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Current and Emerging Potential of Magnetoencephalography in the Detection and Localization of High-Frequency Oscillations in Epilepsy

Eleonora Tamilia1*, Joseph R. Madsen2, Patricia Ellen Grant1, Phillip L. Pearl3 and Christos Papadelis1

1 Fetal-Neonatal Neuroimaging and Developmental Science Center, Division of Newborn Medicine, Department of Medicine, Boston Children’s Hospital, Harvard Medical School, Boston, MA, USA, 2 Division of Epilepsy Surgery, Department of Neurosurgery, Boston Children’s Hospital, Harvard Medical School, Boston, MA, USA, 3 Division of Epilepsy and Clinical Neurophysiology, Department of Neurology, Boston Children’s Hospital, Harvard Medical School, Boston, MA, USA

Up to one-third of patients with epilepsy are medically intractable and need resective surgery. To be successful, epilepsy surgery requires a comprehensive preoperative evaluation to define the epileptogenic zone (EZ), the brain area that should be resected to achieve seizure freedom. Due to lack of tools and methods that measure the EZ directly, this area is defined indirectly based on concordant data from a multitude of presurgical non-invasive tests and intracranial recordings. However, the results of these tests are often insufficiently concordant or inconclusive. Thus, the presurgical evaluation of surgical candidates is frequently challenging or unsuccessful. To improve the efficacy of the surgical treatment, there is an overriding need for reliable biomarkers that can delineate the EZ. High-frequency oscillations (HFOs) have emerged over the last decade as new potential biomarkers for the delineation of the EZ. Multiple studies have shown that HFOs are spatially associated with the EZ. Despite the encouraging findings, there are still significant challenges for the translation of HFOs as epileptogenic biomarkers to the clinical practice. One of the major barriers is the difficulty to detect and localize them with non-invasive techniques, such as magnetoencephalography (MEG) or scalp electroencephalography (EEG). Although most literature has studied HFOs using invasive recordings, recent studies have reported the detection and localization of HFOs using MEG or scalp EEG. MEG seems to be particularly advantageous compared to scalp EEG due to its inherent advantages of being less affected by skull conductivity and less susceptible to contamination from muscular activity. The detection and localization of

Abbreviations: ECG, electrocardiography; EEG, electroencephalography; EMG, electromyography; EOG, electrooculography; EZ, epileptogenic zone; fMRI, functional magnetic resonance imaging; HFO, high-frequency oscillations; iEEG, intracranial electroencephalography; EEG, intracranial electroencephalography; IED, interictal epileptiform discharge; iEEG, intracranial electroencephalography; MEG, magnetoencephalography; MEM, maximum entropy on the mean; MRE, medically refractory epilepsy; MRI, magnetic resonance imaging; MSI, magnetic source imaging; MUSIC, MUltiple SIgnal Classification; PET, positron emission tomography; SAM, synthetic aperture magnetometry; SNR, signal-to-noise ratio; SOZ, seizure-onset zone; SPECT, single photon emission computed tomography; wMEM, wavelet maximum entropy on the mean.
HFOs with MEG would largely expand the clinical utility of these new promising biomarkers to an earlier stage in the diagnostic process and to a wider range of patients with epilepsy. Here, we conduct a thorough critical review of the recent MEG literature that investigates HFOs in patients with epilepsy, summarizing the different methodological approaches and the main findings. Our goal is to highlight the emerging potential of MEG in the non-invasive detection and localization of HFOs for the presurgical evaluation of patients with medically refractory epilepsy (MRE).

Keywords: high-frequency oscillations, magnetoencephalography, epilepsy, epileptogenic zone, epilepsy surgery, source localization

INTRODUCTION

Epilepsy is one of the most common neurological disorders affecting children and adults (1, 2). About 65 million people currently carry the diagnosis of epilepsy all over the world (3), and in the US only, epilepsy causes about $9.6 billion of medical expenditures and indirect care (4). Although most of the patients are treated successfully with antiepileptic drugs, approximately 30% of patients suffer from medically refractory epilepsy (MRE), i.e., they have unsatisfactory control and continue having seizures (5, 6). For these patients, epilepsy surgery represents the most promising treatment to pursue seizure freedom.

The objective of epilepsy surgery is the complete resection or disconnection of the epileptogenic zone (EZ), the area of the cortex necessary for the generation of habitual seizures and the smallest amount of tissue that can be removed to achieve a seizure-free outcome (7, 8). The only way to evaluate the success of the epilepsy surgery is by looking at the postsurgical outcome: if the EZ has been correctly identified and resected with no damage to the functionally relevant eloquent cortex (i.e., the region that is necessary for defined cortical functions), the patient will be seizure free with minimal or no functional deficits. Hence, the success of epilepsy surgery strongly depends on the successful delineation of the EZ. However, there is no diagnostic modality able to unambiguously delineate this zone (8). The EZ is a theoretical construct, and to date, there is no established marker that definitively determines its location and extent. The EZ can only be estimated through a variety of diagnostic tests that point out different cortical zones that are considered more or less precise indicators of the EZ (8–10) (see Figure 1):

- The seizure-onset zone (SOZ), i.e., the area where the clinical seizures originate on ictal recordings.
- The irritative zone, i.e., the area of the cortex that generates interictal epileptiform discharges (IEDs) in the electroencephalography (EEG) or magnetoencephalography (MEG).
- The epileptogenic lesion, i.e., a structural brain abnormality that is causally related to the epilepsy.
- The functional deficit zone, i.e., the area of the cortex that is functionally abnormal during the interictal period, as indicated by neurological examination, neuropsychological testing and functional imaging or non-epileptiform EEG or MEG abnormalities.

The SOZ is generally considered the best estimate of the EZ, and it is removed in the majority of epilepsy surgeries. However, the identification of the SOZ is difficult because clinical seizures are unpredictable in nature and thus can be difficult to be captured by EEG. In addition, seizures captured during patient's monitoring may not represent the full extent of the EZ. This is why the removal of the entire SOZ does not always lead to successful outcome. Another indicator of the EZ is the irritative zone, which is defined by the localization of IEDs that occur more frequently than seizures. The big advantage of the irritative zone is that it can be evaluated during the interictal period independently from the occurrence of seizures. This reduces the required recording time and the associated cost and patient's discomfort. However, the irritative zone is often more widespread than the actual EZ and thus less specific to it (9). Furthermore, the presence of an epileptogenic lesion in the proximity of the irritative zone or the SOZ can provide an additional indicator of the location of the EZ.

Along with the localization of the EZ, the mapping of the eloquent cortex is essential to determine viability and strategy of the epilepsy surgery. This region is determined during the patient's presurgical evaluation by identifying the essential functional areas that subserve motor, memory, language, and visual functions, which need to be preserved from resection. Thus, the ultimate goal of epilepsy surgery is to achieve seizure freedom by removing the EZ and also avoiding functional deficits caused by any damage to the eloquent cortex (10).

