Brainstem reflex physiology

Investigation of the Brainstem Blink Reflex Circuitry in Patients with Juvenile Myoclonic Epilepsy

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Introduction: Juvenile myoclonic epilepsy (JME) which is among idiopathic generalized epilepsies presents with myoclonus and/or generalized tonic-clonic seizures during puberty. Electrophysiological studies showed findings at several levels including cortex, thalamus and spinal cord in JME. Given these information, we aimed to analyze electrophysiological findings at the level of brainstem using blink reflex (BR) and blink reflex recovery cycle (BR-RC).

Patients and method: Eighteen JME patients and age and sex matched 18 healthy volunteers are included in the study. BR and BR-RC with interstimulus intervals of 200 ms and 400 ms were recorded over orbicularis oculi on the nondominant side under the same conditions.

Results: JME patients had longer onset latencies (p=0.046) and higher amplitudes (p=0.022) of R2 component of BR. Recovery of responses after double stimulation with interstimulus interval of 400 ms was higher in JME patients (p=0.040). Use of valproate did not have an impact on reflex responses.

Conclusion: Our results support increased excitability of BR pathway in JME without relevant pathological findings regarding BR.

Brainstem reflex physiology

Ocular Vestibular Evoked Myogenic Potentials to Head Tap and Cervical Vestibular Evoked Myogenic Potentials to Air-conducted Sounds in Isolated Internuclear Ophthalmoplegia

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Objective. The central pathways responsible for ocular vestibular evoked myogenic potentials (VEMPs) to forehead tapping remain to be determined. This study aimed to determine whether the medial longitudinal fasciculus (MLF) carries the signals for ocular VEMPs (oVEMPs) in response to this mode of stimulation.

Methods. Twelve patients with isolated unilateral internuclear ophthalmoplegia (INO) due to brainstem infarction underwent evaluation of the ocular tilt reaction (ocular torsion and skew deviation), tilt of the subjective visual vertical (SVV), cervical VEMPs (cVEMPs) in response to tone burst sound, and oVEMPs induced by tapping the forehead.

Results. Eight (67%) patients showed abnormal oVEMPs that included no wave formation (n=4) and decreased amplitude (n=3) in the lesion side, and bilaterally absent responses in the remaining patient. Furthermore, the patients showed diminished oVEMPs responses in the lesion side compared with normal side (6.0 ± 5.6 vs. 11.7 ± 5.5 uV, paired *t*-test, *p*=0.001) and increased IADamp(%) of the oVEMPs compared with normal controls (43.6 ± 41.2 vs. 9.1 ± 6.2 , *t*-test, *p*=0.018). In contrast, cVEMPs were abnormal in only three (25%) patients, decreased (n=2) or no response in the lesion side. Eleven (92%) patients showed contraversive ocular tilt reaction or SVV tilt.

Conclusion. Patients with INO frequently show impaired formation of ipsilesional oVEMPs in response to forehead tapping. The occasional abnormality and decreased amplitude of ipsilesional cVEMPs also suggest a modulatory pathway for the inhibitory saculocollic reflex descending in the MLF.

Significance. This study suggests that the MLF contains the fibers for the otolith-ocular reflex from the contralateral ear.

Brainstem reflex physiology

Somatosensory and Auditory Startle Reaction in Patients with Movement Disorders

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Objective: Startle reflex (SR) is a generalized defense reaction which is elicited by unexpected stimuli. To elicit SR, auditory and electrical stimulations are used in electrophysiological investigations. Somatosensory startle reflex (SS-SR) is recently described in healthy volunteers after median and tibial stimulation. Studies in patients with brainstem lesions showed importance of upper brainstem in development of SS-SR

Here, we aim to investigate SS-SR systematically in various movement disorders to address its characteristics in comparison to ASR.

Patients and Method: We have examined ASR and SS-SR in patients with dystonia (n=12), multiple system atrophy (n=8), corticobasal degeneration (n=5), restless legs syndrome (n=14), progressive supranuclear palsy (n=11), essential tremor (n=18) and idiopathic PD (n=16) and healthy volunteers (n=35) under the same conditions. ASR and SS-SR were recorded over orbicularis oculi (o.oc), sternocleidomastoid (SCM), and biceps brachii (BB) after bilateral auditory and median nerve electrical stimulations, respectively.

Results: The pattern and probability rates over each three muscle of ASR were similar in MSA, RLS, ET, PD and healthy individuals. Probability rates of responses over each muscle and total ASR probability were the highest in dystonia group whereas they were the lowest in CBD and PSP ($p_{o.oc}=0.016$, $p_{scm}=0.036$, $p_{bb}=0.000$, $p_{total}=0.009$). Onset latencies of O.oc responses were also longer in CBD and PSP groups (p=0.001). Presence of SSR was also the highest in dystonia and lowest in PSP group similar to ASR. Latency of O.oc response was longest in CBD group. Pattern of SS-SR was similar to healthy individuals in all disease groups except dystonia and MSA in which BB responses were more common than SCM responses.

Discussion: The findings of ASR parallel the previous findings. Dystonia patients are known to have exaggerated ASRs. The decreased response in PSP was previously suggested as a support to show extent of pathology. Findings regarding SS-SR also parallel ASR. SS-SR is also exaggerated in dystonia. Different pattern of response appears to be a reflection of overflow phenomenon. However, development of withdrawal reaction is a possibility. Shorter onset latencies in dystonia group probably also reflect increased excitability. Absence of SS-SR or longer latencies in PSP reflects impairment of its pathway which supports the opinion that it may share pathway with ASR and its generator is possibly in the upper brainstem.

Brainstem reflex physiology

Effect of high frequency repetitive transcraneal magnetic stimulation on brainstem excitability in SCI

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Background: We studied the effect of one session of high-frequency repetitive transcraneal magnetic stimulation (rTMS)on brainstem stem excitability in patients with incomplete SCI.

Methods: The study was a randomized, double-blind, sham-controlled trial. We recruited 27 patients with SCI to be randomly distributed in two study groups: active or sham rTMS.

We recorded the blink reflex (BR) induced with electrical stimulation at 15 times sensory threshold (ST) to the supraorbital nerve (SON). A prepulse electrical stimulus on BR (PP-BR), was delivered with ring electrodes to the the right index finger at 2 times ST and applied 100 ms before the SON stimulus. We also examined the excitability recovery curve of the blink reflex (RC-BR), with pairs of stimuli at interstimuli intervals (ISI) of 160, 300, 500, and 1000 ms.

Patients were examined at baseline, and after high-frequency vertex rTMS (20Hz; 40 pulses/burst, 1800 pulses total over 20 minutes).

Results: Fifteen patients received active rTMS and 12 sham rTMS. The age between groups was not significant. At baseline, R2 area of BR and percentage changes (%) in RC-BR at all intervals were similar in both groups. Active but not sham rTMS significantly reduced the area of R2 in BR and decreased the % inhibition with respect to baseline (p≤ 0.05), RC-BR did not change with any rTMS stimulation at any time.

Discussion: High frequency-rTMS over motor cortex induces a decrease in brainstem reflex excitability and reduces inhibitory brainstem activity in patients with SCI.

Brainstem reflex physiology

Startle reflex in neurocritical brainstem patients

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Question: Patients in a neurocritical condition with lesions involving the brainstem cannot cooperate with electrophysiological examination and, therefore, it may be difficult to determine what are the main functions affected. We have used two tests to characterize the deficit in the motor domain, transcranial magnetic stimulation (TMS) to induce motor evoked potentials (MEPs) and loud auditory stimulation (LAS) to induce the startle reflex (SR).

Methods: The study was done in 5 patients who were in their acute/subacute phase after brainstem involvement because of neoplastic tumors (1 cavernoma and 1 meningioma), vascular lesions (1 ictus and 1 haemorrhage) and meningitis with cranial nerve involvement (1 patient). TMS was applied with a round coil or a double-cone coil over the vertex. LAS was applied by discharging the coil flat on top of a metallic platform.

Results: MEPs were obtained in three patients at normal latency while they were absent in the remaining two patients. Attempts of voluntary contraction to facilitate responses did not change the results. The SR was present in two patients (one of them with normal MEPs). Three patients had no startle response. Normal MEPs and SRs were observed only in the patient with meningitis.

Conclusion: There was no clear association between presence or absence of SRs and the standard neurological examination in our patients. Apart from direct lesions in nuclei or tracts, we should also consider alterations in excitability as one of the explanations for absence of SR or MEPs.

