

Review

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Keywords: Brain Computer Interface; data acquisition modalities; functional near-infrared spectroscopy; electroencephalography; functional magnetic resonance imaging; multimodal integration; signal processing



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Review of Multimodal Data Acquisition Approaches for Brain-Computer Interfaces

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Abstract: There have been multiple technological advancements that promise to gradually enable devices to measure and record signals with high resolution and accuracy in the domain of Brain Computer Interfaces (BCI). Multi-modal BCIs have been able to gain significant traction given the potential to enhance signal processing by integrating different recording modalities. In this review, we explore the integration of multiple neuroimaging and neurophysiological modalities including Electroencephalography (EEG), Magnetoencephalography (MEG), Functional Magnetic Resonance Imaging (fMRI), Electrocorticography (ECoG), and Single-Unit Activity (SUA). This multimodal approach leverages the high temporal resolution of EEG and MEG with the spatial precision of fMRI, the invasive yet precise nature of ECoG, and the single-neuron specificity provided by SUA. The paper highlights the advantages of integrating multiple modalities, such as increased accuracy and reliability, and discusses the challenges and limitations of multimodal integration. Furthermore, we explain the data acquisition approaches for each of these modalities. We also demonstrate various software programs that help in extracting, cleaning, and refining the data. We conclude this paper with discussion on the available literature highlighting recent advances, challenges, and future directions for each of these modalities.

Keywords: Brain computer interface; data acquisition modalities; functional near-infrared spectroscopy; electroencephalography; magnetoencephalography; electrocorticography; functional magnetic resonance imaging; multimodal integration; signal processing; multimodal BCI

1. Introduction

The Brain Computer Interfaces (BCIs) have emerged as transformative systems that establish a direct communication link between the human brain and external devices, such as computers or prosthetic limbs. By harnessing the power of neural activity, BCIs empower users to control these devices through their cognitive intentions, obviating the necessity for conventional input devices. [1] The applications enable a wide spectrum, ranging from assisting individuals afflicted with paralysis or locked-in syndrome to regain communication and control over their surroundings, to enabling users to manipulate intricate machinery or engage in video games using cognitive intent. However, despite their immense potential, BCIs remain in the nascent stages of development, beset by numerous technical and ethical challenges that demand scrutiny and resolution before widespread integration. [2] This innovation holds the promise to revolutionize the interface between humans and

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machines, particularly for individuals whose mobility or speech has been impaired due to injuries or disabilities.

The functioning of BCIs hinges upon the ability to capture and interpret the brain's intricate electrical signals. These signals arise from the brain's dynamic activity and serve as a reflection of its cognitive processes. [3] Several methodologies have been employed to measure and decode these signals, including electroencephalography (EEG), which records electrical activity from the scalp, and functional magnetic resonance imaging (fMRI), which captures changes in cerebral blood flow. Additionally, BCIs can harness other modalities such as functional near-infrared spectroscopy (fNIRS) and intracortical recordings, each contributing unique insights into neural activity.

This review explores the swiftly evolving landscape of BCIs, delving into their diverse modalities and the burgeoning possibilities the implementation of multi-modalities has to offer. While each modality boasts distinct advantages, they also bear inherent limitations in terms of resolution, spatial specificity, user invasiveness, robustness and generalization. The integration of multiple neuroimaging modalities, such as EEG, MEG, ECoG, and fMRI, has been proposed to overcome the limitations of single-modality systems. [4] The fusion of multiple modalities presents a solution to surmount these limitations, rendering BCIs more precise, dependable, and robust. For instance, EEG excels with high temporal resolution and non-invasiveness, albeit at the cost of spatial resolution, whereas fMRI affords superior spatial resolution but struggles with temporal precision. Integrating both modalities can bridge these gaps, harnessing their individual strengths synergistically.

The overarching objective of this review is to provide an extensive exploration of the integration of EEG/MEG + fMRI/fNIRS, ECoG + MEG/EEG, MUA/SUA + MEA/ECoG, and fMRI + ECoG/MEA modality combinations within the realm of BCIs. This exploration will encompass an evaluation of the strengths and limitations of each modality in isolation, [5] while delving into the transformative potential unlocked by their multimodal fusion. The review aims to illuminate the trajectory of research in this domain, delineating the challenges and opportunities, and thereby contributing to the ongoing advancement of BCIs. Through an in-depth analysis of modalities, their integration, and their applications, this review endeavors to provide a comprehensive insight into the present and future landscape of advanced BCIs.

2. BCI Modalities under Consideration

There have been multiple technological advancements that promise to gradually enable devices to measure and record signals with high resolution and accuracy in the domain of Brain Computer Interfaces (BCI), which were then utilized for building working models that can be used in various functionalities. Such utilities are as varied as academic endeavors to commercial memory storage devices currently initiated by Neuralink [6] to understand and abet in curing diseases and disabilities of the brain. We provide a review of various BCI modalities that provide an understanding of their working mechanisms. Subsequently, we embark upon exploring the data acquisition techniques for each of those modalities along with the limitations that each of the standalone setups pose.

2.1. Micro-Electrode Array (MEA)

Micro-electrode array techniques, or MEA, are a class of invasive brain-computer interfaces that utilize small, multi-electrode arrays to record the activity of individual neurons in the brain. MEA techniques provide a high spatial resolution and temporal precision, allowing for precise monitoring of neural activity at the single-cell level. MEA techniques use micro-fabricated arrays of electrodes that are capable of recording extracellular electrical signals from neurons in the brain. These electrodes are typically made of metal or conductive polymer and are very small, with diameters ranging from a few micrometres to a few hundred micrometres. The electrodes are placed in specific regions of the brain, such as the cortex, where they can record the activity of individual neurons or small groups of neurons.

MEA techniques offer several advantages over other invasive BCI methods, such as intracortical microelectrode arrays. MEA systems are less invasive, as the electrodes are smaller and can be

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inserted through smaller openings in the skull. Additionally, MEA systems can be easily fabricated using microfabrication techniques, making them more accessible and cost-effective than other invasive BCI methods. MEA systems have a wide range of potential applications, including the development of prosthetic devices that can restore motor function in individuals with paralysis. MEA techniques have also been used to study the basic mechanisms of neural function and to develop new treatments for neurological disorders such as epilepsy and Parkinson's disease.

Figure 1 displays MEA implants. A typical MEA biosensor has a sensory area of 700µm square with a length of 5 mm. Within this area, an alignment of 60 electrodes is designed in an 8x8 grid. The electrodes are usually placed across with separation of either 100, 200, or 500 µm. Planar TiN (titanium nitride) electrodes are available in sizes of 10, 20, and 30 µm that are used in variegated scenarios. However, it should be noted that there are variations in geometries that coincide with the anatomical properties. Such MEAs are present with a reference electrode that has been pre-integrated with the requisite substrate. [8] All electrodes have dual usability both for recording and for stimulation. Within the context of either cell or slice cultures, standard coatings are mandatorily applied on MEAs prior to their use for better cellular fixation and extension. Spike signals might be sensed at separations of up to 100µm from a neuron. Ideally, sites for signal emission are found in an area of 30µm² around the electrode. Closer proximities guarantee greater signal reception, and spatial resolution is inversely proportional to the signal units picked up by a single electrode, and that translates to lower work towards spike sorting. Multi-channel systems enable MEAs that in turn enable higher spatial resolutions. High-Dense-MEAs are another class of MEAs that have electrodes having a diameter of only 10µm housed within a space of only 30µm apart. There has been an inherent challenge in the processing of extra tiny electrodes whereas maintaining low levels of impedance and the noise simultaneously. This impediment has been eradicated by incorporation of TiN as the compound for manufacturing of the electrodes.

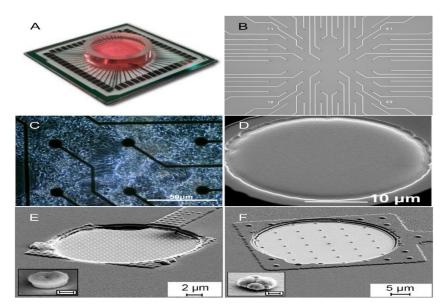


Figure 1. MEA implant at the cortical area. (A) MEA with 60 electrodes. (B) Figure showing arrangement of electrodes. (C) Neuronal growth on MEA. (D) SEM of MEA (E&F) SEM of nano-MEA. [Source: Reproduced under terms of the CC-BY license. [7] Copyright 2021, Authors..]

2.1.1. Data Acquisition Approaches and Limitations with a Standalone Setup

Data acquisition in standalone MEA systems is a critical process that involves capturing and processing neural signals directly from the electrodes implanted in the brain. In a standalone setup, the MEA operates independently, typically interfaced with a data acquisition system that amplifies, filters, and digitizes the recorded signals for further analysis. One of the primary advantages of this setup is its capacity for real-time monitoring and data processing, which is essential for applications such as BCIs and neurological research. [9]

However, there are some limitations to MEA techniques. One of the main challenges is the low signal-to-noise ratio of the recorded signals, which can make it difficult to distinguish individual neurons from the background noise. Additionally, the small size of the electrodes can make it difficult to maintain stable recordings over long periods of time, as the electrodes can shift or become damaged.

A significant challenge is the low signal-to-noise ratio (SNR) inherent in the recordings. [10] The small size of the electrodes, while beneficial for reducing invasiveness and increasing spatial resolution, can result in weak signals that are easily masked by background noise. This low SNR complicates the isolation of individual neuronal spikes, which is crucial for accurate neural decoding. [11] Additionally, the standalone nature of these systems means they are often constrained by power and processing capabilities, limiting their ability to handle large volumes of data or to perform complex real-time analysis. [12]

Another limitation is the long-term stability of the recordings. The electrodes, being in direct contact with the neural tissue, are susceptible to degradation over time due to biological responses such as gliosis, where glial cells proliferate around the electrode site, potentially leading to signal attenuation or loss. [13] Furthermore, electrode drift, caused by slight movements or shifts of the electrodes within the brain tissue, can disrupt the consistency of the recorded signals, making it difficult to maintain reliable long-term data acquisition. [14] These challenges necessitate ongoing research into materials and techniques that can enhance the durability and stability of MEA systems, as well as improvements in signal processing algorithms to better handle the inherent noise and variability in neural recordings. [15]

2.2. Electrocorticography (ECoG)

Electrocorticography (ECoG) is a technique for recording the electrical activity of the brain. ECoG involves placing a grid of electrodes directly on the surface of the brain, underneath the skull, and over the cortical tissue. This technique provides high spatial and temporal resolution, making it a valuable tool for BCI research.

ECoG has been used to decode motor and speech-related signals for BCI applications. The technique has been successful in decoding the movement of individual fingers and the intention to perform specific movements. ECoG can also be used to decode speech production and comprehension, with applications in speech rehabilitation for patients with speech disabilities.

There are some limitations to ECoG, including the invasiveness of the procedure, which requires a craniotomy to place the electrode grid on the surface of the brain. This increases the risk of complications and limits the number of potential patients. However, recent advances in electrode designs and implantation techniques have reduced the risks and made ECoG more feasible for human applications. [16] The major benefit of this approach is that although itself being an invasive modality, this method does not lead to permanent neuron damage or perfusion issues. Moreover, since the modality is localized in nature, most of the noise is automatically cancelled out. Nevertheless, as the modality cannot be coupled together with any other modality to provide cross cutting simulations, a complete coverage of all relevant brain events is not possible while conducting a study using this modality, thus limiting its adaptation.

However, ECoG provides a much better resolution than EEG and other non-invasive modalities. In one study, voltage spectra derived from auto-regressive spectral analyses between 0 and 200Hz were analyzed. [17,18] Each task was compared to the inactive period, and the R² values were calculated to determine the change in mu, beta and gamma rhythms associated to signals captured by electrodes strongly correlated with directional movements. It was demonstrated in this study that ECoG has a far broader frequency range than that of EEG. ECoG has been successfully applied to subjects suffering from instances of drug-refractory epilepsy. [16] Figure 2 shows ECoG study for finger activation.

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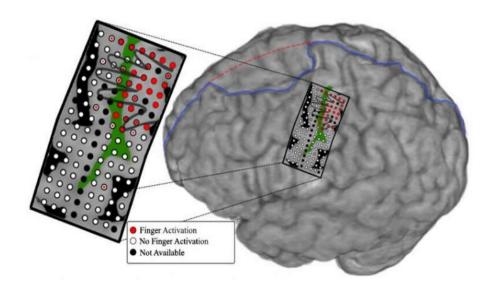


Figure 2. Showing various brain mappings of an ECoG study showing finder activation. [Source: Public Domain. Guy Hotson.].

2.2.1. Data Acquisition Approaches and Limitations with a Standalone Setup

ECoG is a method of acquiring electrical signals directly from the surface of the brain, providing high spatial and temporal resolution that is crucial for BCI research. In a standalone setup, ECoG involves the placement of a grid of electrodes on the cortical surface, under the skull, to record neural activity associated with motor and cognitive functions. This setup allows for the real-time decoding of motor intentions and speech-related signals, making it particularly valuable for applications such as the control of prosthetic devices and speech rehabilitation in patients with communication impairments. [19,20]

Despite its advantages, ECoG faces several limitations when used in a standalone configuration. The most significant challenge is the invasiveness of the procedure, which requires a craniotomy to place the electrodes on the brain's surface. [21] This surgical requirement increases the risk of complications, such as infections and hemorrhage, and limits the number of potential candidates for this technique. Furthermore, while ECoG provides better spatial resolution and frequency range compared to non-invasive modalities like EEG, its inability to be combined with other modalities restricts the scope of neural data that can be captured. This limitation poses challenges in obtaining a comprehensive understanding of brain activity, especially when studying complex cognitive tasks that may require multi-modal data integration. Additionally, electrode drift and long-term stability issues, similar to those encountered in other invasive recording methods, can affect the consistency of signal acquisition over time. [22]

Recent advancements in electrode design and implantation techniques have mitigated some of these risks, making ECoG a more viable option for human applications. [23] However, the standalone nature of the setup means that the data processing and analysis capabilities are constrained by the hardware's limitations, which may not always be sufficient for complex, real-time BCI applications. [24]

2.3. Electroencephalography (EEG)

Electroencephalography (EEG) stands as a prominent non-invasive brain imaging modality, characterized by an array of electrodes adorning the scalp, diligently capturing potential differences across distinct neuronal enclaves. Within the realm of Brain Computer Interfaces (BCIs), EEG assumes a pivotal role, with its operational architecture often tailored to specific frequency bands.

