Denoising task-correlated head motion from motor-task fMRI data with multi-echo ICA

Neha A. Reddy[1,2], Kristina M. Zvolanek[1,2], Stefano Moia[3,4,5], César Caballero-Gaudes[3], Molly G. Bright[1,2]

Author Affiliations:
[1] Department of Physical Therapy and Human Movement Sciences, Feinberg School of Medicine, Northwestern University, Chicago, IL, United States
[2] Department of Biomedical Engineering, McCormick School of Engineering and Applied Sciences, Northwestern University, Evanston, IL, United States
[3] Basque Center on Cognition, Brain and Language, Donostia, Gipuzkoa, Spain
[4] Neuro-X Institute, École polytechnique fédérale de Lausanne, Geneva, Switzerland
[5] Department of Radiology and Medical Informatics (DRIM), Faculty of Medicine, University of Geneva, Geneva, Switzerland

Abstract
Motor-task functional magnetic resonance imaging (fMRI) is crucial in the study of several clinical conditions, including stroke and Parkinson's disease. However, motor-task fMRI is complicated by task-correlated head motion, which can be magnified in clinical populations and confounds motor activation results. One method that may mitigate this issue is multi-echo independent component analysis (ME-ICA), which has been shown to separate the effects of head motion from the desired BOLD signal, but has not been tested in motor-task datasets with high amounts of motion. In this study, we collected an fMRI dataset from a healthy population who performed a hand grasp task with and without task-correlated amplified head motion to simulate a motor-impaired population. We analyzed these data using three models: single-echo (SE), multi-echo optimally combined (ME-OC), and ME-ICA. We compared the models' performance in mitigating the effects of head motion on the subject level and group level. On the subject level, ME-ICA better dissociated the effects of head motion from the BOLD signal; both ME models led to increased activation extent and t-statistics in brain motor regions. In scans with high levels of motion, ME-ICA additionally mitigated artifacts and led to potentially more reliable beta coefficient estimates, compared to ME-OC. On the group level, all three models produced activation clusters in expected motor areas in scans with both low and high motion, indicating that group-level averaging may also sufficiently resolve motion artifacts that vary by subject. These findings demonstrate that ME-ICA is a useful tool for subject-level analysis of motor-task data with high levels of task-correlated head motion. The improvements afforded by ME-ICA are critical to improve reliability of subject-level activation maps for clinical populations in which group-level analysis may not be feasible or appropriate, for example in a chronic stroke cohort with varying stroke location and degree of tissue damage.

Keywords: BOLD fMRI, motor task, task-correlated head motion, multi-echo independent component analysis

Corresponding Author:
Neha A. Reddy
645 N. Michigan Ave., Suite 1100
Chicago, IL 60611
USA
nreddy@northwestern.edu

1. Introduction

Assessing motor function with functional magnetic resonance imaging (fMRI) is critical in several clinical and research applications, including surgical planning (Ciavarro et al., 2021; Heilbrun et al., 2001; Vysotski et al., 2018; Wilkinson et al., 2003), tracking clinical outcomes after interventions (Péran et al., 2020; Stephan et al., 2001; Ward et al., 2003a, 2003b), the study of motor neurophysiology and rehabilitation in stroke and cerebral palsy (Araneda et al., 2021; Cramer, Mark, et al., 2002; Hannanu et al., 2020; Hermsdörfer et al., 2003; J.P. et al., 2007), and the study of Parkinson's disease (Martin et al., 2019; Sabatini et al., 1998; Wu & Hallett, 2005). However, head motion poses a serious problem during motor-task fMRI, since motion confounds interpretation of fMRI results by causing false positive and negative activation in task-fMRI (Friston et al., 1996) and spurious correlations in functional connectivity data (Power et al., 2012; Satterthwaite et al., 2012; van Dijk et al., 2012). In particular, certain clinical populations may have amplified head motion during motor actions; for example, stroke participants were observed to have twice the amount of head motion during hand grasp and ankle flexion tasks compared to healthy individuals (Seto et al., 2001). Head motion is also increased in participants who may be uncomfortable in the MRI environment, such as children (Byars et al., 2002; Kotsoni et al., 2006).

Several strategies have been employed to address the issue of motion correction for fMRI data (Caballero-Gaudes & Reynolds, 2017). Most commonly, volume registration is performed to align images across the scan time points; however, it has been shown that this technique is not sufficient to eliminate motion effects (Freire & Mangin, 2001; Grootoonk et al., 2000). Many studies take a further step of adding the motion parameters and their derivatives calculated during volume registration as regressors during modeling (Friston et al., 1996), though motion-related confounds may still be present in the data and result in spurious findings (Power et al., 2012), particularly when the motion is task-correlated (Johnstone et al., 2006). Censoring and interpolation are approaches that remove large spikes in motion above a set threshold, but drawbacks are disruption of a signal's temporal correlation and challenges in deciding the correct threshold (Caballero-Gaudes & Reynolds, 2017). Additionally, censoring and interpolation do not correct for smaller motions throughout the scan, which may be more prevalent in clinical populations and can also lead to false results. Other commonly used tools to separate effects of noise from the desired Blood Oxygenation Level Dependent (BOLD) signal include approaches based on Principal Component Analysis (PCA), such as CompCor (Behzadi et al., 2007; Muschelli et al., 2014) and GLMdenoise (Kay et al., 2013) and Independent Component Analysis (ICA), such as FIX-ICA (Griffanti et al., 2014; Salimi-Khorshidi et al., 2014) and ICA-AROMA (Pruim et

al., 2015), that identify noise components of the signal and regress out their corresponding timecourses.

Additional challenges arise when head motion is correlated with the task stimulus (Field et al., 2000; Hajnal et al., 1994; Johnstone et al., 2006; Moia et al., 2021; Soltysik & Hyde, 2006; Xu et al., 2014) and the impact of task-correlated motion on fMRI activation results is not easily corrected for (Mumford et al., 2015). For example, some of the motion correction strategies described above have been shown to lose efficacy when motion is task-correlated; such strategies include volume realignment (Field et al., 2000; Morgan et al., 2007), adding motion regressors to the model (Johnstone et al., 2006), and PCA (Patriat et al., 2017). Of note, motor tasks have been associated with increased task-correlated head motion in a motor-impaired population (Seto et al., 2001), suggesting that this issue may be amplified in certain clinical cohorts.

An efficient method to identify task-correlated confounds in fMRI data involves separating the effects of changes in the transverse relaxation parameter T_2^*, which are related to the BOLD response, from the effects of changes in the net magnetization S_0, which include motion, pulsation, and inflow effects (Menon et al., 1993; Ogawa et al., 1993). This approach builds on the physical principles of the T_2^*-weighted fMRI signal and the BOLD response. In initial studies, a short echo time (TE) closely following the radiofrequency (RF) excitation was used to approximate S_0 effects and remove them from the T_2^*-related BOLD fMRI signal. Several studies used dual-echo (Glover et al. 1996; Ing and Schwarzbauer 2012; Bright and Murphy 2013; Buur, Poser, and Norris 2009) and multi-echo (Barth et al. 1999) acquisitions to perform this noise separation technique in fMRI data. Kundu et al. (2012, 2013) proposed a related method to separate BOLD and non-BOLD components using multi-echo information and ICA (ME-ICA) to calculate two parameters, kappa and rho, that quantify the T_2^*- and S_0-weighting of the components, respectively. Along with other features, these kappa and rho parameters can be used to understand the likeliness that each component is attributed to the true BOLD signal, and subsequently better classify the component (Dupre et al., 2021). ME-ICA has been used with success in various resting-state (Cohen, Chang, et al., 2021; Cohen, Yang, et al., 2021; Dipasquale et al., 2017) and task-based MRI (Cohen et al., 2018; Cohen, Jagra, Visser, et al., 2021; Cohen, Jagra, Yang, et al., 2021; Cohen & Wang, 2019; Evans et al., 2015; Gonzalez-Castillo et al., 2016; Lombardo et al., 2016; Moia et al., 2021) applications.

However, studies have not yet evaluated the use of ME-ICA in denoising task-correlated head motion in motor-task fMRI data, where it could be critically useful for the clinical applications described previously, and other correction methods may fail. Motor areas in healthy individuals

have been identified in cortical (Gordon et al., 2023) and cerebellar (Ashida et al., 2019; Spencer et al., 2007; Stoodley, 2012) regions. Multi-echo information might be additionally useful for improving our detection in non-cortical regions implicated in human movement, such as the cerebellum: several studies have observed that multi-echo and ME-ICA methods lead to signal improvements in subcortical regions (Gonzalez-Castillo et al., 2016; Guediche et al., 2021; Lombardo et al., 2016; Lynch et al., 2020).

In this study, we apply ME-ICA to a motor-task dataset including a wide range of task-correlated head motion amplitudes, to evaluate its utility in improving the interpretability of motor-task fMRI in clinical populations. To focus on the effects of task-correlated head motion, distinct from other neural or vascular confounds present in clinical populations, we simulate amplified task-correlated head motion in healthy participants as done in previous studies (Bright & Murphy, 2013; Buur et al., 2009; Ing & Schwarzbauer, 2012; Kochiyama et al., 2005). We evaluate the motion characteristics of this dataset, including average head motion and task-correlation of motion, and compare these characteristics to those of two representative stroke participants. Then, we probe the effects of ME and ME-ICA analysis on subject- and group-level activation results, including parameter estimates and t-statistics in relevant brain motor areas.

2. Methods

2.1. Data Collection

This study was approved by the Northwestern University Institutional Review Board, and all participants provided written, informed consent. Eight right-handed, healthy participants with no history of neurological or vascular disorders (4M, 26 ± 2 years) were scanned on a Siemens 3T Prisma MRI system with a 64-channel head coil. A structural T1-weighted multi-echo MPRAGE image was collected using parameters adapted from Tisdall and colleagues (2016): TR=2.17 s, TEs=1.69/3.55/5.41 ms, TI=1.16 s, FA=7°, FOV=256x256 mm^2, and voxel size=1x1x1 mm^3. The three echo MPRAGE images were combined using root-mean-square. Two functional scans were collected using a multi-band multi-echo gradient-echo echo planar imaging sequence provided by the Center for Magnetic Resonance Research (CMRR, Minnesota): TR=2 s, TEs=10.8/28.03/45.26/62.49/79.72 ms, FA=70°, MB factor=4, GRAPPA=2 voxel size=2.5x2.5x2 mm^3 (Moeller et al., 2010; Setsompop et al., 2012). Tape placed across the forehead provided tactile feedback of head motion, as found to reduce translational and rotational motion (Krause et al., 2019). During the scans, CO_2 was measured via a nasal cannula and gas analyzer at a sampling rate of 20 Hz (PowerLab, ADInstruments).

2.1.1. Motor task

Hand grasp tasks were performed with a device that interfaced with the scanner bed and contained two load cells (Interface Inc., Scottsdale, AZ, USA). This device provided support to both arms, positioning them at an elbow flexion of approximately 30 degrees, and provided a comfortable grasping interface with the load cells for each hand. Prior to the scan, grasp forces at maximum voluntary contraction (MVC) were calculated for the right and left hand as the average of three unilateral maximum grasp force trials. During the task trials, participants performed an isometric unimanual grasping task at 40% MVC. Each task trial was a 10-s "squeeze" and 15-s "relax", alternating 4 trials per hand, for a total of 16 trials. Four participants began with the right-hand grasp, and four participants began with the left hand. Task instructions and real-time force feedback from the load cells were viewed on a screen to facilitate force targeting. Text on the screen instructed the start and end of the hand grasp periods. During the hand grasp periods, a target box of 35-45% MVC indicated the target force level. A moving bar indicated the participants' real-time force level and turned green when the force was within the target range.

During the first functional scan *(Limited)*, participants were instructed to keep their head as still as possible. During the second functional scan *(Amplified)*, participants were instructed to nod their head slightly during each grasp to add task-correlated motion. Out of the six possible directions of head rotation and translation, nodding was chosen as it was a primary direction of motion observed in preliminary data in healthy and stroke participants, as well as feasible to be performed within the limitations of the MRI environment. Right- and left-hand grasp force data from the load cells were measured continuously during the scans at a sampling rate of 20 Hz and synchronized with CO_2 recordings and the scanner trigger (PowerLab, ADInstruments).

