

Titolo Tesi

Federico

May 2, 2022

Contents

1	Introduction: fMRI, Autism, ML	1
2	Dataset: ABIDE I & II	3
3	Image preprocessing	4
3.1	General overview	4
3.2	CPAC, preprocessing pipeline, atlases	5
4	Data: Pearson and Wavelets	7
4.1	Correlation and Z-Fisher transform	8
4.2	Wavelet analysis	8
5	ANN, deep learning	11
5.1	Random forest ??	12
5.2	Activation functions	12
5.3	loss functions	13
5.4	Backpropagation	13
5.5	How to assess network's performances	13
6	Harmonization	14
6.1	Harmonization - theory	14
6.2	Harmonization - results	15
7	Domain-adversarial NN	15
8	Results of deep learning	18
9	Explainable AI (XAI) with SHAP	19
9.1	SHAP - theory	19
9.2	SHAP - results	20
A	Principles of MRI imaging	20
A.1	spin-echo method	20
A.2	Image acquisition and k-space	21
A.3	fMRI	21
A.3.1	Gradient Echo sequences	22
A.3.2	EPI	22

1 Introduction: fMRI, Autism, ML

Autism spectrum disorder (ASD) is becoming a growing social issue, drawing more and more attention especially during the last decades, because of the raise in number of diagnosed cases. ASD is a neurodevelopmental disorder and refers to a broad range of conditions manifesting as deficits in social communication and interaction, repetitive behaviours, even speech difficulties. Early signs start appearing by the age of 2 or 3, but since there isn't a definitive medical test to diagnose it, often it is discovered even after adolescence.

An early diagnosis then, would provide the help and support they need, and, treating it at its early stage would hopefully lead to the regression of this condition. The main factors that bring to the developing of autism aren't very clear so far, and is becoming clearer that there isn't just a cause, but a concatenation and coexistence of various of them, the most strongly suspected causes are genetic and environmental causes. Genetic is associated with a rare gene's mutation, such as a deletion, duplication or inversion, and even though most of the mutation that increase the risk of developing autism are not been traced, it has been assessed that it has a percentage of inheritability around 90%, Ambiental factors such as prenatal air pollution or certain pesticides are also suspected to be associated with a increase of the risk to develop this disorder, even though the etiology is way more complex and it can't be assessed so accurately. Currently ASD is diagnosed by symptoms-based tests concerning behaviour assessment, From a neurological point of view, ASD, manifests in a reduced information processing owing to synaptic dysfunction that manifest in a reduced or altered brain activity. This is confirmed by several but controversial studies where both hyper functional connectivity and hypo-FC were detected in asd patients, in particularly this is age-dependent, over connectivity is usually associated to young children while under connectivity, with adolescences, and adults

In the last decades, different approaches to study this problem have been carried on, trying different diagnostic tool to investigate this matter, such as magnetic resonance imaging (MRI), functional-MRI (f-MRI), or electroencephalography (EEG). Among them, f-MRI is what is drawing more interest, and in particular resting state f-MRI was proven to be suitable for examining functional connectivity among brain regions that different highlighted to be more affected by autism rather than local brain regions.

f-MRI is a diagnostic tool that investigate blood flow variations across the whole brain structure. The measure the blood flow is directly linked with the neuronal activity, an increase of neuronal activity leads to a boost in blood flow in that specific brain area, because of the bigger demand of oxygen and other nutrients to that brain area. Blood vessels also increase their size The different neural connections are identified by measuring the BOLD fluctuations, (acronym for blood-oxygen-level dependent) from different areas of the brain, using the signal's difference between oxy (diamagnetic) and deoxyhemoglobin (paramagnetic). Oxyhemoglobin appears then brighter than deoxyhemoglobin The spontaneous fluctuations in the BOLD fluctuations during resting state are considered a strong indicator for the assessment of the properties of the brain system. During the study of a functional connectivity pattern, different traits are involved and characterise the signal in a strong and evident way, among these: sex and age of the patient are some of the most important both for structural and functional brain properties. Limiting our focus just on functional data, other factor conditioning the final outcome of a f-MRI data acquisition are the FIQ (Full intellectual quotient) or the eye status at scan, about this, functional connectivity with open steady eyes is different from closed eyes, and furthermore, closed eyes patient may be fallen asleep during the scan acquisition and this would heavily marking and modify the functional patterns.

It appears clear that the distinction of asd patient among normal ones it's not a simple and straightforward task. One of the most promising and powerful tool to tackle non-linear problems is machine learning with artificial neural network. ANN belong to artificial intelligence tools and make use of an algorithm that learns from data, and modify its parameters with the aim of recognise distinctive properties from the data and make prediction or decision on new unseen data. An relevant aspect that makes machine learning and artificial neural networks so popular is the ability to deal with non-linear problems introducing several non linear non-linear functions during training acting on their inputs or on the sum of them. A restraint of machine learning though, is that to perform well, an ANN needs to be trained on a big dataset, and the bigger the dataset, the better its performances are, especially for deep neural networks. To this end, particularly in medical field, several data from different acquisition centers need to be put together, and in the last years, several multicenter medical dataset such as the Human Connectome Project, the Alzheimer's Disease Neuroimaging Initiative (ADNI) or Autism Brain Imaging Data Exchange (ABIDE) were created. Unfortunately, one drawback of multicenter datasets is the unavoidable bias towards the site the data belongs to, resulting from hardware and scan's procedure differences, and it wouldn't be possible to uniform them to a single common acquisition protocol. To tackle this problem and try to uniform inter-scan variability, in this work, two different approaches are proposed: analytical harmonization and a deep learning approach. Harmonization is a procedure that aims to remove inter-site related effects in multi-site data, while trying to preserve all other information as biological-related features. The second approach, the deep learning one, tries to extract from data both disease and site information and use site information to remove the bias toward that site before making a prediction.

2 Dataset: ABIDE I & II

Data we are working with belong to the ABIDE dataset (Autism Brain Images Data Exchange): a project aiming to investigate autism using resting state fMRI scans acquired from different medical centers, and putting them together to collect as many data as possible.

The whole ABIDE dataset was published in two releases: ABIDE I released in August 2012 and containing 1112 patient scans, and ABIDE II released in June 2016 containing 1114 scans. ABIDE I includes scans collected from 17 different sites, and the 1112 patients consist of 539 patients with ASD and 573 typical control patients. ABIDE II includes scans collected from 19 different sites, and the 1114 patients consist of 593 patients with ASD and 521 typical control patients. Not every site belonging to ABIDE II is different from those of ABIDE I, but, even though some medical centers are the same, the scanner type, or acquisition pipeline and parameters may have been changed during the time interval between the two releases, so in the following analysis, they are regarded as different acquisition sites.

Besides scan images, ABIDE provides every information related to each patient, as age, sex, intelligence quotient (FIQ), eye status during the scan (open or closed), and every additional information provided by patients.

The vast majority of patients are males as shown in the histogram 1b, for a total amount of 1804 males and 422 females, while control/asd patient number, for each site is almost for each site as while the control/asd case per site is more or less balanced for almost every site, with the exception of KKI-1 that provided two times more controls than asd patients, and KUL-3 and NYU-2 that only provided ASD cases. For a visual comparison the number of controls/asd for each site is displayed on the histogram in figure 1a

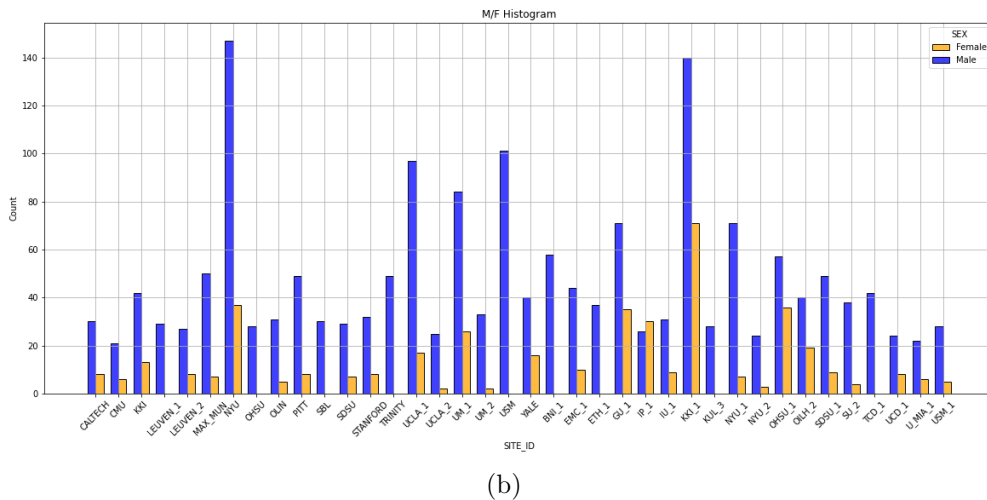
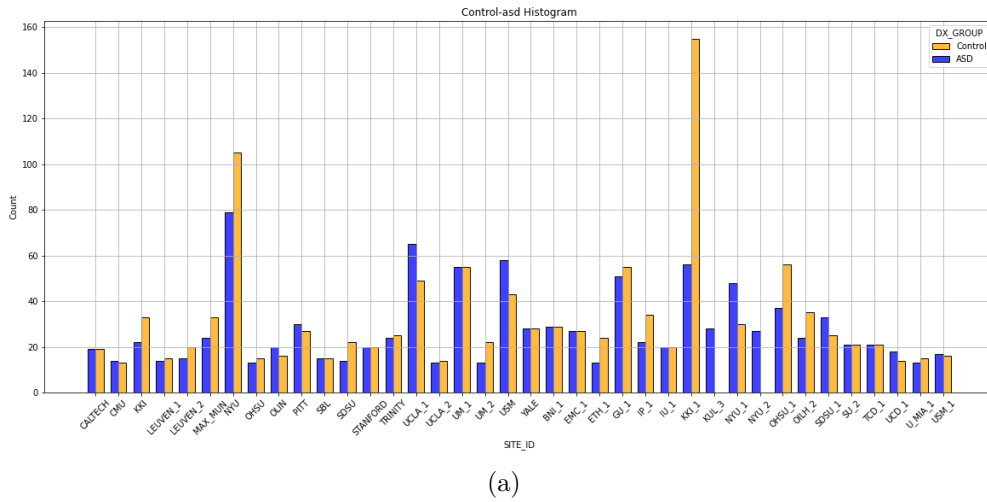
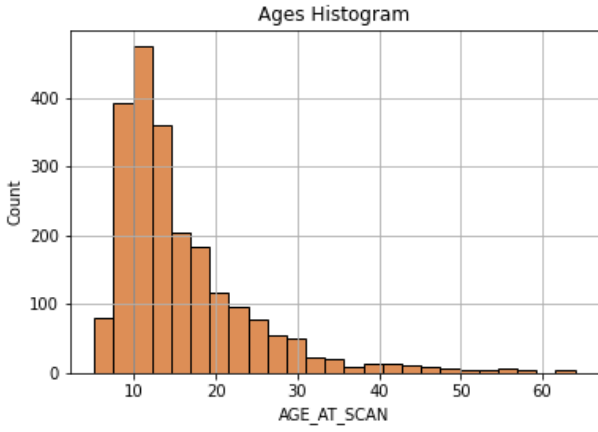
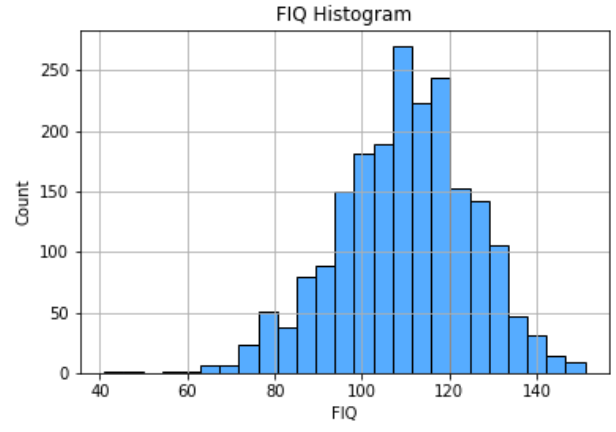


Figure 1

Patients in ABIDE dataset have ages ranging from 4 to > 50 years old, but as shown in figure 2a the vast majority of participant are younger than 40, precisely > 97 % of participant are under 40 y.o.

