## Anexo\_II\_A\_Parametros Simulacion

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ANEXO II. A) Parámetros de la simulación de datos fMRI.

José Pedro Manzano Patrón

Lenguaje: Matlab

Este anexo solo contiene el fichero de configuración de los parámetros para la simulación de datos de fMRI con SimTB [http://mialab.mrn.org/software/simtb/documentation.html]. El experimento simulado consiste en:

- Un paradigma de tarea por bloques con una sola tarea de activación (ON-OFF -> TC\_block\_n = 2;)
- La tarea dura 20 TRs y el intervalo entre tareas (ISI) dura 30 TRs.
- El tiempo de adquisición TR = 2 seg.
- Se simulan 24 componentes independientes (sources).
- Se suponen datos pre-procesados, por lo que se han excluido:
  - Componente 6 (Sinus signal dropout), CSF (ICs 14 y 15) y sustancia blanca (ICs 16,17).
     Las componentes de CSF serán modificadas en el siguiente anexo por una señal sinusoidal y la misma señal al cuadrado (por lo que habrá 26 ICs en total)
- Las componentes 4, 5 (frontal), 18 (Dorsal Atention Network), 27 y 28 (sistema auditivo) se activan levemente (amplitud = 0.5) al escuchar la orden de mover o no.
- Las componentes 22 y 23 (regiones motoras) están fuertemente relacionadas (amplitud por encima de 0.7) con la tarea de mover. Presentan así correlación alta con las componentes 4, 5, 18, 27 y 28 en esos casos. La componente 7 (precuneo) se coactiva con 22 y 23, pero es mucho mas ruidosa.
- La componente 8 (DMN) está inversamente relacionada con la tarea -> En estado de reposo se activa y en los momentos de actividad o de recibir ordenes está inversamente correlacionada.
- Las componentes 7 (ruidosa), 22 y 23 (componentes motoras) están correlacionadas entre sí y se activan solo en algunos casos (cuando tienen que pulsar el boton).
- La generación de las TC se realiza a partir de la convolución del diseño por tareas con la HRF. Ver apartado de %%TC GENERATION para conocer los detalles.
- Por simplificación para el análisis se ha supuesto:

- Todas las ICs aparecen en todos los sujetos.
- Las tareas siguen el mismo orden en todos los sujetos.
- Mismo tejido (sustancia gris) y baseline para todos los sujetos y componentes.
- Para que la simulación sea más similar a datos fMRI reales:
  - Se ha añadido ruido (rician noise) sin spikes de CSF en todas las componentes.
  - Se han añadido artefactos por head-motion.
  - El CNR está distribuido uniformemente de 0.65 a 2 entre los sujetos.