Brainstem reflex physiology

Paired neurophysiological and clinical approach to brainstem assessment in Parkinson's Disease.

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Question: Recent research has highlighted the role of brainstem (BS) structures in the early spread of Parkinson's Disease (PD) pathological process. Vestibular Evoked Myogenic Potentials (VEMPs) corresponding to the Vestibulo-Ocular (VOR), Vestibulo-Masseteric (VMR) and Vestibulo-Collic (VCR) reflexes can provide information on BS function. Aims: to test the aforesaid set of VEMPs in a cohort of PD patients and healthy controls and to correlate it with presence of symptoms ascribable to BS dysfunction.

Methods: 19 PD patients (age 66.9 \pm 5.4 years; 12 males; mean disease duration 6.16 \pm 3.54 years) and 15 age and sex matched controls underwent bilateral recording of VOR, VMR and VCR from inferior oblique, masseter and sternocleidomastoid active muscles, respectively. PD patients were additionally administered a series of clinical scales used for evaluation of brainstem-integrated activities, namely sleep disorders (Epworth Sleepiness Scale, Parkinson's Disease Sleep Scale and REM Sleep Behavior Disorder-Screening Questionnaire or RBD-SQ), postural instability (MiniBESTest) and depression (Geriatric Depression Scale). Groups' comparisons were performed with χ 2 test and Mann-Whitney U-test; Sperman's *rho* test was used for correlation analysis.

Results: VEMPs were significantly impaired in patients compared to controls, absence being the main pattern of alteration. When the set of the 3-VEMP battery was analyzed, both number of altered reflexes (p=0.017) and severity of alteration (p=0.001) were significantly higher in patients than controls. As for each single VEMP, only the VOR and the VMR were significantly altered (VOR: p=0.022; VMR: p=0.005; VCR: p=0.056). Clinical scales revealed the presence of some degree of depression in 36.8% of patients, sleep disturbances in 68.4%, REM sleep disorder in 26.3% and postural instability in 36.8% of PD patients. A significant correlation with VEMP alterations was found only for high scores on RBD-SQ (p=0.554; p=0.014).

Conclusions: Combined assessment of VOR, VMR and VCR was able to detect BS dysfunction in a rostro-caudal extension in PD. This may prove interesting in the perspective of identifying neurophysiological markers of BS dysfunction in early stages of the disease.

Brainstem reflex physiology

Somatosensory and Auditory Startle Reflex in Patients with Stroke and Spinal Cord Injury

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Background: Somatosensory startle reflex (SSSR) was recently studied in healthy subjects. Following corticospinal tract lesions caused by stroke or spinal cord injury (SCI), auditory startle reflex (ASR) has been reported to enhance due to reorganization of circuits rostral and caudal to the lesion. To further understand changes in SSSR and ASR, we investigated both responses in patients with spinal cord injury (SCI) and stroke.

Methods: We examined characteristics of ASR and SSSR in 14 SCI and 40 stroke patients (16 brainstem and 24 cerebral hemispheric infarctions) and 39 age and gender matched healthy subjects. ASR was obtained after eight auditory stimuli and SSSR was elicited after median nerve stimulation at the wrist. Surface electromyographic recordings were obtained from orbicularis oculi (O.oc), sternocleidomastoid (SCM), biceps brachii (BB) and abductor pollicis brevis (APB) muscles.

Results: Total ASR probabilities at distal muscles were significantly higher in patients with SCI and in stroke patients especially with brainstem infarctions. Similarly SSSR rates were increased in both patient groups compared to controls (for APB p<0.05). SSR latency of O.oc in SCI patients were similar to controls whereas it was markedly prolonged in stroke patients (p<0.05).

Conclusion: In conclusion, we have found that SSR and ASR were enhanced in stroke and SCI and this enhancement was more prominent in distal muscles. Secondly, the properties of ASr and SSSR differed according to the lesion site.

Brainstem reflex physiology

An excitatory reflex between R1 and R2 responses of the blink reflex to supraorbital nerve stimuli.

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The typical pattern of the blink reflex to ipsilateral electrical stimuli to the supraorbital nerve (SN) is composed by an early, short duration, well synchronized, R1 response and a late, long duration, polyphasic R2 response. We examined whether the period between R1 and R2 results from lack of excitatory inputs or active inhibition.

Tests were performed in 5 healthy subjects and 10 patients of various disorders affecting blink reflex excitability, including postparalytic facial syndrome and hemifacial spasm. All subjects wore surface electrodes attached to the orbicularis oculi and stimuli were applied to the SN. After obtaining the conventional blink reflex, healthy subjects were requested to close their eyes at various levels of force while applying again the same stimulus.

In healthy subjects closing the eyes with force, SN stimulation induced a response of smaller amplitude and shorter latency than the R2 (25.2 +/- 1.3 ms). A short and incomplete silent period emerged after R1 and after R2. Patients with postparalytic facial syndrome and essential hemifacial spasm showed activity interfering with the silence between R1 and R2, with often expanding between the two responses, repeating in successive trials.

Our results show that there can be reflex responses between R1 and R2 in healthy subjects during contraction and in patients with abnormal reflex excitability. This suggests that the absence of activity between the two responses in normal conditions is the consequence of an active inhibition that reaches the facial motoneurons at that point.

Brainstem reflex physiology

Brainstem Reflexes in Patients with Sleep Bruxism : Masseter Inhibitory Reflex Responses and Auditory Startle Reaction

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Ouestion: Sleep bruxism (SB) is a stereotyped movement disorder characterized by grinding or clenching of the teeth during sleep. Neurophysiologic methods offer approaches to study the excitability in SB by recording the motor potentials evoked by masseter muscles and masseter inhibitory reflex (MIR). Auditory startle reaction (ASR) has been found to be increased in restless leg syndrome but it has not been studied in SB. We aimed to show the hyperexcitability in the central jaw motor pathways in patients with SB by studying MIR and ASR responses

Methods: 10 patients (mean age $48,4\pm 10,2$ years; 8 male and 2 female) and 20 healthy subjects (mean age $34,4\pm 11,2$ years; 6 male and 14 female) were enrolled in the study. The EMG activity was recorded from two masseter muscles by surface electrodes. The duration and latency of early and late silent periods and suppression ratios obtained from patient and control groups were compared. ASR was recorded from o. oculi, masseter, sternocleidomastoid, biceps brachii and tibialis anterior muscles and probability and latencies were evaluated. SPSS for Windows 11.5 package was employed

Results: The differences in silent period onset latencies between patients and normal subjects were not significant but durations of early and late silent periods were shorter in SB group (p

Conclusions: The durations of the early and late silent periods were significantly shorter in SB group. The absence of late silent period in 2 of 10 patients and its presence in all normal subjects is noteworthy. These results may indicate a reduced capacity for inhibition by the circuits responsible for the late period in SB and this increased excitability in central motor pathways could derive from an impaired modulation of subcortical structures especially brainstem inhibitory circuits and not from altered cortical mechanisms. Since ASR did not differ between the control and patient groups, it can be possible to say that there's no abnormality in brainstem and reticulospinal circuits in SB patients.

Brainstem reflex physiology

Comparison of brainstem reflex abnormalities in patients with Multiple Sclerosis, Behçet and Stroke and its topodiagnostic value

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Ouestion : Our current understanding of brainstem reflex physiology comes chiefly from the classic anatomicalfunctional correlation studies that traced the central circuits underlying brainstem reflexes and establishing reflex abnormalities as markers for specific areas of lesion. Our aim is to investigate the correlation between the brainstem reflex abnormalities and lesion localization in three different diseases with brainstem lesions.

Methods: Masseter inhibitory and blink reflexes were studied in 21 multipl sclerosis, 15 stroke and 17 Behçet's disease patients with brainstem lesions on brain magnetic resonance imagings. The reflex responses were compared with those of 20 healthy subjects of comparable age . SPSS for Windows 11.5 package was employed for statistical processing.

