



UNIVERSITY OF SÃO PAULO  
RIBEIRÃO PRETO MEDICAL SCHOOL (FMRP)

*DOCTORAL THESIS*

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Bioinformatic tool to integrate and understand aberrant epigenomic and genomic changes associated with cancer

*Methods, development and analysis*

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By

TIAGO CHEDRAOUI SILVA

February 2018



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Bioinformatic tool to integrate and understand aberrant  
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development and analysis

**Tiago Chedraoui Silva**

*Advisor: Prof. Dr. Houtan Noushmehr*

Doctoral dissertation submitted to the *Ribeirão Preto Medical School* (FMRP/USP), in partial fulfillment of the requirements for the degree of the Doctorate Specialization in Normal and Neoplastic Cell Differentiation. *FINAL VERSION.*

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Prof. Dr.: \_\_\_\_\_ Institution: \_\_\_\_\_

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*To my family who always loved me and gave me all the support to achieve my dreams.*



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First of all, I would like to thank all my friends, from my undergraduate degree at UNICAMP, from my double diploma at Telecom ParisTech, from my year of doctorate sandwich at Cedars-Sinai, from the handball teams. I ask my sincere excuses if I have failed them during these last years, but I will eternally thank for all the support, for all the laughter, for all the hugs, for all understanding in those my depressive days.

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*“Our greatest weakness lies in giving up.  
The most certain way to succeed is always  
to try just one more time.”*

*(Thomas A. Edison)*



# Resumo

SILVA, T. C.. **Ferramenta de bioinformática para integrar e compreender as mudanças epigenômicas e genômicas aberrantes associadas com câncer: métodos, desenvolvimento e análise.** 2018. 93 f. Tese Doutorado (Doutorado – Especialização em Diferenciação celular normal e neoplásicas) – Faculdade de medicina de Ribeirão Preto, Ribeirão Preto – SP.

O câncer configura uma das maiores causas de mortalidade no mundo, caracterizando-se como uma doença complexa orquestrada por alterações genômicas e epigenômicas capazes de alterar a expressão gênica e a identidade celular. Nova evidência obtida por meio de um estudo genômico em larga escala e cujos dados encontram-se disponíveis no banco público do TCGA sugere que um em cada dez pacientes portadores de câncer pode ser classificado com maior eficácia tendo como base a taxonomia molecular quando comparada à histologia. Dessa maneira, nós hipotetizamos que o estabelecimento de mapas genômicos exibindo a localização de sítios de ligação de fatores de transcrição combinada à identificação de regiões diferencialmente metiladas e perfis alterados de expressão gênica possa nos auxiliar a caracterizar e explorar, ao nível molecular, fenótipos associados ao câncer.

Avanços tecnológicos e bancos de dados públicos a exemplo do The Cancer Genome Atlas (TCGA), The Encyclopedia of DNA Elements (ENCODE) e o NIH Roadmap Epigenomics Mapping Consortium (Roadmap) têm proporcionado um recurso inestimável para interrogar o (epi)genoma de linhagens de células tumorais em cultura, bem como de tecidos normais e tumorais em alta resolução. Todavia, a informação biológica encontra-se armazenada em diferentes formatos e não há ferramentas computacionais para integrar esses dados, evidenciando um cenário atual que requer, com urgência, o desenvolvimento de ferramentas de bioinformática e softwares capazes de direcionar a solução deste obstáculo. Nesse contexto, o objetivo principal deste estudo consiste em implementar o desenvolvimento de ferramentas de bioinformática, na linguagem de programação R que, ao final do estudo, será submetido à comunidade científica do projeto Bioconductor sob a licença de código aberto GNU GPL versão 3. Além disso, ajudaremos nossos colaboradores com o aperfeiçoamento do ELMER, um pacote R/Bioconductor que identifica elementos reguladores usando dados de expressão gênica, de metilação do DNA e análise de motivo.

Nossa expectativa é que essas ferramentas possam automatizar com acurácia a pesquisa, o download e a análise dos dados (epi)genômicos que se encontram atualmente disponíveis nas bases de dados públicas dos consórcios internacionais TCGA, ENCODE e Roadmap, além de integrá-los facilmente aos dados genômicos e epigenômicos gerados por pesquisadores por meio de experimentos em larga escala. Além disso, realizaremos também o processamento e a análise manual dos dados que serão automatizados pelas ferramentas, visando validar sua capacidade em descobrir assinaturas epigenômicas que possam redefinir subtipos de câncer. Por

fim, as usaremos para investigar as diferenças moleculares entre dois subgrupos de gliomas recentemente descobertos por nosso laboratório.

**Palavras-chave:** Câncer, metilação do DNA, redes reguladoras de genes, melhoradores, interações de cromatina, sitios de ligação do fator de transcrição, epi-genética, ferramentas computacionais.

# Abstract

SILVA, T. C.. **Bioinformatic tool to integrate and understand aberrant epigenomic and genomic changes associated with cancer: Methods, development and analysis.** 2018. 93 f. Doctoral dissertation (Doctorate Candidate – Specialization in Normal and Neoplastic Cell Differentiation) – Ribeirão Preto Medical School (FMRP/USP), Ribeirão Preto – SP.

Cancer, which is one of the major causes of mortality worldwide, is a complex disease orchestrated by aberrant genomic and epigenomic changes that can modify gene regulatory circuits and cellular identity. Emerging evidence obtained through high-throughput genomic data deposited within the public TCGA international consortium suggests that one in ten cancer patients would be more accurately classified by molecular taxonomy versus histology. Therefore, we have hypothesized that the establishment of genome-wide maps of the de novo DNA binding motifs localization coupled with differentially methylated regions and gene expression changes might help to characterize and exploit cancer phenotypes at the molecular level.

Technological advances and public databases like The Cancer Genome Atlas (TCGA), The Encyclopedia of DNA Elements (ENCODE), and The NIH Roadmap Epigenomics Mapping Consortium (roadmap) have provided unprecedented opportunities to interrogate the epigenome of cultured cancer cell lines as well as normal and tumor tissues with high resolution. Markedly however, biological information is stored in different formats and there is no current tool to integrate the data, highlighting an urgent need to develop bioinformatic tools and/or computational softwares to overcome this challenge. In this context, the main purpose of this study is the development of bioinformatics tools in R programming language that will be submitted to the larger open-source Bioconductor community project under the GNU GPL3 (General Public License version 3). Also, we will help our collaborators improve of the R/Bioconductor ELMER package that identifies regulatory enhancers using gene expression, DNA methylation data and motif analysis.

Our expectation is that these tools can effectively automate search, retrieve, and analyze the vast (epi)genomic data currently available from TCGA, ENCODE, and Roadmap, and integrate genomics and epigenomics features with researchers own high-throughput data. Furthermore, we will also navigate through these data manually in order to validate the capacity of these tools in discovering epigenomic signatures able to redefine subtypes of cancer. Finally, we will use them to investigate the molecular differences between two subgroups of gliomas, one of the most aggressive primary brain cancer, recently discovered by our laboratory.

**Key-words:** Cancer, DNA methylation, gene regulatory networks, enhancers, chromatin interactions, transcription factor binding sites, epi-genetics, computational tools.



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# LIST OF ABBREVIATIONS AND ACRONYMS

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5mc .....	5-methylcytosine
ANOVA ..	Analysis of variance
AUC .....	Area Under the Curve
AWGs .....	Analysis working groups
BAM .....	Binary Alignment/Map
CBS .....	Circular Binary Segmentation
CCG .....	Center for Cancer Genomics
ChIP-seq .	ChIP-sequencing
CNP .....	Copy-number polymorphism
CNV .....	copy number variation
COAD ...	Colon adenocarcinoma
DEA .....	differential expression analysis
DEGs .....	differentially expressed genes
DMCs .....	differentially methylated CpGs
DMR .....	Differentially methylated regions
DNA .....	deoxyribonucleic acid
Dnmts .....	DNA methyltransferases
ELMER ..	Enhancer Linking by Methylation/Expression Relationship
ENCODE .	The Encyclopedia of DNA Elements
EPIC .....	MethylationEPIC BeadChip
FC .....	fold change
FDR .....	false discovery rate
FIMO .....	Find Individual Motif Occurrences
FOXA1 ...	Forkhead box protein A1
FPKM .....	Fragments Per Kilobase of transcript per Million mapped reads
FWER ...	Family-wise error rate
GATA3 ...	GATA-binding protein 3
GDC .....	Genomic Data Commons
GNU GPL3	GNU General Public License version 3
GRanges .	GenomicRanges
GRCh36 ..	Genome Reference Consortium Human Build 36

GRCh37 ... Genome Reference Consortium Human Build 37  
GRCh38 ... Genome Reference Consortium Human Build 38  
GSEA ..... gene set enrichment analysis  
GUI ..... Graphical User Interface  
H3K36 .... histone H3 lysine 9  
H3K4me3 H3K4 trimethylation  
H3K9 .... histone H3 lysine 9  
HM450 ... HumanMethylation450 BeadChip  
HOCOMOCO HOmo sapiens COmprehensive MOdel Collection  
HOXD13 . Homeobox D13  
HOXD3 .. Homeobox D3  
KW ..... Kruskal-Wallis test  
LGG ..... lower-grade glioma  
MAE ..... MultiAssayExperiment  
MAF ..... Mutation Annotation Format  
MWU .... Mann-Whitney U-test  
NGS ..... next-generation sequencing  
NHGRI ... National Human Genome Research Institute  
NIH ..... National Institute of Health  
OR ..... Odds Ratio  
P63 ..... tumor protein p63  
PI3K ..... Phosphatidylinositol 3-kinase  
PPIN ..... Protein-protein interaction networks  
PPIs ..... Protein-protein interactions  
PTEN .... Phosphatase and tensin homolog  
PWM .... position weight matrix  
qCML .... quantile-adjusted conditional maximum likelihood  
RAS ..... Rat Sarcoma  
RB ..... retinoblastoma-associated  
ROADMAP The NIH Roadmap Epigenomics Mapping Consortium  
ROC ..... Receiver Operator Characteristics  
RPM ..... Reads per Millions  
SAM ..... S-adenyl methionine  
SE ..... SummarizedExperiment  
SNP ..... single nucleotide polymorphism  
SOX2 .... Sex determining region Y-box 2  
TARGET . Therapeutically Applicable Research to Generate Effective Treatments

TCGA .... The Cancer Genome Atlas  
TF ..... Transcription Factor  
TFBSs ... Transcription factor binding sites  
TSSs .... transcription start sites  
WGBS ... whole-genome bisulfite sequencing



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## LIST OF SYMBOLS

$H_0$  — Null hypothesis

$H_1$  — Alternative hypothesis

$\alpha$  — Significance level

$\beta$  — rate of the type II error

$S(t)$  — Kaplan–Meier estimator



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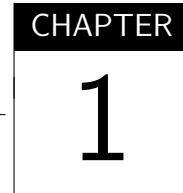
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# INTRODUCTION

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## 1.1 Objectives

The main goal of this project is to develop tools for searching, retrieving and analyzing pan-cancer genomic data from several databases, such as the NCI's Genomic Data Commons (GDC), which contains data from the The Cancer Genome Atlas (TCGA) and Therapeutically Applicable Research to Generate Effective Treatments (TARGET), The Encyclopedia of DNA Elements (ENCODE), and The NIH Roadmap Epigenomics Mapping Consortium (ROADMAP). For a better transparency, all tools will be open source and for their scalability and interoperability they will be published in the Bioconductor project, an environment that provides a broad range of powerful statistical and graphical methods for the analysis of genomic data. Furthermore, We also aim to investigate the intergenic epigenomic changes associated with distinct biological and clinical subgroups of gliomas first discovered by our laboratory (CECCARELLI *et al.*, 2016b). Specifically, using the tools developed, we will integrate DNA methylation, gene expression, mutation and copy number data as well as important epigenomic marks defined by histone modifications in normal samples in order to identify candidate regulatory elements associated with glioma progression.

### 1.1.1 *Specific aims*

1. Download and process transcription factor (TF) ChIP-seq data for each cancer cell and tissue type through the ENCODE dataset;
2. Download and process DNA methylation data for both cancer and non-tumor control cases through the TCGA consortium via HM450K platform;
3. Identify statistically Differentially methylated regions (DMR) at the single CpG resolution;
4. Determine statistically enriched proximal transcription factor binding sites (TFBSs) to altered DNA-methylated regions at the level of individual DNA/protein site interaction;

5. Within known DMRs, classify and identify statistically known and novel DNA binding motifs;
6. Download and process RNA-seq data from both cancer and non-tumor control cases;
7. Use standard data structure to organize the data and the metadata;
8. Correlate the DNA methylation status of Transcription factor binding sites (TFBSs) with target gene RNA-seq expression in order to determine regulatory networks that might alter the pan-cancer genome;
9. Use learning machine algorithms for classifying an independent set of gliomas based on newly identified regulatory networks as related to pan-cancer deregulation;
10. Develop tools to automate the previous steps;
11. Use those tools to investigate the intergenic epigenomic changes associated with distinct biological and clinical subgroups of gliomas g-cimp-low and g-cimp-high discovered by our laboratory and collaborators;
12. Compare the automated results with ones found manually in order to validate the package capacity in providing searching, retrieving and downstream biological analysis to discover pan-cancer epigenomic signatures able to redefine subgroups of gliomas;
13. Submit those set of tools to be freely available in the open-source Bioconductor environment (available at <http://www.bioconductor.org>).

## 1.2 Motivations

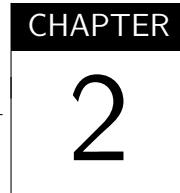
Unravelling the genomic, epigenomic, and proteomic features using high-throughput methodologies is a central question for understanding regulatory gene networks in cancer. In this line of evidence, thousands of tumor and normal samples have been massively sequenced and a large amount of data are publicly deposited by the three main international consortia: TCGA, ENCODE, and Roadmap. However, a major challenge is the fact that the biological information necessary for a complete gene regulatory analysis is spread over different databases that store data in different formats. Moreover, to date there is a lack of computational tools and methods that can integrate and interpret such information. Consequently, the process of analysis is performed manually by the end user, who must access all databases, select and process the data necessary to the project, and integrate that data using multiple downstream analysis tools to extract and interpret the relevant biological information. To overcome these limitations, here we propose to implement tools for searching genomic and epigenomic data acquired from several biological databases, and to provide key scientific analysis steps and methodology, thereby allowing other researchers to apply strategies for in-depth bioinformatics analysis. These tools

will be submitted to Bioconductor, and then, our expectation is that researchers may integrate all relevant data from the most important international consortia in the genomics field with their own experiments. Providing our package through Bioconductor, enables us to access a broader scientific community of advanced informatics users and developers worldwide, who can test the package with their own microarray and next-generation sequencing data, submit bug reports, criticize the methodology, provide new contributions, and ensure the quality of our package in terms of code and documentation. In addition, storing the outputs inside an open-source software like R, allows one to utilize the many available statistical and analytical packages commonly used by researchers (Editorial Nature Genetics, 2014). Then, we expect that these tools will provide insights and novel discoveries into unanticipated regulatory circuits in complex disease and normal developmental biology, which will be verified through the molecular analysis between the newly identified groups G-CIMP-low and G-CIMP-high.

## 1.3 Organization of the Remainder of the Document

This paper is organized into chapters as follows: In Chapter 2, fundamental concepts required to the development of this project are introduced. We review the concepts of cancer and their epigenetic and genetic alterations, followed by the description of the biological data generated through experiments conducted to identify these changes and the data structure used to store it, finally we reviewed some of the data analysis methods used in the genetics field. Chapters 3, 4, and 5 highlights the results of this work. Specifically, in Chapter 3, we detail the methods and computational tools developed, while in Chapter 4 we show their application in the real world. Enclosing this thesis, in Chapter 5, we draw the main conclusions of this work, as well as the scientific contributions derived from this project, possible future works and the main papers that have been published during the Doctorate period.





# REVIEW OF ESSENTIAL CONCEPTS

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This chapter explains some of the fundamental concepts used in the development in this thesis. In section 2.1 the concept of cancer is introduced. The characteristics of cancerous cells compared to normal cells, the molecular mechanisms responsible for their onset and those that facilitate their maintenance and proliferation. In section 2.2 the genetic and epigenetic changes in cancer cells are detailed. In section 2.3 the clinical, genetic and epigenetic data publicly available used for cancer data analysis are described. Their computational representation and data structures available to manipulate large collections of data are detailed. Finally, in section 2.4 several techniques of data analysis used to extract biological knowledge from this data are presented.

## 2.1 Cancer

Tumors are complex tissues composed of multiple distinct cell types that participate in heterotypic interactions with one another. For the onset of tumorigenesis, cancer cells create a "tumor microenvironment" recruiting normal cells which are active participants in tumorigenesis rather than passive bystanders. If compared to normal cells, cancer cells acquired capabilities which includes sustaining proliferative signaling, evading growth suppressors, resisting cell death, enabling replicative immortality, inducing angiogenesis, activating invasion and metastasis, reprogramming of energy metabolism and evading immune destruction (HANAHAN; WEINBERG, 2011). Some examples of these acquired capabilities are described below.

**Sustaining Proliferative Signaling** While normal tissues carefully control the production and release of growth-promoting signals the cancer cells deregulate these signals in ways that promote proliferation. Cancer cells can acquire the capability to sustain proliferative signaling in a number of alternative ways: They may produce growth factor ligands themselves, they may send signals to stimulate normal cells within the supporting tumor-associated stroma, to supply them with various growth factors (CHENG *et al.*, 2008), or they may

elevate the levels of receptor proteins displayed on the cancer cell surface, rendering such cells hyperresponsive to otherwise-limiting amounts of growth factor ligand. Cancer cell also disrupts Negative-Feedback Mechanisms that Attenuate Proliferative Signaling, for example, loss-of-function mutations or promoter methylation of the Phosphatase and tensin homolog (PTEN), a tumor suppressor, amplify Phosphatidylinositol 3-kinase (PI3K) signaling and promote tumorigenesis in a variety of experimental models of cancer (JIANG; LIU, 2009). A second example is the oncogene Rat Sarcoma (RAS), some mutations affecting it compromise Ras GTPase activity, a protein that normally regulates the inactivation of Ras. In addition to the oncogenic mutations in Ras, another mechanism affecting this gene showed silence of the regulatory Ras GTPase proteins through CpG methylation (JIN *et al.*, 2007).

**Evading growth suppressors** Cancer cells must also circumvent actions of tumor suppressor genes that negatively regulate cell proliferation. One example of tumor suppressor proteins is the retinoblastoma-associated (RB), that integrates signals from diverse extracellular and intracellular sources and, in response, decides whether or not a cell should proceed through its growth-and-division cycle (BURKHART; SAGE, 2008). Cancer cells with defects in RB pathway function are thus missing the services of a critical gatekeeper of cell-cycle progression whose absence permits persistent cell proliferation. Another example is the TP53 proteins that receives inputs from stress and abnormality sensors that function within the cell's intracellular operating systems. This protein can activate apoptosis if it received a signal of irreparable damage to the cellular subsystems, or it can interrupt the cell-cycle progression if it receives signals that the degree of damage to the genome is excessive, or if the levels of nucleotide pools, growth-promoting signals, glucose, or oxygenation are suboptimal (HANAHAN; WEINBERG, 2011). Moreover, disruption of such self-attenuating signaling may contribute to the development of adaptive resistance toward drugs targeting mitogenic signaling. To evade those growth suppressors proteins tumor cells developed mechanisms of contact inhibition and its evasion.

**Resisting Cell Death** The loss of TP53 tumor suppressor function, which eliminates this critical damage sensor from the apoptosis-inducing circuitry, is a strategy of tumor cells to limit or circumvent apoptosis (HANAHAN; WEINBERG, 2011).

Those capabilities, which enable tumor growth and metastatic dissemination, were acquired mainly due to a genetic diversity generated by genome instability (HANAHAN; WEINBERG, 2011). A multistep tumor progression can be portrayed as a succession of clonal expansions, which can be triggered by non mutational changes (epigenetic mechanisms such as DNA methylation and histone modifications (BERDASCO; ESTELLER, 2010) or mutational changes affecting the regulation of gene expression.

To increase the rates of mutation, cancer cells often breakdown components of the genomic maintenance machinery that normally monitor genomic integrity and force genetically damaged cells into either senescence or apoptosis (JACKSON; BARTEK, 2009), more specifically those involved in detecting DNA damage and activating the repair machinery, in directly repairing damaged DNA, and in inactivating or intercepting mutagenic molecules before they have damaged the DNA (NEGRINI; GORGOULIS; HALAZONETIS, 2010). Another major source of tumor-associated genomic instability is the loss of telomeric DNA in many tumors that generates karyotypic instability and associated amplification and deletion of chromosomal segments (ARTANDI; DEPINHO, 2009). Recurrence of these amplifications and deletions aberrations at particular sites in the genome indicates that they are likely to harbor genes whose alteration favors neoplastic progression (KORKOLA; GRAY, 2010).

## 2.2 Genetics and epigenetics alterations

While genetics is the study of heritable changes in gene activity or function due to the direct alteration of the DNA sequence such as point mutations, deletions, insertions, and translocation, epigenetics is the study of those not associated with any change of the DNA sequence (MOORE; LE; FAN, 2013). Even though all cells in an organism contain the same genetic information, this epigenetic mechanism controls which gene are expressed enabling the existence of a diversified gene expression profiles in a variety of cells and tissues in multicellular organisms.

### 2.2.1 DNA methylation

DNA methylation is an epigenetic mechanism involving the transfer of a methyl group to the C5 position of the cytosine to form 5-methylcytosine, which regulates gene expression by either blocking the binding of Transcription factors or recruiting proteins involved in gene repression (MOORE; LE; FAN, 2013). The pattern of DNA methylation in the genome can change by DNA methylation and demethylation process, these processes are involved in cell differentiation as unique DNA methylation pattern is able to regulate tissue-specific gene transcription (MOORE; LE; FAN, 2013).

The DNA methylation process is mediated by the family of DNA methyltransferases (Dnmts) proteins which catalyze the transfer of a methyl group from S-adenyl methionine (SAM) to the fifth carbon of cytosine residue to form 5-methylcytosine (5mc). The main proteins responsible for this transfer are the Dnmt3a and Dnmt3b, which are involved in establishing a new methylation pattern to unmodified DNA, and Dnmt1, which is responsible to the methylation of a daughter strand during the DNA replication process.

The majority of DNA methylation occurs on cytosines that precede a guanine nucleotide (CpG sites), however it has already been reported a significant percentage of methylated non-CpG

sites (XIE *et al.*, 2012).

The role of DNA methylation varies in different genomic regions. Within intergenic regions, the DNA methylation represses the expression of potentially harmful genetic elements, while within promoter regions containing CpG islands, stretches of DNA roughly 1000 base pairs long that have a higher CpG density than the rest of the genome and often are not methylated (BIRD *et al.*, 1985), results in stable silencing of gene expression (MOHN *et al.*, 2008). Also, within the gene body, region of the gene past the first exon, the methylation of the first exon leads to gene silencing (BRENET *et al.*, 2011).

## 2.2.2 Histone modifications

This process of transcription regulation involving DNA methylation, works in association with histone modifications and noncoding RNAs.

