Prominent Activation of the Putamen during Essential Palatal Tremor: A Functional MR Imaging Case Study

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CASE REPORT

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SUMMARY: Palatal tremor (PT), also known as palatal myoclonus, is defined by short rhythmic contractions of the palatal musculature. Functional MR imaging (fMRI) revealed prominent bilateral neuronal activation in the putamen associated with essential palatal tremor (EPT) in a 41-year-old man. This implies a central role of the putamen in EPT, most likely as a consequence of diminished inhibition in an afferent pathway. Because fMRI primarily detects activations, dysfunctional areas remain obscure. The present functional study complements previous pathologic studies, which associated PT with lesions to dentate nucleus, red nucleus, and the inferior olive (Guillain-Mollaret triangle).

Palatal tremor (PT), formerly also called palatal myoclonus, is defined by short, mostly rhythmic contractions of the palatal musculature and may occasionally be stimulus-sensitive. PT can be associated with synchronous movements of adjacent structures, including the pharynx, larynx, face, and diaphragm.1,2 PT has been subdivided into an essential form (EPT),3 where no origin can be found and MR imaging is usually normal,4 and symptomatic forms (SPT).5 In SPT, stroke (46%) is the most common cause, followed by trauma (11%) and demyelinating lesions (10%).6,7 Early pathologic studies of PT2,4 outlined an important role of lesions affecting the dentatorubral-olivary pathway, or Guillain-Mollaret triangle.9 This circular pathway connects the red nucleus to the inferior olivary nucleus and the contralateral dentate nucleus of the cerebellum via central tegmental tract (red nucleus–dentate nucleus), and superior cerebellar peduncle (dentate nucleus– Inferior olive), inferior cerebellar peduncle (inferior olive–dorsal olive) fibers, but not in the olivodentate fibers.10 Further, hypertrophic degeneration of the inferior olive was reported in SPT.7,11 Common radiologic changes in SPT are an increase in T2 or proton attenuation MR imaging signal intensity and hypertrophy within the olivary nucleus.10 There are generally no structural lesions in EPT, and, consequently, the MR imaging is usually normal.10

The pathomechanism generating the rhythmic contractions per se in PT remains unexplained. In the present study, we used functional MR imaging (fMRI) to identify neuronal activations associated with PT in a patient with stimulus-sensitive EPT.

Case Report
A 41-year-old man was admitted to our service because of progressive contractions of the palatal and neck musculature. He reported being in a minor motor vehicle crash without loss of consciousness 2 years prior to admission. A pulsatile tinnitus of the left ear evolved several months later and progressed into a rhythmic ear click. Approximately 2 months before admission, contractions in the palatal musculature occurred. The further spread of the contractions with involvement of the larynx led to a first neurologic consultation. Physical examination showed intermittent bilateral short rhythmic contractions of the palatal and inframandibular musculature, compatible with segmental myoclonus. The segmental myocloni occurred in clusters of 5 rhythmic contractions, followed by a variable silent period. The contractions could be triggered by external sensory stimuli (eg, by touching his left arm). Head movement to the left transiently suppressed the myoclonus. The myoclonus was associated with ear clicks in the left ear and bilateral tinnitus. Otoscopy revealed a high-frequency myoclonus of the M. levator tympani that was not synchronous with breathing. The remainder of the general, psychiatric, neurologic, and otorhinolaryngologic examinations—as well as MR imaging scans of the skull, brain, and brain stem and EEG recordings—were normal. Time-resolved MR projection angiography showed no signs of dural AV fistula. The PT did not respond to any medication, including diazepam and gabapentin in the further course of the disease, which led to the diagnosis of an EPT. The patient took no medication before admission and gave written informed consent before inclusion into the study.