Results: Latency abnormalities of the blink and the masseter inhibitory reflexes were the most prominent in MS group. No significant differences were found in duration and degree of suppression among the groups in MIR responses. The highest abnormality percentages in MS group were in R1component of blink reflex (% 71, 4) and S2 component of masseter inhibitory reflex (% 90,5). And also R1 of BR and S2 of MRI were the most abnormal responses in the all groups compared to other parameters

Conclusions: Distinct reflex abnormalities indicate lesions at specific sites. A number of lesions suspected on clinical data may be confirmed by reflex findings only and not by imaging studies. Reflex testing can be utilized to demonstrate multiple lesions and evaluate dissemination of central nervous involvement in patients with brainstem lesions. Increased abnormality percentage in MS group may be explained by the fact that supratentoriel lesions affect brainstem reflex responses. Since lesions in Behçet's disease are localised in mesencephalon commonly, superior to MIR and BR circuits, the abnormality in this group is much less relatively.

Brainstem reflex physiology

From PET-MR Fusion Imaging to Tractography

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From PET-MR Fusion Imaging to Tractography

The last 4 to 5 years has been one of the most exciting periods for Ultra High Field (UHF) MRI research and development, especially for 7.0T.

As many expected, some of the most exciting results came from anatomical and tractographic studies, much owed to the enhanced T2 images as well as T1 images at high field. We now can routinely visualize many small tractographic image, such as the medical forebrain bundle.

Another new progress has been in the area of PET and MRI fusion molecular imaging with UHF MRI. This area, we have also made substantial progresses, such as the visualization of the brainstem raphe nuclei hitherto unable to visualize.

Brainstem reflex physiology

Startle reaction evoked by kinematic stimuli

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1.Question

Kinematic stimuli are used for both assessment and treatment in neurorehabilitation. A patient's voluntary or reflex response may be affected by a startle reaction. We therefore explored whether certain kinematic stimuli are able to elicit a startle reaction.

2. Methods

Eleven healthy subjects were suspended in a Lokomat system and were exposed to unexpected passive left knee flexion at 3 velocities (6, 60, 240°/s). Subjects were asked to perform a right wrist extension as soon as they felt their leg move (conditions: 6-React, 60-React, 240-React, respectively). In some 240°/s trials movement onset was preceded by a low-intensity electrical pre-pulse to the left index finger (240-Prep-React). We recorded EMG activity from right orbicularis oculi and sternocleidomastoid muscles to assess startle responses, from left quadriceps muscle to obtain stretch reflexes, and from right wrist extensors to assess reaction time.

3. Results

Startle responses were present in most 240-Reacttrials, as evidenced by (1) EMG activity in orbicularis oculi and/or sternocleidomastoid, (2) significant reaction time shortening in wrist extensors, and (3) stretch reflex latency shortening in quadriceps, as compared to responses without startle reaction. Only few trials at lower angular velocities resulted in startle responses. In 240-Prep-React trials no startle responses ocurred.

4. Conclusions

Kinematic stimuli of high angular velocity, used to assess muscle stiffness, may elicit a generalized startle reaction, which in turn may modulate stretch reflex latencies of the muscle tested in a passive movement paradigm.

Motor control

Decreased Spreading Depression Susceptibility in Parkinson Rat Model

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Question:Parkinson disease (PD) is known by a major loss of dopaminergic nigrostriatal neurons and by an increased turnover of neurotransmitter by surviving neurons of the nigrostriatal tract. The clinical diagnosis of PD is based on the identification of some combination of the cardinal motor signs of bradykinesia, rigidity, tremor, and postural instability. Spreading depression (SD) known as an evoked neuronal activity and changes in ionic, metabolic and hemodynamic characteristics of the brain. Pronounced release of dopamine during SD and the probable role of dopamine in SD process suggests that disruption of dopaminergic pathway in PD may cause SD to behave differently.

Methods:To test this possibility, we induced dopaminergic lesion by bilateral intracerebral stereotactic injection of 6 µL of 6-hydroxydopamine in the medial forebrain bundle (MFB). After 4 days, SD was induced by the injection of 3M KCI and SD propagation was followed using two ion-sensitive microelectrodes placed in the parietal and occipital cortex.

Results:Eliciting SD in rat model was associated with a significant increase in the threshold of SD and a decrease in the propagation velocity and duration of accompanying extracellular DC changes.

Conclusions: The present data show that rat model of Parkinson's disease are less prone to SD.

Motor control

Preparation for Reaction Time Tasks in Multiple Sclerosis

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Question

Sensorimotor integration can be assessed using simple reaction time tasks to somatosensory stimuli (ssRT). Tasks performed to stimuli applied to the contralateral hemibody would reveal interhemispheric transfer of information (IHT) accross corpus callosum, while tasks performed when a startling-auditory stimulus (SAS) is presented together with the somatosensory cue would reveal the level of subcortical motor preparation. We studied both preparation and IHT in MS patients in whom callosal pathways and subcortical structures may be particularly involved.

Methods

In 13 controls and 20 mildly disabled relapsing-remitting MS patients, we studied ssRT for voluntary wrist-extension to ipsilateral and contralateral low intensity electrical stimuli, applied to the index finger. Percentage delay (PD) of contralateral vs ipsilateral responses was calculated as a measure of IHT. StartReact trials were performed applying randomly a SAS in ssRT trials (20%). Percentage shortening (SAS-Short) with respect to ssRT was calculated as a measure of the effectiveness of preparation. Motor conduction time (MCT) was assessed by recording the motor-evoked potentials to TMS.

Results

In controls, mean PD of contralateral responses was 105.7% longer than for ipsilateral responses. In StartReact trials, the mean SAS-Short was 29.7% for ipsilateral and 23.3% for contralateral responses. With respect to controls, patients had no differences in PD (105.4%) but showed significantly lower SAS-Short for ipsilateral (23.4%;p=0.02) and contralateral trials (10.9%;p=0.005). Contralateral SAS-Short was inversely correlated with MCT (r=-0.294). Two patients had abnormally delayed orbiculari-oculi responses to SAS.

Conclusions

Patients with MS have defective subcortical motor preparation that is worse when IHT sensorimotor integration is required.

Motor control

Essential Tremor, the Olivocerebellar System and Motor Timing- an fMRI study

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Question:

Essential tremor (ET) is the most common tremor disorder. It has been repeatedly shown that the olivocerebellar system is involved in ET, consisting of the inferior olive nucleus (ION), dentate nucleus (DN) and cerebellar cortex.¹ Impairment of the central timing mechanism in ET has been suggested.² The olivocerebellar system plays an important role in motor timing.³ In order to study the functional involvement of this system in ET, we employed a task involving motor timing during functional MRI (fMRI).

Methods:

Thirty propranolol sensitive ET patients with familial upper limb tremor and 30 healthy controls were included. T2*weighted EPI sequences were acquired (180 volumes, TR: 2s, voxel size: 3.5x3.5x3.5mm, 3TMRI). The task consisted of alternating rest and finger tapping (rate of 2Hz) blocks. Task performance was measured by electromyography. Analysis was performed in SPM8 (standard preprocessing, normalization according to the spatially unbiased infra-tentorial template⁴, 4mm smoothing kernel). The left ION was localized with a conjunction analysis of patients and controls. For this abstract, quantified tapping performance is not yet incorporated.

Results:

Within-group random-effects analysis restricted to the brainstem and cerebellum showed activations throughout the cerebellum (bilateral lobules V, VI, VIIIA) and in both left and right ION. Preliminary between-group analysis of the ION and CB right lobule VI showed significantly higher activations in controls compared to ET patients (figure 1).

Conclusions:

Exploratory analysis suggests altered activity in the olivocerebellar system in ET patients compared to controls during a motor timing task. Inclusion of task performance and effective connectivity analysis will be carried out to aid us in discriminating between altered activity due to task performance and pathological motor timing related to ET. Understanding the role of the olivocerebellar system in ET is of great importance for unraveling the mechanism of tremor genesis.

figure 1



Motor control

Correlation of tau pathology in eye movement related brainstem nuclei in cases of progressive supranuclear palsy (PSP) and a proposed role of perineuronal nets

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Background: Progressive supranuclear palsy (PSP) is characterized by hyperphosphorylated tau protein (HTP) accumulations in different brain regions. A hallmark of clinical symptoms is a vertical gaze palsy, which may be accompanied by horizontal gaze palsy, whereas the VOR is rather preserved until late in the disease. Based on monkey studies functional cell groups of the oculomotor system including transmitters have been identified in the human brainstem¹.