Normally, the DNA is wrapped around the histone proteins forming small, packaged sections called nucleosomes. This association is capable of inhibiting gene expression since a more compact region hinders the accessibility of transcription factors.

Those histones might have chemical modifications (methylation, acetylation, ubiquitination, and phosphorylation) added to three specific amino acids on their N-terminal tail which influences how DNA strands are packaged and their transcriptional activity. Those modifications that loosen DNA association with histones generally provide a permissive environment for transcription, whereas the ones that tighten them repress gene expression (MOORE; LE; FAN, 2013).

These modifications involved in adding and/or stripping histone markers in order to impose a repressive state on a gene region, are led by histone-modifying enzymes and are in general in cooperation with the Dnmts proteins which are involved in the mechanism of DNA methylation. For example, Dnmt1 and Dnmt3 are known to bind to the histone methyltransferase SUV39H1 that restricts gene expression by methylation on histone H3 lysine 9 (H3K9) (FUKS *et al.*, 2003). Other examples are Dnmt1 and Dnmt3b that can both bind to histone deacetylases that remove acetylation from histones to make DNA pack more tightly and restrict access for transcription (FUKS *et al.*, 2000; GEIMAN *et al.*, 2004). Also, histone H3 lysine 9 (H3K36) trimethylation, a repressive histone mark, stimulates the methyltransferase activity, the H3K4 trimethylation (H3K4me3) prevents it (OOI *et al.*, 2007; ZHANG *et al.*, 2010).

The miRNAs have emerged as another important epigenetic mechanism that influences gene expression. It can repress gene expression by inhibiting translation or inducing RNA degradation (BEREZIKOV, 2011) and it not only can regulate histone modifications and Dnmt expression thus regulating DNA methylation (BENETTI *et al.*, 2008; SINKKONEN *et al.*, 2008), but also the DNA methylation can regulate the expression of miRNAs (HAN *et al.*, 2007; LUJAMBIO *et al.*, 2008).

Overall, miRNAs, DNA methylation, and histone modifications work closely together to regulate gene expression (MOORE; LE; FAN, 2013).

### **2.2.3 *Genetics alterations***

Although the haploid human genome consists of 3 billion nucleotides, changes in even a single base pair might result in dramatic physiological malfunctions. A mutation, defined as any alteration in the DNA sequence, can be a germ-line mutation, which occurs in a germ-line cell (one that will give rise to gametes) and can be passed to a descendant of the organism, and somatic mutations, which occurs in a somatic cell (one that develops into the body tissues) and are never transmit to their descendants (CLANCY, 2008).

A single base pair alteration that is present in at least 1% of the population is called single nucleotide polymorphism (SNP) and is generally used to refer to a normal variation (does not directly cause disease) (CLANCY, 2008).

An example of a point mutations that is able to alter proteins is the insertion or the deletion of a single base, called frameshift mutations. This change of nucleotides alters the codons (a group of three nucleotides) read by the ribosomes, which affects the protein resulted during the translation process. Table 1 shows a classification of translational effect of variant allele (INSTITUTE, 2017; GDC, 2017).

Table 1 – Translational effect of variant allele. (INSTITUTE, 2017)

<b>Variant_Classification</b>	<b>Group</b>
Frame_Shift_Del	Deletion that moves the coding sequence out of frame
Frame_Shift_Ins	Insertion that moves the coding sequence out of frame
Missense_Mutation	The point mutation alters the protein structure by one amino acid
Nonsense_Mutation	A premature stop codon is created by the variant
Silent	Variant is in coding region of the chosen transcript, but protein structure is identical
Splice_Site	The variant is within two bases of a splice site.
In_Frame_Del	Deletion that keeps the sequence in frame
In_Frame_Ins	Insertion that keeps the sequence in frame
Translation_Start_Site	Point mutation, insertion or deletion that overlaps the start codon
Nonstop_Mutation	variant removes stop codon.
3'UTR	The variant is on the 3'UTR for the chosen transcript
3'Flank	The variant is downstream of the chosen transcript (within 3kb)
5'UTR	The variant is on the 5'UTR for the chosen transcript
5'Flank	The variant is upstream of the chosen transcript (within 3kb)
IGR	Intergenic region. Does not overlap any transcript.
Intron	The variant lies between exons within the bounds of the chosen transcript.
De_novo_Start_InFrame	New start codon is created in frame relative to the coded protein.
De_novo_Start_OutOfFrame	New start codon is created out of frame relative to the coded protein.
RNA	The variant lies on one of the RNA transcripts.
lincRNA	The variant lies on one of the lincRNAs.

Not only changes in DNA can occur on a single nucleotide but they can also occur at the level of the chromosome, in which large segments of chromosomes are altered (deleted, duplicated, inverted, translocated to different chromosomes, rearranged) resulting in changes in the level of gene expression, the complete absence of genes, or the alteration of gene sequence. Feuk, Carson and Scherer (2006) define structural variants as genomic alterations that involve segments of DNA that are larger than  $1kb$  which can be microscopic ( $> 3Mb$ ) or submicroscopic ( $1kb$  to  $3Mb$ ) variants, and also define smaller ( $< 1kb$ ) variations or polymorphisms that involve the copy-number change of a segment of DNA as insertions or deletions (indels). A type of structural variant is the copy number variation (CNV) which refers to a segment of DNA that is  $1kb$  or larger and is present at a variable copy number in comparison with a reference genome, while Copy-number polymorphism (CNP) are CNVs that occurs in more than 1% of the population. Also, it is common to refer to changes in copy number that have arisen in somatic tissue as copy number alterations/aberrations (CNAs or SCNAs) and changes in copy number in germline cells as copy number variations (CNVs). Table 2 summarizes the types of mutations.

Table 2 – Types of DNA Mutations and Their Impact (COMISC, 2017; CLANCY, 2008; FEUK; CARSON; SCHERER, 2006)

<b>Class of Mutation</b>	<b>Type of Mutation</b>	<b>Description</b>
Point mutation	Substitution	One base is incorrectly added during replication and replaces the pair in the corresponding position on the complementary strand.
	Insertion	One or more extra nucleotides are inserted into replicating DNA, often resulting in a frameshift.
	Deletion	One or more nucleotides is "skipped" during replication or otherwise excised, often resulting in a frameshift.
Chromosomal mutation	Inversion	A segment of DNA that is reversed in orientation with respect to the rest of the chromosome.
	Deletion	A region of a chromosome is lost, resulting in the absence of all the genes in that area.
	Duplication or low-copy repeat	A segment of DNA < 1kb in size that occurs in two or more copies per haploid genome, with the different copies sharing > 90% sequence identity.
	Translocation	A change in position of a chromosomal segment within a genome that involves no change to the total DNA content. Translocations can be intra- or inter- chromosomal.
Copy number variation	Gain/Amplification	A single-copy gain or a multi-copy amplification of the entire gene.
	Loss/Deletion	A loss for part or all of a coding region within the gene footprint in a single allele/copy (hemizygous deletion) or deletion for part or all of a coding region within the gene footprint in both alleles/copies (homozygous deletion)(COMISC, 2017) .

## 2.3 Databases and data structure

Recent technological developments allowed the deposition of large amounts of genomic and epigenomic data, such as gene expression, DNA methylation, and genomic localization of transcription factors, into freely available public international consortia like The Cancer Genome Atlas (TCGA), The NCI's Genomic Data Commons (GDC), The Encyclopedia of DNA Elements (ENCODE), and The NIH Roadmap Epigenomics Mapping Consortium (Roadmap) (HAWKINS; HON; REN, 2010). Subsection 2.3.1 presents an overview of those consortia and subsection 2.3.2 presents some of the computational data formats and data structures used to represent the biological data.

### 2.3.1 Databases

**The Cancer Genome Atlas (TCGA):** The TCGA consortium, a National Institute of Health (NIH) initiative, made publicly available molecular and clinical information for more than 30 types of human cancers including exome (variant analysis), single nucleotide polymorphism (SNP), DNA methylation, transcriptome (mRNA), microRNA (miRNA) and proteome. Sample types available at TCGA are primary solid tumors, recurrent solid tumors, blood-derived normal and tumor, metastatic, and solid tissue normal (WEINSTEIN *et al.*, 2013). This project ended in mid-2016 and its data was transferred to the NCI's Genomic Data Commons (GDC).

**The Genomic Data Commons (GDC):** The NCI's Genomic Data Commons (GDC) provides the cancer research community with a unified data repository that enables data sharing across cancer genomic studies in support of precision medicine. It supports numerous cancer genome programs at the NCI Center for Cancer Genomics (CCG), including The Cancer Genome Atlas (TCGA) and Therapeutically Applicable Research to Generate Effective Treatments (TARGET) (NCI, 2017d). Those data is harmonized against GRCh38 (hg38) using GDC Bioinformatics Pipelines which provides methods to the standardization of biospecimen and clinical data, the re-alignment of DNA and RNA sequence data against a common reference genome build GRCh38, and the generation of derived data. Also, this repository has a Legacy Archive to provide access to a unmodified copy of data that was previously stored in CGHub (WILKS *et al.*, 2014) and in the TCGA Data Portal hosted by the TCGA Data Coordinating Center (DCC), whose data were aligned to the reference genomes GRCh37 (hg19) and GRCh36 (hg18).

**The Encyclopedia of DNA Elements (ENCODE):** Founded in 2003 by the National Human Genome Research Institute (NHGRI), the project aims to build a comprehensive list of functional elements that have an active role in the genome, including regulatory elements that govern gene expression. Biosamples include immortalized cell lines, tissues, primary cells and stem cells (CONSORTIUM *et al.*, 2011).

**The NIH Roadmap Epigenomics Mapping Consortium:** Roadmap was launched with the goal of producing a public resource of human epigenomic data to catalyze basic biology and disease-oriented research (BERNSTEIN *et al.*, 2010). Roadmap maps DNA methylation, histone modifications, chromatin accessibility, and small RNA transcripts in stem cells and primary ex vivo tissues (FINGERMAN *et al.*, 2011; BERNSTEIN *et al.*, 2010).

Briefly, these consortia provide large-scale epigenomic data onto a variety of microarrays and next-generation sequencing (NGS) platforms. Each consortium encompasses specific types of biological information on specific type of tissue or cell and, when analyzed together, it provides an invaluable opportunity for research laboratories to better understand the developmental progression of normal cells to cancer state at the molecular level and importantly, correlate these phenotypes with tissue of origins.

### **2.3.2 Data and data structure**

If previously the main bottleneck to scientific progress in cellular biology was data collection, now this bottleneck shifted to analysis of data (MCPHERSON, 2009). The large-scale genomic data mining, which is the process of using many diverse datasets to address a specific biological question, involves three main tasks: establishing methodology for efficiently querying large data collections; assembling data from appropriate repositories, and integrating information from a variety of experimental data types (HUTTENOWER; HOFMANN, 2010). In data science, raw data refers to data that have not been changed since acquisition and after its editing, cleaning or modification results in processed data. In genomics the raw data, sequencing read data, from next-generation sequencing machines is stored in a FASTQ, a common file format data that combines both the sequence and an associated per base quality score (COCK *et al.*, 2009). Those stored reads are then aligned to a reference genome sequence with software like Bowtie 2 (LANGMEAD; SALZBERG, 2012), generating SAM files, the accepted standard for storing short read alignment data, which are subsequently compressed to binary format (Binary Alignment/Map (BAM)) via SAMtools (LI *et al.*, 2009). It is common to consider this set of SAM, BAM, and FASTQ files as raw data. Some databases classify this raw data as level 1, protected level or raw level. It is important to highlight that due to privacy concerns, the access to this raw data is controlled because it could be used in identity tracing attacks (ERLICH; NARAYANAN, 2014; AYDAY *et al.*, 2014) as generally includes individually identifiable data such as low-level genomic sequencing data, germline variants, SNP6 genotype data, and certain clinical data elements (NCI, 2017e). This raw data are processed through bioinformatics pipelines generating processed data files or matrices (usually in the form of tab-delimited text files). Some databases classify this processed data as level 2 or higher or as open. For example, the previously stored data in CGHub, TCGA Data Portal and Broad Institute's GDAC Firehose, were provided as different levels or tiers that were defined in terms of a specific combination of both processing level (raw, normalized, integrated) and access level (controlled or open access).

Level 1 indicated raw and controlled data, level 2 indicated processed and controlled data, level 3 indicated Segmented or Interpreted Data and open access and level 4 indicated region of interest and open access data.

In the next subsections, we will focus on the processed data and their data structures used in this project.

### 2.3.2.1 Data

**Array-based DNA methylation data:** DNA methylation data files contain information on raw and normalized signal intensities, detection confidence and calculated beta values for methylated and unmethylated probes. The processed data from DNA methylation array-based platforms such as GoldenGate, Infinium Human Methylation27, and the Infinium HD 450K methylation array, are presented in the form of beta-values that uses a scale ranging from 0.0 (probes completely unmethylated) up to 1.0 (probes completely methylated). In each sample file, each row represents a 1 bp region (probe) and its correspondent beta-value. Also, probes overlapping with SNPs are normally masked as they are based largely on ad hoc assumptions and subjective criteria (ZHOU; LAIRD; SHEN, 2016).

**Somatic Variant data:** The DNA-Seq Somatic Variant Analysis algorithms, such as MuSE (FAN *et al.*, 2016), Mutect2 (CIBULSKIS *et al.*, 2013), SomaticSniper (LARSON *et al.*, 2011) and Varscan2 (KOBOLDT *et al.*, 2012), identify and characterizes somatic mutations by comparing reference alignments from tumor and normal samples from the same case. The results of the variant calls are stored in a Mutation Annotation Format (MAF), which are filtered to remove any potentially erroneous or germline variant calls. Each row of these represents a mutation in the genome of a sample (NCI, 2017a).

**RNA-Seq Gene Expression data:** The processed data contains read counts measured on a gene level, which might be normalized using some methods such as the Fragments Per Kilobase of transcript per Million mapped reads (FPKM) (NCI, 2017b) and FPKM Upper Quartile (FPKM-UQ) (NCI, 2017c).

**miRNA-Seq data:** The processed data contains expression levels measured and normalized post-alignment. Normalization is performed using the Reads per Millions (RPM) method. Expression levels of known miRNAs and observed miRNA isoforms are generated for each sample.

**Copy Number Variation data:** The processed data is a table that associates contiguous chromosomal segments with genomic coordinates, mean array intensity (log2 ratio segment means), and the number of probes that binds to each segment. This information might be masked to remove segments with probes known to contain germline mutations. From the Circular Binary Segmentation (CBS) methods output described previously, algorithms

Table 3 – Example TCGA barcode

TCGA	02	0001	01	C	01	D	0182	01
Project	Tissue source site	Participant	sample	vial	portion	analyte	place	center

such as GISTIC2 (MERMEL *et al.*, 2011) are used to identify significantly altered regions of amplification or deletion across sets of patients.

**Histone ChIP-Seq data:** The processed data contains either the fold change over control and signal p-value stored in a bigWig format or as peaks in a bed and bigBed format (ENCODE, 2017). The ENCODE consortium highlight that these peak files are not meant to be interpreted as definitive binding events, but are rather intended to be used as input for subsequent statistical comparison of replicates (ENCODE, 2017). This data is used as input for software to infer chromatin-state such as ChromHMM (ERNST; KELLIS, 2012) and StatePaintR (COETZEE *et al.*, 2017).

To identify each of these samples, each database uses a unique identifier, which is able to help users to find a certain sample in the database or to find associated information. The TCGA database, for example, created an ID in two forms, a human-readable version called barcode and a non-human readable called UUID. A TCGA barcode is composed of 28 alphanumeric elements, representing the sample information. How those elements are grouped and which information they have is shown in table 3, a code table for each field value is found in the GDC website (NCI, 2017f). The barcode identifies if a sample is a primary solid tumor sample or a normal tissue sample, but if there was a mistake in the metadata and the sample which was normal was set to a tumor sample, the barcode is then invalid. Due to this issue, the TCGA UUID, composed of 32 randomly generated digits, was created. The TCGA barcode was the primary identifier of biospecimen data since the pilot project began. However, since for any one sample, the barcode can change as the meta-data associated with it changes, the TCGA project transitioned to using UUIDs as the primary identifier.

### 2.3.2.2 Data structures

Although there exists a wealth of possibilities (KANNAN *et al.*, 2015) in accessing cancer associated data, Bioconductor (<https://www.bioconductor.org/>) represent the most comprehensive set of open source, updated and integrated professional tools for the statistical analysis of large-scale genomic data. It uses the R statistical programming language and has been developing several data structures to handle genetics, epigenetics and clinical data. Some of the main data structures from the R/Bioconductor used in this project are described below.

**DataFrame:** A data frame is used for storing data tables. It is a list of vectors of equal length.

**GenomicRanges (GRanges):** This data structure was created to represent and manipulate genomic intervals and variables defined along a genome. GRanges is a vector of genomic locations, which are composed of the sequence names (chr1, chr2), the sequence range (start and end) and the strand information, and associated annotations (metadata) (LAWRENCE *et al.*, 2013). An example of GRanges object is shown in Figure 1.

```
GRanges object with 10 ranges and 2 metadata columns:
  seqnames      ranges strand |      score          GC
  <Rle>    <IRanges>  <Rle> | <integer>      <numeric>
  a      chr1 [101, 111]     - |      1              1
  b      chr2 [102, 112]     + |      2 0.8888888888888889
  c      chr2 [103, 113]     + |      3 0.7777777777777778
  .      ...   ...   . | ...
  h      chr3 [108, 118]     + |      8 0.2222222222222222
  i      chr3 [109, 119]     - |      9 0.1111111111111111
  j      chr3 [110, 120]     - |     10              0
  -----
  seqinfo: 3 sequences from an unspecified genome; no seqlengths
```

Figure 1 – Example of a Granges object.

**SummarizedExperiment (SE):** This data structure, exemplified in Figure 2, is a matrix-like container where rows represent ranges of interest (as a GRanges object) and columns represent samples (with sample data summarized as a DataFrame). It can contain one or more assays, each represented by a matrix-like object of numeric or other mode.

**MultiAssayExperiment (MAE):** This data structure is an integrative environment where multiple assays are managed and preprocessed for genomic data analysis. It is composed of at least two metadata matrices, one with the phenotype metadata (i.e. clinical data) and one table mapping each data column from different assays to an entry in the phenotype metadata. The data stored by this data structure can be several SummarizedExperiment objects (i.e DNA methylation, gene expression, copy number, histone modification signals).

## 2.4 (Epi)Genomic data analysis

### 2.4.1 Integrative analysis

Often the onset and progression of cancerous diseases are linked to the aberrant function of proteins and alterations in gene expression, which has led research in genetics in search of the molecular alterations responsible for such aberrant behavior. Among the types of alterations genetic and epigenetic that can impact gene function are, gene copy number (CN), DNA methylation, single nucleotide variations (SNV), and indels (small insertions and deletions). These

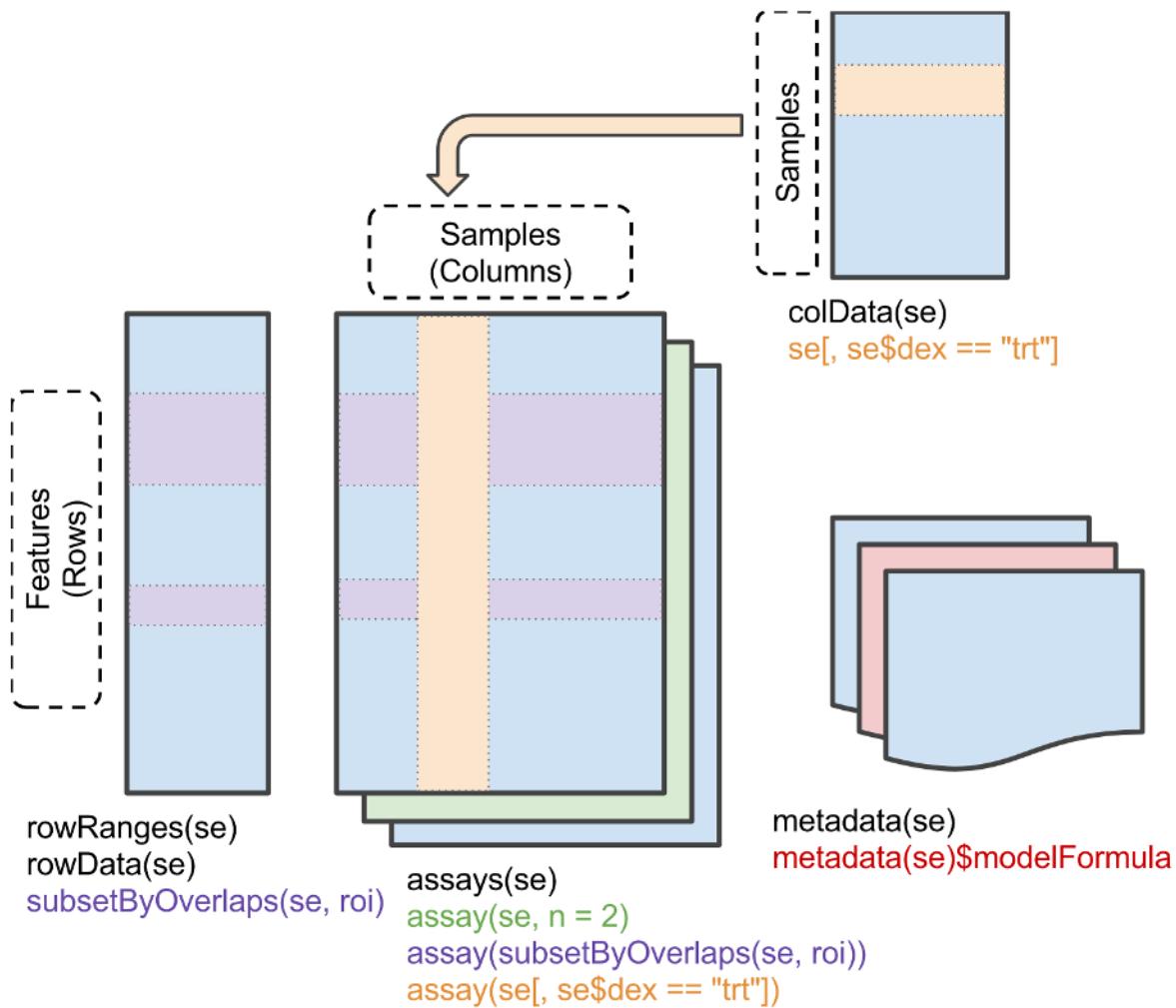


Figure 2 – Example of Summarized Experiment object. Figure reproduced from *SummarizedExperiment* manual (MORGAN *et al.*, 2017).

variations can have a direct effect by modifying the function of the gene product, for example, an indel in a coding region, or an indirect effect such as the modification of regulatory regions which can interfere with gene expression by inhibiting transcription (THINGHOLM *et al.*, 2016).

Furthermore, recent technological developments have enabled the creation of genome-wide data for multiple types of variations. Although individual analyzes of these variations helped increasing knowledge of the genome and of complex disorders, integrative analyses that evaluate cancer transcriptome data in the context of other data sources are often capable of extracting deeper biological insight from the data (RHODES; CHINNAIYAN, 2005).