All the aforementioned cortical zones are initially defined using a battery of non-invasive diagnostic tests, such as scalp EEG, MEG, magnetic resonance imaging (MRI), function MRI (fMRI), positron emission tomography, and single photon emission computed tomography. The results of these non-invasive tests can be insufficiently concordant and fail to derive a clear hypothesis about the location of the EZ. In these cases, long-term monitoring with intracranial EEG (iEEG) is needed in order to evaluate the possibility of surgery (11–14). However, iEEG recordings face several limitations not only related to their invasiveness and cost but also to their limited spatial sampling that might lead to misleading conclusions (15, 16). The intracranial electrodes have a field of view of only several millimeters and, if their placement is not optimal, areas with relevant epileptic activity may be overlooked. Therefore, seizures that originate from areas not covered by the iEEG electrodes, but propagate to the area where the electrodes are placed, might lead to misleading results. In order to
reduce or improve the use of such invasive, time-consuming, and costly investigations, and to improve the postsurgical outcome of patients who can undergo surgery, non-invasive presurgical tools with high sensitivity and specificity to the epileptogenic focus are being sought (17). Utilization of a non-invasive biomarker may help in various clinical scenarios that are typical in MRE (18), particularly in children. These include, for example, (i) patients with early onset severe epilepsies in which both generalized clinical semiology and EEG abnormalities can manifest yet with the presence of a definitive localized epileptogenic focus (19); (ii) tuberous sclerosis with multiple tubers in which resection of the most epileptogenic tuber can greatly improve seizure outcome (20); (iii) multifocal cortical dysplasia in which resection of the most epileptogenic lesion can lead to seizure freedom (21); (iv) bilateral migrational anomalies, e.g., polymicrogyria, in which partial lesionectomy may lead to seizure control and also enable sparing of eloquent cortex; and (v) temporal lobe epilepsy, whose large proportion of failed surgeries indirectly shows many underlying epileptogenic networks (18).

New Promising Trends for the Delineation of the EZ

Given the lack of an unambiguous marker of the EZ and the limitations of the invasive recordings, the presurgical delineation of the EZ is complicated and often unsuccessful. As a result, a large proportion of patients undergoing epilepsy surgical resection [34–73% depending on the resection type (22)] continue to have seizures after the surgery. In addition, epilepsy surgery remains the most underutilized of all accepted medical interventions (23, 24): in United States, less than 1% of patients with MRE are referred to epilepsy centers for surgery (25). In order to improve the safety and efficacy of the epilepsy surgical treatment, there is an overriding need to identify and validate reliable biomarkers that can determine the extent and location of the EZ with high precision and accuracy.

During the last decade, high-frequency oscillations (HFOs) above 80 Hz have emerged as a new promising biomarker of epileptogenicity (26, 27). Recent studies have shown that the resection of the tissue generating HFOs may improve presurgical diagnosis (28, 29) and surgical outcome of patients with MRE (30–38). The HFO zone has been recently proposed as a further piece of information to circumscribe the EZ (see Figure 1). Its correlation with the other cortical zones is an area of active research (26, 27). Despite the promising findings, HFOs are not yet suited for the diagnosis or monitoring of epilepsy in clinical practice (39, 40). One of the major reasons is the difficulty to detect and localize them with non-invasive methods (26, 27, 40). However, as the clinical use of MEG has
been increasing for patients with epilepsy (41–47), an emerging body of literature has provided evidence that HFOs can be also recorded non-invasively (48–52). These recent findings pave the way for the translation of these new biomarkers in clinical practice.

The use of a non-invasive biomarker of epileptogenicity might limit long-term monitoring and invasive intracranial recordings leading to a significant improvement of the presurgical evaluation procedure in MRE patients. Such a biomarker would not only help to identify the epileptogenic tissue for surgery but also permit definitive differential diagnosis of epilepsy from acute symptomatic seizures so treatment can begin immediately. It might also make it possible to assess the efficacy of therapeutic interventions without waiting for another seizure to occur, which could be associated with significant morbidity or mortality (40).

MEG IN THE PRESURGICAL EVALUATION OF PATIENTS WITH EPILEPSY

Magnetoencephalography is being increasingly used during the first phase of the presurgical evaluation of patients with MRE (41–47), helping to evaluate whether surgery is viable and to plan the surgical strategy. MEG is rapidly becoming invaluable for the presurgical evaluation of patients with epilepsy (17) due to its several strengths and possible applications:

- **Intrinsic properties:** MEG presents a unique set of intrinsic properties: (i) MEG signals are not distorted by the skull or the intervening soft tissues between brain and scalp (53–55); this property is particularly important in patients with large lesions, anatomical malformations, or patients who already had a resection and undergoing second surgery; (ii) MEG, unlike EEG, is contactless and thus patient’s preparation is faster; this is particularly important for pediatric patients who are difficult to stay still for long time and do not always follow instructions; (iii) MEG signals can be recorded with a high density of sensors avoiding the problem of salt bridge between EEG electrodes, a problem often faced particularly in children due to their small heads (41); and (iv) high-frequency activity is considered to be less susceptible to contamination from muscular activity in MEG than in scalp EEG (56, 57).

- **Functional mapping:** MEG has the ability to localize functional areas of the brain (58–63), such as primary sensory areas (i.e., visual, somatosensory, and auditory cortices), as well as areas responsible for higher and more complex cognitive functions [i.e., language (64, 65) and memory (66, 67)]. MEG is advantageous compared to alternative methods, such as the intraoperative awake surgery (68) and the fMRI (69–71).

- **Optimization of iEEG:** MEG can guide the placement of iEEG electrodes (44, 72–74) by providing information regarding the localization of epileptic activity and functional areas. Such information regarding the electrodes placement is important since iEEG cannot be repeated easily due to the local scar following implantation. This may be particularly beneficial for patients with normal MRI findings (61, 75) or with multifocal/diffuse disease (45, 76).

In summary, MEG has several significant inherent strengths that make it a valuable tool to localize non-invasively the epileptic activity, to map the eloquent cortex, and to guide the placement of subdural strips and grids and/or depth electrodes. The advantages of MEG in the preoperative evaluation of patients with MRE have been reported in several studies that showed that the concordance of the MEG localization with the resected area correlated with postsurgical outcome (61, 77–79). The major limitation of MEG is the expense: MEG use is associated with high costs, first in terms of initial equipment purchase (~$1.5–2 million) and second in terms of daily operations (liquid helium and personnel expertise required) (41). The operational costs are expected to decrease significantly in the near future thanks to the availability of liquid helium recycling systems (41).

BASIC PRINCIPLES OF MEG

Magnetoencephalography records the magnetic activity generated by electrical currents in active neurons of the human brain using the phenomenon of electromagnetic induction (41, 80, 81). Simultaneous recordings of epileptiform activity by MEG and iEEG have demonstrated that 4 cm² of synchronously active cortex is needed to generate a spike that can be detected by MEG (82). The magnetic signals generated by the human brain are extremely weak compared to background electromagnetic activity. Thus, MEG recordings are performed inside specially designed rooms, called magnetically shielded rooms, which minimize the electromagnetic noise from external sources such as the power line or electrical devices. Figure 2 shows a conventional adult MEG system and its basic principles. During MEG recordings, the patient lays down on a bed and places his/her head inside the MEG sensor helmet. An illustrative complete description of MEG recordings can be found elsewhere (51). The MEG helmet contains specially designed detection coils that are able to record changes in the magnetic field (83, 84). These coils are connected to superconducting quantum interference devices, which convert the magnetic field passing through the detection coils into voltage changes.

