td>−0.36 (0.25)*</td>
<td>1063 (189)</td>
</tr>
<tr>
<td>Perfect pitch (n = 11)</td>
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<td>−0.57 (0.21)**</td>
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</tr>
<tr>
<td>No perfect pitch (n = 19)</td>
<td>26 (4)</td>
<td>−0.23 (0.17)</td>
<td>1043 (183)</td>
</tr>
<tr>
<td>Nonmusicians (n = 30)</td>
<td>26 (3)</td>
<td>−0.23 (0.24)</td>
<td>896 (236)</td>
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</table>

*Negative values indicate leftward asymmetry of the PT (18). 0.001 compared to musicians without perfect pitch (21).

G. Schlaug, Y. Huang, H. Steinmetz, Department of Neurology, Heinrich-Heine-Universität Düsseldorf, D-40225 Düsseldorf, Germany.
L. Jäncke, Institute of General Psychology, Section of Cybernetical Psychology and Psychobiology, Heinrich-Heine-Universität Düsseldorf, D-40225 Düsseldorf, Germany.

*To whom correspondence should be addressed.
†Present address: Department of Neurology, Beth Israel Hospital, Boston, MA 02215, USA.

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*Negative values indicate leftward asymmetry of the PT (18). 0.001 compared to musicians without perfect pitch (21).
We found that the PT was more lateralized to the left in musicians (P = 0.028). Possession of perfect pitch explained most of the variation in the degree of PT asymmetry among musicians (P < 0.001) (19–21). Musicians with perfect pitch showed stronger leftward PT asymmetry compared to other musicians, whereas musicians without perfect pitch did not differ from controls (Table 1 and Fig. 1).

Our finding of increased leftward PT asymmetry among musicians should be seen in the following context. First, PET has demonstrated that the posterior superior temporal region, including the PT, is involved in music perception (5). Second, in one postmortem myeloarchitectonic study of a musician with melody deafness after circumscribed brain injury, the lesion was centered on the left PT, sparing the primary auditory and inferior parietal cortex (22). Third, gross left-right asymmetry of the PT, as measured in our study, reflects cytoarchitectonic asymmetries of auditory association areas located on the PT (12). Thus, our morphometric findings in musicians may suggest that the functional capacity of cortex shown to preserve musical functions increases with leftward structural asymmetry of this neural system. This result lends anatomical support to behavioral and electrophysiological evidence of a difference in lateralization of musical processing between musicians and nonmusicians, with more left-lateralized representation in musicians (6, 7). Our data concur with the general concept that, because of time constraints of interhemispheric transfer, efficiency of neuronal assemblies is expected to increase with the number of elements clustered in one hemisphere (23). In fact, this principle may be the essence of hemispheric specialization (23).

Our study does not reveal the mechanism creating structural asymmetry. Leftward PT asymmetry usually appears in the human fetus between the 29th and 31st gestational week (24), so that prenatal factors are likely to play a role. Nevertheless, considering that the maturation of fiber tracts and intracortical neuropil, two presumed determinants of gyral shape (25), are still progressing by the age of seven (26), it remains uncertain whether gross anatomy may also be susceptible to some postnatal plastic change, such as in response to specific stimulation (20). Our study demonstrates that individual variability in cognitive performance can co-vary with features of external brain morphology.

REFERENCES AND NOTES

2. Since the 19th century, a number of investigators have ascribed amusia to left or right hemisphere dysfunction. Both the occurrence of amusia without aphasia and of aphasia without amusia have been described (double dissociation), suggesting at least partial independence of the neural structures subserving the respective functions. For overviews, see N. Wertheim, in Handbook of Clinical Neurology, P. J. Vinken and G. W. Bruyn, Eds. (Elsevier, New York, 1969), vol. 4, pp. 190–230; A. R. Damasio and H. Damasio, in Language and Memory, M. Critchley and R. A. Henson, Eds. (Heinemann, London, 1977), pp. 141–155; and J. Siergent, Trends Neurosci. 296, 168 (1986).
5. J. C. Mascott, M. E. Phelps, R. E. Carson, D. E. Kuhl, Neurology 32, 921 (1992). In this study, subjects who later described themselves as having "specific, highly organized" analytic approaches consistently displayed left greater than right temporal activation during a tonal memory task. Right greater than left activation during the same task was found in individuals reporting no specific strategy.
6. T. G. Bever and R. J. Chiarello, Science 185, 537 (1974). After comparing ear advantages for monaural presented melodies in musicians (right-ear advantage) and nonmusicians (left-ear advantage), the authors concluded that "the brain auditory system that has real neuroanatomical concomitants, permitting the utilization of a different strategy of musical apprehension that calls on left hemisphere functions." For an opposing view see R. J. Zatorre, Neuropsychologia 17, 607 (1979).
16. In vivo magnetic resonance morphometry was performed as described and validated previously in human cadaver brains (H. Steinmetz et al., Brain 118, 39, 357 [1990]) through use of a Siemens 1.5 T magnet and a 22-min fast-low-angle-shot magnetic resonance sequence covering the entire brain, yielding a 3-mm by 3-mm by 3-mm magnetization-echo size. Each dataset consisted of 26 contiguous, 1.17-mm-thick sagittal slices. For off-line morphometry, the datasets obtained from musicians and nonmusicians were mixed and coded. They were analyzed independently by two observers (G.S. and H.-J. Freund) using the blinded observer method.
17. The professional musicians were recruited through announcements in three well-known music schools in Germany as well as through personal contacts. All were classical musicians who had either finished their education or were receiving formal training at a music school. No amateur musicians were included. All string players (n = 14) were keyboard players as well but had a preference for string instruments; the others were keyboard players (n = 16). The age-, sex-, and handedness-matched nonmusicians were recruited through announcements in the local medical school. Most of them were medical students or young faculty members in university hospitals. None of them had ever played a musical instrument or received formal musical training. All subjects gave informed consent.
18. Handedness was assessed with the 12-item questionnaire of M. Amossen, Br. J. Psychol. 61, 202 (1970). Consistent right-handedness was defined as performance of all 12 tasks with the right hand, with up to two "either" preferences being acceptable. These male and female musicians were nonconsistent righthanders; all other subjects were consistent righthanders. Also, non-
mammalian Vestibular Hair Cell Regeneration

