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# Optical-flow analysis toolbox for characterization of spatiotemporal dynamics in mesoscale optical imaging of brain activity



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#### ABSTRACT

Wide-field optical imaging techniques constitute powerful tools to investigate mesoscale neuronal activity. The sampled data constitutes a sequence of image frames in which one can investigate the flow of brain activity starting and terminating at source and sink locations respectively. Approaches to the analyses of information flow include qualitative assessment to identify sources and sinks of activity as well as their trajectories, and quantitative measurements based on computing the temporal variation of the intensity of pixels. Furthermore, in a few studies estimates of wave motion have been reported using optical-flow techniques from computer vision. However, a comprehensive toolbox for the quantitative analyses of mesoscale brain activity data is still lacking. We present a graphical-user-interface toolbox based in Matlab® for investigating the spatiotemporal dynamics of mesoscale brain activity using optical-flow analyses. The toolbox includes the implementation of three optical-flow methods namely Horn-Schunck, Combined Local-Global, and Temporospatial algorithms for estimating velocity vector fields of flow of mesoscale brain activity. From the velocity vector fields we determined the locations of sources and sinks as well as the trajectories and temporal velocities of flow of activity. Using simulated data as well as experimentally derived sensory-evoked voltage and calcium imaging data from mice, we compared the efficacy of the three optical-flow methods for determining spatiotemporal dynamics. Our results indicate that the combined local-global method we employed, yields the best results for estimating wave motion. The automated approach permits rapid and effective quantification of mesoscale brain dynamics and may facilitate the study of brain function in response to new experiences or pathology.

# Introduction

Mesoscale optical imaging of brain activity allows the study of physiological models of brain functions at the network and system levels. The activity of thousands of neurons can be recorded simultaneously to study how networks of neurons communicate with each other. Important regional-scale information has been revealed with mesoscale optical imaging and it is increasingly being used to further our understanding of cortical functional connectivity and flow of information over the cortical surface (Frostig, 2009). In rodent brain imaging, mesoscale assessments correspond to spatial dimensions ranging from hundreds of micrometers to a few millimeters. Various methods have been developed to monitor mesoscale brain activity including optical imaging approaches and multielectrode electrophysiology (Nunez and Srinivasan, 2006; Hillman, 2007; Townsend et al., 2015; Grinvald et al., 2016; Obien et al., 2015). In optical imaging, the brain is illuminated with light and imaged to detect neuronal activity at the population level. This neuronal activity is transduced into an optical signal by either intrinsic or extrinsic reporters and interpreted as changes in the intensity of captured light (Frostig, 2009; Grinvald et al., 2016; Kalatsky and Stryker, 2003; Scanziani and Häusser, 2009). In intrinsic imaging, neuronal activity is inferred from either changes in reflectance of light from tissue due to oxygenation states of hemoglobin (Nemoto et al., 1997; Biswal et al., 2011) or from changes in auto-fluorescence of endogenous molecules such as flavoproteins (Reinert et al., 2004; Husson and Issa, 2009; Shibuki et al., 2009). In extrinsic imaging on the other hand, exogenous fluorescent molecules are introduced into the brain and neuronal activity is inferred from fluorescence changes modulated by either neuronal membrane voltage (voltage imaging), or concentration of an intracellular ion (e.g. calcium imaging) or an extracellular molecule (e.g. glutamate) (Xie et al., 2016; Storace et al., 2015; Abdelfattah et al., 2016; Akemann et al., 2012; Chen et al., 2013). Whichever method is selected, the data obtained with optical imaging is a set of images whereby the intensity of each pixel is indicative of the summed activity of nearby neurons. Mesoscale brain activity is also recorded with multielectrode electrophysiology in

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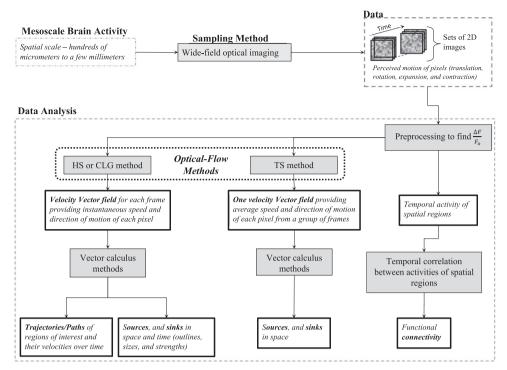


Fig. 1. Flow diagram summarizing mesoscale brain activity data acquisition and analysis. The data consists of perceived motion of pixels i.e. traveling waves originating at sources and vanishing at sink locations. The most common analysis includes observing temporal activity of pixels or regions of interest and determining functional connectivity between spatial regions using temporal correlations. The less common analysis is estimating wave motion with "optical-flow methods" (borrowed from computer vision) in which the spatiotemporal dynamics in brain activity are characterized for determining source/sink locations and properties of traveling waves i.e. trajectories and temporal speeds of pixels. We compare the performance of three optical flow methods; Horn-Schunk (HS), Combined local-global (CLG), and Temporospatial (TS) for analyzing optical imaging data. We present a graphical user interface based Matlab toolbox for mesoscale brain activity data analysis. Shaded boxes show methods included in the toolbox and bold outlined boxes show output entities of interest.

which electrodes are spread over surface of the brain to enable spatially discrete sampling of neuronal activity. The electrophysiology data can also be visualized as a set of 2D images where the intensity of each pixel would represent the voltage measured by an electrode (Buzsáki et al., 2012). Thus, mesoscale sampling of brain activity can capture the spatiotemporal dynamics of networks of neurons resulting in data with one temporal and two spatial dimensions (Fig. 1).

There are a variety of features that can be observed from optical imaging of brain activity. For example, when a group of neurons become active in response to a stimulus or from ongoing spontaneous activity, a source-like structure can be observed whereby pixel intensities corresponding to the site of activation gradually increase. The observed shape of the source-like activity in 2D can vary and may take the form of a point, line, circle, ellipse, or a combination of these shapes depending upon the spatial organization of the active neurons (Huang et al., 2010; Takagaki et al., 2011; Mohajerani et al., 2013). As time passes from the moment of activation, the source activity might weaken and disappear or sink in the same spatial region it originated from. In contrast, it might travel like a wave (Lilly, 1954) to another region following a continuous path/trajectory before weakening and disappearing into a sink where pixel intensities gradually decrease. The sink might also have shapes in 2D similar to the source activity. For propagating activity, the paths might be a mixture of translational, rotational, expansion or compression trajectories. If multiple brain areas are simultaneously active, multiple sources, travelling waves, and sinks would be observable. The observation of brain activity at the mesoscale hence mimics the motion or flow of waves.

The most common analysis used for the extraction of physiological features from optical imaging of brain activity is the quantitative assessment of temporal variation of intensity by plotting the average intensity of pixels versus time for defined regions of interest. More recently, the temporal correlations of optical signals were used to determine the functional connectivity between different regions (Fig. 1) (Mohajerani et al., 2013; Vanni and Murphy, 2014; Chan et al., 2015;

McVea et al., 2012; White et al., 2011; Minderer et al., 2012; Kuhn et al., 2008). To identify sources, sinks, and activity trajectories, visual inspection was often used (Mohajerani et al., 2013; Han et al., 2008). However, few studies have reported estimates of neuronal activity spread using "optical-flow" methods which were first developed in the field of computer vision. In "optical-flow" approaches, velocity vector flow fields are calculated to determine speeds and directions of motion (Paragios et al., 2006; Sun et al., 2010). From the velocity vector fields, locations of sources and sinks are estimated using vector calculus methods. Inouye et al. (1994), Inouye et al. (1994) and Inouye et al. (1995) reported the use of Horn-Schunk (HS) method (Horn and Schunck, 1981) to determine the flow of brain oscillations over the human scalp in flattened (3D surface to 2D) electroencephalography data. Takagaki et al. (2011) developed and reported a new method (Temporospatial - TS) in which temporal correlation of a given pixel with neighboring pixels was used to estimate local motion in voltage sensitive dye (VSD) imaging data. Recently, Mohajerani et al. (2013) used the combined local-global (CLG) algorithm (Bruhn et al., 2002; Jara-Wilde et al., 2015) to determine velocity vector fields of flow in wide-field VSD imaging data. They also manually determined the location of sources and sinks in space and time. Following Mohajerani et al., Townsend et al. (2015) also used the CLG method to estimate wave patterns in cortical activity recorded with multielectrode arrays.

Only HS, CLG, and TS optical flow methods have been used for the analysis of brain activity but there are numerous other optical flow algorithms that have been developed (Paragios et al., 2006; Sun et al., 2010; Otte and Nagel, 1994; Tavakoli et al., 2008; Liu et al., 2015; Kroeger et al., 2016). However, a quantitative and comparative evaluation of the performance and analytical efficacy of optical-flow methods for determining velocity vector fields in brain activity remains lacking. There also has been a scarcity of quantitative analysis for extracting spatiotemporal dynamics in brain activity from velocity vector fields. Off the shelf tools are also missing for streamlining data

analysis i.e. estimating velocity vector fields followed by automatically identifying sources and sinks, and calculating trajectories and velocities of brain activity with respect to time. In addition to identifying sources and sinks in space-time, it would also be useful to quantitatively characterize their properties such as outlines (what is the shape of a source or a sink), sizes (how big a source or sink is in space) and strengths (how much activity outflow or inflow is there in time).

In this paper, we present for the neuroscience community, a graphical user interface based Optical-Flow Analysis Toolbox in Matlab® (Mathworks Inc.) for investigating the spatiotemporal dynamics of mesoscale brain activity (OFAMM). We compared the performance and analytical efficacy of three optical-flow methods namely, Horn-Schunck (HS), combined local-global (CLG), and temporospatial (TS) for determining velocity vector fields of perceived flow in brain activity monitored using voltage and calcium imaging. Also, we compared the performance of HS and CLG method-based analyses in determining sources and sinks in space-time as well as trajectories of brain activity waves and their temporal speeds. These three optical flow methods were selected from numerous others (Paragios et al., 2006), primarily, due to their previous use with brain activity. Since ground truth values are unknown in real experimentally derived data sets of brain activity, simulated data was first used to investigate the accuracy of our analysis and its sensitivity to the addition of noise. Later, we tested and validated the application of our analysis on real data acquired with wide-field voltage and calcium imaging from mouse cortical activity. Similar to previous findings (Berger et al., 2007), higher instantaneous and temporal speeds were estimated with voltage imaging as compared to calcium imaging. The results are consistent with voltage signals reporting predominantly subthreshold activity and calcium signals reporting suprathreshold spiking activity (Berger et al., 2007).

#### Materials and methods

The General framework

The ultimate goal of analyzing sampled brain activity with opticalflow methods is to characterize and study the perceived motion of activity (in space-time). To do so, the set of two-dimensional (2D) images collected over time are first preprocessed to filter noise and determine the percentage change in fluorescence from a baseline ( $\Delta F$ / F<sub>o</sub>) for each pixel. Next using optical-flow methods, "velocity vector fields" are determined by estimating the displacement of pixels over time. Thus, a velocity vector is estimated for each pixel whose magnitude and direction represents its speed and direction of motion. Here, we used three different optical flow methods HS, CLG, and TS for determining the velocity vector fields (Fig. 1). With the TS method, only one vector field is estimated for a group of image frames whereas with the CLG and HS methods vector fields are obtained for all pairs of consecutive image frames. From the vector fields, the locations of sources and sinks i.e. the regions of origin and termination of activity, can be determined using vector calculus methods. Because only one vector field is estimated for a group of frames when employing the TS method, the location of sources and sinks can be estimated only in space but not in time (Takagaki et al., 2011). However, from the vector fields estimated by the CLG and HS methods the location of sources and sinks can be estimated in both space and time and more importantly, the trajectories of pixels or regions of interest and their temporal velocities can also be calculated using vector calculus methods (Fig. 1). The details of the optical-flow and vector calculus methods that we use in this work are given below.