2.2. Data Analysis

2.2.1. Creation of motor task and end-tidal CO_2 regressors

Using MATLAB, the force traces from the right and left hands during the scans were normalized to the participant's maximum grip force, then convolved with a canonical hemodynamic response function. The resulting traces were rescaled to the range of the normalized force traces, then demeaned. The demeaned force trace was then downsampled to the resolution of the functional MRI images (TR=2s) in order to generate the corresponding RGrip and LGrip regressors.

End-tidal peaks were detected in the CO_2 data using an automatic peak-finder in MATLAB, then manually inspected. The end-tidal peaks were interpolated to form an end-tidal CO_2 trace that was convolved with a canonical hemodynamic response function, rescaled to the range of

the unconvolved timeseries, then demeaned. The demeaned trace was then downsampled to the resolution of the functional MRI images (TR=2s).

2.2.2. Structural MRI pre-processing

T1-weighted images for each subject were processed with FSL's (Jenkinson et al., 2012) fsl_anat, which performs bias field correction and brain extraction.

2.2.3. Functional MRI pre-processing

FSL (Jenkinson et al., 2012) and AFNI (Cox J.S., 1996) tools were used for fMRI preprocessing of single-echo and multi-echo datasets as follows.

Single-echo (SE)

The second echo data (TE=28.03 ms) were used as a proxy for conventional single-echo (SE) fMRI analysis. The first 10 volumes of each scan were removed to allow the signal to achieve steady-state magnetization. Head-motion realignment was computed with reference to the Single Band reference image taken at the start of the scan (3dVolreg, AFNI). After brain extraction (bet, FSL), the SE timeseries were converted to signal percentage change for further analysis.

Multi-echo (ME)

The first 10 volumes of each echo were removed to allow the signal to achieve steady-state magnetization. Head-motion realignment was estimated for the first echo with reference to the Single Band reference image taken at the start of the scan (3dVolreg, AFNI) and then applied to all echoes (3dAllineate, AFNI). All images were brain extracted (bet, FSL). Tedana (Dupre et al., 2021) was used to calculate a T_2*-weighted combination of the five echo datasets, producing the optimally combined (ME-OC) fMRI dataset. The ME-OC timeseries was converted to signal percentage change for further analysis.

As tedana leads to a tighter brain mask through removal of additional voxels with insufficient signal, for the subject-level analyses described subsequently, this tighter brain mask is applied to both SE and ME analysis for more appropriate comparison between models.

Multi-echo Independent Component Analysis (ME-ICA)

Multi-echo independent component analysis (ME-ICA) was performed on the ME-OC fMRI data (prior to conversion to signal percentage change) using tedana. The resulting components were manually classified (accepted as signal of interest or rejected as noise), using criteria adapted

from (Griffanti et al., 2017), and aided by Rica (Uruñuela, 2021). Components were *rejected* if any of the following applied:

1. Spatial maps showing
    a. Larger amplitudes mostly in white matter or CSF voxels;
    b. Small and scattered clusters
    c. Striped alternating positive and negative regions (indicative of multi-band or edge artifacts)
2. Time series showing
    a. Sudden, abrupt changes
3. Power spectrum
    a. Main frequency above 0.1 Hz
4. Multi-echo information
    a. High rho or low kappa

When classification based on these criteria was unclear, the component was accepted.

2.2.4. General linear models for denoising

The SE and ME-OC images from each functional dataset were processed using AFNI's 3dREMLfit, applied to the signal from each voxel *(Y_{SE} for single-echo data and Y_{OC} for multi-echo optimally combined data)*.

1. The *SE model* included six motion parameters from volume realignment *(Mot)*, up to fourth-order Legendre polynomials *(Poly)*, the end-tidal CO_2 regressor (CO_2), and left- and right-hand grip regressors *(RGrip* and *LGrip)*.

$$Y_{SE} \sim \beta_{Mot}X_{Mot} + \beta_{Poly}X_{Poly} + \beta_{CO_2}X_{CO_2} + \beta_{RGrip}X_{RGrip} + \beta_{LGrip}X_{LGrip} + \varepsilon$$

2. The *ME-OC model* included 6 motion parameters from volume realignment *(Mot)*, up to fourth-order Legendre polynomials *(Poly)*, the end-tidal CO_2 regressor (CO_2), and left- and right-hand grip regressors *(RGrip* and *LGrip)*.

$$Y_{OC} \sim \beta_{Mot}X_{Mot} + \beta_{Poly}X_{Poly} + \beta_{CO_2}X_{CO_2} + \beta_{RGrip}X_{RGrip} + \beta_{LGrip}X_{LGrip} + \varepsilon$$

3. The *ME-ICA model* included 6 motion parameters from volume realignment, up to fourth-order Legendre polynomials, the end-tidal CO_2 regressor (CO_2), left- and right-hand grip regressors *(RGrip* and *LGrip)*, and the rejected ME-ICA components *(IC_{rej})*.

$$Y_{OC} \sim \beta_{Mot}X_{Mot} + \beta_{Poly}X_{Poly} + \beta_{CO_2}X_{CO_2} + \beta_{RGrip}X_{RGrip} + \beta_{LGrip}X_{LGrip} + \beta_{IC_{rej}}X_{IC_{rej}} + \varepsilon$$

For each of the three models, a denoised dataset was also calculated by subtracting the fitted timeseries associated with the nuisance regressors (i.e., the product of the corresponding beta-

coefficient maps and the regressor time series) from the Y_{SE} and Y_{OC} data. Nuisance regressors are defined here as all the model regressors other than *RGrip* and *LGrip*.

2.2.5. Quantification of head motion

Framewise displacement (FD) was calculated from the volume realignment parameters as the sum of the absolute difference of the realignment parameters between samples (Power et al., 2012). The motion parameters are defined as X (anterior-posterior axis), Y (left-right axis), Z (inferior-superior axis), Roll (rotation around X), Pitch (rotation around Y), and Yaw (rotation around Z). As described in *2.2.3*, for SE data, these volume realignment parameters were calculated with respect to the second echo Single Band reference image; for ME data, these volume realignment parameters were calculated with respect to the first echo Single Band reference image. To see how many data points may have been removed if censoring was applied, the number of data points with a FD > 0.5 was calculated. While several censoring thresholds have been used previously, FD > 0.5 is within the range of previous fMRI studies (Power et al., 2014, 2017; Siegel et al., 2014), and is used here as an example. The left- and right-hand grasp force traces were summed to create an overall task timeseries. Pearson correlation coefficients comparing this task timeseries with each of the six motion parameters from volume realignment were calculated.

2.2.6. Assessment of head motion dissociation from Blood Oxygenation Level Dependent (BOLD) signal

As done by Moia et al. (2021), the denoised datasets described in *2.2.4* were used to calculate DVARS, the spatial root mean square of the first derivative of the signal (Smyser et al., 2010) using 3dTto1D (AFNI). For SE, ME-OC, and ME-ICA models, DVARS was calculated within the tighter brain extraction mask created after tedana. A dataset with high association between head motion and BOLD signals would display greater DVARS with increasing FD, while a dataset with low association between head motion and BOLD signals would display consistent DVARS with increasing FD. To quantify this association, the slopes of the DVARS vs. FD relationship were calculated for each scan (DVARS ~ FD). For the Limited and Amplified motion conditions, the slopes of the DVARS vs. FD relationship were modeled at the group level as |Slope| ~ Model + (1|Subject).

2.2.7. ROI analysis

To facilitate quantitative comparisons of motor activation across subjects and scans, several ROIs known to be involved with hand grasping force generation were identified using manual and automated methods.

The right- and left-hand knob areas of the motor cortex, corresponding to hand motor activation (Yousry et al., 1997), were manually drawn for each participant in anatomical space, then non-gray matter voxels removed by finding overlap with a gray matter mask created using fsl_anat (thresholded at 0.5). This mask was dilated by two voxels, then linearly transformed to the functional space for each subject (epi_reg, FLIRT, FSL).

Right and left motor cerebellum ROIs were created from the SUIT probabilistic cerebellum atlas in MNI space (J. Diedrichsen et al., 2011; Jörn Diedrichsen et al., 2009). Hand and finger motor tasks have been associated with activity in ipsilateral cerebellum lobules V/VI and VIIIa/b (Ashida et al., 2019; Spencer et al., 2007; Stoodley, 2012). Consequently, right lobules V/VI and VIIIa/b from the SUIT atlas were thresholded at 50%, binarized, and combined to form a right motor cerebellum ROI; the same was done with the left motor lobules to create a left motor cerebellum ROI. These masks in MNI space were then non-linearly transformed to functional space for each subject (FNIRT, FSL).

The number of positive activated voxels (defined as voxels showing a statistical significance of $p_{FDR} < 0.05$), median positive beta coefficient, and median positive t-statistic corresponding to the *RGrip* regressor were calculated in the left-hand knob and right motor cerebellum ROIs for each scan in functional space. The number of positive activated voxels ($p_{FDR} < 0.05$), median positive beta coefficient, and median positive t-statistic corresponding to the *LGrip* regressor were calculated in the right-hand knob and left motor cerebellum ROIs.

Significant differences in the parameters calculated by each model were modeled as Parameter ~ Model + (1|Subject). Significant differences in the median beta coefficient calculated for the Limited and Amplified conditions were modeled as BetaCoefficient ~ Condition + (1|Subject).

2.2.8. Correlation of motor task regressors with end-tidal CO_2 regressor

As subjects might modify their breathing during the hand grasps, which in turn alters the level of arterial CO_2 and thus influences blood flow and BOLD fMRI signals throughout the brain due to vasodilation (Birn et al., 2006; Farthing et al., 2007), we accounted for this potential confounding effect by including an end-tidal CO_2 regressor in our subject-level models (*2.2.4*). End-tidal CO_2 recordings are a reasonable non-invasive estimate of arterial CO_2 levels (McSwain et al., 2010).

We assessed the task-correlation of respiratory effects in our datasets by calculating Pearson correlation coefficients comparing the left- and right-hand grip regressors and the sum of these grip regressors with the end-tidal CO_2 regressor. To better understand the effect of adding this CO_2 regressor to our models, we analyzed the datasets with additional subject-level general linear models that did not include the CO_2 regressor.

2.2.9. Motor activation group analysis

Group-level activation maps were calculated for the Limited and Amplified motion scans to compare motor activation results across subjects with low and high amounts of motion, respectively.

Beta coefficient and t-statistic maps for *RGrip* and *LGrip* for each subject were converted to MNI space by applying a single concatenated spatial transformation of the functional images to the subject's T1-w structural image (epi_reg, FSL) and from this image to the MNI template (FLIRT and FNIRT, FSL). Group-level analysis was performed using AFNI's 3dMEMA, using contrasts for *RGrip* > 0 and *LGrip* > 0. Group-level maps were thresholded at p < 0.005 and clustered at α < 0.05, using a bi-sided t-test (3dFWHMx, 3dClustSim, 3dClusterize, AFNI).

Hand-knob and cerebellum motor area ROIs used for subject-level quantitative analysis (*2.2.8*) were similarly created for group-level ROI analysis in MNI space. The right- and left-hand knob areas of the motor cortex were manually drawn in MNI space, non-gray matter voxels removed by finding overlap with a gray matter mask, then dilated by two voxels. Motor cerebellum areas were created from the SUIT atlas as described in *2.2.8*.

2.3. Stroke datasets

Our head motion simulation for the Amplified condition aimed to approximate the head motion characteristics of motor-impaired participants. Thus, we analyzed head motion data from two participants with chronic hemiparetic stroke and unilateral hand impairment to compare with our healthy participant simulation results. Stroke Subject 1 (M, 65yrs) experienced impairment of the left hand, and Stroke Subject 2 (M, 65yrs) experienced impairment of the right hand. Two fMRI datasets were collected from each participant and used only to measure head-motion realignment parameters. No further analysis was performed on these datasets, as it was outside the scope of the current study.

The first participant's functional scans were single-echo acquisitions, with parameters: (Scan 1) TR=1.3 s, TE=34.4 ms, FA=62°, MB factor=4, voxel size=2x2x2 mm³; (Scan 2) TR=2 s, TE=20 ms, FA=90°, GRAPPA=2, voxel size=2x2x2 mm³. The second participant's functional

scans fully matched the healthy participant datasets (i.e. collecting multi-echo fMRI data). The stroke participants performed the same motor task described above and were instructed to keep their head as still as possible (note, the first participant performed 7 left hand grips and 9 right hand grips). Head-motion realignment parameters were computed with reference to the Single Band reference images taken at the start of each scan (3dVolreg, AFNI).