In this section we will highlight several integrative approaches, including meta-analysis for extracting robust profiles from independent data sets, enrichment analysis for identifying coordinately regulated functional gene modules, protein interaction networks for detecting interaction complexes deregulated in cancer, transcriptional networks for inferring regulatory mechanisms in cancer and analyses of model system profiles with human tumor profiles for inferring activity of oncogenic pathways.

#### 2.4.1.1 Functional enrichment analysis of cancer signatures

A gene set enrichment analysis (GSEA) or functional enrichment analysis is a method to identify classes of genes or proteins that are over-represented in a large set of genes or proteins and may have an association with disease phenotypes. This type of analysis integrates differentially expressed genes identified by a differential expression analysis (DEA) with external functional information which is necessary for interpreting and summarizing large cancer signatures. Most approaches use external annotation databases such as Gene Ontology which is a database of controlled vocabulary gene annotations describing the biological processes, molecular functions and cellular localizations of genes as a resource for enrichment analysis in cancer signatures.

Let  $L$  be a list of genes ranked by their differential expression between the classes, the goal of GSEA is to determine whether members of a gene set  $S$  tend to occur toward the top (or bottom) this list, which might correlate with the phenotypic class distinction (SUBRAMANIAN *et al.*, 2005).

#### 2.4.1.2 Protein interaction networks and cancer signatures

A major objective of systems biology is to organize molecular interactions as networks (VINAYAGAM *et al.*, 2014). Protein-protein interactions (PPIs) are essential to almost every process in a cell and are not only crucial for understanding cell physiology in normal and disease states, since the disruption of protein-protein interactions may result in disruption of the cell component or process to which they contribute compromising the cell viability or even leading to cell death, but also for the drug development, since drugs can affect PPIs (EMBL-EBI, 2017; ALZATE, 2009). These interactions are represented as Protein-protein interaction networks (PPIN), a mathematical representation of the physical contacts between proteins in the cell. The totality of PPIs that happen in a cell, an organism or a specific biological context is called interactome.

A detailed human interactome network that captures the entire cellular network is invaluable in interpreting cancer signatures, allowing one to infer activated subnetworks and specific proteins that are most important to a subnetwork. For example, for a given set of overexpressed proteins in the same subnetwork, one specific protein that interacts with the entire subnetwork might be the key to control the expression the entire subnetwork.

A lot of this information is available through molecular interaction databases such as IntAct (<http://www.ebi.ac.uk/intact>) (ORCHARD *et al.*, 2013).

#### 2.4.1.3 Transcriptional targets and cancer signatures

Similar to the Protein-protein interaction networks, global transcriptional networks, which defines directional pathways (i.e. which gene actives are activated by a given gene), have the potential to improve the interpretation of cancer signatures. For example, if the targets of all

transcription factors were known, then one could easily infer which transcription factors must be activated in a tumor to yield the observed cancer signature. With that information, it is possible to reduce a complex cancer signature to a small number of activated transcriptional factors which will be potential therapeutic targets.

The identification of transcription factor-binding sites (TFBSs) is done using high throughput experimental methods such as ChIP-Chip and ChIP-sequencing (ChIP-seq) which identifies a region of 100–1000 base pairs (b.p.) in which the TFBS (typically 9-15 b.p.) resides (JAYARAM; USVYAT; MARTIN, 2016). In-silico sequence-based methods were developed to predict TFBSs. These methods scan a DNA sequence of interest with a position weight matrix (PWM), a  $4 \times n$  matrix of scores for each of the 4 bases across each position in the binding motif, for a transcription factor of interest and perform a pattern-matching. PWM models can be obtained from a number of resources including the open access database JASPAR (PORTALES-CASAMAR *et al.*, 2009), HOCOMOCO (KULAKOVSKIY *et al.*, 2013) and HOMER (HEINZ *et al.*, 2010). Among the existing software that scans a sequence database for individual matches to each of the motifs are Find Individual Motif Occurrences (FIMO) (GRANT; BAILEY; NOBLE, 2011), HOMER (HEINZ *et al.*, 2010) and PATSER (TURATSINZE *et al.*, 2008).

#### 2.4.1.4 Human Epigenomes

The human body contains more than 200 different cell types each one has an identical copy of the genome but expresses a distinct set of genes, due to their epigenome which in each cell regulates gene expression in a number of ways - by organizing the nuclear architecture of the chromosomes, restricting or facilitating transcription factor access to DNA, and preserving a memory of past transcriptional activities (RIVERA; REN, 2013). The integrative analysis of epigenomic maps, which references to the collection of DNA methylation state and covalent modification of histone proteins along the genome (BONASIO; TU; REINBERG, 2010), has been shown to be important in the study of the gene regulatory programs (GIFFORD *et al.*, 2013; HAWKINS *et al.*, 2010; RADA-IGLESIAS *et al.*, 2012). Table 4 shows a summary of the epigenomic marks and their associate role.

Table 4 – Core set of five histone modification marks and other epigenomic marks

Histone marks	Role
Histone H3 lysine 4 trimethylation (H3K4me3)	Promoter regions (HEINTZMAN <i>et al.</i> , 2007; BERNSTEIN <i>et al.</i> , 2005)
Histone H3 lysine 4 monomethylation (H3K4me1)	Enhancer regions (HEINTZMAN <i>et al.</i> , 2007)
Histone H3 lysine 36 trimethylation (H3K36me3)	Transcribed regions
Histone H3 lysine 27 trimethylation (H3K27me3)	Polycomb repression (BONASIO; TU; REINBERG, 2010)
Histone H3 lysine 9 trimethylation (H3K9me3)	Heterochromatin regions (PETERS <i>et al.</i> , 2003)
Histone H3 acetylated at lysine 27 (H3K27ac)	Increase activation of genomic elements (HEINTZMAN <i>et al.</i> , 2009; RADA-IGLESIAS <i>et al.</i> , 2011; CREYGHTON <i>et al.</i> , 2010)
Histone H3 lysine 9 acetylation (H3K9ac)	Transcriptional activation (NISHIDA <i>et al.</i> , 2006)
DNase hypersensitivity	Regions of accessible chromatin (THURMAN <i>et al.</i> , 2012)
DNA methylation	Repressed regulatory regions (CEDAR; BERGMAN, 2009; MOORE; LE; FAN, 2013)

Using these set of epigenetic marks, it is possible to discover epigenomic states which would aid in understanding “non-coding” genomic elements (COETZEE *et al.*, 2017). Several tools can use those marks to infer chromatin-state such as ChromHMM (ERNST; KELLIS, 2012) and StatePaintR (COETZEE *et al.*, 2017). A 15-state model created from ROADMAP data using ChromHMM is presented as follows: active TSS-proximal promoter states (TssA, TssAFlnk), a transcribed state at the 5' and 3' end of genes showing both promoter and enhancer signatures (TxFlnk), actively-transcribed states (Tx, TxWk), enhancer states (Enh, EnhG), and a state associated with zinc finger protein genes (ZNF/Rpts). The inactive states consist of constitutive heterochromatin (Het), bivalent regulatory states (TssBiv, BivFlnk, EnhBiv), repressed Polycomb states (ReprPC, ReprPCWk), and a quiescent state (Quies) (KUNDAJE *et al.*, 2015). Those precomputed chromatin-state segmentation from public databases such as Blueprint, ENCODE and ROADMAP using ChromHMM (<http://compbio.mit.edu/ChromHMM/>) are available at [http://egg2.wustl.edu/roadmap/web\\_portal/chr\\_state\\_learning.html#core\\_15state](http://egg2.wustl.edu/roadmap/web_portal/chr_state_learning.html#core_15state) and using StatePaintR are available at [www.statehub.org](http://www.statehub.org).

These data are extremely useful to characterize each different region of the genome. For example, using those information Kundaje *et al.* (2015) were able to identify enhancer regions containing strong H3K27ac signal which showed a higher DNA accessibility, lower methylation, and higher TF binding than enhancers regions lacking H3K27ac. Those regions with the strong H3K27ac signal are more likely to have a regulatory role.

## 2.4.2 Statistical analysis

### 2.4.2.1 Hypothesis Testing

Human genetic studies aim to identify if a phenotype is related to the genotypes at various loci, that is, if genetic variations have an influence on the risk of disease or other health-related phenotypes. Statistical analysis is a crucial to present the findings in an interpretable and objective manner (SHAM; PURCELL, 2014).

The most popular hypothesis testing approach used to test if genotypes and phenotypes are related is the frequentist significance testing approach. This is a classical approach that involves setting up two competing hypothesis: a null hypothesis ( $H_0$ ) and an alternative hypothesis ( $H_1$ ).

Computing the statistical significance can be done using a one-tailed or a two-tailed test. A two-sided test is appropriate to evaluate both directions of the test, for example, is the estimated value smaller or higher than the reference, which actually tests if the estimated value is different from the reference. A one-sided test is appropriate to evaluate only one direction of the test, for example, is the estimated p-value smaller than the reference. An example in genetic studies for a two-sided test would be  $H_0$  hypothesis that genotypes have no effect on the phenotypes while the  $H_1$  hypothesis is that there is an effect. Table 5 shows other examples.

Table 5 – Example of three hypothesis tests about the population mean  $\mu$ . In genetics it could be the mean level of expression of a gene.

Type	Null	Alternative
Right-tailed	$H_0 : \mu = 0$	$H_1 : \mu > 0$
Left-tailed	$H_0 : \mu = 0$	$H_1 : \mu < 0$
Two-tailed	$H_0 : \mu = 0$	$H_1 : \mu \neq 0$

Table 6 – Type I and II Errors.  $\alpha = P(\text{Type I Error})$ ,  $\beta = P(\text{Type II Error})$

Decision	$H_0$ is True	$H_0$ is False
Do Not Reject $H_0$	Correct Decision	Incorrect Decision ( $1 - \beta$ )
Reject $H_0$	Incorrect Decision ( $1 - \alpha$ )	Correct Decision

#### 2.4.2.2 Making a decision: P-value approach

The decision to reject or accept  $H_0$  is made based on the calculation of a test statistic (T) from the observed data. As the value of T depends on particular individuals in the population, repeating the study using different random samples from the population would provide of many different values for T. These set of T can be summarized as a probability distribution.

Even though, the decision made to reject or accept  $H_0$  just state that we had enough evidence to behave one way or the other. The rejection of the null hypothesis does not prove that the alternative hypothesis is true as the acceptance the null hypothesis does not prove that the null hypothesis is true. It might happen that null hypothesis was rejected when it was true, or it was not rejected when it was false. The first error in statistics is called a Type I error ("false positive"), while the second is called a Type II error ("false negative"). Table 6 shows the relations between truth/falseness of the null hypothesis and outcomes of the test.

The rate of the type I error or significance level  $\alpha$  is the probability of having a false positive. Normally, the significance level is set to 5%, implying that it is acceptable to have a 5% probability of incorrectly rejecting the null hypothesis. With the same logic, the rate of the type II error is denoted by  $\beta$ .

To make a decision whether to reject or accept the null hypothesis, the concept of probability value was introduced. A p-value ( $p\text{-value} \in [0, 1]$ ) is the probability under a specified statistical model, constructed under a set of assumptions (normally "null hypothesis") that a statistical summary of the data (e.g., the sample mean difference between two compared groups) would be equal to or more extreme than its observed value (WASSERSTEIN; LAZAR, 2016). That means, the smaller the p-value, the greater the statistical incompatibility of the data with the null hypothesis and greater the p-value more compatible is the data with the null hypothesis. In summary, if P-value is small (e.g.  $P\text{-value} \leq \alpha$ ) then the null hypothesis is rejected, otherwise, it is not rejected.

It is important to highlight that a p-value does not measure the size of an effect or the importance of a result. It might happen that a very small effect produces smaller p-values if the sample size is big or measurement precision is high. On the other hand, a large effect might produce higher p-values if the sample size is small or measurements are imprecise.

#### 2.4.2.3 Correcting for multiple testing

When performing a set of statistical inferences simultaneously more likely erroneous inferences are to occur. For example, if 100 tests are carried out, then 5% of them (that is 5 tests) are expected to have  $P - value < 0.05$  by chance when  $H_0$  is, in fact, true for all the tests. Compared to a single test (equations 2.1a and 2.1b), the probability of having a type 1 error multiple test is given by the equations 2.1c and 2.1d (ŠIDÁK, 1967).

$$P(\text{Making an error}) = \alpha \quad (2.1\text{a})$$

$$P(\text{Not making an error}) = 1 - \alpha \quad (2.1\text{b})$$

$$P(\text{Not making an error in } m \text{ tests}) = (1 - \alpha)^m \quad (2.1\text{c})$$

$$P(\text{Making at least 1 error in } m \text{ tests}) = 1 - (1 - \alpha)^m \quad (2.1\text{d})$$

To handle this multiple statistical testing problems, some techniques to re-calculating probabilities obtained from a statistical test which was repeated multiple times have been developed to prevent the inflation of false positive rates.

Among the different approaches to control type I errors we have Family-wise error rate (FWER) which controls the probability of at least one type I error, and False discovery rate (FDR) which controls the expected proportion of Type I errors among the rejected hypotheses. Compared to FDR, controlling FWER is extremely conservative approach as the power to detect  $H_1$  gets very small.

Among the different adjustment methods to control FWER includes the Bonferroni correction in which the p-values are multiplied by the number of comparisons ( $M * P_i < \alpha$ ) and the Holm correction,  $P - adjusted_i = P_i * (M + 1 - i)$ , where  $i \in \{1, 2, \dots, n\}$  and smaller the p-value is smaller will the index  $i$  be (AICKIN; GENSLER, 1996). The Benjamini-Hochberg (BH) method to control FDR procedure will identify the largest  $k$ , such that  $P_k \leq \frac{k}{m} \alpha$ , all null hypotheses  $H_i$  for  $i \in \{1, \dots, k\}$  are rejected.

These methods make the assumption that the tests are independent tests, which often is not valid for genomics data. For dependent tests, permutation methods are often used to calculate p-values. This approach recalculates a p-value comparing the P-value calculated from the real data test with random ones, which are performed by randomly shuffling the case-control (or phenotype) labels. All  $M$  tests are recalculated on the reshuffled data set, with the smallest P value of these  $M$  tests being recorded. The procedure is repeated for many times to construct an

empirical frequency distribution of the smallest P values. This empirical adjusted P value ( $P_*$ ) is given by:

$$P_* = \frac{r+1}{n+1}$$

where  $n$  is the number of permutation carried out, and  $r$  is the number of permuted p-values smaller than P-value calculated from the real data.

For example, considering P-value = 0.1 and the permuted p-values

$$P_{perm} = \{0.001, 0.01, 0.02, 0.03, 0.05, 0.2, 0.5, 0.6, 1\}$$

the first 5 permuted p-values are smaller than the original p-value, which would give us  $r = 5$ , resulting in:

$$P_* = \frac{r+1}{n+1} = \frac{5+1}{9+1} = 0.6$$

It is important to highlight that a high number of permutations is required in order to produce reliable permuted p-value adjusted. (DAVISON; HINKLEY, 1997; NORTH; CURTIS; SHAM, 2002; NORTH; CURTIS; SHAM, 2003; SHAM; PURCELL, 2014).

#### 2.4.2.4 Nonparametric and parametric tests

Statistical procedures can be classified into two groups: Parametric and nonparametric. Parametric statistical procedures rely on assumptions about the shape of the distribution (i.e., assume a normal distribution) in the underlying population and about the form or parameters (i.e., means and standard deviations) of the assumed distribution. While nonparametric statistical procedures don't rely or rely on only a few assumptions about the shape or parameters of the population distribution from which the sample was drawn. Some of these procedures are summarized in Table 7.

The most used parametric test for comparing the means of two independent groups is the t-test, which assumes that the data are normally distributed, that samples from different groups are independent and that the variances between the groups are equal (KITCHEN, 2009). The most commonly used nonparametric test for independent samples is the Mann-Whitney U-test (MWU), which assumes that observations from the different groups are random samples (i.e. independent and identically distributed) from their respective populations, are mutually independent and are ordinal or continuous measurements. If matched or dependent samples are compared the nonparametric Wilcoxon signed-rank test is used (WHITLEY; BALL, 2002). When there are more than two groups being compared, the nonparametric test used is the Kruskal-Wallis test (KW), a generalization of the MWU and the parametric test used is the Analysis of variance (ANOVA) (PARAB; BHALERAO, 2010).

It is also important to highlight that for larger sample sizes (greater than 20 or 30) P values can be calculated using a Normal distribution, on other words parametric tests could be used instead of the nonparametric ones (VICKERS, 2005).

Table 7 – Summary of parametric and nonparametric procedures.

Analysis Type	Example	Parametric	Nonparametric
Compare means between two distinct/independent groups	Is the mean TP53 gene expression for control group different from the mean for treatment group?	Two-sample t-test	Wilcoxon ranksum test
Compare two quantitative measurements taken from the same individual	Was there a change in gene expression after the treatment?	Paired t-test	Wilcoxon signedrank test
Compare means between three or more distinct/independent groups	For a given three groups (e.g., placebo, drug #1, drug #2), is the TP53 gene expression different among the three groups?	Analysis of variance (ANOVA)	Kruskal-Wallis test
Estimate the degree of association between two quantitative variables	Is age related to the TP53 gene expression?	Pearson coefficient of correlation	Spearman's rank correlation

### 2.4.3 Survival analysis

In the medical sciences, in several circumstances, it is necessary to evaluate if a treatment had a beneficial effect on the survival of patients. For this, it is measured the fraction of patients living for a certain amount of time after treatment.

It is called "Survival times" data that measure follow-up time from a defined starting time to the occurrence of a given event (e.g. from the diagnosis of a disease to death) (BEWICK; CHEEK; BALL, 2004). These survival times are "censored" when there is a follow-up time but the event has not yet occurred or is not known to have occurred. That might happen if a patient drops out of the study before its end, or if you are studying a treatment and it is not over yet.

#### 2.4.3.1 Kaplan-Meier method: Estimating the survival curve

It is defined survival function  $S(t)$  is defined as the probability of surviving at least to time  $t$ , while a graph of  $S(t)$  against  $t$  is called the survival curve. To estimate this curve there exists the Kaplan-Meier method:

$$S(t) = \prod_{i:t_i \leq t} \left(1 - \frac{d_i}{n_i}\right),$$

where  $d_i$  are the number of events and  $n_i$  the total individuals at risk at time  $i$ . Table 8 shows an example of survival data and Table 9 shows an example of applying Kaplan-Meier method. Figure 3 shows a more real data example. It is important to highlight that for a censored time the proportion surviving will be 1, that means an individual is considered to be at risk of dying in the next event of the censoring but not in subsequent events (BLAND; ALTMAN, 2004).

Table 8 – Survival time and status for a group of patients.

Patient ID	Survival times (in days)	Status
1	1	Dead
2	1	Dead
3	2	Alive (censored)
4	3	Dead

Table 9 – Kaplan-Meier method example for table 8

Interval	$n_i$ : patient risk at time $t_i^-$	$d_i = \text{deaths at time } t_i$	$c_i = \text{censored at time } t_i$	$S(t)$
[0, 1)	4	0	0	1
[1, 3)	4 - 0 = 4	2	1	$1 - \frac{2}{4} = 0.5$
[3, End of study]	4 - 2 - 1 = 1	1	0	$0.5 * (1 - \frac{1}{1}) = 0$

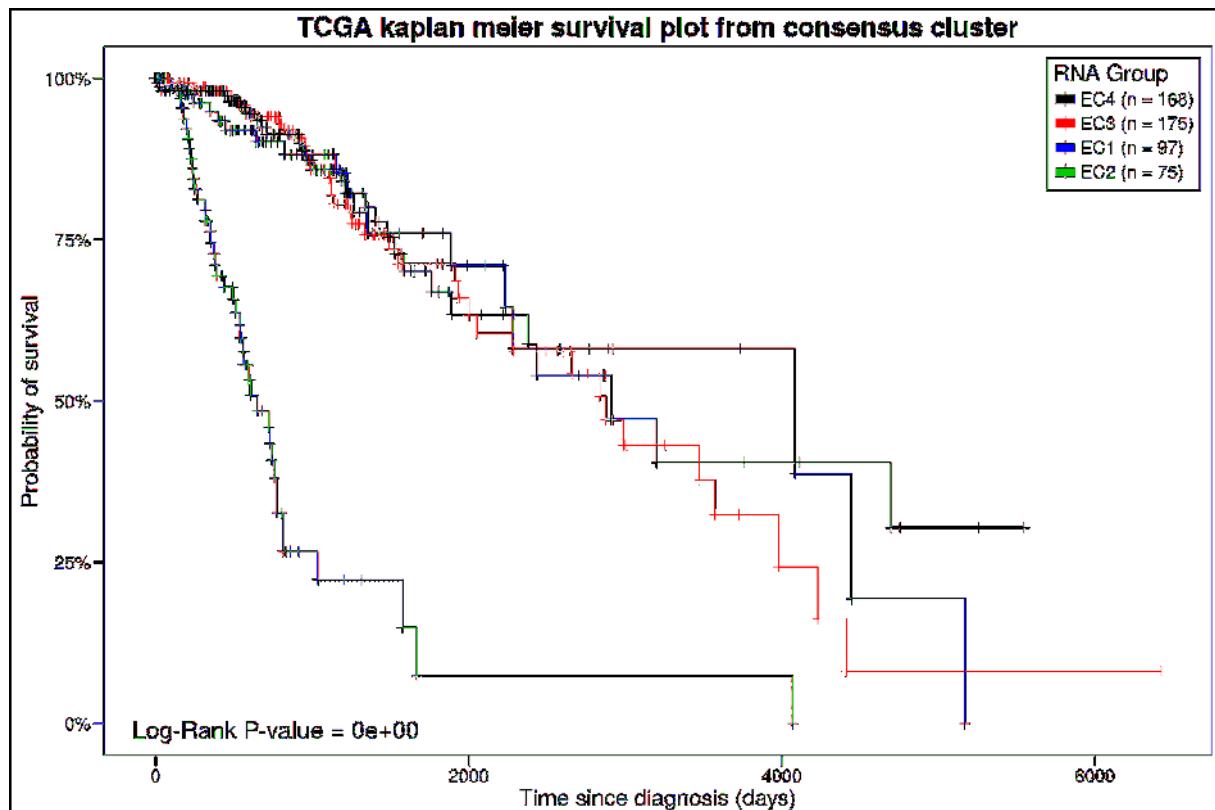


Figure 3 – Example of survival curve for TCGA samples clustered by RNA expression levels. Group 2 has the worst survival. Censored data is marked (+) in the plot.

#### 2.4.3.2 Comparing survival curves of two groups using the log-rank test

To compare the survival distributions of two or more groups, the hypothesis test log-rank test is used to test the null hypothesis that there is no difference between the populations in the probability of an event at any time point. The approximated statistics used for comparison purposes for  $k$  groups is

$$T = \sum_{i \in \{1, \dots, k\}} \frac{(O_i - E_i)^2}{E_i},$$

where  $O_i$  is the observed numbers of death in group  $i$  while  $E_i$  is its expected numbers of deaths. If the null hypothesis is true,  $T$  is distributed approximately as a  $\chi^2_{k-1}$  (MATTHEWS; FAREWELL *et al.*, 1996). If  $T$  calculated is 9.44, and  $k = 2$ , evaluating the quantile function (also known as “inverse CDF” or “ICDF”) of the chi-squared distribution the significance level of these data is equal to  $P_r(\chi^2_1 \geq 9.44) = 0.002$  (YAU, 2012). For a given cut-off, normally 0.05, the results with a p-value smaller are considered significant.