Optical-flow estimation methods

Horn-Schunck (HS) method

The HS method (Horn and Schunck, 1981; 2016, http://

www.mathworks.com/help/vision/ref/opticalflowhs-class.html) operates on two consecutive frames and estimates the motion of pixels from one frame to the other by iteratively solving an optimization problem (stated below) formulated from two constraints. The first constraint called "brightness constancy" assumes a pixel to have the same brightness level in both frames after movement i.e.

$$I(x + u, y + v, t + dt) = I(x, y, t)$$
 (1)

where I(x,y,t) is the pixel brightness in the first frame at time t and spatial location (x,y) and I(x+u,y+v,t+dt) is the pixel brightness in the second frame at t+1 after (u,v) displacements from (x,y) in x and y directions respectively. The second constraint called "spatial smoothness" prevents discontinuities in the flow field. Mathematically,

$$|\nabla u|^2 = \left(\frac{\partial u}{\partial x}\right)^2 + \left(\frac{\partial u}{\partial y}\right)^2 \text{ and, } |\nabla v|^2 = \left(\frac{\partial v}{\partial x}\right)^2 + \left(\frac{\partial v}{\partial y}\right)^2$$
 (2)

These two constraints are combined to get a minimization problem given by the following equation

$$\min_{u, v} \left\{ \int (I_t + I_x u + I_y v)^2 dx dy + \alpha \int (|\nabla u|^2 + |\nabla v|^2) dx dy \right\}$$
(3)

where  $I_t$ ,  $I_x$ , and  $I_y$  are derivatives of I(x,y,t) with respect to time, spatial direction x, and spatial direction y, respectively (Akemann et al., 2012) and  $\alpha$  is the ratio of the weights of spatial smoothness integral to that of the brightness constancy. This equation is solved numerically to estimate the values of u and v. As the integral is calculated on the whole image field, HS method is termed a global method. A numerical implementation of the HS method in Matlab\* was downloaded from

(https://www.mathworks.com/matlabcentral/fileexchange/22756horn-schunck-optical-flow-method) and used here for analysis. Briefly, in the numerical implementation, image derivatives are calculated first with respect to x, y, and t using convolution filter (see computeDerivatives.m file in the toolbox). Then, the vector field is calculated iteratively solving the numerical version of Eq. 3 (see HS.m file in the toolbox). The parameters  $\alpha$ , and the number of iterations can be set in the graphical-user-interface of the toolbox. The larger the number of iterations, the better will be the solution convergence but at the cost of computational time. For increasing values of  $\alpha$ , the spatial smoothness constraint will have a larger contribution in the minimization problem and hence the estimated velocity vector field will be smoother. The initial values of u and v for all pixels in images were set to 0. The reader is also referred to the "opticalFlowHS.m" function in Matlab® 2015a (or later) and related help on Mathworks Inc., website for more details of numerical implementation of the HS method (2016,http://www.mathworks.com/help/vision/ref/ opticalflowhs-class.html). Users of Matlab® 2015a or later are encouraged to use the built-in function for computing optical flow with the HS method. We however do not expect to see significant difference between computational times and estimated optical flow with the two implementations as the underlying numerical computations are quite similar.

We chose  $\alpha=0.1$  and the number of iterations = 2000 for our subsequent analysis by trial and error. We estimated velocities of pixels with the HS method for the travelling plane and circular waves (simulated data) while tweaking parameter values to minimize deviation from actual velocities. Alternately,  $\alpha$  and the number of iterations were also determined analytically by using simulated data with known values of trajectories and velocities for the three Gaussian waves (see Supplementary Movie 1- last clip). Different combinations of the values of  $\alpha$  (ranging between 0.05 and 100), and the number of iterations (ranging between 100 and 9000) were used to estimate the best combination that gave the minimum error for estimated velocities. The best combination of  $\alpha=0.35$  and number of iterations = 2000 was used for analysis for generating supplementary Figs. S3-S7.

Supplementary material related to this article can be found online

at http://dx.doi.org/10.1016/j.neuroimage.2017.03.034.

#### Combined local-global (CLG) method

The CLG method (Bruhn et al., 2002; Jara-Wilde et al., 2015) combines the HS method and a second method called Lucas-Kanade (LK). The LK method assumes the constancy of motion in the neighborhood of a pixel i.e. the velocity and direction of motion of pixels around the pixel of interest are equal. Mathematically, LK method formulates the following minimization problem

$$\min_{u, v} \left\{ \sum_{p} (I_{t}(p) + I_{x}(p)u + I_{y}(p)v)^{2} \right\}$$
(4)

where p is selected from an  $n \times n$  neighborhood of (x, y) pixel. In a more general formulation of the LK method the optimization argument can be convolved with a Gaussian window to filter noise in the images is

$$\min_{u, v} \left\{ \sum_{p} G_{\sigma}^{*}(I_{l}(p) + I_{x}(p)u + I_{y}(p)v)^{2} \right\}$$
 (5)

where  $G_{\sigma}$  is a two dimensional normalized Gaussian window with standard deviation of  $\sigma$  and \* denote the convolution operation. As the LK method assumes that the motion (speed and direction) is constant in the neighborhood of a pixel, it is called a local method and works better while dealing with small displacements.

The combination of HS and LK methods proposed by Bruhn et al. Bruhn et al. (2002) constitutes the CLG method and is formulated as below:

$$\min_{u, v} \left\{ \sum_{p} G_{\sigma}^{*}(I_{l}(p) + I_{x}(p)u + I_{y}(p)v)^{2} + \alpha \int (|\nabla u|^{2} + |\nabla v|^{2}) dx dy \right\}$$
(6)

The value of  $\alpha$  here determines the contribution of the second term related to spatial smoothness constraint to the overall minimization problem. Similar to the HS method, for increasing values of  $\alpha$ , the estimated velocity vector field will be smoother. We used a numerical implementation of the CLG method in Matlab (Liu, 2009) with additional parameters that need to be set. The implementation uses a hierarchical method which finds solutions gradually from coarse to refined levels. Coarse solutions are found from spatially down sampled image sequences. The down sampling factors for finding coarse solutions are multiples of a parameter termed "ratio" and the minimum accepted size of a down sampled image is specified by the 'minWidth' parameter. The number of hierarchical levels depend upon the size of images, and the parameters, ratio and minWidth. For an image size of 128x128 pixels, ratio = 0.5, and minWidth = 32, final solution is found gradually by finding solutions for image sizes of 32×32, 64×64, and 128×128 pixels. At each level of hierarchy, the vector field is estimated using results from previous level as an initial value. Other parameters in the numerical implementation of the CLG method include number of iterations for different hierarchical levels. The larger the number of iterations, the better will be convergence of the numerical solution to the actual value but at the cost of computational time. The values of parameters can be set via the graphical-user interface of the toolbox. The reader is referred to Liu C. 2009 for further details of the numerical implementation of the CLG method (Liu, 2009).

We chose the values indicated below following (Liu, 2009), and trial and error in order to minimize the error in the estimation of velocities for the travelling plane and circular waves (simulated data).

 $\alpha$ =0.03, ratio=0.5, minWidth = ImageSize × ratio × 0.5, nOuterFPIterations=7, nInnerFPIterations=1, nSORIterations=30

Alternately, the following values were used for generating supplementary Figs. S3-S7.

 $\alpha$ =0.04, ratio=0.5, minWidth = ImageSize × ratio × 0.5, nOuterFPIterations=15, nInnerFPIterations=3, nSORIterations=30

These values were obtained analytically by using trial and error application of CLG method onto simulated data same as that used for the HS method (see Supplementary Movie 1 – last clip). First, a large number of iterations was used and the values of  $\alpha$  and ratio were determined which gave the minimum error in the estimated velocities. Later, the number of iterations were reduced to a desired number which kept the same error to a minimum.

#### Temporospatial (TS) method

The TS method proposed and demonstrated by Takagaki et al. Takagaki et al. (2011) estimates motion of pixels by identifying neighboring pixels whose temporal activity is highly correlated and hence was named Temporospatial method. To determine the speed and direction of motion of a pixel, the correlation of its activity with that of neighboring pixels is calculated to determine the highest correlation coefficient and thus estimate the next location of the pixel. The speed of motion is then determined by dividing the displacement of pixel from one location to the other by the best temporal lag (delay) that provides the highest correlation coefficient. Finding temporal correlation of all pairs of pixels is a computationally intensive process. To reduce computational time, Takagaki et al. defined spatial motion templates of pixels which represented different types of motion (e.g. expansion, contraction, translation, and rotation) as bases along which the motion of pixels could be resolved. The motion of a pixel was then estimated by finding the components of its motion along the bases (see Takagaki et al. 2011 for details). Since the whole temporal activity of a pixel in a group of frames is used to determine temporal correlations with neighboring pixels, only one vector is obtained for a pixel representing the average movement of that pixel in time. Thus for a group of image frames only one vector field is obtained. The implementation of TS method we used in this work was written in Matlab® by the first author based on Takagaki et. al (Takagaki et al., 2011). In this implementation, four parameters can be set via the graphical-user interface of the toolbox. The parameter "target frame" specifies the frame of interest for which we seek to estimate the average velocity vector field. "Corr win frames" is the number of frames around the target frame for calculating the correlation between time series of intensity values for pixels. Increasing the value of this parameter will increase the accuracy of finding correlation. "Max Lag" is the maximum allowed correlation lags (both positive and negative) used for correlation calculations. This parameter controls the permissible duration between events happening on pixels whose correlation is being calculated. Finally, "Corr win pixels" determine the number of pixels forming a circular window around a pixel of interest whose motion we estimate by determining correlations between time series of intensity values of pixels. Larger values of this parameter would allow to estimate higher speeds but at the cost of computational time.

#### Vector calculus methods

Determination of the locations of sources and sinks from velocity vector fields

The locations of sources and sinks (points of outward or inward flow respectively) from velocity vector fields were calculated using three methods. In the first method, divergence of the vector field was calculated to obtain a scalar field with positive and negative numbers indicating locations of sources and sinks respectively while their magnitude indicates the strength. Contours of constant divergence values were also calculated. In the second method, Poincare index was calculated for a pixel by finding the sum of differences of angles of velocity vectors of neighboring pixels. Poincare index when normalized by  $2\pi$ , has values of +1, 0, and -1 for a source or a sink, plane flow, and saddle points, respectively. In the third method, trace and determinant

values of the Jacobian matrix of the flow were calculated for each pixel. A pixel is a source if the determinant and trace values are positive and is a sink if the determinant is positive with a negative trace value. Following Townsend et. al., a pixel was only considered a source (or a sink) if all the three methods identified it as a source (or a sink) and it had at least two closed contours surrounding it (Townsend et al., 2015). The contour maps were calculated from the divergence and the contour levels were automatically selected by Matlab® in a range from minimum to maximum values of input divergence matrix. This range is automatically divided into 10 contour levels. A point was detected as a source if it was surrounded by at least two positive level closed contours and a sink if it was surrounded by at least two negative level closed contours. All three methods were implemented in Matlab<sup>®</sup>. The size and strength of a source or a sink was determined from the closest (innermost) contour. The size was defined as the number of pixels inside the contour while the strength was defined as the contour value. The contour borders defined the shape of the source or sink.

#### Determination of trajectories of pixels and their temporal speeds

In order to determine trajectories of pixels, streamlines were estimated from the velocity vector fields using the built-in "stream3" function of Matlab. A streamline represents the curve to which a subset of velocity vectors in the field are tangent and thus it represents path of the flow of a pixel. The stream3 function calculates a temporal streamline from the velocity vector fields for a pixel of interest defined in space-time. The temporal velocity of the pixel along its trajectory was then calculated from the streamline.

#### Simulated data and addition of noise

Simulated data were generated in Matlab\* with known shapes of sources and sinks, and motion profiles (velocities and directions of movement) to compare the analytical efficacy of TS, CLG, and HS methods in determining velocity vector fields and source/sink locations, trajectories of pixels and their temporal speeds. Three image sequences (explained in the results section) were generated; Traveling half-sinusoid plane wave, traveling half-sinusoid circular wave originating at the center (single source), and complex multisource/multisink traveling Gaussian waves. To add noise to the simulated image sequences, the average power of all pixels (over space and time) was calculated. The amplitude of Gaussian white noise added to each pixel was proportional to the average power. The code for generating simulated data and the addition of noise is provided with the toolbox.