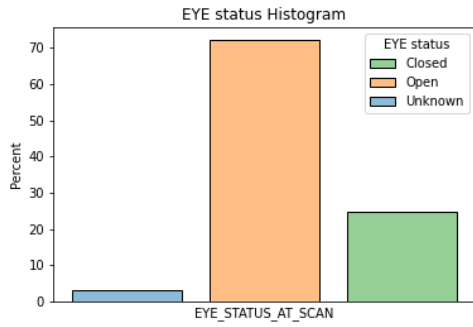


(a)

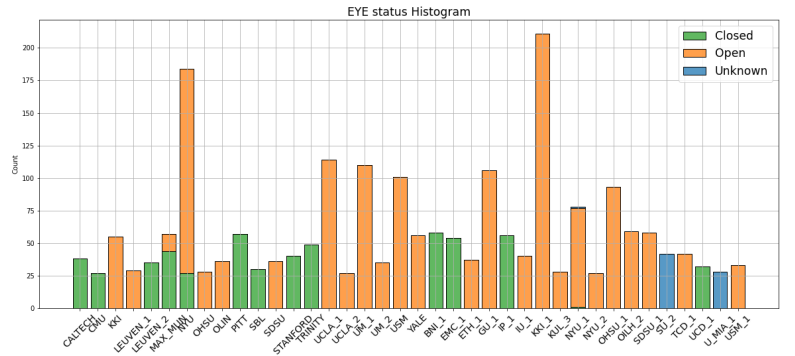


(b)

Figure 2



(a)



(b)

Figure 3

so we can restrict our further analysis to patients belonging to this group. Intelligence quotient, whose distribution is shown in figure 2b, was not provided for every participant, in fact 171 out of 2226 values were not provided, and before proceeding they were replaced by the mean of all the other values. The lack of a common acquisition protocol is also evident from the eye status at scan feature: as showed in figure 3b each site acquired scans either with open eyes or with closed, without a common method, and sometimes this information is not even specified. In its entirety, the whole dataset consists of more than 70% of patient acquired with open eyes, as showed in figure 3a.

For each patient in ABIDE both structural and functional images were provided.

3 Image preprocessing

3.1 General overview

For a better understanding of how MRI and f-MRI and signal detection work, we refer to appendix.. (talk briefly about echo time and TR ?)

During an MRI and f-MRI scan session, data are usually acquired slice by slice, and the thinner is the slice, the more spatial resolution we can accomplish. But there's a trade-off between spatial and temporal resolution: decrease the slice's thickness would lead to a better space resolution but at the cost of increasing repetition time to maintain the same Signal to Noise ratio (SNR). This is because the signal is proportional to the number of hydrogen nuclei, which is proportional to the slice volume.

In order to obtain a strong BOLD signal, echo time plays a significant role as well: to obtain the maximum strength signal it has to be set $T_E = T_2^*$ so in the case of BOLD signal it should be around 30 ms. Images acquired using a shorter echo time have a weaker BOLD signal because of the lack of signal to detect. [?]

Acquired data suffer from different source of noise and artifacts deriving from both hardware and physiological sources. As an instance, breath rate can affect BOLD signal because of the induced local motion of the brain's vessels, or the change in blood oxygenation and pressure

One of the main artifact intrinsic to the signal's nature is distortion and field inhomogeneities, deriving from making the scan sensitive to the BOLD signal which intrinsically is the detection of a signal loss due to field distortion. This could be corrected by employing small coils inside the scan to smooth magnetic field differences, this process called shimming still left some artifacts, therefore common data preprocessing pipelines use extra acquisitions to create a field map of the remaining field inhomogeneity and a shift and stretching voxel to correct for them.

Head motion is due to the physical movement of the patient inside the scanner. It results in a misalignment from one acquired volume to the next. Physiological noise is caused by the cardiac cycle and the breathing of the patient. They respectively provoke pulsatile motion of arteries, CSF and tissues and small head motion. As a second effect, the variable amount of air in the chest affects the magnetic field B_0 . To correct for head motion, motion correction steps are performed at the beginning. It works by spatially applying transformations as rotation or translation volume by volume, aiming to overlap every acquired slice to a chosen reference volume, like the first or that in the middle.

For EPI data, slice timing correction is usually performed as well. It aims to correct artifacts deriving from the sequentiality of acquisition for each slice of the brain is acquired at different time. The entire time elapsed to acquire all the slices is called repetition-time, and it usually is from 1 to 3 seconds. Slice timing correction uses interpolation in time to shift the BOLD timeseries of each voxel, in order to align them to a reference starting time. The use of interpolation, though can lead to a slight loss of high frequencies information.

A further common step is spatial smoothing of both structural and functional data. It operates calculating a weighted average of each pixel over neighbored voxels. To this end, a gaussian kernel with a chosen FWHM is applied to create the weights. Spatial smoothing is useful to avoid abrupt changes of signal between two neighbouring voxels.

Band-pass temporal filtering is commonly applied to BOLD data, aiming to reduce artifacts from hardware like the slow changing in the baseline of the BOLD signal over time. A low pass filter removes high frequencies above a cut off frequency, it is commonly applied in processing resting state fmri data because the physiological signal is driven by low frequencies oscillation while high ones are associated to noise.

A common further step is nuisance regression, which aims to reduce the structured noise: it works computing timecourses called nuisance regressors, These signals include motion and BOLD signal fluctuation from white matter or cerebro spinal fluid. From them, the variance is computed and this value is removed from the data using multi linear regression analysis.

When we need to run a group analysis, meaning analyze and compare different patient's images, one of the most relevant step is *registration*: structural (T1) data are aligned over a standard coordinates system space to universally describe location of the different brain parts, to make sure that the same voxel coordinate corresponds to the same brain area for all the subjects. The most common spaces is Talairach and Montreal Neurological Institute's MNI-152, obtained from linearly coregistering 152 T1 brain scans. A second registration step occurs within each patient to align EPI (functional) data to the structural image of that subject.

Looking at picture 4 we can see an example of step by step functional and structural image registration to a MNI152 template: first the functional image is registered to the structural and next, they are both registered to MNI template.

Once the structural and functional images were registered to a standard space, it's possible to extract brain region information using an atlas. Atlases are in the same space as the template image (MNI space for example) and are brain images where different brain areas are marked with different color intensities to associate each area to a label. This division into areas and the subsequent labeling, is called parcellization.

One of the most popular atlas is Harvard Oxford atlas obtained by manually labelling 37 MRI scans, aligning them to the standard MNI template and finally averaging each transformed label [?].

3.2 CPAC, preprocessing pipeline, atlases

In our work, data were preprocessed using CPAC (Configurable Pipeline for the Analysis of Connectomes): a configurable, open source pipeline, based on Nipype platform. CPAC was run on a Docker container installed on a computer assembled with:

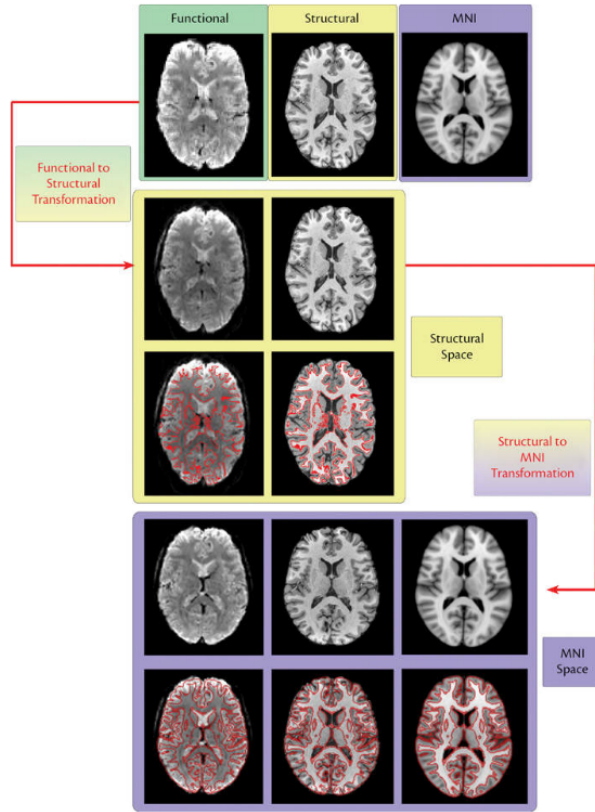


Figure 4: Registration steps: starting from the first row there are the acquired functional and structural images and the MNI template respectively. The second row shows the functional image registered on the structural space. The third row shows the same images as above, but overlaid with red boundaries extracted from the structural image. The fourth row shows the final images both registered to the MNI 152 template.

- 16-core Intel i7-5960X processor
- 64 Gb RAM

This allowed us to run 3 patients in parallel, reserving 4 cores for each participant and up to 12 Gb memory to each patient, necessary to save intermediate-steps outputs.