## 2.4.4 Machine Learning

Machine learning is a field of computer science focused on the development and application of algorithms that improve with experience (MITCHELL *et al.*, 1997). In genetics and genomics, it has been applied for the interpretation of large genomic datasets and annotation of a wide variety of genomic sequence elements. For example, for the detection of transcription start sites (TSSs) locations, which have proven hard to detect in silico due to the complexity and the fairly diffuse structures of Eukaryotic promoters, Down and Hubbard (2002) developed a machine-learning method is able to build useful models of promoters for more than 50% of the human transcription start sites (DOWN; HUBBARD, 2002).

The machine learning techniques can be classified into two main categories: supervised and unsupervised learning (MITCHELL *et al.*, 1997). The supervised learning, which aims to infer data labels by learning from already labeled data, has three stages: design, model, and test. The first stage refers to the selection of a learning algorithm used to learn from data (e.g. choose between support vector machines or random forest algorithms) and its training data. The second stage is the creation of a model from labeled data using the algorithm selected previously. The last stage uses this generated model to predict the labels of unlabeled data. The unsupervised learning methods, on the other hand, cluster the data without using labels, which requires an additional step in which semantics must be manually assigned to each cluster. As this discovery is not tied to previously defined classes, these methods have as benefit the ability to identify potentially novel types of genomic elements.

### 2.4.4.1 Supervised learning

Supervised learning is a type of machine learning algorithm that identifies patterns in data to make predictions. Specifically, the algorithm takes a known set of input data and known responses to the data (labels) and trains a model to generate predictions for the response to new data. The supervised learning algorithms can be classified into two categories: classification and regression.

In classification, the goal is to assign a label to an observation. That is, responses are categorical variables. For example, given dataset known to be an enhancer or not enhancer, one could want to predict the locations of enhancers in the genome.

In regression, the goal is to predict a continuous measurement for an observation. That is, the responses variables are real numbers. In genomics, these methods are used in genetic epidemiology to detect association of genetic variants with a trait or disease of interest (DASGUPTA *et al.*, 2011).

#### 2.4.4.2 Unsupervised learning

Unsupervised learning is a type of machine learning algorithm used to draw inferences from datasets consisting of input data without labeled responses. The most common unsupervised learning method is cluster analysis, which is used for exploratory data analysis to find hidden patterns or grouping in data. These algorithms seek to group a set of objects into clusters (groups) such that those in the same group are more similar to each other than to those in another cluster. The major unsupervised learning techniques used in bioinformatics are described below.

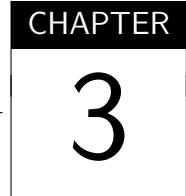
**Hierarchical clustering:** This clustering method builds a hierarchy of clusters, through either a agglomerative ("bottom-up") or a divisive ("top-down") procedures. In the agglomerative procedures, each  $n$  observation starts in its own cluster and until only one cluster remains the groups with the smallest dissimilarity are merged. On the other hand, in the divisive procedures, all observations start in one cluster and until all observations are in their own cluster the group is split into two groups with the biggest dissimilarity. To decide which clusters to merge or divide a metric to measure of dissimilarity between sets of observations is required. Among the existing agglomerative techniques are the single linkage, also known as the nearest-neighbour technique, which defines the smallest dissimilarity between two points in each group as the dissimilarity between two groups (FLOREK *et al.*, 1951), the complete linkage which defines the largest dissimilarity between two points in each group as the dissimilarity between two groups (DEFAYS, 1977), the average linkage which defines the average dissimilarity overall points as the dissimilarity between two groups (SOKAL, 1958) and the Ward's method which at each merge step minimizes the increase in the total within-cluster error sum of squares, which means that the groups leading to a minimum increase in total within-cluster variance are merged (JR, 1963; EVERITT *et al.*, 2011). The results of hierarchical clustering are usually represented in a dendrogram a branching diagram in which the objects are represented in one of the axes while the similarity between clusters are represented in the other axis by the length of the connection which joins them (MANNING; SCHÜTZE *et al.*, 1999).

**Optimization clustering techniques:** These are a set of non-hierarchical clustering techniques that cluster individuals into a specified number of groups, by either minimizing or maximizing some numerical criterion (EVERITT *et al.*, 2011). One of these techniques is the k-means algorithms, which iteratively updates a partition by simultaneously relocating each object to the group to whose mean it was closest and then recalculating the group means until the groups no longer change. More explicitly, let  $S = \{S_1, S_2, \dots, S_k\}$  be the

set of  $k$  clusters, this algorithm tries to minimize the squared distances of the elements from the cluster center:

$$N(n, k) = \min \sum_{g=1}^k \sum_{j \in S_g} \|x_j - \mu_g\|^2,$$

where  $x_j$  is a data element that belongs to a group  $g$ , and  $\mu_g$  is the center of points of the  $g$  group. A counterpoint of these techniques also requirement to ‘estimate’ the number of clusters in the data, for which a variety of methods have been suggested and most are relatively informal and involve, essentially, plotting the value of the clustering criterion against the number of groups in which large changes of levels in the plot are usually taken as suggestive of a particular number of groups. Some formal methods were also proposed such as the Silhouette index by Rousseeuw (1987). And the second counterpoint of this technique is the choice of initial starting values which might lead to a local optimal solution, some solution to this problem such as a bootstrap-like approach to ‘refine’ initial seeds are suggested (STEINLEY, 2003).



# DEVELOPMENT OF METHODS AND SOFTWARES FOR CANCER DATA ANALYSIS

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This chapter introduces three R/Bioconductor packages developed during the doctoral project. These tools are complementary and together they are able to perform a deep integrative analysis of genomic and epigenomic data from different databases, from the data acquisition stage to the integrative analysis.

First, to deal with the problem of data acquisition and analysis we created the R/Bioconductor TCGAbiolinks package which searches, downloads and organizes data and metadata from the National Cancer Institute's (NCI) Genomic Data Commons (GDC) into the latest R/Bioconductor data structures. Second, to deal with the problem of integrative analysis, we worked closely with Dr. Benjamin P. Berman's group to create a new version of the R/Bioconductor ELMER package, which infers regulatory element landscapes and transcription factor networks from cancer methylomes (YAO *et al.*, 2015). This version not only was greatly improved in terms of stability, performance, and extensibility, but it also added a number of new features. Finally, to improve the usability of the tools developed, we used the R web application framework shiny (CHANG *et al.*, 2016) to create the R/Bioconductor TCGAbiolinksGUI package, which provides a graphical interface for our packages and others of the Bioconductor project.

## 3.1 TCGAbiolinks: An R/Bioconductor package to download and analyze data from GDC

The aim of TCGAbiolinks is four-fold: (i) to facilitate data retrieval via GDC's API; (ii) to prepare the data using the appropriate preprocessing strategies; (iii) to provide a means to conduct different standard analyses and advanced integrative analyses and (iv) to allow the user to easily reproduce earlier research results. We introduce public methods used in several marker papers to integrate DNA methylation and gene expression data. In addition, our tool extracts published molecular subtype information for each TCGA sample within a tumor type (generally

embedded in supplementary tables, PDFs or external websites). The tool was developed in the R language specifically for integration within the Bioconductor project, thus we have provided most of the data objects as the Bioconductor specified ‘SummarizedExperiment’ class (HUBER *et al.*, 2015), thereby allowing easy integration with other data types and statistical methods that are common in the Bioconductor repository.

### 3.1.1 The *TCGAbiolinks* package

TCGAbiolinks is an R package, which is licensed under the General Public License (GPLv3), and is freely available through the Bioconductor repository (GENTLEMAN *et al.*, 2004). By conforming to the strict guidelines for package submission to Bioconductor, we were able to utilize and incorporate existing R/Bioconductor packages and statistics to assist in identifying differentially altered genomic regions defined by mutation, copy number, expression or DNA methylation; to reproduce previous TCGA marker studies; and to integrate data types both within TCGA and across other data types outside of TCGA. TCGAbiolinks consists of functions that can be grouped into three main levels: Data, Analysis and Visualization. More specifically, the package provides multiple methods for the analysis of individual experimental platforms (e.g. differential expression analysis or identifying differentially methylated regions or copy number alterations) and methods for visualization (e.g. survival plots, volcano plots and starburst plots) to facilitate the development of complete analysis pipelines. In addition, TCGAbiolinks offers in-depth integrative analysis of multiple platforms, such as copy number and expression or expression and DNA methylation, as demonstrated and applied in our recent TCGA study of 1122 gliomas (CECCARELLI *et al.*, 2016b). These functions can be used independently or in combination to provide the user with fully comprehensible analysis pipelines applied to TCGA data. A schematic overview of the package is presented in Figure 4. The next subsections describe each of the three main levels (Data, Analysis and Visualization), highlighting the importance and utility of each associated function and sub-function.

#### *Data*

TCGA data is accessible via the [NCI Genomic Data Commons \(GDC\) data portal](#), [GDC Legacy Archive](#) and [the Broad Institute’s GDAC Firehose](#). The GDC Data Portal provides access to the subset of TCGA data that has been harmonized against Genome Reference Consortium Human Build 38 (GRCh38) (hg38) using GDC Bioinformatics Pipelines which provides methods to the standardization of biospecimen and clinical data, the re-alignment of DNA and RNA sequence data against a common reference genome build GRCh38, and the generation of derived data. Whereas the GDC Legacy Archive provides access to an unmodified copy of data that was previously stored in CGHub (WILKS *et al.*, 2014) and in the TCGA Data Portal hosted by the TCGA Data Coordinating Center (DCC), in which uses as references Genome Reference Consortium Human Build 37 (GRCh37) (hg19) and Genome Reference Consortium Human

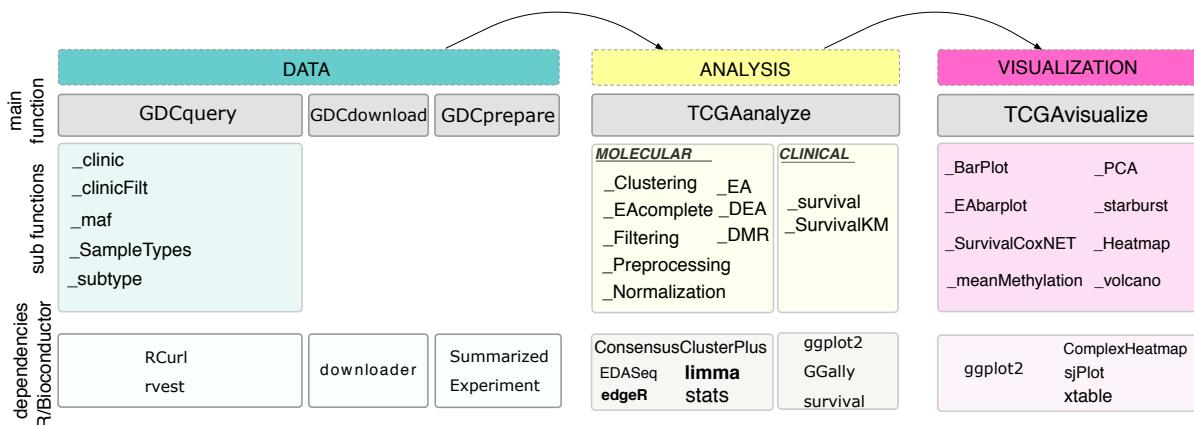


Figure 4 – Overview of TCGAbiolinks functions. TCGAbiolinks is organized in three categories. In the first category (Data), functions to query the GDC database, to download the data and to prepare it are made available. The second category (Analysis) contains functions that allow the user to carry out different types of analyses; these include clustering (*TCGAanalyze\_Clustering*), differential expression analysis (*TCGAanalyze\_DEA*) and enrichment analysis (*TCGAanalyze\_EA*). Finally, the obtained results can be visualized using the functions in the third category (Visualization): these include principal component analysis (*TCGAvizualize\_PCA*), starburst plots (*TCGAvizualize\_starburst*) and survival curves (*TCGAvizualize\_SurvivalCoxNET*). The different dependencies to other R/Bioconductor packages are specified in the last row of the figure.

Build 36 (GRCh36) (hg18).

The previously stored data in CGHub, TCGA Data Portal and Broad Institute's GDAC Firehose, were provided as different levels or tiers that were defined in terms of a specific combination of both processing level (raw, normalized, integrated) and access level (controlled or open access). Level 1 indicated raw and controlled data, level 2 indicated processed and controlled data, level 3 indicated Segmented or Interpreted Data and open access and level 4 indicated region of interest and open access data. While the TCGA data portal provided level 1 to 3 data, Firehose only provides level 3 and 4. An explanation of the different levels can be found at TCGA Wiki (<https://wiki.nci.nih.gov/display/TCGA/Data+level>). However, the GDC data portal no longer uses this based classification model in levels. Instead a new data model was created, its documentation can be found in GDC documentation at <https://gdc.nci.nih.gov/developers/gdc-data-model/gdc-data-model-components>. In this new model, data can be open or controlled access. While the GDC open access data does not require authentication or authorization to access it and generally includes high level genomic data that is not individually identifiable, as well as most clinical and all biospecimen data elements, the GDC controlled access data requires dbGaP authorization and eRA Commons authentication and generally includes individually identifiable data such as low level genomic sequencing data, germline variants, SNP6 genotype data, and certain clinical data elements. The process to obtain access to controlled data is found in GDC web site at <https://gdc.nci.nih.gov/access-data/obtaining-access-controlled-data>.

TCGAbiolinks divides the GDC data retrieval into three main functions: *GDCquery*, *GDCdownload* and *GDCprepare*. *GDCquery* allows the user to query data from the NCI's Genomic Data Commons (GDC) data portal or GDC Legacy Archive by accessing the

database API. Up to the moment, the GDC data portal provides data from two programs The Cancer Genome Atlas (TCGA) and Therapeutically Applicable Research to Generate Effective Treatments (TARGET) on more than 38 diseases type cancer types and 5 different molecular data types (Transcriptome Profiling, Raw Sequencing Data, Copy Number Variation, DNA Methylation) as well as 3 different types of clinical reports (clinical tables, pathology reports and histology image slides). The pathology reports and histology image slides are not prepared but are downloaded from the GDC Legacy Archive to a directory if requested by the user. *GDCdownload* receives the output of *GDCquery*, a file manifest with all the metadata necessary to download it. In order to organize the files downloaded, this function will save the data with the following pattern "Root directory/project/source/data\_category/data\_type/file\_id/file\_name" (i.e Example: GDCdata/TCGA-GBM/harmonized/DNA\_Methylation/Methylation\_Beta\_Value/079fcaff-3ae6-4150-b2e6-2b7330ffbcd9/jhu-usc.edu\_GBM.HumanMethylation450.10.lvl-3.TCGA-19-A6J5-01A-21D-A33U-05.gdc\_hg38.txt) If a file was previously downloaded it will not be re-downloaded. *GDCprepare* is a function that reads open processed data and prepares them for downstream analysis. Specifically, the objects are organized in a *SummarizedExperiment* object to allow easy integration with other Bioconductor packages, such as GRanges (LAWRENCE *et al.*, 2013), IRanges (LAWRENCE *et al.*, 2013), limma (RITCHIE *et al.*, 2015) and edgeR (ROBINSON; MCCARTHY; SMYTH, 2010). The samples are always referred to by their given TCGA/TARGET barcode. If the user prefers the data not to be prepared in a *SummarizedExperiment*, there is an option to set the argument *SummarizedExperiment* to FALSE; the data are then prepared as a standard data frame object (rows and columns).

## Analysis

The analysis functions and subfunctions are designed to analyze TCGA data through both common and novel methods. The main function, called *TCGAanalyze*, comprises two distinct types of analysis: molecular analysis and clinical analysis. Once the data are prepared into data matrices (genes/loci in rows and samples in columns) or a *SummarizedExperiment*, the downstream analysis can be divided into (i) supervised analysis: differential expression analysis, enrichment analysis and master regulator analysis or (ii) unsupervised analysis: inference of gene regulatory network, clustering, classification, Receiver Operator Characteristics (ROC) (SONEGO; KOCSOR; PONGOR, 2008), Area Under the Curve (AUC), feature selection and survival analysis. *TCGAanalyze\_Normalization* allows users to normalize mRNA transcripts and miRNA using the EDASeq package (RISSO *et al.*, 2011). This function uses within-lane normalization procedures to adjust for GC-content effects (or other gene-level effects) on read counts: LOESS robust local regression and global-scaling, full-quantile and between-lane normalization procedures to adjust for distributional differences between lanes (e.g. sequencing depth). *TCGAanalyze DEA* allows the user to identify differential expression or regions between two populations or conditions. In particular, we used the edgeR package from Bioconductor, which uses the quantile-adjusted conditional maximum likelihood (qCML) method for experiments

with a single factor to detect differentially expressed genes (DEGs) (ROBINSON; MCCARTHY; SMYTH, 2010). Compared to several other estimators, qCML is the most reliable in terms of bias on a wide range of conditions; specifically, qCML performs best in situations involving many small samples with a common dispersion (ROBINSON; SMYTH, 2007). The P-values generated from the analysis are sorted in ascending order and corrected using the Benjamini & Hochberg procedure for multiple testing correction (BENJAMINI; HOCHBERG, 1995a). After running *TCGAanalyze\_DEA*, it is possible to filter the output by fold change and/or significance and to use the *TCGAanalyze\_LevelTab* function to create a table of DEGs, including fold change (FC), false discovery rate (FDR), gene expression levels of samples under conditions of interest and delta values (the difference in gene expression multiplied by logFC). *TCGAanalyze\_DMR* allows the user to identify differentially methylated regions (DMRs) between two groups with a DNA methylation difference above a certain threshold. To calculate P-values, this subfunction uses the Wilcoxon ranksum statistical non-parametric test and adjusts the values using the FDR method. *TCGAanalyze\_Clustering* allows the user to perform a hierarchical cluster analysis through two methods: ward.D2 and ConsensusClusterPlus (WILKERSON; HAYES, 2010).

### Visualization

The visualization section allows the user to visualize the results generated by the analysis sections using heatmap, cluster, plots with incremental layers (ggplot2), pathway enrichment analysis and PCA. Furthermore, we provide methods to generate a starburst plot, which integrates gene expression and DNA methylation data (NOUSHMEHR *et al.*, 2010).

### 3.1.2 Comparisons

Recently, several tools to retrieve TCGA data sets have been made available, as summarized in Table 10. These tools include TCGA-Assembler (ZHU; QIU; JI, 2014), CGDS-R (GAO *et al.*, 2013), canEnvolve (SAMUR *et al.*, 2013), Firehose (DENG *et al.*, 2017), RTCGAToolbox (SAMUR, 2014), and cBioPortal (CERAMI *et al.*, 2012). These tools can be divided into three representative categories. The first category comprises tools mainly used to download cancer genomics data, such as TCGA-Assembler and CGDSR. The second category includes tools that focus mainly on data analysis and integration, such as canEnvolve. The third category comprises tools to download and analyze data, such as RTCGAToolbox, Firehose and cBioPortal.

RTCGAToolbox is a tool that systematically accesses the Broad GDAC Firehose (<https://gdac.broadinstitute.org/>) preprocessed data and performs basic analysis and visualization of an individual data type (expression, mutation or DNA methylation). Despite the existence of TCGA specific software packages, none of these tools perform the integrative analysis harnessing methodologies designed by TCGA Analysis working groups (AWGs), such as identifying epigenetically silenced genes (represented in a starburst plot (NOUSHMEHR *et al.*, 2010) or functional copy number identification (CECCARELLI *et al.*, 2016b). Although RTCGAToolbox

Table 10 – Each column represents a software tool compared with TCGAbiolinks, and each row represents a feature. The cells checked with X indicates features that exists in the tool. Available platform abbreviations are defined as: R (R script); C (R package deposited in CRAN); B (Bioconductor package); W (available only as a web portal);

		Packages							
		TCGAbiolinks	TCGAAssembler	canEnvolve	TCGA2stat		Firehose-FirebrowserR	RTCGAtoolbox	cBio Portal CGDS-R
Features	Sub-features								
Availability	Platform	B	R	W	C	CW	B	CW	
Genome of reference	Access to data aligned against the GRCh38/hg38	X	X						
	Access to data aligned against the GRCh37/hg19	X	X	X	X	X	X	X	X
Query TCGA Cases	Individual TCGA samples (e.g. TCGA-01-0001)	X	X				X		
Download	All TCGA platforms	X							
Data type analysis	mRNA	X		X	X	X	X	X	X
	miRNA	X		X	X	X	X	X	X
	copy number	X		X	X	X	X	X	X
	DNA methylation	X			X	X	X	X	X
	Clinical	X		X	X	X	X	X	X
	Protein			X		X			X
	Mutatation	X		X	X	X	X	X	X
Integrative analysis	DNA methylation and gene expression	X				X			
Other	Extensible to other BioC packages	X							

can download and analyze Firehose-generated data, neither tool can provide the downloaded data as a "SummarizedExperiment" object, which is critical for allowing the full integration and use of other popular Bioconductor packages, an integral aspect of Bioconductor (GENTLEMAN *et al.*, 2004, 2004). Briefly, the SummarizedExperiment class is a matrix-like container in which rows represent ranges of interest (as a GRanges or GRangesList object) and columns represent samples (with sample data summarized as a DataFrame). A Summarized Experiment contains one or more assays, each represented by a matrix-like object of numeric or other mode. Finally, TCGAbiolinks is able to access data aligned against the Genome Reference Consortium Human Build 38 (hg38) via the [NCI Genomic Data Commons \(GDC\) data portal](#), and data aligned against the Genome Reference Consortium Human Build 19 (hg19) via the [GDC Legacy Archive](#). This feature is only available using TCGA-Assembler.

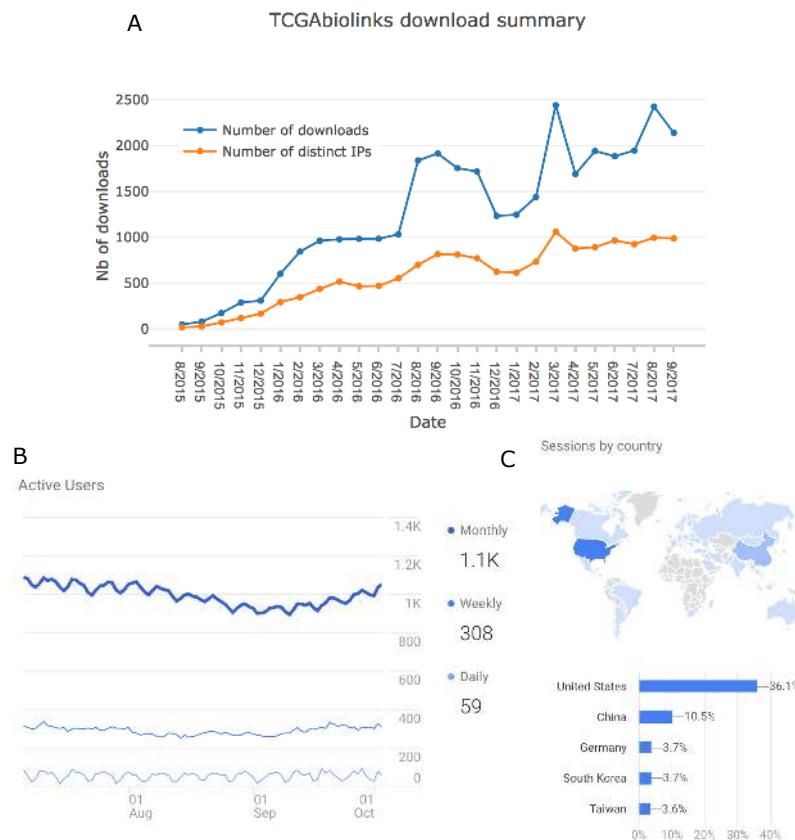


Figure 5 – TCGAbiolinks download summary. A) Number of downloads per month. B) Number of active users. C) Percentage of sessions connect to the documentation by country.