Methods: To study the spread of the disease through the oculomotor premotor network sections of 8 archival PSP cases with different eye movement deficits were immunostained for HTP. The analysis involved the glutaminergic rostral interstitial nucleus of the medial longitudinal fascicle (RIMLF) and paramedian pontine reticular formation (PPRF), the glycinergic saccadic omnipause neurons (OPN), the cholinergic oculomotor nucleus (nIII) and the vestibular nuclei (VN). Since the RIMLF and nIII contain well-developed perineuronal nets (PN)², we further stained sections of the PSP cases for the presence of aggrecan (ACAN) and HTP to prove a possible protective role of PNs as suggested from Alzheimer's disease cases³.

Results: All PSP-cases showed HTP-staining in neurons and glia of RIMLF, PPRF and OPNs, in late cases in motonuclei as well. Thereby a correlation between eye movement deficits and degeneration of premotor gaze centers and the complete pathways to motoneurons was found, whereas the VN are less affected.

The analysis of PNs revealed that ACAN-based PNs were mainly found around HTP-negative neurons, and only few HTP-positive neurons showed weak ACAN staining.

Conclusion: The early and more severe degeneration of premotor centers compared to nIII support a hypothesis that in PSP the pathology progresses along neuronal chains in an anterograde fashion, and it is not confined to specific transmitters. The severe tau pathology in RIMLF and nIII in late PSP cases implies that the presence of PNs may not protect them from degeneration, but on a cellular basis the integrity of PNs is affected by tau-pathology.

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Motor control

Oscillatory activity in primate reticular formation

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Piper rhythmicity (35-60 Hz) has been reported in EMGs since 1907. These oscillations are often assumed to originate in the cortex, as they are coherent with sensorimotor EEG (Brown et al 1998). However, we have recently seen similar oscillations in a slice preparation of primate brainstem reticular formation. These oscillations were decreased by the GABA-A antagonist bicuculline, and completely suppressed by the GAP junction blocker carbenoxolone, suggesting generation by a complex intrinsic network. Given our recent work showing the importance of the reticular formation for recovery after corticospinal tract lesion (Zaaimi et al 2012); we were interested in whether such oscillations might become more visible during functional recovery.

Macaque monkeys were trained to pull a lever, which opened a door allowing them to retrieve a food reward. In two animals, after training we made a unilateral lesion of the pyramidal tract by thermocoagulation, and allowed recovery, which reached plateau after ~ 4 months. The recovery differed between the two lesioned monkeys: one was moderately impaired and could still take the food with the paretic hand, whilst the second lost the ability to perform precision grip but was still able to pull a lever. Recordings were then made of local field potential (LFP) from the reticular formation and EMG. Recordings from one unlesioned animal served as a control. In the control and in the less impaired monkeys reticulomuscular coherence was observed in the beta band (20-30Hz); this was not seen in the more impaired monkey. By contrast, the more impaired animal showed ~70Hz oscillations in reticular formation LFP around the time of movement. These preliminary results suggest that intrinsic rhythmicity around 40-60 Hz is present in the reticular formation and could be unmasked after lesion of pyramidal tract.

Motor control

Modulation of force and velocity for accurate tasks by a startle.

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1. Question

Some skilfulfast activities require control of trajectory and force to fulfill the task. It is known that voluntary reactions can be speeded up by a startling auditory stimulus (SAS) delivered at the same time as the imperative signal (IS), a phenomenon termed Start-React (Valls-Solé et al., 1999). The aim of the present study was to examine how the performance of a task that demands accuracy, mainly in terms of velocity during the trajectory or force control when holding an object, is changed by a SAS timed to occur either at the time of IS or at a pre-determined time interval when movement has already begun and motor control of accurate reaching is expected.

2. Methods

Nine subjects performed an accurate elbow extension between two spots placed at an angular distance of 30 degrees when holding with their right hand a pen that monitored, through two strain gauge systems, the force of the subject's fingers during the hold, and the force at which pen-tip was exposed with table contact. According to block of trials, this end spot had three possible diameters: 5, 10, and 20 mm. Kinematic and force parameters were measured for three conditions, control, with SAS at IS or SAS delivered when movement has already started.

3. Results

In SAS trials, there were no evident differences with respect to control trials in the general outline but there were differences in the timing of the events. In SAS-IS trials time to peak velocity was 166 ms (SD= 41 ms), time to peak force was 95 ms (SD= 43 ms) both shorter than in control or SAS delivered when movement began. Peak velocity (315±56 $^{\circ}$ /s) and peak force (8±2 arbitr. units) were larger for SAS-IS trials than for control or SAS delivered when movement started.

4. Conclusions

Startle can advance a task that requires accuracy with velocity and force modulations, but when task onset is already launched there are not evident changes in velocity or force parameters, that seem already prepared.

Motor control

Is a multi array-contact lead able to improve STN-DBS in Parkinson's Disease?

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Introduction: Multi-directional electrodes in STN-DBS give the possibility to steer the high frequency current into the direction for clinical improvement and moreover recording of multi-array local field potential recordings (LFPs) give the possibility to determine the spatiotemporal distribution of oscillatory power across the STN that is related to the symptomatology in PD patients. Ultimately this information can be used as a feedback to optimize the stimulation parameters.

Methods: Eight PD patients were included in the study. After micro-electrode recording (MER), to determine the boundaries of the STN, temporarily a prototype of a new 32-contact DBS lead was inserted during the DBS procedure. Different modes of current steering were employed to improve motor symptoms and patients were scored on beneficial and adverse effects. Prior to and immediately after stimulation the distribution of spectral power was recorded by LFPs and spatiotemporal changes were evaluated with Fourier analysis.

Results: Thresholds of the effects of spherical stimulation were comparable to thresholds of stimulation through the conventional electrode at the same location in 89% of the cases. In eight of fourteen side effects, steering stimulation current increased the threshold for side effects by ≥1mA compared to spherical stimulation. The size of the therapeutic window could be widened in patient 6, 7 and 8 by steering stimulation in the posterior, posterior and anterior direction, respectively. LFP recordings prior to stimulation yielded the boundaries of the STN by showing increased spectral power particularly in the 13-40 Hz range. Recordings across all directions showed distinct spatiotemporal patterns of neuronal activity, which were related to the pattern of stimulation and the PD symptoms of the patient.

Conclusions: With a new DBS lead it is possible to steer stimulation current in STN-DBS such that it leads to a larger therapeutic window than with conventional spherical stimulation. Simultaneous LFP recordings across the entire STN provide spatial information about the location of the STN and its disease-related electrical activity. This may potentially be of benefit in predicting how to steer the current towards the sensorimotor part, while avoiding adverse effects.

Motor control

Parietal transcranial direct current stimulation affects primary motor cortex excitability in humans.

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The posterior parietal cortex is part of the cortical network involved in motor learning and is connected with the primary motor cortex, as shown in animal models, and experiments in humans. Neuroplastic alterations of neuronal connectivity are an important basis for learning processes. These have however not been explored for parieto-motor cortical connections. We aimed to explore plastic alterations of these connections via non-invasive brain stimulation with transcranially applied direct currents (transcranial direct current stimulation, tDCS) in healthy humans. Thirteen subjects received anodal, cathodal and sham tDCS over the left posterior parietal cortex (P3) for 15 min (stimulation intensity 500 mA, electrode size 15 cm²). As a control, they also received anodal and cathodal tDCS over the left parietal cortex but 3 cm posterior or lateral to P3. Subsequent neuroplastic changes of the excitability of the ipsilateral M1 were monitored via TMS single pulse-induced motor evoked potentials (MEP) applied before, immediately after the stimulation, and every five minutes (0, 5, 10, 15, 20, 25, 30 min), then every 30 minutes (60, 90, 120 min). The results show tDCS-polarity-dependent alterations of motor cortex excitability after parietal tDCS. MEPs were enhanced by anodal, but reduced by cathodal tDCS. For anodal tDCS, these effects lasted for at least 120 min after tDCS. These results hint to plasticity of interregional connectivity between P3 and M1 in the human cerebral cortex. Further studies are necessary to explore the functional relevance of these effects.

figure 1



Motor control

Acrylamide Disrupts the Ontogeny of Neurobehaviour in Albino Rats

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The present study aimed to elucidate abnormalities in the ontogeny of sensorimotor reflexes in developing rats after prenatal and perinatal acrylamide or saline intoxication of pregnant rats. Acrylamide was used as an experimental probe to investigate neurobehavioural and morphological changes in developing rats after administration of the toxin to pregnant mothers. Acrylamide was administered to non-anaesthetised pregnant rats by gastric intubation at a dose of 10 mg/kg/day. Rat pups were assigned to one of three groups: Group A, which comprised pups whose mothers were treated with saline (control group); Group B, which comprised pups whose mothers were treated with acrylamide from day D7 of gestation to birth (prenatal intoxication); and Group C, which comprised pups whose mothers were treated with acrylamide from D7 of gestation to D28 after birth (perinatal intoxication). This study has been conducted recently in Beni-Suef University (Egypt) and Kig Saud University (Saudi Arabia) by May 2012. Acrylamide-induced morphological changes (CNS aberration) and neurobehavioural changes (sensorimotor reflex retardation) were studied. The reflexes tested included rooting, forelimb (FL) grasping, hind limb (HL) grasping, surface body righting, air body righting, FL hopping, HL hopping, chin tactile placing and visual placing. These reflexes were tested in newborns in all groups from postnatal day 2 (D2) until reflex maturation. The appearance of select external features was recorded. administration of acrylamide in pregnant albino rats disrupts the ontogeny of sensorimotor reflexes and morphological changes in the CNS of developing albino rats.