# Optical imaging - Data acquisition and analysis

#### Animals and surgery

Two adult (~25 g) male C57BL/6J mice were used for voltage sensitive dye imaging experiments and one "Emx-GCaMP6f" animal was used for calcium imaging. The "Emx-GCaMP6" mouse was produced by crossing homozygous B6.129S2-Emx1 <sup>tm1(cre)Krj</sup>/J strain (Jax no. 005628) with B6;129S-Gt(ROSA)26Sor tm95.1(CAG-GCaMP6f)Hze/ J strain (Jax no. 024105, Ai95) (Madisen et al., 2015). The presence of GCaMP6f expression was determined by genotyping each animal before each surgical procedure. Mice were housed in clear plastic cages in groups of two to five under a 12 h light, 12 h dark cycle. Mice were given ad libitum access to water and standard laboratory mouse diet at all times. The animal protocols were approved by the University of Lethbridge Animal Care Committee and were in accordance with guidelines set forth by the Canadian Council for Animal Care. At approximately 3 months of age, mice were given an acute craniotomy. Mice were anesthetized with isoflurane (1.0-1.5%) for induction and during surgery, and a reduced maintenance concentration of isoflurane (0.5%) was used later during data collection. Mice were placed on a metal plate that could be mounted onto the stage of the upright macroscope, and the skull was fastened to a steel plate. A 7 × 6 mm

unilateral craniotomy (bregma 2.5 to -4.5 mm, lateral 0 to 6 mm) was made, and the underlying dura was removed, as described previously (Mohajerani et al., 2013; Kyweriga and Mohajerani, 2016). Throughout surgery and imaging, body temperature was maintained at 37 °C using a heating pad with a feedback thermistor.

#### Optical imaging

For voltage-sensitive dye based wide-field optical imaging (VSDI), the dye RH-1691 (Optical Imaging, New York, NY) (Shoham et al., 1999) was dissolved in 4-(2-hydroxyethyl)-1-piperazineethanesulfonic acid (HEPES)-buffered saline solution (0.4 mg ml<sup>-1</sup>) and applied to the exposed cortex for 30-60 min, as previously described (Mohajerani et al., 2010; Mohajerani et al., 2011; Lim et al., 2012; Lim et al., 2013). After staining, rest of the dye was washed out and the cortical surface was covered with 1.5% agarose in HEPES-buffered saline and a coverslip on top to reduce movement artifacts. The dye was excited by a red LED (Luxeon K2, 627 nm center) and the excitation and emission optical filters for imaging were 630  $\pm$  15 nm and 688  $\pm$ 15 nm (Semrock, New York, NY), respectively. For calcium imaging, mice were anesthetized with isofluorane and their skull was exposed (left hemisphere). Thinning of the skull bone was then performed followed by affixing of a glass coverslip with cyanoacrylate glue and implantation of a metal head plate. The fluorophore GCaMP6f was excited with a blue LED (Luxeon, 470 nm center) filtered by an excitation filter 470 ± 40 nm (Semrock, New York, NY). The emitted light was filtered by a 542 ± 27 nm emission filter (Semrock, New York, NY). 12-bit CCD camera (1M60 Pantera, Dalsa, Waterloo, ON) was used to acquire images at a rate of 150 and 30 frames per second for VSD and calcium imaging. Because animal brain states show spontaneous change, we averaged 10-45 trials of stimulus presentation to reduce these effects. To correct for time-dependent changes in fluorescent signals that accompany all imaging, we also collected a number of nonstimulation trials that were used for normalization of the stimulated data. A 10-s interval between each sensory stimulation was used. The camera was focused 0.5-1 mm below cortical surface to avoid distortion of signal due to movement of superficial blood vessels. The preprocessed data is provided with the toolbox data residing in subfolder "SampleData" in the main toolbox folder.

# Sensory stimulation

Sensory forelimb stimulation was done by passing a current (0.2–1 mA for 1 ms) through a thin needle (0.14 mm) inserted into the forepaw. For auditory stimulation, a 12 kHz pure tone at 80 dB was played using a Tucker-Davis Technologies (TDT) RX6 and delivered to animal's right ear while the animal was sitting in a sound proof booth in the laboratory. The speaker (TDT, ES1 electrostatic loudspeaker) was calibrated to emit uniformly distributed amplitude of all frequencies.

## Data preprocessing

To reduce regional bias in caused by uneven dye staining or brain curvature, the VSD and calcium responses were expressed as a percentage change relative to baseline ( $\Delta F/F_0 \times 100\%$ ) using Matlab\*. The baseline ( $F_0$ ) consisted of intensity values obtained from nostimulation trials. Individual  $\Delta F/F_0$  image sequences were then filtered by spatial Gaussian filter with sigma equal to 67  $\mu$ m. To reduce the effect of heartbeat in our calcium imaging data, the calcium signal was temporally filtered with a finite-impulse-response low-pass filter (cutoff frequency 5 Hz) before spatial filtering. For all real data, we determined signal-to-noise ratios (SNRs) before and after filtering (see Supplementary Material). For VSDI, SNR was ~5 (7 dB) and ~42 (16 dB) before and after filtering respectively, while for CaI, these values were ~13 (11 dB) and ~31 (15 dB).

#### Statistical analysis

All statistical analysis was done in Matlab®. Two sample t-test was

used for statistical comparison and a p-value of 0.05 was used to determine significance.

#### Results

We first compared the performance of the three optical-flow (OF) methods using simulated data for which the ground truth values of wave motion are known. Later, we demonstrated the use of OF methods on real experimentally derived voltage and calcium optical imaging data.

Optical-flow characterization of simulated data

Travelling plane wave (no source and sink): a half-sinusoid moving in one direction

One of the most commonly observed wave pattern in brain activity is a travelling plane wave (Townsend et al., 2015; Takagaki et al., 2011; Mohajerani et al., 2013; Sato et al., 2012; Ermentrout and Kleinfeld, 2001; Muller and Destexhe, 2012; Frostig et al., 2008) which is

encountered when activity has emerged from a source and is travelling towards a sink. It would also be seen in situations where the imaging field of view is limited. We mimicked such activity by a half-sinusoid travelling plane wave at a constant speed (1 pixel per frame - p/f) in the horizontal direction (angle = 0, see Fig. 2A and Supplementary Movie 1).

Determination of velocity vector fields. With the three algorithms (TS, CLG, and HS), optical-flow of the travelling half-sinusoid was determined i.e. velocity vector fields were estimated where velocity vectors obtained for all pixels represent their speeds and directions of motion (see Fig. 2A). Note that with the TS method only one vector field for the whole image sequence representing average speeds and directions is obtained. With the CLG and HS methods however, vector fields are estimated from all pairs of consecutive frames and thus their number is one less than the number of frames in the image sequence. The velocity vectors for the CLG and HS methods thus represent instantaneous speeds and directions. The estimated vector fields determined from all three methods matched very closely with the

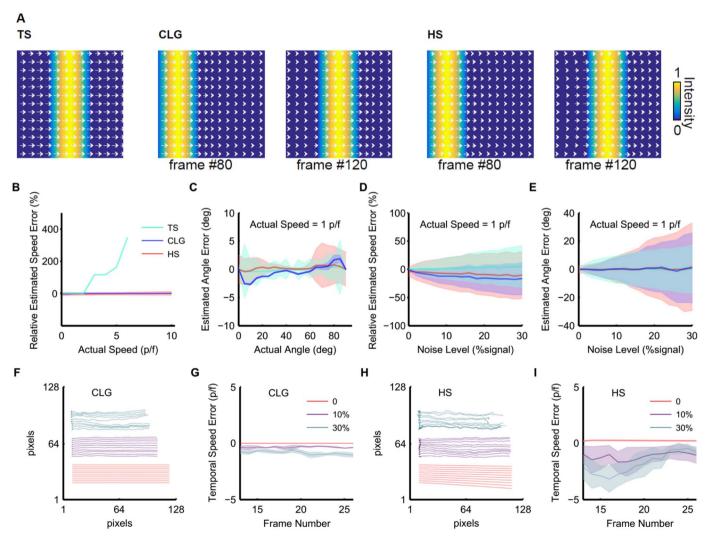


Fig. 2. Comparison of the performance of HS, CLG, and TS methods for estimating dynamics of a simulated traveling plane wave. **A.** Representative frames from the image sequence with overlaid velocity vector fields. Note that with the TS method only one vector field is obtained for the whole image sequence while with the HS and CLG methods, a vector field is obtained for each pair of consecutive frames (see text) **B.** Instantaneous speed estimation error  $(E_{is})$  versus actual speeds of plane wave for angle = 0°. Performance of the TS method depends on the size of correlation window in space (see text) and shows high error for speeds where displacements of pixels are beyond the size of the correlation window. HS shows an increase in the standard deviation of the error with increasing speeds but CLG is robust. **C.** Instantaneous angle estimation error  $(E_{ia})$  versus actual angles (direction of motion) of plane wave with speed = 1 p/f. TS method shows large standard deviation of error. HS and CLG methods show similar performance with the CLG method showing smallest standard deviation. **D, E.** Instantaneous speed (D) and angle (E) estimation error versus noise level. The performance for three methods degrades with increasing noise levels. **F-I.** Trajectories of pixels for 0% (red), 10% (purple) and 30% (green) noise level shown in single frame (F) for CLG, and (H) for HS methods. Temporal speed estimation error  $(E_{ts})$  versus time (frame number) for the three noise levels (G) for CLG, and (I) for HS methods. Overall the best performance is that of CLG.

actual flow (Fig. 2A). Moreover, flow fields calculated with the CLG and HS methods were similar to each other but HS underestimated the speed of flow at the top and bottom boundaries. The time required for the calculation of velocity vector fields was longest for the TS method (~79 s for estimating 1 velocity vector field from 242 frames each having 128×128 pixels). CLG and HS methods required 0.52 s and 0.59 s respectively for generating 1 velocity vector field from two consecutive frames. These times were obtained on a 64-bit PC with the following specifications; Processor Intel ® Core™ i7-4770K CPU @ 3.50 GHz, RAM 32 GB, and with our toolbox running in the Matlab® R2013b environment.

Performance of HS, CLG, and TS methods for determining velocity vector fields. In order to gauge the efficacy of the three methods in determining velocity vector fields (speeds and angles), we compared estimated and actual instantaneous velocity vectors by determining the errors for a range of values of speeds and angles (directions of motion) of the sinusoids:

$$E_{is} = \frac{Estimatedspeed(S_e) - Actualspeed(S_a)}{Actualspeed}$$

and

 $E_{ia} = Estimatedangle(A_e) - Actualangle(A_a)$ 

First, Eis was measured for all pixels from one velocity vector field (frame) and the trend of means and standard deviations of Eis was calculated for values of actual speeds ranging between 0 and 10 pixels/ frame - p/f (Fig. 2B) while keeping the angle = 0 (horizontal direction of motion). The TS method performed well for actual speeds less than 2p/f but its performance steeply declined for greater speeds because of limited size of the time window used for determining temporal correlations (see methods). The TS method was therefore sensitive to the size of window chosen for analysis. A larger window would produce better estimates for higher actual speeds but with an added computational cost (Supplementary Fig. S1B). The performance of CLG and HS methods was better than TS for higher speeds with CLG performing better than HS (small standard deviation for CLG in Fig. 2B). Similarly, Eia was determined for all pixels and the trend of mean and standard deviations of Eia was observed for values of angles ranging between 0 and 90° (Fig. 2C) while keeping the speed constant = 1p/f. Eia remained within ±5° for all the three methods with individual variations in the values of means and standard deviations. Although the mean value of Eia calculated with the TS method is closer to zero

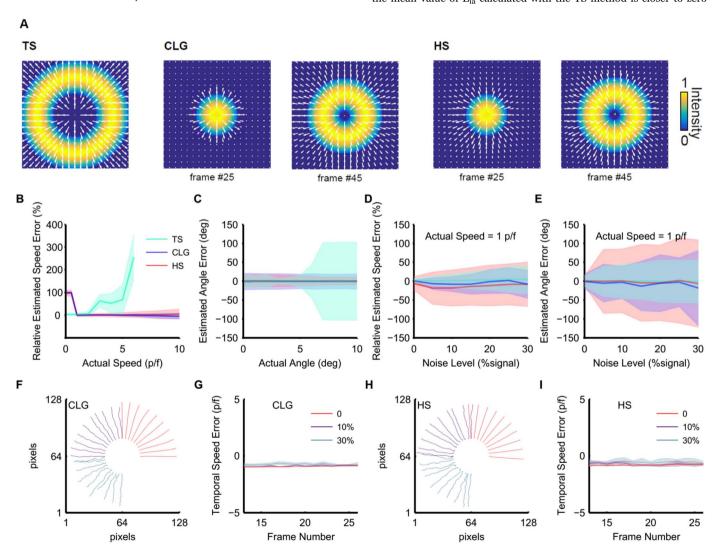


Fig. 3. Comparison of performance of HS, CLG, and TS methods for estimating dynamics of a simulated traveling circular wave. A. Representative frames from the image sequence with overlaid velocity vector fields. B, C. Instantaneous speed estimation error  $(E_{is})$  (B) and instantaneous angle estimation error  $(E_{ia})$  (C) versus actual speeds. D, E. Instantaneous speed estimation error (D) and instantaneous angle estimation error (E) versus noise levels. F-I. Trajectories of pixels for 0% (red), 10% (purple) and 30% (green) noise level shown in single frame (F) for CLG, and (H) for HS methods. Temporal speed estimation error  $(E_{ts})$  versus time (frame number) for the three noise levels (G) for CLG, and (I) for HS methods. Overall the best performance is that of CLG.

than the means calculated using the HS and CLG methods, the standard deviation of  $E_{ia}$  calculated with the TS method was much higher. The performance of CLG and HS in estimating the angles was similar with a very small standard deviation of error with the CLG method. Based on these results, the CLG method should be preferred over TS and HS methods for determining velocity vector fields for travelling plane waves.