Functional Imaging
Blood oxygenation level–dependent (BOLD) fMRI was performed to take advantage of the stimulus sensitivity of the EPT, by using a whole-body 1.5T MR scanner (Sonata; Siemens, Erlangen, Germany). PT occurred spontaneously or could be evoked by tactile stimulation of the patient’s left arm by a physician inside the scanning room. During the functional scanning, 22 clusters of PT occurred, lasting on average 23.4 seconds (minimum, 5 seconds; maximum, 60 seconds). Functional T2*-weighted images were obtained based on echo-planar single-shot pulse sequence (EPI)—matrix size, 64 × 64; field of view (FOV), 220 mm × 220 mm; 25 sections; 4-mm section thickness; 1-mm gap covering the whole brain; flip angle, 90°; repetition time (TR), 2.5 seconds; echo time (TE), 40 milliseconds; 400 measurements lasting 16 minutes 40 seconds. After functional scanning, a high-resolution 1-mm isovoxel T1-weighted magnetization-prepared rapid gradient echo (MPRAGE) (matrix 256 × 256, 176 sections) was acquired. To confirm the findings of the first imaging session, a second functional scanning was performed 3 days from the point of view of the patient.
later. Appropriate frequency and duration of PT occurred only during the first half of the measurement. Therefore, only the first 200 scans were analyzed, which included 19 clusters of PT lasting on average 9.9 seconds (minimum, 2.5 seconds; maximum, 30 seconds).

Anatomic and functional images were analyzed by using BrainVoyager QX (Brain Innovation, Maastricht, the Netherlands). Preprocessing included 3D motion correction, section-time correction, Gaussian spatial filtering (full width half maximum 4 mm), and high-pass temporal filtering of 3 cycles in time course (removal of low-frequency drifts). The functional images were coregistered to anatomic MPRAGE images. Normalization was not performed. The PT time courses were convolved with a hemodynamic reference function to create basis regressors. All reported activations are based on a fixed-effects general linear model (GLM) of the first run with statistical threshold of $P < .01$ (corrected Bonferroni; corresponding to $t > 5.3$) and extent threshold of 500 mm$^3$. The first run was re-evaluated with additional motion-correction predictors. No motion-related activation was found in the putamen. The results of the second run are not shown.

Peak activation associated with PT was identified in basal ganglia bilaterally and closely resembled the anatomic boundaries of the putamen. Additional activations were present in the precentral gyrus bilaterally and right superior temporal and angular gyrus (Fig 1). The activation in the putamen bilaterally could be reproduced in a second functional scan at day 3. The level of significance was, however, lower because of different alternation of myoclonus clusters and silent periods (not illustrated). No focal activation could be observed in prefrontal motor cortex, the brain stem, or the cerebellum.

**Discussion**

fMRI was used to identify neuronal activations associated with PT in a 41-year-old patient with stimulus-sensitive EPT. PT was associated with peak neuronal activation in the putamen bilaterally. No focal activation was detected in structures of the Guillain-Mollaret triangle, which were previously associated with PT. Additionally, no focal activation could be observed in prefrontal motor cortex, the brain stem, or the cerebellum. The analysis is based on a fixed-effects GLM with statistical threshold of $P < .01$ (corrected Bonferroni) and spatial threshold of 500 mm$^3$. Radiologic convention: right hemisphere is depicted on the left-hand side.

Additional neuronal activations were present bilaterally in the precentral gyrus, which might be attributed to the motor control of the palatine muscles. In line with involuntary activity, no activity was registered in prefrontal areas. Further, activation was present in the right superior temporal and angular gyrus. This activation may reflect auditory stimulation, which occurs simultaneously with PT in the form of audible ear clicks.

Previous fMRI investigations of PT were limited to the brain stem, including the Guillain-Mollaret triangle, and did not cover the putamen in the investigated volume. Dysfunctional activation was found in the dentate nuclei, the left inferior olivary nucleus and the left red nucleus in a 56-year-old patient with PT. This investigation demonstrated dysfunction on the basis of MR images sensitized to changes in cerebral blood oxygenation state. In contrast, the present investigation primarily shows activations. In another functional study, voluntary PT was associated with hyperactivation of the inferior olives, a brain stem region, and the cerebellar dentate nuclei in a brother of a patient with EPT who was able to elicit, modulate, and stop rhythmic contractions of the soft palate associated with ear clicks voluntarily. This voluntary control of rhythmic contractions obviously differs from the patient in the present investigation with involuntary EPT.

One limitation of the present study is that it is only a single case description, so further investigations are necessary to confirm the presented findings. A prerequisite for fMRI is several repetitions of activation interleaved with rest periods within a few minutes, which is rarely encountered in PT. Putative motion-related pseudoactivations are unlikely, because...
of the reproducibility of activations with and without nonexplanatory motion regressors and after 3 days. Also, no significant motion regressor–associated activations were present in the putamen.

**Conclusion**

The present study revealed prominent bilateral activation of the putamen associated with EPT. These results imply a central role of the putamen in the generation of EPT. Further studies will be needed to confirm a putative disinhibition of the putamen in EPT.

**References**