We choose to employ CPAC because, basing on machine learning classification results, [?] CPAC prove to be the most efficient preprocessing pipeline to preprocess ABIDE dataset. CPAC employs tools like AFNI, FSL and ANTS to perform image correction of structural MRI and rs f-MRI. Data were preprocessed following the same steps as the pipeline employed to create the free-access dataset ABIDE-preprocessed. This pipeline includes both anatomical and functional preprocessing. Anatomical pipeline speps consist in:

- Skull removal using AFNI's 3dSkullStrip
- Tissue segmentation using FSL-FAST, to separate gray matter, white matter and CSF, using a thresholding probability map whose threshold's values were set the same as ABIDE-preprocessed pipeline values
- Registration to a standard template using ANTS, with a spatial resolution of 2mm

Functional pipeline consists of the following steps

- Slice timing correction using AFNI-3dTshift
- Motion estimate and correction using AFNI-3dvolreg
- Distortion correction using PhaseDiff and AFNI 3dQWarp
- Create a brain-only mask of the functional data using AFNI 3DAUTOMASK

At the end of functional preprocessing steps, timeseries are extracted from each patient, making use of different atlases such as Automated Anatomical Labeled (AAL), Harvard-Oxford (HO), CC200, CC400, DesikanKilliany. An atlas is a 3D standard brain template, each one including a different parcelization of the brain, which is obtained dividing it into N labeled ROIs (Regions Of Interest) for anatomical and/or functional analysis. For example the DesikanKlein atlas employed in c-pac consists of 94 ROIs of which 32 cortical regions each side, 3 ROIs belonging to cerebellar vermis and 29 subcortical regions. According to previous studies, the atlas that gave best classification performances was HO, so this was the atlas we choose to employ for our work. The Harvard-Oxford atlas consists of a subcortical and a cortical atlas, with a total of 117 ROIs of which 21 subcortical and 96 corticals (48 for each emishpere). The HO atlas employed by CPAC is provided by FSL library ¹, and as includes both subcortical and cortical atlases, even though there's only 111 ROIs, namely 14 subcortical and 97 corticals. The lacking ROI in the subcortical areas are L/R cerebral white matter, L/R cerebral cortex, L/R lateral ventricle and the brain stem ². In this unified HO atlas (subcortical and cortical) the entire cerebral cortex area was removed because it's replaced by the finer parcellization of the cortical areas from the cortical atlas.

The extra cortical ROI in this H-O atlas is present in the 83th position and it's named 3455; but there are no information about this ROI neither on Harvard-Oxford documentation nor on CPAC's/FSL's documentation, and because it's only made of 2 voxels it was excluded for our further analysis.

4 Data: Pearson and Wavelets

Once all 110 timeseries were extracted from each patient, we need to create a correlation matrix, comparing each timeseries with all the others. The total number of combination that is possible to obtain from n timeseries is given by

$$N_{comb} = \frac{n \cdot (n - 1)}{2} \quad (1)$$

So in the case of HO atlas with 110 ROIs we obtain 5995 combination each one expressed by a correlation coefficient.

¹<https://fsl.fmrib.ox.ac.uk/fsl/fslwiki/>

²Subcortical ROIs <https://neurovault.org/images/1700/> Cortical ROIs <https://neurovault.org/images/1705/>

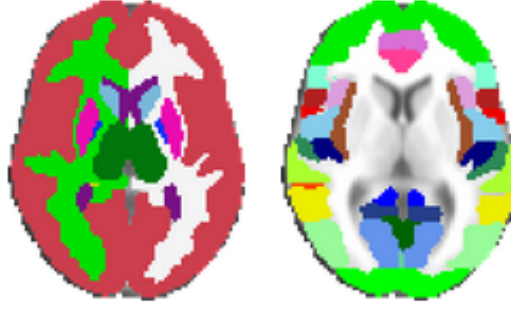


Figure 5: Harvard-Oxford subcortical and cortical atlas

4.1 Correlation and Z-Fisher transform

Pearson correlation often simply called correlation coefficient is the measure of a linear relation lying between two sets of data x_1 and x_2 , it's defined as the covariance of (x_1 , x_2) over the product of the standard deviation of the two sets and has the important properties to be scale-invariant in magnitude.

$$Corr(x, y) = r_{xy} = \frac{Cov(x, y)}{\sigma_x \sigma_y} = \frac{E[(x - \mu_x)(y - \mu_y)]}{\sigma_x \sigma_y} \quad (2)$$

This correlation coefficient can assume values between -1 and +1 where the extremes correspond to exact anti-correlation or correlation respectively, so that, if a linear relation lies between the two sets, a high absolute value indicates that the two series tend to be simultaneously greater or lower than their respective means

Given two series x and y , of length n correlation can be easily computed by

$$r_{xy} = \frac{\sum_{i=1}^n (x_i - \mu_{x_i})(y_i - \mu_{y_i})}{\sqrt{\sum_{i=1}^n (x_i - \mu_{x_i})^2} \sqrt{\sum_{i=1}^n (y_i - \mu_{y_i})^2}} \quad (3)$$

For each patient the Pearson correlation coefficient was computed for each timeseries' pair. Before further analysis, a common way to proceed is to transform each coefficient with Fisher z-transformation. The reason behind this is obtaining a variable which is normally distributed even when correlation coefficients are leaning toward the extremes.

$$z = \frac{1}{2} \ln \left(\frac{1+r}{1-r} \right) = \text{arctanh}(r) \quad (4)$$

At the end of this analysis, we obtain correlation matrices like that showed as an example in fig 6, referred to patient 20243 from ABIDE I dataset.

4.2 Wavelet analysis

A different approach to compute a correlation coefficient is by making use of wavelet analysis. Wavelet is a mathematical tool for analyzing time series or images, and provides a comprehensive way for investigating the bivariate relationship between timeseries both in time (or space) and frequency domain. To understand wavelet transformation it could be helpful to compare them with Fourier transform analysis.

The FT is a mathematical transform that takes a function from time or space domain into spatial or temporal frequency domain. It's a complex function that works decomposing the function $f(x)$ into series (ideally infinite sums) of sines and cosine of different frequencies, which are multiples (harmonics) of the fundamental frequency of the initial function. The discrete fourier transform, acts on a function defined on a finite domain returning a sequence of samples of the discrete fourier transform sampled at an interval equals to the reciprocal of the duration of the input sequence.

$$F(\omega) = \int_{-\infty}^{+\infty} f(t) e^{-i\omega t} dx \longrightarrow F(k) = \sum_{n=0}^{N-1} x_n e^{\frac{-2\pi i k n}{N}} \quad (5)$$

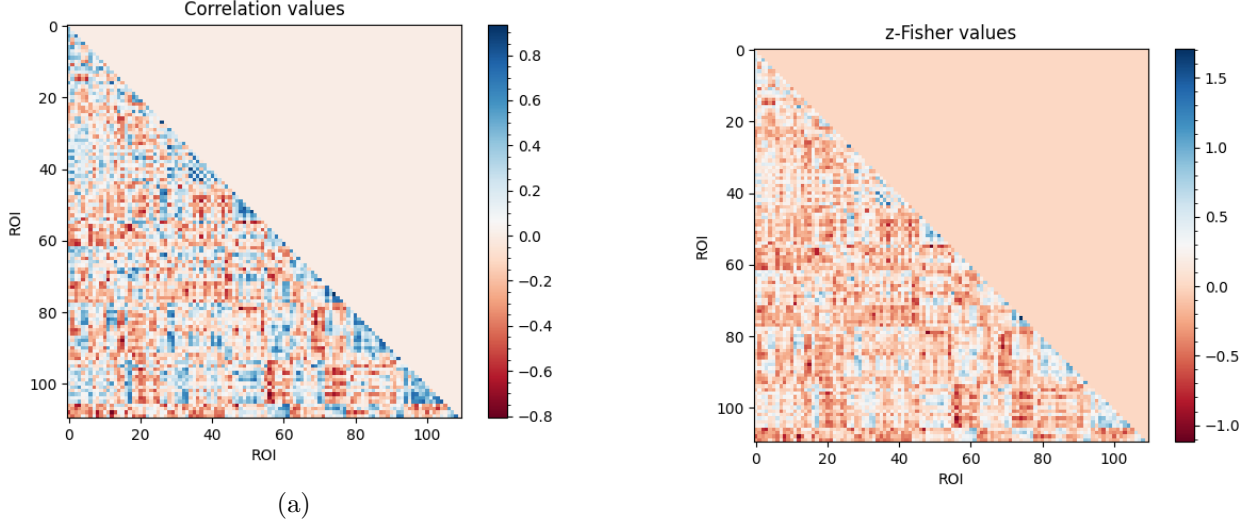


Figure 6: Correlation and z-values matrix computed from timeseries extracted using Harvard-Oxford atlas, from patient 20243 belonging to ABIDE I dataset

where N is the total length of the timeseries.

One of the main drawback of FT is the lacking of space information. Wavelet analysis provides for this lacking computing both frequency and spatial information from a signal.

A wavelet, literally “small wave ”is a wave function limited both in space and period: begins at zero, grows and decrease in a limited time period and returns to zero. So that it’s a function local in both the temporal and frequency domain. A wavelet is required to satisfy two properties: to have zero mean and unitary squared norm.

$$\text{zero\&mean} \quad (6)$$

It is defined over the space L^2 of Lebesgue measurable functions that are both absolutely integrable and square integrable. In this space the two properties can be satisfied. A wavelet transform is then formally a mapping from $L^2(\mathbb{R}) \rightarrow L^2(\mathbb{R}^2)$.

There are different wavelets that satisfy properties 6, such as Morlet, Meyer or Mexican hat.

One of the most convenient for our analysis is the Morlet wavelet given by eq 7

$$\psi(t) = e^{iw_0 t} e^{-t^2/2} \quad (7)$$

(Inserire immagini ?)

Two main type of analysis can be performed using wavelets: continuous wavelet transform (CWT) and discrete wavelet transform (DWT)

Continuous wavelet transform decomposes a time series in time-frequency domain by successively convolving the timeseries with several scaled and translated version of a mother wavelets $\psi_{a,b}(t)$

$$\psi_{a,b}(t) = \frac{1}{\sqrt{a}} \left(\frac{t-b}{a} \right) \quad (8)$$

where a is a positive real number and defines the scale, while b is any real number and defines the shift parameters: it specifies the location of the wavelet in space or time.

So the CWT given by the convolution of $\psi_{a,b}(t)$ with the function f (assumed to be real in the equation below) is

$$W_{a,b}(f) = \int_{-\infty}^{+\infty} f(t) \cdot \psi_{a,b}^*(t) dt \quad (9)$$

It can be useful to visualize this as: given a and b , a wavelet is slided across the signal and for each step a coefficient is extracted by multiplying the wavelet and the signal, and this operation is repeated for different values of a and b .

To computationally implement CWT, a discretized version was implemented on MATLAB tools Wavelet toolbox.