3. Results

All structural and functional scans were successfully collected as described above. The Amplified condition scan from Subject 1 was excluded from ROI and group activation analysis because the average head motion (FD) of the scan was identified as an outlier (**Supplemental Figure 1**).

3.1. Motion characteristics of the datasets

The forces produced by all healthy and stroke subjects are shown in **Figure 1A** and **Figure 1B.** All healthy participants had high task accuracy, demonstrated by force traces that stayed within the green shaded area representing the target force range (35-45% MVC). The stroke participants were also able to successfully perform the task with both paretic and non-paretic hands. Stroke Subject 2 demonstrated some concurrent grasping with the hand that was not instructed by the task, and also missed one grasping period during the first scan. A non-zero baseline force, seen in some of the subjects' force traces, may have been due to the participant lightly grasping the load cell during the rest periods.

Figure 1C displays the motion parameters from volume realignment and calculated FD for one representative healthy subject (Subject 2) and one stroke subject (Stroke Subject 1, Scan 1) during the hand grasp task. Head motion was observed to be highly related to the motor task timings, which are indicated by the gray shaded regions. This task-correlation is visible even in the Healthy Limited condition, though with lower amplitudes of motion. During the Amplified condition, the healthy subject demonstrated increased task-correlated head motion in the Z and pitch directions. In contrast, the stroke subject primarily displayed task-correlated motion in the Y and Z directions.

These observations are quantified in **Figure 2**, which summarizes the average FD, percent of volumes that would be censored (FD > 0.5), and the correlation between the estimated realignment parameters and hand grasping task for each scan. Every healthy participant had an increase in FD and the percent of potentially censored volumes during their Amplified motion scan compared to the Limited motion scan. Generally, the healthy participants showed an increase in task-correlated head motion in the Amplified motion scan, particularly in the X, Z, and Pitch

directions that are associated with the added, self-induced head nodding. Note that participants with a high increase in FD between scans did not necessarily have a high increase in task-correlated head motion, and vice versa, demonstrating the important distinction between these two characteristics of head motion during fMRI (ex. Subject 1 Amplified, Subject 4 Amplified). The two scans for Stroke Subject 1 have similar average head motion and percent potentially censored volumes to that of the healthy Amplified scans, while Stroke Subject 2 had much higher average head motion. Both Stroke Subjects 1 and 2 displayed task-correlated head motion, though the correlations were somewhat more similar to the healthy Limited scans. FD calculations across the scans showed high similarity between the SE and ME analyses (**Supplemental Figure 2**).

Figure 1. (A) Force traces from healthy subjects 1, 3, 5, and 7 and Stroke Subject 1, normalized to each subject's hand grasp at maximum voluntary contraction (MVC). Green shaded areas indicate target force range (35-45% MVC) displayed in visual task instructions. (B) Force traces from healthy subjects 2, 4, 6, and 8 and Stroke Subject 2, normalized to each subject's MVCs. Green shaded areas indicate target force range (35-45% MVC) displayed in visual task instructions. (C) Motion parameters from volume realignment and calculated Framewise Displacement (FD) from one representative healthy subject (Subject 2) in the Limited motion and

Amplified motion conditions and one Stroke subject (Stroke Subject 1, Scan 1) with mild motor impairment. Shaded areas represent periods when the subjects were instructed to perform the grasping task.

Figure 2. Average Framewise Displacement (FD), percent of data points censored at FD > 0.5, and correlation of the task regressor (LGrip + RGrip) with the timeseries of the realignment parameters for each functional scan. For healthy participants, the Limited and Amplified motion tasks are shown. For stroke participants, the two scans acquired in each participant are shown, during which the instructions were to keep head motion minimal.

3.2. Assessment of head motion dissociation from Blood Oxygenation Level Dependent (BOLD) signal

The ability of each model to dissociate the effects of head motion from the BOLD fMRI signal was assessed by examining the relationship between DVARS, a measure of fMRI signal change, and FD, a measure of head motion, at each fMRI timepoint. An ideal denoising strategy would minimize the magnitude of this relationship, meaning large changes in BOLD signal would not occur at the same times as large changes in head position. **Figure 3A** shows DVARS vs. FD plots

for each scan, calculated for all three models, in the Limited and Amplified conditions. The ME-ICA models consistently had lower DVARS than the SE and ME-OC models, even during volumes with high FD. **Figure 3B** summarizes the DVARS vs. FD relationship across scans. Within both the Limited and Amplified motion conditions, the ME-ICA model had significantly lower slopes than the SE and ME-OC models ($p < 0.05$, Bonferroni-corrected), indicating a better ability to dissociate the effects of head motion from the BOLD signal. ME-OC also led to lower slopes than SE, significant only in the Amplified condition. This suggests a trend of ME-OC in decreasing the relationship between head motion and BOLD signal changes, compared to SE; further improvements are seen with the addition of ME-ICA.

Figure 3. (A) Relationship between DVARS and Framewise Displacement (FD) for each scan. A more horizontal fit line indicates lower correlation of the change in BOLD signal with the amount of head motion at each time point of the scan. Note difference in X-axis scale between Limited and Amplified conditions. (B) Summary of slopes from DVARS vs. FD graphs in (A). Bars indicate significant difference between models ($p < 0.05$, Bonferroni-corrected).

3.3. Subject-level activation

We analyzed subject-level activation maps to understand how task-correlated head motion affects our ability to visualize motor activation and evaluate how ME preprocessing (ME-OC and ME-ICA) can reduce these confounding effects. **Figure 4** shows subject-level activation maps for the right grip task, with location of slices chosen to depict activation in the hand motor area. Four representative scans with low, moderate, high, and very high levels of average head motion (average FD) are depicted. In the low motion scan, typical of healthy participants in the Limited motion dataset, there is minimal difference between the three models; activation in the motor cortex is similar across models and noise is slightly reduced in the ME models, seen as apparent smoothing across the brain and reduced artifacts at the edges of the brain and cerebellum. In the moderate motion scan, the SE model shows noise at the edges of the brain, which is mitigated using the ME models. In the high motion scans, the SE model shows both noise at the edges of the brain and a banding artifact that may be related to the interaction between the multi-band slice acquisition and head motion. The ME-OC model does not remove these artifacts, while the ME-ICA model is able to better mitigate them. This same effect is observed to a greater degree in the very high motion scan, which was an outlier for the healthy Amplified datasets in terms of the average FD, though comparable to the motion observed in Stroke Subject 2. Activation in the motor cortex, indicated by arrows in **Figure 4**, is present in all scans and models; however, in the high and very high motion scans, motor cortex activation is most easily visually differentiated from the surrounding region when using ME-ICA. The moderate and high head motion scans are more similar to the motion of the stroke participant scans, and ME-ICA may have particular utility in reducing the larger motion-related artifacts in these scans. Similar results were observed for the rest of the scans.

Figure 4. Subject-level maps in functional space of percent BOLD signal change related to the right grip task. Four representative scans with low, moderate, high, and very high levels of head motion are shown (Subject 3 Limited, Subject 4 Limited, Subject 8 Amplified, and Subject 1 Amplified respectively). Average FD and correlation of task with motion in the X direction for each scan are listed. Arrows indicate expected regions of activation in the motor cortex. As head motion increases, artifacts increase at the edges of the brain and banding patterns appear. These artifacts are mitigated by ME-OC and ME-ICA.

3.4. Subject-level ROI analysis

To quantify the differences in motor task activation across scans and analyses, activation characteristics were specifically probed in brain regions related to hand motor tasks: the hand-knob regions of the motor cortex and cerebellum lobules V/VI and VIIIa/b. The manually drawn hand-knob masks for each subject are shown in **Supplemental Figure 3**. For each motor task (left- or right-hand targeted grasping), activation results were analyzed in the contralateral hand knob and ipsilateral cerebellum. **Figure 5** shows the activation extent, median positive beta coefficient, and median positive t-statistic in each region, summarized across the Limited and Amplified motion conditions and the three models. Activation extent was defined as the number of positive activated voxels ($p_{FDR} < 0.05$) in each ROI.

In the hand-knob regions, the ME models had significantly greater activation extent than SE ($p < 0.05$). In the cerebellum, both ME models had greater activation extent than SE, with only the ME-OC achieving significance. This trends are seen in both the Limited and Amplified motion conditions.

In terms of beta coefficients, with Limited motion, the ME models had greater median positive beta coefficients than SE in the hand-knob regions, but lower beta coefficients than SE in the cerebellum. However, with Amplified motion, ME-OC leads to a higher beta coefficient than SE or ME-ICA; this trend is significant in the left hand knob, though also present in the right hand knob. Given that the hand grasp task was the same across the Limited and Amplified motion scans, the activation represented by the beta coefficient within the hand-knob region is expected to be similar. In the left hand knob, the median beta coefficient is significantly different between the Amplified and Limited conditions for only the ME-OC model (SE: $p = 0.07$; ME-OC: $p = 0.03$; ME-ICA: $p = 0.38$). In the right hand knob, the median beta coefficient is significantly different between the Amplified and Limited conditions for SE and ME-OC (SE: $p = 0.03$; ME-OC: $p = 0.04$; ME-ICA: $p = 0.16$). For the Amplified motion scans, the ME-OC model's higher beta coefficients suggest a possible inflation of activation effect size using ME-OC, mitigated by the addition of ME-ICA. The Amplified to Limited condition comparisons are more mixed for left cerebellum (SE: $p = 0.002$; ME-OC: $p < 0.0001$; ME-ICA: $p = 0.02$) and right cerebellum (SE: $p = 0.13$; ME-OC: $p = 0.16$; ME-ICA: $p = 0.01$), with no clear pattern across models.

The median positive t-statistic was larger in the ME conditions compared to SE. During Amplified motion scans, ME-ICA had comparable t-statistics to ME-OC in the hand-knob regions, though the beta coefficient was smaller. Therefore, the comparable t-statistic may reflect a large decrease in the variance of the residuals in the ME-ICA model compared to ME-OC, suggesting a better ability of the ME-ICA model to mitigate uncertainty in the model.

Figure 5. Summary activation extent, beta coefficient, and t-statistic values for each scan associated with right-hand grip in the left-hand knob (LHK) and right cerebellum hand motor areas (RCB) and values associated with left-hand grip in the right-hand knob (RHK) and left cerebellum

hand motor areas (LCB). Bars indicate significant difference between models ($p < 0.05$, Bonferroni-corrected).

3.5. Correlation of motor task regressors with end-tidal CO_2 regressor

To understand and quantify the potential confounding effect of task-correlated CO_2 changes in our dataset, we visualized the interactions of the end-tidal CO_2 regressor with the task regressors and calculated the correlations of an end-tidal CO_2 regressor with the right, left, and right + left task regressors; three representative datasets with high correlation values are shown in **Figure 6**. In these datasets, increases in end-tidal CO_2 are seen in the periods between tasks (**Figure 6A**). This observation is likely due to more shallow breathing during the hand grasp periods. The datasets were then modelled with and without the CO_2 regressor to assess the impact of accounting for the effects of CO_2 variations during the scan. Particularly in the ME-OC condition, the addition of a CO_2 regressor reduced some regions with negative t-statistics (**Figure 6B**), which are likely attributed to the predominantly negative correlation of CO_2 regressor with the task (**Figure 6C**). Across all scans, the correlation values varied widely, ranging from $|r| = 0.063$ to $|r| = 0.64$. Most correlation values between the CO_2 regressor and the task regressors (R, L, and R+L) were negative due to the increase in CO_2 that occurred after each hand grasp ended.

Figure 6. Three representative scans with high negative correlations of the CO_2 regressor with the summed task regressors are shown. (A) Timeseries of the end-tidal CO_2 regressor and

summed right and left task regressors, normalized to unit variance. End-tidal CO_2 increases in the period between tasks. (B) T-statistic maps from the right grip regressor are shown when running each model with and without a CO_2 regressor. Particularly in the ME-OC model, the addition of a CO_2 regressor reduced areas of negative t-statistics (indicated by arrows). (C) Correlations of the CO_2 regressor with the right (R), left (L), and summed (R+L) task regressors for each scan show primarily negative correlations across the dataset.