Efficacy of HS, CLG, and TS methods in determining vector fields from noisy data. Since experimental derived data representing brain activity will inherently contain noise, we tested the performance of the three optical-flow methods by adding noise to the simulated data (see methods). By adding various levels of noise to each pixel in the image sequence, the errors Eis and Eia in speed and angle estimation respectively were calculated in the same way as described previously. The addition of noise adversely affected the estimates of speeds and angles with larger errors in estimation for larger noise levels (Figs. 2D and 2E). The estimate of angles however, were more sensitive to greater noise (E<sub>ia</sub> ~30° for 30% noise level). From all the three methods tested, the temporospatial method was most robust in the estimation of angle and velocity from noisy data. This robustness resulted from improved signal to noise ratio that arises from the temporal correlation between pixels. The performance of CLG and HS methods was similar to each other for all noise levels with CLG having smaller standard deviations of errors. Our results thus suggest that filtering imaging data to remove noise is preferable before determining vector fields.

Determination of trajectories of pixels and their speeds versus time. To determine underlying anatomical and physiological communication pathways between different brain regions, it is imperative to be able to follow the movement pathway of activity. Hence, using vector calculus methods, the trajectories of travel (streamlines) of pixels or regions of interest were determined from velocity vector fields calculated with CLG and HS methods. Note that this analysis cannot be done with the single velocity vector field obtained with the TS method. Estimation of trajectories from velocity vector fields determined with the CLG method were better than that of the HS method (compare red lines in Figs. 2F and 2H). However, when noise (10% and 30%) was added to the simulated data, the performance of estimation declined similarly for both methods. Temporal speeds (St) of pixels along the trajectory were also estimated and the error ( $E_{ts}$  = estimated  $S_t$  - actual  $S_t$ ) was calculated. The error was zero for simulated data with no noise but increased for 10% and 30% noise levels (purple and green lines in Figs. 2G and 2I). The error was worse for analysis based on the HS method (Fig. 2I). Note that the displacement (direct distance from start to finish) of a pixel was smaller at higher noise levels. This is mainly because of error in the estimate of angles resulting in zig-zag shaped trajectories of pixels. The zig-zag effect grows larger with increasing noise levels. Our results suggest that the trajectories of pixels and their speeds are better estimated from velocity vector fields determined with the CLG method after noise reduction.

Travelling circular wave: a half-sinusoid originating at a source and propagating in all directions. Another important pattern of activity found in imaging data is that of the propagation of a signal originating from a source such as those seen after peripheral sensory stimulation (Mohajerani et al., 2013; Frostig et al., 2008; Ferezou et al., 2007; Polack and Contreras, 2012). In order to mimic this type of activity, a circular ring of half-sinusoid was generated which originated at the center of a frame and spread outwards to the edges with a constant velocity of 1 p/f (Supplementary Movie 1). The optical-flow was estimated with the three methods to obtain velocity vector fields

(Fig. 3A). Errors in the estimates of instantaneous speeds and angles ( $E_{\rm is}$  and  $E_{\rm ia}$ ) were determined for propagation speeds ranging between 0 and 10 p/f (Fig. 3B-C). Note that in the simulated data all directions of motion i.e. all angles are concurrent and hence only the propagation speed was changed. Similar to the results above, performance of the TS method degraded for estimation of higher speeds (Fig. 3B) because of the limited size of window used for finding temporal correlations between pixels (see methods). The TS method also failed to estimate the directions of motion for higher speeds for the same reason (Fig. 3C). The performance of the CLG and HS methods was similar to each other with CLG's performance better than that of HS for estimating speeds but not angles (smaller standard deviations of errors for CLG speed curve and larger for CLG angle curve).

The efficacy of the three methods for determining velocity vector fields from noisy data was assessed by adding different noise levels to all pixels and observing the errors in estimates of instantaneous speeds and angles. The propagation speed was set to 1p/f for this analysis. The TS method was most robust to noise as computing temporal correlations averages out some noise (Fig. 3D-E). However, all three methods were better in estimating speeds rather than angles from noisy data. With the addition of even small noise levels, the performance of the HS method in estimating angles degraded quickly. These results suggest that it is preferable to filter raw data before applying any of the three methods for determining velocity vector fields.

Next, trajectories of selected pixels and their velocities were calculated from velocity vector fields that were estimated with the CLG and HS methods with and without adding noise to all pixels. Without the addition of noise, the estimated trajectories were accurately determined (red lines in Fig. 3F and H). However, with the addition of 10% noise level, the estimated trajectory is such that the total displacement of pixels is shorter than the actual (purple lines). This is because the estimation of angles of velocity vector fields was adversely affected with the addition of noise and thus the estimated path is zig-zag. The zig-zag effect becomes more pronounced with the addition of 30% noise level (green lines). The effect of noise on the estimation of temporal speeds was negligible (Fig. 3G and I) with the CLG method based analysis showing smaller standard deviations of error. Combined these results also favor the use of CLG method for determining velocity vector fields.

Travelling Gaussian waves: Complex simulated data containing three Gaussian sources, travelling waves, and sinks

When multiple brain regions are active simultaneously (Huang et al., 2010; Mohajerani et al., 2013; Mohajerani et al., 2010; Ferezou et al., 2007), multiple travelling waves are observed (see Fig. 1b in (Mohajerani et al., 2013)). In order to gauge the effectiveness of our analyses in estimating the spatiotemporal dynamics of such activity, we generated simulated data that incorporated these characteristics by forming an image sequence in which three two-dimensional Gaussian sources (with same amplitude = 1) originated at three distinct spacetime locations i.e. the spatial points of origin are distinct as well as the frames in which they start (Supplementary Movie 1). Based on their shapes, these activity-like events were named C1 (circular), E1 (elliptical), and C2 (circular). The temporal order of their appearance was C1, E1, and C2. All events consisted of an initiation and expansion to reach their final size and when fully expanded, these events underwent translation and/or rotation. The trajectories of C1 and E1 formed arcs of different circles while the trajectory of C2 was linear. E1 while translating also rotated 30° counter-clockwise about its center (like a spiral wave). Finally, all events contracted and sunk into three distinct space locations. For each event, the full expansion and contraction occurred within 15 frames (~0.1 secs). For C1 and E1 during translations while following arcs of circles, the variation in angular speed followed a half-sinusoid while for C2, the linear translation

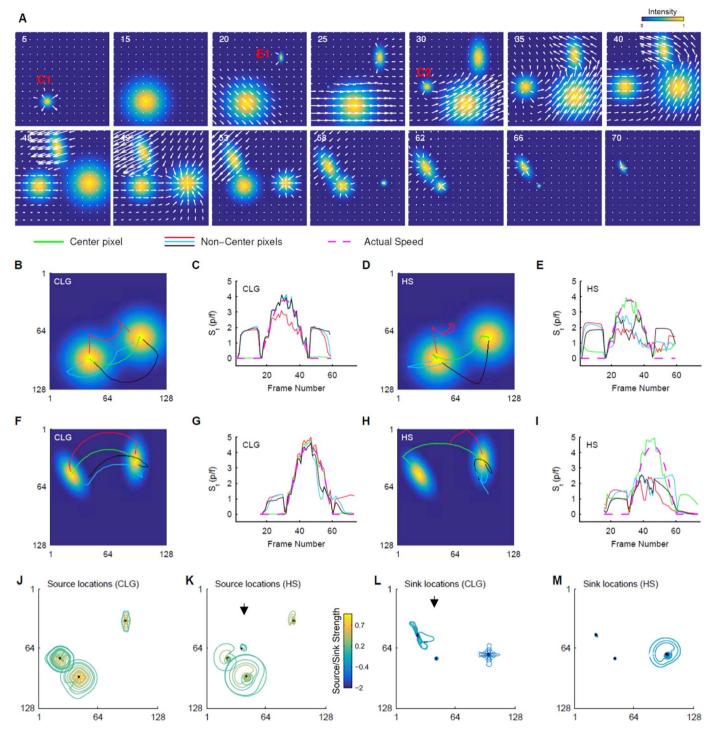


Fig. 4. Comparison of the performance of HS and CLG methods for estimating dynamics of an image sequence simulating three sources, traveling events, and sinks (see text). A. Montage of frames from the image sequence overlaid with velocity vector fields determined with the CLG method. B. Trajectories of selected pixels of C1 event (see text), center pixel (green), non-center pixels (blue, black, and red), calculated with the CLG method based analysis. C. Estimated temporal speeds of pixels in (B) - same colors, versus time (frame numbers). Magenta line shows actual speed of the center pixel. D-E. Same as B-C but with the HS method based analysis. F-G, and H-I Same as B-C and D-E respectively for E1 event. J-K. Estimation of source locations (black dots) and shapes/outlines (contours) with the CLG and HS methods. L-M. Estimation of sink locations with the CLG and HS methods. Contour colors show source/sink strengths. The CLG method wins in estimating pixel trajectories and temporal speeds and works favorably for source/sink analyses.

occurred at constant speed. By generating this more complex multi-site simulated activity data, our goal was to determine how accurately our analysis would estimate the underlying dynamics and to compare the performance of CLG and HS methods. The TS method was not used here because it requires large computational time and provides only a single vector field for the whole image sequence.

Velocity vector fields were first estimated (Fig. 4A for the CLG method) and later using methods of vector calculus, the trajectories of

pixels of interest and their speeds were calculated (Fig. 4B-I for C1 and E1). The analysis based on the CLG method accurately captured the dynamics of C1 and E1 (with a few exceptions for E1 – see below). However, the analysis based on the HS method however was not as efficient. For example, the center of C1 was static while the Gaussian is expanding, follows a half sinusoid speed curve (i.e. the speed changes sinusoidally) while translating along the arc, and was static again while the Gaussian is contracting (see dashed magenta line for actual speed

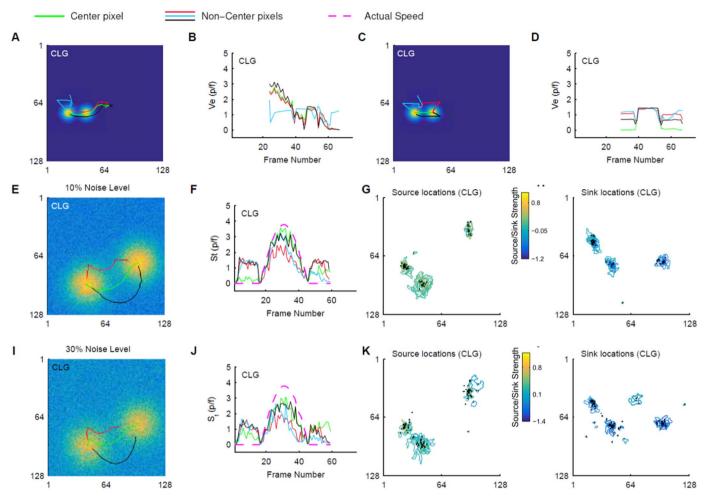


Fig. 5. Limitations of the CLG method based analysis – errors due to the addition of noise and/or overlap of simulated events in space-time. A. Erroneous estimation of trajectories of selected pixels of C2 starting frame 24 because at this time, C2 is in the wake of C1 (see text). Center pixel (green), non-center pixels (blue, black, and red). B. Erroneous estimated temporal speeds of pixels in (A) - same colors, versus time (frame numbers). C-D. Same as B-C but correct estimation of trajectories and temporal speeds with analysis starting frame 29. E-H, Same as Fig. 4 B-C, J, & L but with 10% noise level added to the image sequence. I-L. Same as E-H with 30% noise level added to the image sequence. The addition of noise reduces the performance of the CLG method. Caution is advised for selecting pixels and regions of interest in space-time for determining trajectories and temporal speeds.

of center in Fig. 4C). The estimated speed of the center of C1 using the CLG method closely matched the actual speed (see green lines in Figs. 4B and 4C for trajectory and speed respectively). However, the HS method based analysis failed to capture the dynamics accurately during the expansion and contraction phases (green lines in Fig. 4D-E). The actual speed of non-center points of C1 was ~2p/f when C1 was expanding. After expansion, their speeds increased sinusoidally to a peak value of ~4p/f, and then decreased sinusoidally to 0 p/f. During the contraction phase of C1, their speeds were ~2p/f. The estimated speeds of these non-center points also closely matched the actual speeds (see blue, black, and red lines in Fig. 4B-C for CLG method based analysis). The HS method based analysis failed to accurately capture the dynamics of non-center points (Fig. 4D-E). Note, that there was a momentary pause of the C1 Gaussian for two frames between expansion and translation, and for two frames between translation and contraction. This feature was accurately captured in the estimates of speeds of non-center pixels of C1 especially with the CLG method based analysis.