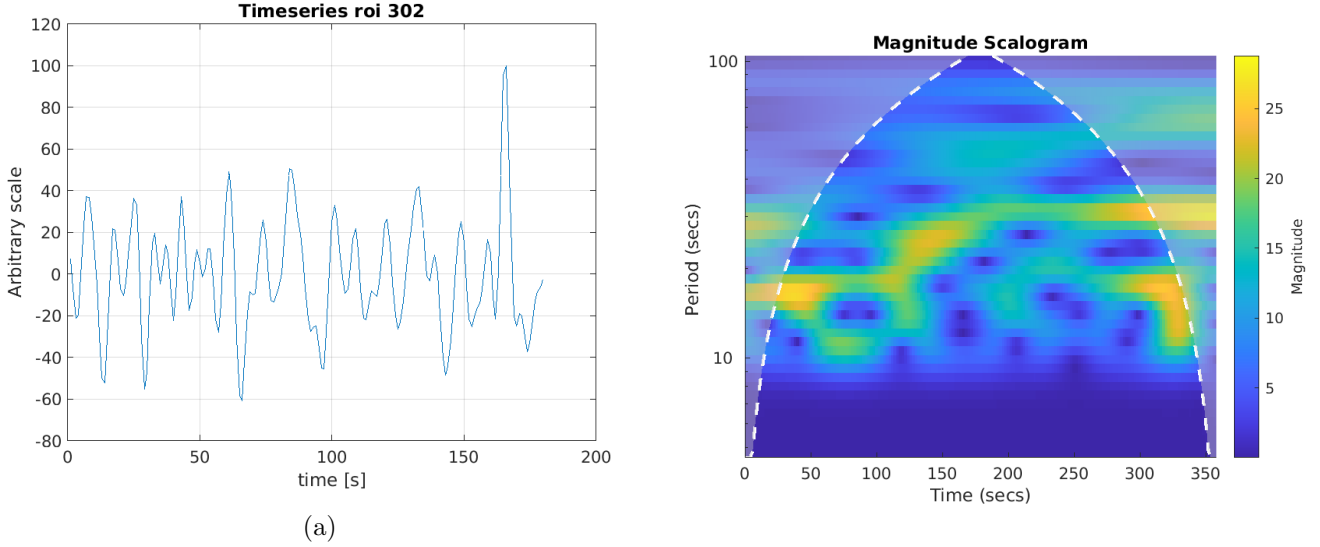


Figure 7

The difference between the continuous wavelet transform implemented on MATLAB and discrete wavelet transform lies in how finely stretching and shifting parameters are sampled: the CWT discretizes scale more finely than the discrete wavelet transform.

In the CWT, parameters are discretised based on a fractional power of two, by setting $a = 2^{1/\nu}$ and $b = \tau 2^{1/\nu}$ where ν and τ are integers. The ν parameter is often referred to as the number of “voices per octave” because increasing the scale by an octave (a doubling) requires ν intermediate steps, for example from $2^{\nu/\nu}$ to $2^{2\nu/\nu}$, ν steps are required. The larger the value of ν , the finer the discretization of the scale parameter. In the DWT, the scale parameter is always discretized to integer powers of 2, 2^j , $j=1,2,3,\dots$, so that the number of voices per octave is always 1

Using CWT it's possible to obtain signal's information on a matrix: a time-frequency plot also called scalogram by convolving the signal, using different wavelets parameters a and b from a grid.

One problem that arise from working with a signal with a finite support, is that when a wavelet is located at the beginning of at the end of a signal it extends itself outside the boundary of our data, and the convolution requires nonexistent values beyond the boundary. The confidence value obtained is then less accurate than other obtained for central values of time/space location. To overcome this problem it would be possible to accept this data information loss and truncate the values beyond boundaries; while an other approach could be artificially extend data using methods such as zero paddind which assumes that the signal is zero outside its original support, or symmetrization which extend the signal symmetrically outside the boundaries or smooth padding which recovers a signal by extrapolating values from the first derivative values or from the signal itself. Symmetrization method is the one employed for the subsequent analysis. Areas of the scalogram affected by these edge effects are indicated as outside the Cone of influence (COI).

Figure shows the timeserie belonging to ROI 308: left superior frontal gyrus of patient 51056 and its corresponding continuous wavelet transform.

In cwt figure, x axis corresponds to space or time points and on the y axis frequencies or periods from "a" parameter are showed. Values outside the COI (showed as a dotted white line) are shown with a lighter faded.

Continuous wavelet transform can only be used to analyze one signal at a time; to analyze and compare two signals an analysis called wavelet coherence has to be carried on. From two CWTs it's possible to compute the Cross Wavelet Transform (XWT) which allow us to examine their cross-wavelet power and relative phases.

Using XWT it's possible to compute the Wavelet Coherence.

Denoting as $W^X(a,b)$ the continuous wavelet transform of a signal X , it's defined the *wavelet power spectrum* of a signal $X(n)$ as $W^{XX}(a,b) = W^X(a,b) (W^X(a,b))^*$ where the $*$ represent the conjugate transpose. Similarly, the cross-wavelet spectrum of two time series $X(n)$ and $Y(n)$ is defined as $W^{XY}(a,b) = W^X(a,b) W^Y(a,b)$ whose module $|W^{XY}(a,b)|$ represents the amount of joint power between the two time series, and it's called *cross wavelet power* - From it it's possible to compute the the complex argument

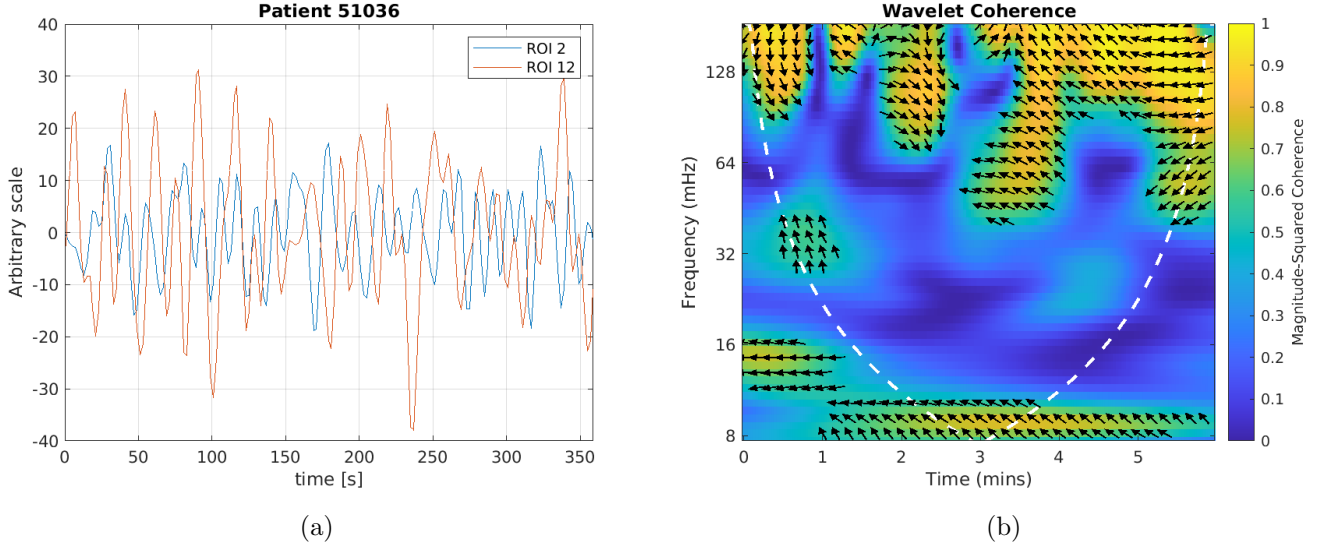


Figure 8

$\Delta\phi(a, b) = \arctan\left(\frac{\text{Im}[W^{XY}(a, b)]}{\text{Re}[W^{XY}(a, b)]}\right)$ which represents the phase between $X(n)$ and $Y(n)$.

A squared cross-wavelet coherence $R^2(n, s)$ is defined as reported in equation 10. This coefficient ranges in the interval $(0, 1)$ and represents the localized correlation coefficient between x and y in the time-frequency domain.

$$R^2(a, b) = \frac{|S(W^{XY}(a, b))|^2}{S(|W(a, b)^X|^2)S(|W(a, b)^Y|^2)} \quad (10)$$

To assess the level of significance over the noise, the statistical significance level of wavelet coherence has to be estimated using Monte Carlo methods generating a large (1000) samples of red noise for each time series, with the same 1-lag autoregressive coefficients (AR1 coefficient). For each pair of noise, a wavelet coherence matrix is calculated and stacked to obtain a $n \times m \times x \times 1000$ 3D noise matrix to extract the distribution of the entries. This null distribution of wavelet coherence coefficient is used to estimate the threshold to 5% significance level for the subsequent analysis.

To obtain a correlation-like coefficient from every two timeseries we are going to analyze, we use wavelet coherence coefficients from the matrix and their phase information to compute the time of in-phase coherence which can be seen as the percentage of time synchronicity between the two ROIs, defined as

$$c_{ij} = \frac{100}{N} \sum_{a, b} I\{R_{ij}^2(a, b) > a_{95}\} \cdot I\left\{-\frac{\pi}{4} < \arg_{ij}(W^{XY}(a, b)) < \frac{\pi}{4}\right\} \quad (11)$$

Where N is the total number of points inside the cone of influence (COI) $I\{\dots\}$ is either 0 or 1 depending on if the condition inside is satisfied; a_{95} is the threshold value above which the computed wavelet coherence coefficient is regarded as significant.

Similarly to this coefficient is the time of counter-phase coherence modifying from the formula above the phase condition into $I\left\{\arg_{ij}(W^{XY}(a, b)) < -\frac{3\pi}{4} \vee \arg_{ij}(W^{XY}(a, b)) > \frac{3\pi}{4}\right\}$

In short with this analysis, we are just considering coefficients with an high significance level (above 95%) and with a small, or big phase shift (between $-\frac{\pi}{4}$ and $\frac{\pi}{4}$ and the opposite).

Wavelet coherence maps were calculated using MATLAB's Wavelet Toolbox, which employs the Morlet wavelet, and analyze the frequency range using 12 subscales per octave and 9 octave.

Figure 8 represents the wavelet coherence scalogram obtained from two ROIs of patient 51056.

From equation 11 both coefficients in-phase and anti-phase matrix coefficients were computed. The correlation coefficients matrix for each subject, were then flattened and used as input to the neural networks.