3.6. Group-level activation

Group-level activation maps were calculated for the Limited and Amplified motion conditions to determine the difference in the models' abilities to identify primary regions of hand motor activity in datasets with low and high levels of head motion. Group activation maps for the three models are shown for the contrasts of Right Grip > 0 and Left Grip > 0 (**Figure 7**). In both Limited and Amplified motion conditions and with all three models, there are clusters in the hand-knob regions contralateral to the hand grasp. There are two expected areas of activation in the ipsilateral cerebellum, the V/VI and VIIIa/b lobules. The R > 0 Limited and L > 0 Amplified contrasts show significant clusters in both lobule regions, while the R > 0 Amplified and L > 0 Limited contrasts only show one cluster in one lobule region. The ME models result in slightly more robust activation in the cerebellum compared to SE, as demonstrated by larger clusters in the cerebellum, indicated by pink arrows in **Figure 7**.

Other, primarily negative, clusters are seen in the occipital and frontal lobes and at the edges of the brain, which may be artifacts, highlighted with white arrows in **Figure 7**. The clusters in the inferior frontal lobes in the Amplified condition are reduced in the ME-OC and ME-ICA models compared to SE. The clusters in the occipital lobe are somewhat inconsistent across motion conditions and contrasts, with the ME-ICA model resulting in the lowest prevalence of occipital lobe clusters in all but the Right Grip > 0 contrast in the Limited condition.

Quantitative comparisons of the clusters across Limited and Amplified conditions and the three models for hand knob and cerebellum motor ROIs are shown in **Figure 8**. Similar to the subject-level results, group-level activation extent increases in the ME models compared to the SE model. The hand-knob clusters seen in the ME models have greater median beta coefficients than the SE model. In cerebellar regions, the median positive beta coefficient and median positive t-statistic have an inconsistent effect between models and ROIs.

Figure 7. Group-level beta coefficient maps for the contrasts Right Grip > 0 and Left Grip > 0, shown for each analysis model and head motion condition. Opacity of beta coefficients is modulated by the t-statistic, as recommended by Taylor and colleagues (2023). Significant clusters are outlined in black, found by thresholding at p < 0.005 and clustering at α < 0.05. Pink arrows highlight cerebellar regions with robust activation in the ME models. White arrows highlight regions of negative clusters, which may be artifacts.

Figure 8. Summary activation extent, beta coefficient, and t-statistic values for clustered group activation maps associated with right hand grip in the left hand knob (LHK) and right cerebellum hand motor areas (RCB) and values associated with left hand grip in the right hand knob (RHK) and left cerebellum hand motor areas (LCB).

4. Discussion

In this study, we tested ME-OC and ME-ICA against SE methods on a motor-task dataset with a range of task-correlated head motion amplitudes. In a population of healthy adults, we collected a set of scans with typical, limited head motion and a set of scans with amplified task-correlated motion to simulate the challenges potentially experienced in a motor-impaired population. We examined the motion and breathing characteristics of these datasets to better understand the inherent artifacts and modeling challenges, and we showed our simulation was a reasonable approximation of motion challenges observed in stroke participants performing a similar task (**Figure 2**). Then, we applied SE, ME-OC, and ME-ICA models to identify activity in the brain associated with the hand grasp task. ME models demonstrated a better ability to dissociate head motion from the BOLD signal compared to SE. Additionally, ME-ICA models were better able to mitigate motion-related artifacts and provide potentially more reliable activation estimates on a subject-level when task-correlated head motion was magnified. On the group level, all three models resulted in similar activation clusters in the motor areas of interest, with ME-OC and ME-ICA reducing some presumed artifactual negative clusters in other brain regions in certain instances.

4.1. Magnitude and task-correlation of head motion are distinct phenomena

Through our simulation, we aimed to increase task-correlated head motion during a motor task; as expected, this simulation increased both average FD and the correlations of various directions of motion with the task (**Figure 1**). However, an increase in average FD was not always associated with a concurrent increase in task-correlation, and vice versa. For example, Subject 1's Amplified scan had a much higher average FD than the other Amplified motion scans (FD = 1.85), but the degree of task correlation of the motion was similar. The same trend can be seen in the stroke datasets. Stroke Subject 1 had an average FD similar to that of the Healthy Amplified scans, as well as similar task-correlation in the Z direction. Stroke Subject 2 had much higher average FD, but lower task-correlations of motion. These observations suggest that two distinct, but related, complications may arise when dealing with increased head motion in motor-task fMRI: greater average head motion across the scan and greater task-correlation of head motion.

The analysis of potentially censored volumes showed that, using a threshold of FD > 0.5 that is considered 'lenient' by some studies (Power et al., 2014, 2017), 20-50% of volumes may be removed by censoring in our simulated "patient" data with Amplified movement (**Figure 2**). This level of data removal may be unacceptable in task-fMRI studies, and ME-ICA may be an important alternative to reduce strong motion-related signal changes and reduce loss of data.

4.2. ME-ICA mitigates motion-related artifacts in subject-level activation maps

To assess the ability of each model to reduce motion-related artifacts in the brain, we analyzed the relationship between DVARS and FD in each subject's data. The relationship between DVARS and FD has been used in several previous studies to understand how well a model is able to dissociate the effects of head motion from changes in the BOLD signal (Kundu et al., 2013; Moia et al., 2021; Muschelli et al., 2014; Power et al., 2019). We found that the ME-ICA model led to a significantly lower correlation between DVARS and FD compared to the SE and ME-OC models, particularly in the Amplified motion condition (**Figure 4**). Moia and colleagues (2021) observed the same trends across models when analyzing fMRI data during a breath-hold task, where task-correlated motion is also typically enhanced (Bright et al., 2009).

We then observed how these differences in dissociation of head motion from BOLD signal manifested in subject-level activation maps. Subject-level activation maps are particularly important in populations that may exhibit subject-level vascular and neural impairment differences. For example, reliable subject-level activation maps can aid in analysis in a stroke population that stratifies participants by motor impairment level. Vascular differences in this population due to the stroke lesion may also make group-level analysis less feasible. We analyzed subject-level activation maps in the Limited and Amplified datasets to understand how the increased amplitude and task-correlation of head motion, as seen in motor-impaired populations, may influence our ability to reliably visualize motor activation.

In a scan with low average motion, we observed the primary effect of ME-OC and ME-ICA to be an apparent smoothing across the brain compared to SE (**Figure 4**). As average head motion increased in a scan, more artifacts became apparent in the SE and ME-OC conditions, primarily as noise at the edges of the brain and a banding artifact across the brain, possibly due to the interaction between the multi-band acquisition and head motion. ME-ICA largely mitigated these artifacts; Olafsson and colleagues (2015) also observed improvements when implementing ME-ICA with a simultaneous multi-slice protocol. The moderate, high, and very high motion examples shown in **Figure 4** are more representative of the level of motion seen in a motor-impaired population (**Figure 2**) and display the artifacts that might be encountered during a motor-task scan in these populations. At higher levels of motion, SE and ME-OC models may not sufficiently remove artifacts at the subject level, while ME-ICA models allow for cleaner subject-level maps.

Of note, a previous SE study also analyzed the effect of ICA on removing the effects of task-correlated motion during a motor task (Kochiyama et al., 2005). The authors found that their ICA approach led to fewer false negative errors than a traditional voxel-wise regression-based

approach. However, they note a limitation of their method that reduces its efficacy when the head motion follows a similar block design to the task (i.e., movement at the beginning of the task, remaining at the same position, and a return to the original position at the end of the task). In our study, this was the type of movement we simulated in the Amplified condition, and we observed similar characteristics of movement in the Limited condition and in the Stroke Subject scans. Even with this type of problematic motion in our scans, we found the subject-level improvements described above; these improvements were potentially aided by the added benefits of ME acquisition and ME rho and kappa parameter information used to classify ME-ICA components.

4.3. ME-ICA results in more reliable beta coefficient estimates in the presence of large amounts of task-correlated head motion

We quantified motor activation in the subject-level maps through ROI analysis in regions associated with hand grasping: the hand-knob area of the motor cortex and cerebellum lobules V/VI and VIIIa/b. The association of hand grasp with activity in the hand knob of healthy participants has been well established (Cramer, Weisskoff, et al., 2002; Ehrsson et al., 2000; Keisker et al., 2009, 2010; Kuhtz-Buschbeck et al., 2008; Ludman et al., 1996; Thickbroom et al., 1998, 1999). The cerebellum lobules V/VI and VIIIa/b are associated with upper extremity activity and are also activated during hand and finger motor tasks (Ashida et al., 2019; Spencer et al., 2007; Stoodley, 2012). Therefore, we chose these regions to analyze the activation extent, median positive beta coefficient, and median positive t-statistic for each scan, comparing across models (**Figure 5**).

The hand-knob ROIs were manually drawn for each participant to isolate the hand motor region. Since our Amplified motion condition involved voluntary head motion, we sought to exclude potential confounds of activity in the head motor area. The hand knob region used in this study has been widely implemented by other groups to identify hand motor activity (Carlstedt et al., 2009; Cramer et al., 2001; Davare et al., 2008; Nakajima et al., 2020; Newton et al., 2005; Qiu et al., 2014). This definition of the hand motor area is also compatible with more recent understandings of the organization of the motor homunculus described by Gordon and colleagues (2023). This new homunculus model describes functional motor areas with distal body regions (e.g., hand) in the center and proximal body regions (e.g., shoulder) surrounding; this description still describes distal hand motor activity in one specific region of the motor cortex, in a comparable location to classical models of the homunculus. In the cerebellum, head motion is associated with activity in vermal lobule VI (Mottolese et al., 2013), which is an inter-hemispheric region that does not overlap with our defined motor cerebellum ROI.

Our hand-knob ROIs were drawn on each subject's anatomical image, which had high spatial contrast to allow for proper delineation of the region. ROIs were then transformed to each subject's functional space using a linear transformation (FLIRT). Since SE and ME analysis used different echoes for registration (second and first echo, respectively), the transformations may be slightly different. However, the echoes used in both SE and ME analyses had sufficient contrast to properly perform the registration steps.

In our study, we found activation extent in the selected hand-knob and cerebellum ROIs to be greater using the ME models compared to SE, likely due to increased sensitivity to activation aided by increased contrast-to-noise. Gonzalez-Castillo and colleagues (2016) had similar findings in their analysis of several types of task-fMRI data, showing a significant increase in activation extent from SE to ME models.   The beta coefficient results varied across ROIs. In the cerebellum, ME models demonstrated a significantly lower median beta coefficient than the SE model. This result is in line with the results of Gonzalez-Castillo and colleagues (2016), who observed a decrease in average effect size across different types of tasks.

Notably, our study found different trends for the beta coefficient in the hand-knob regions. With an ideal model, the beta coefficient would be similar between the Limited and Amplified conditions because the hand grasp was similar during both scans (**Figure 1A**) and therefore the magnitude of BOLD activation should also be similar. The Limited condition scan, performed by young, healthy individuals, with head motion minimized by tactile feedback of tape on their foreheads, is our best approximation of ground truth activation for this dataset. We expect that a reliable measure of activation from the Amplified condition scans would have a beta coefficient more similar to the Limited condition scans. In particular, the hand-knob regions are heavily studied to measure hand motor activity, and accounting for the potential confounding effects of head motion there is critical. For example, the increase in hand-knob beta coefficient values in the Amplified compared to Limited conditions observed when using the ME-OC model may be an artifactual increase caused by the added head motion. The ME-ICA model has more comparable beta coefficients between motion conditions, suggesting it does a better job at representing the true activation in the motor cortex during the hand grasp when motion is high. A related finding was observed in a visual retinotopy task fMRI experiment performed by Steel and colleagues (2022), who also reported an increase in reliability of population receptive field parameter estimates when using ME-ICA compared to ME-OC, evaluated through correlation of parameter values across several datasets from the same subject. We also found that ME-ICA leads to a lower relationship of beta coefficient with both average FD and task-correlation of movement

across all Limited and Amplified scans (**Supplemental Figure 4**), another observation that suggests improved performance compared to SE and ME-OC models.