Magnetoencephalography offers excellent temporal resolution in the range of sub-milliseconds and very good spatial resolution of few millimeters (85–89). Under specific requirements, MEG can localize superficial sources with an accuracy that reaches the cytoarchitectonic level of the cortex (90). Several studies reported that MEG and scalp EEG are comparable in terms of localization accuracy but their combination has better yield than either technique alone (89, 91). In epilepsy patients, some studies showed that MEG is more sensitive for spike sources, especially from the neocortex, because of the higher signal-to-ratio than EEG (89, 92), but no conclusions can be drawn yet. The relative orientation of the active source with respect to the skull significantly affects the strength of the recorded MEG signals. Sources with tangential orientation produce the maximum measurable magnetic field outside the scalp, while sources with radial orientation produce the minimum measurable magnetic field and thus are almost undetectable (93–95).
Fig 2 | Basic principles of signal generation in a conventional adult magnetoencephalography (MEG) system. (A) A conventional MEG system. The patient places his/her head in a special helmet that accommodates a high number of sensors (typically >250). (B) Schematic representation of the electromagnetic induction phenomenon: a changing magnetic field (B) generates a measurable electrical current (I) that is recorded outside of the head by special magnetic field pick-up coils. The coils are connected with the superconducting quantum interference device (SQUIDs), which convert weak changes in the magnetic field into measurable voltage (V). (C) Schematic representation of the MEG signal generation. An intracellular electrical current (blue arrow) generates a magnetic field (red ring) around the apical dendrite. The magnetic field is picked up by the detection coils. Sources in the brain sulci cause tangential fields that can be detected by MEG.

**MAGNETIC SOURCE IMAGING (MSI)**

Magnetoencephalography records the magnetic brain activity that is measurable outside the scalp using extracranial sensors. Thus, the number, location, and strength of the generators that produce the MEG signal are unknown. In order to define the underlying generators and estimate their characteristics, the inverse problem should be solved (81) (Figure 3). The inverse problem does not have unique solution because an infinite number of sources within the brain can produce similar extracranially recorded fields. Thus, a priori assumptions about the number and the nature of the underlying sources are necessary to constrain the inverse problem solution. Assumptions should also be made regarding the nature of the volume conductor, what is called the forward problem, by postulating a priori head models (96) (Figure 3). The forward problem has unique solution: an intracranial source of known location, orientation, and magnitude generates a mathematically well-defined extracranial magnetic field.

Magnetic source imaging is the method that combines MEG with MRI by projecting the magnetic activity recorded with the MEG on the patient's anatomic image after solving the inverse problem (97). The MSI is a model-based imaging technique that integrates temporal and spatial components of MEG to identify the generating source of the magnetic fields recorded by the physical MEG sensors (see Figure 3). MSI significantly improves the interpretation of MEG raw data, which consist of the magnitudes of magnetic fields measured at different recording sites, since it enables to estimate the three-dimensional intracerebral location, orientation, and time activity of the underlying neuronal sources. MSI has both technical and clinical validity in the localization of the sources of IEDs (45, 97, 98).

**HIGH-FREQUENCY OSCILLATIONS AS A NEW BIOMARKER OF EPILEPSY**

High-frequency oscillations in epilepsy are generally described as short spontaneous EEG patterns in the frequency range from 80 to 500 Hz, consisting of at least four oscillations that stand out of the background activity (27). HFOs can be sub-classified according to their frequency content in “ripples” (80–250 Hz) and “fast ripples” (250–500 Hz) (99). Recently, very-high-frequency oscillations (>1,000 Hz) have been also observed in patients with epilepsy (100). Ripples and fast ripples seem to be generated by different pathophysiological mechanisms, but it is still debated whether their relation with the EZ is different.
In literature, different recording methodologies (micro- vs. macro-electrodes) have been adopted (101), which led to discrepant conclusions (26) on the importance of discriminating between the two frequency ranges for the delineation of the EZ. Further research is needed to determine whether this discrimination is crucial or not. Furthermore, it may be challenging to reliably distinguish between physiological and pathological activity when looking at epileptic HFOs, since they may overlap with physiological oscillations in both ripple and fast ripple frequency bands (102, 103).

**HFOs IN THE PRESURGICAL EVALUATION OF EPILEPSY**

High-frequency oscillations are considered a valuable piece of information that could enable a direct definition of the EZ (27, 104) (Figure 1). Recent and ongoing research has been investigating the correlation of the HFO generating area with respect to the other cortical zones [for reviews, see Ref. (26, 27, 105, 106)]. HFOs have proved to be a reliable interictal indicator of the SOZ: they have been observed in the SOZ with higher rates than outside during interictal periods (107–110). Also, HFOs seem to be more specific and accurate markers of the SOZ than interictal spikes (110). Intercital HFOs can overlap spikes but can also occur independently from them in space and time (109). The clinical significance of the distinction between HFOs with and without spikes has not been established yet. However, it has been suggested that considering only HFOs that overlap with spikes would be particularly beneficial in the clinical context because it allows (i) reducing the time needed for the inspection of EEG or MEG data and (ii) excluding physiological HFOs that would represent false detections of epileptogenic biomarkers (111). Furthermore, spikes with HFOs have shown to be more closely related to the SOZ than spikes in general (109). This has led to hypothesize that the presence of HFOs may be crucial to distinguish between “red” and “green” spikes, i.e., between pathological and non-pathological spikes originating, respectively, from inside or outside the EZ. The occurrence of HFOs may help to perform such distinction and identify the most epileptogenic spikes (112, 113). Further research is needed to investigate this HFO-spike concurrence in relation to epileptogenicity.

High-frequency oscillations can also help in the delineation of the EZ independently of lesional boundaries. HFOs have proved to be a more reliable marker of the SOZ than the epileptogenic lesion, since their occurrence within a lesion is more closely linked to the SOZ than to pathologic tissue changes (114). Furthermore, in patients with focal cortical dysplasia, HFOs have been found mainly in the SOZ rather than in other areas of pathologic tissue (115). Thus, they represent a marker of the SOZ independent of the underlying pathology.

Looking at the postsurgical outcome, HFOs have proved to be promising markers of epileptogenesis for a successful surgery: removal of the HFO generating tissue is an indicator of good surgical outcome (31–38), while residual HFOs after surgery can predict poor outcome (37). Furthermore, the surgical removal of all the HFO generating areas, both within and outside the SOZ, correlates with good postsurgical outcome, suggesting that HFOs can also point toward epileptogenic areas outside the SOZ (31).

In summary, HFOs seem to be a reliable interictal biomarker of tissue capable of generating seizures. However, the evidence...
for an effective clinical use of HFOs for epilepsy surgery decision making is still weak (116) and the application of HFOs for presurgical evaluation is at an early stage (117, 118). There are still significant challenges for the clinical translation of HFOs as epileptogenic biomarkers. One of these challenges is the difficulty to detect and localize them non-invasively. This is reflected in the current literature that mostly investigated HFOs using invasive intracranial recordings (26, 27).