5 ANN, deep learning

Machine learning algorithms can be divided into two macro areas: supervised and unsupervised learning. From a dataset, containing features, an unsupervised algorithm's goal is trying to learn the principal prop-

erties from the dataset's structure attempting to learn the probability distribution of all the features in it. On the other hand, in supervised learning each input data is associated with a label, specifying the class that data belongs to, so the algorithm is to learn the common features for each class trying to predict the label given the corresponding input. A perceptron is the simplest model of a neural network, corresponding to a single neuron which given an n-dimensional input $\mathbf{x} = \{x_1, x_2, \dots, x_n\}$ gives an m-dimensional output $\mathbf{y} = \{y_1, y_2, \dots, y_m\} = \sum w_{ij}x_j$. The output is given by a function called *activation function*. A deep neural network is a hierarchical organization of neurons layer, connected with other neurons. The input data are passed to the first input layer where each neuron produces an output using the activation function, which will be the input to the next neuron layer and so on. These middle layers are called *hidden layer* and each neuron on every layer is linked to each neuron of the previous and next layer -insert figure?-. each connection between two neurons is weighted, these weights are randomly set at the initial set up of the network. The output of each neuron i will therefore be $z_i = \sum w_{ij}x_j + b_i$. To fully define a neural network we just need some parameters called hyperparameters, the principals of which are:

- number of layers
- number of neuron for each layer
- activation function for each layer
- number of epochs
- learning rate

When reach the end of this process, we obtain the output: a numerical value indicating which class that input belongs to. The goal of a machine/deep learning algorithm is to learn using a train dataset, and generalize it in order to perform well on an unseen dataset called Test dataset. Performing well means to produce a low test error on the test dataset after minimizing the error on train dataset. This theme leads to one of the most important issue in machine learning: the concept of underfitting and overfitting. Underfitting occurs when the model isn't able to learn the required amount of informations during train; this usually happens when the model has a small number of parameters in regard to the number needed to explain the input features. On the contrary, overfitting occurs when the model has too many parameters in regard to those required to learn the input features as a result, the model performs well on the train dataset and has low performances in the test one. As an example in a two-dimensional space, if we aim to fit a dataset of 9 points sampled from a quadratic curve, if we try to fit it using a 9-th grade (or more) polynomial, we find out that infinite curves are possible to perfectly fit that dataset, but when chosen one, it will not be suited to describe an other dataset sampled from the same original quadratic curve. - better if I put a figure - To monitor the evolution of the model's performances during train is a common practise to split the train dataset in two parts, namely a real and actual train dataset and a smaller validation dataset used to make constant checkups on what the model is learning with the train set. The validation dataset is used for the fine tuning of hyperparameters and it typically consists of 10-30% of the whole train dataset. However, when dealing with small datasets, dividing it into Train and Test can be a non trivial issue because of the shortage of data for a proper train procedure. To work this out, a procedure called k-fold cross validation is often employed, at the cost of increasing computational costs. This consists in the creation of k different partition of the main whole dataset (namely in advance of dividing it into train + test). From these partitions, k-1 are used as train, and the last acts as test. Moving forward each combination of partitions is used so that each different partition is used as test, and the others as train. Then for each run k there are two subsets (a train and a test) from the original dataset. This way, each fold's train set is different from the others and same goes for test dataset. In practise what is usually done, is not a sequential partitioning the dataset, to avoid creating folder containing all the same label if the dataset was ordered, but randomly picking data in order to create subsets containing the same proportion between classes as the main dataset.

5.1 Random forest ??

5.2 Activation functions

The activation function of a layer defines the output of each neuron belonging to that layer, there are different activation function, linear or non-linear. A linear activation function outputs a value $f(x) = b + x^T W$. The

sigmoid function, or more correctly logistic function is the following

$$f(x) = \frac{1}{1 + e^{-x}} \quad (12)$$

- or the ReLU functions $\phi(z) = \max(0, z)$
- the softmax function for multiclass classification, gives the probability of the input belonging to each (mutually exclusive) class. SO the softmax can't be applied independently to each output z_i , since it depends on all elements of \mathbf{z} . The probability of belonging to the i -th class over a total of M classes is $p(i = i|z) = \phi(z) = \frac{e^{z_i}}{\sum_{j=1}^M e^{z_j}}$ being M the total number of classes

5.3 loss functions

This predicted value is then compared with the actual label value and the loss is computed between these two values. Computing the loss, returns a quantity value of how much the prediction is far from the actual label. Then an optimization function called gradient descent is applied to backwardly modify and tuning weights in order to minimize the loss. This algorithm employed by a feedforward network for weights' fine tuning is called *backpropagation* (backward propagation of errors). The function to be minimized is the *loss function* and the main technique to do so, is the likelihood maximization. The most common loss function is the cross-entropy loss, this function relies on the concept of entropy and is then defined as $H(y, \hat{y}) = -\sum_i y_i \cdot \log(\hat{y}_i)$. Where the sum is intended over all the possible class for a classification problem. In case of multi-class classification we call this loss categorical crossentropy loss, while if we only have two classes, (we are then performing a binary classification), the loss is (if we explicit the sum for two $i = \{0, 1\}$)

$$H_p(q) = -\frac{1}{N} \sum_{i=1}^N y_i \log(p(y_i)) + (1 - y_i) \log(1 - p(y_i)) \quad (13)$$

minimizing this function means maximizing the likelihood that for a classification with only 0 and 1 possible outcomes is

$$L = \prod_i \quad (14)$$

5.4 Backpropagation

After the computation of the loss value, the next step is the process called backpropagation, a process whereby network's weights are updated. Let's denote the loss function as $J(w, b) = \frac{1}{n} \sum_{i=1}^n L(y_i, \hat{y}_i)$. To find the right variation to the weights we can use the gradient descent algorithm, so we compute the gradient of the cost function $\nabla J(w)$ so that we can compute the new weight $w = w + \Delta w$ where $\Delta w = -\eta \nabla J(w)$ which expressed in component becomes $\Delta w_j = -\eta \frac{\partial J(w)}{\partial w_j} = -\eta \cdot \sum (\phi(z^{(i)}) - y^{(i)}) x_j^{(i)}$. A drawback of this gradient descent algorithm is that the weights are updated after computing a summation, namely, after the whole dataset is seen, so that optimization moves with big leaps. A common implementation of this gradient descent is an optimized version called stochastic gradient descent, whose difference is that the upgrade of the weights is sequential, for each training sample, so that the summation is not computed. Each step of this weight updating is then $\eta (\phi(z^{(i)}) - y^{(i)}) x_j^{(i)}$. Even for this implementation though, there's a drawback due to the fact that the steps towards the minimum are small and are influenced by all the unavoidable fluctuations in data training (some of them results in a move downward and some upward). A compromise between these two opposite way of work is mini batch gradient descent, which compute a step only after computed the gradient over a small subset of the training data. An other important weights optimization algorithm made from an extension of SGD is the Adam algorithm. It makes use of two different optimization algorithms: AdaGrad and RMSProp. It adapts the learning rate based on the first moment (the average) of the recent gradient magnitudes, and of the second moments (the uncentered variance) as well.

5.5 How to assess network's performances

Once the model has been trained, it is finally applied to the test set to assess the performances of the model to an previously unseen dataset. A common metric for model performances' evaluation is Accuracy:

it is simply defined as the ratio between the total number of correct prediction and the total number of predictions (total number of the test's data), so, using T and F for true and false, and P and N for positive and negative, accuracy is so defined

$$Accuracy = \frac{TP + TN}{TP + TN + FP + FN} \quad (15)$$

Even though accuracy may seem a good parameter, it isn't the most accurate one, when we're dealing with a unbalanced test dataset, if for example, in a binary classification with A and B classes, if just 10 out of 100 samples belongs to a different class, let's say A and B, and the model predicts every data belonging to the class B, we will get a score of 90 % accuracy, which is not a truthful result because the model isn't making any distinction between the two classes. There are then other ways for quality assessment that overcome this problem. One of this is based on the introduction of two quantities: precision and recall. Precision is the ratio between true positive and total positive cases (TP + FP), which is a measure of how many positive predicted cases were actually positive, while recall measures how many actual positives were correctly classified, namely it represents the true positive rate.

$$Precision = \quad \quad \quad Recall = \quad \quad \quad (16)$$

Unfortunately precision and recall are linked in a sense that often enhancing one decreases the other and vice-versa. There's therefore a trade off between recall and precision. If we want for instance a model to score more true positive, risking to have more false positives, we can set a threshold. To reach less the 10% positive missing means set the recall to 90%, this operation is called setting the operating point. Often though is not always established at the beginning, what operating point will be the best suited for our classification goals, so what is done is study the model under all the possible thresholds at once. This is achieved by studying the precision-recall curve. The closer a curve lies on the upper right corner (high precision and high recall), the more correctly the model is working. Besides the precision-recall curve, a similar curve is usually employed to study different thresholds' effect, it's called the receiver operating characteristics curve, usually referred as ROC curve. The ROC curve is build from the true positive rate (TPR, previously called recall) and the false positive rate (FPR) and shows the evolution of TPR vs FPR, and the ideal curve would be close to the top left (high tpr and low fpr). To summarize the model's performances with a single number using ROC's curve information, we use the Area under the curve AUC. The reference value for an AUC is 0.5 which is obtained when a model is just randomly predicting. In other words, the AUC can be explained as the probability that randomly picked point from the positive class, will have an higher score (according to the model) than a randomly picked point from the negative class.

6 Harmonization

6.1 Harmonization - theory

The issue of collecting a large dataset from multiple sources has a positive side because it facilitate the generalization and the robustness of the model, but, as a drawback, a level out procedure is necessary to avoid the model to learn differences deriving from inhomogeneity of data because of multiple site differences in data acquisition procedures. To try to reduce multi-site and multi-scan variety, procedures called harmonization are become necessary. The state-of-art procedure is called ComBat (meaning of combat?), specifically adapted for functional connectivity to mitigate scanner effects, is based on a previous method initially proposed [reference to johnson, fortin] in 2007, for gene expression studies to compute batch effect corrections. ComBat technique is based on the empirical Bayes method: it aims to reduce inter-site variability while preserving biological variability such as differences due to sex, age.. it assumes that the errors introduced in the imaging features can be standardized by adjusting the mean and the variance across the batches Let y_{ijf} be the numeric value of the feature f for the patient i acquired with the scan or from the site j, so that $i = 1, 2, \dots, K$ indexes the scanner, $j = 1, 2, \dots, N_i$ indexes the total number of subjects acquired for each scanner i, and f ranges from 1 to F being F the total number of features. Besides, let's assume that this value y, can be written as

$$y_{ijf} = \alpha_f + x_{ij}^T \beta_f + \gamma_{if} + \delta_{if} \epsilon_{ijf} \quad (17)$$

where α_f is the mean value for the feature f, x_{ij} is the entry of the matrix X for the covariates of interest such as age, gender, β_f is the vector of regression coefficients corresponding to X, the terms γ_{if} and δ_{if} represent

the additive and multiplicative terms for site- i effects related to feature f respectively, and ϵ_{ijf} is a error term which is assumed to follow a normal distribution with zero mean and variance σ_f^2 . To estimate feature-dependent parameters, data are standardized feature-wise, so that every feature has similar overall mean and variance. A least square regression, is then performed to determine parameters $\hat{\alpha}_f, \hat{\beta}_f, \hat{\gamma}_{if}$ and subsequently, $\hat{\sigma}_f^2 = \frac{1}{N} \sum_i j \left(y_{ijf} - \hat{\alpha}_f - X\hat{\beta}_f - \hat{\gamma}_{if} \right)^2$ being N the total number of patients. The standardized data are then calculated by 18

$$Z_{ijf} = \frac{y_{ijf} - \hat{\alpha}_f - X\hat{\beta}_f}{\hat{\sigma}_f} \quad (18)$$

Is furthermore assumed that standardized data follow a normal distribution $Z_{ijf} \sim N(\gamma_{if}, \delta_{if}^2)$ and that the distribution of batch normal distribution follow a normal distribution: $\gamma_{if} \sim N(Y_i, \tau_i^2)$ and an inverse gamma distribution $\delta_{if}^2 \sim \frac{\beta^\alpha}{\Gamma(\alpha)} (1/x)^{\alpha+1} \cdot e^{-b/x}$

The hyperparameters Y_i, τ_i^2 , are then estimated using the moment methos Under the distribution assumption above, the empirical batched parameters for batch effect $\gamma_{if}, \delta_{if}^2$ are estimaterg leadind to the estimate of the empirical batch adjusted data: y_{ijf}^{ComBat}

$$y_{ijf}^{ComBat} \quad (19)$$

6.2 Harmonization - results

Harmonization procedure was tested on the whole dataset to have a visual representation of how features were modified once the harmonization was applied

To test harmonization procedure and evaluate its performances in making sites' feature uniform, we used NeuroHArmonize package on 1494 patients coming from ABIDE I and ABIDE II of which 722 controls and 772 cases, We splitted them into train and test groups, then, for each train dataset, using NeuroHarmonize, the harmonization model was estimated on control subjects only, so these subjects were harmonized to remove site-dependent features and the chosen covariates were the age of the subject and the full intellectual quotient. The model was then applied to the case subjects belonging to train, and to both case and controls subjects of the test set. A random forest regressor was trained on this dataset attempting to make a binary classification of the site, for each pair of sites.