Guarde el siguiente fichero con el nombre que desee, pero no olvide modificarlo cuando vaya a realizar la simulación. Para ejecutar la simulación y ver los resultados, acuda al **ANEXO I. B) SIMULACIÓN DATOS fMRI** 

```
In [ ]: %-----
     % To create the simulation parameter structure:
     % >> sP = simtb_create_sP('experiment_params_block', M, nC);
        Simulation can be executed with any number of subjects, M, or components, nC,
        though nC should be >= 4 given task modulation amplitudes (see Lines 46-49).
     % To run the simulation:
     % >> simtb_main(sP)
     %-----
     %% OUTPUT PARAMETERS
     %------
     % Directory to save simulation parameters and output
     out_path = '/Users/hose/Desktop/TFM_TECI/simulated_data/';
     % Prefix for saving output
     prefix = 'block';
     % FLAG to write data in NIFTI format rather than matlab
     saveNII_FLAG = 0;
     % Option to display output and create figures throughout the simulations
     verbose_display = 1;
     %-----
     %% RANDOMIZATION
     %-----
     seed = 3571;
                        % choose seed for repeatable simulation
     %% SIMULATION DIMENSIONS
     %-----
     M = 5; % number of subjects
     % nC is the number of components defined below, nC = length(SM_source_ID);
     nV = 148; % number of voxels; dataset will have [nV x nV] voxels.
     nT = 250; % number of time points
```

```
TR = 2; % repetition time
% number of different connectivity states
%nStates = 2;
% probability of unique events
%pU = 0.5;
% amplitude of unique events (relative to module-specific events)
%aU = .5;
% probability of state specific events
%pState = .5;
%Module membership for each state
%ModMem = zeros(nC, nStates);
% Number of event types (i.e., number of different modules)
%nE = 3;
%% SPATIAL SOURCES
%______
% Choose the sources. To launch a stand-alone GUI:
% >> simtb_pickSM
SM_source_ID = [ 2 3 4 5 7 8 9 10 11
              12 13 18 19 20 21 22 23 24 ...
               25 26 27 28 29 30]; % all but (1, 6, 14, 15, 16, 17)
% Sources 1 (general non-brain), 6 (Sinus signal dropout), 14,15 (CSF) and 16, 17 (WM) a
% as if the signal is already pre-processed
% LABEL COMPONENTS
%%
%nonbrain = find(SM_source_ID == 1);
s2
                         = find(SM_source_ID == 2);
s3
                         = find(SM_source_ID == 3);
% Frontal: 1 second temporal delay from bilateral frontal
         = find(SM_source_ID == 4);
comp_F1
comp_F2
          = find(SM_source_ID == 5);
% Medial Frontal: has lower baseline intensity (signal dropout)
%comp_MF = find(SM_source_ID == 6);
% Precuneus: activation only to targets
```

```
= find(SM_source_ID == 7);
comp_P
% DMN: negative activation to task events
comp_DMN
           = find(SM_source_ID == 8);
%
s9
                    = find(SM_source_ID == 9);
s10
                    = find(SM_source_ID == 10);
s11
                    = find(SM_source_ID == 11);
                    = find(SM_source_ID == 12);
s12
s13
                    = find(SM_source_ID == 13);
% Dorsal Attention Network: activation to novels more than targets
comp_DAN = find(SM_source_ID == 18);
s19
                      = find(SM_source_ID == 19);
s20
                      = find(SM_source_ID == 20);
s21
              = find(SM_source_ID == 21);
% (Sensori) Motor: activation to targets and novels (weakly)
         = find(SM_source_ID == 22);
comp_M1
comp_M2
          = find(SM_source_ID == 23);
% Bilateral frontal: positive activation to for targets and novels
         = find(SM_source_ID == 24);
comp_BF
s25
                       = find(SM_source_ID == 25);
s26
                       = find(SM_source_ID == 26);
% Here, we label components or component groups that may be used later
% Auditory: strong positive activation for all task events
comp_AUD1 = find(SM_source_ID == 27);
comp_AUD2 = find(SM_source_ID == 28);
% Hippocampus: activation only to novels
          = find(SM_source_ID == 29);
comp_H1
comp_H2
          = find(SM_source_ID == 30);
% compile list of all defined components of interest
complist = [s2 s3 s9 s10 s11 s12 s13 s19 s20 s21 s25 s26 ...
                        comp_AUD1 comp_AUD2 comp_DMN comp_BF comp_F1 comp_F2 ...
                      comp_DAN comp_H1 comp_H2 comp_M1 comp_M2 ...
            ];
%% COMPONENT PRESENCE
```

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_____
% \ [M \ x \ nC] \ matrix for component presence: 1 if included, 0 otherwise
% For components not of interest there is a 90% chance of component inclusion.
%SM\_present = (rand(M,nC) < 0.9); Some not present in some subjects
% Components of interest (complist) are included for all subjects.
SM_present(:,complist) = ones(M,length(complist));
               -----
%% SPATIAL VARIABILITY
%-----
% Variability related to differences in spatial location and shape.
SM\_translate\_x = 0.1*randn(M,nC);  % Translation in x, mean 0, SD 0.1 voxels.
SM_translate_y = 0.1*randn(M,nC); % Translation in y, mean 0, SD 0.1 voxels.
SM_theta
         = 1.0*randn(M,nC); % Rotation, mean 0, SD 1 degree.
           Note that each 'activation blob' is rotated independently.
%
SM_spread = 1+0.03*randn(M,nC); % Spread < 1 is contraction, spread > 1 is expansion.
%-----
%% TC GENERATION
Y-----
% Choose the model for TC generation. To see defined models:
% >> simtb_countTCmodels
TC_source_params = cell(M,nC);  % initialize the cell structure
% Use the same HRF for all subjects and relevant components
P(1) = 8; % delay of response (relative to onset)
P(2) = 16; % delay of undershoot (relative to onset)
P(3) = 1; % dispersion of response
P(4) = 1;  % dispersion of undershoot
P(5) = 8; % ratio of response to undershoot
P(6) = 6;  % onset (seconds)
P(7) = 24; % length of kernel (seconds)
[TC_source_params{:}] = deal(P);
% Implement 1 second onset delay for components comp_F1 and comp_F2, for instance
P(6) = P(6) + 1; \% delay by 1s
[TC_source_params{:,[comp_F1 comp_F2]}] = deal(P);
%-----
%% EXPERIMENT DESIGN
%-----
% BLOCKS
```