Other

EARLY ABNORMAL SPONTENEOUS ACTIVITY FINDINGS IN AXONAL VARIANT OF GUILLAIN BARRE SYNDROME: Is this a new variant of Guillain Barre Syndrome.

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Objective: To evaluate the earliest denervation potentials in axonal variant of GBS.

Background:Guillain Barre Syndrome (GBS) is one of the most important acute neurolological emergency. Denervation potentials on needle EMG is the hallmark of axonal damage. Usually this is a time dependent phenomenon. The degeneration of axon depends upon length of axon to be degenerated. Studies claim variable time duration for denervation potentials from two weeks to three weeks.

Material/Methods: This is a cross-sectional survey of patients admitted and referred for neurophysiologic assessment. Clinical and neurophysiological data of GBS patients over a period of three years and three months and ten days was collected. NCS/EMG performed by a qualified neurophysiologist. Diagnosed cases of GBS with available data of NCS/EMG were included. Patients with history of Diabetes Mellitus, previous history of any sort of neuropathy and demyelinating variants after diagnosis were excluded as well. Clinical and Neurophysiologic data were collected on Performa for analysis.

Result: Total forty three patients were diagnosed as GBS and those with axonal variants were finally included. Out of forty three, eighteen had axonal variant of GBS and rest of them demyelinating variants. Twelve patients of axonal variant (31%) showed fibrillation potentials, positive Sharp Waves and increased insertional activity within 4-12 days of symptoms onset and six (69%) beyond that period. Total twelve patients were finally included. Active denervation in the form of fibrillation potentials and positive sharp waves were noted frequently and decreased interference pattern in almost all patients. NCS were performed before EMG examination.

Conclusion: Fibrillation Potentials, Positive Sharp Waves and decreased interference pattern were noted in early course of disease in GBS patients interestingly before two weeks of symptoms onset. This study raises the query for a possible new Hyperacute or Fulminant variant of GBS. These findings need further histopathology and etiologic correlation as well as further prognostic importance.

Key words: GBS GuillainBarre Syndrome, NCS Nerve Conduction Study, EMG Electromyography.

Other

PREVALENCE OF NEUROPHYSIOLOGIC DISCOMFORT AMONG EXTENSIVE AND NORMAL CELLULAR PHONE USERS

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The neurophysiologic discomfort (NPD) appears as a side effect of certain drugs or stress and its analysis is very complex. May the extensive or normal cellular phone usage be a reason of NPD? In this study, an attempt has been made to find the risk of NPD among normal and extensive cellular phone users.

Methodology: Information gathering chronological model (Kumar et.al. 2012) was applied to collect the information among cellular phone users. Following the modified methodology of Interphone study (2008) and Balikci et al.(2005), Prevalence of NPD among 659 CP users was analyzed within 4 individual groups i.e., LU, NU, MU and HU. The NPD was self reported by 33 CP users.

Results:

The cellular phone users belonging to LU, NU, MU & HU were 217 (32.9%), 140 (21.2%), 209 (31.7%) & 93 (14.1%) respectively. The overall 33 CP users were reported NPD and its association with individual groups were found in increasing order except HU. The significant association was not found within these individual groups in overall analysis for NPD. Within the individual groups of the male CP users, there were no significant association found for the NPD symptom but a trend in female subjects of NU (p=0.096) and MU (p=0.066) group was observed when compared to LU of female population. None of the children participants in this study were found to have the NPD symptom.

Conclusion: Conclusively, cellular phone's usage is not a cause of NPD among extensive and normal users but in females, a trend for the risk of NPD was observed among those belonging to NU and MU group when compared to LU.

Other

Analyses of the changes in Electroencephalogram induced by Working Memory tasks

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Introduction: The working memory, essential cognitive function and with limited capability, allows temporary storage and the manipulation of information used in complex cognitive tasks such as language, problem solving, understanding of new instructions and reasoning. In turn, the Fluid Intelligence (IF) is the ability to engage and respond to new situations, regardless of previously knowledge acquired, evoking logical reasoning and concept formation.

Objective: This study aimed to verify the brain areas activated during the execution of working memory and fluid intelligence tasks, determining the variations at the level of electroencephalography activity (EEG quantitative (qEEG)).

Sample: Belonged to the sample 31 students of the Technology and Health School of Porto (ESTSP), of both sexes, ages between 18 and 25 years, right-handed, with no psychiatric or neurological problems.

Methodology: Have been made an EEG to each participant, during which they were subjected to two tests of working memory (Direct Evocation of Digits (DED) and Inverse Evocation of Digits (IED)) and to one task of fluid intelligence (Raven Matrices). Afterwards, proceeded to the analysis of qEEG at different times of the tasks.

Results: Have been observed statistically significant differences in the comparison between the beginning and end of both the DED and IED, as the Raven Matrices, and the relationship of these tests to the baseline record. In fact, in a more specific analysis of the data, have been verified the activation of frontal and parietal areas in the WM tasks and a fronto-parietal network in IF tests, as well as a decrease in EEG power right parietal alpha, in other words, an exacerbation of the alpha band in this region in 4 of the 5 series of RAVEN matrices when compared to baseline.

Conclusion: It was shown that during WM tasks are exacerbated prefrontal and parietal regions, while the testing of IF evoke the activation of a fronto-parietal network, occurring variations of power in delta, theta, alpha and beta bands. It was found that the qEEG is useful in the evaluation of electroencephalographic changes in cognitive tests for this sample and people group. However, the possible association with other types of tests may supplement data, making them more accurate in terms of the veracity of results and reproducibility of tasks.

Keywords: Working Memory, Fluid Intelligence, Direct Evocation of Digits, Inverse Evocation of Digits, RAVEN Matrices, qEEG.

Other

The role of auditory brainstem response in diagnosing auditory impairments of Dejerine-Sottas

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Dejerine-Sottas disease is classified as hereditary motor sensory neuropathy (HMSN) type III and shows evidence of Friedreich's ataxia, significant reduction in nerve conduction velocity (NCV), hypomyelination and demyelination of the nerve fibers.

In this study, a 10-years-old girl with Dejerine-Sottas disease is presented in which routine clinical signs (ataxia and reduced NCV) are seem with significant impairments of auditory brainstem pathway. It is indicated that pure tone audiometry and standard tympanometry (tympanometry and acoustic reflex) showed normal results in both ears (normal peripheral auditory system). In contrast, auditory brainstem responses (ABRs) indicated abnormal findings in absolute latencies of I, III, and V and inter-peak latencies of I-III and I-V. These findings suggested auditory brainstem involvement especially in low and mid regions.

Based on our evidence, ABR test is very important for revealing any impairment(s) in auditory brainstem pathway of patients with Dejerine-Sottas.

figure 1



Other

Visualization of human brainstem substructures using gray matter nulling 3D-MPRAGE at 7 Tesla

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Question

The human brainstem is one of the most complex neural entities, in both, structural and functional domains. Its plurality of small and densely packed substructures results in demanding settings for in-vivo MR visualization. We addressed this issue applying ultra-high field MRI at 7 Tesla focusing on enhanced differentiation of brainstem substructures. This approach is an essential prerequisite breaking down the complex brainstem anatomy.

Methods

A modified 3D-MPRAGE-Sequence in the sagittal and transverse plane was applied in a 7T MR-system. To compensate for B1-inhomogeneities, a highly adiabatic inversion prepulse was implemented. Five healthy volunteers were examined. The resulting images were visually correlated to histological plates and post-mortem MRI images from Duvernoy's Atlas.