### 3.1.3 Software availability

TCGAbiolinks is available under the GNU General Public License version 3 (GNU GPL3). Its source code is available at <https://github.com/BioinformaticsFMRP/TCGAbiolinks> and a binary version for windows, macOSX and Linux is freely available through the Bioconductor repository at <http://bioconductor.org/packages/TCGAbiolinks/>. To execute this tool, it is required to have installed a R version  $\geq 3.3$ .

### 3.1.4 Public reception

TCGAbiolinks had a good reception the research community being among the top 5% of the most downloaded tools of the Bioconductor project. In October 2017, the tool already had more than 17 thousand downloads, with an average of visits to the documentation pages of a thousand users per month. The Figure 5 shows a summary since the beginning of the project.

## 3.2 TCGAbiolinksGUI: A graphical user interface to analyze GDC cancer molecular and clinical data

Although TCGAbiolinks is a suitable R package for most data analysts with a strong knowledge and familiarity with R specifically those who can comfortably write R commands, we developed TCGAbiolinksGUI to enable user access to the methodologies offered in TCGAbiolinks and to give users the flexibility of point-and-click style analysis without the need to enter specific arguments. TCGAbiolinksGUI takes in all the important features of TCGAbiolinks and offers a graphics user interface (GUI) thereby eliminating any need to familiarize TCGAbiolinks' key functions and arguments.

### 3.2.1 *Infrastructure*

The TCGAbiolinksGUI user interface was created using Shiny, a Web Application Framework for R, and uses several packages to provide advanced features that can enhance Shiny apps, such as shinyjs to add JavaScript actions (ATTALI, 2017), shinydashboard to add dashboards (CHANG; Borges Ribeiro, 2017) and shinyFiles (PEDERSEN, 2016) to provide access to the server file system.

The following R/Bioconductor packages are used as back-ends for the data retrieval and analysis: TCGAbiolinks (COLAPRICO *et al.*, 2016) which allows to search, download and prepare data from the NCI's Genomic Data Commons (GDC) data portal into an R object and perform several downstream analysis; ELMER (Enhancer Linking by Methylation/Expression Relationship) (YAO *et al.*, 2015; SILVA *et al.*, 2017a) which identifies DNA methylation changes in distal regulatory regions and correlate these signatures with the expression of nearby genes to identify transcriptional targets associated with cancer; ComplexHeatmap (GU; EILS; SCHLESNER, 2016) to visualize data as oncoprint and heatmaps; pathview (LUO; BROUWER, 2013) which offers pathway based data integration and visualization; and maftools (MAYAKONDA; KOEFFLER, 2016) to analyze, visualize and summarize Mutation Annotation Format (MAF) files.

### 3.2.2 *Graphical user interface design*

The user interface has been divided into three main Graphical User Interface (GUI) menus. The first menu defines the acquisition of GDC data. The second defines the analysis steps which subdivides according to the molecular data types. And the third is dedicated to harnessing integrative analyses. We present below a brief description of each menu and their features that can be accessed through a side panel (see figure 6):

- **Data:** Provides a guided approach to search for published molecular subtype information, clinical and molecular data available in GDC. In addition, it downloads and processes

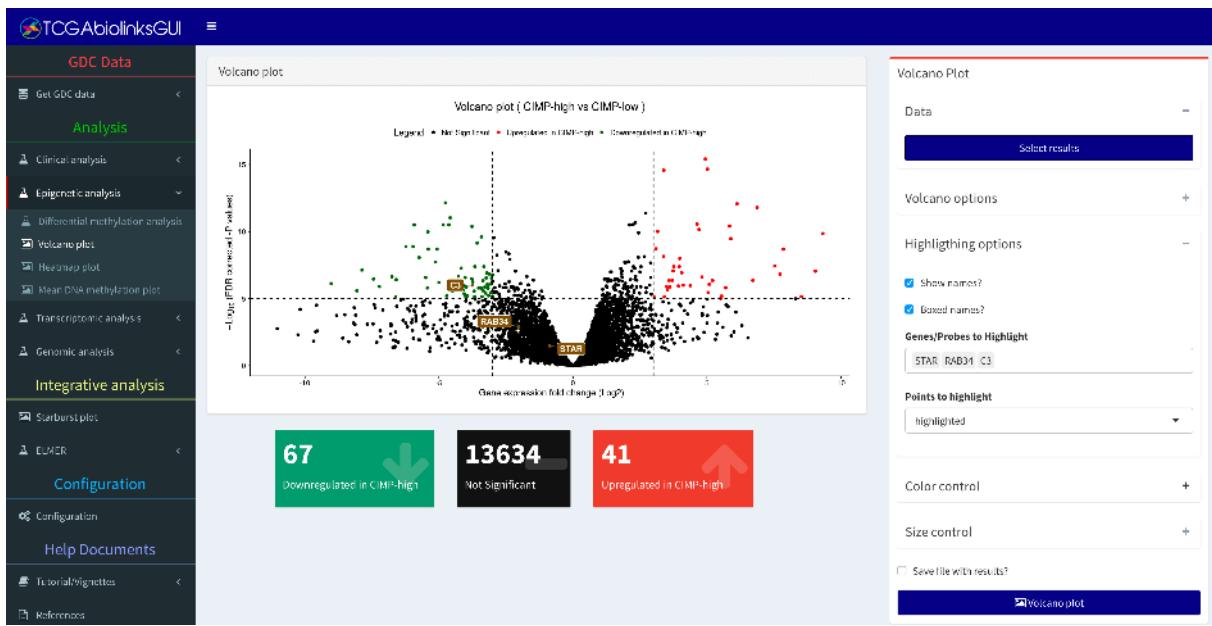


Figure 6 – The volcano plot menu of TCGAbiolinksGUI: The panel on the left shows the menus divided by different analyses, the panel on the right shows the controls available for the menu selected. In the center is a volcano plot window from the analysis menu. It is possible to control the colors, to change cut-offs and to export results into a CSV document.



Figure 7 – Molecular data download: Gene expression Search, download and prepare into an R object of gene expression data for two TCGA-LUSC samples ("TCGA-34-5231-01", "TCGA-77-7138-01"). The Summarized Experiment object created is saved as an RData file (TCGA-LUSC-Gene\_Expression\_Quantification\_hg38.rda).

the molecular data into an R object that can be used for further analysis. For raw DNA methylation data obtained in the form of Intensity Data (IDAT) files, we provide a pipeline using the R/Bioconductor minfi package (<http://bioconductor.org/packages/minfi/>) to prepare the data for subsequent bioinformatics analysis (ARYEE *et al.*, 2014) performing a background and dye-bias correction with the `preprocessnoob` function followed by a detection P-value quality masking (sample-specific) (MORRIS; BECK, 2015) and probes overlapping repeats or single nucleotide polymorphisms masking (non-sample specific) (ZHOU; LAIRD; SHEN, 2016) (Figure 9).

- **Clinical analysis:** Performs survival analysis to quantify and test survival differences between two or more groups of patients and draws survival curves with the 'number at risk' table, the cumulative number of events table and the cumulative number of censored subjects table using the R/CRAN package survminer (KASSAMBARA; KOSINSKI, 2017).
- **Epigenetic analysis:** Performs a differentially methylated regions (DMR) analysis, visualizes the results through both volcano and heatmap plots, and visualizes the mean DNA methylation level by groups or subtypes. For certain tumor types like Glioma, we have added a function to classify non TCGA derived DNA methylation data into one of the 7 published epigenomic subtypes (CECCARELLI *et al.*, 2016b) using a RandomForest trained model derived from DNA methylation signatures available at [https://tcga-data.nci.nih.gov/docs/publications/lggbm\\_2015/](https://tcga-data.nci.nih.gov/docs/publications/lggbm_2015/) (Figure 10). Details of how these RandomForest were created are available in the `TCGAbiolinksGUI.data` vignette (<http://bioconductor.org/packages/TCGAbiolinksGUI.data/>).
- **Transcriptomic analysis:** Performs a Differential Expression Analysis (DEA), and visualizes the results through both volcano and heatmap plots. Pathway analysis can be performed on a list of differentially expressed genes (LUO; BROUWER, 2013).
- **Genomic analysis:** Visualize and summarize the mutations from MAF (Mutation Annotation Format) files through summary plots and oncoplots using the R/Bioconductor maftools package (GU; EILS; SCHLESNER, 2016; MAYAKONDA; KOEFFLER, 2016) (Figures 8 and 14).
- **Integrative analysis:** Integrate the DMR and DEA results through a starburst plot. Integrate clinical and mutation data by way of a Kaplan-Meier survival analysis for groups of mutated samples vs non-mutated for a given gene (Figure 13). DNA methylation and gene expression data can be further analyzed using the R/Bioconductor ELMER package to discover functionally relevant genomic regions associated with cancer (YAO *et al.*, 2015; SILVA *et al.*, 2017a).



Figure 8 – TCGAbiolinksGUI: Visualizing mutation summary. This maftools plot available thought TCGAbiolinks-GUI shows a summary of the MAF file. Highlighting the most mutated genes, SNV class and variant classification distributions.

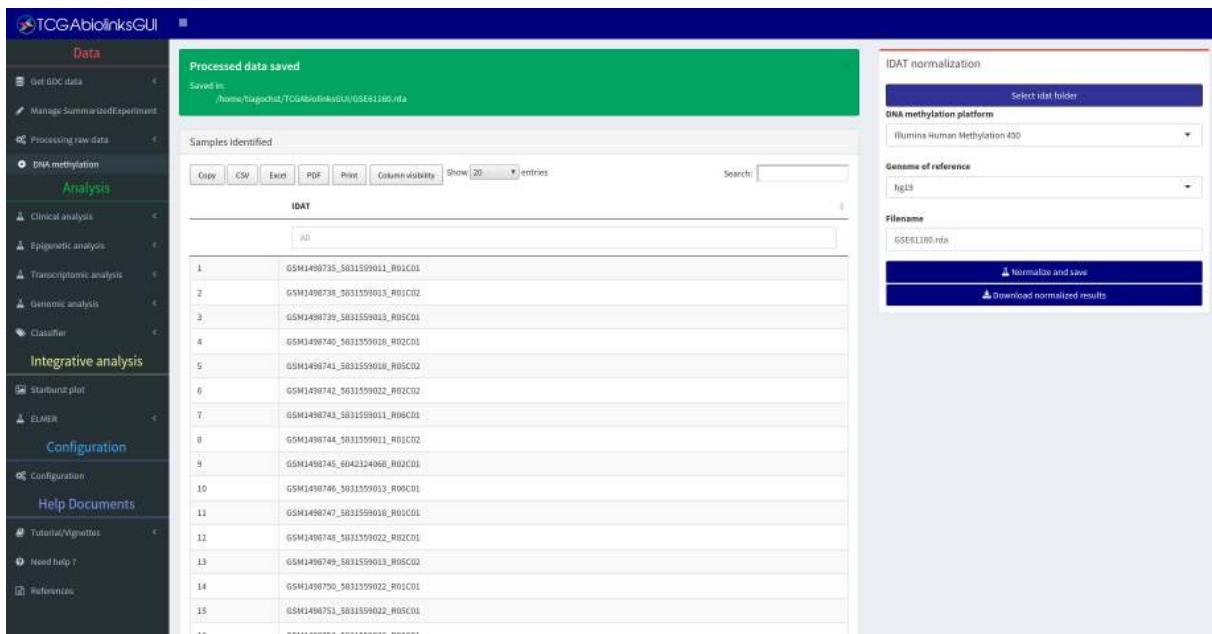


Figure 9 – TCGAbiolinksGUI: IDAT normalization. Table shows files identified that will be processed. Data retrieved from GEO (accession GSE61160).

### 3.2.3 Documentation

We provide a guided tutorial for users via an online vignette document (<http://bit.do/TCGAbiolinksDocs>) which details each step and menu function. Printable PDF ([http://bit.ly/TCGAbiolinks\\_PDFTutorial](http://bit.ly/TCGAbiolinks_PDFTutorial)) and YouTube video instructions ([http://bit.ly/TCGAbiolinksGUI\\_videoTutorial](http://bit.ly/TCGAbiolinksGUI_videoTutorial)) are also provided to help users utilize TCGAbiolinksGUI. A demonstration version of the tool is available at <http://tcgabiolinks.fmrp.usp.br:3838/>. To help improve and

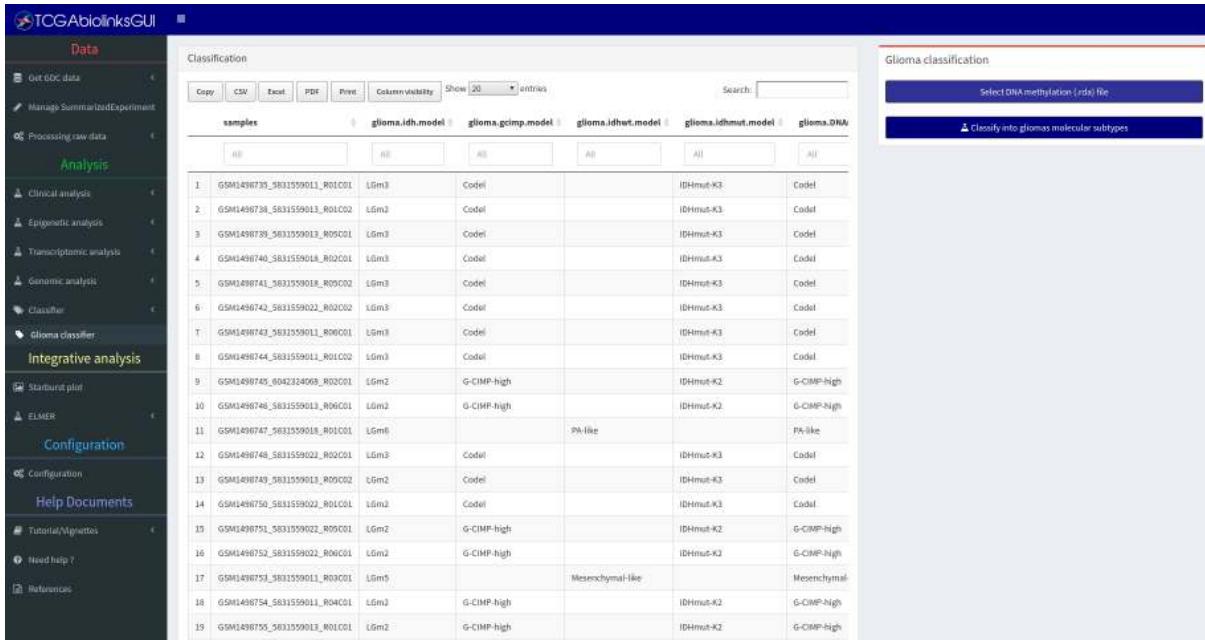


Figure 10 – TCGAbiolinksGUI: glioma classifier. Predicting molecular subtypes based on DNA methylation using data from GEO (accession GSE61160).

expand our tool over time, users are encouraged to report and file bug reports or feature requests via our GitHub repository (<https://github.com/BioinformaticsFMRP/TCGAbiolinksGUI/issues>).

### 3.2.4 Docker container

We recognize that one of the possible drawbacks of using our tool is the arduous process of installing R/Bioconductor environment and all of the required dependencies. To resolve this, a web version of our tool would suffice however the demand on data storage, computational processing, and multiple user access would not be justifiable and instead, we encourage users to use their own computers. However, to further simplify the usability and accessibility of our tool, we provide a Docker image file. This file is compatible with most popular operating systems and it is available online at <https://hub.docker.com/r/tiagochst/tcgabiolinksgui/>. It contains the complete R/Bioconductor environment configured to use TCGAbiolinksGUI. By providing a Docker image, we anticipate users with very little knowledge of R/Bioconductor to run TCGAbiolinksGUI without the need to install associated dependencies or configure system files, common steps required to run R installations and load R/Bioconductor packages. This image can be easily downloaded and deployed through the Kitematic tool (<https://kitematic.com/>), a simple application for managing Docker containers for Mac, Linux, and Windows. A detailed documentation on how to get and use the docker image via the Kitematic application is available at [http://bit.ly/TCGAbiolinksGUI\\_docker](http://bit.ly/TCGAbiolinksGUI_docker). The Docker image will be updated as regularly as TCGAbiolinksGUI thus providing several types of access for end-users interested in analyzing TCGA data from GDC.



Figure 11 – TCGAbiolinksGUI: Enrichement analysis of genes. TCGAbiolinks uses Gene Ontology which defines concepts/classes used to describe gene function, to perform an enrichment analysis. The plots shows molecular function, biological process, cellular components, and pathway gene Ontology classes. Each barplot represents a class with the numnber of genes in the class. The width of the barplot represents the significance, and the red line represents the percentage genes inputed that is found class, for example of all genes in the biological process regulation of endothelial cell migration our 2 genes inputed represents 20% of all genes in this process.



Figure 12 – TCGAbiolinksGUI: Visualizing DMR results as heatmap. Plot shows the DMR results: hypermethylated probes in Solid tissue normal samples compared to the Primary Solid tumor samples. Each column is a sample, while each row is a probe. Blue colors represents probes with low levels of DNA methylation and red the ones with high level.

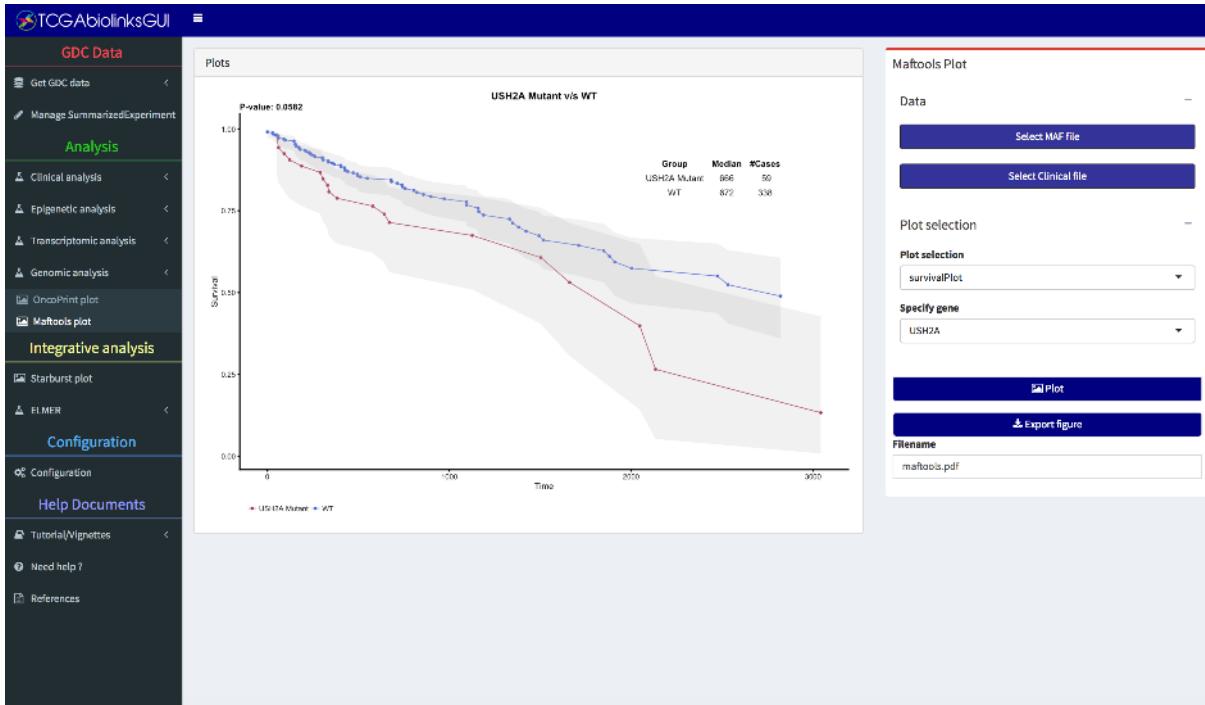


Figure 13 – TCGAbiolinksGUI: Integrating clinical data and mutation data. Performing Kaplan-Meier survival analysis for groups of samples with a mutation in gene USH2A vs the WT ones using the R/Bioconductor maftools package.

### 3.2.5 Comparison of alternative software

Web tools used for cancer data analysis might be classified into two broad groups. The first group only provides an interface to existing software analysis tools. The Galaxy project (<https://galaxyproject.org/>), which is an open, web-based platform for accessible, reproducible, and transparent computational biomedical research, is an example of such a tool that belongs to this group. The other group is composed of exploratory tools mainly focused on the visualization of processed data and pre-computed results. The cBioPortal project (GAO *et al.*, 2013; CERAMI *et al.*, 2012), by providing several visualizations for mining the TCGA data, is an example of a tool that falls within this classification.