In addition, for the E1 event, the estimated trajectory and speed of the center pixel (green line in Fig. 4F-G) closely matched the actual trajectory and speed (dotted magenta line) respectively with the CLG method based analysis. However, the estimated trajectories and speeds were slightly erroneous for non-center pixels (see black, blue, and red lines in Fig. 4F-G). This error could in part be due to discrete spatial sampling and rotation of the E1 event during translation. Here also, the

analysis captured the momentary pause before translation along the arc. With the HS method based analysis, the estimation of event dynamics (trajectories and speeds) was not as accurate (Fig. 4H-I). Furthermore, we generated simulated data with an image sequence of a rotating pinwheel-like sinusoidal wave (see Supplementary Movie 1) mimicking the rotation of a spiral wave (Takagaki et al., 2011). We estimated its optical flow with the CLG and HS methods and observed that CLG performed better (Fig. S8). These results thus demonstrate the superiority of the CLG method based analysis over that of HS in tracking pixel trajectories and determining temporal speeds.

Since simulated data had sources and sinks associated with three simulated activity patterns, we applied vector calculus methods on each velocity vector field determined by the CLG and HS methods to identify the location, outlines, and strengths of sources and sinks in space-time (see methods). Both CLG and HS method based analyses performed well in identifying the location of sources and sinks (see black dots in Fig. 4J-M). Although there were only three events (C1, E1, and C2), they were detected as sources in different frames (times) while expanding. Similarly, these events were detected as sinks in different frames while contracting. Most of the identified source and sink locations except a few, were very close (within a few pixels) to the actual locations. A weak source (relatively smaller contour value) identified with the HS method based analysis and a weak sink identified with the CLG method based analysis were a little farther from the actual locations (arrows in Fig. 4K and L). These fake sources

and sinks are perceived in regions where events are spatially close and move away or towards each other respectively. Hence, caution is advised when interpreting whether a detected location is a real source or a sink when multiple events are present and approach or move away from each other in experimental data.

The characteristics of sources and sinks i.e. their outlines (shapes), sizes, and strengths were also estimated by finding the contours from the divergence of vector fields (see methods). The size of a source (or a sink) was defined as the area of the contour detected around it while the contour's value (divergence) was its strength. Both CLG and HS methods based analyses identified the outlines (contours) of sources and sinks with the former performing better qualitatively (see contours in Fig. 4J-M). With both types of analyses, the stronger sources (yellower contours in Fig. 4J-K) and sinks (bluer contours in Fig. 4L-M) were correctly detected close to the center location. Additionally, larger sizes were detected gradually with increasing time (frame numbers) as C1, E1, and C2 expanded. We also observed the product of size and strength (strength weighed size or vice versa) as a useful parameter to determine the significance of a source or sink i.e. larger

product would indicate a significant source or sink activity (data not shown). The quantification of parameters for sources and sinks discussed here will be useful for pin-pointing the origin and termination of activity spread in real image sequences as discussed below in Section *Wide-field optical imaging of mouse brain with forelimb sensory stimulation*.

Limitations of the CLG method based analysis in estimating trajectories and speeds of pixels; a word of caution. The CLG method based analysis failed to accurately estimate the dynamics of pixels which were shared by multiple dynamic events in space-time. For example, the C2 activity starts and ends in frames 24 and 68 of the image sequence respectively (see Supplementary Movie 1). The estimated trajectories and speeds of all C2 pixels between these frames were erroneous (representative pixels in Fig. 5A-B). The actual speeds of all C2 pixels except the center were ~1.43p/f during expansion and contraction with a momentary pause and linear translation at 1.3p/f in between. The center pixel only moves during translation. The estimated trajectories and speeds however, were

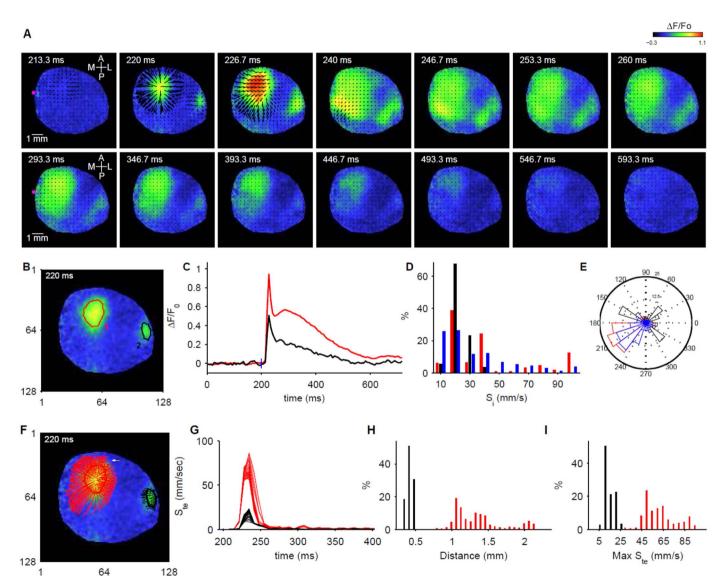


Fig. 6. Optical flow characterization (with the CLG method) of forelimb stimulation-evoked VSD activation in anesthetized mouse. **A.** Montage of selected frames from the image sequence with overlaid velocity vector fields. Magenta dot in the first (top left) frame indicates bregma location. **B.** Regions of interest (ROIs) on the primary (red) and secondary (black) forelimb areas. **C.**  $\Delta F/F_0$  vs time for ROIs in (B). **D.** Distribution of instantaneous speeds taken across all image frames of pixels in the ROIs in B (same colors) and for pixels within the mask (blue bars). **E.** Similar to (D) but distribution of instantaneous angles taken across all image frames of pixels within ROIs and the mask. **F.** Trajectories of pixels (shown only for some pixels) in ROIs from (B) – same colors as in (B). **G.** Estimated temporal speeds of all pixels in the ROIs. **H.** Distribution of lengths of trajectories. **I.** Distribution of maximum of temporal speeds. Blue lines (at 200 ms) in C and G indicate the stimulus onset.

nowhere close to the actual ones because the existence of the C2 wave in frame 24 is concurrent and within the boundaries of the C1 event (see Supplementary Movie 1). The CLG method thus captured the motion dynamics of the pixels of C1 and continued to follow its wake in subsequent frames. For the center pixel of C2, the estimated trajectory was that of pixel of C1. When the center C2 pixel appeared in time (in frame 24), it caused a discontinuity in the flow field which was ignored by the CLG method due to the spatial smoothness constraint (Eq. 2 see methods) and perhaps due to the motion constancy constraint as well (Eq. 4). When the C2 was estimated starting in frame 29 after coming out of the wake of C1, the trajectories and speeds of both center and non-center pixels were accurate compared to previous values (Fig. 5C-D) with the exception of some pixels which towards the end entered into the wake of E1 activity and thus the estimates of their trajectories and speeds were erroneous (see blue line in Fig. 5C-D). These results suggest that if events are overlapped in space-time, the estimation of trajectories and speeds can be erroneous and caution has to be taken in identifying the pixels of interest in both space and time.

We further tested the errors in optical-flow estimation (with both CLG and HS methods) due to overlapping events by changing the amplitude of individual Gaussian events (C1 = 2.5, E1 = 10, and C2 = 5; see Supplementary movie 1 and Fig. S2). With these amplitudes, there was a larger spatiotemporal overlap between the values of three motion patterns. Particularly in frames, where E1, C1, and C2 appeared concurrently, the amplitudes of C1 and C2 were comparable to the intensity values of neighboring E1 pixels. Hence, the estimation of motion was erroneous. Implementing optical-flow analyses on this dataset revealed degraded performance of both CLG and HS methods. However, the CLG method based analyses still performed better than the HS method in determining pixel trajectories and temporal speeds (compare Fig. S2 B,C and F,G with D,E and H,I respectively). The locations of sources determined from velocity vector fields estimated with the CLG method were qualitatively better than those estimated with the HS method while the opposite was true for determining sink locations (Fig. S2 J-M).

Limitations of the CLG method based analysis - sensitivity to the addition of noise. The efficacy of the CLG method based analysis in determining velocity vector fields, trajectories of pixels and temporal speeds from noisy data was assessed by adding 10% and 30% noise levels to all pixels. The addition of noise adversely affected the estimates of trajectories and speeds of pixels and identification of the location of sources and sinks with larger effects observed for larger noise levels (Fig. 5E-L shown only for C1; compare with Figs. 4B-C, 4J, and 4L). For example, the trajectory of the center pixel (green line in Figs. 5E and 5I) was not estimated to be purely the arc of a circle but has a translation component as well. For non-center pixels, the estimated trajectories were incomplete (red, blue, and black lines) with dislocated final pixel points. Similarly the estimated speed was erroneous for center and non-center pixels (Fig. 5F & J) but the overall trend of motion was successfully captured i.e. expansion, translation, and contraction. These errors in the estimates of pixel trajectories and temporal speeds were due to errors in the estimates of velocity vector fields by the CLG method i.e. the errors in the estimates of instantaneous speeds and angles (see Fig. 2B-E and 3B-E). The estimates of characteristics of sources and sinks were also adversely affected e.g. multiple sources and sinks were estimated near the actual source/sink points (Fig. 5G-H & K-L). However, non-existing sources and sinks that were estimated were spatiotemporally located very close to the actual space-time locations. Estimates of outlines of sources and sinks were also adversely affected with the addition of noise however, the strengths were still the highest near the origin point of sources and sinks (yellower and bluer contours respectively near the center). These results thus suggest that noise adversely affects the analysis and

filtering of raw data is advisable before determining velocity vector fields.

Optical-flow characterization of real optical imaging data

Wide-field optical imaging of mouse brain with forelimb sensory stimulation

Wide-field imaging of mouse brain was performed using voltagesensitive dye imaging while mouse forepaw was electrically stimulated. An increase in the intensity of the fluorescence signal was observed 10-15 ms after forelimb stimulation in the contralateral primary forelimb (FLS1) area of the sensory cortex (Fig. 6A and Supplementary movie 2). The activity expanded initially after appearing and translated towards the medio-caudal direction. Sequentially, another source of activity appeared in the contralateral secondary forelimb sensory area (FLS2) that was smaller in size and exhibited a small translation towards the medio-caudal direction. Thus, two regions of interest (ROIs) were defined on the primary and secondary forelimb cortical areas (Fig. 6B) after visual inspection of the image sequence. The average intensity of pixels in the ROIs sharply increased around frame 34 and then decreased in two steps (Fig. 6C). First, a sharp decrease to a value greater than half of the peak value was observed and then a gradual decrease to the baseline. This is in accordance with previously reported trends of FLS1 and FLS2 evoked responses to forelimb stimulation in isofluorane anesthetized mice (Mohajerani et al., 2013; Mohajerani et al., 2011).