Results

We identified 22 structures (sagittal), clearly assignable to histology data. In the brainstem 12 substructures were manually outlined for all five subjects, also based on histology. Adopting histology nomenclature clearly identifiable structures were labeled (Figure 1 - 2).

Conclusions

Optimized 3D-MPRAGE-Imaging in the gray matter nulling regime at 7 Tesla provides enhanced image contrast between substructures in the human brainstem, giving rise to better understanding anatomy and potentially related pathology.

References

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Legend Figure 1:

- a 7T in-vivo gray matter nulled sagittal MPRAGE images.
- **b** Sagittal histological section from the Duvernoy's Atlas.
- 19 Spinal trigeminal nucleus (CN V)
- 20 Lateral cuneate nucleus
- 21 Inferior olivary nucleus
- 26 Pons (corticospinal tract)
- 28 Medial lemniscus
- 29 Superior cerebellar peduncle
- 30 Oculomotor nerve (CN III)
- 31 Substantia nigra
- 32 Red nucleus
- 33 Inferior colliculus
- 34 Superior colliculucs
- 36 Prechiasmal optical nerv
- 39 Anterior commissure
- 42 Dorsomedial thalamic nucleus
- 43 Centromedian thalamic nucleus
- 44 Pulvinar
- 49 Basilar artery

- 60 Quadrigeminal cistern
- 61 Splenium
- 62 Premedullary cistern
- 63 Cerebellomedullary cistern
- 98 Zona incerta



Legend Figure 2:

Four 7T in vivo axial slices through the Brainstem showing the olivary nucleus (dotted circles) detailed.

Other

Shunting during stapes surgery

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Question. The possibility to use intraoperative shunting of tympanum is reflected in this work when performing stapedoplasty for prevention of cochleovestibular dysfunction for patients with otosclerosis.

Materials and Methods. Under our supervision there were 37 patients with otosclerosis aged from 28 till 59. 19 people- were the main group of patients that were installed a tymphanostomy tube during the stapes surgery for 7 days and 18 patients- control group of patients with classic piston stapedoplasty with a titanic artificial limb. Examination of vestibular and acoustical functions was carried out before and after the surgery. Vestibular function was registered and evaluated with the help of video occulography, acoustical function was evaluated by occulometry and by threshold voice-frequency audiometry (in Hz) and intensity of noise was analyzed by audio-noise meter.

Results. In a main group of 19 people: the air-bone gap (ABG) for 18 patients did not exceed 10-15 dB. On the control audiograms (p>0,05) on early stage after the surgery (7th day). Subjective feelings of violation of vestibular functions (dizziness, nausea, balance disorder) on the early stage after the surgery (1st day) had 2 patients. The speed of a slow component of spontaneous nystagmus on the 1st and 3rd day after the surgery changed from 0,05 °/s till 13,7 °/s with frequency from 0,1 Hz till 2,7 Hz, at reference average values 0.1- 1.8 °/s and frequency 0.1-1.9 Hz on 7th day came nearer to reference values or did not change at all (p>0,05). Tonality change and decrease in intensity of noise was noted in both groups (p>0,05).

Conclusions. Shunting of tympanum cavity during stapes surgery promotes passage of wound detaches, supports adequate function of an acoustical pipe. Air-bone gap is reduced on early stages, the risk of development of negative cochleovestibular symptomatology decreases.

Other

Assessment of brainstem reflexes improves the diagnostic sensitivity of multimodal evoked potentials, MRI and clinical testing in the investigation of brainstem function in multiple sclerosis

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Question

Brainstem (BS) functions are conventionally studied by multimodal evoked potential (EP) recordings, MRI and clinical examination (CLIN). In Multiple Sclerosis (MS), increasing evidence accounts for a BS involvement, often undetected by standard testing. Recently, brainstem reflexes (BSRs) have drawn attention in evaluating BS dysfunction in MS especially the vestibulocollic (VCR) and vestibuloocular reflexes. In contrast, the vestibulomasseteric (VMR), acousticmasseteric (AMR) and trigeminocollic (TCR) reflexes have never been studied systematically in MS. Aims: to investigate whether the diagnostic sensitivity of CLIN, EPs and MRI can be improved adding the assessment of VMR, AMR, TCR and VCR either as single reflexes or in a 4-BSR battery.

Methods

The 4-BSR battery was recorded in 60 patients (33.3±8.3 yrs) with relapsing-remitting MS (illness duration 8.2±6.4 yrs). EP set included standard BAEPs, median and tibial SEPs Conventional MRI scans were focused on the BS lesion load. Group differences and correlations between variables were analysed with Mann-Whitney U test and Mc Nemar test.

Results

Distribution of BSR and EP abnormality frequencies in MS was: VMR 62.1%, AMR 55.1%, TCR 58.6%, VCR 25.9%; BAEPs 37.3%, median SEPs 60.3% and tibial SEPs 58.6%. Overall, BS dysfunction was detected as follows: BSRs 86.9%, EPs 82.7%, MRI 71.7%, CLIN 37.7%. While the performance of BSRs and EPs, taken separately, was not significantly higher than that of combined MRI/CLIN testing (70%), the paired use of BSRs/EPs had a sensitivity of 93.3%, which was significantly superior (p=0.007), in a subset of patients with a disease duration \leq 6.4 yrs.

Conclusions

BSRs revealed brainstem lesions otherwise undetected by CLIN and MRI, thus providing additional evidence of BS dysfunction in MS. Noteworthy, BSRs could effectively complement the usual EP testing in early detection of clinically and radiologically silent lesions. This may encourage EPs/BSRs paired use in newly diagnosed.

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Other

Longitudinal assessment of brainstem reflexes in multiple sclerosis compared to multimodal evoked potentials, MRI and clinical evaluations

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Question

We have previously shown in patients with relapsing-remitting Multiple Sclerosis (MS) that: *i*) the vestibulomasseteric (VMR), acousticmasseteric (AMR), trigeminocollic (TCR) and vestibulocollic (VCR) reflexes are able to spot brainstem (BS) dysfunctions undetected by clinical and MRI examinations; *ii*) the combined use of these Brainstem Reflexes (BSRs) with multimodal Evoked Potentials (EPs) is more valuable than each single test in the early years after onset. Our aim was to document BS changes over time by BSRs, EPs, MRI and BS signs/symptoms (CLIN) before and after at least one year follow up, in MS.

Methods

Forty-five MS patients (34.8±8.6 yrs old; disease duration 8.9±6.6 yrs) underwent BSRs, EPs (namely Brainstem Auditory Evoked Potentials - BAEPs, median and tibial Somatosensory Evoked Potentials -mSEPs and tSEPs), MRI and CLIN examination. BSR and EP data were ranked and summed up to obtain a cumulative score expressing severity of neurophysiological impairment. Before-after changes were tested with Wilcoxon test.

Results

After 15.1±4.2 months from initial evaluation, no relapses had been reported by any patient. This was in line with the stability of the frequency of CLIN and MRI abnormalities (37.3% and 71.1%, respectively) at the follow up. Despite this, BSRs and EPs revealed a worsening of BS function. In particular, although the proportion of altered BSRs did not change significantly (80.6% vs 90.3%; p=0.180), a significant worsening of scores was observed for VMR (p=0.001), AMR (0.018) and TCR (p=0.013). Similarly to BSRs, the incidence rate of EP abnormalities did not increased significantly (84.4% vs 86.7%, p=0.564), but the analysis of cumulative score showed a significant worsening for the whole EP set (p=0.03) as well as for median SEP (68.9% vs 75.6%, p=0.03), P14 mSEP (33.3% vs 51.1%, p=0.005), tibial SEP (60% vs 66.7%, p=0.03).

Conclusions

BSRs and EPs were able to reveal a significant worsening of BS functions in spite of any variation of both BS signs/symptoms and of MRI BS lesion load. This is in agreement with previous reports on BSR/EP ability to detect clinically and radiologically silent BS lesions. Further studies are needed in a larger cohort of patient to assess BSR clinical usefulness in a longitudinal perspective.

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Other

The spectrum of ocular motor abnormalities as the only clinical sign of brainstem lesions

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Background and aims: The widespread use of magnetic resonance imaging (MRI) revealed an increasing variety of ocular motor abnormalities as the only clinical sign of (mainly ischemic) brainstem lesions. This paper reviews the variety of such abnormalities.