HFO Automatic Detection

To date, the visual detection of HFOs has led to a good understanding of the relationship between HFOs and epilepsy (26, 27, 108). However, visual marking is time consuming and subjective, since there is not a well-established definition of HFOs that enables their straightforward identification and each recorded signal should be inspected with an extended time scale display (i.e., 250 mm/s). The development of automatic HFO detectors has received much interest during the last decade and represents an area of active research (50, 51, 119–129). Such automatic detectors are crucial for the investigation of HFOs as biomarkers of epileptogenicity. They are likely necessary to propel future clinical applications, since they would enable an objective and consistent identification of HFOs in large-scale recordings. Given the lack of a quantitative definition of HFOs, the criteria used for automated detection differ from study to study, as well as the reported range of HFO amplitude (10–1,000 μV) and duration (30–100 ms) (120). Several algorithms have been proposed in the last decade for the automatic detection of HFOs (for reviews, see Ref. (119, 120)): they commonly define an HFO event as characterized by at least four oscillations standing out from the ongoing background activity in the frequency range of interest (80–500 Hz) and having at least 10 ms inter-event interval. In order to distinguish real HFOs from the filtering effects of a sharp transient (130) or from other EEG activity, recent advances in the automatic detection algorithms have also incorporated information from (i) the time–frequency domain, assuming that an HFO must appear as a short-lived event with incorporated information from (i) the time–frequency domain, as well as to the lack of consideration of peripheral recordings, such as electrocardiography (ECG), electromyography (EMG), or electrooculography (EOG), which can help the identification of cardiac, muscular, or ocular artifacts (145) resembling HFOs.

High-Frequency Oscillations and MEG

The non-invasive detection and localization of HFOs with MEG would significantly expand the clinical utility of these new promising biomarkers to an earlier stage in the diagnostic process and to a wider range of patients with epilepsy. Some recent studies have shown that epileptic HFOs can be detected non-invasively with scalp EEG (51, 110, 132–136), MEG (49–52, 137–142), or simultaneously with both techniques (51). Figure 4 shows simultaneous MEG and scalp EEG data with interictal HFOs co-occurring with epileptic spikes in a pediatric patient with focal epilepsy (51).

Detection of Epileptic HFOs with MEG

Given the increasing interest in the non-invasive localization of HFOs using MEG, recent studies have proposed novel methods for the automatic or semi-automatic detection of HFOs and their distinction from artifacts in MEG recordings (49–51). MEG data contain more high-frequency artifacts than iEEG (143), thus a careful visual inspection of the data is required to guarantee that the detected HFOs are indeed of cerebral origin. A purely visual HFO detection has been used when a limited number of virtual sensors were investigated (49, 52) (more details about “virtual” sensors are provided in Section “Non-Invasive Localization of Epileptic HFOs with MEG”). Yet, given the lower rates of HFOs in the MEG data compared to the iEEG and the high number of physical sensors (>300), a purely visual approach is impractical when analyzing the MEG signals from all the physical sensors as in Ref. (50, 51). A semi-automated approach for the HFO detection is the most appropriate in these cases. This approach consists of a highly sensitive automatic detection followed by the visual review of the detected events by an EEG/MEG expert in order to increase the specificity of the detection method (28, 126, 127, 135).

The automatic detector used in Ref. (50) was designed to identify HFOs in all the MEG channels and to keep only the events occurring in the channel with the highest HFO rate. The proposed detector disregarded any HFO that (i) occurred simultaneously in more than 100 channels and in different frequency bands, since muscle activity and movement artifacts usually involve many channels and have a broad frequency content (144); (ii) occurred in channels located at the edge of the MEG helmet, since these channels are mostly affected by movement (143); or (iii) showed a power increase in the ripple band lower than in channels with no HFOs. These criteria were proposed to exclude any HFO occurring simultaneously with other high-frequency physiologic activity, instrumental noise, or artifacts, since they could affect the MSI results. Then, the visual review was performed to inspect the time–frequency map and to discard artifacts visible in the MEG signals in the clinical frequency range (0.3–70 Hz). Limitations of the proposed method are possibly due to the subjectivity of the time–frequency map inspection, as well as to the lack of consideration of peripheral recordings, such as electrocardiography (ECG), electromyography (EMG), or electrooculography (EOG), which can help the identification of cardiac, muscular, or ocular artifacts (145) resembling HFOs.

Papadelis and colleagues (51) proposed a semi-automatic method to identify the HFOs that occurred on both MEG and scalp EEG data and that overlapped IEDs. The proposed detector identifies HFOs in the EEG signals not only looking at the time–frequency domain but also verifying the presence of an isolated island in the time–frequency map in order to automatically disregard possible artifacts (122, 130). Then, the visual review of the detected events was performed to keep only the HFOs that are also visible in the MEG data and that concur with the IEDs. In addition, to exclude any possible artifacts, the authors followed the guidelines proposed in Ref. (135) and excluded any oscillation: (i) with very high amplitude compared to the background; (ii) showing irregular morphology or large frequency variability (as shown in Figure 5); or (iii) co-occurring with muscle, movement, and electrode artifacts as identified with EOG, ECG, and EMG data at the timing of the detected HFOs. The main limitation of the proposed detection method is the lack of an automatic HFO
FIGURE 4 | High-frequency oscillations (HFOs) on simultaneous scalp electroencephalography (EEG) and magnetoencephalography (MEG) data. (A) Interictal spikes from a pediatric patient (15-year-old girl with encephalomalacia of the right middle cerebral artery region) with medically refractory epilepsy simultaneously recorded with scalp EEG (left) and MEG (right). Ten seconds of data filtered from 1 to 70 Hz are shown. (B) Interictal HFOs co-occurring with spikes. Extended timescale display of 2 s of data highlighted with gray background in (A). Data are filtered from 80 to 150 Hz for the HFO visualization. HFOs are underlined with red lines. (C) Time–frequency map of two representative HFOs marked with black squares in (B). We can observe the isolated peak in the ripple frequency band. Source: adapted from Papadelis et al. (51).

detection in the MEG signals: the EEG/MEG expert should verify the presence of simultaneous HFOs in the EEG/MEG signals through visual inspection, which can be subjective and possibly biased.

Further studies on simultaneous EEG/MEG recordings (51), possibly accompanied by simultaneous iEEG as well (135), may allow a better understanding of the scalp HFOs. Further knowledge and agreement about the morphology and structure of the HFOs in MEG data may lead to the design of highly reliable detection algorithms. A fully automated approach for the HFO detection would enable to analyze large databases minimizing the selection bias and human labor.

Non-Invasive Localization of Epileptic HFOs with MEG

Given the high resistivity of the skull and the assumption that a large extent of the cortex needs to be active in order to observe an HFO on the scalp, HFOs have been mostly investigated with intracranial depth electrodes and grids/strips on the cortical surface (26, 27, 40, 120). However, a few recent studies reported HFOs in the ripple frequency band recorded with scalp EEG in patients with epilepsy (51, 110, 132–135), showing lower HFO rates than in the iEEG. A recent scalp EEG study also reported fast ripples (250–500 Hz) recorded with subdermal electrodes in patients with focal epilepsy (136). Given the low number of
electrodes used, source localization techniques have not been used to localize the HFO generators in all these scalp EEG studies, except for the one of Papadelis and colleagues (51) who localized the HFO sources from high-density scalp EEG data of children with MRE. High-density scalp electrode distributions indeed are necessary to obtain a solid sampling and accurate localization of HFOs on the scalp, as showed by Zelmann and colleagues (135), since such oscillations represent the sum of activity of multiple spatially distributed focal and coherent sources.