In the conventional BCI configuration, subjects are presented with stimuli at precise intervals. These stimuli elicit potentials within targeted cerebral regions, constituting the renowned event-related potentials (ERPs). The deliberate repetition of these ERPs enhances the signal-to-noise ratio,

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attenuating extraneous interference. Noise, discerned as spurious signals originating from artifacts such as cardiac activity, cranial structures, and respiratory processes, is systematically minimized. Recently, biometric authentication systems have been also explored with EEG modality banking on unique behavioral patterns to identify the user. [25]

The versatile applications of EEG-based BCI systems traverse a myriad of domains, ranging from prosthetic innovations to immersive video game experiences. This owes much to the modality's non-invasiveness, rendering it conducive to widespread use. With streamlined setup procedures and the absence of surgical prerequisites, EEG modalities emerge as the vanguard for forthcoming BCI explorations across multifarious scenarios. Also, EEG has been implemented in automated driving setups that check for driver drowsiness. [26] Within the realm of neuroimaging techniques, Electroencephalography (EEG) stands as a beacon of distinct advantages, setting it apart from its counterparts. A cascade of benefits distinguishes EEG, rendering it a quintessential tool for deciphering the intricacies of brain function.

Foremost is the unparalleled temporal resolution that EEG confers. This modality operates on a millisecond timescale, unrivalled by other neuroimaging techniques such as fMRI. This temporal acuity empowers EEG to unravel the chronology and dynamics of neural processes with exquisite precision. It excels in capturing the intricate temporal orchestration of cognitive phenomena and the precise timing of sensory perceptions. EEG's low cost and portability amplify its prowess. In the financial realm, EEG equipment remains economical, facilitating its widespread adoption across research settings. The added dimension of portability imbues EEG with remarkable flexibility, allowing seamless transitions between disparate locations. From hospital wards to research laboratories, and even field studies, EEG offers a versatile lens through which to examine brain function in diverse contexts. Furthermore, machine learning algorithms have found quantitative applications [27] for outcome predictions in traumatic brain injury, [28,29] eating disorder [30] and autism spectrum disorder [31] cases.

Safety emerges as a concern in neuroimaging, and EEG's non-invasiveness and radiation-free nature allay these apprehensions. Its innocuous profile renders it suitable for an extensive spectrum of subjects, including children and expectant mothers. This safety assurance complements EEG's unobtrusiveness, fostering an environment conducive to exploring neural dynamics. A hallmark attribute of EEG lies in its capability to discern neural responses to specific stimuli or events. This attribute is pivotal in dissecting cognitive processes like attention, memory, and language. By mapping brain activity onto cognitive functions, EEG empowers researchers to unveil the intricate interplay between neural patterns and cognitive operations. EEG shines as a formidable neuroimaging tool, armed with an arsenal of advantages that demarcate its uniqueness. Its temporal acumen, portability, and responsiveness to stimuli-driven dynamics set the stage for insightful inquiries into brain function. Amid the intricate landscape of neuroimaging modalities, EEG holds a distinctive position, offering a vantage point to decode the symphony of neural activity.

Among the software options available, we have found python and R to be rich in libraries that aid in data filtering and analysis in the course of this review, such as MNE-Python [32] and nin-Py [33] modules with a multitude of functionalities. Figure 3 shows calibration of EEG readings in R workspace.

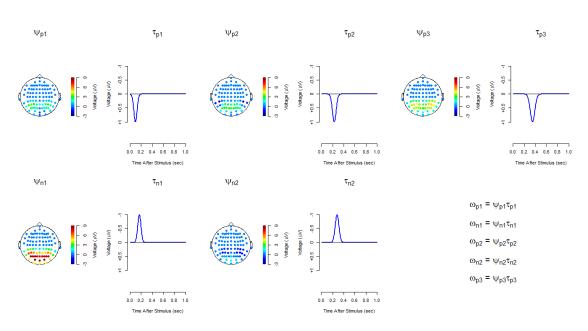


Figure 3. EEG action potential readings in R using eegkit library. Spatial Maps: The left column (A, C, E) shows the spatial distribution of component functions (ψ) across EEG channels in a 2D spatial layout. Color scales are limited to a range of -3 to 9 to standardize visual representation. Temporal Dynamics: The right column (B, D, F) depicts the temporal evolution of component functions (τ) over a time sequence ranging from 0 to 1 second, with y-axis limits set from -1 to 1. [Source: Self drawn with R-server hosted at http://sayantan.xyz:7779/].

2.3.1. Data Acquisition Approaches and Limitations with a Standalone Setup

EEG-based Brain Computer Interfaces (BCIs) epitomize a revolutionary paradigm, channeling EEG signals to govern external devices such as computers or prosthetic counterparts. This dynamic symbiosis hinges upon the discernment of distinct brain activity patterns aligned with specific commands. While EEG-based BCIs yield manifold advantages, their evolution is also marked by intrinsic constraints warranting scrutiny. Non-invasiveness and high temporal resolution stand as hallmarks of EEG-based BCIs. These attributes empower seamless interaction with the brain's cognitive canvas. However, a caveat arises in the form of spatial resolution. EEG's measurement of electrical activity on the scalp distills activity remote from the neural core. The resultant signals traverse a labyrinth of influences, from skull thickness to scalp conductivity, subtly tinging measurement accuracy.

Figure 4 shows a raw EEG data profile as available from Brain Compute Software. For the sake of consistency, we have retraced their steps with EEG signal data as they did with fNIRS data. We observed that a dedicated library such as eegUtils might also have been used to attain similar results. It must be noted that such dedicated libraries are also capable of providing additional topologies as regards the dataset versus its mapping in the brain. However, such niche results come at the cost of specialising the framework for a specific methodology and lead to loss of general capabilities in the BCI system.

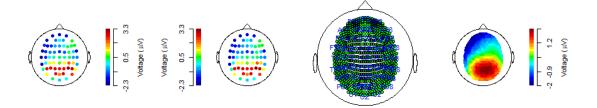


Figure 4. The spatial map of predicted EEG voltages mapped using eegkit library, obtained after spatial smoothing and using a dense cap configuration. This map highlights the estimated voltage distribution at the same time point, providing insights into the spatial patterns of EEG activity. [Source: Self drawn with R-server hosted at http://sayantan.xyz:7779/].

Susceptibility to artifacts emerges as a thorny challenge in EEG-based BCIs. The susceptibility to contamination by diverse noise sources – be it motion artifacts, eye blinks, or muscular contractions – complicates the demarcation between genuine brain signals and confounding interference. This reality introduces an intricacy that demands judicious signal processing.

Effective command control in EEG-based BCIs hinges upon rigorous user training. Mastering the generation of specific neural patterns linked to designated commands, be it cursor movement or prosthetic limb control, demands substantial effort and time investment. This learning curve underscores the cognitive effort required to harmonize intent with action. A critical limitation lies in EEG-based BCIs' reduced information content compared to counterparts like fMRI. While fMRI unveils brain region localization and activation specifics, EEG captures merely scalp-level electrical dynamics. This restricted information domain can hinder the decipherment of intricate cognitive processes or precise identification of implicated brain regions.

An ICA Model has been considered for the purposes of this review with source signals designated by columns of S, the mixing matrix is denoted by columns of M, and the noise signals are represented by E whereas zero mean columns are denoted by X. Its aim is to usually compute the unmixing matrix W in a manner that tcrossprod(X, W) denoted by Column S are as independent from other columns as possible. The goal is to find the orthogonal rotation matrix R such that the source signal is able to estimate S (= Y %*% R) [34] and are as independent as possible. [35] Additionally, the Infomax approach is able to calculate the orthogonal rotation matrix R that nearly maximizes the joint entropy of a nonlinear function. Also, the orthogonal rotation matrix R is computed using FastICA algorithm that again nearly maximizes the negentropy of the estimated source signals. Figure 5 displays brain maps after signal analysis and noise filtering activities.

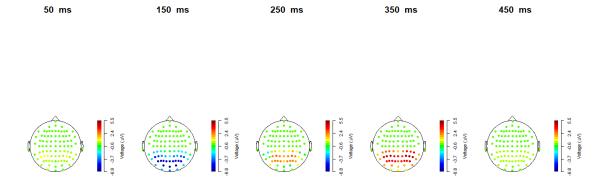


Figure 5. EEG signal analysis and noise filtering in R using eegkit library. The 2D plot represents the average control voltage at the specified time point. Channels related to ears, nose, and the reference electrode (Cz) have been excluded for clarity. The spatial distribution is shown using EEG coordinates in the 2D plane. [Source: Self drawn with R-server hosted at http://sayantan.xyz:7779/].

In summation, EEG-based BCIs marshal a transformative narrative by harmonizing brain and machine. Nonetheless, their trajectory is shadowed by limitations. The challenge of spatial precision, the battle against artifacts, the exigent training requirements, and the information ceiling – these facets underscore the need for a judicious balance between potential and constraints. Despite these limitations, EEG-based BCIs hold their ground as potent tools across diverse domains, propelling scientific inquiry in medicine, neuroscience, and human-computer interaction. The pursuit of innovation in this intersection remains unwavering, as researchers strive to chart new vistas and conquer existing limitations in the landscape of EEG-based BCIs.

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MEG [18] works by recording magnetic fields produced by potential differences in neurons using ultra-sensitive magnetic field sensors. (Proudfoot et al., 2014) This modality has a better spatial resolution than other non-invasive modalities. [36] The working mechanism involves passage of electric current within a strong magnetic field that interferes with the intracellular current of the dendrites thus mapping their structure and features. Figure 6 shows a schematic diagram of the workings of MEG.

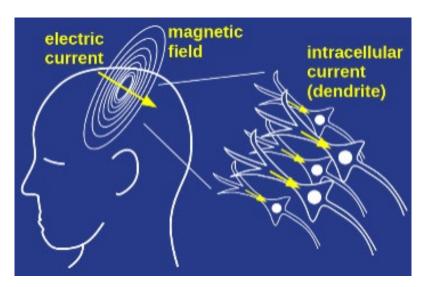


Figure 6. Schematic diagram of workings of MEG [Source: Reproduced under terms of the CC0 license. [37] Public Domain..]

However, high external magnetic noise in a general setting prevents application of this modality in a real-life scenario. Nevertheless, active research is being conducted to ensure viability of this modality as a high-resolution imaging system that might help in creating better commercial BCI products. Not much data analytics has been performed on this modality at the moment, and simple spatial filtering methods have been incorporated that take the properties from a geometric viewpoint of signal transmission in MEG into account and process artefacts specifically encountered in MEG-based BCI. Such techniques have enabled researchers to achieve real time synchronized feedback of stimulus in real time against actions performed. [38]

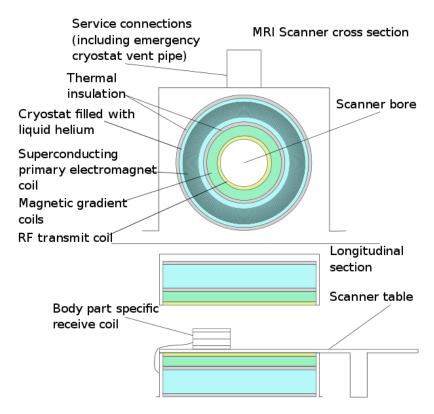


Figure 7. Schematic diagram of a conventional MRI scanner. Both MRI and fMRI machines have same architecture. [Source: Reproduced under terms of the CC-BY-SA license. [39] Copyright 2017, Author..]

More recently, EEG [40] and MEG [41] are also being analyzed together with fMRI to produce better temporal resolutions in results. A combination of results from all three is used in a study of memory processing wherein the details of localization, activation, and time course of a specific area of activation could be determined accurately. [42] Moreover, to counter the slow speed of the readings, acceleration techniques are applied to shallow learning. [43] deep learning [44] and ensemble learning [45–47] setups. A software implementation for data classification of fMRI has been attempted within Grid Workflow environment. In this paper, we attempt to review a modality encompassing grid workflow setup that can be easily adapted and scaled with any modality added with AI/ML capabilities.

2.4.1. Data Acquisition Approaches and Limitations with a Standalone Setup

Data acquisition in MEG involves capturing the weak magnetic fields generated by neuronal activity using highly sensitive sensors, typically superconducting quantum interference devices (SQUIDs). These sensors are arranged in an array around the subject's head to ensure comprehensive spatial coverage of the brain's magnetic activity. The process requires meticulous calibration and environmental control to minimize interference from external magnetic sources and to enhance the fidelity of the acquired data.

Given the extreme sensitivity of MEG sensors, data acquisition in a standalone setup often necessitates the use of magnetically shielded rooms (MSRs). These rooms are designed to attenuate external magnetic fields that could otherwise overwhelm the weak signals originating from neuronal activity. The effectiveness of shielding is paramount in ensuring the quality of the data, particularly when the setup is intended for use outside controlled laboratory environments.

Synchronizing the data acquisition process with external stimuli or other neuroimaging modalities (e.g., EEG or fMRI) is essential for capturing precise temporal correlations in brain activity. In standalone MEG setups, this often involves the use of high-precision clocks and synchronization

protocols to ensure that the timing of data acquisition is accurately aligned with the experimental conditions.

The standalone operation of MEG systems requires a high degree of technical expertise to manage the calibration, data acquisition, and noise mitigation processes. This complexity can be a barrier to the widespread adoption of MEG outside of specialized research institutions. Additionally, the need for constant monitoring and adjustment during data acquisition increases the operational burden, potentially limiting the feasibility of using MEG in routine clinical or commercial settings.

2.5. Functional-Magnetic Resonance Imaging (fMRI)

Functional Magnetic Resonance Imaging (fMRI) stands as a preeminent neuroimaging methodology, unravelling the intricate tapestry of brain activity through the lens of hemodynamic response. This technique capitalizes on the interplay between neural excitation and metabolic demands, offering insights into the neural symphony underpinning cognitive and motor functions. At its core, fMRI captures the dance of blood flow changes within the brain, reflective of underlying neural activity. The catalyst is the heightened metabolic requirements of activated neurons, prompting a surge in oxygen and nutrient delivery. This metabolic upsurge translates to an augmentation in blood flow, an effect termed the hemodynamic response.