Results: Ocular motor abnormalities as the only clinical sign of MRI-documented brainstem lesions include complete and partial 3rd, 4th and 6th nerve palsies, vertical gaze palsies, crossed vertical gaze palsy, monocular elevation paresis, internuclear ophthalmoplegia, horizontal gaze palsy, horizontal gaze palsy with facial palsy, 1½-syndrome, 1½-syndrome with facial palsy ("8½-syndrome"), upbeat nystagmus, horizontal-rotatory nystagmus, horizontal nystagmus, skew deviation, and ocular tilt reaction.

Conclusions: Brainstem lesions causing isolated ocular motor abnormalities may be divided into 4 main groups. (A) Lesions involving infranuclear ocular motor nerve segments cause complete and partial 3rd, 4th and 6th nerve palsies. (B) Lesions affecting nuclei related to eye movements such as 3rd and 6th nerve nucleus, rostral interstitial nucleus of the medial longitudinal fasciculus, interstitial nucleus of Cajal, nucleus intercalatus Staderini are followed by horizontal and vertical gaze palsies, upbeat nystagmus, horizontal-rotatory nystagmus. (C) Lesions interrupting internuclear connections lead to internuclear ophthalmoplegia, monocular elevation paresis, skew deviation and ocular tilt reaction. (D) Combined lesions of nuclear and internuclear or infranuclear structures are the anatomical basis of 1½-syndrome, 8½-syndrome, or horizontal gaze palsy with facial palsy. The clinical significance of these disorders is not known.

Other

Optokinetic nystagmus of Chiari Malfromation. The aim of our study is to determine the special characteristics of disorders of the Optokinetic nystagmus in the early stages of Chiari Malfromation.

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Optokinetic nystagmus of Chiari Malfromation.

Diagnosis and treatment of cochleovestibular dysfunction becomes complicated in case of genetically based combination of pathology.

The Chiari Malformation (CM) is a congenital development defect and associated with the caudal displacement of cerebellum and brain stem.

The aim of this study is to determine the special characteristics of disorders of the Optokinetic nystagmus in the early stages of CM.

At the otorhinolaryngology department of North-West State Medical University named after I.I.Mechnikov (Russia, St Petersburg), 54 cases (patients aged 16 to 69) were studied. All patients showed the symptoms of cochleovestibular dysfunction.

The Optokinetic nystagmus was taken in a dark-room using VNGULMAR system manufactured by the German company "Heinemann Medizintechnik GmBH" running software developed by the French company Synapsys. The fixed speed horizontal and vertical Optokinetic stimulus on screen (20 o/s) was applied with distracted and focused attention of the patients.

All the studies were conducted before the calloric test (CT) and after CT. The intensity of Optokinetic nystagmus was assessed as per SSC (speed of slow component) of nystagmus. Later on the relative asymmetry in Optokinetic nystagmus was calculated for each same name pair of horizontal and vertical Optokinetic reactions.

The studies showed that in order to diagnose CM in the early stages it is important not only to identify the asymmetry in horizontal and vertical Optokinetic nystagmus but to mark the value and the sign of asymmetry as well. The sign of asymmetry is especially important after CT. Patients with the low placement of cerebellar tonsil Optokinetic reactions are more susceptible to CT.

Therefore, in the early stages of CM and "in the risk group", i.e. the patients with low placement of cerebellar tonsil, before the neurological symptoms have taken place and we have only vestibular dysfunction, the videonystagmography with quantitative evaluation of Optokinetic reaction becomes a voluble source of important diagnostic information and an indication for MRI studies.

Other

Topology of Brainstem Lesions Causing Subjective Visual Vertical Tilt

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Objectives

The subjective visual vertical (SVV) is a sensitive sign of a vestibular otolith tone imbalance in the roll plane. Recently, there are increasing evidences suggesting the existence of crossed and uncrossed anatomical pathways in the brainstem for verticality perception. In this study, we aimed to determine the topology of the anatomical pathway in the brainstem for verticality perception.

Methods

We measured the SVV in 82 patients with acute unilateral infarction involving the brainstem only. The topology of the brainstem lesions responsible for pathological SVV tilt were determined using MRI-based voxel-wise lesion-behavior mapping, and the probabilistic lesion maps were constructed.

Results

Fifty percent (41/82) of patients with acute unilateral brainstem infarction showed abnormal tilts of the SVV, of which 76% (31/41) patients showed ipsiversive and 24% (10/41) had contraversive tilt. Patients with contraversive SVV tilt showed overlapping of the lesions in the rostral medial vestibular nucleus, medial longitudinal fasciculus, and rostral interstitial medial longitudinal fasciculus and interstitial nucleus of Cajal. In contrast, the lesions in patients with ipsiversive SVV tilt with ocular tilt reaction were mostly involved the medial and inferior vestibular nuclei in the caudal medulla and lesions with purely verticality tilt involves the medial side of the medial lemniscus.

Conclusions

Our data support the evidence that there is a pathway transmitting ipsiversive otolithic signals bypassing the oculomotor system at the medial side of the medial lemniscus called the ipsilateral vestibulo-thalamic tract (IVTT).

Other

Complexities in Interpretation of Stimulated Cranial Nerve EMG During Brainstem Surgery

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Question:

Stimulated electromyography (EMG) is often used to identify cranial nerves (CN) that have a motor component during brainstem surgeries. Often multiple CN are monitored as lesions extend over large areas of the brainstem. Several cranial nerves supply muscles that are adjacent, and often multiple muscles appear to be activated with stimulation of a single CN. This may make identification of individual cranial nerves difficult. In this study we analyzed how often single CN stimulation resulted in activation of muscles innervated by multiple CN and how best to better determine which cranial nerve has been stimulated.

Methods:

Neurophysiologic intraoperative monitoring (NIOM) data from 6 subjects that underwent brainstem surgery in which two or more motor CN monitoring were monitored was analyzed. The CN monitored included CN V, VI, VII, IX, X, XI, XII. The patterns of EMG activation were noted, and the CN most likely stimulated to elicit that pattern was determined.

Results:

A few patterns of apparently simultaneous CN activation were noted. CN IX and X; CN VII and IX; CN VII, IX and X; and CN V and VIII were simultaneously activated in one or more patients. Independent activation was noted with the following CN stimulation: V, VII, IX, X. When multiple cranial nerves appeared to be stimulated simultaneously, closer analysis revealed one nerve that was primarily activated. Attention to latency and morphology of the waveform assisted with determination of the abnormal response. Lack of a clear peak and longer latency suggested a volume conducted response from nearby muscles. When CN IX and X were activated simultaneously, activation was most likely of CN X; simultaneous activation of CN VII and IX was most likely of CN IX; simultaneous activation of CN VII, IX and X was most likely of CN X; and simultaneous activation of CN V and VII was sometimes from CN V stimulation and sometimes from CN VII stimulation.

Conclusions:

Stimulation of a single CN can produce EMG activity in muscle groups that are seemingly supplied by other CN. Knowing which muscle groups appear to be commonly co-activated and analyzing the morphology and latency of the responses allows for correct identification of the CN stimulated.

Other

Diagnostic test study of Indonesian version of The Neurological Disorders Depression Inventory for Epilepsi In adult Epilepsy patients with major depression disorders

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Background.Depression is a common psychiatric disorder in epilepsy. The prevalence is 20-80%. The depression is not routinely assessed in neurology clinics, because the assestment takes a long time. Therefore many patients are under diagnosed and untreated. The Neurological Disorders Depression Inventory for-Epilepsy (NDDI-E) is a depression screening examination consist of only 6-aitem.

Purpose. To determine the accuracy and cut-off point of NDDI-E Indonesian version as a screening depression examination for adult epilepsy patients.

Method. Diagnostic test study was conducted at epilepsy clinic on Ciptomangunkusumo hospital. All the epilepsy patient who met the inclusion criteria was examined. The patient took the NDDI-E Indonesian version as a self assessment. Then they were assest used the International Neuropsychiatric Interview Mini ICD-10 (MINI-ICD10) as a gold standar.

Results. From the 105 subjects, there were 23 people suffered from major depression by MINI-ICD10. Receiver Operating Characteristic (ROC) curve obtained which is close to 100%, cut-off point at 11, with Sensitivity 91.3% Specificity 89% PPV 70% and NPV of 97.3%. It was statistically classified as strong because the value of Area Under the Curve (AUC) is 97.5% with a confidence interval (95% CI 95% -99%).