Over the last years, different research studies (48–52, 137–142) have focused on the possibility of recording HFOs with MEG and localizing their generators at the source level, given the significant inherent advantages of MEG compared to scalp EEG. The first evidence that HFOs can be non-invasively detected and localized using MEG with an accuracy of few millimeters was provided by Papadelis et al. (48). In this study, the authors used a head-shaped construction to generate artificial signals that resemble the human HFOs and used MSI with different source methods to localize the underlying generators (see Figure 6). Four different source localization methods were used: (i) the equivalent current dipole (ECD), which describes the underlying source as an infinitesimal small line current element (81), was applied at the peak activity of HFOs on averaged data; (ii) the MUltiple SIgnal Classification (MUSIC), which scans a single ECD through a three-dimensional head volume and computes projections onto an estimated signal subspace (146); (iii) the synthetic aperture magnetometry (SAM), which is a beamformer designed to detect signals from a specified location and attenuate signals from all other locations (147); and (iv) the magnetic field tomography (MFT), which relies on a non-linear algorithm with optimal properties for tomographic analysis of the MEG signal (148). The authors showed that weak transient signals, resembling the human HFOs, can be detected with an accuracy of few millimeters at cortical and subcortical regions. All localization methods showed high accuracy (localization error <5 mm) even when only few (six) trials were included in the analysis. In particular, the localization accuracy was unaffected by the low number of trials (six) for the beamformer (i.e., the SAM) and the MFT, while decreased for the MUSIC and ECD on averaged data. Although this study was the first one to provide evidence that human HFOs can be non-invasively localized with MEG, it did not report source localization data for single events. Epileptic HFOs are non-phase-locked spontaneous events and their localization analysis must be performed on single trials. Furthermore, the confounding effect of background activity unrelated to HFOs and its effect on the source localization accuracy were not considered.

The source localization of extended generators of oscillatory activity in specific frequency bands requires dedicated methodologies. Given the low amplitude and low SNR of HFOs, the source localization techniques commonly used for the analysis of

![Figure 5](image-url)
epileptiform discharges with MEG (98, 149) (e.g., the ECD) are not recommended for the localization of HFOs (150, 151). This is because HFOs are not phase-locked events, and thus, their exact onset timing is unknown. As a result, different events cannot be averaged before solving the inverse problem to improve the SNR, as is typically done in stimulation studies (48, 152) or with epileptic spikes of similar morphology (41). Thus, alternative methods optimized for the source localization of single oscillatory events are necessary.

To improve the SNR of HFOs, beamforming techniques have been used (49, 52). Beamformers have the potential to distinguish HFOs from noisy background activity (48, 137, 153). The beamformer method (154–156) reconstructs the neuronal activity for specific locations within the brain, so-called virtual sensors, as the weighted contribution from different MEG physical sensors. Such beamformer weights act like spatial filters and allow attenuating noise from distant sources (154–157). In a pioneer work, Van Klink et al. (49) used virtual sensors, whose location was defined based on the localization of the interictal spikes, in order to improve the SNR of the MEG signals in the time domain and to confirm the presence of HFOs (>80 Hz). The results of this study showed that the less noisy beamformer virtual sensors enabled visual detection of epileptic HFOs that could not be identified in the signals recorded by the physical MEG sensors (see Figure 7). The HFOs were detected more often in the irritative zone than in the contralateral “mirror” region and overlapped with other clinical findings. This study showed that the proposed non-invasive method enables visual identification of areas with epileptic HFOs using MEG. However, this study presented a methodological drawback: a limited number of virtual sensors were selected based on the location of spikes in the physical MEG sensors missing possible HFOs which may occur outside the area under investigation. The same approach was used by Nissen et al. (52) to show that brain areas with interictal spikes and HFOs were functionally isolated from the rest of the interictal epileptic network. The automatic placement of a larger number of virtual electrodes, along with automatic HFO detection, would make the proposed method more suitable for application in clinical setting (52).

Another practical approach for the source localization of ripple HFOs in MEG signals was proposed by von Ellenrieder et al. (50). This was the first study that detected interictal HFOs as visible events standing out from the background MEG signal in the time domain and localized their generators at the source level. The authors used the wavelet maximum entropy on the mean (wMEM) method for the source localization. The wMEM is an extension of the maximum entropy on the mean (MEM) method (158–160) that is particularly well suited for this application since it was developed for the localization of oscillatory activity (161). The wMEM decomposes the signal into a discrete set of wavelet bases functions; it then performs MEM source localization on each time–frequency box (161). The study of von Ellenrieder and colleagues (50) is the first one performing MSI of spontaneous, and likely pathological, HFOs the ripple band (80–160 Hz) in patients with epilepsy. They detected HFOs from the signals of the MEG physical sensors, independently of the interictal spikes, and used the wMEM method to determine the location and extent of their generators on the cortex. For the validation of the HFO localization, the authors evaluated the spatial concordance of the HFO sources with the epileptogenic region defined by two specialists based on the available clinical information for each patient (i.e., resected region, ictal and interictal iEEG findings, visible lesion in the MRI, ictal, and interictal scalp EEG findings) (see Figure 8). The validation did not show fully concordant results suggesting the need for further studies in a larger cohort of patients. Furthermore, such a validation approach does not prove that non-invasively localized HFOs provide the same results as invasively localized HFOs. To this

![Figure 6](image_url)
purpose, comparisons of MEG results with the iEEG findings from the same subject are necessary to verify that MEG HFO localization coincides with the location obtained with invasive methods.

The same method for the localization of ripple HFOs was also used by Papadelis and colleagues (51) who were the first to investigate interictal HFOs with simultaneous MEG and high-density scalp EEG data from pediatric patients with MRE. In this study, the authors identified the HFOs occurring with interictal spikes in both MEG and scalp EEG and performed source localization using wMEM on the data from both modalities. The results of this study showed that the HFO localization was concordant between MEG and scalp EEG as well as concordant with other clinically relevant zones, such as the irritative zone and the SOZ (see Figure 9). The HFO localization for one patient was further validated against the ground truth given by the findings obtained from the HFOs detected in the iEEG data (see Figure 9, bottom). Such validation confirmed the source localization reliability of both MEG and scalp EEG. However, given the limited sample size, these promising findings need further validation.

In addition to the studies described so far, a series of MEG studies investigated the power in high-frequency bands of patients with epilepsy using accumulated source imaging (137–142). The accumulated source imaging is a source localization method that was developed to localize and quantify spontaneous brain activity (162). Researchers showed that MEG high-frequency components were localized within the EZ in pediatric patients with epilepsy (137–142). Xiang and co-workers (139, 140) also reported that MEG source localization of ictal and interictal high-frequency components was concordant with the iEEG results. The main limitation of these studies (137–142) is that they did not investigate HFOs as individual events in the time domain, as typically done with iEEG. Thus, it is not possible to establish whether the reported high-frequency components have the typical HFO morphology in the time domain and if they represent the same phenomenon observed in the time domain signals (27, 120, 135).