Supplementary material related to this article can be found online at http://dx.doi.org/10.1016/j.neuroimage.2017.03.034.

Determination of velocity vector fields. The CLG method was used to estimate velocity vector fields of the image sequence, which captured the flow of activity spread (montage Fig. 6A). To analyze the global dynamics of pixel intensity changes in the whole imaged region, instantaneous speeds and angles were determined from velocity vectors for all pixels (in space and time). The distribution of these instantaneous speeds (black bars in Fig. 6D) showed a peak around 20 mm/s (10 mm/s = 1 p/f) with a mean  $\pm$  std value of 33.3  $\pm$ 24.7 mm/s and some pixels (~5%) moving as fast as 100 mm/s. The distribution of angles showed a peak in the medio-caudal direction (black line in Fig. 6E) which is in accordance with the visually observed features of the largest activity in the FLS1 area. Further, analysis was done for FLS1 and FLS2 ROIs to observe local dynamics and compare these functionally connected regions. The distributions of speeds indicated that the activity propagation was faster in the primary as compared to the secondary forelimb area (compare cyan and magenta in Fig. 6D). Mean ± std values of speeds for FLS1 and FLS2 ROIs were  $42.0 \pm 32.3$  mm/s and  $23.2 \pm 6.2$  mm/s respectively and the two distributions were significantly different from each other (two-sample t-test, p < 0.001). The distribution of angles for the FLS1 ROI showed the largest peak in the medio-caudal direction (Fig. 6E) whereas the FLS2 ROI did not prefer this direction.

Next, to determine the flow of activity of individual pixels, trajectories of pixels, their speeds over time, and lengths of trajectories were calculated for all pixels in both FLS1 and FLS2 ROIs (Fig. 6G-I). Pixels in the FLS1 ROI covered longer distances at higher speeds as compared to pixels in the FLS2 ROI. However, the distribution of distances traveled by pixels in the FLS1 ROI was bimodal indicating two peaks around 1.1 mm and 2.1 mm. As can be seen from the trajectories, the longer distances covered were in the medio-caudal direction (Fig. 6F) whereas the smaller travel was in the opposite and orthogonal directions. Furthermore, pixels which initially moved in the opposite direction, moved back in the medio-caudal direction later in the image frames (white arrow in Fig. 6F). The distribution of the

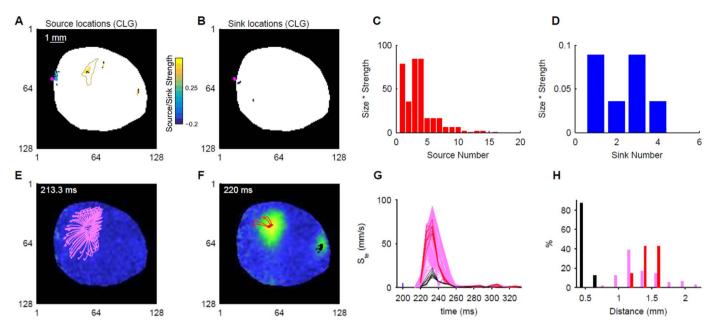


Fig. 7. Semi-automatic optical flow analysis (with CLG method). A-B. Identified source (A) and sink (B) locations (black dots) and associated contours of VSD imaging data presented in Fig. 6. Contour colors indicate strength of sources and sinks. C. Size times Strength versus source number for identifying significant sources. D. Same as (C) for sinks. E-F. Pixel trajectories shown for three selected sources (1 – light pink, 2 - red, and 3 - black) with high size × strength values. Note that source 1 was identified in 213.3 ms whereas sources 2 and 3 were identified in 220 ms. G-H. Distribution of temporal speeds (G) and length of trajectories (H) of pixels of sources 1, 2, and 3 (see E-F). The blue line (at 200 ms) in G shows the stimulus onset.

maximum value of speeds (Fig. 6I) indicated that most of the pixels in the FLS1 and FLS2 ROIs traveled at speeds of 50 mm/s and 20 mm/s respectively, which agrees with visually observed values.

Identification of sources and sinks. Using vector calculus methods, the locations of sources and sinks both in space and time were determined from the velocity vector fields (Fig. 7A-B). Their strengths and sizes were also calculated from contours. Significant sources i.e. ones generating observable activity, were identified by calculating the product of sizes and strengths. Three noteworthy sources [1, 3, and 4 in Fig. 7C] were further analyzed which collocated in space and time with the FLS1 and FLS2 ROIs selected in the analysis shown above (see Fig. 7E-F). For these sources, pixel trajectories for pixels within the contours were determined and their speeds and distances covered. The distributions of distances (Fig. 7H) and max speeds (not shown) were very similar to ones obtained from manual analysis.

Validation of analysis. The CLG method based analysis allows the study of spatiotemporal dynamics of brain activity sampled with voltage sensitive dye optical imaging. However, since the true velocity vector fields are unknown, we validated the values of speeds and angles of some randomly chosen pixels by determining them

manually. Manual validation of results for the FLS1 activity was satisfactory as pixel trajectories and temporal speeds were faithfully estimated. However, the estimated pixel trajectories seemed to be smaller than actual for the FLS2 activity perhaps because of overlapping smaller activities within the same region and smaller sampling rate i.e. discretization in time.

Quantitative comparison of voltage and calcium imaging data

To further validate the efficacy of our analysis, the spatiotemporal dynamics estimated from calcium and voltage imaging were compared (see methods for VSD and Ca imaging with GCaMP6f). With the assumption that the underlying biological activity in the brain is similar for mice stimulated with similar tone stimulus (i.e. frequency and amplitude), we hypothesized that the imaged data using different fluorophores would capture similar spatial dynamics from the auditory cortical area with differences originating from the temporal response dynamics of fluorophores. Since the temporal dynamics of the VSD signal are much faster than that of the GCaMP6f fluorescent protein, we expected the observation of faster velocities in VSD compared to GCaMP6f signals from the auditory cortex in response to sensory stimulation (Xie et al., 2016; Berger et al., 2007 May).

After collecting image sequences from the two imaging modalities,

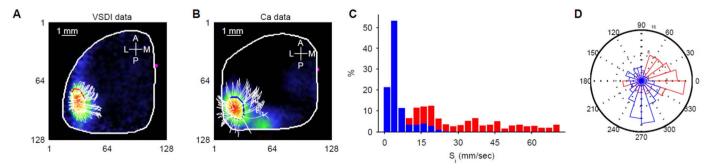


Fig. 8. Quantitative comparison of voltage and calcium imaging data. A-B. Representative images showing extent of activity over the auditory cortical area sampled with VSD (A) and calcium (B) imaging. White lines show travel trajectories for some pixels. Magenta dots indicate bregma location. C. Distribution of instantaneous speeds from all pixels within the regions of interest shown in A (red) and B (blue). D. Same as C but distribution of instantaneous angles.

preprocessing was done similarly to obtain the changes in fluorescence from the baseline  $(\Delta F/F_0)$ . For both datasets, cortical responses were observed in the primary auditory cortex (AC) and other cortical areas as well. In both image sequences, the latency of response in the AC area was similar (10-20 m s for VSDI and 20-30 m s for Ca imaging). Both activities expanded from primary auditory cortex and after some translation started to sink in the same area. Velocity vector fields for both image sequences were obtained as the output of optical flow analysis with the CLG method. To compare the instantaneous speeds and angles for pixels in the auditory cortical area of the two image sequences, we manually chose regions encompassing the auditory cortex (Fig. 8A-B). As expected, higher instantaneous speeds (Fig. 8C) were sampled with the VSDI (red) as compared to calcium imaging (blue). The distributions of angles (Fig. 8D) qualitatively were different but the largest peaks are in the mediocaudal direction (fourth quadrant) suggesting that the largest flow of activity was comparable in the two datasets.

Optical-Flow Analysis Toolbox in Matlab\* for investigating the spatiotemporal dynamics of Mesoscale brain activity (OFAMM)

We present version 1.0 of the OFAMM toolbox for the analysis of brain activity to be freely used by the research community under the GNU General Public License, version 3 (GPL-3.0) (2016, https://opensource.org/licenses/GPL-3.0). The toolbox and sample data including simulated and real optical imaging data can be downloaded from [http://lethbridgebraindynamics.com/ofamm/]. The toolbox contains a graphical user interface (GUI) shown in Fig. 9. This GUI

follows the optical brain activity data analysis framework described in Fig. 1. The detailed procedure to use the toolbox is provided in the supplementary material. Briefly, one can load an image sequence and mask (if applicable) into the GUI. On the image sequence, all or any of the three optical flow methods, HS, CLG, and TS, can be executed to estimate velocity vector fields which can be viewed in the GUI. The distributions of instantaneous speeds of all pixels in space-time within a region of interest can be plotted. Source/sink locations and trajectories of pixels of interest can then be obtained from the velocity vector fields by applying vector calculus analyses. Graphs of source/sink locations and those of pixel trajectories and speeds can be plotted. All the results are stored in mat files and the user can plot additional graphs at will (see supplementary material).

#### Discussion

Brain activity sampled with optical imaging consists of travelling (Lilly, 1954) and spiral waves which originate at sources and terminate at sinks. These waves are perceived in sampled image sequences as neurons in different brain regions become sequentially active (Townsend et al., 2015; Mohajerani et al., 2013; Sato et al., 2012; Benucci et al., 2007) e.g. source is the first region where neurons become active while neurons in the sink region are last to respond. Aquino et al., predicted similar waves on the cortical surface with modelling of the fMRI BOLD signal and experimentally demonstrated their presence in the visual cortex (Aquino et al., 2012). It is physiologically very relevant to quantify characteristics of such waves, sources, and sinks that might be used to specify the functionality of a

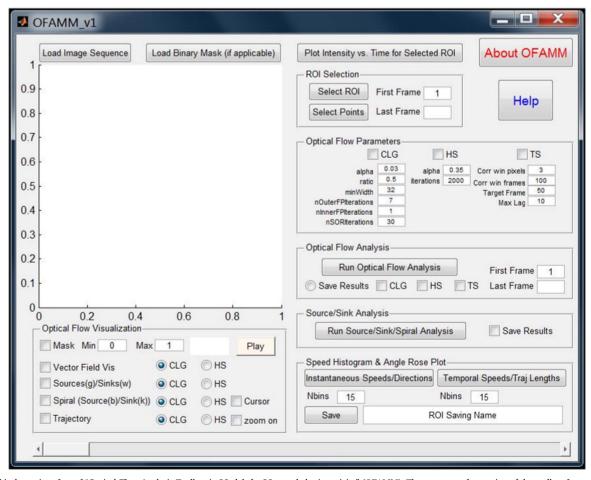


Fig. 9. Graphical user interface of "Optical Flow Analysis Toolbox in Matlab for Mesoscale brain activity" (OFAMM). The sequence of operation of the toolbox for analyzing optical imaging data parallels the flow chart of data analysis shown in Fig. 1i.e. first finding velocity vector fields and then identifying sources, sinks, trajectories of regions or pixels of interest, and their temporal speeds.

brain area and its connectivity with other brain areas (Takagaki et al., 2011; Mohajerani et al., 2013; Chan et al., 2015; Inouye et al., 1995; Lefèvre et al., 2007; Khan et al., 2014) during different brain states and various health states (Mohajerani et al., 2011).

It is known that many brain disorders result in part from circuit dysfunction. The impairment developed in neural circuits and network activity cause instability in neural communication (Busche and Konnerth, 2016; Kuljiš, 2010; Lim et al., 2014; Connor et al., 2016; Busche et al., 2015) which might increase or decrease the cortical dynamics of signal flow or due to circuit remodeling displace the location of sources or sinks. Thus, using these characteristics, normal and diseased brain circuits and their connectivity could be compared (Risacher and Saykin, 2013; Zhang et al., 2016; Visser et al., 2010).

Different methods used for the analysis of brain activity in optical imaging datasets have been reported previously but a consolidated toolbox for analyses is missing. Here we present a Matlab\* based software toolbox for the optical flow analyses of brain activity datasets. We have incorporated three optical flow methods into the toolbox namely HS, CLG, and TS. Using all three optical flow methods, we have estimated the characteristics of traveling waves, sources, and sinks in brain activity dynamics.