In practice, fMRI unfolds as a collaborative endeavour between the subject and the scanner. The subject engages in tasks while the scanner, fortified by a potent magnetic field and radio waves, tracks blood flow oscillations. These oscillations metamorphose into vivid images of neural engagement, portraying the choreography of brain activity. This visual revelation lays bare the precise regions orchestrating a given task and unfurls the intricate interplay between these enclaves. Moreover, to counter the slow speed of the readings, acceleration techniques are applied on shallow learning, [48] deep learning [49] and ensemble learning [50–52] setups. A prominent asset of fMRI lies in its exquisite spatial resolution, a hallmark trait that unfurls the neural canvas in high definition. With spatial precision bordering on a few millimeters, fMRI excels in cartography of brain regions. This capability assumes particular importance in decoding neural circuits engaged in cognitive and motor tasks. Insights gleaned from fMRI have been pivotal in deciphering neural circuitry in various contexts, be it cognition, motion, or the perturbations induced by afflictions.

The non-invasive nature of fMRI emerges as another gem in its crown, rendering it a widely embraced and secure tool for probing the human brain. Unlike invasive techniques, fMRI circumvents surgical interventions, cementing its status as a low-risk procedure. This attribute equips researchers to navigate the neural landscapes of healthy individuals and patients grappling with neurological or psychiatric maladies. In concert with its spatial precision and safety, fMRI unfolds its versatility by embracing real-time exploration of brain function across a diverse spectrum of cognitive and motor activities. This expansive reach empowers fMRI to stand as a versatile sentinel, casting light upon the neural substrates of behavior and cognition.

More recently, EEG [53] and MEG [54] are also being analyzed together with fMRI to produce better temporal resolutions in results. A combination of results from all three is used in a study of memory processing wherein the details of localization, activation, and time course of a specific area of activation could be determined accurately. A software implementation for data classification of fMRI has been attempted within Grid Workflow environment. [55] In this paper, we attempt to review a modality encompassing grid workflow setup that can be easily adapted and scaled with any modality added with AI/ML capabilities. In essence, fMRI wields its magnetism to chart the ebbs and flows of neural choreography, revealing the hidden symphony of brain activity. Its spatial precision, non-invasiveness, and versatility weave a tapestry of insights that illuminate the contours of health and ailment within the human brain. As the human saga unfolds, fMRI remains a cornerstone, offering a non-invasive glimpse into the intricate drama of neural performance.

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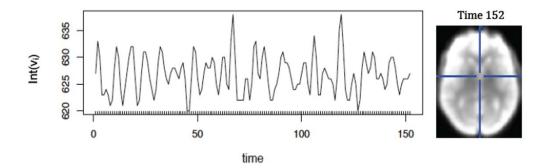


Figure 8. Conventional fMRI analysis: BOLD response to target stimuli. (a) healthy controls (Talairach coordinates x = 5, y = 36, z = 13). (b) subjects at clinical high risk for psychosis (Talairach coordinates x = 5, y = 36, z = 10). Random effects analysis, P < .001 (Bonferroni corrected for multiple comparisons). SAG: sagittal; COR: coronary; TRA: transversal; A: anterior; P: posterior; R: right; L: left. [Source: Reproduced under terms of the CC-BY license. [56] Copyright 2016, Authors...]

2.5.1. Data Acquisition Approaches and Limitations with a Standalone Setup

Functional magnetic resonance imaging (fMRI) stands as a pioneering tool in forging brain-computer interfaces (BCIs) that bridge the chasm between mental states and external device control. While fMRI based BCIs unveil a constellation of strengths, they also tread a path punctuated by nuanced limitations. The fusion of fMRI with BCIs heralds a realm of promise, underscored by remarkable advantages. Paramount among these is fMRI's exceptional spatial resolution, unveiling neural orchestrations with exquisite detail. [57] Additionally, fMRI's unique prowess in peering into the depths of intricate brain structures enriches our understanding beyond the purview of alternative techniques.

However, this is accompanied by constraints warranting judicious contemplation. The temporal resolution of fMRI emerges as a limitation, capturing neural dynamics at a deliberate pace spanning seconds. Consequently, rapid mental state transitions, such as those underpinning motor control or language processing, prove challenging for fMRI based BCIs. Further, fMRI's susceptibility to motion artifacts introduces a layer of complexity that perturbs certain BCI applications. The landscape of fMRI based BCIs is further shaped by the complexity and costs they entail. The procurement and upkeep of fMRI scanners entail substantial financial investments, alongside the need for specialized personnel to navigate their operation. This amalgam renders the translation of fMRI based BCIs into everyday environments, like homes or workplaces, an intricate endeavor.

Ethical considerations add a profound layer of discourse. The ability of fMRI to divulge an individual's mental state raises ethical dilemmas about privacy and consent. The potential misuse of this sensitive information poses concerns of unwarranted inference into personal thoughts or intentions. This interplay of technological prowess and ethical stewardship underscores the need for a conscientious balance. [58] Amid these contours, fMRI based BCIs showcase encouraging strides in areas ranging from motor control to communication and cognitive enhancement. The trajectory of relentless research and development augurs a future where fMRI-based BCIs will seamlessly intertwine with clinical and research realms, enriching our understanding and shaping human-computer interaction.

The alliance between fMRI and BCIs offers an avenue of exploration, charting the fusion of neural states and external control. However, the expedition is not devoid of challenges. Careful calibration of potentials and limitations navigates a course toward a future where fMRI based BCIs coalesce with ethical diligence, technical ingenuity, and transformative application. As this journey unfolds, the horizon of fMRI based BCIs shines with the promise of shaping a dynamic nexus between human cognition and technological advancement.

There is a push towards development of non-invasive techniques for functional imaging of the brain. fMRI displays contrast differences in the Blood Oxygen Level Dependent (BOLD) signal wherein the differential contrast is observed due to variable blood flow in the grey and the white

matter, where the latter receives greater blood flow. Figure 9 represents fMRI-readings from a 3D-image acquired for 152 time points.

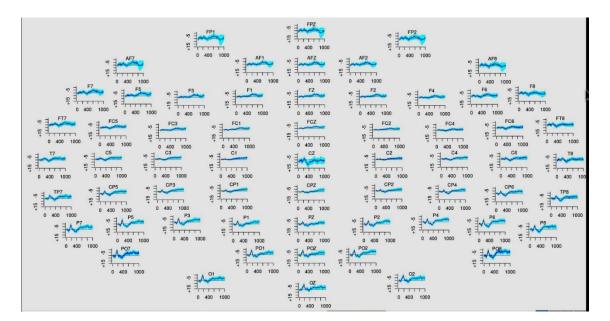


Figure 9. fMRI signal analysis in R. [Source: Self drawn with R-server hosted at http://sayantan.xyz:7779/].

2.6. functional-Near Infrared Spectroscopy (fNIRS)

Functional Near-Infrared Spectroscopy (fNIRS) emerges as a pivotal non-invasive neuroimaging modality, sculpting insights through the dance of near-infrared light and cerebral blood flow. Rooted in the tenet that neural activity is entwined with hemodynamic shifts, fNIRS deciphers the neural script by monitoring the absorption of light within the brain. fNIRS is another non-invasive modality that is performed by imaging hemodynamic signals of brain tissues using near-infrared spectrum. It works by measuring haemoglobin concentration changes in the blood stream. [59] Distinct virtues mark fNIRS as a noteworthy contender in the neuroimaging arena. Its non-invasive nature, devoid of ionizing radiation, resonates as a harbinger of safety, making it universally palatable. The harmony between neural activity and hemodynamics finds resonance in fNIRS, forging a pathway to trace the ebb and flow of blood. This orchestration is unveiled by detecting fluctuations in the concentrations of oxygenated and deoxygenated hemoglobin, unraveling the neural narrative. Moreover, fNIRS is able to provide a better temporal resolution. [60,61] Figure 10 shows various components of a fNIRS device.

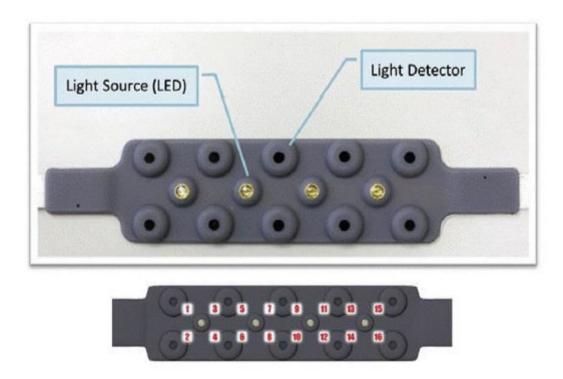


Figure 10. fNIRS device with light sources, detectors and (optode) channel measurement locations. [Source: Reproduced under terms of the CC-BY license. [62] Copyright 2015, Authors..]

The tapestry of fNIRS is embroidered with advantages that set it apart. Foremost is its temporal prowess, furnishing researchers with the ability to capture the nuances of brain activity in real-time. This dynamic window into neural choreography elucidates the symphony of cognition as it unfolds. The portability of fNIRS amplifies its utility, extending its reach from clinical domains to field studies, fostering flexibility in research endeavors. This modality is used in locomotory and ambulatory studies including yoga, [63] meditation, [64] robotic [65] and emotional [66] therapies. fNIRS is a virtuoso in unveiling the intricacies of deep brain structures that often evade other neuroimaging methodologies. The prefrontal cortex and the superior temporal gyrus, enclaves of cognitive richness, become accessible realms under fNIRS's gaze. Additionally, its robustness against movement artifacts renders it a preferred choice in studying infants and young children, circumventing the constraints that encumber EEG or fMRI. fNIRS stands out against fMRI in its portability and ability to filter out noise. Although, it provides a lower spatial resolution, fNIRS offers higher temporal resolutions and allows measurements of hemodynamic concentration changes with changing exertion of muscles. [67]

In concert, fNIRS emerges as a luminary, unraveling neural narratives in real-time with exquisite temporal fidelity and plunging into the enigmatic depths of the brain. Its non-invasive and transportable profile positions it as a versatile tool, finding its stride in clinical investigations, neurofeedback training, and sculpting the landscape of brain-computer interfaces. Amidst the symphony of neuroimaging methodologies, fNIRS's melodic notes resonate, tracing the cerebral rhythms that shape our cognitive tapestry. Each of these modalities have various stages of data acquisition, pre-processing and feature selection steps that allow the setup to filter and classify the input data from noise and amplify the same.

Figure 11 shows fNIRS workflow. The fNIRS workflow involves several key steps. First, researchers formulate their research question and design the study. Participants are then recruited and consented to that. In the pre-experiment phase, fNIRS probes are placed on the participant's scalp and calibrated if needed. During the experiment, data is collected while participants perform tasks or experience stimuli. Subsequently, collected fNIRS data is preprocessed to remove noise and transformed into hemoglobin concentration changes. Statistical analyses are applied to compare these changes between conditions. Interpreting the results involves relating hemodynamic patterns to the

research question and existing literature. The implications and limitations of findings are discussed before drawing conclusions and considering future research directions.

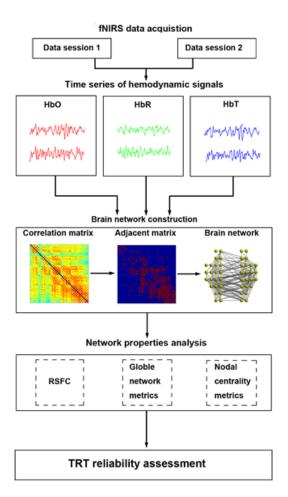


Figure 11. Schematic diagram showing fNIRS workflow. [Source: Reproduced under terms of the CC-BY license. [68] Copyright 2013, Authors..]

2.6.1. Data Acquisition Approaches and Limitations with a Standalone Setup

Functional Near-Infrared Spectroscopy (fNIRS) emerges as a burgeoning modality in the realm of Brain-Computer Interfaces (BCIs), leveraging its hallmark attributes of exceptional temporal resolution and portability. In the harmonious interplay between fNIRS and BCIs, insights into neural dynamics are harnessed, propelling interactions between cognition and external devices.

fNIRS-based BCIs harness the dynamic landscape of blood oxygenation changes within the brain to decipher distinct mental states. This intimate connection between hemodynamics and cognition serves as the foundation for translating these changes into control signals, enabling users to seamlessly navigate computer interfaces and devices. The allure of fNIRS-based BCIs rests in their non-invasive nature, imbuing comfort and accessibility. Their portability further amplifies their utility, flexibly extending from clinical settings to diverse environments.

We have subsequently reviewed data processing by taking publicly loaded datasets against these action points and pushed that over R engine to visualize the datasets as shown in Figure 12. One of the limitations of this library is this package can only read raw csv files generated by Hitachi ETG-4000. [69] However, the package is under active development and further development for additional support of different file types that looks promising. Both R and python provide libraries with similar syntax that give almost equivalent output. However, python versions are more mature and are able to provide finer results as compared to its R counterpart. The performance of both the software packages is equivalent to each other without any notable difference. However, it should also

be noted that for graphical user interfaces, R programs written in Shiny are much faster than the python graphical interface programs written in wx-widgets or tcl-Tk.

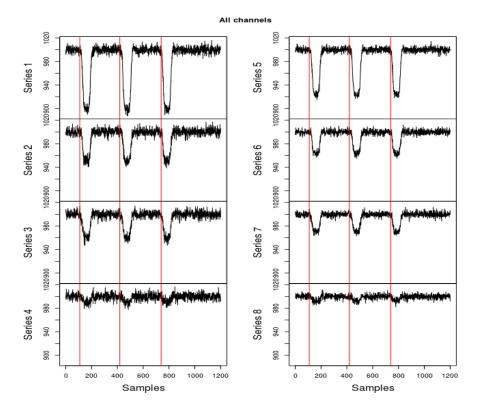


Figure 12. Raw fNIRS data using erzk/fnirsr library for R showing all channels in separate facets. This should enable spotting outliers. [Source: Self drawn with R-server hosted at http://sayantan.xyz:7779/].

The data is subsequently segregated and normalized in the R-workbench. We have used SparklyR functions and fnirsr library designed by Eryk Walczak [69] to review BCI analytics workflow. However, alternate functions can also be used. The data is further subjected to cleaning and pre-processing to obtain noise-filtered clean channels as observed in Figure 13. fNIRS signal is likely to show a linear trend which can be removed. The linear trend can be removed from all channels (recommended) or from a single channel.