Across both motion conditions, the median positive t-statistic increased in the ME models compared to SE, with no significant difference between ME-OC and ME-ICA. While Gonzalez-Castillo and colleagues (2016) similarly observed an increase in average t-statistic using ME models, they also found a significant increase from ME-OC to ME-ICA. As discussed above, in our study, the beta coefficient was lower in the ME-ICA versus ME-OC model during the Amplified condition. T-statistics increase with a larger beta coefficient and/or smaller model residuals; therefore, it is important to realize that the ME-ICA and ME-OC models had comparable t-statistics *despite* a lower ME-ICA beta coefficient. This indicates that the ME-ICA model likely had lower residuals of the model, in addition to a more reliable beta coefficient estimate in the hand-knob regions, during the Amplified condition. Although we cannot determine the accuracy of our activation estimates without complementary imaging modalities, these collective results suggest that ME-ICA is beneficial in our single-subject analysis of high-motion data.

4.4. In addition to head motion, end-tidal CO_2 is also correlated with the motor task
Previous studies have found that task-correlated breathing may be an additional complication in motor-task fMRI, causing artifactual activation through variations in CO_2 (Birn et al., 2006; Farthing et al., 2007). Shallow breathing during the task can cause an increase in CO_2, which leads to widespread vasodilation in the brain with varying timings (Bright et al., 2009; Chang et al., 2008). Although not the main focus of this study, we assessed the magnitude of this issue in our dataset by calculating the correlations between the end-tidal CO_2 regressors and the hand grasp regressors. Correlations varied considerably across subjects and scans, with some correlations greater than $|r| = 0.5$ (**Figure 6C**). The direction of these correlations was primarily negative; end-tidal CO_2 increased after the grasp periods (**Figure 6A**), likely due to shallower breaths during the task periods that lead to a gradual increase in arterial CO_2 (Abbott et al., 2005).

Our approach to mitigating the confounds of task-correlated breathing was to add an end-tidal CO_2 regressor to each model to account for the CO_2 variation during the scan. To understand the effect of adding this regressor to our model, we analyzed our datasets with models that did not include the CO_2 regressor. On a subject level, the primary difference was that adding a CO_2 regressor reduced some areas of apparent negative activation (**Figure 6B**). As the CO_2 regressor was negatively correlated with the task regressors, this difference suggests that some variance in the data that was originally anti-correlated with the task regressors can be attributed to the CO_2 regressor after its addition to the model. Future work focused on understanding these

physiological confounds during motor tasks may consider using a lagged general linear model approach that has been used previously to account for spatially-varying temporal alignment between the CO_2 regressor and the voxelwise BOLD fMRI timeseries (Moia et al., 2020, 2021; Stickland et al., 2021; Zvolanek et al., 2023). However, this exploration was outside the scope of this study, which focused on the use of ME-ICA to mitigate the effects of head motion.

Our findings emphasize the importance of recording and accounting for the effects of breathing during motor-task scans. Notably, non-motor tasks, such as working memory tasks, can similarly entrain breathing (Bright et al., 2020). If collecting CO_2 data is not feasible, a respiration belt can also be used to measure breathing; a respiratory variance per time (RVT) regressor derived from respiration belt data has been shown to explain similar variance in the BOLD signal compared to an end-tidal CO_2 regressor, when large breathing modulations are present (Zvolanek et al., 2023). In resting-state data, where breathing modulations are presumably smaller, RVT and end-tidal CO_2 are potentially complementary in explaining BOLD signal changes (Chang & Glover, 2009; Golestani et al., 2015). Hence, collection of both CO_2 and respiratory belt data may be useful depending on the study goals, magnitude of breathing variability, and feasibility of physiological signal acquisition.

4.5. Group-level activation maps are similar across models, even with increased head motion

Group-level analysis is commonly performed in motor-task fMRI studies to identify clusters of activation across subjects (Ehrsson et al., 2000; Keisker et al., 2009, 2010; Kuhtz-Buschbeck et al., 2008). To compare the three models' performance in datasets with low and high motion, we did group-level analyses within the Limited and Amplified conditions (**Figure 7**). Unlike with the subject-level maps, there were no large visual differences between the conditions and models. All maps showed motor activation in the expected hand-knob and one or both of the cerebellum motor regions. Lombardo and colleagues (2016) compared ME-ICA and ME-OC models during group-level task-fMRI analysis and found that effect size increased with ME-ICA; therefore, while we did not observe consistent differences at the group-level when using ME-ICA, this model may lead to group-level improvements in other instances. In this study, we found that SE models were robust to detecting motor activation in the motor cortex and cerebellum, even with datasets that have high motion. In contrast to the findings by Lombardo et al., we found inconsistent changes in beta coefficients between models. While the ME models showed an increase in median beta coefficient in the hand-knob regions compared to SE, there was no clear difference between the ME-OC and ME-ICA models, and this result was not seen in the cerebellum. Based on these

findings, it is difficult to make a conclusion about the quantitative differences between the SE, ME-OC, and ME-ICA models in group analysis of our dataset.

Given our observations, if a study aims to detect results at the group level, SE fMRI data may be sufficient to map motor activation during a simple motor task even in high motion datasets. However, in clinical populations that may exhibit high task-correlated motion, subject-level analysis is particularly important: clinical research questions and patient diagnosis may motivate achieving insight into single-subject neural activity patterns, and it may not even be feasible or appropriate to implement group analyses because of pathological differences in anatomy and vascular physiology between subjects. In these cases, our study demonstrates that ME-ICA is critical to disentangling the effects of head motion from true motor activation, resulting in more reliable subject-level conclusions.

4.6. Limitations

Our motion simulation used an instructed head nod to amplify the task-correlated head motion during the scans. However, this is not a perfect substitute for the increased head motion exhibited by clinical populations. The added voluntary head motion in our Amplified scans may add a *neural* confound to the motor activation analysis. To limit our sensitivity to the neural effects of the voluntary head motion, we manually drew a hand knob ROI to isolate the region of the motor cortex involved in hand motion from the region involved in head motion.

Another challenge is identifying a ground truth in our study. While there are expected areas of activation in the hand knob and cerebellum motor areas that we used as our a priori ROIs for analysis (J. Diedrichsen et al., 2011; Jörn Diedrichsen et al., 2009; Yousry et al., 1997), we did not have a clear ground truth for the magnitude of the beta coefficient and activation extent within these ROIs. The Limited motion condition in our study involved young, healthy individuals performing a task with low levels of motion; this was our best approximation of accurate beta coefficient measurements using fMRI methods. Electroencephalography (EEG) may be another method to determine beta coefficient accuracy in the cortical hand knob areas, though it would not be able to investigate subcortical cerebellum motor areas.

Additionally, ME-ICA is unlikely to salvage datasets with very large amounts of head motion. In our study, the Amplified motion dataset approximated the motion characteristics of Stroke Subject 1. However, Stroke Subject 2 exhibited much higher head motion, and we have not thoroughly tested whether ME-ICA would be effective in mitigating the effects of this extreme degree of head motion. However, our findings with our outlier with very high motion (Subject 1 Amplified dataset) suggest that ME-ICA may demonstrate similar improvements in datasets with

FD up to 1.8 (**Figure 4**). Future work should probe the limits of ME-ICA in dealing with large amounts of head motion.

4.7. Considerations for future studies

The hand grasping task studied here can be associated with high levels of task-correlated head motion, but other motor tasks may have lower associated head motion. For example, commonly used finger tapping tasks involve more isolated and limited movements and likely will not encounter the same challenges with head motion. However, tasks such as hand grasping, and more proximal or complicated upper extremity movements, are important in the study of different neurological pathways and clinical conditions such as stroke (Ellis et al., 2016; McPherson et al., 2018; Wilkins et al., 2020). Also, individuals with motor impairments might not be able to perform finger individuation tasks, but could perform hand grasping or other less precise movements for the study of brain motor activity.

Other external methods of minimizing head motion may be possible, such as using a customized head mold (Power et al., 2019) or applying tape across the participant's forehead to add tactile feedback of head motion (Krause et al., 2019). In this study, we implemented the addition of tape across the forehead to further reduce head motion during our Limited motion condition and allow for a better approximation of ground truth activation. Additionally, real-time motion tracking for prospective motion correction techniques can be implemented to reduce false activations (Maclaren et al., 2013; Schulz et al., 2014). Each of the previously described techniques can potentially be used in combination with ME-ICA to further minimize motion artifacts.

Although there are clear benefits to ME acquisition and ME-ICA analysis, there are also certain limitations. ME acquisition requires longer TRs in order to collect multiple echoes, which may not be feasible or desirable in every application. Manual classification of ICA components for ME-ICA analysis also takes some training for consistent classification across scans, and additional hands-on analysis time to perform the classification. We pre-determined a list of classification rules (*2.2.3*) to limit the variation in component classification. For large datasets, in which it may not be feasible to manually classify components, a strict set of rules similar to those used in this study can be applied to automatically classify components with the decision tree in tedana (Dupre et al., 2021).

5. Conclusion

Motor-task fMRI is prone to task-correlated head motion that confounds activation results, particularly in certain clinical populations. In this study, we collected an fMRI dataset from a healthy population who performed a hand grasp task with Limited and Amplified task-correlated head motion to simulate a motor-impaired population. We analyzed these data using three models: single-echo (SE), multi-echo optimally combined (ME-OC), and multi-echo independent component analysis (ME-ICA). On the subject level, ME models better dissociated the effects of head motion from the BOLD signal and led to increased activation extent and t-statistics in brain motor regions. In scans with high levels of motion, ME-ICA additionally mitigated artifacts and led to more reliable beta coefficient estimates. Overall, we show that ME-ICA is a useful tool for analyzing motor-task data with high levels of task-correlated head motion, which is particularly important in studies of clinical populations where subject-level analysis is the goal.

Acknowledgements

N.A.R. and K.M.Z. were supported by the National Institutes of Health under a training program (T32EB025766). K.M.Z was supported by the National Heart, Lung, and Blood Institute of the National Institutes of Health under Award Number F31HL166079. The content is solely the responsibility of the authors and does not necessarily represent the official views of the National Institutes of Health. This work was supported by the Center for Translational Imaging at Northwestern University. This research was also supported by the Spanish Ministry of Economy and Competitiveness (Ramon y Cajal Fellowship, RYC-2017- 21845), the Basque Government (BERC 2018-2021 and PIBA_2019_104), and the Spanish Ministry of Science, Innovation and Universities (MICINN; PID2019-105520GB-100).

References

Abbott, D. F., Opdam, H. I., Briellmann, R. S., & Jackson, G. D. (2005). Brief breath holding may confound functional magnetic resonance imaging studies. *Human Brain Mapping*, *24*(4), 284–290. https://doi.org/10.1002/hbm.20086

Araneda, R., Dricot, L., Ebner-Karestinos, D., Paradis, J., Gordon, A. M., Friel, K. M., & Bleyenheuft, Y. (2021). Brain activation changes following motor training in children with unilateral cerebral palsy: An fMRI study. *Annals of Physical and Rehabilitation Medicine*, *64*(3), 101502. https://doi.org/10.1016/j.rehab.2021.101502

Ashida, R., Cerminara, N. L., Edwards, R. J., Apps, R., & Brooks, J. C. W. (2019). Sensorimotor, language, and working memory representation within the human cerebellum. *Human Brain Mapping*, *40*(16), 4732–4747. https://doi.org/10.1002/hbm.24733

Behzadi, Y., Restom, K., Liau, J., & Liu, T. T. (2007). A component based noise correction method (CompCor) for BOLD and perfusion based fMRI. *NeuroImage*, *37*(1), 90–101. https://doi.org/10.1016/j.neuroimage.2007.04.042

Birn, R. M., Diamond, J. B., Smith, M. A., & Bandettini, P. A. (2006). Separating respiratory-variation-related fluctuations from neuronal-activity-related fluctuations in fMRI. *NeuroImage*, *31*(4), 1536–1548. https://doi.org/10.1016/j.neuroimage.2006.02.048

Bright, M. G., Bulte, D. P., Jezzard, P., & Duyn, J. H. (2009). Characterization of regional heterogeneity in cerebrovascular reactivity dynamics using novel hypocapnia task and BOLD fMRI. *NeuroImage*, *48*(1), 166–175. https://doi.org/10.1016/j.neuroimage.2009.05.026