If one were to correctly classify TCGAbiolinksGUI, it would belong to the first group. Compared to the Galaxy project, TCGAbiolinksGUI offers an open platform which improves the accessibility of R/Bioconductor packages, allowing users an advantage to integrate their features with existing Bioconductor packages. Unlike the Galaxy project, which requires the interface elements to be structured through XML files (TURAGA *et al.*, 2016), TCGAbiolinksGUI can resolve this simply because it was built within the R/Shiny frameworks. cBioPortal and TCGAbiolinksGUI provides users with access to raw and processed data, however TCGAbiolinksGUI allows users to perform in-depth integrative analysis which cBioPortal currently lacks. For example, if a user is interested in defining differentially expressed genes or DNA methylation events between two populations of tumors (i.e. FOXA1 mutants and FOXA1 wildtypes), by using cBioPortal, a user would have to download the gene expression, DNA methylation and mutation

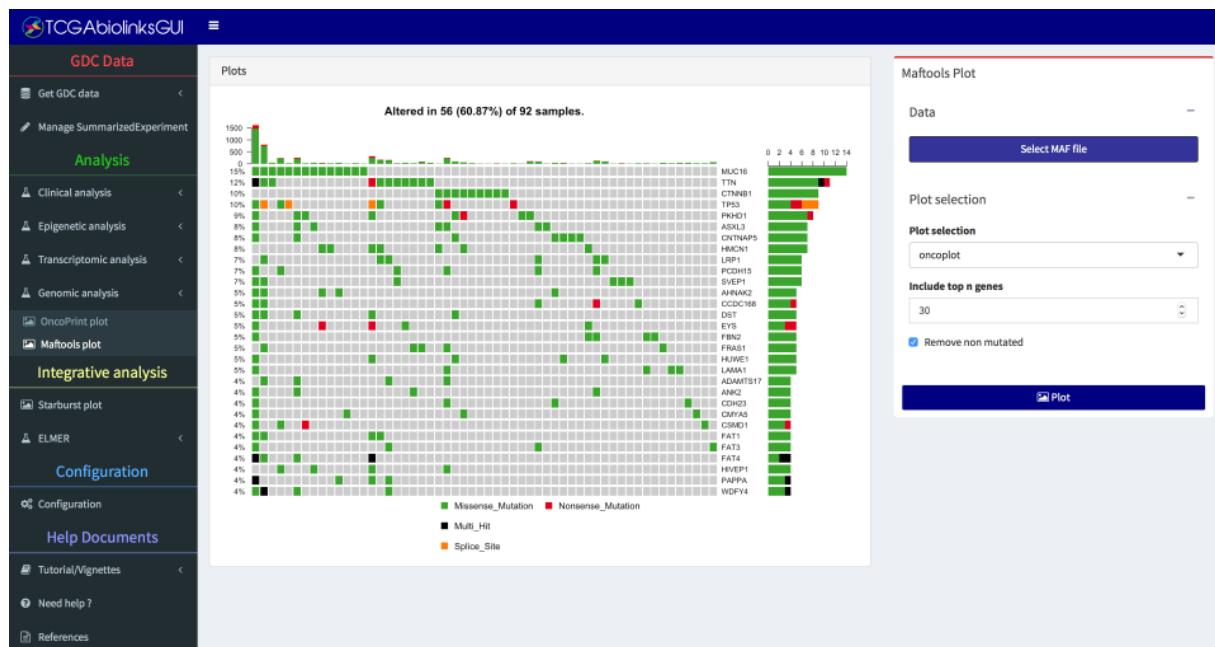


Figure 14 – TCGAbiolinksGUI: Visualizing mutation as an oncplot. Each column represents a sample and each row a different gene. The top barplot has the frequency of mutations for each patient, while the right barplot has the frequency of mutations for each gene. The plot by default is ordered by the most mutated genes.

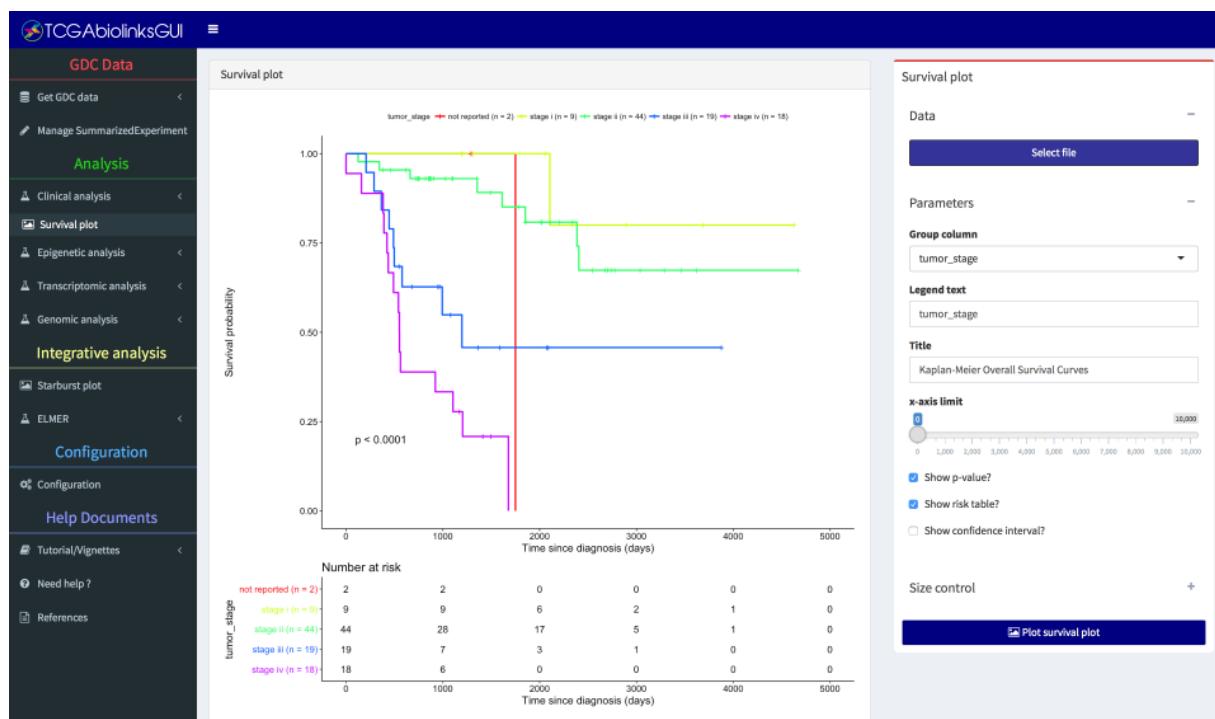


Figure 15 – TCGAbiolinksGUI: Survival analysis. The plot shows the Kaplan-Meier Overall Survival Curves for ACC (Adrenocortical carcinoma) stratified by tumor stage. As expected, higher levels which are more aggressive have a lower survival.

data, define the samples per group, and import the data into their favorite statistical tool to identify their list of differentially expressed or methylated genes. Whereas, within TCGAbiolinksGUI, we developed the platform so that the user can define the sample groups based on their mutation spectrum, perform any type of supervised analysis that can then define differentially expressed or methylated gene list and this can be directly ported into pathway analysis. In addition, if the user is interested in observing survival differences between the groups, this can also be done within TCGAbiolinksGUI and thereby reducing the need to exit a specific data platform or having to massage the downloaded data to fit some other statistical platform to achieve the same goals.

### 3.3 Enhancer Linking by Methylation/Expression Relationships (ELMER)

Motivated by the discovery of transcriptional enhancers in tissue DNA methylation data (BERMAN *et al.*, 2012), and subsequent approaches to linking these enhancers to transcriptional targets using a chromQTL approach (ARAN; HELLMAN, 2013) (reviewed in Yao *et al.*), Yao *et al.* developed the R/Bioconductor *ELMER* (Enhancer Linking by Methylation/Expression Relationships) package, a tool which infers regulatory element landscapes and transcription factor networks from cancer methylomes.

This tool combined DNA methylation and gene expression data from human tissues to infer multi-level cis-regulatory networks through several steps which included the identification of distal enhancer probes with significantly altered DNA methylation levels in primary tumor tissues compared to normal tissues, followed by the identification of putative target genes, and a comprehensive gene regulatory network analysis which combined transcription factor motifs at the altered enhancers with TF expression to identify the underlying master regulators. This approach identified several known and unknown master regulators in TCGA data, such as GATA-binding protein 3 (GATA3) and Forkhead box protein A1 (FOXA1) in breast cancer, and tumor protein p63 (P63) and Sex determining region Y-box 2 (SOX2) in squamous cell lung carcinoma (YAO *et al.*, 2015; SILVA *et al.*, 2016).

Based on user feedback and a full review of the source code, we identified and implemented a number of software improvements, which are summarized in table 11: (i) The original package contained no standard data structure to handle multiple assays (DNA methylation, gene expression, and clinical data), which would be required for an integrative genomic data analysis. Recently, the Bioconductor team provided such a data structure through the [Multi-AssayExperiment](#) package. (ii) All auxiliary databases (human TF list, classification of TF in families, gene annotation, DNA methylation annotation and motif occurrences within probe sites) used in the package were created and maintained manually, thereby making the upgrade process laborious; thus, we automated this process. (iii) The package was developed to analyze primary tumor tissue samples compared to normal tissues samples, thus not allowing arbitrary

Table 11 – Main differences between ELMER old version (v.1) and the new version (v.2)

Features	ELMER Version 1	ELMER Version 2
Primary data structure	mee object (custom data structure)	MAE object (Bioconductor data structure)
Auxiliary data	Manually created	Programmatically created
Number of human TFs	1,982	2,014 (UniProt database)
Number of TF motifs	91	771 (HOCOMOCO v11 database)
TF classification	78 families	82 families and 331 subfamilies (TFClass database, HOCOMOCO)
Analysis performed	Normal vs tumor samples	Group 1 vs group 2
Statistical grouping	Unsupervised only	Unsupervised or supervised using labeled groups
TCGA data source	The Cancer Genome Atlas (TCGA) (not available)	The NCI's Genomic Data Commons (GDC)
Genome of reference	GRCh37 (hg19)	GRCh37 (hg19)/GRCh38 (hg38)
DNA methylation platforms	HM450	EPIC and HM450
Graphical User Interface (GUI)	None	TCGAbiolinksGUI
Automatic report	None	HTML summarizing results
Annotations	None	StateHub

subgroups to be compared (for instance mutants vs. non-mutants, treated vs. untreated, etc.) (iv) The first approach used known epigenomic markers for enhancers to constrain the genomic regions searched for differential methylation. However, this selection could limit our algorithm to identifying regulatory networks for tissue types that exist in the epigenomic databases; we found this constraint problematic, and thus now search *all* distal regulatory regions without any such filter. (v) The function used to download data from The Cancer Genome Atlas (TCGA) data portal (TOMCZAK *et al.*, 2015) broke when the TCGA site was shutdown and its data transferred to The NCI's Genomic Data Commons (GDC) (GROSSMAN *et al.*, 2016); we now have a more general data provider interface that supports GDC as the default provider. (vi) The package only supported data aligned to Genome Reference Consortium GRCh37 (hg19), and we now provide support for Genome Reference Consortium GRCh38 (hg38). (vii) There was no support to the recent HumanMethylationEPIC (EPIC) array. In addition to the specific improvements listed above, we substantially re-wrote most of the code to be more efficient and maintainable, also most of the output plots generated were improved.

In this section, we present a new version of the R *ELMER* package, which addresses all the issues described above.

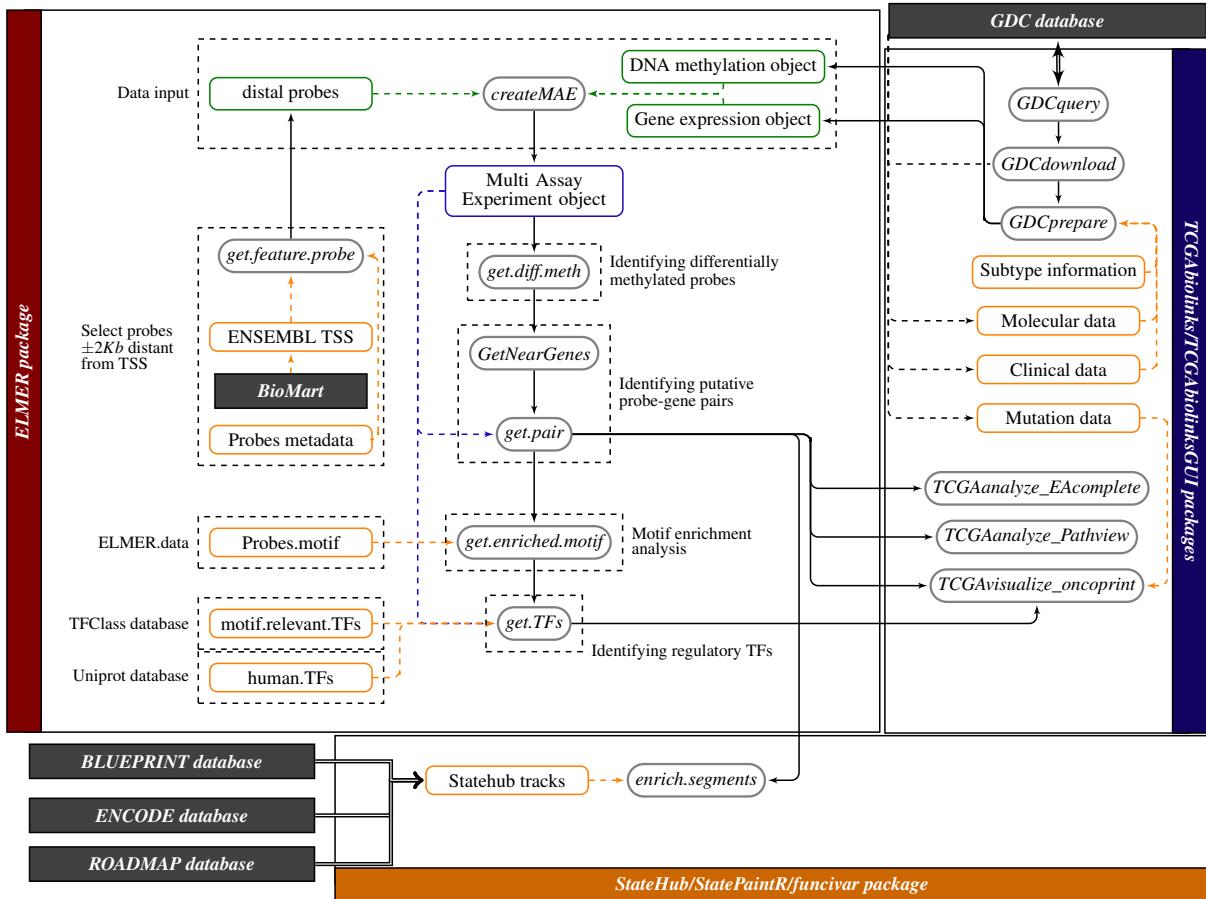


Figure 16 – ELMER workflow: ELMER receives as input a DNA methylation object, a gene expression object (a matrix or a `SummarizedExperiment` object) and a Genomic Ranges (`GRanges`) object with distal probes to be used as filter which can be retrieved using the `get.feature.probe` function. The function `createMAE` will create a Multi Assay Experiment object keeping only samples that have both DNA methylation and gene expression data. Genes will be mapped to genomic position and annotated using ENSEMBL database (AKEN *et al.*, 2016), while for probes it will add annotation from ZHOU; LAIRD; SHEN (<http://zwdzwd.github.io/InfiniumAnnotation>). This MAE object will be used as input to the next analysis functions. First, it identifies differentially methylated probes followed by the identification of their nearest genes (10 upstream and 10 downstream) through the `get.diff.meth` and `GetNearGenes` functions respectively. For each probe, it will verify if any of the nearby genes were affected by its change in the DNA methylation level and a list of gene and probes pairs will be outputted from `get.pair` function. For the probes in those pairs, it will search for enriched regulatory Transcription Factors motifs with the `get.enriched.motif` function. Finally, the enriched motifs will be correlated with the level of the transcription factor through the `get.TFs` function. In the figure green Boxes represents user input data, blue boxes represent output object, orange boxes represent auxiliary pre-computed data and gray boxes are functions.

### 3.3.1 Implementation

Here we describe each of following analysis steps shown in Figure 16: i) Organize data as a `MultiAssayExperiment` object; ii) Identify distal probes with significantly different DNA methylation level when comparing two sample groups; iii) Identify putative target genes for differentially methylated distal probes, using methylation vs. expression correlation; iv) Identify enriched motifs for each probe belonging to a significant probe-gene pair; v) Identify master regulatory Transcription Factors (TF) whose expression associate with DNA methylation changes at multiple regulatory regions.

### 3.3.1.1 Organization of data as a *MultiAssayExperiment* object

To facilitate the analysis of experiments and studies with multiple samples the Bioconductor team created the *SummarizedExperiment* class (HUBER *et al.*, 2015), a data structure able to store data and metadata for a single experiment but not for data spanning several experiments for the same sample. To overcome this problem, recently, the MultiAssay SIG (Special Interest Group) created the *MultiAssayExperiment class* (SIG, 2017) a data structure to manage and preprocess multiple assays for integrated genomic analysis. This data structure is now an input for all main functions of *ELMER* and can be generated by the *createMAE* function.

To perform *ELMER* analyses, users need to populate a *MultiAssayExperiment* with a DNA methylation matrix or *SummarizedExperiment* object from HumanMethylation450 BeadChip (HM450) or MethylationEPIC BeadChip (EPIC) platform; a gene expression matrix or *SummarizedExperiment* object for the same samples; a matrix mapping DNA methylation samples to gene expression samples; and a matrix with sample metadata (i.e. clinical data, molecular subtype, etc.). If TCGA data are used, the last two matrices will be automatically generated. If using non-TCGA data, the matrix with sample metadata should be provided with at least a column with a patient identifier and another one identifying its group which will be used for analysis, if samples in the methylation and expression matrices are not ordered and with same names, a matrix mapping for each patient identifier their DNA methylation samples and their gene expression samples should be provided to the *createMAE* function. Based on the genome of reference selected, metadata for the DNA methylation probes, such as genomic coordinates, will be added from Zhou, Laird and Shen (2016); and metadata for gene expression and annotation is added from ENSEMBL database (YATES *et al.*, 2015) using *biomaRt* (DURINCK *et al.*, 2009).

### 3.3.1.2 Selecting distal probes

Probes from HumanMethylationEPIC (EPIC) array and Infinium HumanMethylation450 (HM450) array are removed from the analysis if they have either internal SNPs close to the 3' end of the probe; non-unique mapping to the bisulfite-converted genome; or off-target hybridization due to partial overlap with non-unique elements (ZHOU; LAIRD; SHEN, 2017). This probe metadata information is included in *ELMER.data* package, populated from the source file at <http://zwdzwd.github.io/InfiniumAnnotation> (ZHOU; LAIRD; SHEN, 2017). To limit ELMER to the analysis of distal elements, probes located in regions of  $\pm 2kb$  around transcription start sites (TSSs) were removed.

### 3.3.1.3 Identification of differentially methylated CpGs (DMCs)

For each distal probe, samples of each group (group 1 and group 2) are ranked by their DNA methylation beta values, those samples in the lower quintile (20% samples with the lowest methylation levels) of each group are used to identify if the probe is hypomethylated in group 1 compared to group 2, using an unpaired one-tailed t-test. The 20% is a parameter to the *diff.meth*

function called *minSubgroupFrac*. For the (ungrouped) cancer case, this is set to 20% as in Yao *et al.* (2015), because we typically wanted to be able to detect a specific molecular subtype among the tumor samples; these subtypes often make up only a minority of samples, and 20% was chosen as a lower bound for the purposes of statistical power (high enough sample numbers to yield t-test p-values that could overcome multiple hypothesis corrections, yet low enough to be able to capture changes in individual molecular subtypes occurring in 20% or more of the cases.) This number can be set arbitrarily as an input to the *diff.meth* function and should be tuned based on sample sizes in individual studies. In the *Supervised* mode, where the comparison groups are implicit in the sample set and labeled, the *minSubgroupFrac* parameter is set to 100%. An example would be a cell culture experiment with 5 replicates of the untreated cell line, and another 5 replicates that include an experimental treatment.

To identify hypomethylated differentially methylated CpGs (DMCs), a one-tailed t-test is used to rule out the null hypothesis:  $\mu_{group1} \geq \mu_{group2}$ , where  $\mu_{group1}$  is the mean methylation within the lowest group 1 quintile (or another percentile as specified by the *minSubgroupFrac* parameter) and  $\mu_{group2}$  is the mean within the lowest group 2 quintile. Raw p-values are adjusted for multiple hypothesis testing using the Benjamini-Hochberg method (BENJAMINI; HOCHBERG, 1995b), and probes are selected when they had adjusted p-value less than 0.01 (which can be configured using the *pvalue* parameter). For additional stringency, probes are only selected if the methylation difference:  $\Delta = \mu_{group1} - \mu_{group2}$  was greater than 0.3. The same method is used to identify hypermethylated DMCs, except we use the *upper* quintile, and the opposite tail in the t-test is chosen.

### 3.3.1.4 Identification of putative target gene(s)

For each differentially methylated distal probe (DMC), the closest 10 upstream genes and the closest 10 downstream genes are tested for inverse correlation between methylation of the probe and expression of the gene. To select these genes, the probe-gene distance is defined as the distance from the probe to the transcription start site specified by the ENSEMBL gene level annotations (YATES *et al.*, 2015) accessed via the R/Bioconductor package [biomaRt](#) (DURINCK *et al.*, 2009; DURINCK *et al.*, 2005). By choosing a constant number of genes to test for each probe, our goal is to avoid systematic false positives for probes in gene rich regions. This is especially important given the highly non-uniform gene density of mammalian genomes. Thus, exactly 20 statistical tests were performed for each probe, as follows.

For each probe-gene pair, the samples (all samples from both groups) are divided into two groups: the *M* group, which consisted of the upper methylation quintile (the 20% of samples with the highest methylation at the enhancer probe), and the *U* group, which consists of the lowest methylation quintile (the 20% of samples with the lowest methylation.) The 20% ile cutoff is a configurable parameter *minSubgroupFrac* in the *get.pair* function. As with its usage in the *diff.meth* function, the default value of 20% is a balance, allowing for the identification of

changes in a molecular subtype making up a minority (i.e. 20%) of cases, while also yielding enough statistical power to make strong predictions. For larger sample sizes or other experimental designs, this could be set even lower.

For each candidate probe-gene pair, the Mann-Whitney U test is used to test the null hypothesis that overall gene expression in group M is greater than or equal than that in group U. This non-parametric test was used in order to minimize the effects of expression outliers, which can occur across a very wide dynamic range. For each probe-gene pair tested, the raw p-value  $P_r$  is corrected for multiple hypothesis using a permutation approach as follows. The gene in the pair is held constant, and  $x$  random methylation probes are chosen to perform the same one-tailed U test, generating a set of  $x$  permutation p-values  $P_p$ . We chose the  $x$  random probes only from among those that were "distal" (farther than 2kb from an annotated transcription start site), in order to draw these null-model probes from the same set as the probe being tested (SHAM; PURCELL, 2014). An empirical p-value  $P_e$  value was calculated using the following formula (which introduces a pseudo-count of 1):

$$P_e = \frac{\text{num}(P_p \leq P_r) + 1}{x + 1} \quad (3.1)$$

Notice that in the *Supervised* mode, no additional filtering is necessary to ensure that the *M* and *U* group segregate by sample group labels. The two sample groups are segregated by definition, since these probes were selected for their differential methylation, with the same directionality, between the two groups (Figure 17).

### 3.3.1.5 Characterization of chromatin state context of enriched probes using StateHub

Unlike version 1 of *ELMER*, we now consider *all* distal probes in the identification of regulatory elements. DNA methylation is known to affect several different classes of distal chromatin state element, including active enhancers, poised enhancers, super enhancers, and insulators. In order to provide a functional interpretation of the regulatory elements identified by *ELMER*, we perform a chromatin state enrichment analysis of the probes within significant probe-gene pairs, using the *statePaintR* tools from the [statehub.org](http://statehub.org) (COETZEE *et al.*, 2017), along with the new *FunciVar* package (in preparation on <https://github.com/Simon-Coetzee/funcivar>). Enrichment of the putative pairs within chromatin states is calculated against a background model that uses the distal probe set that the putative pairs are drawn from.