Birds and mammals are born with a full complement of inner ear hair cells, which were thought to be irreversibly lost when damaged (1). It is now well known that birds have the capacity to regenerate hair cells in their auditory and vestibular organs after damage by acoustic trauma or ototoxic drugs (2) and that these new cells can mediate functional recovery (3). Recent studies by A. Forge et al. (4) and by M. E. Warchol et al. (5) suggest that the vestibular epithelium of the mature mammalian inner ear may also have the ability to produce new hair cells by renewed mitotic activity in response to aminoglycoside injury in vivo (4) and in vitro (5). However, these reports do not provide convincing evidence that the DNA labeling, seen at a low frequency in vitro, is the source of the apparent recovery of hair cell apical surfaces observed in vivo.

Our study was undertaken to determine if cell division can be shown to give rise to new hair cells in normal mature mammalian vestibular epithelium or during the first 6 weeks after aminoglycoside ototoxicity. Three groups of young mature albino Hartley guinea pigs were used. The experimental animals in each group were treated with a single transtympanic injection of the ototoxic aminoglycoside, gentamicin, in the left ear (6). Animals in each control group were given an identical volume of 0.9% saline. The first group of animals was killed after 1 to 6 weeks and used for light microscopic evaluation of damage produced in the sensory epithelium of the utricle (7). The second group, killed after 1 to 16 weeks, was used for scanning electron microscopy (SEM) (8) in order to compare our results with those of Forge et al. (4). In animals of the third group, an osmotic pump filled with [H]thymidine was implanted under the skin of the back with its output leading to a cannula inserted into the perilymphatic space before treatment with aminoglycoside (9). These animals were killed after 1 to 16 weeks (10).

Hair cell damage and loss was evident in the light microscopic sections and SEM analyses of tissue from gentamicin-treated animals (Fig. 1). Experimental animals had fewer hair cells than controls, particularly in the striolar region. Other signs or damage observed by light microscopy of SEM included nuclear pyknosis, nuclear swelling, vacuolization, cytoplasmic extrusion, and stereocilia fusion. The extent of damage was variable at all survival times. At 1 to 2 weeks after gentamicin treatment, hair cell injury was limited primarily to the striolar region in 10 of 16 animals examined by SEM. In three of the animals damaged was observed over a larger area, extending from the striola toward the periphery of the organ. Complete destruction of the sensory hair cells was observed in the remaining three animals. Four weeks after gentamicin administration, one animal displayed hair cell damage extending out from the striolar region; in the other animal bleeding and fusion of stereocilia were seen over the entire surface of the sensory epithelium. In the animal killed 4 months after gentamicin, the surface of the utricle continued to show damaged stereocilia bundles throughout the entire sensory epithelium. The average length of the sensory epithelium and the linear support cell density remained constant between the control and experimental animals (Table 1) (11). However, the linear hair cell density was 51 to 85% lower in experimental animals than controls (P < 0.001).

Table 1. Results of treatment with gentamicin on guinea pig utricle: Length of sensory epithelium, hair cell density, and support cell density. Measurements are averages ± standard deviation.

<table>
<thead>
<tr>
<th>Animal number</th>
<th>Treatment (weeks)</th>
<th>Sensory epithelium length (μm)</th>
<th>Hair cell density (μm²)</th>
<th>Support cell density (μm²)</th>
</tr>
</thead>
<tbody>
<tr>
<td>94-01</td>
<td>1</td>
<td>7.2 (±2.0)</td>
<td>1.6 (±0.7)</td>
<td>11.6 (±4.6)</td>
</tr>
<tr>
<td>94-02</td>
<td>1</td>
<td>9.3 (±2.1)</td>
<td>3.4 (±0.8)</td>
<td>9.9 (±3.1)</td>
</tr>
<tr>
<td>94-03</td>
<td>1</td>
<td>7.2 (±1.6)</td>
<td>2.6 (±1.4)</td>
<td>10.4 (±1.5)</td>
</tr>
<tr>
<td>94-04</td>
<td>4</td>
<td>7.4 (±1.5)</td>
<td>1.3 (±1.5)</td>
<td>8.4 (±1.5)</td>
</tr>
<tr>
<td>94-05</td>
<td>4</td>
<td>8.7 (±2.0)</td>
<td>1.6 (±0.6)</td>
<td>7.1 (±1.6)</td>
</tr>
<tr>
<td>94-06</td>
<td>6</td>
<td>7.2 (±1.9)</td>
<td>4.3 (±0.8)</td>
<td>10.7 (±2.8)</td>
</tr>
<tr>
<td>94-07</td>
<td>6</td>
<td>7.4 (±1.6)</td>
<td>8.7 (±1.7)</td>
<td>12.6 (±5.3)</td>
</tr>
<tr>
<td>94-08</td>
<td>6</td>
<td>8.9 (±2.1)</td>
<td>6.6 (±1.9)</td>
<td>10.2 (±1.2)</td>
</tr>
</tbody>
</table>

*Control group received no gentamicin.

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