In summary, evidences from phantom and human studies (48–52, 137–141, 143) have shown that human HFOs can be non-invasively localized using MEG with an accuracy of few mm. Proper methods should be used for their localization because HFOs are weak oscillatory transients, which can be otherwise undesirably removed if localization methods are applied on averaged or coregistered events, as HFOs are not necessarily mutually phase locked. To date, there is no MEG study that has evaluated the clinical value of the HFO source localization in patients with MRE. To this purpose, further studies are needed to correlate the localization of the HFO zone with the resected area and the postsurgical outcome of patients with MRE. In addition, systematic investigations of simultaneous MEG, high-density scalp EEG, and iEEG would allow a direct comparison of the two

![Figure 7](https://www.frontiersin.org/landgraf/files/ microbes.png)
Figure 8 | Output of the detection and localization of high-frequency oscillations (HFOs) with magnetoencephalography (MEG) in three patients with focal epilepsy. Upper panels: topographic maps of the HFO detection in the MEG sensors. Each sensor is represented by a dot whose color reflects the number of detected HFOs at the time of the HFOs in the highest rate channel. Lower panels: source localization (wavelet maximum entropy on the mean) of HFOs in the ripple band for the same three patients. Sources are displayed in a normalized color scale. The epileptogenic zone, determined based on the available clinical information, is delineated with a red square. For patient 1 and 2, the sources of HFOs were totally or partially concordant with the epileptogenic region, whereas for patient 3, they were discordant. Source: adapted from von Ellenrieder et al. (50).

non-invasive modalities with the invasive gold standard for an optimal validation of the HFO source localization results.

DISCUSSION

Significant advances in diagnostic technology have largely improved efficacy and safety of epilepsy surgery. Yet, there is still an overriding need to identify and validate new reliable biomarkers that can precisely delineate the extent and location of the EZ. The identification of a more specific biomarker of epileptogenicity would (i) limit the need for long-term monitoring, (ii) enable a smaller and more precise resection of the EZ, (iii) improve the postsurgical outcome, and (iv) reduce the neurological deficits due to excessive surgical resection volume (37). To this purpose, HFOs have been the focus of investigations in the last decade given their emerging potential as a new precise and reliable biomarker of epileptogenicity (26, 27). The possibility to record such biomarkers with non-invasive techniques is crucial to expand their clinical utility (40). The non-invasiveness of the acquisition technique would (i) limit the need for presurgical long-term monitoring with iEEG; (ii) enable an optimized planning of the intracranial investigations; (iii) allow the assessment of the efficacy of therapeutic interventions without waiting for a seizure to occur, which can be associated with significant morbidity or mortality; (iv) permit definitive differential diagnosis of epilepsy from acute symptomatic seizures so treatment can begin immediately; and (v) enable the study of the differentiation between physiologic and pathologic HFOs by allowing the non-invasive investigation of healthy controls. This strong interest in localizing HFOs combined with MEG’s increasing role in the presurgical evaluation of patients with MRE (41–47) has led to increasing interest in detecting and localizing epileptic HFOs using MEG (48–52, 137–142).

In this review, we provided an overview of recent MEG studies that investigated HFOs in patients with epilepsy, discussing the proposed methodologies and the main findings reported in the current literature. The reported results are promising, but also foster the need for further MEG studies aiming at: (i) evaluating the accuracy of the HFO source localization against a ground truth given by simultaneous iEEG recordings in a large sample of patients, (ii) evaluating the clinical value of MEG in the HFO localization by looking at the patient’s postsurgical outcome and its correlation with the resection of the HFO generating areas; and (iii) establishing a definition of HFOs that is appropriate for
automatic detection in the context of MEG studies (as this could require different criteria compared to the iEEG literature), and possibly validating it against the patient’s outcome, rather than visual detection (118). Moreover, since high SNR is critical for the accurate localization of HFOs (48), the use of virtual channels (46, 49) should be further investigated. This will allow the localization of HFOs without relying on the spike location and avoiding biased results. Source localization could also be improved by merging simultaneous scalp EEG and MEG recordings: the complementary information of these two modalities might improve the source localization reliability (163), in particular for the MEM method (164).

CONCLUSION

The emerging potential of MEG for the detection and localization of HFOs in patients with epilepsy has been reported in several recent studies. The promising findings emphasize the need for large and robust studies necessary to establish and widespread the clinical use of HFOs in MEG data as a non-invasive biomarker of epileptogenic tissue. The possibility to detect and localize HFOs with MEG would open new possibilities of diagnosing and monitoring epilepsy. Non-invasively recorded HFOs could provide clinicians with an early epileptogenic marker in all patients with epilepsy, as well as in patients at high-risk of epilepsy, and could be also used to evaluate the effect of antiepileptic drug therapy. Finally, further studies are needed to address the question of whether the resection of the HFO zone localized with MEG can predict the postsurgical outcome of patients with MRE.

AUTHOR CONTRIBUTIONS

ET designed and drafted the work. She approved the version to be published and agreed to be accountable for all aspects of the work. JM contributed to the conception of the work and revised it critically for important intellectual content. He approved the version to be published and agreed to be accountable for all aspects of the work. PG contributed to the conception of the work and revised it critically for important intellectual content. She approved the version to be published and agreed to be accountable for all aspects of the work. PP contributed to the conception of the work and revised it critically for important intellectual content.
intellectual content. He approved the version to be published and agreed to be accountable for all aspects of the work. CP contributed significantly to the conception of the work and revised it critically for important intellectual content. He approved the version to be published and revised.

REFERENCES

1. Hirtz D, Thurman DJ, Gwinn-Hardy K, Mohamed M, Chaudhuri AR, Zulutsky R. How common are the common neurologic disorders? Neurology (2007) 68(5):326–37. doi:10.1212/01.wnl.0000278071.91524.4d
2. Sinha B, Mekh E, Moos K. Neuroimaging and neurogenetics of epilepsy in humans. Neuroscience (2009) 164(1):164–73. doi:10.1016/j.neuroscience.2009.08.056
3. Thurman DJ, Beghi E, Begley CE, Berg AT, Buchhalter JR, Ding D, et al. Standards for epidemiologic studies and surveillance of epilepsy. Epilepsia (2011) 52(7):2–26. doi:10.1111/j.1528-1167.2011.03121.x
4. Yoon D, Frick KD, Carr DA, Austin JK. Economic impact of epilepsy. Familial Career Development Fellowship of Harvard Medical School, Office for Faculty Development.