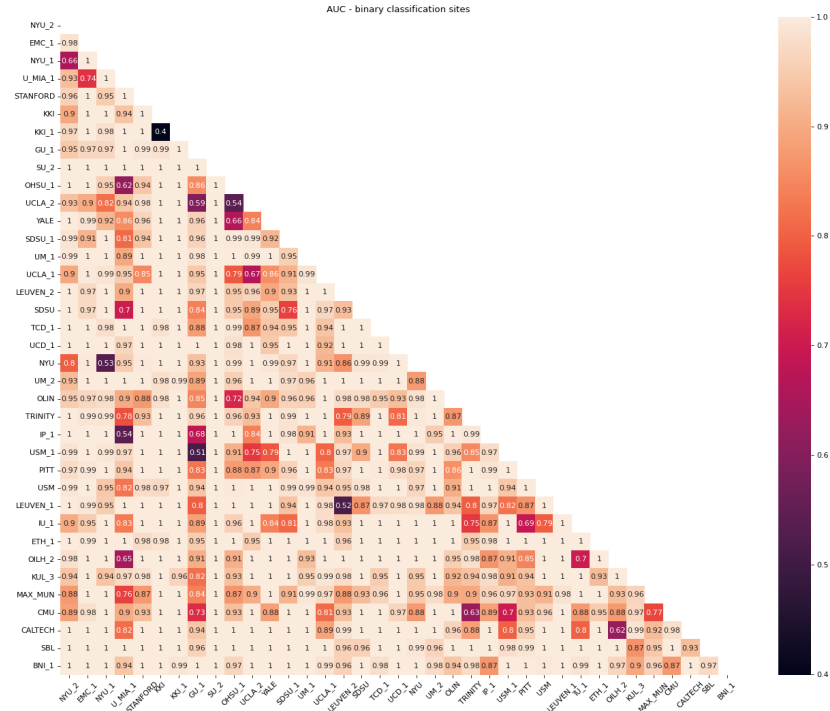
This approach, allows train and test dataset to be still independent, because harmonization model was only trained on Train subjects and once learnt the parameters, is applied to the Test dataset.

Figure 9 shows the result of classification on the dataset described above. Scores are reported in terms of AUC for the classification of each pair of sites, and sites along the x and y axes are ordered by incresging average age of patients coming from that site. Specifically, fig 9b shows the AUC score of the model trained with the raw, not harmonized dataset, it is clear from this binary classification that almost every site is well distinguishable from the others by the value of AUC, almost uniformly near to 1. On the contrary, in fig 9a values are lower, meaning that sites are not so distinguishable anymore, however, a certain distinctive features are still present for sites presenting high difference in mean age. This is a consequence of choosing age as covariate, which preserves differences in age-dependent features.

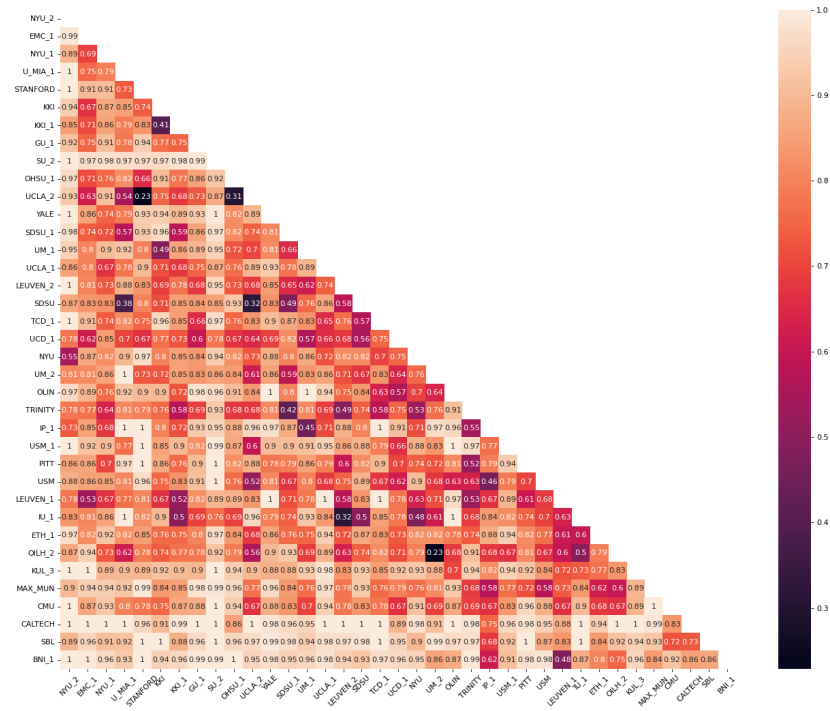
[?] To have a visual representation of how much each feature is modified in respect of its sourced site, in fig 10 are shown two features among 5995: *feature 324* and *feature 2800* as a function of sites. For a clearer idea of what is shown, feature 324 represents the correlation between right and left inferior frontal gyrus, and feature 2800 that between left precuneous cortex and the left inferior frontal gyrus Their values are plotted as a box along the y axis and sites are indicated along the x-axis, it is clear how harmonization is a powerful tool to level up feature and remove site-dependent information from them.

7 Domain-adversarial NN

An other approach is trying to train a neural network to recognise discriminative features, making train as much site-independent as possible. The genral idea underlying this network comes from a reviewed and readapted version of the solution of the problem tackled in Ganin - Ustonova article Their work takes place when we possess two different datased let's call them a Source and a Target dataset, containing features following two different distributions. Moreover, source dataset is labeled while Target is unlabeled, so their



(a)



(b)

Figure 9: Binary classification

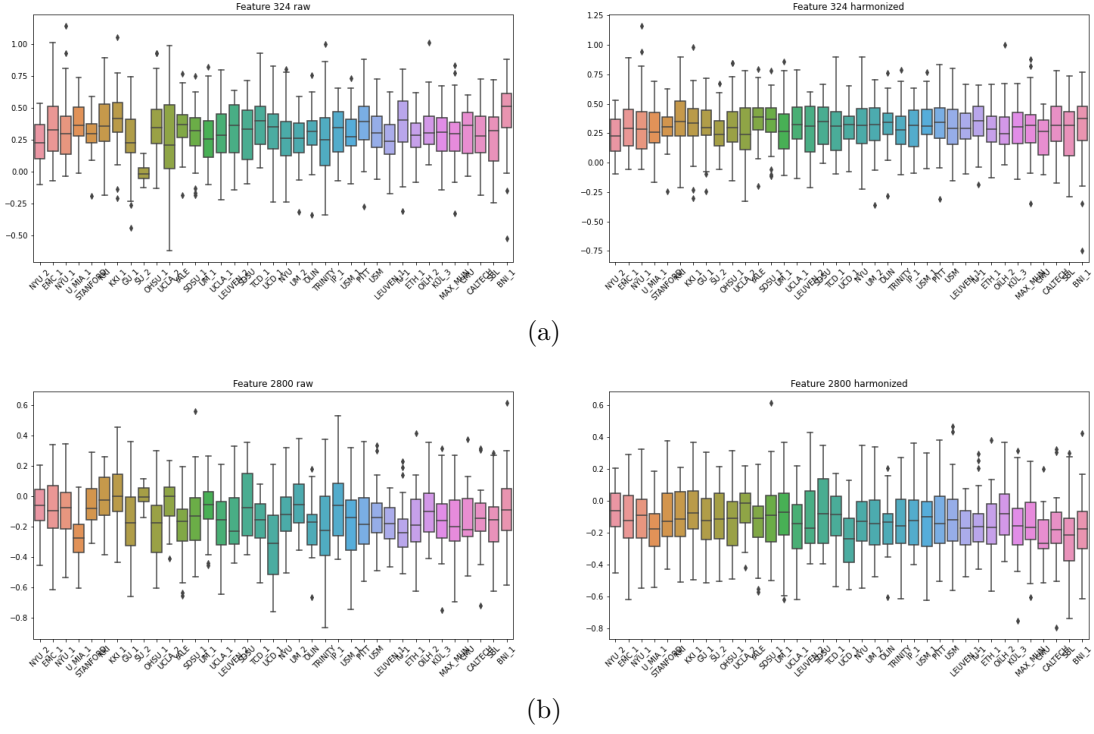


Figure 10: Features

goal was to create a network able to distinguish relevant features from the labeled dataset and make them domain independent. We illustrate the main aspects of this problem using a shallow neural network with a single hidden layer consisting of D nodes. Let's suppose the network takes as input an m -dimensional features vector; the hidden layer is then a function $G_f : R^m \rightarrow R^D$ of the type $G_f(x; \mathbf{W}, \mathbf{b}) = f(W \cdot x + b)$, where \mathbf{W} and \mathbf{b} are the matrix and vector parameters to be optimized. Similarly the output will be a function $G_y : R^D \rightarrow [0, 1]$ of the type $G_y(G_f(x; W, b); V, c) = f'(V \cdot G_f(x) + c)$. Where V and c are parameters to be optimized: a vector and a scalar respectively. A usual train carried on only on the (labeled) Source domain, will therefore bring to the minimization of the quantity

$$\min_{W, b, V, c} \left[\frac{1}{n} \sum_{i=1}^n L_y(G_y(G_f(x_i)), y_i | W, b, V, c) \right] \quad (20)$$