Bright, M. G., & Murphy, K. (2013). Removing motion and physiological artifacts from intrinsic BOLD fluctuations using short echo data. *NeuroImage*, *64*(1), 526–537. https://doi.org/10.1016/j.neuroimage.2012.09.043

Bright, M. G., Whittaker, J. R., Driver, I. D., & Murphy, K. (2020). Vascular physiology drives functional brain networks. *NeuroImage*, *217*. https://doi.org/10.1016/j.neuroimage.2020.116907

Buur, P. F., Poser, B. A., & Norris, D. G. (2009). A dual echo approach to removing motion artefacts in fMRI time series. *NMR in Biomedicine*, *22*(5), 551–560. https://doi.org/10.1002/nbm.1371

Byars, A. W., Holland, S. K., Strawsburg, R. H., Bommer, W., Dunn, R. S., Schmithorst, V. J., & Plante, E. (2002). Practical aspects of conducting large-scale functional magnetic resonance imaging studies in children. *Journal of Child Neurology*, *17*(12), 885–890. https://doi.org/10.1177/08830738020170122201

Caballero-Gaudes, C., & Reynolds, R. C. (2017). Methods for cleaning the BOLD fMRI signal. *NeuroImage*, *154*(December 2016), 128–149. https://doi.org/10.1016/j.neuroimage.2016.12.018

Carlstedt, T., Hultgren, T., Nyman, T., & Hansson, T. (2009). Cortical activity and hand function restoration in a patient after spinal cord surgery. *Nature Reviews Neurology*, *5*(10), 571–574. https://doi.org/10.1038/nrneurol.2009.137

Chang, C., & Glover, G. H. (2009). Relationship between respiration, end-tidal CO2, and BOLD signals in resting-state fMRI. *NeuroImage*, *47*(4), 1381–1393. https://doi.org/10.1016/j.neuroimage.2009.04.048

Chang, C., Thomason, M. E., & Glover, G. H. (2008). Mapping and correction of vascular hemodynamic latency in the BOLD signal. *NeuroImage*, *43*(1), 90–102. https://doi.org/10.1016/j.neuroimage.2008.06.030

Ciavarro, M., Grande, E., Pavone, L., Bevacqua, G., De Angelis, M., di Russo, P., Morace, R., Committeri, G., Grillea, G., Bartolo, M., Paolini, S., & Esposito, V. (2021). Pre-surgical fMRI Localization of the Hand Motor Cortex in Brain Tumors: Comparison Between Finger Tapping Task and a New Visual-Triggered Finger Movement Task. *Frontiers in Neurology*, *12*(May). https://doi.org/10.3389/fneur.2021.658025

Cohen, A. D., Chang, C., & Wang, Y. (2021). Using Multiband Multi-echo Imaging to Improve the Robustness and Repeatability of Co-activation Pattern Analysis for Dynamic Functional Connectivity. *NeuroImage*, *243*(September), 118555. https://doi.org/10.1016/j.neuroimage.2021.118555

Cohen, A. D., Jagra, A. S., Visser, N. J., Yang, B., Fernandez, B., Banerjee, S., & Wang, Y. (2021). Improving the Breath-Holding CVR Measurement Using the Multiband Multi-Echo EPI Sequence. *Frontiers in Physiology*, *12*(February), 1–14. https://doi.org/10.3389/fphys.2021.619714

Cohen, A. D., Jagra, A. S., Yang, B., Fernandez, B., Banerjee, S., & Wang, Y. (2021). Detecting Task Functional MRI Activation Using the Multiband Multiecho (MBME) Echo-Planar Imaging (EPI) Sequence. *Journal of Magnetic Resonance Imaging*, *53*(5), 1366–1374. https://doi.org/10.1002/jmri.27448

Cohen, A. D., Nencka, A. S., & Wang, Y. (2018). Multiband multi-echo simultaneous ASL/BOLD for task-induced functional MRI. *PLoS ONE*, *13*(2), 1–21. https://doi.org/10.1371/journal.pone.0190427

Cohen, A. D., & Wang, Y. (2019). Improving the Assessment of Breath-Holding Induced Cerebral Vascular Reactivity Using a Multiband Multi-echo ASL/BOLD Sequence. *Scientific*

Reports, *9*(1), 1–12. https://doi.org/10.1038/s41598-019-41199-w

Cohen, A. D., Yang, B., Fernandez, B., Banerjee, S., & Wang, Y. (2021). Improved resting state functional connectivity sensitivity and reproducibility using a multiband multi-echo acquisition. *NeuroImage*, *225*, Neuroimage. https://doi.org/10.1016/j.neuroimage.2020.117461

Cox J.S., R. W. . H. (1996). AFNI: Software for analysis and visualization of functional magnetic resonance neuroimages. *Computers and Biomedical Research*, *29*(29), 162–173. https://ac-els-cdn-com.ezp-prod1.hul.harvard.edu/S0010480996900142/1-s2.0-S0010480996900142-main.pdf?_tid=c22bae7a-b8f5-4a8b-9dac-91c1ed841d53&acdnat=1549393398_e37181b8933a2ac88c2d7dc0eab14413

Cramer, S. C., Mark, A., Barquist, K., Nhan, H., Stegbauer, K. C., Price, R., Bell, K., Odderson, I. R., Esselman, P., & Maravilla, K. R. (2002). Motor cortex activation is preserved in patients with chronic hemiplegic stroke. *Annals of Neurology*, *52*(5), 607–616. https://doi.org/10.1002/ana.10351

Cramer, S. C., Nelles, G., Schaechter, J. D., Kaplan, J. D., Finklestein, S. P., & Rosen, B. R. (2001). A Functional MRI Study of Three Motor Tasks in the Evaluation of Stroke Recovery. *Neurorehabilitation and Neural Repair*, *15*(1), 1–8. https://doi.org/10.1177/154596830101500101

Cramer, S. C., Weisskoff, R. M., Schaechter, J. D., Nelles, G., Foley, M., Finklestein, S. P., & Rosen, B. R. (2002). Motor cortex activation is related to force of squeezing. *Human Brain Mapping*, *16*(4), 197–205. https://doi.org/10.1002/hbm.10040

Davare, M., Lemon, R., & Olivier, E. (2008). Selective modulation of interactions between ventral premotor cortex and primary motor cortex during precision grasping in humans. *Journal of Physiology*, *586*(11), 2735–2742. https://doi.org/10.1113/jphysiol.2008.152603

Diedrichsen, J., Maderwald, S., Küper, M., Thürling, M., Rabe, K., Gizewski, E. R., Ladd, M. E., & Timmann, D. (2011). Imaging the deep cerebellar nuclei: A probabilistic atlas and normalization procedure. *NeuroImage*, *54*(3), 1786–1794. https://doi.org/10.1016/j.neuroimage.2010.10.035

Diedrichsen, Jörn, Balsters, J. H., Flavell, J., Cussans, E., & Ramnani, N. (2009). A probabilistic MR atlas of the human cerebellum. *NeuroImage*, *46*(1), 39–46. https://doi.org/10.1016/j.neuroimage.2009.01.045

Dipasquale, O., Sethi, A., Laganà, M. M., Baglio, F., Baselli, G., Kundu, P., Harrison, N. A., & Cercignani, M. (2017). Comparing resting state fMRI de-noising approaches using multi- and single-echo acquisitions. *PLoS ONE*, *12*(3), 1–25.

https://doi.org/10.1371/journal.pone.0173289

Dupre, E., Salo, T., Ahmed, Z., Bandettini, P. A., Bottenhorn, K. L., Caballero-Gaudes, C., Logan, T. D., Gonzalez-Castillo, J., Heunis, S., Kundu, P., Laird, A. R., Markello, R., Markiewicz, C. J., Moia, S., Staden, I., Teves, J. B., Uruñuela, E., Vaziri-Pashkam, M., Whitaker, K., & Handwerker, D. A. (2021). TE-dependent analysis of multi-echo fMRI with tedana. *The Journal of Open Source Software*, *6*(59), 1–5. https://doi.org/10.21105/joss.03103

Ehrsson, H. H., Fagergren, A., Jonsson, T., Westling, G., Johansson, R. S., & Forssberg, H. (2000). Cortical activity in precision- versus power-grip tasks: An fMRI study. *Journal of Neurophysiology*, *83*(1), 528–536.

Ellis, M. D., Lan, Y., Yao, J., & Dewald, J. P. A. (2016). Robotic quantification of upper extremity loss of independent joint control or flexion synergy in individuals with hemiparetic stroke: a review of paradigms addressing the effects of shoulder abduction loading. *Journal of NeuroEngineering and Rehabilitation*, *13*(1), 1–11. https://doi.org/10.1186/s12984-016-0203-0

Evans, J. W., Kundu, P., Horovitz, S. G., & Bandettini, P. A. (2015). Separating slow BOLD from non-BOLD baseline drifts using multi-echo fMRI. *NeuroImage*, *105*, 189–197. https://doi.org/10.1016/j.neuroimage.2014.10.051

Farthing, J. P., Cummine, J., Borowsky, R., Chilibeck, P. D., Binsted, G., & Sarty, G. E. (2007). False activation in the brain ventricles related to task-correlated breathing in fMRI speech and motor paradigms. *Magnetic Resonance Materials in Physics, Biology and Medicine*, *20*(3), 157–168. https://doi.org/10.1007/s10334-007-0077-z

Field, A. S., Yen, Y. F., Burdette, J. H., & Elster, A. D. (2000). False cerebral activation on BOLD functional MR images: Study of low-amplitude motion weakly correlated to stimulus. *American Journal of Neuroradiology*, *21*(8), 1388–1396.

Freire, L., & Mangin, J. F. (2001). Motion correction algorithms may create spurious brain activations in the absence of subject motion. *NeuroImage*, *14*(3), 709–722. https://doi.org/10.1006/nimg.2001.0869

Friston, K. J., Williams, S., Howard, R., Frackowiak, R. S. J., & Turner, R. (1996). Movement-related effects in fMRI time-series. *Magnetic Resonance in Medicine*, *35*(3), 346–355. https://doi.org/10.1002/mrm.1910350312

Golestani, A. M., Chang, C., Kwinta, J. B., Khatamian, Y. B., & Jean Chen, J. (2015). Mapping the end-tidal CO2 response function in the resting-state BOLD fMRI signal: Spatial specificity, test-retest reliability and effect of fMRI sampling rate. *NeuroImage, 104*, 266–

277. https://doi.org/10.1016/j.neuroimage.2014.10.031

Gonzalez-Castillo, J., Panwar, P., Buchanan, L. C., Caballero-Gaudes, C., Handwerker, D. A., Jangraw, D. C., Zachariou, V., Inati, S. J., Roopchansingh, V., Derbyshire, J. A., & Bandettini, P. A. (2016). Evaluation of multi-echo ICA denoising for task based fMRI studies: Block designs, rapid event-related designs, and cardiac-gated fMRI. *NeuroImage*, *141*, 452–468. https://doi.org/10.1016/j.neuroimage.2016.07.049

Gordon, E. M., Chauvin, R. J., Van, A. N., Rajesh, A., Nielsen, A., Newbold, D. J., Lynch, C. J., Seider, N. A., Krimmel, S. R., Scheidter, K. M., Monk, J., Miller, R. L., Metoki, A., Montez, D. F., Zheng, A., Elbau, I., Madison, T., Nishino, T., Myers, M. J., … Dosenbach, N. U. F. (2023). A somato-cognitive action network alternates with effector regions in motor cortex. *Nature*, *617*(May). https://doi.org/10.1038/s41586-023-05964-2

Griffanti, L., Douaud, G., Bijsterbosch, J., Evangelisti, S., Alfaro-Almagro, F., Glasser, M. F., Duff, E. P., Fitzgibbon, S., Westphal, R., Carone, D., Beckmann, C. F., & Smith, S. M. (2017). Hand classification of fMRI ICA noise components. *NeuroImage, 154*(June 2016), 188–205. https://doi.org/10.1016/j.neuroimage.2016.12.036