### 3.3.1.6 Motif enrichment analysis

In order to identify enriched motifs and potential upstream regulatory TFs, all probes with occurring in significant probe-gene pairs are combined for motif enrichment analysis. HOMER (Hypergeometric Optimization of Motif EnRichment) (HEINZ *et al.*, 2010) is used to find motif occurrences in a ±250bp region around each probe, using HOCOMOCO (HOmo sapiens

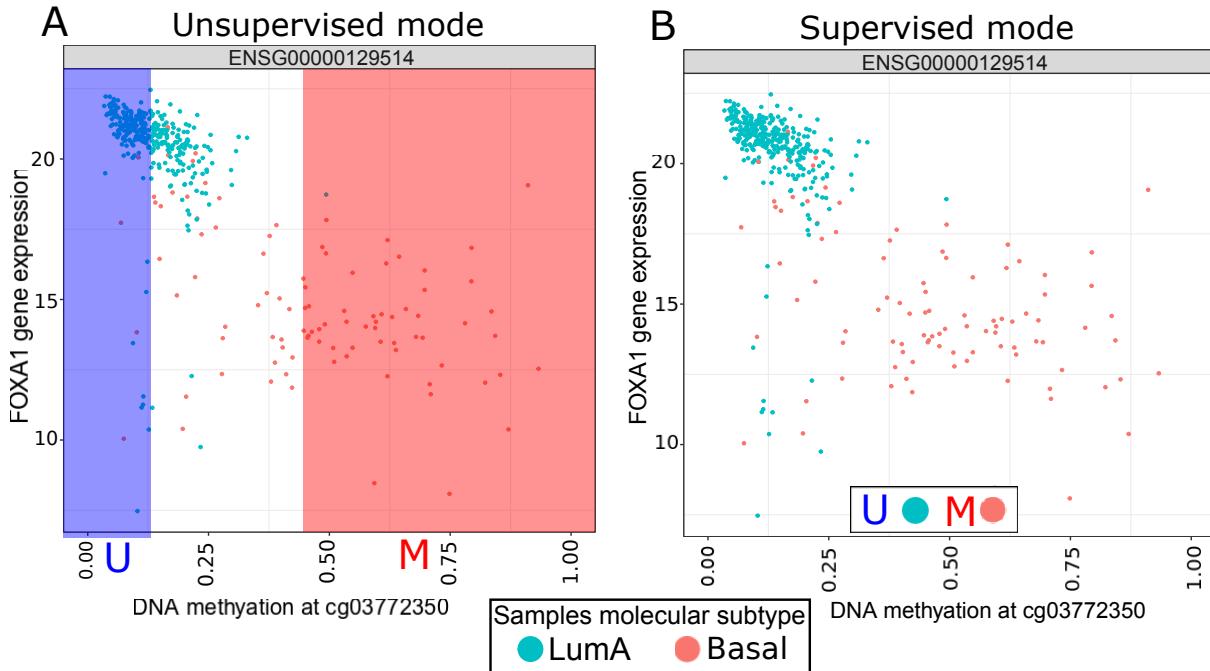


Figure 17 – Supervised mode maximizes statistical power. Difference of groups *U* and *M* definition in *supervised* and *unsupervised* mode. A: *unsupervised* mode; when minSubgroupFrac argument is set to 40%, the methylated group is defined as the highest quintile and the unmethylated group as the lowest quintile; B: *supervised* mode; methylated and unmethylated group are defined as one of the known molecular subtypes. For example, the unmethylated group is represented by all the LumA samples while the methylated group is represented by all the Basal samples. The t-test p-value achieved for the Unsupervised mode is  $8.3E - 25$ , while the Supervised mode is :  $1.43E - 43$ .

COmprehensive MOdel COllection) v11 (KULAKOVSKIY *et al.*, 2016) . Transcription factor (TF) binding models are available at <http://hocomoco.autosome.ru/downloads>. HOCOMOCO is the most comprehensive TFBS database and is consistently updated, marking an improvement over ELMER version 1. We have also improved the statistical analysis performed to use Fisher's exact test and Benjamini-Hochberg multiple hypothesis testing correction (see Supplemental Methods and Note). Importantly, we now characterize the Confidence Interval for the enrichment of each motif, and only select motifs that have an *FDR* < 0.05 and where the lower bound of the Odds Ratio 95% CI is greater than 1.1 (specified by option *lower.OR*).

#### *Identification of master regulator TFs*

When a group of enhancers is coordinately altered in a specific sample subset, this is often the result of an altered upstream *master regulator* transcription factor in the gene regulatory network. *ELMER* identifies master regulator TFs corresponding to each of the TF binding motifs enriched from the previous analysis step. For each enriched motif, *ELMER* takes the mean DNA methylation of all distal probes (in significant probe-gene pairs) that contain that motif occurrence (within a  $\pm 250bp$  region) and compares this mean DNA methylation to the expression of each gene annotated as a human TF. This step has been updated to work differently for the new *Supervised* and *Unsupervised* modes, using the same approach as described for the identification of putative target gene(s) step. This also can result in greater statistical power when

using the *Supervised* mode (see Supplemental Methods and Note for full details).

As in ELMER version 1, all human TFs are ranked by their  $-\log_{10}(Pr)$  values, and those falling within the top 5% of this ranking were considered candidate Master Regulators (see Figure 21, below). In addition to the top 5% of all TFs, we also highlight all TFs from the same DNA binding domain family. In ELMER version 1, these families were defined by a hard-coded association file that was hand-curated. A major improvement of ELMER 2.0 is that we now use the classifications from the TFClass database (WINGENDER; SCHOEPS; DÖNITZ, 2013), based on UniProt domains. Each TF is assigned to a family, as well as a more specific sub-family, of DNA binding domain proteins. For each significant motif, our output reports the most significant master regulator at both the family and sub-family level.

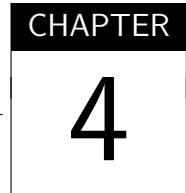
### **Data availability**

The TCGA data was downloaded from the NCI Genomic Data Commons (GDC) data portal (GROSSMAN *et al.*, 2016) using TCGAbiolinks R/Bioconductor package (COLAPRICO *et al.*, 2015; SILVA *et al.*, 2016). Gene annotations were retrieved from ENSEMBL (YATES *et al.*, 2015) database via biomaRt R/Bioconductor package (DURINCK *et al.*, 2005; DURINCK *et al.*, 2009). DNA methylation microarrays metadata were retrieved from <http://zwdzwd.github.io/InfiniumAnnotation> (ZHOU; LAIRD; SHEN, 2017). Transcription factor (TF) binding models can be downloaded at HOCOMOCO database (<http://hocomoco.autosome.ru/>) (KULAKOVSKIY *et al.*, 2016). The list of human TF can be accessed at <http://www.uniprot.org/> (APWEILER *et al.*, 2004). The classification of human transcription factors (TFs) can be viewed at <http://tfclass.bioinf.med.uni-goettingen.de/tfclass> (WINGENDER; SCHOEPS; DÖNITZ, 2013).

### **Software availability**

ELMER source code is available at <https://github.com/tiagochst/ELMER> and the auxiliary data files are available <https://github.com/tiagochst/ELMER.data>. *ELMER* is available under the GNU General Public License version 3 (GNU GPL3).





# CANCER DATA ANALYSIS

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This chapter presents results of the analysis of cancer data using the software and algorithms presented in the previous chapter. In section 4.1 we present some use cases using mainly the TCGAbiolinks package. In section 4.2 we present some data analysis using the ELMER package. In section 4.3, using the previous tools described we perform a glioma analysis focused specially in the two molecular subtypes G-CIMP-low and G-CIMP-high discovered by our laboratory and collaborators.

## 4.1 Use cases using TCGAbiolinks

In this section, we introduce and describe the utility and application of TCGAbiolinks through some use cases. In subsection 4.1.1 we show a lower-grade glioma downstream analysis with gene expression. In subsection 4.1.2 we show a downstream analysis integration of gene expression and methylation data of colon adenocarcinoma data, describing how to generate a starburst plot (NOUSHMEHR *et al.*, 2010), which was introduced to illustrate the results of integrating DNA methylation and gene expression data.

### 4.1.1 Lower-grade glioma downstream analysis with gene expression

For this case study, we used the recently available lower-grade glioma (LGG) data to investigate gene expression differences between the reported molecular subtypes (IDHmutant, IDHwildtype and IDHmutant codels) (PLATFORMS, 2015). In particular, we used TCGAbiolinks to download 293 samples profiled using messenger RNA expression (IlluminaHiSeq RNASeqV2) with available molecular subtypes. The data was normalized using the *TCGAanalyze\_Normalization* function and we applied three filters to remove features/mRNAs with low signals across samples, obtaining 4578, 4284 and 1187 mRNAs, respectively. A clustering analysis was then applied using the ConsensusClusterPlus package (WILKERSON; HAYES, 2010) which identified four distinct groups of samples (EC1-EC4) (Figure 18A). The survival curves for each cluster were generated using *TCGAanalyze\_survival* and are shown in Figure 18B. As expected,

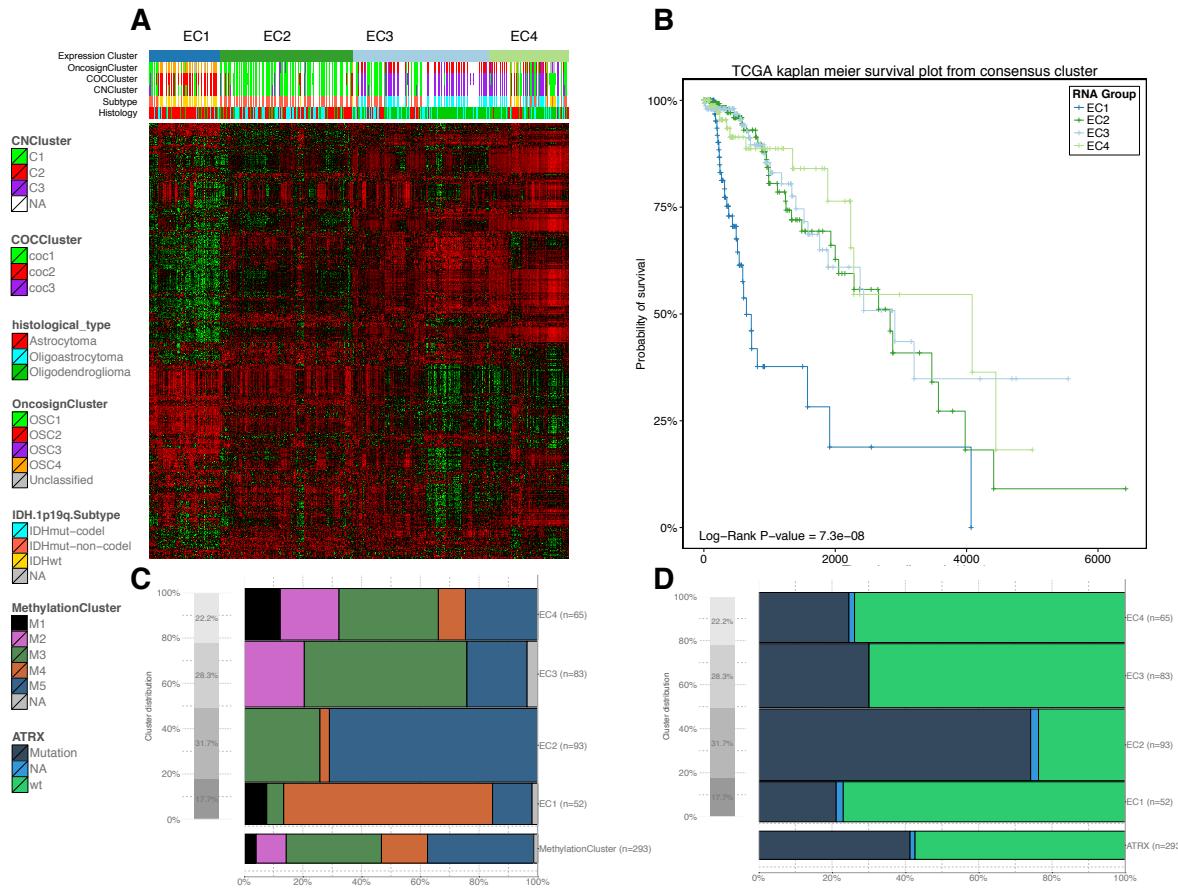


Figure 18 – Case study - Integrative (or Downstream) analysis of gene expression and clinical data from LGG disease with unsupervised clustering and crossing expression clusters with clinical and molecular information. **(A)** Heatmap of 1187 more variables genes clustered with tree  $k = 4$  in EC1, EC2, EC3, EC4. **(B)** Kaplan Meier survivals plot for EC clusters. **(C and D)** Distribution of the DNA Methylation clusters and ATRX mutation within the EC clusters.

each cluster effectively separated IDHwildtype tumors (EC1) from IDHmutant-non-codel (EC2) and IDHmutant-codel tumors (EC3 and EC4) (Figure 18). Additional biological subtypes (DNA methylation subtypes) were reproduced as expected (Figure 18D) (PLATFORMS, 2015).

#### 4.1.2 Downstream analysis integration of gene expression and methylation data

The DNA methylation of specific promoter CpG islands has the potential to influence gene expression. In this case study, we used TCGAbiolinks to examine the biological relationship between DNA methylation and gene expression in Colon adenocarcinoma (COAD). Using *GDCquery*, *GDCdownload* and *GDCprepare*, we obtained DNA methylation data (Infinium HumanMethylation450 and Infinium HumanMethylation27 platforms) and gene expression data (IlluminaGA RNASeqV2 platform) for the same TCGA COAD samples (NETWORK *et al.*, 2012). A supervised analysis was performed on the molecular subtypes CIMP-Low [CIMP.L] and CIMP-High [CIMP.H]. The gene expression analysis started by the identification of outliers,

followed by the normalization methods. Using *TCGAanalyze\_DEA*, 34 DEGs ( $\log_2FC \geq 3.0$  and  $FDR \leq 10^{-4}$ ) were identified. The result of this analysis is represented in a volcano plot (Figure 19A) created using *TCGAVisualize\_volcano*. For the DNA methylation analysis, using *TCGAanalyze\_DMR* we identified 73 CpG-methylated probes ( $\Delta\bar{\beta} \geq 0.25$  and  $FDR \leq 10^{-5}$ ; (Figure 19B). The DNA methylation and gene expression results were integrated as in the previous TCGA marker paper (NOUSHMEHR *et al.*, 2010; NETWORK *et al.*, 2012), by generating a starburst plot (Figure 19C) in which the x-axis is the  $\log_{10}$  of the correct P-value for DNA methylation and the y-axis is the  $\log_{10}$  of the correct P-value for the expression data. The starburst plot highlights nine distinct quadrants. To incorporate the DNA methylation difference cut-off into the graph, we highlighted genes that might have the potential for silencing due to epigenetic alterations. We highlighted five genes, EYA1, SIX2, ACSL6, OGDHL and SLC30A2, that showed a  $\Delta\bar{\beta} \geq 0.25$  and a  $\log_2FC \geq 3.0$  between CIMP.L and CIMP.H.

## 4.2 Use cases using ELMER

To exemplify the use of the new version we performed *ELMER* analysis on TCGA BRCA (Breast Invasive Carcinoma) data retrieved directly from the GDC server by TCGAbiolinks. Use Case 1 compares tumor samples to normal samples using the *Unsupervised* analysis mode, while Use Case 2 compares known BRCA molecular subtypes using the *Supervised* mode. The HTML output report containing the code and results of both analysis are available as supplemental material and at [https://github.com/tiagochst/ELMER\\_supplemental/raw/master/supplemental\\_files.zip](https://github.com/tiagochst/ELMER_supplemental/raw/master/supplemental_files.zip).

### 4.2.1 Use Case 1: Breast Invasive Carcinoma (Unsupervised mode)

We performed *ELMER* analysis comparing 778 Primary solid Tumor and 83 Solid Tissue Normal BRCA samples. In this use case we wanted to be able to detect non pre-determined molecular subtypes among the tumor samples, so the percentage of samples used to identify the differentially methylated probes in function *get.diff.meth* was set to 20% and the mode in function *get.pair* and in function *get.TFs* which was set to "unsupervised".

This analysis showed that the set of differentially methylated CpG probes (DMCs) hypomethylated in the tumors and linked to the expression of a nearby gene had an enrichment for TFBS motifs for FOX family transcription factors (FOXA2, FOXA3, FOXA1, etc.) (Figure 20). For the most highly enriched motif FOAX3, the master regulator analysis identified *FOXA1* as the top candidate among all TFs in the human genome (Figure 21), with the collaborating factors GATA3 and ESR1 as the next best candidates (Figure 22). This illustrates the important point that *in vitro* defined motifs from public TFBS databases are not always bound by the same TF family member *in vivo*. This was the same result of the analysis in YAO *et al.*, where we showed that ELMER identification of FOXA1, GATA3, and ESR1 were driven specifically by

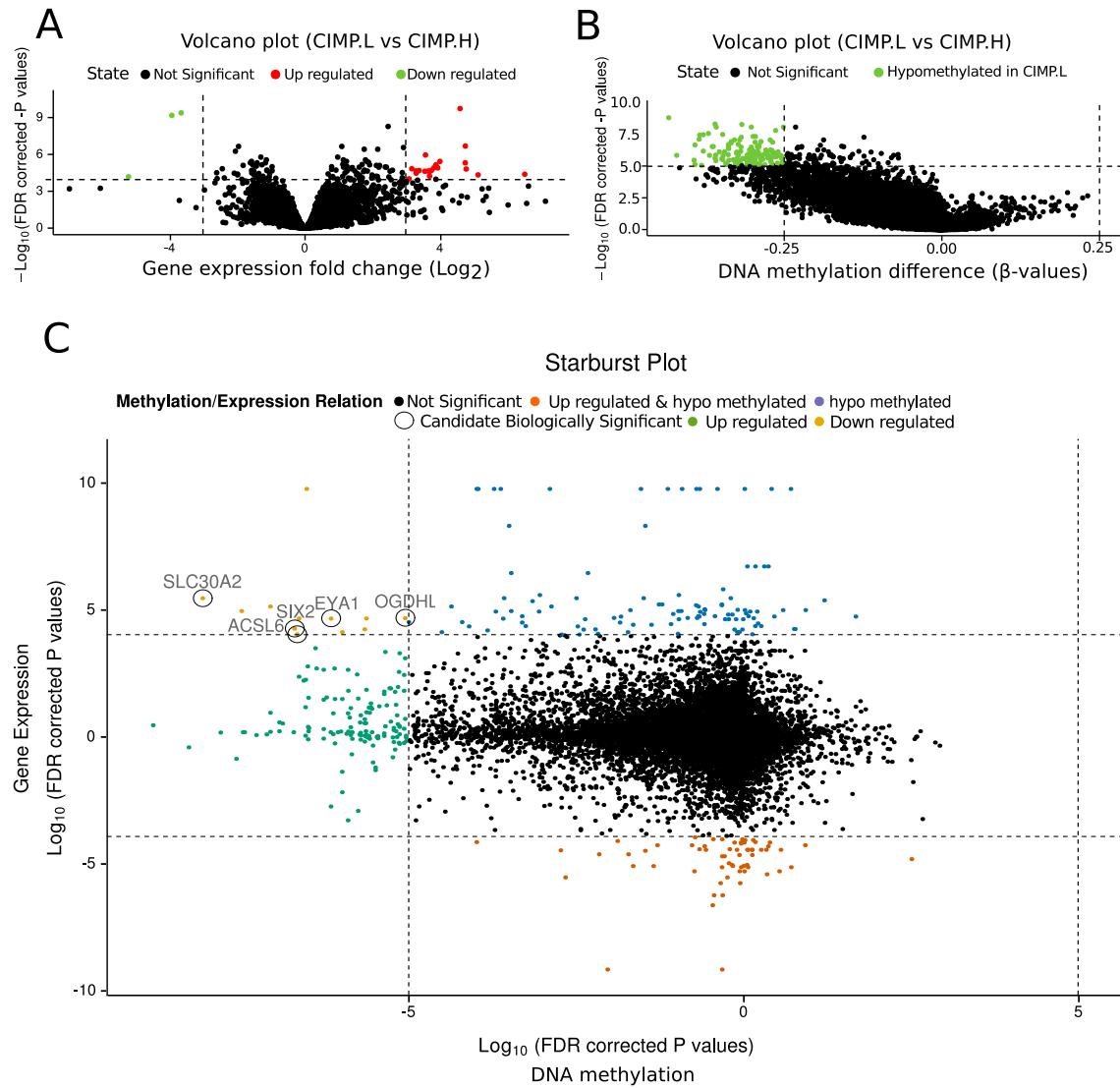


Figure 19 – Case study - Integrative analysis of gene expression and DNA methylation data from COAD disease, comparing groups CIMP.L and CIMP.H. **(A)** Expression volcano plot: fold change of expression data versus significance. **(B)** DNA methylation volcano plot: difference of DNA methylation versus significance. **(C)** Starburst plot: DNA methylation significance versus gene expression significance.

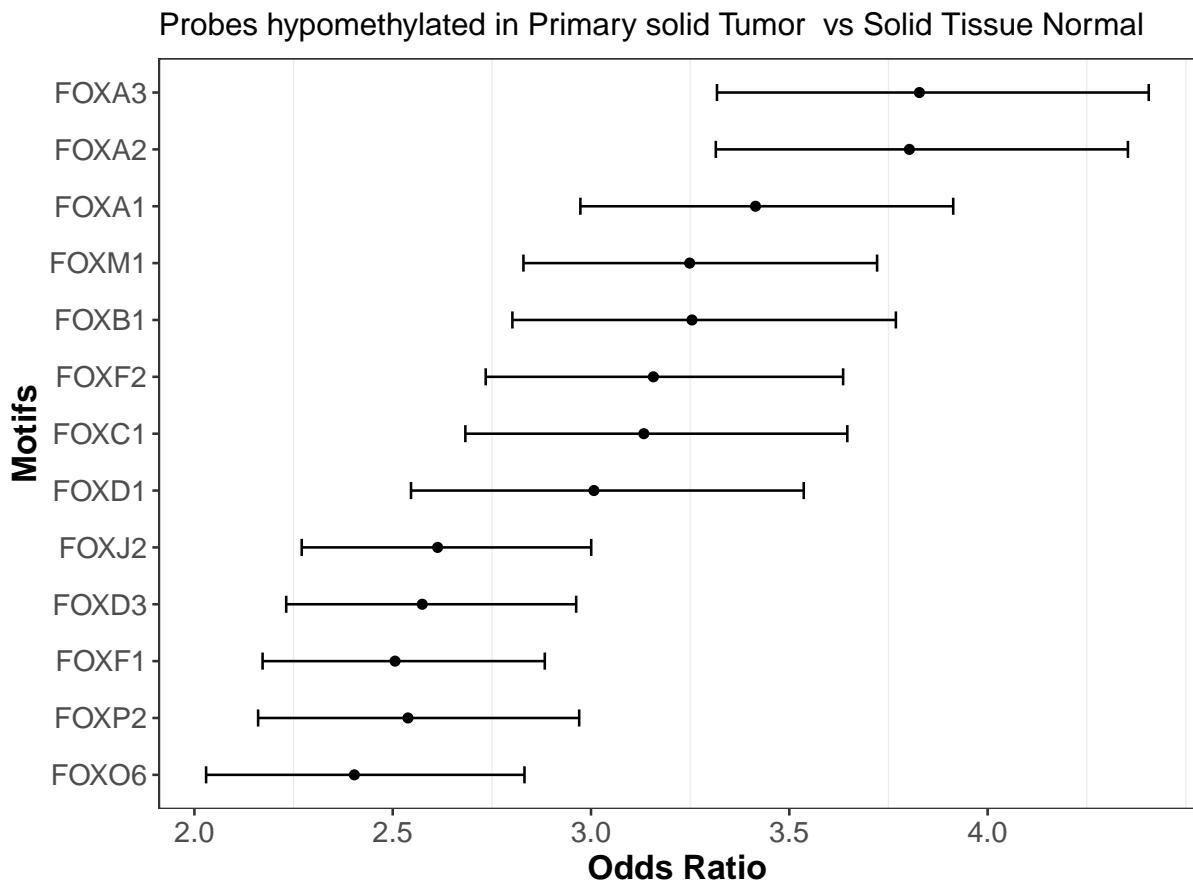


Figure 20 – Motif enrichment plot shows the enrichment levels ( $OR \geq 2.0$ ) for the most significant motifs based on the TCGA Breast Cancer Unsupervised analysis. A number of less significant motifs meet our default OR threshold of 1.1 ( $lower.or = 1.1$ ), which can be browsed in our full Supplemental output report.

the ER+ (luminal A and luminal B) tumors. However, our unsupervised analysis (both in YAO *et al.* and here) did not reveal Master Regulators for the other Breast Cancer molecular subtypes, such as Basal-like, HER2+, etc.