```
TC_block_same_FLAG = 1; % 1 = block structure same for all subjects
                 % 0 = block order will be randomized
TC_block_ISI
          = 30; % length of OFF inter-stimulus-intervals (in samples)
TC_block_amp = zeros(nC, TC_block_n); % initialize [nC x TC_block_n] matrix
% task-state 1: OFF
TC_block_amp([comp_AUD1 comp_AUD2],
                                  1) = 1.0; % moderate task-modulation
TC_block_amp([comp_BF comp_F1 comp_F2 comp_DAN], 1) = 0.7; % mild
TC_block_amp([comp_DMN],
                                    1) =-0.5; % negative weak
% task-state 2: ON
TC_block_amp([comp_AUD1 comp_AUD2],
                                   2) = 1.2; % strong
TC_block_amp([comp_BF comp_F1 comp_F2],
                                   2) = 1.0; % moderate
TC_block_amp([comp_DAN],
                                    2) = 0.8; \% mild
TC_block_amp([comp_P],
                                    2) = 0.5; \% weak
TC_block_amp([comp_M1 comp_M2],
                                   2) = 1.0; % moderate
TC_block_amp([comp_DMN],
                                    2) =-0.5; % negative weak
%______
%% UNIQUE EVENTS
Y------
TC_unique_FLAG = 1; % 1 = include unique events
TC_unique_prob = 0.2*ones(1,nC); % [1 x nC] prob of unique event at each TR
TC_unique_amp(:,[comp_AUD1 comp_AUD2])
                                     = 0.35;
TC_unique_amp(:,[comp_BF comp_F1 comp_F2])
                                     = 0.3;
TC_unique_amp(:,[comp_DAN])
                                      = 0.5;
TC_unique_amp(:,[comp_P])
                                      = 0.5;
TC_unique_amp(:,[comp_M1 comp_M2])
                                     = 0.2;
TC_unique_amp(:,[comp_H1 comp_H2])
                                      = 0.4:
TC_unique_amp(:,[comp_DMN])
                                      = 0.3;
%-----
%% DATASET BASELINE
Y______
% [1 x M] vector of baseline signal intensity for each subject
D_baseline = 800*ones(1,M); % [1 x M] vector of baseline signal intensity
Y------
%% TISSUE TYPES
%-----
% FLAG to include different tissue types (distinct baselines in the data)
                     % if 0, baseline intensity is constant
D_TT_FLAG = 0;
D_TT_level = [1.15, 0.8, 1, 1.2]; % TT fractional intensities
% To see/modify definitions for tissue profiles:
% >> edit simtb_SMsource.m
```

```
%% PEAK-TO-PEAK PERCENT SIGNAL CHANGE
%-----
D_pSC = 3 + 0.25*randn(M,nC); % [M x nC] matrix of percent signal changes
% To make statistical moments of data look more like real data
%D_pSC(:,comp_CSF1) = 1.2*D_pSC(:,comp_CSF1);
%D_pSC(:,comp\_CSF2) = 1.2*D_pSC(:,comp\_CSF2);
%D_pSC(:,comp_WM1) = 0.5*D_pSC(:,comp_WM1);
%D_pSC(:,comp_WM2) = 0.5*D_pSC(:,comp_WM2);
%-----
%% NOISE
%------
% [1 x M] vector of contrast-to-noise ratio for each subject
% CNR is distributed as uniform between 0.65 and 2.0 across subjects.
minCNR = 0.65; maxCNR = 2;
D_CNR = rand(1,M)*(maxCNR-minCNR) + minCNR;
%-----
%% MOTION
Y______
D_motion_FLAG = 0;
                  % 1=motion, 0=no motion
D_motion_ROTmax = 5;  % max rotation, in degrees
D_motion_deviates = ones(M,3); % proportion of max each subject moves
D_motion_deviates(1,:) = 0.5;  % Subject 1 moves half as much
%-----
% END of parameter definitions
```