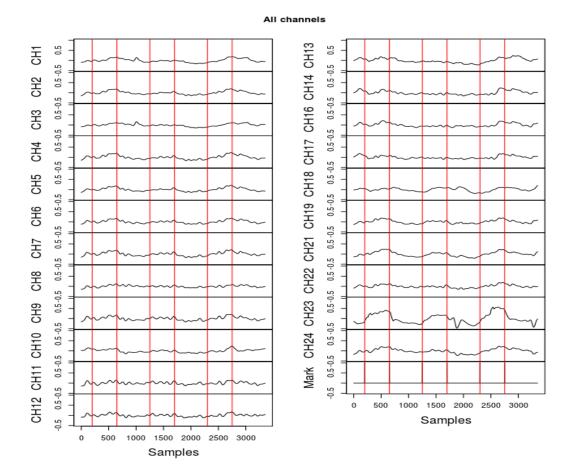


Figure 13. Segregated and normalised fNIRS data using erzk/fnirsr library for R. [Source: Self drawn with R-server hosted at http://sayantan.xyz:7779/].

Additionally, the library also supports HOMER2 datasets that provide additional information that can help a researcher visualize the data in faceted time series plots. Using the functions, a graph can be denoised and merged to create a comparative study as shown in Figure 14. Profiling methods with publicly available EEG data are available just like it was performed for fNIRS dataset. [70] HOMER2 [71] and OptoNet II [72] provide MATLAB scripts used for analyzing fNIRS data. The libraries are able to provide estimates and maps of brain activation areas. The application has been in a constantly evolving state since the early 1990s. It started off as the Photon Migration Imaging toolbox, that was subsequently reshaped into HOMER1. The application has undergone considerable changes to be rebranded as HOMER2. The application has a GUI interface similar to its older version but with easier and better support for group analyses and re-configuration of the processing stream. Also, it allows users to integrate their custom algorithms into the processing stream.

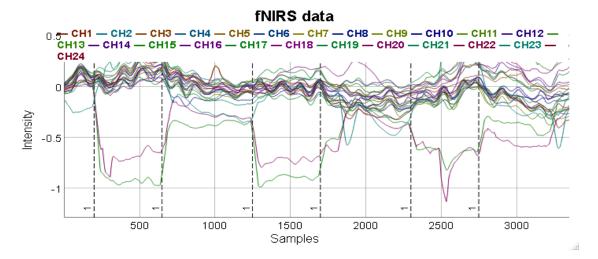


Figure 14. fNIRS noise filtered clean and merged data using erzk/fnirsr library for R. [Source: Self drawn with R-server hosted at http://sayantan.xyz:7779/].

The allure of fNIRS-based BCIs lies in their portability, non-invasiveness, and real-time insight into neural dynamics. However, the path forward is illuminated by the quest to augment specificity, accuracy, and resilience against noise. Future research endeavors must traverse the labyrinth of technological and methodological challenges, with the aim of unraveling the intricate symphony of cognition through fNIRS-based BCIs. In the evolving narrative of fNIRS-based BCIs, each limitation is a clarion call for innovation. As the precision of spatial localization and classification accuracy is refined, and as techniques to mitigate noise and artifacts advance, the symbiotic relationship between fNIRS and BCIs will burgeon, nurturing a landscape where thought meets action with unprecedented clarity and fidelity.

2.7. Multi-/Single-Unit Activity (MUA/SUA)

Multi-Unit Activity (MUA) and Single-Unit Activity (SUA) are critical components in the study of neural signals, providing insights into the collective and individual behavior of neurons, respectively. MUA refers to the electrical activity recorded from a group of neurons in proximity to an electrode, capturing the summed action potentials from multiple neurons. [73] This modality is particularly useful for understanding the overall neural activity within a specific brain region, as it provides a broad view of neuronal ensemble dynamics. On the other hand, SUA focuses on the activity of individual neurons, offering a more granular perspective by isolating and recording the action potentials from a single neuron. This is typically achieved through the use of high-impedance microelectrodes that can discern the firing patterns of individual neurons from the surrounding neural noise. The combination of MUA and SUA in neural recordings, often facilitated by microelectrode arrays (MEA), allows researchers to study both the macro-level population dynamics and micro-level neuronal specificity, contributing to a more comprehensive understanding of brain function. [74] These modalities are crucial in various applications, including brain-computer interfaces (BCIs) and the study of neural circuits in both normal and pathological state.

3. Integrated Setup for Advanced BCI

As the field of advanced BCIs continues to evolve, we can expect to see exciting applications that not only enhance our understanding of the brain but also improve the lives of individuals with neurological conditions, create more immersive entertainment experiences, and revolutionize human-computer interaction. The future of BCIs is filled with promise, and ongoing research and development will play a critical role in unlocking their full potential. Creating an integrated setup for advanced BCIs that combines various standalone BCI modalities involves careful planning, specialized equipment, and robust synchronization methods. Neurovascular coupling is the

phenomenon of a rise in cortical blood flow. It is a multi-step activity that encompasses all actions from stimulation of neurons to release transmitters that finally leads to vasoconstriction/dilation. Although associated to the same activity, EEG and fNIRS cover separate events in this cascade. The collaboration of these two methods offers a unique chance to examine cortical activity in an elaborate manner. EEG and fNIRS have very different but complementary temporal and spatial resolution. On one hand, cortical responses having high temporal resolution are detected by Evoked Potentials to a given stimulus, on the other hand fNIRS depends on that localization of changes in metabolism of hemoglobin with oxygen post activation. The combination of these modalities allows for a comprehensive examination of neurovascular coupling. For instance, EEG-fNIRS and MEG-fMRI setups enable researchers to simultaneously capture the fast electrical activity of neurons and the slower hemodynamic changes, offering a detailed view of the timing and spatial distribution of brain activity. These multimodal approaches are particularly powerful in understanding how neuronal and vascular components interact in both normal and pathological conditions. Placement sensors can be achieved in different manners. The first method is called adjacent positioning whereas the second method is called co-located measures. This method is limited to ring electrodes and requires transparent gel, which usually are non-conductive. As fNIRS and EEG measurements are recorded independently, it is important for simultaneous trigger synchronization with both data streams. [75] One of the key advantages of combining these modalities is the complementary nature of their spatial and temporal resolutions. EEG and MEG provide millisecond-scale temporal resolution, capturing the immediate neuronal responses to stimuli. In contrast, fMRI and fNIRS offer millimeter-scale spatial resolution, mapping the ensuing hemodynamic changes across the cortex. Together, they provide a multi-faceted view of brain activity, linking neuronal events to vascular responses. The combination of MEA, ECoG, EEG, MEG, fMRI, and fNIRS provides a powerful toolkit for investigating neurovascular coupling. Each modality offers unique strengths, and their integration allows for a more comprehensive understanding of the complex interactions between neuronal activity and blood flow in the brain.

3.1. EEG/MEG-fMRI/fNIRS Integration

Combining electroencephalography (EEG) and magnetoencephalography (MEG) with functional magnetic resonance imaging (fMRI) or functional near-infrared spectroscopy (fNIRS) provides a comprehensive view of brain activity by leveraging the strengths of each modality. EEG and MEG offer excellent temporal resolution, capturing neural dynamics in real-time, while fMRI and fNIRS provide superior spatial resolution, allowing for detailed mapping of brain regions involved in various cognitive tasks.

The integration of these modalities is particularly valuable in studying complex brain functions such as memory processing, language, and motor control. For instance, EEG/MEG can detect fast neural oscillations that occur during cognitive tasks, while fMRI/fNIRS can localize the brain regions responsible for these activities. This multimodal approach enables researchers to determine not only where brain activity occurs but also when it happens, offering a more complete picture of brain function. An exact synchronous action is attained with the NIRx Parallel Port Replicator, that is used to stimulate the incoming signal via USB ports and splits a single DB-25 (parallel port) input to four or more outputs. A general setup for a representative fMRI-EEG hybrid setup has been shown in Figure 15.

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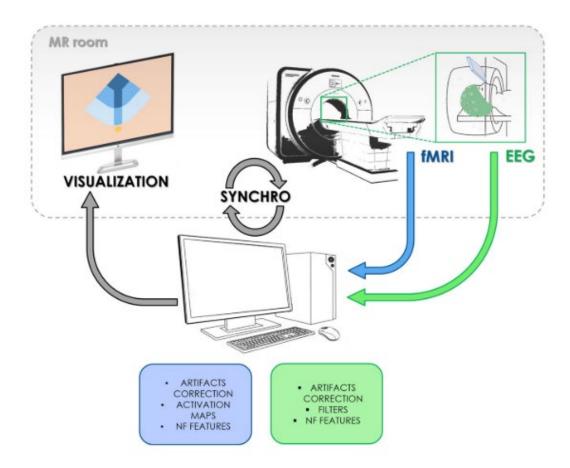


Figure 15. Schematic visualisation of the bimodal EEG-fMRI neurofeedback platform. [Source: Reproduced under terms of the CC-BY license. [76] Copyright 2020, Authors.

Similarly, EEG records brain activity via external electrodes placed on the scalp whereas fnirs measures chigs in hemoglobin concentrations using light sensors albeit placed at similar location on the scalp. EEG signals are filtered to remove electrical noise mostly using high and low pass filters. On the other hand, fNIRS filters correct for motion artifacts and physiological noise such as cardiac and respiratory rhythms. During the feature extraction step, EEG extracts feature like spectral power, event-related potentials (ERPs), or connectivity measures whereas fNIRS extracts features such as changes in oxygenated and deoxygenated hemoglobin levels. All these show that combining EEG and fNIRS does not lead to any conflicts, however, can help alleviate shortcomings of both. EEG and fNIRS can be combined to capture complementary information about neural and hemodynamic activity. Fusion techniques can also be applied that include concatenation, weighted averaging, or using them as inputs to a machine learning model. A combined schematic has been represented in Figure 16.

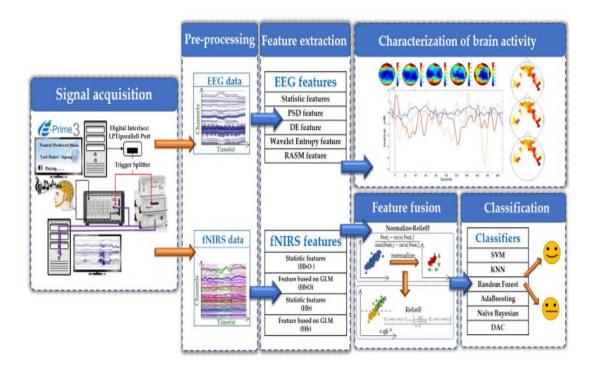
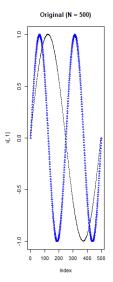
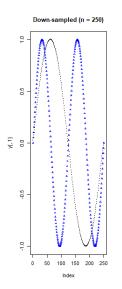


Figure 16. EEG-fNIRS hybrid setup. [Source: Reproduced under terms of the CC-BY license. [77] Copyright 2022, Authors..]

3.1.1. Data Processing in a Hybrid Setup

eegkit [78] is an R library that is powerful and provides precise results similar to its Python counterpart. It is a toolkit for Electroencephalography Data and uses a fast-discrete Fourier transform (eegfft) to calculate the power spectral density of EEG data and generates a plot of the power estimate using the plot (single channel) or image bar (multi-channel) function. The function employs stratigraphic series for bandpass filtering. Generally, the first column is designated as the location (e.g., depth), whereas the second column ought to be data value. Additionally, there is a plethora of other functions that perform multiple functionalities. Padfac function is used to Pad a specific column with zeros to (padfac x npts) points, where npts is the original number of data points. Flow gives the lowest frequency to bandpass, and fhigh provides the highest frequency to bandpass. win designates the Window type for bandpass filter: 0 translates to rectangular, 1 as Gaussian and 2 as Cosinetapered window or otherwise known as the Tukey window. Similarly, alpha that translates as inverse of standard deviation and acts as a measure of the width of the Dirichlet kernel, gives the Gaussian window parameter. [79] Also, p provides the Cosine-tapered (Tukey) window parameter, where p is the percent of the data series tapered parameter. The demean function is used to remove mean from data series whereas detrend function removes linear trend from data series. Similarly, addmean function is used to add mean value to bandpass result and the output function is used to show the output for either filtered series or bandpass filter window. The smallest frequency for plotting id determined using xmin function whereas xmax function gives the opposite, that is the largest frequency for plotting. A summary of plots can be visualized using the genplot function. Figure 17 displays band pass filtered and normalized EEG data in R. The EEG signals were filtered using a 4thorder Butterworth band-pass filter with cutoff frequencies set between 0.5 Hz and 30 Hz to remove noise and retain relevant brain activity frequencies. The filtered data was subsequently normalized to a z-score, providing a mean of 0 and a standard deviation of 1. Different colors represent signals from various EEG channels, showing the dynamic changes in brain activity across time.





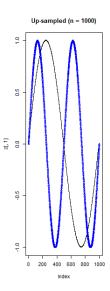


Figure 17. Band pass filtered and normalized EEG data in R. [Source: Self drawn with R-server hosted at http://sayantan.xyz:7779/].

Classification models can be trained to use the integrated EEG-fNIRS features using labelled data from cognitive tasks to classify brain states or conditions. Similarly, characterization of brain activity can be done by obtaining spatial information from EEG about localized neural activity. Then fNIRS can be used to capture changes in oxygenation, indicating an increase or decrease in brain activity. This combined data can provide a more comprehensive view of cognitive processes.

3.1.2. Challenges with a Hybrid Setup

While fNIRS-based BCIs unveil a promising horizon, they are not devoid of limitations. One pivotal constraint lies in the modest spatial resolution of fNIRS, rendering the precise localization of neural activity a challenge, in contrast to more robust techniques like fMRI. This intricacy poses a hurdle in the endeavor to attain specificity within BCIs.

Classification accuracy emerges as another arena of contention. While fNIRS-based BCIs have demonstrated potential in detecting mental states like concentration and relaxation, achieving high accuracy for intricate tasks such as speech recognition or motor control remains elusive. The interplay of individual neural variations constrained spatial and temporal resolution, and task complexity collectively contribute to this challenge. Noise and artifacts cast a further veil of complexity. [80] Motion artifacts and physiological noise introduce potential distortions that can erode the accuracy and reliability of fNIRS signals. Although signal processing and machine learning techniques offer mitigation strategies, the intrinsic vulnerability persists, necessitating ongoing vigilance.

3.2. fMRI-fNIRS Integration

The integration of fMRI and fNIRS data culminates in a detailed comprehension of brain function, harmonizing profound insights into neural dynamics furnished by fMRI with hemodynamic revelations from fNIRS. This union finds applications in cognitive exploration, clinical diagnostics, and propelling our insights into the symbiosis of brain and behavior. A synergy in signal acquisition can be achieved as fMRI captures neural activity with the aid of powerful magnetic fields whereas fNIRS quantifies hemoglobin concentrations by leveraging near-infrared light sensors. The alignment of fMRI and fNIRS data in spatial dimensions ensures their correspondence within specific brain regions. Co-registering these data enhances the fidelity of interpreting combined observations.