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Van’t Klooster MA, van Klink NE, Leijten FS, Zelmann R, Gebbink TA, Akiyama T, McCoy B, Go CY, Ochi A, Elliott IM, Akiyama M, et al. Focal resection of fast ripples on extraoperative intracranial EEG improves seizure outcome in pediatric epilepsy. *Epilepsia* (2011) 52(10):1802–11. doi:10.1111/j.1528-1167.2011.03199.x

Ibrahim GM, Fallah A, Snead OC III, Drake JM, Rutka JT, Bernstein M. Functional magnetic source imaging and mapping: a novel technique to assess language areas with epileptic high frequency oscillations are functionally isolated in nonlesional epilepsy. *Epilepsy Res* (2011) 21(10):743–7. doi:10.1016/j.eplepsyres.2011.07.006

Rutka JT, Go CY, Ochi A, Weiss SK, Rutka JT, et al. MEG predicts outcome following surgery for intractable epilepsy in children with normal or nonfocal MRI findings. *Epilepsia* (2007) 48(1):149–57. doi:10.1111/j.1528-1167.2006.00910.x

Benke T, Koylu B, Visani P, Karner E, Brenneis C, Bartha L, et al. Language lateralization in temporal lobe epilepsy: a comparison between fMRI and BOLD contrast functional MRI in stroke patients. *Epilepsia* (2006) 48(6):715–23. doi:10.1111/j.1528-1167.2005.00549.x

Murata Y, Sakatani K, Hoshino T, Fujiwara N, Kato Y, Nakamura S, et al. Effects of cerebral ischemia on evoked cerebral blood oxygenation responses and BOLD contrast functional MRI in stroke patients. *Stroke* (2006) 37(10):2514–20. doi:10.1161/01.wnl.0000239698.50656.3b

Benke T, Koylu B, Visani P, Karner E, Brenneis C, Barthel L, et al. Language lateralization in temporal lobe epilepsy: a comparison between fMRI and the Wada test. *Epilepsia* (2006) 47(1):1308–19. doi:10.1111/j.1528-1167.2006.00549.x

Knowlton RC, Razdan SN, Limdi N, Elgavish RA, Killen J, Blount J, et al. Does magnetoencephalography add to scalp video-EEG as a diagnostic tool in epilepsy surgery? *Epilepsy* (2004) 62(6):943–8. doi:10.1212/01.wnl.0000115122.81621.FE

Paulini A, Fischer M, Rampp S, Scheler G, Hofpagnet R, Kaltenhäuser M, et al. Lobar localization information in epilepsy patients: MEG—a useful tool in routine presurgical diagnosis. *Epilepsys Res* (2007) 76(2–3):124–30. doi:10.1016/j.eplepsres.2007.07.006

Khan S, Lefevre J, Balleit S, Michmizos KP, Ganesan S, Kitzbichler MG, et al. Encoding cortical dynamics in sparse features. *Front Hum Neurosci* (2014) 118:18. doi:10.3389/fnhum.2014.00175

Seo H, Holland K, Rose D, Rozhkov L, Fujiwara H, Byars A, et al. Multimodality imaging in the surgical treatment of children with nonlesional epilepsy. *Neurology* (2011) 76(1):41–8. doi:10.1212/WNL.0b013e318204a380

Tovar-Spinozo ZS, Ochi A, Rutka JT, Go CY, Ochsu H. The role of magnetoencephalography in epilepsy surgery. *Neurosur* Focus (2008) 25(3):E16. doi:10.3171/FOC.2008/25/9/E16

Tovar-Spinozo ZS, Ochi A, Rutka JT, Go CY, Ochsu H. The role of magnetoencephalography in epilepsy surgery. *Neurosurg Focus* (2008) 25(3):E16. doi:10.3171/FOC.2008/25/9/E16

Papanicolaou AC, Simos PG, Breier JI, Zouridakis G, Willmore LJ, Whelle JW, et al. Magnetoencephalographic mapping of the language-specific cortex. *J Neurosurg* (1999) 90(1):85–93. doi:10.3171/ns.1999.90

Simos PG, Papanicolaou AC, Breier JI, Whelle JW, Constantinou JE, Gormley WB, et al. Localization of language-specific cortex by using magnetic source imaging and electrical stimulation mapping. *J Neurosurg* (1999) 91(5):787–96. doi:10.3171/ns.1999.91.5.0787

Ver Hoe JW, Sawrie S, Killen J, Knowlton RC. Left mesial temporal sclerosis and verbal memory: a magnetoencephalography study. *J Clin Neurophysiol* (2008) 25(1):1–6. doi:10.1097/WNPB.0b013e3181636a6c

Lutzenberger W, Ripper B, Busse L, Birbaumer N, Kaiser J. Dynamics of gamma-band activity during an audiovisual working memory task in humans. *Clin Neurophysiol* (2002) 22(13):1935–44. doi:10.1016/s1388-2457(02)00618-6

Duffau H, Capelle L, Denvil D, Sichez N, Gatiogol P, Taillander L, et al. Usefulness of intraoperative electrical subcortical mapping during surgery for low-grade gliomas located within eloquent brain regions: functional results in a consecutive series of 103 patients. *J Neurosurg* (2003) 98(4):764–78. doi:10.3171/NS.2003.98.4.0764

Sakatani K, Murata Y, Fujiwara N, Hoshino T, Nakamura S, Kano T, et al. Comparison of blood-oxygen-level dependent functional magnetic resonance imaging and near-infrared spectroscopy recording during functional brain activation in patients with stroke and brain tumors. *J Biomed Opt* (2007) 12(6):062110. doi:10.1117/1.2823036

Murata Y, Sakatani K, Hoshino T, Fujiwara N, Kato Y, Nakamura S, et al. Effects of cerebral ischemia on evoked cerebral blood oxygenation responses and BOLD contrast functional MRI in stroke patients. *Stroke* (2006) 37(10):2514–20. doi:10.1161/01.STR.0000239698.50656.3b

Benke T, Koylu B, Visani P, Karner E, Brenneis C, Barthel L, et al. Language lateralization in temporal lobe epilepsy: a comparison between fMRI and the Wada test. *Epilepsia* (2006) 47(1):1308–19. doi:10.1111/j.1528-1167.2006.00549.x

Knowlton RC, Razdan SN, Limdi N, Elgavish RA, Killen J, Blount J, et al. Effects of epilepsy magnetic source imaging on intracranial electrode placement. *Ann Neurol* (2009) 65(6):715–23. doi:10.1002/ana.21660