Griffanti, L., Salimi-Khorshidi, G., Beckmann, C. F., Auerbach, E. J., Douaud, G., Sexton, C. E., Zsoldos, E., Ebmeier, K. P., Filippini, N., Mackay, C. E., Moeller, S., Xu, J., Yacoub, E., Baselli, G., Ugurbil, K., Miller, K. L., & Smith, S. M. (2014). ICA-based artefact removal and accelerated fMRI acquisition for improved resting state network imaging. *NeuroImage*, *95*, 232–247. https://doi.org/10.1016/j.neuroimage.2014.03.034

Grootoonk, S., Hutton, C., Ashburner, J., Howseman, A. M., Josephs, O., Rees, G., Friston, K. J., & Turner, R. (2000). Characterization and correction of interpolation effects in the realignment of fMRI time series. *NeuroImage*, *11*(1), 49–57. https://doi.org/10.1006/nimg.1999.0515

Guediche, S., de Bruin, A., Caballero-Gaudes, C., Baart, M., & Samuel, A. G. (2021). Second-language word recognition in noise: Interdependent neuromodulatory effects of semantic context and crosslinguistic interactions driven by word form similarity. *NeuroImage*, *237*(May), 118168. https://doi.org/10.1016/j.neuroimage.2021.118168

Hajnal, J. V., Myers, R., Oatridge, A., Schwieso, J. E., Young, I. R., & Bydder, G. M. (1994). Artifacts due to stimulus correlated motion in functional imaging of the brain. *Magnetic Resonance in Medicine*, *31*(3), 283–291. https://doi.org/10.1002/mrm.1910310307

Hannanu, F. F., Goundous, I., Detante, O., Naegele, B., & Jaillard, A. (2020). Spatiotemporal patterns of sensorimotor fMRI activity influence hand motor recovery in subacute stroke: A longitudinal task-related fMRI study. *Cortex*, *129*, 80–98.

https://doi.org/10.1016/j.cortex.2020.03.024

Heilbrun, M. P., Lee, J. N., & Alvord, L. (2001). Practical application of fMRI for surgical planning. *Stereotactic and Functional Neurosurgery*, *76*(3–4), 168–174. https://doi.org/10.1159/000066715

Hermsdörfer, J., Hagl, E., Nowak, D. A., & Marquardt, C. (2003). Grip force control during object manipulation in cerebral stroke. *Clinical Neurophysiology*, *114*(5), 915–929. https://doi.org/10.1016/S1388-2457(03)00042-7

Ing, A., & Schwarzbauer, C. (2012). A dual echo approach to motion correction for functional connectivity studies. *NeuroImage*, *63*(3), 1487–1497. https://doi.org/10.1016/j.neuroimage.2012.07.042

J.P., P., K.J., S., P.A., B., A., C., B., P., & A., B. (2007). Ankle dorsiflexion fMRI in children with cerebral palsy undergoing intensive body-weight-supported treadmill training: A pilot study. *Developmental Medicine and Child Neurology*, *49*(1), 39–44.

Jenkinson, M., Beckmann, C. F., Behrens, T. E. J., Woolrich, M. W., & Smith, S. M. (2012). FSL. *NeuroImage*, *62*, 782–790. https://doi.org/10.1016/j.neuroimage.2011.09.015

Johnstone, T., Ores Walsh, K. S., Greischar, L. L., Alexander, A. L., Fox, A. S., Davidson, R. J., & Oakes, T. R. (2006). Motion correction and the use of motion covariates in multiple-subject fMRI analysis. *Human Brain Mapping*, *27*(10), 779–788. https://doi.org/10.1002/hbm.20219

Kay, K. N., Rokem, A., Winawer, J., Dougherty, R. F., & Wandell, B. A. (2013). GLMdenoise: A fast, automated technique for denoising task-based fMRI data. *Frontiers in Neuroscience*, *7*(7 DEC), 1–15. https://doi.org/10.3389/fnins.2013.00247

Keisker, B., Hepp-Reymond, M. C., Blickenstorfer, A., & Kollias, S. S. (2010). Differential representation of dynamic and static power grip force in the sensorimotor network. *European Journal of Neuroscience*, *31*(8), 1483–1491. https://doi.org/10.1111/j.1460-9568.2010.07172.x

Keisker, B., Hepp-Reymond, M. C., Blickenstorfer, A., Meyer, M., & Kollias, S. S. (2009). Differential force scaling of fine-graded power grip force in the sensorimotor network. *Human Brain Mapping*, *30*(8), 2453–2465. https://doi.org/10.1002/hbm.20676

Kochiyama, T., Morita, T., Okada, T., Yonekura, Y., Matsumura, M., & Sadato, N. (2005). Removing the effects of task-related motion using independent-component analysis. *NeuroImage*, *25*(3), 802–814. https://doi.org/10.1016/j.neuroimage.2004.12.027

Kotsoni, E., Byrd, D., & Casey, B. J. (2006). Special considerations for functional magnetic resonance imaging of pediatric populations. In *Journal of Magnetic Resonance Imaging*

(Vol. 23, Issue 6, pp. 877–886). https://doi.org/10.1002/jmri.20578

Krause, F., Benjamins, C., Eck, J., Lührs, M., van Hoof, R., & Goebel, R. (2019). Active head
motion reduction in magnetic resonance imaging using tactile feedback. *Human Brain
Mapping*, *40*(14), 4026–4037. https://doi.org/10.1002/hbm.24683

Kuhtz-Buschbeck, J. P., Gilster, R., Wolff, S., Ulmer, S., Siebner, H., & Jansen, O. (2008). Brain
activity is similar during precision and power gripping with light force: An fMRI study.
NeuroImage, *40*(4), 1469–1481. https://doi.org/10.1016/j.neuroimage.2008.01.037

Kundu, P., Brenowitz, N. D., Voon, V., Worbe, Y., Vértes, P. E., Inati, S. J., Saad, Z. S.,
Bandettini, P. A., & Bullmore, E. T. (2013). Integrated strategy for improving functional
connectivity mapping using multiecho fMRI. *Proceedings of the National Academy of
Sciences of the United States of America*, *110*(40), 16187–16192.
https://doi.org/10.1073/pnas.1301725110

Lombardo, M. V., Auyeung, B., Holt, R. J., Waldman, J., Ruigrok, A. N. V., Mooney, N.,
Bullmore, E. T., Baron-Cohen, S., & Kundu, P. (2016). Improving effect size estimation and
statistical power with multi-echo fMRI and its impact on understanding the neural systems
supporting mentalizing. *NeuroImage*, *142*, 55–66.
https://doi.org/10.1016/j.neuroimage.2016.07.022

Ludman, C. N., Cooper, T. G., Ploutz-Synder, L. L., Potchen, E. J., & Meyer, R. A. (1996). Force
of voluntary exercise does not affect sensorimotor cortex activation as detected by
functional MRI at 1.5 T. *NMR in Biomedicine*, *9*(5), 228–232.
https://doi.org/10.1002/(SICI)1099-1492(199608)9:5<228::AID-NBM420>3.0.CO;2-1

Lynch, C. J., Power, J. D., Scult, M. A., Dubin, M., Gunning, F. M., & Liston, C. (2020). Rapid
Precision Functional Mapping of Individuals Using Multi-Echo fMRI. *Cell Reports*, *33*(12),
108540. https://doi.org/10.1016/j.celrep.2020.108540

Maclaren, J., Herbst, M., Speck, O., & Zaitsev, M. (2013). Prospective motion correction in brain
imaging: A review. *Magnetic Resonance in Medicine*, *69*(3), 621–636.
https://doi.org/10.1002/mrm.24314

Martin, J. A., Zimmermann, N., Scheef, L., Jankowski, J., Paus, S., Schild, H. H., Klockgether,
T., & Boecker, H. (2019). Disentangling motor planning and motor execution in
unmedicated de novo Parkinson's disease patients: An fMRI study. *NeuroImage: Clinical*,
22(March), 101784. https://doi.org/10.1016/j.nicl.2019.101784

McPherson, J. G., Chen, A., Ellis, M. D., Yao, J., Heckman, C. J., & Dewald, J. P. A. (2018).
Progressive recruitment of contralesional cortico-reticulospinal pathways drives motor
impairment post stroke. *Journal of Physiology, 596*(7), 1211–1225.

https://doi.org/10.1113/JP274968

McSwain, S. D., Hamel, D. S., Smith, P. B., Gentile, M. A., Srinivasan, S., Meliones, J. N., & Cheifetz, I. M. (2010). End-tidal and arterial carbon dioxide measurements correlate across all levels of physiologic dead space. *Respiratory Care*, *55*(3), 288–293.

Menon, R. S., Ogawa, S., Tank, D. W., & Uğurbil, K. (1993). 4 Tesla gradient recalled echo characteristics of photic stimulation-induced signal changes in the human primary visual cortex. *Magnetic Resonance in Medicine*, *30*(3), 380–386. https://doi.org/10.1002/mrm.1910300317

Moeller, S., Yacoub, E., Olman, C. A., Auerbach, E., Strupp, J., Harel, N., & Ugurbil, K. (2010). Multiband multislice GE-EPI at 7 Tesla, with 16-fold acceleration using partial parallel imaging with application to high spatial and temporal whole-brain FMRI. *Magn. Reson. Med.*, *63*(5), 1144–1153. https://doi.org/10.1002/mrm.22361.Multiband

Moia, S., Stickland, R. C., Ayyagari, A., Termenon, M., Caballero-Gaudes, C., & Bright, M. G. (2020). Voxelwise optimization of hemodynamic lags to improve regional CVR estimates in breath-hold fMRI. *Proceedings of the Annual International Conference of the IEEE Engineering in Medicine and Biology Society, EMBS*, *2020-July*, 1489–1492. https://doi.org/10.1109/EMBC44109.2020.9176225

Moia, S., Termenon, M., Uruñuela, E., Chen, G., Stickland, R. C., Bright, M. G., & Caballero-Gaudes, C. (2021). ICA-based denoising strategies in breath-hold induced cerebrovascular reactivity mapping with multi echo BOLD fMRI. *NeuroImage*, *233*(March), 1–18. https://doi.org/10.1016/j.neuroimage.2021.117914

Morgan, V. L., Dawant, B. M., Li, Y., & Pickens, D. R. (2007). Comparison of fMRI statistical software packages and strategies for analysis of images containing random and stimulus-correlated motion. *Computerized Medical Imaging and Graphics*, *31*(6), 436–446. https://doi.org/10.1016/j.compmedimag.2007.04.002

Mottolese, C., Richard, N., Harquel, S., Szathmari, A., Sirigu, A., & Desmurget, M. (2013). Mapping motor representations in the human cerebellum. *Brain*, *136*(1), 330–342. https://doi.org/10.1093/brain/aws186

Mumford, J. A., Poline, J. B., & Poldrack, R. A. (2015). Orthogonalization of regressors in fMRI models. *PLoS ONE*, *10*(4), 1–11. https://doi.org/10.1371/journal.pone.0126255

Muschelli, J., Nebel, M. B., Caffo, B. S., Barber, A. D., Pekar, J. J., & Mostofsky, S. H. (2014). Reduction of motion-related artifacts in resting state fMRI using aCompCor. *NeuroImage*, *96*, 22–35. https://doi.org/10.1016/j.neuroimage.2014.03.028

Nakajima, R., Kinoshita, M., & Nakada, M. (2020). Motor Functional Reorganization Is Triggered

by Tumor Infiltration Into the Primary Motor Area and Repeated Surgery. *Frontiers in Human Neuroscience*, *14*(August), 1–13. https://doi.org/10.3389/fnhum.2020.00327

Newton, J. M., Sunderland, A., & Gowland, P. A. (2005). fMRI signal decreases in ipsilateral primary motor cortex during unilateral hand movements are related to duration and side of movement. *NeuroImage*, *24*(4), 1080–1087. https://doi.org/10.1016/j.neuroimage.2004.10.003

Ogawa, S., Menon, R., Tank, D., Kim, S., Merkle, H., Ellermann, J., & Ugurbilt, K. (1993). Functional brain mapping by blood oxygenation level-dependent. *Biophys*, *64*(3), 803–812.