. To tackle the domain independence problem, an other term is added to the summation: a term coming from a domain classifier layer: $G_d : R^D \rightarrow [0, 1]$ of the type $G_d(G_f(x; W, b); u, z) = f'(V \cdot G_f(x) + c)$ that learns the vector and scalar parameters u and z , by trying to classify the domain. With this addition, the complete quantity to be optimized becomes

$$E(W, b, V, c, u, z) = \frac{1}{n} \sum_{i=1}^n \quad (21)$$

So the purpose of this kind of training is to find a saddle point by finding minimizing parameters for feature loss and maximizing those related to the site loss, namely

$$(\hat{\theta}_f, \hat{\theta}_y) = \underset{\theta_f, \theta_y}{\operatorname{argmin}} E(\theta_f, \theta_y, \hat{\theta}_d) \quad \hat{\theta}_d = \underset{\theta_d}{\operatorname{argmax}} E(\hat{\theta}_f, \hat{\theta}_y, \theta_d) \quad (22)$$

This could be achieved by training a network in an adversarial way, trying to minimize the category label loss and maximize site label loss. this is achieved by adding a domain classifier branch just after a main feature extractor branch. The domain classifier branch is characterized by a *gradient reversal layer* at the top of it. This layer has no parameters and don't act during forward propagation, but, during backpropagation, it invert all the weight from the subsequent layer before passing them to the preceding one. So as shown in figure -insert figure- the network is forked after the feature extractor branch, on one side, he starts the label predictor branch, and on the other, there's the domain classifier branch with the

gradient reversal layer at the beginning. The setup of this work, differs from Ganin’s because we don’t have a labeled Source and an unlabeled Target domain, but they are both labeled, so there’s no need to keep them separated and also, in our case, the Target domain is not just made of two domains, but of as many sites our data belong to.

In this work, the label classifier branch ends with a sigmoid function which return only one class label $y_i \in [0, 1]$ while the domain classifier branch ends with a multi-node layer of a number equal to the total sites’ number. An adversarial loss is also implemented, aimed to avoid that if, for instance, the domain classifier, correctly classify one input, the computed error would be low, and it would have a low contribution during the back propagation, and leaving the weights almost unmodified after the backpropagation, and therefore leaning towards that site.

8 Results of deep learning

In our work we employed a simple deep neural network architecture illustrated in figure .. with three hidden made up of 64, 64 and 32 nodes respectively activated by a ReLU function and each layer separated from the next by a Batch-Normalization and a 0.3-Dropout layer. After the 32-node layer we used a single output layer with a single node activated by a sigmoid function, for classification. As loss function we used binary crossentropy and the neural network’s weights were optimized via Adam optimizer with a learning rate of 10^{-4} and model’s performances were assessed with a 10 or 5 k-fold Cross validation, collecting the mean and std deviation of the results for each fold.

The other neural network employed was the site-adversarial neural network: following a similar structure as the DNN, it can be divided into three main components: the first feature extraction branch comprising the DNN up to the 32-node layer; from this point the network is splitted into two branches: one categorical classifier, along the lines of the DNN consists of the last output layer with a sigmoid activation function, and the other branch, the site classifier consisting of a gradient reversal layer as the first layer followed by a layer with 16 nodes, and after a batch Batch-Normalization layer, the multi-output layer with N nodes, being N the total number of sites input data belong to. This final layer is activated by a softmax function, and the loss function employed for site classification is the categorical crossentropy implemented using a custom function to deal with a possible low crossentropy score when by chance the site classifier, categorize one or more input data’s sitelabel correctly, which would have low impact on the whole branch classifier making the adversarial strategy useless.

We mainly tried three different kind of analysis using this DNNs: 1) classification of raw (not harmonized) data, 2) classification on the whole harmonized dataset, creating the harmonization model on all control subjects and applying the model on all ASD subjects, and 3) classification with harmonization implemented inside a K-fold cross-validation, creating the model on controls and applying it to ASD, and 4) classification using the domain adversarial neural network.

Following this procedure we implement a k-fold cross validation using stratifiedKFold provided by sklearn library. Data were splitted into a train and a test dataset, therefore using train data we computed the harmonization model only on control patients and then apply the model to asd patient and both Control and ASD Test data. This is the best way to maintain train and test data as much separated as possible and avoid data leakage between train and test.

Analysis were carried out using different selection criterias and thresholds: The first classification attempt was using the whole dataset from ABIDE 1 and 2, consisting of only male patients aged between 5 and 40 years old, without any further constraint on eye status at scan. Results 1) 2) 3)

The second analysis was performed under the same constraint as above, but with an additional condition on eye status choosing to select only patient who kept eyes open throughout the entire scan time.

The following analysis were carried out with the same constraint of the first two, but separately on ABIDE 1 and ABIDE 2 dataset.

A further analysis concerns the feature’s dimensionality reduction, that is the reduction of features from the input dataset using PCA For each one of the analysis above, PCA was implemented to restrain the effect of overfitting.

From the initial input data consisting of 5995 features, we tried different to classify using 20, 50, 100 and 200 principal components

9 Explainable AI (XAI) with SHAP

9.1 SHAP - theory

SHAP (SHapley Additive exPlanation) is a technique based on game theory which provides a method for model explainability. Let's say we have a coalition game: N players, and different subsets created from these players. Each subset of players S gain a payoff at the end of the game. A function ν maps every subset to its payoff. $\nu(S_i) = \text{payoff}_i$. Based on which player had more influence in the final score, we ask how to split this payoff in a fair way between each player of the subset. To this end we use **shapley values**: a solution concept (i.e. a formal rule to predict how a game will be played) $\phi_i(\nu)$. It represents the incremental benefit of including player i in the subset and average over all subsets that include player i . They are specific for each player i and represents the marginal contribution of that player to the final score, in other words they are the difference between the group's score with player i and the group's score without player i . To apply this concept to a machine learning model we just substitute the payout is the model prediction and each player is the value of a single feature. Shap value is then a weighted average of the marginal contribution of a feature (player) across all possible coalitions

For each player i and a map function ν shap values are defined as:

$$\phi_i(\nu) = \frac{1}{N} \sum_{S \subseteq N \setminus \{i\}} \binom{N-1}{|S|}^{-1} (\nu(S \cup \{i\}) - \nu(S)) \quad (23)$$

To get the idea of how we can explain a model, we start asking: how much is a prediction driven by the fact that a feature has a certain value A , instead of its baseline value \bar{A} ? the baseline value \bar{A} is the result of To answer this question we introduce a coalition vector $z' = \{0, 1\}^F$: where F is the total number of features; we have then a binar vector made by zeros and ones where zero means that we withheld the corresponding variable from our analysis, and the one that we are including it. To map the z' vector to the corresponding feature values vector a function $h_x(z') = x$ which returns the actual features' value for the corresponding input vector x . Thus we aim to study a simplified model $g(z')$ which is equal to $f(h_x(z'))$ if $z' \approx x'$. The most important properties of this models is called local accuracy and states that g and f must give the same result when passing all the features: g has to match f when $x = h_x(x')$. We attribute a contribution ϕ_i to each simplified vector z'_i

$$g(z') = \phi_0 + \sum_{i=0}^N \phi_i z'_i \quad (24)$$

The solution to this problem exists and is unique, it's shapley values. To assign the importance value to a feature we need to train two different models: one with that feature included and the second one with that feature excluded and thereafter compute the difference between the two models' prediction. computing this differences for all possible subsets $S \subseteq F \setminus \{i\}$. We are then going to use them to explain a deep learning model with the SHAP method. There are different way of approximating the computation of SHAP values like KernelSHAP which is a model agnostic explanation method, or Deep SHAP which employs shapley values into DeepLIFT algorithm. DeepLIFT aims to compute the difference in output deriving with an input different from some "reference" input chosen depending on the data we are working on. Let t represent the output of some inner neuron and let x_1, x_2, \dots, x_n be some preceding neurons necessary to compute t , if we label t_0 the reference output, we can compute the value $\Delta t = t - t_0$ and use them to define the DeepLIFT values $C_{\Delta x_i \Delta t}$ so that $\sum_i 1^n C_{\Delta x_i \Delta t} = \Delta t$. For a given input neuron x with difference from reference Δx and a target neuron t , is defined a multiplier $m_{\Delta x \Delta t} = \frac{C_{\Delta x \Delta t}}{\Delta x}$ which represents the contribution of Δx to Δt . Deep SHAP combines shap values for smaller components of a network to compute value for the whole network, it does so recursively passing Deep LIFT multipliers during backpropagation: we can indeed re-writing multiplier values in terms of shap values, for a target neuron f_3 we have $m_{x_j f_3} = \frac{\phi_i(f_3, x)}{x_j - E[x_j]}$ obtaining $\phi_i(f_3, y) \approx m_{y_i, f_3} (y_i - E[y_i])$

To express the significance of a feature we plotted them using SHAP feature importance, based on the relation between feature importance and shap value: feature with high absolute shapley value are important. The importance value for a feature j is then calculated from $I_j = \frac{1}{n} \sum_{i=1}^n \|\phi_j^{(i)}\|$

9.2 SHAP - results

A Principles of MRI imaging

Nuclear magnetic resonance imaging, is a widespread imaging technique It's widely employed in radiology to obtain anatomical images or study physiological processes, taking advantage of its flexible sensitivity to different tissues It's based on the interaction of a nuclear spin of any tissue's molecule, with an external magnetic field \mathbf{B}_0 . The principal molecule we are referring to is water, and subsequently, hydrogen is the dominant nucleus in MRI. When a nucleus undergoes an external magnetic field, the interaction would make the angular moment $\vec{\mu}$ align with the magnetic field, but, since protons have a thermal energy associated with the temperature T , they are not static, so they start precessing around the symmetry axes given by the direction of the field \vec{B}_0 . The force applied to a spinning proton under a magnetic field is given by

$$\vec{\tau} = \vec{\mu} \times \vec{B} \quad (25)$$

The rotation frequency ω_0 of the spinning proton is given by the Larmor equation

$$\omega_0 = \gamma B_0 \quad (26)$$

where γ is a constant related to the proton, in wates it has a value of $\gamma = 2.7 \cdot 10^8 \frac{rad \cdot s}{T}$ and B_0 is the external magnetic field. For a typical scanner with a magnetic field of 2-3 T, Larmor frequency assumes values of $\sim MHz$ just below radio waves. proton spins can assume two values: up and down in relation to the external field direction ; after a body underwent a magnetic field, its total magnetization is given by the net difference between spins up and down This net magnetization causes a flux which can be detected by an external coil only if it is perturbed from its equilibrium state. This is achieved by using another external magnetic field under the form of a radiofrequency impulse, with a frequency resonating with the Larmor frequency of preprocessing. This flips the spin in a direction usually perpendicular to the original B_0 one, which is supposed to be along \hat{z} axis. Thus protons will start rotating in the x-y plane, still continuing preceding around their symmetry axis. Doing so, they gradually lose their initial energy and tend to realign along z-axis. This process is called relaxation and it follows an exponential decay with a time constant T_1 called longitudinal relaxation (longitudinal in relation to the original $B_0\hat{z}$ direction), or spin-lattice relaxation, referring to the loss of thermal energy through interaction with surrounding lattice. The total magnetization along z-axis during time, after the rf impulse, will then regrow along z-axis following:

$$M_z(t) = M_0(1 - e^{-t/T_1}) \quad (27)$$

. An other effect to be take into account is the interaction of spin with a the local field where the external field and the neighbor's one coexist. This spin-spin interaction causes the dephasing of spin. This brings to two different relaxation velocities along z-axis and x-y plane called transversal relaxation which occurs without energy exchange, and it's characterized by a time constant T_2 . The two relaxation processes are described by Bloch equations and once integrated bring to the equation of the evolution of magnetization over time.

$$M_{x,y}(t) = M_{x,y}(0) \cdot e^{-t/T_2} \quad (28)$$

In a real system, however, there's and additional dephasing source, coming from external field inhomogeneities. This effect is often taken into account by the introduction of a different decay time T_2' which along with T_2 bring to a overall time costant given by $\frac{1}{T_2^*} = \frac{1}{T_2} + \frac{1}{T_2'}$. Sometimes, though, this additional T_2' is so small compared to T_2 , that dominates over it, resulting in a rapid loss of information. This can be avoided employing a specific rf pulse sequence called spin-echo method.