3.2.1. Data Processing in a Hybrid Setup

Achieving temporal synchrony between fMRI and fNIRS data is crucial to accurately timestamp events. The amalgamation of fMRI attributes (e.g., BOLD activation maps) and fNIRS traits (e.g., oxygenation dynamics) generates a more nuanced perspective during the Feature Extraction phase. [81] Employing advanced analytical techniques uncovers correlations and interplays between fMRI and fNIRS signals. Interpreting the unified dataset yields insights into the interplay between neural dynamics and hemodynamic alterations. Integrative fMRI-fNIRS configurations enable real-time neurofeedback, Brain-Computer Interfaces (BCIs), and examination of cerebral states during tasks. This also helps to address disparities in the spatial and temporal resolutions inherent in fMRI and fNIRS methodologies. Figure 18 displays a workflow of the hybrid fMRI-fNIRS setup.

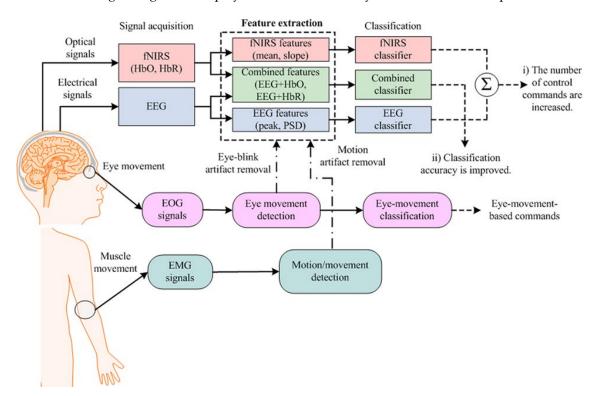


Figure 18. Schematic diagram of fMRI-fNIRS hybrid setup. [Source: Reproduced under terms of the CC-BY license. [82] Copyright 2017, Authors..]

3.2.2. Challenges with a Hybrid Setup

While the integration of fMRI and fNIRS offers significant advantages in capturing comprehensive brain activity, several challenges must be addressed to optimize the efficacy of this hybrid approach. One of the primary challenges is the disparity in spatial resolution between fMRI and fNIRS. fMRI provides high spatial resolution through the detection of BOLD signals, which can localize brain activity to specific cortical regions with millimeter precision. In contrast, fNIRS, while offering valuable hemodynamic information, generally has lower spatial resolution due to its reliance on near-infrared light passing through the scalp and skull. [83] This discrepancy can complicate the accurate co-registration of data, potentially leading to challenges in interpreting the spatial localization of brain activity.

The temporal resolution of fMRI and fNIRS also presents a challenge. fMRI measures BOLD signal changes with a temporal resolution on the order of seconds, which may not capture rapid neural fluctuations effectively. fNIRS, although it can track changes in hemodynamic responses with a finer temporal resolution, still does not match the millisecond precision of EEG or MEG. The integration of these differing temporal scales requires sophisticated synchronization techniques to ensure accurate alignment of data across modalities.

Both fMRI and fNIRS are susceptible to various types of noise and artifacts. fMRI is particularly prone to motion artifacts, physiological noise (e.g., heartbeats and respiration), and scanner-related

artifacts. fNIRS faces challenges related to motion artifacts, skin contamination, and physiological noise, such as variations in heart rate and respiration. [84] The presence of these artifacts can degrade the quality of the data and complicate the fusion process. Effective preprocessing and noise reduction techniques are essential to mitigate these issues.

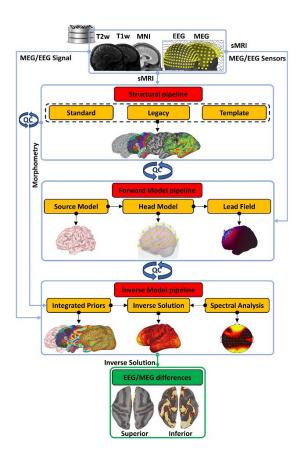
The fusion of fMRI and fNIRS data involves complex data integration processes. These processes include co-registration of spatial data, alignment of temporal signals, and the application of analytical techniques to extract meaningful correlations. The complexity of these tasks demands advanced computational tools and methodologies to ensure accurate and reliable integration of multimodal data.

3.3. ECoG-MEG/EEG Integration

ECoG, an invasive method that records electrical activity directly from the cortical surface, can be integrated with non-invasive modalities such as MEG and EEG to enhance the spatial and temporal resolution of brain activity monitoring. This multimodal approach leverages the strengths of each technique, providing a comprehensive view of neural dynamics. [85]

ECoG offers high spatial resolution and excellent signal-to-noise ratio due to its proximity to the neural tissue, making it particularly effective for localizing brain regions involved in specific functions. [86] However, its invasive nature limits its application to clinical settings, particularly in patients undergoing epilepsy surgery. In contrast, MEG and EEG are non-invasive techniques with lower spatial resolution but provide critical information on the temporal dynamics of neural activity. By integrating ECoG with MEG/EEG, researchers can combine the high spatial precision of ECoG with the broader coverage and non-invasiveness of MEG/EEG, leading to more accurate localization and timing of brain events.

Figure 19 shows the ECoG-EEG/MEG hybrid setup. Co-registering the data enhances the fidelity of interpreting combined observations. ECoG provides high spatial resolution and signal quality, particularly for cortical areas, which can complement the broader coverage of MEG/EEG. Combining invasive ECoG with non-invasive MEG/EEG can result in richer datasets, improving BCI system robustness. [88]



3.3.1. Data Processing in a Hybrid Setup

In a synchronous setup, ECoG, MEG, and EEG data are acquired simultaneously, allowing for the direct correlation of signals across different modalities. [89] This approach is particularly useful in clinical settings where real-time monitoring of brain activity is required. The challenge lies in the synchronization of the data streams from the different modalities, [90] which often have different sampling rates and data formats. Advanced signal processing techniques, such as time-frequency analysis and coherence analysis, are typically employed to align and integrate the data.

In some cases, ECoG and MEG/EEG data are acquired asynchronously, either due to technical constraints or the need to avoid excessive patient burden. While this approach may reduce the complexity of the experimental setup, it requires careful post-processing to align the data temporally. This often involves using external markers or events, such as auditory or visual stimuli, to synchronize the data retrospectively.

The combination of ECoG's high spatial resolution with MEG/EEG's excellent temporal resolution provides a detailed view of neural activity. This is particularly valuable in studies of brain function where both the precise location and timing of neural events are critical, such as in the investigation of seizure dynamics in epilepsy or the study of sensorimotor processes. [89,91]

ECoG provides localized information from specific brain regions, while MEG/EEG covers a broader area of the cortex. The multimodal approach allows for the investigation of interactions between local and distributed brain networks, offering a more holistic understanding of brain function.

ECoG's proximity to the cortical surface results in a higher signal-to-noise ratio compared to scalp-recorded EEG. Integrating ECoG with MEG/EEG can help validate findings from non-invasive recordings, reducing the likelihood of artifacts and improving the reliability of the results.

3.3.2. Challenges with a Hybrid Setup

The primary limitation of this multimodal approach is the invasiveness of ECoG, which requires surgical implantation of electrodes. This restricts its use to clinical populations, particularly patients undergoing neurosurgical procedures. [92] The risks associated with surgery, such as infection and hemorrhage, also limit the widespread application of ECoG.

Combining data from multiple modalities with different sampling rates, spatial resolutions, and noise characteristics presents significant technical challenges. Advanced computational tools are required to align, synchronize, and integrate the data, which can be computationally intensive and time-consuming.

ECoG requires surgical implantation, limiting its use to clinical settings or specific patient groups. Moreover, integrating invasive and non-invasive data involves complex signal processing and interpretation challenges.

The multimodal setup is resource-intensive, requiring specialized equipment, software, and expertise in both invasive and non-invasive neuroimaging techniques. This can limit the accessibility of this approach to well-funded research institutions and clinical centers. [93]

While MEG and EEG can be used for long-term monitoring of brain activity, the invasive nature of ECoG typically restricts its use to shorter periods. This limits the ability to study long-term neural dynamics, such as those involved in learning and memory, using the multimodal approach. [94]

$3.4.\ MUA/SUA\text{-}MEA/ECoG\ Integration$

The integration of MUA and SUA with MEA and ECoG provides a powerful framework for studying neural activity with high spatial and temporal resolution. This hybrid approach allows for a comprehensive examination of neuronal dynamics, facilitating insights into both single-neuron and population-level activity. The following subsections detail the data processing techniques and challenges associated with this integration. [95]

Integrating MUA/SUA with MEA/ECoG involves capturing neural signals from different sources with varying spatial resolutions. MUA and SUA are typically recorded using high-density electrodes or fine-tipped microelectrodes, while MEA provides a broader spatial coverage with multiple electrode sites. ECoG, positioned directly on the cortical surface, offers high-resolution spatial and temporal data. Achieving temporal synchronization between these modalities is crucial for accurate data fusion. Techniques such as timestamp alignment and event-based triggering are employed to ensure that data from MUA/SUA, MEA, and ECoG are temporally coherent. Figure 20 displays the various points on the skull where a hybrid setup can be arranged.

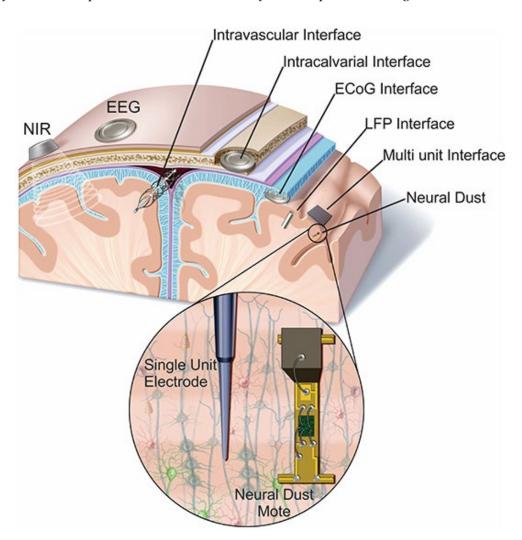


Figure 20. Schematic diagram of ECoG-EEG/MEG hybrid setup. [Source: Reproduced under terms of the CC-BY license.: [96] Copyright 2021, Authors..]

3.4.1. Data Processing in a Hybrid Setup

Data preprocessing is essential for minimizing noise and enhancing the quality of the integrated signals. For MUA and SUA, preprocessing steps include spike sorting, artifact removal, and filtering to isolate neural activity from background noise. MEA data often requires de-noising and correction for drift or signal attenuation. ECoG data, which may be affected by physiological artifacts such as muscle activity or electrical interference, requires filtering and artifact rejection. The preprocessing phase ensures that the signals are cleaned and standardized for subsequent analysis.

Feature extraction involves identifying relevant neural features from each modality. For MUA and SUA, features such as spike rates, firing patterns, and interspike intervals are extracted. MEA provides data on local field potentials (LFPs), which reflect population-level activity. [97] ECoG signals are analyzed for broadband oscillatory activity and event-related potentials. Integrating

features from these diverse sources involves aligning spatial and temporal domains, creating a unified dataset that captures both individual neuron activity and broader cortical dynamics.

Advanced analytical techniques are employed to fuse and interpret data from MUA/SUA, MEA, and ECoG. Methods such as cross-correlation, coherence analysis, and dimensionality reduction are used to explore relationships between single-unit and population-level activity. Machine learning algorithms and statistical models facilitate the integration of complex datasets, enabling researchers to identify patterns and interactions across different scales of neural activity. [98] Visualization tools are also utilized to represent the combined data, providing insights into the functional connectivity and spatial organization of neural networks.

3.4.2. Challenges in a Hybrid Setup

A significant challenge in integrating MUA/SUA with MEA and ECoG is the disparity in spatial and temporal resolution. MUA and SUA provide high-resolution data at the level of individual neurons, whereas MEA offers a broader view with moderate resolution, and ECoG captures activity from larger cortical areas. Aligning these data sources requires careful consideration of their respective resolution capabilities to avoid potential mismatches in spatial localization and temporal timing.

The complexity of integrating data from multiple sources is a major challenge. Combining MUA, SUA, MEA, and ECoG data involves sophisticated data processing and synchronization techniques. The need to align signals from different spatial scales and coordinate various preprocessing steps can complicate the integration process. Advanced computational methods and robust data handling protocols are necessary to manage and interpret the combined datasets effectively.

Artifact management is crucial when dealing with multiple recording modalities. Each method has its own sources of noise and interference, which can affect the quality of the integrated data. Ensuring effective artifact removal and noise reduction is essential to maintain the accuracy and reliability of the results. The presence of artifacts can obscure meaningful neural signals and hinder the integration process.

The practical aspects of setting up and maintaining a hybrid MUA/SUA-MEA/ECoG system pose additional challenges. Coordinating the use of different equipment, ensuring compatibility, and managing the physical setup can be complex. The integration process must accommodate the requirements of each modality while maintaining participant comfort and minimizing potential interference between recording systems.

3.5. fMRI-MEA/ECoG-EEG Integration

The integration of fMRI with MEA and ECoG combines the strengths of high-resolution spatial imaging from fMRI with the detailed temporal and spatial activity insights provided by MEA and ECoG. This hybrid approach enables a comprehensive analysis of brain function, enhancing our understanding of neural dynamics across different scales. The following subsections explore the data processing methods and challenges associated with fMRI-MEA/ECoG integration.

Combining fMRI with MEA and ECoG requires precise synchronization of data acquisition to align neural activity with the BOLD (Blood-Oxygen-Level Dependent) signals captured by fMRI. fMRI provides high spatial resolution images of brain activity but has limited temporal resolution. MEA and ECoG offer high temporal resolution and local cortical activity data. Achieving temporal alignment involves using event markers or triggers to synchronize data streams from fMRI, MEA, and ECoG systems. This may include integrating hardware solutions or software tools that ensure accurate timestamping across all modalities. Figure 21 displays an representation for interaction of fMRI and EEG modalities.