To compare their performance, we generated simulated data with known speeds and directions of movement (data which other investigators can use). The performance of HS, CLG, and TS for determining velocity vector fields was assessed by comparing the estimation errors in speeds and angles with and without the addition of noise (2nd row in Figs. 2 and 3). The results suggest that for noisy data the TS method is quite robust. However, since the TS method relies on determining temporal correlations in a group of frames, only one vector field is obtained where each vector represents average speed and direction. From this single vector field, the location of sources and sinks are determined only in space and not in time. The computational time for executing TS method is also considerable (Supplementary Fig. S1). In contrast, HS and CLG methods are computationally fast and provide vector fields for pairs of consecutive frames and thus can provide information regarding instantaneous speeds and directions. From multiple vector fields location of sources and sinks can be determined in space as well as time. The trajectories of pixels and their temporal speeds can also be determined. For the complex simulated dataset containing Gaussian propagating waves, we compared the performance of HS and CLG methods for determining activity trajectories, their temporal speeds, sources, and sinks. We demonstrated that the CLG method based analyses provide favorable estimates compared to the analyses based on the HS method (Fig. 4). This is in agreement with previously reported studies where the CLG method performed better in estimating speed of large moving objects in videos and displacement of fluorescently labeled proteins (Patel and Shukla, 2013; Delpiano et al., 2011). For this reason, we performed the analysis of real imaging data using only the CLG method. The user of the toolbox is however encouraged to experiment with all three optical flow methods for their simulated as well as experimentally obtained data.

The parameters for applying optical-flow methods e.g. window size for the TS method, and α and number of iterations for HS and CLG methods can also be set via the graphical user interface before executing the algorithms on an image sequence. Different values of these parameters could affect the estimated velocity vector fields, source and sink locations, and calculated pixel trajectories as well as temporal speeds. For our simulated and real optical imaging data, we also used alternate values of parameters (see methods) to compare their performance (see Supplementary Figs. S3–S7). With the use of alternate parameters, the best parameters that improved the performance of HS and CLG algorithms were found (e.g. compare Fig. S5 with Fig. 4). However, using these alternate parameters resulted in similar results in that (Frostig, 2009) the CLG method performs better than HS and (Nunez and Srinivasan, 2006) filtering data before using optical-flow algorithms is desirable. Since the application of optical-

flow methods is sensitive to the choice of algorithm and associated parameters whose values can be set subjectively, we recommend that manual validation of results should be performed.

The CLG method, although most favorable, has some limitations. When two or more events overlap in space-time, the CLG method provides erroneous results for the estimation of trajectories of pixels as well as their speeds (Fig. 5 and S2). The errors in estimation were worse with the HS method (data not shown). Caution is thus advised in selecting the pixels or regions of interest in space-time to correctly trace the pathways of activity patterns. Additionally, when noise was added to the signal, the errors in estimation with the CLG method increased. Thus it is very important to filter the real imaging data before performing the CLG method based analysis. We recommend permissible signal-to-noise ratios (SNRs) of up to 20 (~13 dB) as with SNR of 10, the errors in estimates of trajectories and temporal speeds were evident (Fig. 5E). For de-noising wide-field optical imaging data and improving SNR, we propose a similar procedure as mentioned in the methods Section Data Preprocessing. For the ease of the user, we have also included a separate GUI for preprocessing the wide-field imaging data (see Supplementary material for help on procedure for using the GUI).

We demonstrate how the CLG optical-flow method can be applied to real optical imaging data and that the distributions of instantaneous speeds and angles, temporal speeds and lengths of trajectories are useful parameters for studying population based cortical activity (Fig. 6). Additionally, with this semi-automatic analysis, significant sources can be identified and one can observe their spatial distributions (Fig. 7). We validated our optical flow analysis by using two sets of imaging data collected from two different animals with the same stimulation protocol but different fluorophores (VSD and GCaMP6f) for imaging (Fig. 8). As expected, higher instantaneous speeds were captured with VSD as compared to GCaMP6f since the response characteristics of VSD are much faster than GCaMP6f. Finally we present the optical flow analysis toolbox for characterizing the spatiotemporal dynamics in brain activity measured using optical imaging methods (Fig. 9).

#### **Conflicts of interest**

none.

#### **Author contribution statement**

MHM, MM, NA, and SI designed the study. NA and SI wrote Matlab\* code for the toolbox and designed the simulated data. MHM, and NA performed the experiments. NA and SI analyzed the data. SI, NA, and MHM wrote the manuscript.

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#### Supplementary\_Movie\_1\_simulated\_WFI\_Data.avi.

The movie shows all simulated image sequences that we used for comparing the performance of TS, HS, and CLG optical flow methods. It also contains an image sequence that we used for parametric analysis in which we determined the alternative values of parameters for HS and CLG methods. The name of each image sequence appears before its presentation in the movie. The Matlab files that generate these

simulated image sequences are provided in the supplementary sample

#### Supplementary\_Movie\_2\_Real\_WFI\_Data.avi.

The movie displays three image sequences that were collected in real wide-field imaging experiments with voltage-sensitive dye and genetically encoded calcium sensor (GCaMP6). Color of each pixel in an image sequence represents the respective percentage  $\Delta F/F_0$  value based on the scale in the colorbar. The text "Stimulus onset" appears in the frame in which the stimulus was applied and the name of each image sequence appears before its presentation in the movie. The supplementary sample data contains Matlab files of these image sequences as well as related files containing information about the experiments e.g. frame rate, pixels in ROIs, the mask file etc.

### Appendix A. Supporting information

Supplementary data associated with this article can be found in the online version at doi:10.1016/j.neuroimage.2017.03.034.

#### References

- Frostig, R.D. (Ed.), 2009. In Vivo Optical Imaging of Brain Function [Internet]2nd ed.. CRC Press/Taylor & Francis, Boca Raton (FL), (cited 2016 Oct 31)(. (Frontiers in Neuroscience)(. Available from)(http://www.ncbi.nlm.nih.gov/books/NBK20234/).
- Nunez, P.L., Srinivasan, R., 2006. A theoretical basis for standing and traveling brain waves measured with human EEG with implications for an integrated consciousness. Clin. Neurophysiol. J Int Fed. Clin. Neurophysiol. 117 (11), 2424–2435.
- Hillman, E.M.C., 2007. Optical brain imaging in vivo: techniques and applications from animal to man. J Biomed. Opt. 12 (5), 051402.
- Townsend, R.G., Solomon, S.S., Chen, S.C., Pietersen, A.N.J., Martin, P.R., Solomon, S.G., et al., 2015. Emergence of complex wave patterns in primate cerebral cortex. J Neurosci. 35 (11), 4657–4662.
- Grinvald, A., Omer, D.B., Sharon, D., Vanzetta, I., Hildesheim, R., 2016. Voltage-sensitive dye imaging of neocortical activity. Cold Spring Harb. Protoc., (Jan;2016(1) (1) and top(89367).
- Obien, M.E.J., Deligkaris, K., Bullmann, T., Bakkum, D.J., Frey, U., 2015. Revealing neuronal function through microelectrode array recordings. Front. Neurosci. [Internet], (Jan 6 [cited 2016 Jul 24)(;8. Available from)(http://www.ncbi.nlm.nih. gov/pmc/articles/PMC4285113/).
- Kalatsky, V.A., Stryker, M.P., 2003. New paradigm for optical imaging: temporally encoded maps of intrinsic signal. Neuron 38 (4), 529–545.
- Scanziani, M., Häusser, M., 2009. Electrophysiology in the age of light. Nature 461 (7266), 930–939.
- Nemoto, M., Nomura, Y., Tamura, M., Sato, C., Houkin, K., Abe, H., 1997. Optical imaging and measuring of local hemoglobin concentration and oxygenation changes during somatosensory stimulation in rat cerebral cortex. Adv. Exp. Med. Biol. 428, 521–531
- Biswal, N.C., Xu, Y., Zhu, Q., 2011. Imaging tumor oxyhemoglobin and deoxyhemoglobin concentrations with ultrasound-guided diffuse optical tomography. Technol. Cancer Res. Treat. 10 (5), 417–429.
- Reinert, K.C., Dunbar, R.L., Gao, W., Chen, G., Ebner, T.J., 2004. Flavoprotein autofluorescence imaging of neuronal activation in the cerebellar cortex in vivo. J. Neurophysiol. 92 (1), 199–211.
- Husson, T.R., Issa, N.P., 2009. Functional imaging with mitochondrial flavoprotein autofluorescence: theory, practice, and applications. In: Frostig, R.D. (Ed.), In Vivo Optical Imaging of Brain Function [Internet]2nd ed.. CRC Press/Taylor & Francis, Boca Raton (FL), (cited 2017 Feb 8)(. (Frontiers in Neuroscience)(. Available from) (http://www.ncbi.nlm.nih.gov/books/NBK20240/).
- Shibuki, K., Hishida, R., Tohmi, M., Takahashi, K., Kitaura, H., Kubota, Y., 2009. Flavoprotein fluorescence imaging of experience-dependent cortical plasticity in rodents. In: Frostig, R.D. (Ed.), In Vivo Optical Imaging of Brain Function [Internet] 2nd ed.. CRC Press/Taylor & Francis, Boca Raton (FL), (cited 2017 Feb 8)(. (Frontiers in Neuroscience)(. Available from)(http://www.ncbi.nlm.nih.gov/books/NBK20239/).
- Xie, Y., Chan, A.W., McGirr, A., Xue, S., Xiao, D., Zeng, H., et al., 2016. Resolution of high-frequency mesoscale intracortical maps using the genetically encoded glutamate sensor iGluSnFR. J. Neurosci. J. Soc. Neurosci. 36 (4), 1261–1272.
- Storace, D.A., Braubach, O.R., Jin, L., Cohen, L.B., Sung, U., 2015. Monitoring brain activity with protein voltage and calcium sensors. Sci. Rep. [Internet]., (May 13 [cited 2016 Jul 23)(;5. Available from)(http://www.ncbi.nlm.nih.gov/pmc/articles/ PMC4429559/).
- Abdelfattah, A.S., Farhi, S.L., Zhao, Y., Brinks, D., Zou, P., Ruangkittisakul, A., et al., 2016. A bright and fast red fluorescent protein voltage indicator that reports neuronal activity in organotypic brain slices. J. Neurosci. J. Soc. Neurosci. 36 (8), 2458–2472.
- Akemann, W., Mutoh, H., Perron, A., Park, Y.K., Iwamoto, Y., Knöpfel, T., 2012. Imaging neural circuit dynamics with a voltage-sensitive fluorescent protein. J. Neurophysiol. 108 (8), 2323–2337.
- Chen, T.-W., Wardill, T.J., Sun, Y., Pulver, S.R., Renninger, S.L., Baohan, A., et al., 2013.