Olafsson, V., Kundu, P., Wong, E. C., Bandettini, P. A., & Liu, T. T. (2015). Enhanced identification of BOLD-like components with multi-echo simultaneous multi-slice (MESMS) fMRI and multi-echo ICA. *NeuroImage*, *112*, 43–51. https://doi.org/10.1016/j.neuroimage.2015.02.052

Patriat, R., Reynolds, R. C., & Birn, R. M. (2017). An improved model of motion-related signal changes in fMRI. *NeuroImage*, *144*, 74–82. https://doi.org/10.1016/j.neuroimage.2016.08.051

Péran, P., Nemmi, F., Dutilleul, C., Finamore, L., Falletta Caravasso, C., Troisi, E., Iosa, M., Sabatini, U., & Grazia Grasso, M. (2020). Neuroplasticity and brain reorganization associated with positive outcomes of multidisciplinary rehabilitation in progressive multiple sclerosis: A fMRI study. *Multiple Sclerosis and Related Disorders*, *42*(October 2019), 102127. https://doi.org/10.1016/j.msard.2020.102127

Power, J. D., Barnes, K. A., Snyder, A. Z., Schlaggar, B. L., & Petersen, S. E. (2012). Spurious but systematic correlations in functional connectivity MRI networks arise from subject motion. *NeuroImage*, *59*(3), 2142–2154. https://doi.org/10.1016/j.neuroimage.2011.10.018

Power, J. D., Mitra, A., Laumann, T. O., Snyder, A. Z., Schlaggar, B. L., & Petersen, S. E. (2014). Methods to detect, characterize, and remove motion artifact in resting state fMRI. *NeuroImage*, *84*, 320–341. https://doi.org/10.1016/j.neuroimage.2013.08.048

Power, J. D., Plitt, M., Kundu, P., Bandettini, P. A., & Martin, A. (2017). Temporal interpolation alters motion in fMRI scans: Magnitudes and consequences for artifact detection. *PLoS ONE*, *12*(9), 1–20. https://doi.org/10.1371/journal.pone.0182939

Power, J. D., Silver, B. M., Silverman, M. R., Ajodan, E. L., Bos, D. J., & Jones, R. M. (2019). Customized head molds reduce motion during resting state fMRI scans. *NeuroImage*, *189*(January), 141–149. https://doi.org/10.1016/j.neuroimage.2019.01.016

Pruim, R. H. R., Mennes, M., van Rooij, D., Llera, A., Buitelaar, J. K., & Beckmann, C. F. (2015). ICA-AROMA: A robust ICA-based strategy for removing motion artifacts from fMRI data.

NeuroImage, *112*, 267–277. https://doi.org/10.1016/j.neuroimage.2015.02.064

Qiu, T. ming, Yan, C. gan, Tang, W. jun, Wu, J. song, Zhuang, D. xiao, Yao, C. jun, Lu, J. feng, Zhu, F. ping, Mao, Y., & Zhou, L. fu. (2014). Localizing hand motor area using resting-state fMRI: validated with direct cortical stimulation. *Acta Neurochirurgica*, *156*(12), 2295–2302. https://doi.org/10.1007/s00701-014-2236-0

Sabatini, U., Boulanouar, K., Fabre, N., Chollet, F., Berry, I., Colonnese, C., Manelfe, C., Bozzao, L., & Rascol, O. (1998). Cortical motor reorganization in akinetic Parkinsonian patients: A functional magnetic resonance imaging study. *NeuroImage*, *7*(4 PART II), 394–403. https://doi.org/10.1016/s1053-8119(18)31309-0

Salimi-Khorshidi, G., Douaud, G., Beckmann, C. F., Glasser, M. F., Griffanti, L., & Smith, S. M. (2014). Automatic denoising of functional MRI data: Combining independent component analysis and hierarchical fusion of classifiers. *NeuroImage*, *90*, 449–468. https://doi.org/10.1016/j.neuroimage.2013.11.046

Satterthwaite, T. D., Wolf, D. H., Loughead, J., Ruparel, K., Elliott, M. A., Hakonarson, H., Gur, R. C., & Gur, R. E. (2012). Impact of in-scanner head motion on multiple measures of functional connectivity: Relevance for studies of neurodevelopment in youth. *NeuroImage*, *60*(1), 623–632. https://doi.org/10.1016/j.neuroimage.2011.12.063

Schulz, J., Siegert, T., Bazin, P. L., Maclaren, J., Herbst, M., Zaitsev, M., & Turner, R. (2014). Prospective slice-by-slice motion correction reduces false positive activations in fMRI with task-correlated motion. *NeuroImage*, *84*, 124–132. https://doi.org/10.1016/j.neuroimage.2013.08.006

Seto, E., Sela, G., McIlroy, W. E., Black, S. E., Staines, W. R., Bronskill, M. J., McIntosh, A. R., & Graham, S. J. (2001). Quantifying head motion associated with motor tasks used in fMRI. *NeuroImage*, *14*(2), 284–297. https://doi.org/10.1006/nimg.2001.0829

Setsompop, K., Gagoski, B. A., Polimeni, J. R., Witzel, T., Wedeen, V. J., & Wald, L. L. (2012). Blipped-Controlled Aliasing in Parallel Imaging (blipped-CAIPI) for simultaneous multi-slice EPI with reduced g-factor penalty. *Magn. Reson. Med.*, *67*(5), 1210–1224. https://doi.org/10.1002/mrm.23097.Blipped-Controlled

Siegel, J. S., Power, J. D., Dubis, J. W., Vogel, A. C., Church, J. A., Schlaggar, B. L., & Petersen, S. E. (2014). Statistical improvements in functional magnetic resonance imaging analyses produced by censoring high-motion data points. *Human Brain Mapping*, *35*(5), 1981–1996. https://doi.org/10.1002/hbm.22307

Smyser, C. D., Inder, T. E., Shimony, J. S., Hill, J. E., Degnan, A. J., Snyder, A. Z., & Neil, J. J. (2010). Longitudinal analysis of neural network development in preterm infants. *Cerebral*

Cortex, *20*(12), 2852–2862. https://doi.org/10.1093/cercor/bhq035

Soltysik, D. A., & Hyde, J. S. (2006). Strategies for block-design fMRI experiments during task-related motion of structures of the oral cavity. *NeuroImage*, *29*(4), 1260–1271. https://doi.org/10.1016/j.neuroimage.2005.08.063

Spencer, R. M. C., Verstynen, T., Brett, M., & Ivry, R. (2007). Cerebellar activation during discrete and not continuous timed movements: An fMRI study. *NeuroImage*, *36*(2), 378–387. https://doi.org/10.1016/j.neuroimage.2007.03.009

Steel, A., Garcia, B. D., Silson, E. H., & Robertson, C. E. (2022). Evaluating the efficacy of multi-echo ICA denoising on model-based fMRI. *NeuroImage*, *264*(September), 119723. https://doi.org/10.1016/j.neuroimage.2022.119723

Stephan, K. E., Magnotta, V. A., White, T., Arndt, S., Flaum, M., O'Leary, D. S., & Andreasen, N. C. (2001). Effects of olanzapine on cerebellar functional connectivity in schizophrenia measured by fMRI during a simple motor task. *Psychological Medicine*, *31*(6), 1065–1078. https://doi.org/10.1017/S0033291701004330

Stickland, R. C., Zvolanek, K. M., Moia, S., Ayyagari, A., Caballero-Gaudes, C., & Bright, M. G. (2021). A practical modification to a resting state fMRI protocol for improved characterization of cerebrovascular function. *NeuroImage*, *239*(February), 118306. https://doi.org/10.1016/j.neuroimage.2021.118306

Stoodley, C. J. (2012). The cerebellum and cognition: Evidence from functional imaging studies. *Cerebellum*, *11*(2), 352–365. https://doi.org/10.1007/s12311-011-0260-7

Taylor, P. A., Reynolds, R. C., Calhoun, V., Gonzalez-Castillo, J., Handwerker, D. A., Bandettini, P. A., Mejia, A. F., & Chen, G. (2023). Highlight Results, Don't Hide Them: Enhance interpretation, reduce biases and improve reproducibility. *NeuroImage*, *274*(April 2023), 2022.10.26.513929. https://doi.org/10.1016/j.neuroimage.2023.120138

Thickbroom, G. W., Phillips, B. A., Morris, I., Byrnes, M. L., & Mastaglia, F. L. (1998). Isometric force-related activity in sensorimotor cortex measured with functional MRI. *Experimental Brain Research*, *121*(1), 59–64. https://doi.org/10.1007/s002210050437

Thickbroom, G. W., Phillips, B. A., Morris, I., Byrnes, M., Sacco, P., & Mastaglia, F. L. (1999). Differences in functional magnetic resonance imaging of sensorimotor cortex during static and dynamic finger flexion. *Experimental Brain Research*, *126*(3), 431–438. https://doi.org/10.1007/s002210050749

Tisdall, M. D., Reuter, M., Qureshi, A., Buckner, R. L., Fischl, B., & van der Kouwe, A. J. W. (2016). Prospective motion correction with volumetric navigators (vNavs) reduces the bias and variance in brain morphometry induced by subject motion. *NeuroImage*, *127*, 11–22.

https://doi.org/10.1016/j.neuroimage.2015.11.054

Uruñuela, E. (2021). *Rica*. GitHub Repository.

van Dijk, K. R. A., Sabuncu, M. R., & Buckner, R. L. (2012). The influence of head motion on intrinsic functional connectivity MRI. *NeuroImage*, *59*(1), 431–438. https://doi.org/10.1016/j.neuroimage.2011.07.044

Vysotski, S., Madura, C., Swan, B., Holdsworth, R., Lin, Y., Munoz Del Rio, A., Wood, J., Kundu, B., Penwarden, A., Voss, J., Gallagher, T., Nair, V. A., Field, A., Garcia-Ramos, C., Meyerand, M. E., Baskaya, M., Prabhakaran, V., & Kuo, J. S. (2018). Preoperative FMRI associated with decreased mortality and morbidity in brain tumor patients. *Interdisciplinary Neurosurgery: Advanced Techniques and Case Management*, *13*(November 2017), 40–45. https://doi.org/10.1016/j.inat.2018.02.001

Ward, N. S., Brown, M. M., Thompson, A. J., & Frackowiak, R. S. J. (2003a). Neural correlates of motor recovery after stroke: A longitudinal fMRI study. *Brain*, *126*(11), 2476–2496. https://doi.org/10.1093/brain/awg245

Ward, N. S., Brown, M. M., Thompson, A. J., & Frackowiak, R. S. J. (2003b). Neural correlates of outcome after stroke: A cross-sectional fMRI study. *Brain*, *126*(6), 1430–1448. https://doi.org/10.1093/brain/awg145

Wilkins, K. B., Yao, J., Owen, M., Karbasforoushan, H., Carmona, C., & Dewald, J. P. A. (2020). Limited capacity for ipsilateral secondary motor areas to support hand function post-stroke. *Journal of Physiology*, *598*(11), 2153–2167. https://doi.org/10.1113/JP279377

Wilkinson, I. D., Romanowski, C. A. J., Jellinek, D. A., Morris, J., & Griffiths, P. D. (2003). Motor functional MRI for pre-operative and intraoperative neurosurgical guidance. *British Journal of Radiology*, *76*(902), 98–103. https://doi.org/10.1259/bjr/66817309

Wu, T., & Hallett, M. (2005). A functional MRI study of automatic movements in patients with Parkinson's disease. *Brain*, *128*(10), 2250–2259. https://doi.org/10.1093/brain/awh569

Xu, Y., Tong, Y., Liu, S., Chow, H. M., AbdulSabur, N. Y., Mattay, G. S., & Braun, A. R. (2014). Denoising the speaking brain: Toward a robust technique for correcting artifact-contaminated fMRI data under severe motion. *NeuroImage*, *103*, 33–47. https://doi.org/10.1016/j.neuroimage.2014.09.013

Yousry, T. A., Schmid, U. D., Alkadhi, H., Schmidt, D., Peraud, A., Buettner, A., & Winkler, P. (1997). Localization of the motor hand area to a knob on the precentral gyrus. A new landmark. *Brain*, *120*(1), 141–157. https://doi.org/10.1093/brain/120.1.141

Zvolanek, K. M., Moia, S., Dean, J. N., Stickland, R. C., Caballero-Gaudes, C., & Bright, M. G. (2023). Comparing end-tidal CO2, respiration volume per time (RVT), and average gray

matter signal for mapping cerebrovascular reactivity amplitude and delay with breath-hold task BOLD fMRI. *NeuroImage*, *272*(March), 2022.11.28.517116. https://doi.org/10.1101/2022.11.28.517116