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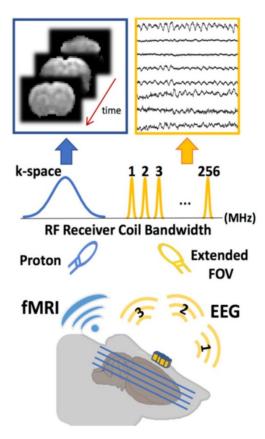


Figure 21. Schematic diagram of fMRI-EEG hybrid setup. [Source: Reproduced under terms of the CC-BY license.: [99] Copyright 2018, Authors..]

3.5.1. Data Processing in a Hybrid Setup

Preprocessing is essential to clean and standardize data from fMRI, MEA, and ECoG. fMRI data preprocessing includes motion correction, spatial normalization, and temporal filtering to remove noise and artifacts. MEA data requires spike sorting, noise filtering, and drift correction, while ECoG data needs artifact removal from physiological sources such as muscle activity. Combining these modalities involves aligning the preprocessed data to account for differences in spatial resolution and noise characteristics, ensuring that the integrated dataset reflects accurate neural and hemodynamic information.

Feature extraction involves identifying relevant metrics from each modality. For fMRI, features include BOLD signal changes and activation maps. MEA data is analyzed for spike rates, firing patterns, and local field potentials (LFPs), while ECoG provides information on oscillatory activity and event-related potentials. Integrating features from these sources requires aligning spatial coordinates and temporal events. Techniques such as spatial normalization, feature concatenation, and multi-modal fusion are used to create a unified representation of brain activity that combines the spatial detail of fMRI with the temporal precision of MEA and ECoG.

The integration of fMRI with MEA and ECoG can be used for real-time applications such as neurofeedback and Brain-Computer Interfaces (BCIs). Real-time processing of fMRI, MEA, and ECoG data allows for dynamic monitoring and modulation of brain activity, enabling feedback mechanisms based on neural and hemodynamic states. This integration supports advanced applications in cognitive training, rehabilitation, and personalized neuromodulation. [100]

3.5.2. Challenges in a Hybrid Setup

A major challenge in integrating fMRI with MEA and ECoG is the discrepancy in spatial and temporal resolution. fMRI provides high spatial resolution but has limited temporal resolution, while MEA and ECoG offer high temporal resolution but with different spatial scales. Aligning these data

sources requires careful consideration of their resolution differences to avoid mismatches in spatial localization and temporal synchronization.

Combining fMRI, MEA, and ECoG data involves complex data processing and integration procedures. The need to align signals from different spatial and temporal domains requires sophisticated computational methods and robust data handling protocols. The integration process must address issues related to data fusion, feature alignment, and dimensionality reduction to create a cohesive dataset.

Managing artifacts is a significant challenge when integrating data from multiple recording modalities. Each modality is susceptible to different types of artifacts, such as motion artifacts in fMRI and electrical noise in MEA and ECoG. Effective artifact removal and noise reduction techniques are necessary to ensure the quality and reliability of the integrated data. The presence of artifacts can obscure meaningful signals and complicate the integration process.

Handling the large volumes of data generated by fMRI, MEA, and ECoG requires substantial computational resources and storage capacity. The integration of high-resolution signals from multiple sources necessitates efficient data management strategies and advanced analytical tools. Researchers must ensure that their infrastructure can support the processing and storage demands of comprehensive datasets.

3.6. fMRI/ECoG-MEG/EEG Integration

Integrating ECoG with MEG or EEG offers a multifaceted approach to studying brain activity. This hybrid setup leverages the high spatial resolution of ECoG with the superior temporal resolution of MEG/EEG, facilitating a comprehensive understanding of neural dynamics. Below, we delve into the data processing methods and challenges associated with ECoG-MEG/EEG integration.

Synchronizing data acquisition from ECoG and MEG/EEG is crucial for accurate integration. ECoG records cortical activity directly from the brain's surface, providing high spatial and temporal resolution, whereas MEG and EEG capture electrical activity from the scalp with varying spatial resolutions. To achieve temporal alignment, precise event markers or triggers are used to coordinate data collection across both modalities. This may involve using synchronized recording systems or software tools to ensure consistent timestamps. Figure 22 shows a schematic flow diagram of fMRI-MEG/EEG hybrid setup.

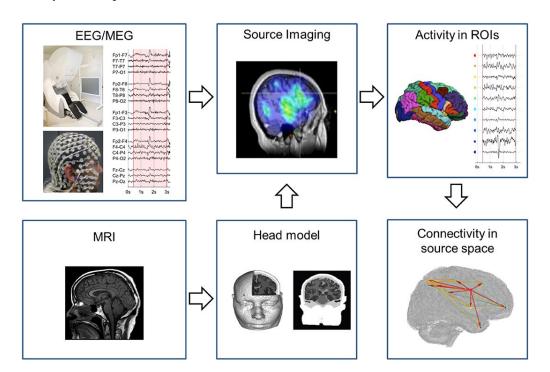


Figure 22. Schematic diagram of fMRI-EEG hybrid setup. [Source: Reproduced under terms of the CC-BY license.: [101] Copyright 2019, Authors..]

3.6.1. Data Processing in a Hybrid Setup

The hybrid setup of ECoG-MEG/EEG can be applied in real-time scenarios such as neurofeedback and Brain-Computer Interfaces (BCIs). Real-time analysis of integrated data allows for dynamic monitoring and modulation of brain activity. This integration supports advanced applications in cognitive training, brain state classification, and personalized neuromodulation, providing immediate feedback based on neural dynamics.

Feature extraction from ECoG includes identifying neural oscillations, event-related potentials (ERPs), and local field potentials (LFPs). MEG and EEG data provide complementary features such as oscillatory patterns and spectral power. Integrating features involves aligning spatial coordinates and temporal events, which can be achieved through methods such as spatial normalization and temporal interpolation. Techniques like multi-modal fusion and concatenation are employed to combine these features, resulting in a unified representation of brain activity. [102]

Preprocessing involves cleaning and standardizing data from both ECoG and MEG/EEG. For ECoG, preprocessing includes removing noise from electrical interference and correcting for signal drift. MEG and EEG preprocessing typically involves filtering to remove artifacts from muscle activity, eye movements, and other sources of noise. Both ECoG and MEG/EEG data are filtered to remove irrelevant frequencies and enhance the signals of interest. Artifacts common to ECoG and MEG/EEG need to be addressed through advanced filtering techniques and artifact correction algorithms.

3.6.2. Challenges in a Hybrid Setup

One of the primary challenges in integrating ECoG with MEG/EEG is the discrepancy in spatial and temporal resolution. ECoG provides high spatial resolution and precise temporal data from the cortical surface, whereas MEG/EEG offers broader spatial coverage but with lower resolution. Aligning these different data types requires addressing issues related to spatial localization and temporal synchronization to ensure accurate representation of brain activity.

The integration of ECoG with MEG/EEG involves complex data processing and fusion. Combining high-resolution signals from ECoG with broader, less localized signals from MEG/EEG requires advanced computational methods. Ensuring that data from both modalities is accurately aligned and integrated poses significant technical challenges, necessitating robust algorithms and processing techniques.

Artifacts can significantly impact the quality of both ECoG and MEG/EEG data. ECoG data may be affected by electrical noise from external sources, while MEG/EEG signals can be contaminated by muscle activity, eye movements, and other physiological artifacts. Effective artifact removal and noise reduction are critical to ensuring the accuracy and reliability of the integrated data. This requires sophisticated preprocessing techniques and careful management of potential sources of interference.

Handling the large volumes of data generated by ECoG and MEG/EEG requires substantial computational resources and storage capacity. The integration process must manage the high-resolution data from ECoG alongside the broader datasets from MEG/EEG. Efficient data management strategies and advanced analytical tools are necessary to process and analyze the combined dataset effectively.

4. Discussion

The integration of multiple brain imaging modalities, such as, fNIRS, EEG, MEG, ECoG, fMRI, MEA and MUA, addresses the limitations of individual modalities while enhancing the overall understanding of brain activity. Each modality offers unique strengths and challenges, and their combination provides a more comprehensive view of neural dynamics and brain function.

[103] Integrating multiple modalities can improve the accuracy and precision of BCI performance. Different modalities can be used together to provide both spatial and temporal information about brain activity. [104] Whereas fNIRS has high temporal resolution (in the order of

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milliseconds) and can be used to measure brain activity in real time. [105] However, fNIRS has limited spatial resolution, and it is susceptible to motion artifacts and noise from systemic physiological activity. While fMRI has high spatial resolution but limited temporal resolution. On the other hand, ECoG provides excellent spatial and temporal resolution by recording directly from the cortical surface. It allows precise localization of brain activity but is invasive and limited to clinical settings. Whereas MEA and MUA further complement these modalities by offering detailed insights into neural activity at the microelectrode level. MEA records activity from multiple neurons simultaneously, providing high spatial and temporal resolution data. MUA, on the other hand, captures the combined activity of multiple neurons within a localized area, offering insights into neural population dynamics.

5. Conclusions

The canvas of BCI applications extends far and wide, encompassing clinical interventions, assistive technologies, and immersive gaming and entertainment domains. These applications epitomize the versatility of BCIs, from restoring motor function in paralysis patients to augmenting cognitive abilities in the neurotypical populace. BCIs also stand poised to revolutionize interactive experiences in the realm of gaming and entertainment. Each modality has a different spatial and temporal resolution, which can be combined to provide a more comprehensive picture of brain activity. [106] We have explored various hybrid modalities in their earlier studies [107–112] out of which EEG-fNIRS stands out [113–116]. This review has outlined the myriad merits stemming from multimodal integration, encompassing heightened spatial and temporal resolution, diminished susceptibility to noise and artifacts, and augmented precision. The integration of fNIRS, EEG, fMRI, MEG, ECoG, and MUA/SUA modalities provides a powerful toolkit for studying brain activity from multiple perspectives. By combining the strengths and addressing the limitations of each modality, researchers can achieve a more comprehensive and accurate understanding of neural processes. This multimodal approach not only enhances spatial and temporal resolution but also improves the ability to study complex brain dynamics and connectivity. As technology advances, the potential for multimodal BCIs will continue to grow, offering new opportunities for both research and clinical applications. By unveiling the benefits and complexities of multimodal fusion and illustrating diverse application scenarios, this paper contributes to the nascent narrative of BCIs. As the trajectory of BCIs continues to evolve, future research endeavors should pivot towards refining multimodal integration methods and exploring new horizons for advanced BCIs across diverse domains.

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References

 Lee, D.Y.; Jeong, J.H.; Shim, K.H.; Kim, D.J. Classification of Upper Limb Movements Using Convolutional Neural Network with 3D Inception Block. In Proceedings of the 8th Int. Winter Conf. Brain-Computer Interface, BCI 2020; 2020.

32

- 2. Shih, J.J.; Krusienski, D.J.; Wolpaw, J.R. Brain-Computer Interfaces in Medicine. *Mayo Clin Proc* **2012**, *87*, 268, doi:10.1016/J.MAYOCP.2011.12.008.
- 3. Guger, C.; Allison, B.; Leuthardt, E.C. Brain-Computer Interface: A State-of-the-Art Summary 2; 2015;
- 4. Khorev, V.; Kurkin, S.; Badarin, A.; Antipov, V.; Pitsik, E.; Andreev, A.; Grubov, V.; Drapkina, O.; Kiselev, A.; Hramov, A. Review on the Use of Brain Computer Interface Rehabilitation Methods for Treating Mental and Neurological Conditions. *J Integr Neurosci* 2024, 23, 125, doi:10.31083/J.JIN2307125/1757-448X-23-7-125/FIG9.PNG.
- 5. Suk, H. II; Fazli, S.; Mehnert, J.; Müller, K.R.; Lee, S.W. Predicting BCI Subject Performance Using Probabilistic Spatio-Temporal Filters. *PLoS One* **2014**, *9*, doi:10.1371/journal.pone.0087056.
- 6. Musk, E.; Neuralink An Integrated Brain-Machine Interface Platform with Thousands of Channels. *bioRxiv* **2019**, 703801, doi:10.1101/703801.
- Schulte, S.; Gries, M.; Christmann, A.; Schäfer, K.H. Using Multielectrode Arrays to Investigate Neurodegenerative Effects of the Amyloid-Beta Peptide. *Bioelectron Med* 2021, 7, 1–7, doi:10.1186/S42234-021-00078-4/FIGURES/2.
- 8. Nicolas-Alonso, L.F.; Gomez-Gil, J. Brain Computer Interfaces, a Review. Sensors 2012, 12, 1211–1279, doi:10.3390/s120201211.
- 9. Obien, M.E.J.; Deligkaris, K.; Bullmann, T.; Bakkum, D.J.; Frey, U. Revealing Neuronal Function through Microelectrode Array Recordings. *Front Neurosci* **2015**, *9*, 423, doi:10.3389/FNINS.2014.00423/BIBTEX.
- 10. Buzsáki, G.; Mizuseki, K. The Log-Dynamic Brain: How Skewed Distributions Affect Network Operations. *Nature Reviews Neuroscience* 2014 15:4 **2014**, 15, 264–278, doi:10.1038/nrn3687.
- 11. Einevoll, G.T.; Franke, F.; Hagen, E.; Pouzat, C.; Harris, K.D. Towards Reliable Spike-Train Recordings from Thousands of Neurons with Multielectrodes. *Curr Opin Neurobiol* **2012**, 22, 11–17, doi:10.1016/J.CONB.2011.10.001.
- 12. Franke, F.; Jäckel, D.; Dragas, J.; Müller, J.; Radivojevic, M.; Bakkum, D.; Hierlemann, A. High-Density Microelectrode Array Recordings and Real-Time Spike Sorting for Closed-Loop Experiments: An Emerging Technology to Study Neural Plasticity. *Front Neural Circuits* **2012**, *6*, 36500, doi:10.3389/FNCIR.2012.00105/BIBTEX.
- 13. Grill, W.M.; Mortimer, J.T. Stability of the Input-Output Properties of Chronically Implanted Multiple Contact Nerve Cuff Stimulating Electrodes. *IEEE Transactions on Rehabilitation Engineering* **1998**, *6*, 364–373, doi:10.1109/86.736150.
- Williams, J.C.; Rennaker, R.L.; Kipke, D.R. Long-Term Neural Recording Characteristics of Wire Microelectrode Arrays Implanted in Cerebral Cortex. Brain Research Protocols 1999, 4, 303–313, doi:10.1016/S1385-299X(99)00034-3.
- 15. Bakkum, D.J.; Frey, U.; Radivojevic, M.; Russell, T.L.; Müller, J.; Fiscella, M.; Takahashi, H.; Hierlemann, A. Tracking Axonal Action Potential Propagation on a High-Density Microelectrode Array across Hundreds of Sites. *Nature Communications* 2013 4:1 **2013**, 4, 1–12, doi:10.1038/ncomms3181.
- Centracchio, J.; Sarno, A.; Esposito, D.; Andreozzi, E.; Pavone, L.; Gennaro, G. Di; Bartolo, M.; Esposito, V.; Morace, R.; Casciato, S.; et al. Efficient Automated Localization of ECoG Electrodes in CT Images via Shape Analysis. *Int J Comput Assist Radiol Surg* 2021, doi:10.1007/s11548-021-02325-0.
- 17. Leuthardt, E.C.; Schalk, G.; Wolpaw, J.R.; Ojemann, J.G.; Moran, D.W. A Brain-Computer Interface Using Electrocorticographic Signals in Humans. *J Neural Eng* **2004**, *1*, 63–71, doi:10.1088/1741-2560/1/2/001.
- 18. Guger, C.; Allison, B.; Leuthardt, E.C. Brain-Computer Interface A State-of-the-Art Summary 2; 2015; ISBN 978-3-319-25188-2.
- 19. Schalk, G.; Leuthardt, E.C. Brain-Computer Interfaces Using Electrocorticographic Signals. *IEEE Rev Biomed Eng* **2011**, *4*, 140–154, doi:10.1109/RBME.2011.2172408.
- 20. Kellis, S.; Miller, K.; Thomson, K.; Brown, R.; House, P.; Greger, B. Decoding Spoken Words Using Local Field Potentials Recorded from the Cortical Surface. *J Neural Eng* **2010**, *7*, doi:10.1088/1741-2560/7/5/056007.
- 21. Lachaux, J.P.; Rudrauf, D.; Kahane, P. Intracranial EEG and Human Brain Mapping. *J Physiol Paris* **2003**, 97, 613–628, doi:10.1016/J.JPHYSPARIS.2004.01.018.
- 22. Chao, Z.C.; Nagasaka, Y.; Fujii, N. Long-Term Asynchronous Decoding of Arm Motion Using Electrocorticographic Signals in Monkeys. *Front Neuroeng* **2010**, *3*, doi:10.3389/FNENG.2010.00003/.
- Slutzky, M.W.; Jordan, L.R.; Lindberg, E.W.; Lindsay, K.E.; Miller, L.E. Decoding Rat Forelimb Movement Direction from Epidural and Intracortical Field Potentials. *J Neural Eng* 2011, 8, 036013, doi:10.1088/1741-2560/8/3/036013.
- 24. Flinker, A.; Korzeniewska, A.; Shestyuk, A.Y.; Franaszczuk, P.J.; Dronkers, N.F.; Knight, R.T.; Crone, N.E. Redefining the Role of Broca's Area in Speech. *Proc Natl Acad Sci U S A* **2015**, *112*, 2871–2875, doi:10.1073/PNAS.1414491112/SUPPL_FILE/PNAS.201414491SI.PDF.
- 25. Rahman, A.; others Multimodal EEG and Keystroke Dynamics Based Biometric System Using Machine Learning Algorithms. *IEEE Access* **2021**, doi:10.1109/access.2021.3092840.