Sutherlin WW, Mamkel A, Thyerel D, Maleeva T, Minarad Y, Philpott L, et al. Influence of magnetic source imaging for planning intracranial EEG in epilepsy. *Neurology* (2008) 71:990–6. doi:10.1212/01.wnl.000026591.29858.1a
113. Engel J Jr, Bragin A, Staba R, Mody I. High-frequency oscillations: what is normal and what is not? Epilepsia (2009) 50:598–604. doi:10.1111/j.1528-1167.2009.01917.x
114. Jacobs J, Levan P, Chitillon CD, Olivier A, Dubeau F, Gotman J. High frequency oscillations in intracranial EEGs mark epileptogenicity rather than lesion type. Brain (2009) 132(4):1022–37. doi:10.1093/brain/awn351
115. Jacobs J, Kerber K, Levan P, Dümppelmann M, Koiranthenberg R, Schulze-Bonhage A. Occurrence of high frequency oscillations depends on pathology in patients with focal cortical dysplasia. Epilepsia (2010) 35:Abst2. doi:10.1016/j.geneurobiol.2012.03.001
116. Gloss D, Nolan SJ, Staba R. The role of high-frequency oscillations in epilepsy surgery planning. Cochrane Database Syst Rev (2014) 1:CD010235. doi:10.1002/14651858.CD010235.pub2
117. van'T Klooster MA, Leijten FS, Huiskamp G, Ronner HE, Baayen JC, van Rijen PC, et al. High frequency oscillations in the intra-operative ECoG to guide epilepsy surgery (“The HFO Trial”): study protocol for a randomized controlled trial. Trials (2015) 16(1):1. doi:10.1186/s13063-015-0932-6
118. Höller Y, Kutil R, Klaffenböck L, Thomschewski A, Höller PM, Bathke AC, et al. High-frequency oscillations in epilepsy and surgical outcome. A meta-analysis. Front Hum Neurosci (2015) 9:574. doi:10.3389/fnhum.2015.00574
119. Zelmann R, Mari F, Jacobs J, Zijlmans M, Dubeau F, Gotman J. A comparison between detectors of high frequency oscillations. Clin Neurophysiol (2012) 123(1):106–16. doi:10.1016/j.clinph.2011.06.006
120. Worrell GA, Jerni K, Kobayashi Y. Recording and analysis techniques for high-frequency oscillations. Prog Neurobiol (2012) 98(3):265–78. doi:10.1016/j.pneurobio.2012.02.006
121. Zelmann R, Mari F, Jacobs J, Zijlmans M, Chander R, Gotman J. Occurrence of scalp-fast oscillations among patients with different spiking rate and their role as epileptogenicity marker. Epilepsia (2013) 106(3):345–56. doi:10.1111/epilepsypres.2013.06.003
122. Burns S, Frauscher B, Zelmann R, Haegeland C, Sarntcheh J, Gotman J. The morphology of high frequency oscillations (HFO) does not improve by automatic time-frequency analysis. PLos One (2014) 9(4):e94381. doi:10.1371/journal.pone.0094381
123. Crépon B, Navarro V, Hasboun D, Clémenceau S, Martinerie J, Baulac M, et al. Mapping interictal oscillations greater than 200 Hz recorded with intracranial macroelectrodes in human epilepsy. Brain (2010) 133(1):33–45. doi:10.1093/brain/awp277
124. Staba RJ, Wilson CL, Bragin A, Fried I, Engel J, Weiss SA, et al. Quantitative analysis of high-frequency oscillations (80–500 Hz) recorded in human epileptic hippocampus and entorhinal cortex. J Neurophysiol (2002) 88(4):1743–52. doi:10.1152/jn.00322.2002
125. Fedele T, van’t Klooster MA, Burns S, Zweiphennig W, van Klink N, Leijten F, et al. Automatic detection of high frequency oscillations during epilepsy surgery predicts seizure outcome. Clin Neurophysiol (2016) 127(9):3066–74. doi:10.1016/j.clinph.2016.06.009
126. Gardner AB, Worrell GA, Marsh E, Dlugos D, Litt B. Human and automated detection of high-frequency oscillations in clinical intracranial EEG recordings. Clin Neurophysiol (2007) 118(S5):1134–43. doi:10.1016/j.clinph.2006.12.019
127. Benar CG, Chauviere L, Bartolomei F, Wendling F. Pitfalls of high-pass filtering for detecting epileptic oscillations: a technical note on “false ripples. Clin Neurophysiol (2010) 121(3):301–10. doi:10.1016/j.clinph.2009.10.019
128. Ioannides AA, Kostopoulos GK, Laskaris NA, Liu L, Shihata T, Schellens M, et al. Timing and connectivity in the human somatosensory cortex from single trial mass electrical activity. Hum Brain Mapp (2002) 15:231–46. doi:10.1002/hbm.10023
129. Baumgartner C, Patai F, Lindinger G, Deecke L. Magnetoencephalography in focal epilepsy. Epilepsia (2000) 3(Suppl 41):S39–47. doi:10.1111/j.1528-1157.2000.tb01533.x
130. Sakuma K, Sekihara K, Hashimoto I. Neural source estimation from a time-frequency component of somatic evoked high-frequency magnetic
oscillations to posterior tibial nerve stimulation. Clin Neurophysiol (1999) 110(9):1585–8. doi:10.1016/S1388-2457(99)00120-0

151. Sekihara K, Nagarajan SS, Poeppel D, Marantz A, Miyashita Y. Reconstructing spatiotemporal activities of neural sources using an MEG vector beamformer technique. IEEE Trans Biomed Eng (2001) 48(7):760–71. doi:10.1109/30.930901

152. Papadelis C, Ahtam B, Nazarova M, Nimec D, Snyder B, Grant PE, et al. Cortical somatosensory reorganization in children with spastic cerebral palsy: a multimodal neuroimaging study. Front Hum Neurosci (2014) 8:725. doi:10.3389/fnhum.2014.00725

153. Guggisberg AG, Kirsch HE, Mantle MM, Barbaro NM, Nagarajan SS. Fast oscillations associated with interictal spikes localize the epileptogenic zone in patients with partial epilepsy. Neuroimage (2008) 39(2):661–8. doi:10.1016/j.neuroimage.2007.09.036

154. Hillebrand A, Barnes GGR. Beamformer analysis of MEG data. Int Rev Neurobiol (2005) 68:149–71. doi:10.1016/S0074-7742(05)68006-3

155. Vrba J, Robinson SE. Signal processing in magnetoencephalography. J Neurosci Methods (2001) 25(2):249–71. doi:10.1006/meth.2001.1238

156. Hillebrand A, Singh KD, Holliday IE, Furlong PL, Barnes GR. A new approach to neuroimaging with magnetoencephalography. Hum Brain Mapp (2005) 25(2):199–211. doi:10.1002/hbm.20102

157. Adjamin S, Worthen SF, Hillebrand A, Furlong PL, Chizh BA, Hobson AR, et al. Effective electromagnetic noise cancellation with beamformers and synthetic gradiometry in shielded and partly shielded environments. J Neurosci Methods (2009) 178(1):120–7. doi:10.1016/j.jneumeth.2008.12.006

158. Chowdhury RA, Lina JM, Kobayashi E, Grova C. MEG source localization of spatially extended generators of epileptic activity: comparing entropic and hierarchical Bayesian approaches. PLoS One (2013) 8:e55969. doi:10.1371/journal.pone.0055969

159. Heers M, Hedrich T, An D, Dubeau F, Gotman J, Grova C, et al. Spatial correlation of hemodynamic changes related to interictal epileptic discharges with electric and magnetic source imaging. Hum Brain Mapp (2014) 35(9):4396–414. doi:10.1002/hbm.22482

160. Heers M, Chowdhury RA, Hedrich T, Dubeau F, Hall JA, Lina JM, et al. Localization accuracy of distributed inverse solutions for electric and magnetic source imaging of interictal epileptic discharges in patients with focal epilepsy. Brain Topogr (2015) 29(1):162–81. doi:10.1007/s10548-014-0423-1

161. Lina JM, Chowdhury RA, Lemay E, Kobayashi E, Grova C. Wavelet-based localization of oscillatory sources from magnetoencephalography data. IEEE Trans Biomed Eng (2014) 61:2350–64. doi:10.1109/TBME.2012.2189883

162. Xiang J, Luo Q, Kotecha R, Korman A, Zhang F, Luo H, et al. Accumulated source imaging of brain activity with both low and high-frequency neuromagnetic signals. Front Neuroinform (2014) 8:57. doi:10.3389/fninf.2014.00057

163. Muravchik CH, Nehorai A. EEG/MEG error bounds for a static dipole source with a realistic head model. IEEE Trans Signal Process (2001) 49(3):470–84. doi:10.1109/78.905859

164. Chowdhury RA, Zerouali Y, Hedrich T, Heers M, Kobayashi E, Lina JM, et al. MEG-EEG information fusion and electromagnetic source imaging: from theory to clinical application in epilepsy. Brain Topogr (2015) 28(6):785–812. doi:10.1007/s10548-015-0437-3

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