A.1 spin-echo method

The spin echo method employs two rf pulses the first with an angle of $\pi/2$ and the second of π .

1. The first radio-frequency pulse is applied with an angle of $\pi/2$ and this produces the spin to flip over a transversal plane and they gradually start dephasing because they undergo small field nonuniformities different from point to point, so they start fanning out, some move faster and some are delayed in relation to the average magnetization vector

2. The second rf impulse is sent, with an angle of π to the transversal magnetization axis y' . This ensures that all the accumulated phases flip and become negative,
3. Late and early phases continue to accumulate delay, but now this delay pushes them towards the main magnetization vector, and the phase will therefore return zero. The elapsed time up to this moment is called *Time to echo* T_E .

A.2 Image acquisition and k-space

For imaging acquisition, both effects must be taken into account. Moreover, it is necessary to relate a signal with a spatial position. In order to do so, in addition to the initial static magnetic field B_0 there's another field with lower intensity than B_0 which is not uniform in intensity, it follows a spatial gradient so that the total magnetic field along \hat{z} -axis is given by the sum of this two contributes, and the signal contains space-varying frequency components according to equation 26 which can be rewritten as $\omega(z) = \gamma B(z)$ being x the spatial coordinate and with B_z now given by the relation $B_z(z, t) = B_0 + z \cdot G(t)$. The physical process underlying signal acquisition is Faraday induction, according to which an electrical potential is related to the variation of a magnetic flux over time $\mathcal{E} = -\frac{d\Phi}{dt}$ being Φ the varying flux through the receiving coil. The varying flux is obtained just after the application of the rf pulse, while tipped spins are precessing and realigning to z -axis, and all this process is called free induction decay (FID). This coil detect only the signal deriving from transversal plane because the one along longitudinal axes would be saturated from the strong magnetic field of the main B_0 coil. Changes in acquisition sequences and rf pulses allow us to emphasise one of the three fundamental parameters T_1, T_2, ρ where ρ is the proton concentration inside the tissue.

An image acquired with a spin echo sequence can be T1, T2 or proton density weighted

The acquired signal is examined by inverse Fourier transform, exploiting the differences from Larmor frequency which cause a phase displacement along z -axis $\phi_G(z, t)$

$$s(t) = \int_0^t dz \rho \quad (9.14) \text{ pag 145} \quad (29)$$

Data are acquired under the form of a matrix called k-space. K-space is a coordinate system used to store spatial frequencies information before applying the inverse fourier transform. Low spatial frequencies, corresponding to large object across the whole real image, are encoded at the center of the matrix and high spatial frequencies corresponding to small objects and finer details, are encoded in the peripheries.

The construction of k-space is done step by step in relation to the gradient applied time by time, producing a trajectory on the k-space. Each spatial line of k-space is acquired after TR, so the signal has to be recreated each time. Starting from $k = 0$, the center of k-space, after a preencoding gradient, we move toward negative position of both k-phase and k-frequency; subsequently a frequency encoding gradient (also called readout gradient) is applied and the signal is collected along the frequency encoding direction. The signal is then reformed and the process is repeated, from $k = 0$, to a new k-phase position and an other line of k-frequency is acquired.

A.3 fMRI

Functional MRI aims to measure brain activity by detecting changes in blood linked to a neuronal activation.

This kind of analysis is referred as blood-oxygen-level dependent (BOLD) imaging.

The process that from a neuronal stimulus leads to a measurable blood signal is governed by the hemodynamic response function (HRF). As shown in figure 11 after a neuronal stimulus on a timescale of ≈ 10 -100 ms, the BOLD signal takes ≈ 6 s to reach a peak and a total of 20-30 s to return to its zero baseline.

Signal comes from changes of blood-oxygen level of hemoglobin: an iron molecule carrying oxygen throughout veins and capillaries. Hemoglobin can be either oxyhemoglobin or deoxyhemoglobin, these two molecules differs in their magnetic susceptibility as oxyhemoglobin is diamagnetic while deoxyhemoglobin is paramagnetic. The presence of deoxy-hemoglobin causes local field inhomogeneities, leading to a reduction of signal from water molecules, because molecules go out of phase with one another more quickly, reducing the total magnetization.

This effect is characterized by T_2^* relaxation time so this is the tuning parameter related to this kind of acquisition; this gives the alternative name of BOLD-images as T_2^* -weighted images.

Three different types of tissues are present in the brain: cerebrospinal fluid (CSF), and gray and white matter.

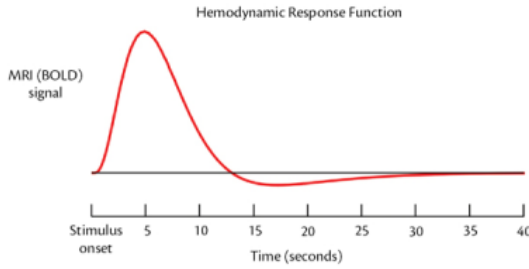


Figure 11

A.3.1 Gradient Echo sequences

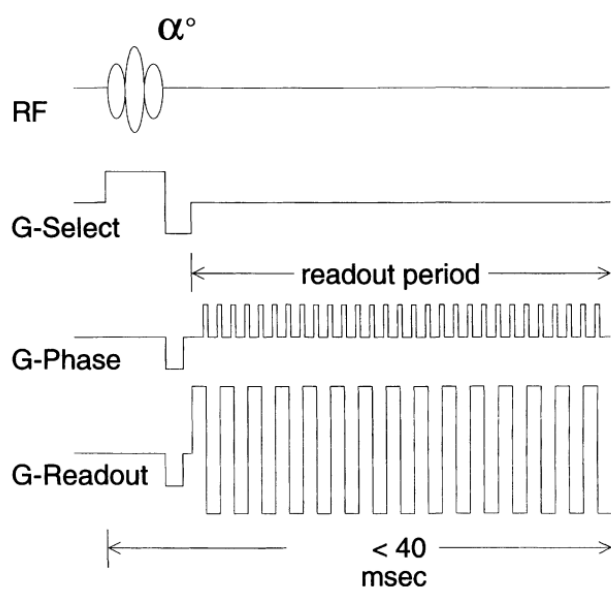
The sequence employed to acquire functional imaging is the Echo Planar Imaging (EPI) which is a particular sequence from the family of gradient echo sequences. Gradient echo sequences differ from spin echo because they allow a faster acquisition time because they use an excitation flip angle less than 90 degrees, this allows a minor time interval to make spins back to their longitudinal component as they are not completely flipped on the transverse plane. The transverse component just created decays and dephases according to T_2^* . If a gradient is applied, they dephase faster, and it is then reversed to make them rephase. In a gradient echo sequence, the echo peak lies on the T_2^* decay curve while in a spin echo, it lies on the T_2 decay curve; and thanks to this it is susceptible to any uniformity variation among which that due to hemoglobin variation. Since TRs are short there is some transversal magnetization left after the acquisition, in *spoiled* Gradient Echo, the residual magnetization is wiped out. Ultra fast gradient echo starts with a 180 degrees pulse, to invert longitudinal magnetization, in order to increase T_1 weighting, and all k-space lines are acquired after this impulse. This first pulse is followed by minor pulses, and k-space is acquired after each echo.

A.3.2 EPI

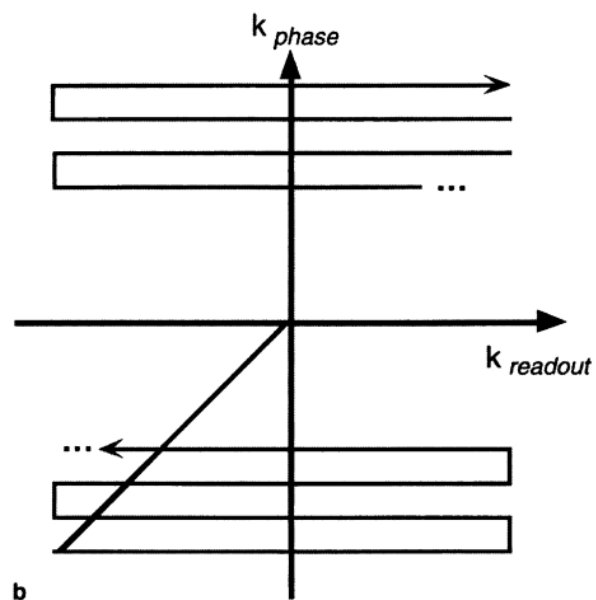
During an EPI session, a single RF impulse of 90 degrees is applied and the whole k-space is acquired in a single acquisition: as shown in figure 12a after the initial RF pulse, together with the gradient for slice selection (G-Select), the acquisition of k-space information starts from the left bottom as in figure ?? and each line along k-frequency corresponding to the readout gradient (G-Readout) is acquired. The period of the G-Readout is the echo time T_E , and it's modulated such that it is sensitive to T_2^* setting $T_E \approx T_2^*$. During the acquisition of the whole k-space each line is shifted from the other along k-phase direction by applying brief pulses of the phase encoding gradient (G-Select) called blip, creating a snake-like path during the acquisition of the k-space matrix.

Pixel-to-pixel frequency difference in the phase-encoding direction emphasizes chemical shift artifact. EPI generates gradient echoes under the FID curve created by the RF flip. As the curve decays according to T_2^* , gradient EPI sequences will be T_2^* weighted. Sensitivity to T_2^* , of course, introduces sensitivity to artifacts caused by changes in magnetic susceptibility (eg, air/tissue interfaces) and imperfect magnet shim.

References



(a)



(b)

Figure 12