- 26. Cui, Y.; Wu, D. EEG-Based Driver Drowsiness Estimation Using Convolutional Neural Networks. In Proceedings of the Lect. Notes Comput. Sci. (including Subser. Lect. Notes Artif. Intell. Lect. Notes Bioinformatics); 2017; Vol. 10635, pp. 822–832.
- 27. Zolfaghari, S.; Rezaii, T.Y.; Meshgini, S.; Farzamnia, A.; Fan, L.C. Speed Classification of Upper Limb Movements Through EEG Signal for BCI Application. *IEEE Access* **2021**, doi:10.1109/access.2021.3102183.
- 28. Noor, N.S.E.M.; Ibrahim, H. Machine Learning Algorithms and Quantitative Electroencephalography Predictors for Outcome Prediction in Traumatic Brain Injury: A Systematic Review. *IEEE Access* **2020**, *8*, 102075–102092, doi:10.1109/ACCESS.2020.2998934.
- 29. Casarotto, S.; others The Rt-TEP Tool: Real-Time Visualization of TMS-Evoked Potentials to Maximize Cortical Activation and Minimize Artifacts 2021.
- 30. Baines, S.; Hensels, I.S.; Talmi, D. An EEG Study on the Effect of Being Overweight on Anticipatory and Consummatory Reward in Response to Pleasant Taste Stimuli 2021.
- 31. Sundaresan, A.; Penchina, B.; Cheong, S.; Grace, V.; Valero-Cabré, A.; Martel, A. Evaluating Deep Learning EEG-Based Mental Stress Classification in Adolescents with Autism for Breathing Entrainment BCI. *Brain Inform* **2021**, doi:10.1186/s40708-021-00133-5.
- 32. Gramfort, C.; Luessi, M.; Larson, E.; Engemann, D.A.; Strohmeier, D.; Brodbeck, C.; Goj, R.; Gramfort, A.; Luessi, M.; Larson, E.; et al. MEG and EEG Data Analysis with MNE-Python. *Front Neurosci* **2013**, *7*, 267, doi:10.3389/fnins.2013.00267.
- 33. Strangman, G.E.; Zhang, Q.; Zeffiro, T. Near-Infrared Neuroimaging with NinPy. *Front Neuroinform* **2009**, 3, doi:10.3389/neuro.11.012.2009.
- 34. Langlois, D.; Chartier, S.; Gosselin, D. An Introduction to Independent Component Analysis: InfoMax and FastICA Algorithms. *Tutor Quant Methods Psychol* **2010**, doi:10.20982/tqmp.06.1.p031.
- 35. Daubechies, I.; others Independent Component Analysis for Brain FMRI Does Not Select for Independence. *Proceedings of the National Academy of Sciences U.S.A.* **2009**, doi:10.1073/pnas.0903525106.
- 36. Ashrafulla, S. EEG and MEG: Functional Brain Imaging with High Temporal Resolution Syed Ashrafulla Electrical Signals in the Brain. **2001**.
- 37. Holroyd, T. Magnetoencephalography Available online: https://commons.wikimedia.org/wiki/File:Magnetoencephalography.png (accessed on 29 December 2022).
- 38. Proudfoot, M.; Woolrich, M.W.; Nobre, A.C.; Turner, M.R. Magnetoencephalography. **2014**, 336–343, doi:10.1136/practneurol-2013-000768.
- 39. Chap, C. File:Mri Scanner Schematic Labelled.Svg Wikipedia Available online https://en.wikipedia.org/wiki/File:Mri_scanner_schematic_labelled.svg (accessed on 28 December 2022).
- 40. Scheeringa, R.; Bonnefond, M.; Van Mourik, T.; Jensen, O.; Norris, D.G.; Koopmans, P.J. Relating Neural Oscillations to Laminar FMRI Connectivity. *bioRxiv* **2020**, 2020.09.18.303263.
- 41. Hiwaki, O. Novel Technique for Noninvasive Detection of Localized Dynamic Brain Signals by Using Transcranial Static Magnetic Fields. *IEEE J Transl Eng Health Med* **2021**, *9*, doi:10.1109/JTEHM.2020.3039043.
- 42. Pandey, S.; Voorsluys, W.; Rahman, M.; Buyya, R.; Dobson, J.; Chiu, K. CONCURRENCY AND COMPUTATION: PRACTICE AND EXPERIENCE A Grid Workflow Environment for Brain Imaging Analysis on Distributed Systems; 2009;
- 43. Islam, R.; Moni, A.; Islam, M.; Mahfuz, R.-A.-; Islam, S.; Hasan, K.; Hossain, S.; Ahmad, M.; Uddin, S.; Azad, A.; et al. Emotion Recognition From EEG Signal Focusing on Deep Learning and Shallow Learning Techniques. *IEEE Access* **2021**, doi:10.1109/access.2021.3091487.
- 44. Xiong, S.; Wu, G.; Fan, X.; Feng, X.; Huang, Z.; Cao, W.; Zhou, X.; Ding, S.; Yu, J.; Wang, L.; et al. MRI-Based Brain Tumor Segmentation Using FPGA-Accelerated Neural Network. *BMC Bioinformatics* **2021**, doi:10.1186/s12859-021-04347-6.
- 45. Suh, S.; Lee, H.; Lukowicz, P.; Lee, Y.O. CEGAN: Classification Enhancement Generative Adversarial Networks for Unraveling Data Imbalance Problems. *Neural Networks* **2021**, 133, 69–86, doi:10.1016/j.neunet.2020.10.004.
- 46. Sharma, M.; Tiwari, J.; Patel, V.; Acharya, U.R. Automated Identification of Sleep Disorder Types Using Triplet Half-Band Filter and Ensemble Machine Learning Techniques with EEG Signals. *Electronics (Basel)* **2021**, doi:10.3390/electronics10131531.
- 47. Zhu, Y.; Li, Y.; Li, P. EEGNet With Ensemble Learning to Improve the Cross-Session Classification of SSVEP Based BCI From Ear-EEG. *IEEE Access* **2021**, doi:10.1109/access.2021.3052656.
- 48. Islam, R.; others Emotion Recognition From EEG Signal Focusing on Deep Learning and Shallow Learning Techniques. *IEEE Access* **2021**, doi:10.1109/access.2021.3091487.
- 49. Xiong, S.; others MRI-Based Brain Tumor Segmentation Using FPGA-Accelerated Neural Network. *BMC Bioinformatics* **2021**, doi:10.1186/s12859-021-04347-6.
- 50. Zhu, Y.; Li, Y.; Lu, J.; Li, P. EEGNet With Ensemble Learning to Improve the Cross-Session Classification of SSVEP Based BCI From Ear-EEG. *IEEE Access* **2021**, doi:10.1109/access.2021.3052656.

- 51. Sharma, M.; Tiwari, J.; Patel, V.; Acharya, U.R. Automated Identification of Sleep Disorder Types Using Triplet Half-Band Filter and Ensemble Machine Learning Techniques with EEG Signals. *Electronics (Basel)* **2021**, doi:10.3390/electronics10131531.
- 52. Lee, B.-H.; Jeong, J.-H.; Lee, S.-W. SessionNet: Feature Similarity-Based Weighted Ensemble Learning for Motor Imagery Classification. *IEEE Access*, doi:10.1109/access.2020.3011140.
- 53. Scheeringa, R.; Bonnefond, M.; Mourik, T. Van; Jensen, O.; Norris, D.G.; Koopmans, P.J. Relating Neural Oscillations to Laminar FMRI Connectivity. *bioRxiv* **2020**, doi:10.1101/2020.09.18.303263.
- 54. Hiwaki, O. Novel Technique for Noninvasive Detection of Localized Dynamic Brain Signals by Using Transcranial Static Magnetic Fields. *IEEE J Transl Eng Health Med* **2020**, doi:10.1109/jtehm.2020.3039043.
- 55. Pandey, S.; Voorsluys, W.; Rahman, M.; Buyya, R.; Dobson, J.; Chiu, K. A Grid Workflow Environment for Brain Imaging Analysis on Distributed Systems. *Concurr Comput* **2009**.
- 56. Leicht, G.; others EEG-Informed FMRI Reveals a Disturbed Gamma-Band-Specific Network in Subjects at High Risk for Psychosis. *Schizophr Bull* **2016**, 42, 239–249, doi:10.1093/schbul/sbv092.
- 57. Scheinost, D.; Noble, S.; Horien, C.; Greene, A.S.; Lake, E.M.R.; Salehi, M.; Gao, S.; Shen, X.; O'Connor, D.; Barron, D.S.; et al. Ten Simple Rules for Predictive Modeling of Individual Differences in Neuroimaging. *Neuroimage* 2019.
- 58. Martisius, I. Data Acquisition and Signal Processing Methods for Brain Computer Interfaces; 2016; ISBN 9786090211977.
- 59. Irani, F.; Platek, S.M.; Bunce, S.; Ruocco, A.C.; Chute, D. Functional Near Infrared Spectroscopy (FNIRS): An Emerging Neuroimaging Technology with Important Applications for the Study of Brain Disorders. *Clinical Neuropsychologist* **2007**, *21*, 9–37, doi:10.1080/13854040600910018.
- 60. Grohol, J.M. What Is Functional Near-Infrared Spectroscopy? 2017.
- 61. Belluscio, V.; Casti, G.; Ferrari, M.; Horschig, J.M.; Vannozzi, G.; Quaresima, V.; Sappia, M.S. Modifications in Prefrontal Cortex Oxygenation in Linear and Curvilinear Dual Task Walking: A Combined FNIRS and IMUs Study. *Sensors* **2021**, doi:10.3390/s21186159.
- 62. Deepeshwar, S.; Vinchurkar, S.A.; Visweswaraiah, N.K.; Nagendra, H.R. Hemodynamic Responses on Prefrontal Cortex Related to Meditation and Attentional Task. *Front Syst Neurosci* **2015**, *8*, doi:10.3389/fnsys.2014.00252.
- 63. Jiang, D.; Liu, Z.; Sun, G. The Effect of Yoga Meditation Practice on Young Adults' Inhibitory Control: An FNIRS Study. *Front Hum Neurosci* **2021**, doi:10.3389/fnhum.2021.725233.
- 64. Annen, J.; others Mapping the Functional Brain State of a World Champion Freediver in Static Dry Apnea. *Brain Struct Funct* **2021**, doi:10.1007/s00429-021-02361-1.
- 65. Kim, D.H.; Lee, K.-D.; Thomas, B.C.; Park, H.-S. Increasing Motor Cortex Activation During Grasping Via Novel Robotic Mirror Hand Therapy: A Pilot FNIRS Study. *Res Sq* **2021**, doi:10.21203/rs.3.rs-810023/v1.
- 66. Tang, T.B.; Chong, J.S.; Kiguchi, M.; Funane, T.; Lu, C.-K. Detection of Emotional Sensitivity Using FNIRS Based Dynamic Functional Connectivity. *IEEE Transactions on Neural Systems and Rehabilitation Engineering* **2021**, doi:10.1109/tnsre.2021.3078460.
- 67. Yücel, M.A.; Selb, J.J.; Huppert, T.J.; Franceschini, M.A.; Boas, D.A. Functional Near Infrared Spectroscopy: Enabling Routine Functional Brain Imaging. *Curr Opin Biomed Eng* **2017**, 4, 78–86, doi:10.1016/j.cobme.2017.09.011.
- 68. Niu, H.; Li, Z.; Liao, X.; Wang, J.; Zhao, T.; Shu, N.; Zhao, X.; He, Y. Test-Retest Reliability of Graph Metrics in Functional Brain Networks: A Resting-State FNIRS Study. *PLoS One* **2013**, *8*, e72425, doi:10.1371/JOURNAL.PONE.0072425.
- 69. Walczak, E. Removing Triggers from Hitachi ETG-4000 FNIRS Recordings 2019.
- 70. Ardali, M.K.; Rana, A.; Purmohammad, M.; Birbaumer, N.; Chaudhary, U. Semantic and BCI-Performance in Completely Paralyzed Patients: Possibility of Language Attrition in Completely Locked In Syndrome. *Brain Lang* **2019**, *194*, 93–97, doi:10.1016/j.bandl.2019.05.004.
- 71. Homer, D. Users' Guide 2012.
- 72. Gihyoun Lee; Ji-Su Park; Jungsoo Lee; Jinuk Kim; Young-Jin Jung; Yun-Hee Kim OptoNet II: An Advanced MATLAB-Based Toolbox for Functional Cortical Connectivity Analysis With Surrogate Tests Using FNIRS. *IEEE Access* 2020, doi:10.1109/access.2020.3042808.
- 73. Einevoll, G.T.; Franke, F.; Hagen, E.; Pouzat, C.; Harris, K.D. Towards Reliable Spike-Train Recordings from Thousands of Neurons with Multielectrodes. *Curr Opin Neurobiol* **2012**, 22, 11–17, doi:10.1016/J.CONB.2011.10.001.
- 74. Buzsáki, G. Large-Scale Recording of Neuronal Ensembles. *Nature Neuroscience* 2004 7:5 **2004**, 7, 446–451, doi:10.1038/nn1233.
- 75. Padmanabhan, P.; Nedumaran, A.M.; Mishra, S.; Pandarinathan, G.; Archunan, G.; Gulyás, B. The Advents of Hybrid Imaging Modalities: A New Era in Neuroimaging Applications. *Adv Biosyst* **2017**, doi:10.1002/adbi.201700019.
- 76. Lioi, G.; others Simultaneous EEG-FMRI During a Neurofeedback Task, a Brain Imaging Dataset for Multimodal Data Integration. *Sci Data* **2020**, 7, 1–15, doi:10.1038/s41597-020-0498-3.