- Ultrasensitive fluorescent proteins for imaging neuronal activity. Nature 499 (7458), 295–300.
- Buzsáki, G., Anastassiou, C.A., Koch, C., 2012. The origin of extracellular fields and currents–EEG, ECoG, LFP and spikes. Nat. Rev. Neurosci. 13 (6), 407–420.
- Huang, X., Xu, W., Liang, J., Takagaki, K., Gao, X., Wu, J.-Y., 2010. Spiral wave dynamics in neocortex. Neuron 68 (5), 978–990.
- Takagaki, K., Zhang, C., Wu, J.-Y., Ohl, F.W., 2011. Flow detection of propagating waves with temporospatial correlation of activity. J Neurosci. Methods 200 (2), 207–218.
- Mohajerani, M.H., Chan, A.W., Mohsenvand, M., LeDue, J., Liu, R., McVea, D.A., et al., 2013. Spontaneous cortical activity alternates between motifs defined by regional axonal projections. Nat. Neurosci. 16 (10), 1426–1435.
- Lilly, J.C., 1954. Instantaneous relations between the activities of closely spaced zones on the cerebral cortex; electrical figures during responses and spontaneous activity. Am. J. Physiol, 176 (3), 493–504.
- Vanni, M.P., Murphy, T.H., 2014. Mesoscale transcranial spontaneous activity mapping in GCaMP3 transgenic mice reveals extensive reciprocal connections between areas of somatomotor cortex. J. Neurosci. 34 (48), 15931–15946.
- Chan, A.W., Mohajerani, M.H., LeDue, J.M., Wang, Y.T., Murphy, T.H., 2015. Mesoscale infraslow spontaneous membrane potential fluctuations recapitulate high-frequency activity cortical motifs. Nat. Commun. 6, 7738.
- McVea, D.A., Mohajerani, M.H., Murphy, T.H., 2012. Voltage-sensitive dye imaging reveals dynamic spatiotemporal properties of cortical activity after spontaneous muscle twitches in the newborn rat. J. Neurosci. J. Soc. Neurosci. 32 (32), 10982–10994.
- White, B.R., Bauer, A.Q., Snyder, A.Z., Schlaggar, B.L., Lee, J.-M., Culver, J.P., 2011. Imaging of functional connectivity in the mouse brain. PloS One 6 (1), e16322.
- Minderer, M., Liu, W., Sumanovski, L.T., Kügler, S., Helmchen, F., Margolis, D.J., 2012. Chronic imaging of cortical sensory map dynamics using a genetically encoded calcium indicator. J. Physiol. 590 (1), 99–107.
- Kuhn, B., Denk, W., Bruno, R.M., 2008. In vivo two-photon voltage-sensitive dye imaging reveals top-down control of cortical layers 1 and 2 during wakefulness. Proc. Natl. Acad. Sci. USA 105 (21), 7588–7593.
  Han, F., Caporale, N., Dan, Y., 2008. Reverberation of recent visual experience in
- Han, F., Caporale, N., Dan, Y., 2008. Reverberation of recent visual experience in spontaneous cortical waves. Neuron 60 (2), 321–327.
- Paragios, N., Chen, Y., Faugeras, O. (Eds.), 2006. Handbook of Mathematical Models in Computer Vision [Internet]. Springer US, Boston, MA, (cited 2016 Jul 26)(. Available from)(http://link.springer.com/10.1007/0-387-28831-7).
- Sun D., Roth S., Black M., 2010. Secrets of optical flow estimation and their principles. In: 2010 IEEE Conference on Computer Vision and Pattern Recognition (CVPR). pp. 2432–2439.
- Inouye, T., Shinosaki, K., Iyama, A., Matsumoto, Y., Toi, S., Ishihara, T., 1994. Potential flow of frontal midline theta activity during a mental task in the human electroencephalogram. Neurosci. Lett. 169 (1–2), 145–148.
- Inouye, T., Shinosaki, K., Iyama, A., Matsumoto, Y., Toi, S., 1994. Moving potential field of frontal midline theta activity during a mental task. Brain Res. Cogn. Brain Res. 2 (2) 87–92
- Inouye, T., Shinosaki, K., Toi, S., Matsumoto, Y., Hosaka, N., 1995. Potential flow of alpha activity in the human electroencephalogram. Neurosci. Lett. 187 (1), 29–32.
   Horn, B.K.P., Schunck, B.G., 1981. Determining optical flow. Artif. Intell. 17 (1), 185–203
- Bruhn A., Weickert J., Schnörr C., 2002. Combining the Advantages of Local and Global Optic Flow Methods. In: Proceedings of the 24th DAGM Symposium on Pattern Recognition [Internet]. London, UK, UK: Springer-Verlag [cited2016 Jul 23]. pp. 454–462. Available from: (http://dl.acm.org/citation.cfm?Id=648287.756376).
- Jara-Wilde, J., Cerda, M., Delpiano, J., Härtel, S., 2015. An implementation of combined local-global optical flow. Image Process Line 5, 139–158.
- Otte, M., Nagel, H.-H., 1994. Optical flow estimation: advances and comparisons. In: Eklundh, J.-O. (Ed.), Computer Vision ECCV '94 [Internet]. Springer Berlin Heidelberg, (cited2016 Jul 26)(. pp. 49–60. (Lecture Notes in Computer Science)(. Available from)(http://link.springer.com.ezproxy.alu.talonline.ca/chapter/10.1007/3-540-57956-7\_5).
- Tavakoli V., Nambakhsh M., Sahba N., Makinian A., 2008. A new variational technique for combining affine registration and optical flow in echocardiography images. Conference Proceedings Annu International Conference IEEE Eng Med Biol Soc IEEE Eng Med Biol Soc Annu Conference 2008, pp. 205–208.
- Liu, T., Merat, A., Makhmalbaf, M.H.M., Fajardo, C., Merati, P., 2015. Comparison between optical flow and cross-correlation methods for extraction of velocity fields from particle images. Exp. Fluids 56 (8), 166.
- Kroeger, T., Timofte, R., Dai, D., Van Gool, L., 2016. Fast optical flow using dense inverse search. ArXiv160303590 Cs [Internet], (Mar 11 [cited2016 Jul 23)(; Available from) (http://arxiv.org/abs/1603.03590).
- Berger, T., Borgdorff, A., Crochet, S., Neubauer, F.B., Lefort, S., Fauvet, B., et al., 2007 May. Combined voltage and calcium epifluorescence imaging in vitro and in vivo reveals subthreshold and suprathreshold dynamics of mouse barrel cortex. J. Neurophysiol. 97 (5), 3751–3762.
- Estimate optical flow using Horn-Schunck method MATLAB [Internet]. [cited 2016 Jul 23]. Available from: http://www.mathworks.com/help/vision/ref/opticalflowhsclass.html.
- Liu C, 2009. Beyond Pixels: Exploring New Representations and Applications for Motion Analysis [Internet]. Available from: http://people.csail.mit.edu/celiu/OpticalFlow/.
- Madisen, L., Garner, A.R., Shimaoka, D., Chuong, A.S., Klapoetke, N.C., Li, L., et al., 2015. Transgenic mice for intersectional targeting of neural sensors and effectors with high specificity and performance. Neuron 85 (5), 942–958.
- Kyweriga, M., Mohajerani, M.H., 2016. Optogenetic approaches for mesoscopic brain mapping. Methods Mol. Biol. Clifton NJ 1408, 251–265.
- Shoham, D., Glaser, D.E., Arieli, A., Kenet, T., Wijnbergen, C., Toledo, Y., et al., 1999.

Imaging cortical dynamics at high spatial and temporal resolution with novel blue voltage-sensitive dyes. Neuron 24 (4), 791–802.

- Mohajerani, M.H., McVea, D.A., Fingas, M., Murphy, T.H., 2010. Mirrored bilateral slow-wave cortical activity within local circuits revealed by fast bihemispheric voltage-sensitive dye imaging in anesthetized and awake mice. J. Neurosci. 30 (10), 3745–3751
- Mohajerani, M.H., Aminoltejari, K., Murphy, T.H., 2011. Targeted mini-strokes produce changes in interhemispheric sensory signal processing that are indicative of disinhibition within minutes. Proc. Natl. Acad. Sci. USA 108 (22), E183–E191.
- Lim, D.H., Mohajerani, M.H., LeDue, J., Boyd, J., Chen, S., Murphy, T.H., 2012. In vivo large-scale cortical mapping using channelrhodopsin-2 stimulation in transgenic mice reveals asymmetric and reciprocal relationships between cortical areas. Front. Neural Circuits 6, 11.
- Lim, D.H., LeDue, J., Mohajerani, M.H., Vanni, M.P., Murphy, T.H., 2013. Optogenetic approaches for functional mouse brain mapping. Front. Neurosci. 7, 54.
- Sato, T.K., Nauhaus, I., Carandini, M., 2012. Traveling waves in visual cortex. Neuron 75 (2), 218–229.
- Ermentrout, G.B., Kleinfeld, D., 2001. Traveling electrical waves in cortex: insights from phase dynamics and speculation on a computational role. Neuron 29 (1), 33–44.
- Muller, L., Destexhe, A., 2012. Propagating waves in thalamus, cortex and the thalamocortical system: experiments and models. J. Physiol. Paris 106 (5–6), 222–238
- Frostig, R.D., Xiong, Y., Chen-Bee, C.H., Kvasnák, E., Stehberg, J., 2008. Large-scale organization of rat sensorimotor cortex based on a motif of large activation spreads. J. Neurosci. J. Soc. Neurosci. 28 (49), 13274–13284.
- Ferezou, I., Haiss, F., Gentet, L.J., Aronoff, R., Weber, B., Petersen, C.C.H., 2007. Spatiotemporal dynamics of cortical sensorimotor integration in behaving mice. Neuron 56 (5), 907–923.
- Polack, P.-O., Contreras, D., 2012. Long-range parallel processing and local recurrent activity in the visual cortex of the mouse. J. Neurosci. J. Soc. Neurosci. 32 (32), 11120–11131.
- GNU General Public License, version 3 (GPL-3.0) | Open Source Initiative [Internet]. [cited 2016 Jul 26]. Available from: https://opensource.org/licenses/GPL-3.0.
- Benucci, A., Frazor, R.A., Carandini, M., 2007. Standing waves and traveling waves distinguish two circuits in visual cortex. Neuron. 55 (1), 103–117.
- Aquino, K.M., Schira, M.M., Robinson, P.A., Drysdale, P.M., Breakspear, M., 2012. Hemodynamic traveling waves in human visual cortex. PLoS Comput. Biol. 8 (3), e1002435.

- Lefèvre, J., Obozinski, G., Baillet, S., 2007. Imaging brain activation streams from optical flow computation on 2-riemannian manifolds. In: Karssemeijer, N., Lelieveldt, B. (Eds.), Information Processing in Medical Imaging [Internet]. Springer Berlin Heidelberg, (cited2016 Jul 24)(. p. 470–481. (Lecture Notes in Computer Science)(. Available from)(http://link.springer.com.ezproxy.alu.talonline.ca/chapter/10.1007/ 978-3-540-73273-0 39).
- Khan, S., Lefèvre, J., Baillet, S., Michmizos, K.P., Ganesan, S., Kitzbichler, M.G., et al., 2014. Encoding cortical dynamics in sparse features. Front. Hum. Neurosci. 8, 338.
- Busche, M.A., Konnerth, A., 2016. Impairments of neural circuit function in Alzheimer's disease. Philos. Trans. R. Soc. Lond. B Biol. Sci. 371, 1700.
- Kuljiš, R.O., 2010. Integrative understanding of emergent brain properties, quantum brain hypotheses, and connectome alterations in dementia are key challenges to conquer Alzheimer's Disease. Front. Neurol. 1, 15.
- Lim, D.H., LeDue, J.M., Mohajerani, M.H., Murphy, T.H., 2014. Optogenetic mapping after stroke reveals network-wide scaling of functional connections and heterogeneous recovery of the peri-infarct. J. Neurosci. J. Soc. Neurosci. 34 (49), 16455–16466
- Connor, S.A., Ammendrup-Johnsen, I., Chan, A.W., Kishimoto, Y., Murayama, C., Kurihara, N., et al., 2016. Altered cortical dynamics and cognitive function upon Haploinsufficiency of the autism-linked excitatory synaptic suppressor MDGA2. Neuron. 91 (5), 1052–1068.
- Busche, M.A., Grienberger, C., Keskin, A.D., Song, B., Neumann, U., Staufenbiel, M., et al., 2015. Decreased amyloid-β and increased neuronal hyperactivity by immunotherapy in Alzheimer's models. Nat. Neurosci. 18 (12), 1725–1727.
- Risacher, S.L., Saykin, A.J., 2013. Neuroimaging Biomarkers of Neurodegenerative Diseases and Dementia. Semin Neurol. 33 (4), 386–416.
- Zhang, J., Cheng, W., Liu, Z., Zhang, K., Lei, X., Yao, Y., et al., 2016. Neural, electrophysiological and anatomical basis of brain-network variability and its characteristic changes in mental disorders. Brain J. Neurol., 14.
- Visser, S., Meijer, H.G.E., Lee, H.C., van Drongelen, W., van Putten, M.J.A.M., van Gils, S.A., 2010. Comparing epileptiform behavior of mesoscale detailed models and population models of neocortex. J. Clin. Neurophysiol. Publ. Am. Electroencephalogr. Soc. 27 (6), 471–478.
- Patel, E., Shukla, D., 2013. Comparison of optical flow algorithms for speed determination of moving objects. Int. J. Comput. Appl. 63, 5.
- Delpiano, J., Jara, J., Scheer, J., Ramírez, O.A., Ruiz-del-Solar, J., Härtel, S., 2011.
  Performance of optical flow techniques for motion analysis of fluorescent point signals in confocal microscopy. Mach. Vis. Appl. 23 (4), 675–689.