Conclusion. NDDI-E Indonesian version has a high accuracy to determine major depressive disorder in adult epilepsy patients with the cut-off point at 11.

Other

Study of locomotor and cognitive disorders in the ovariectomized female Wistar rats

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Objective:

Menopause is accompanied by a cognitive and neurodegenerative dysfunction. These disorders are mainly due to the collapse of the level of estrogen causing a chronic inflammation in the brain associated with a several unbalance in neurotransmission. This study has for objective to study the impact of the deficit in estrogens on the levels of anxiety on ovariectomized female Wistar rat.

Methodology:

Female rats (172 to 226 g), aged 6 months, were used during this study. The animals were randomly divided into two groups: a control group "C" and an ovariectomized group "OVX". Three months later, the levels of anxiety were evaluated by valid behavioral tests, Open Field test (OFT) and elevated plus maze (EPM).

Results:

The parameter values are collected and statically analyzed show **partiality** that:

* In the OFT, the number of central squares visited as well as the time spent in the central squares by the OVX rats are significantly lower than the recorded values among the control.

* In the test EPM test, the time spent in the open arms by the OVX rats is substantially lower than that recorded by the control rats.

Conclusion: These preliminary results suggest that the ovariectomy is associated with an increase in the level of anxiety, which would be due to a decrease in the secretion of ovarian hormones, particularly estrogen.

Keywords: Ovariectomized rat, cognitive disorders, OFT, EPM.

Other

Methamphetamine-induced aberrant neurogenesis of neural progenitor cells

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Adult hippocampal neurogenesis is an important modulator of brain plasticity and plays a key role in learning and memory functions. The neural stem cell (NSC)-derived neurons are incorporated into normal brain structures and participate in learning and memory tasks, i.e. the functions mediated by the hippocampus. As a drug of abuse, methamphetamine (METH) induces working memory deficits, oxidative stress, neuroinflammation, hyperthermia, and disruption of the blood-brain barrier (BBB). While the hippocampus is one of the brain regions particularly susceptible to METH toxicity, the mechanisms underlying METH toxicity are still poorly understood. In particular, the input of METH-induced BBB disruptions on adult brain neurogenesis is not known.

The central hypothesis of the present study is that the disruption of BBB integrity induced by chronic METH exposure is responsible for toxicity to neural stem cells (NSCs), resulting in deficient neurogenesis. To address this hypothesis, C57BL/6 mice were treated with escalating doses of METH 3 times per day for four days. Exposure of to METH reduced immunoreactivity of nestin, the marker protein of NSCs, in the dentate gyrus. Because the samples were collected 5 days after the last injection of METH, these results suggest that aberrant neurogenesis persist even exposure to METH is discontinued. Delayed differentiation was also observed in cultures of mouse neural stem cell line NE-4C and primary mouse embryonic NSCs exposed to METH in vitro. Indeed, METH exposure reduced the length of dendrites in both cell types and inhibited the formation of secondary branching dendrites in NE-4C cells. These results suggest that METH exposure affect differentiation of NSC into the proper neuronal linage, which may contribute to the development of cognitive dysfunction in drug abusers. This work was supported by the National Institutes of Health, grants DA027569, CA133257, MH063022, and MH098891.

Pain

Trigeminal Somatosensory Evoked Potentials in Assessment of Craniofacial Pain: Preliminary Results

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Introduction: Craniofacial pain syndromes include various kinds of primary and secondary headaches, most of which are related to trigeminal system such as trigeminal neuralgia, trigeminal neuropathy, cluster headache or other trigeminal autonomic cephalalgia. Blink reflex, masseter inhibitory reflex, jaw jerk and trigeminal laser-evoked potentials are known to be more reliable in neurophysiological testing of trigeminal system and electrophysiologic testing of trigeminal reflexes accurately distinguishes symptomatic and classic trigeminal neuropathy.

Here, we aimed to evaluate the reliability and stability of trigeminal nerve somatosensory evoked potentials (TSEP) in less frequently encountered craniofacial pain syndromes.

Patients and method: TSEP and blink reflex (BR) were recorded in 10 patients with craniofacial pain during the active disease period and in 25 healthy individuals. The latencies of N2, P2 and N3 waves as well as amplitudes of N2/P1 and N2/P2 were measured.

Results: We could not obtain TSEP in three patients (30%) whereas it was elicited in all healthy subjects (p=0.01). All of the patients without TSEP response had idiopathic etiology and only one of them also lacked BR responses. Latencies appeared to be longer in patients with craniofacial pain compared to healthy subjects. However, BR responses were symmetrical and within normal ranges.

Conclusion: Keeping in mind the low number of participants included, we think that TSEP is able to be obtained in a stable manner in healthy subjects and it may be more capable to reflect changes in pain conditions compared to BR.

Pain

Functional magnetic resonance imaging of pain-related brainstem nuclei in single subjects at 7 Tesla

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The function or dysfunction of pain-related nuclei in the brainstem has only been scarcely studied in humans due to the lack of non-invasive measurement methods. The performance of functional magnetic resonance imaging (fMRI), a standard non-invasive technique, is hampered by the close vicinity of the brainstem to large arteries and ventricles as well as its propensity to spatial distortions caused by the oral cavity. Furthermore, the small average size of brainstem nuclei necessitates higher spatial resolution and accuracy than in studies of the cortex. Here, we present a new approach based on ultra-high field fMRI acquisition at 7 Tesla and a brainstem-optimized analysis method (mICA), which we apply to study pain-related activity and connectivity of brainstem nuclei in single subjects. Following a multimodal imaging approach based entirely on echo-planar imaging, we acquired distortion matched functional, T1weighted structural as well as diffusion-weighted images at a resolution of 1.2 mm isotropic. Five 6min runs of restingstate fMRI were acquired, during two of which subjects received a continuous pressure pain stimulus applied to their lower leg by an inflatable cuff. Pain intensity was percept matched at 50/100 points on a visual analogue scale. Applying our recently developed mICA approach (masked independent component analysis), we were able to detect reproducible resting-state activity for specific brainstem nuclei, like the cuneiform nuclei, periaqueductal grey and as well as brainstem-cortex functional connectivity at the single-subject level. We identified a number of pain-related nuclei that showed distinctive activity changes during pain stimulation (Figure 1). Identification of nuclei was greatly aided by fractional anisotropy (FA) maps created from the diffusion data. Finally, assessing activity and functional connectivity of brainstem nuclei on the single-subject level may soon give us a deeper understanding of disease subtypes, individual differences in pain processing, as well as other functions localized in the brainstem



Figure 1: Results from a single subject obtained at 7 Tesla. A number of known pain-related nuclei show activity changes as a result of the pain stimulation. Abbreviations: Cun = cuneate nucleus, DMX = dorsal motor nucleus of the vagal nerve, VLM = ventrolateral medulla, LC = locus coeruleus, PAG = periaqueductal grey.

Pain

fMRI evidence for a reduction in affective processing of thermal pain in responders of transcutaneous vagal nerve stimulation (TVNS)

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Objectives

Although the transcutaneous vagal nerve stimulation (TVNS) is increasingly used in treatment of chronic pain, the underlying mechanism is unclear. The goal of this study was to analyse the cerebral effects of TVNS under experimental pain in fMRI.

Methods

Twenty healthy volunteers took part in two separate fMRI sessions (3 Tesla) with experimental pain, which was applied to the right forearm with MEDOC Sensory Analyser. During TVNS session the electrical stimulation was applied bilaterally to auricular concha with rectangle impulses (8 Hz frequency; 200 µs wave length); the intensity was maximal but not painful. During placebo session the stimulation device was switched off. The order of the sessions was randomised. Individual sensory thresholds were registered before and after each fMRI session. fMRI data was afterwards processed with standard settings in SPM8 and differences in BOLD effects between verum and placebo session were calculated. We used region of interest (ROI) based analysis on ROIs, which were previously identified in association with thermal pain.

Results

Cerebral pain processing areas were activated bilaterally under thermal stimulation. The group analysis did not show the differences in sensory parameters but revealed stronger activation of right amygdala under TVNS. The group of responders (subjects with increased pain threshold after TVNS, n= 8), showed a reduction in BOLD signal in the right nucleus caudate, the middle frontal cortex and left hypothalamus under TVNS in comparison to placebo.

Conclusion

Thermal stimulation elicited bilateral fMRI activation in pain processing regions of the brain in healthy volunteers. Responders to TVNS showed decreased activation in cerebral areas associated with affective processing of pain.