- 77. Chincarini, M.; Costa, E.D.; Qiu, L.; Spinelli, L.; Cannas, S.; Palestrini, C.; Canali, E.; Minero, M.; Cozzi, B.; Ferri, N.; et al. Reliability of FNIRS for Noninvasive Monitoring of Brain Function and Emotion in Sheep. *NanoScience and Technology*, doi:10.1038/s41598-020-71704-5.
- 78. Helwig, N.E.; Maintainer Package 'Eegkit' Title Toolkit for Electroencephalography Data 2015.
- Haddad, R.A.; Akansu, A.N. A Class of Fast Gaussian Binomial Filters for Speech and Image Processing. IEEE Transactions on Signal Processing 1991, doi:10.1109/78.80892.
- 80. Khan, H.; Khadka, R.; Sultan, M.S.; Yazidi, A.; Ombao, H.; Mirtaheri, P. Unleashing the Potential of FNIRS with Machine Learning: Classification of Fine Anatomical Movements to Empower Future Brain-Computer Interface. *Front Hum Neurosci* **2024**, *18*, 1354143, doi:10.3389/FNHUM.2024.1354143/BIBTEX.
- 81. Yuan, Z.; Ye, J.C. Fusion of FNIRS and FMRI Data: Identifying When and Where Hemodynamic Signals Are Changing in Human Brains. *Front Hum Neurosci* **2013**, *7*, doi:10.3389/FNHUM.2013.00676/ABSTRACT.
- 82. Hong, K.S.; Khan, M.J. Hybrid Brain-Computer Interface Techniques for Improved Classification Accuracy and Increased Number of Commands: A Review. *Front Neurorobot* **2017**, *11*, doi:10.3389/fnbot.2017.00035.
- 83. Klein, F. Optimizing Spatial Specificity and Signal Quality in FNIRS: An Overview of Potential Challenges and Possible Options for Improving the Reliability of Real-Time Applications. *Frontiers in Neuroergonomics* **2024**, *5*, 1286586, doi:10.3389/FNRGO.2024.1286586/BIBTEX.
- 84. Lanka, P.; Bortfeld, H.; Huppert, T.J. Correction of Global Physiology in Resting-State Functional near-Infrared Spectroscopy. *Neurophotonics* **2022**, *9*, doi:10.1117/1.NPH.9.3.035003.
- 85. Vansteensel, M.J.; Jarosiewicz, B. Brain-Computer Interfaces for Communication. *Handb Clin Neurol* **2020**, 168, 67–85, doi:10.1016/B978-0-444-63934-9.00007-X.
- 86. Curot, J.; Busigny, T.; Valton, L.; Denuelle, M.; Vignal, J.P.; Maillard, L.; Chauvel, P.; Pariente, J.; Trebuchon, A.; Bartolomei, F.; et al. Memory Scrutinized through Electrical Brain Stimulation: A Review of 80 Years of Experiential Phenomena. *Neurosci Biobehav Rev* 2017, 78, 161–177, doi:10.1016/j.neubiorev.2017.04.018.
- 87. Areces-Gonzalez, A.; Paz-Linares, D.; Riaz, U.; Wang, Y.; Li, M.; Razzaq, F.A.; Bosch-Bayard, J.F.; Gonzalez-Moreira, E.; Ontivero-Ortega, M.; Galan-Garcia, L.; et al. CiftiStorm Pipeline: Facilitating Reproducible EEG/MEG Source Connectomics. *Front Neurosci* **2024**, *18*, 1237245, doi:10.3389/FNINS.2024.1237245/BIBTEX.
- 88. Fahimi Hnazaee, M.; Wittevrongel, B.; Khachatryan, E.; Libert, A.; Carrette, E.; Dauwe, I.; Meurs, A.; Boon, P.; Van Roost, D.; Van Hulle, M.M. Localization of Deep Brain Activity with Scalp and Subdural EEG. *Neuroimage* **2020**, 223, 117344, doi:10.1016/J.NEUROIMAGE.2020.117344.
- 89. Puce, A.; Hämäläinen, M.S. A Review of Issues Related to Data Acquisition and Analysis in EEG/MEG Studies. *Brain Sci* **2017**, 7, doi:10.3390/BRAINSCI7060058.
- 90. Rashid, M.; Sulaiman, N.; P. P. Abdul Majeed, A.; Musa, R.M.; Ahmad, A.F.; Bari, B.S.; Khatun, S. Current Status, Challenges, and Possible Solutions of EEG-Based Brain-Computer Interface: A Comprehensive Review. *Front Neurorobot* **2020**, *14*, 25, doi:10.3389/FNBOT.2020.00025.
- 91. Mercier, M.R.; Dubarry, A.S.; Tadel, F.; Avanzini, P.; Axmacher, N.; Cellier, D.; Vecchio, M. Del; Hamilton, L.S.; Hermes, D.; Kahana, M.J.; et al. Advances in Human Intracranial Electroencephalography Research, Guidelines and Good Practices. *Neuroimage* **2022**, *260*, 119438, doi:10.1016/J.NEUROIMAGE.2022.119438.
- 92. Chang, E.F. Towards Large-Scale Human-Based Mesoscale Neurotechnologies. *Neuron* **2015**, *86*, 68, doi:10.1016/J.NEURON.2015.03.037.
- 93. Basso, M.A.; Frey, S.; Guerriero, K.A.; Jarraya, B.; Kastner, S.; Koyano, K.W.; Leopold, D.A.; Murphy, K.; Poirier, C.; Pope, W.; et al. Using Non-Invasive Neuroimaging to Enhance the Care, Well-Being and Experimental Outcomes of Laboratory Non-Human Primates (Monkeys). *Neuroimage* **2021**, 228, 117667, doi:10.1016/J.NEUROIMAGE.2020.117667.
- 94. Zhang, Y.D.; Dong, Z.; Wang, S.H.; Yu, X.; Yao, X.; Zhou, Q.; Hu, H.; Li, M.; Jiménez-Mesa, C.; Ramirez, J.; et al. Advances in Multimodal Data Fusion in Neuroimaging: Overview, Challenges, and Novel Orientation. *Inf Fusion* **2020**, *64*, 149, doi:10.1016/J.INFFUS.2020.07.006.
- 95. Lewis, C.M.; Hoffmann, A.; Helmchen, F. Linking Brain Activity across Scales with Simultaneous Optoand Electrophysiology. *Neurophotonics* **2024**, *11*, doi:10.1117/1.NPH.11.3.033403.
- 96. Leuthardt, E.C.; Moran, D.W.; Mullen, T.R. Defining Surgical Terminology and Risk for Brain Computer Interface Technologies. *Front Neurosci* **2021**, *15*, 599549, doi:10.3389/FNINS.2021.599549/BIBTEX.
- 97. Ahmadi, N.; Constandinou, T.G.; Bouganis, C.S. Inferring Entire Spiking Activity from Local Field Potentials. *Sci Rep* **2021**, *11*, 19045, doi:10.1038/S41598-021-98021-9.
- 98. Zhang, H.; Zhou, Q.Q.; Chen, H.; Hu, X.Q.; Li, W.G.; Bai, Y.; Han, J.X.; Wang, Y.; Liang, Z.H.; Chen, D.; et al. The Applied Principles of EEG Analysis Methods in Neuroscience and Clinical Neurology. *Mil Med Res* **2023**, *10*, doi:10.1186/S40779-023-00502-7.
- 99. Mandal, R.; Babaria, N.; Cao, J.; Liu, Z.; IEEE, S.M. Adaptive and Wireless Recordings of Electrophysiological Signals during Concurrent Magnetic Resonance Imaging. *bioRxiv* **2018**, 259762, doi:10.1101/259762.

- Deshpande, G.; Rangaprakash, D.; Oeding, L.; Cichocki, A.; Hu, X.P. A New Generation of Brain-Computer Interfaces Driven by Discovery of Latent EEG-FMRI Linkages Using Tensor Decomposition. *Front Neurosci* 2017, 11, 246, doi:10.3389/FNINS.2017.00246.
- 101. van Mierlo, P.; Höller, Y.; Focke, N.K.; Vulliemoz, S. Network Perspectives on Epilepsy Using EEG/MEG Source Connectivity. *Front Neurol* **2019**, *10*, 451129, doi:10.3389/FNEUR.2019.00721/BIBTEX.
- 102. Saha, S.; Mamun, K.A.; Ahmed, K.; Mostafa, R.; Naik, G.R.; Darvishi, S.; Khandoker, A.H.; Baumert, M. Progress in Brain Computer Interface: Challenges and Opportunities. *Front Syst Neurosci* **2021**, *15*, 578875, doi:10.3389/FNSYS.2021.578875.
- 103. Kübler, A.; Mushahwar, V.K.; Hochberg, L.R.; Donoghue, J.P. BCI Meeting 2005 Workshop on Clinical Issues and Applications. In Proceedings of the IEEE Transactions on Neural Systems and Rehabilitation Engineering; 2006.
- 104. Buccino, A.P.; Keles, H.O.; Omurtag, A. Hybrid EEG-FNIRS Asynchronous Brain-Computer Interface for Multiple Motor Tasks. *PLoS One* **2016**, *11*, doi:10.1371/journal.pone.0146610.
- 105. Yoon, J.W.; Roberts, S.J.; Dyson, M.; Gan, J.Q. 2009 Special Issue Adaptive Classification for Brain Computer Interface Systems Using Sequential Monte Carlo Sampling. *Neural Networks* **2009**, 22, 1286–1294, doi:10.1016/j.neunet.2009.06.005.
- 106. Belkacem, A.N.; Jamil, N.; Palmer, J.A.; Ouhbi, S.; Chen, C. Brain Computer Interfaces for Improving the Quality of Life of Older Adults and Elderly Patients. *Front Neurosci* **2020**, doi:10.3389/fnins.2020.00692.
- 107. Alhudhaif, A. An Effective Classification Framework for Brain-Computer Interface System Design Based on Combining of FNIRS and EEG Signals. *PeerJ Comput Sci* **2021**, 7, 1–24, doi:10.7717/PEERJ-CS.537.
- 108. Amiri, S.; Fazel-Rezai, R.; Asadpour, V. A Review of Hybrid Brain-Computer Interface Systems. *Advances in Human-Computer Interaction* **2013**, 2013, doi:10.1155/2013/187024.
- 109. Pandarinathan, G.; Mishra, S.; Nedumaran, A.; Padmanabhan, P.; Gulyás, B. The Potential of Cognitive Neuroimaging: A Way Forward to the Mind-Machine Interface. *J Imaging* **2018**, doi:10.3390/jimaging4050070.
- 110. Rawji, V.; Kaczmarczyk, I.; Rocchi, L.; Fong, P.Y.; Rothwell, J.C.; Sharma, N. Preconditioning Stimulus Intensity Alters Paired-Pulse TMS Evoked Potentials. *Brain Sci* **2021**, *11*, 1–13, doi:10.3390/brainsci11030326.
- 111. Revell, A.Y.; others White Matter Signals Reflect Information Transmission Between Brain Regions During Seizures. 2021.
- 112. Mareček, R.; others Automated Fusion of Multimodal Imaging Data for Identifying Epileptogenic Lesions in Patients with Inconclusive Magnetic Resonance Imaging. *Hum Brain Mapp* **2021**, 42, 2921–2930, doi:10.1002/hbm.25413.
- 113. Ortega, P.; Faisal, A.A. Deep Learning Multimodal FNIRS and EEG Signals for Bimanual Grip Force Decoding. *J Neural Eng* **2021**, *18*, 0460e6, doi:10.1088/1741-2552/ac1ab3.
- 114. Uchitel, J.; Vidal-Rosas, E.E.; J., R.C.; Zhao, H. Wearable, Integrated EEG–FNIRS Technologies: A Review. *Sensors* **2021**, doi:10.3390/s21186106.
- 115. Hollenstein, N.; others Decoding EEG Brain Activity for Multi-Modal Natural Language Processing. *Front Hum Neurosci* **2021**, doi:10.3389/fnhum.2021.659410.
- 116. Jeong, T. Time-Series Data Classification and Analysis Associated With Machine Learning Algorithms for Cognitive Perception and Phenomenon. *IEEE Access* **2020**, doi:10.1109/access.2020.3018477.

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