#### *Functional annotation of cis-regulatory modules using StateHub*

While ELMER version 1 was limited to searching within annotated enhancer elements, we have since found that this constraint was not necessary to achieve statistical power. Thus ELMER 2.0 by default searches *all* distal elements in the genome (distal elements are those greater than  $\pm 1kb$  from a TSS; By changing ELMER default settings, it is possible to analyze TSS-proximal probes, either together with distal probes or separately).

Because ELMER can now search essentially all probes on the array, it is important to understand the context of the probes that result from an ELMER analysis. Typically these are enhancer probes, but some regulatory changes may involve promoters, insulators, etc. We used the *StateHub* (COETZEE *et al.*, 2017) and *FunciVar* Bioconductor packages, to characterize enrichment of the various cell-type-specific chromatin states in the significant BRCA-hypomethylated probes (Figure 23).

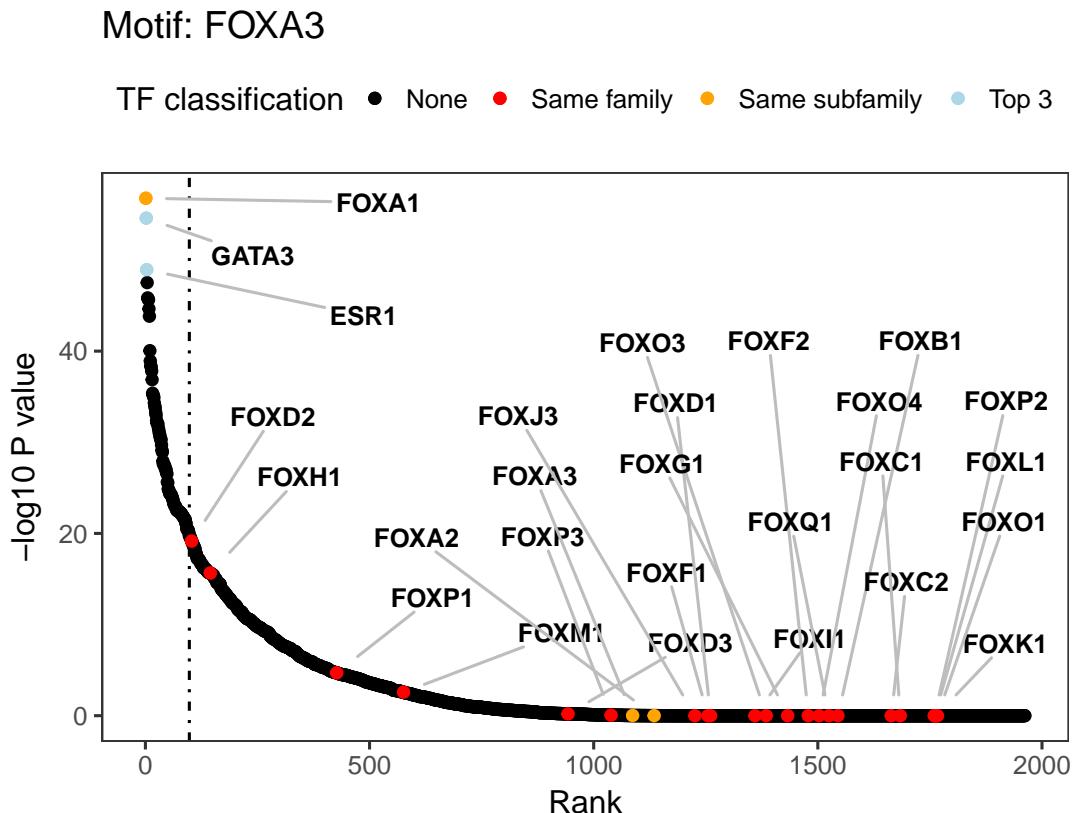


Figure 21 – TF ranking plot shows the statistical  $-\log_{10}(P\text{-value})$  assessing the anti-correlation level of candidate Master Regulator TF expression with average DNA methylation level for sites with the given motif (FOXA3). By default, the top 3 associated TFs (blue dots), and all of the TF family members (red dots) and subfamily members (orange dots) are labeled. The anti-correlation data for the top three candidates (FOXA1, GATA3, and ESR1) are derived from the data shown in Figure 22

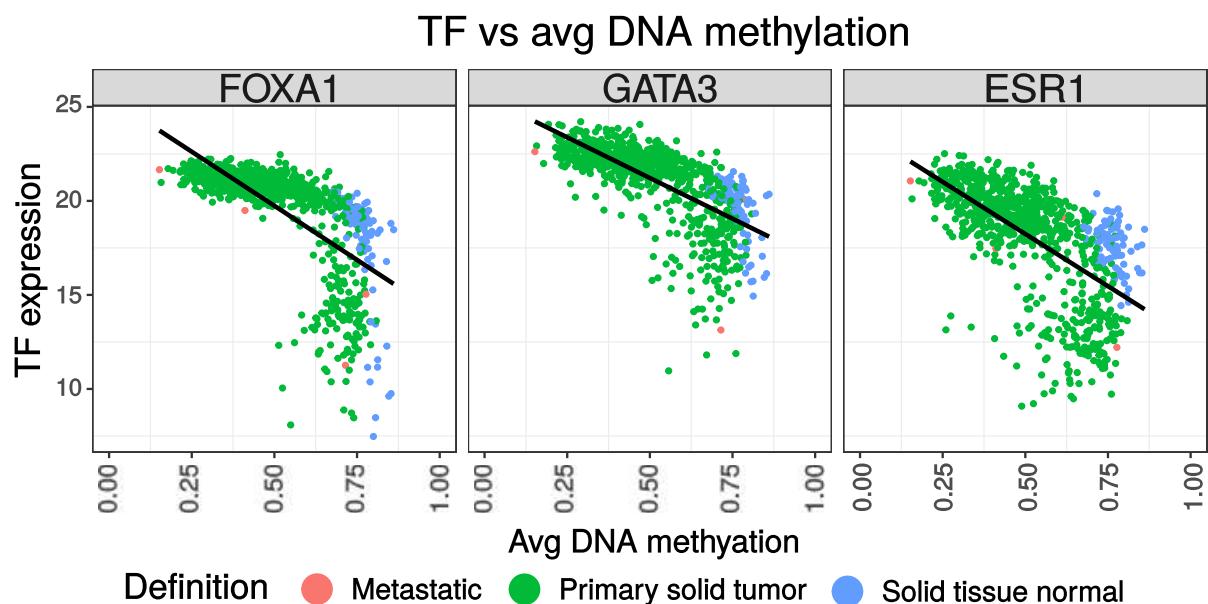


Figure 22 – FOXA1, GATA3 and ESR1 were identified as the most significant Master Regulator candidates for the top motif (FOXA3). All FOX factors belonging to the same TFClass binding family are highlighted.

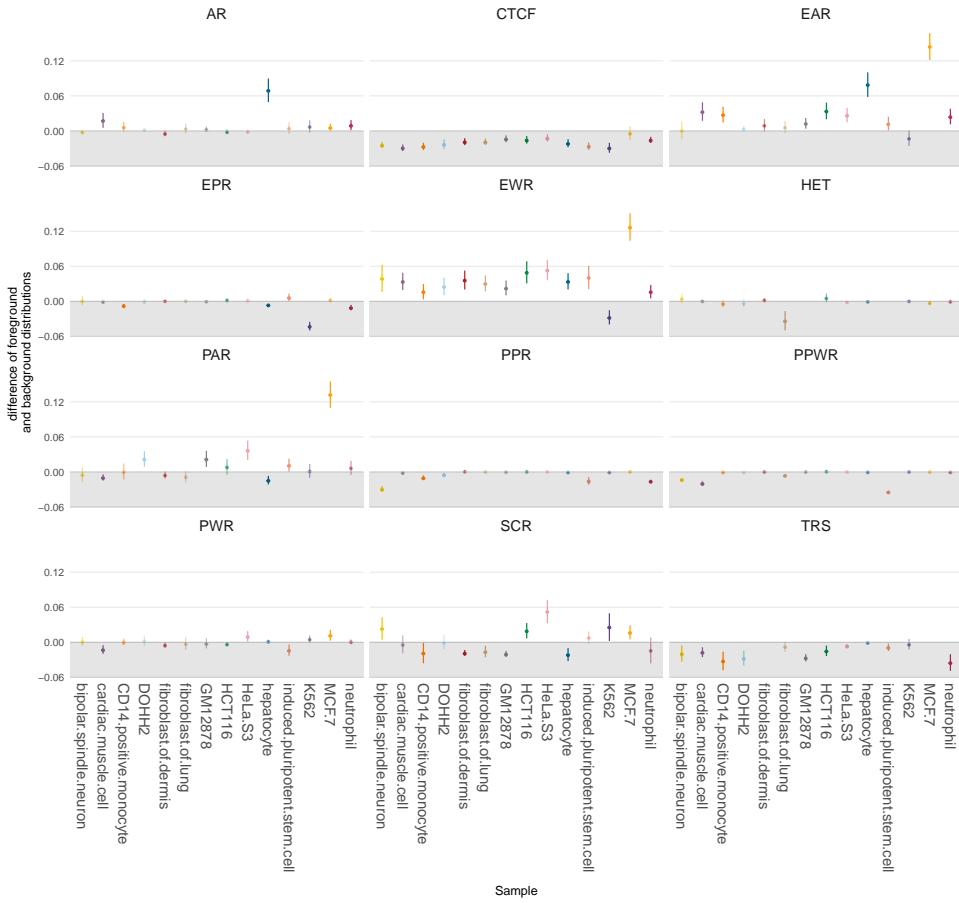


Figure 23 – Enrichment of paired probes and chromatin states of encode cells. The plot shows enrichment for enhancer active region, weak enhancer and active promoter region for MCF-7 cell. Acronyms - AR: Active region, EAR: active enhancer, EWR: Weak Enhancer, EPR: poised enhancer, PAR: active promoter, PWR: Weak Promoter, PPR: poised promoter, PPWR: Weak Poised Promoter, CTCF: architectural complex, TRS: transcribed, HET: heterochromatin, SCR: Polycomb Repressed Silenced. Y-axis shows the probability difference in overlap for the foreground class vs. random probes (Confidence Interval based on beta-binomial distribution).

Importantly, the MCF-7 cell line, and ER-positive breast cancer cell line, is much more strongly enriched for all enhancer and promoter classes than other cell types. As more reference cell types become available, this analysis will be useful in characterizing tumor GRN changes that reflect particular cell types or co-opted developmental programs.

#### Comparing inferred results with MCF-7 chIA-PET

As shown in Yao *et al.* (2015), we compared the putative pairs inferred to the chromatin loops derived from deep-sequenced ChIA-PET data from ER+ Breast Cancer MCF7 cells (LI *et al.*, 2012). First, we identify the number of *ELMER* pairs overlapping the ChIA-PET loops, then we repeat using randomly generated pairs with properties similar to the *ELMER* pairs. For each true *ELMER* probe in a probe-gene pair, we randomly select a different probe from the complete set of distal probes. We then choose the nth nearest gene to the random probe, where n is the same as the adjacency of the true *ELMER* probe (i.e. if the true probe is linked to the second gene upstream, the random probe will also be linked to its second gene upstream). Thus,

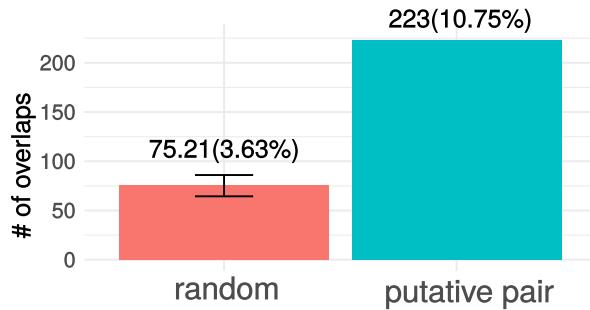


Figure 24 – The graph shows the comparison of the number of probe-gene pairs identified within MCF7 ChIA-PET data using the putative pairs from BRCA vs. random pairs

the random linkage set has both the same number of probes and the same number of linked genes as the true set. One hundred such random datasets were generated to arrive at a 95% CI ( $\pm 1.96 * SD$ ). The result is shown in Figure 24. ELMER pairs were enriched for ChIA-PET loops by roughly 3-fold over random pairs.

#### **4.2.2 Use Case 2: BRCA molecular subtypes analysis (Supervised mode)**

Several studies identified distinct molecular Breast Cancer molecular subtypes including luminal-like (Luminal A and Luminal B) subclasses, which are Estrogen receptor-positive (ER-positive), and the basal-like, ErbB2-positive and normal-like subclasses (ER-negative) (PEROU *et al.*, 2000; YERSAL; BARUTCA, 2014; SØRLIE *et al.*, 2001). We performed ELMER analysis comparing known molecular subtypes (Her2, Luminal A, Luminal B and Basal-like) using the TCGA BRCA dataset and classifications retrieved from (CIRIELLO *et al.*, 2015) using TCGABiolinks.

We performed pairwise analyses between different molecular subtypes. It is important to note that we expect these analyses to have increased statistical power over Unsupervised analyses, as illustrated above in Figure 17. One useful output plot is the comprehensive heatmap, which illustrates the identification of inverse correlated probe-gene pairs (Figure 25) for the LumA vs Basal-like analysis.

The unsupervised analysis of the same sample identified several Luminal type Master Regulators (MRs) such as FOXA1, GATA3, and ESR1. In order to identify MRs for the other subtypes, we created a table (Table 12) of candidate MRs identified by each pairwise ELMER run (complete results can be found in the supplemental HTML file described in the Supplementary Methods section).

Interestingly, several new MRs are identified for the Basal-like group, and these were mostly consistent in comparisons against Luminal and HER2+ subtypes. One group of MRs identified are the SOX10 and SOX9 TF signatures. For these signatures, the regulatory TF candidate identified are the SOX9 (Sry-related HMG box-9) TF and SOX11 (Sry-related HMG box-11)

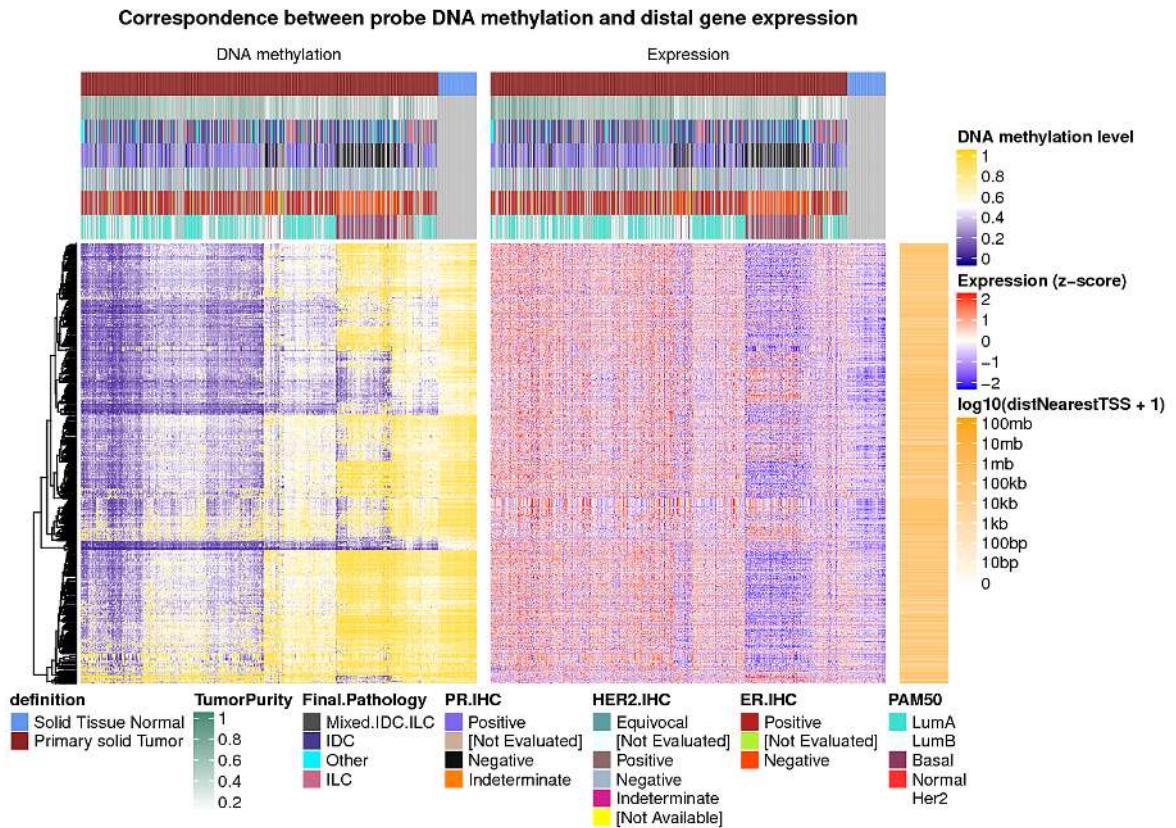


Figure 25 – The comprehensive heatmap view shows all probe / gene pairs identified by ELMER, clustered according to similarity. This plot is based on the Supervised analysis of LumA vs Basal-like Breast Cancer cases. The inverse correlation between methylation and expression can be observed.

TF; this correlation between basal-like and SOX11 was recently described by (SHEPHERD *et al.*, 2016) and *SOX9* was described by (GONG *et al.*, 2015). Most interestingly, we found KLF5 to be a consistently predicted MR for the Basal-like breast subtype. KLF5 is a master pluripotency factor of embryonic stem cells, and has been associated with a number of different cancers. In breast cancer, its overexpression has been linked to aggressive, ER-negative and basal-like breast cancers (BEN-PORATH *et al.*, 2008).

## 4.3 Glioma analysis

Enhancer Linking by Methylation/Expression Relationship (ELMER) analysis was performed to identify the molecular differences between the newly identified G-CIMP-low and G-CIMP-high subtypes of glioma recently described by Dr. Noushmehr and his lab (CECCARELLI *et al.*, 2016b). There is a particular interest in these molecular subtypes of glioma, since they have a great difference in both DNA methylation level and survival rate, with the G-CIMP-high subtype having a higher level of DNA methylation in some regions and a better survival rate. Thus, we seek to understand the molecular biology behind these changes leading to a worsening of survival.

Table 12 – Use case 2 - BRCA supervised analysis: Candidate master regulator TFs (MRs) for each molecular subtype found in a pairwise comparison. This table only includes MRs occurring in multiple pairwise comparisons.

<b>TF</b>	<b>LUMA (vs basal)</b>	<b>LUMB (vs basal)</b>	<b>Basal (vs LumB)</b>	<b>Basal (vs HER2)</b>	<b>HER2 (vs Basal)</b>
<b><i>AR</i></b>	x	x			
<b><i>BCL11A</i></b>			x	x	
<b><i>CEBPB</i></b>			x	x	
<b><i>E2F3</i></b>			x	x	
<b><i>EMX1</i></b>	x	x			
<b><i>ESR1</i></b>	x	x			
<b><i>ETV6</i></b>			x	x	
<b><i>FOXA1</i></b>	x	x			x
<b><i>FOXPI</i></b>	x	x			x
<b><i>GATA3</i></b>	x	x			x
<b><i>GLII</i></b>	x	x			
<b><i>HOXB1</i></b>	x	x			
<b><i>HOXB3</i></b>					x
<b><i>HOXC10</i></b>					x
<b><i>KLF5</i></b>			x	x	
<b><i>LMX1B</i></b>	x	x			
<b><i>NR2E3</i></b>	x	x			
<b><i>PBX1</i></b>		x			
<b><i>RARA</i></b>	x	x			
<b><i>SOX8</i></b>			x	x	
<b><i>SOX9</i></b>			x	x	
<b><i>SOX11</i></b>			x		
<b><i>ZNF467</i></b>	x	x	x	x	
<b><i>ZIC1</i></b>					

For this analysis, TCGA data from the NCI's Genomic Data Commons (GDC) was downloaded using our R/Bioconductor TCGAbiolinks package described in the previous chapter. Table 13 summarizes the number of samples in each group that have both deoxyribonucleic acid (DNA) methylation data for the Illumina HumanMethylation450 platform (HM450) and gene expression data (RNA-Seq) and Table 14 summarizes the main values for the ELMER arguments. This analysis was performed with data aligned against the genome of reference hg38 and in the "supervised" mode, which ensures that in all the steps the comparisons are made between each group using all their samples.

Table 13 – G-CIMP-high vs G-CIMP-low analysis: number of samples with both DNA methylation (HM450) and gene expression (RNA-seq) data.

Group	Number of samples
G-CIMP-high	233
G-CIMP-low	11

The DMCs (differentially methylated CpGs) analysis, which uses a one-tailed t-test to find probes with higher mean level in G-CIMP-high group compared to the G-CIMP-low one.

Table 14 – G-CIMP-high vs G-CIMP-low analysis: ELMER arguments values.

Step	Argument	Value
createMAE	genome	hg38
Pair probe-gene correlation/master TF	Mode	Supervised
All	minSubgroupFrac	100%
All	group1	GGimp-high
All	group2	GGimp-low
All	direction	hypermethylated probes
DMCs	min $\Delta\bar{\beta}$	0.3
DMCs	p-value adj cut-off	0.01
Pair probe-gene correlation	# permutations	10000
Pair probe-gene correlation	raw p-value cut-off	0.001
Pair probe-gene correlation	empirical p-values cut-off	0.001
Motif enrichment	minimum # probes	10
Motif enrichment	lower OR	1.1

The result of this analysis is summarized in Figure 26. The volcano plot shows the difference of DNA methylation ( $\Delta\bar{\beta}$ ) versus significance  $-\log_{10}(\text{FDR corrected P-values})$ . Using as default cutoffs  $\Delta\bar{\beta} \geq 0.3$  and  $-\log_{10}(\text{FDR corrected P-values}) \leq 0.01$ , 7540 probes were identified to be differentially methylated.

Those differentially methylated probes were then paired with their 10 upstream and 10 downstream genes and those significantly anti-correlated were selected (genes have higher expression levels when the DNA methylation levels decreases). Using as cutoffs  $-\log_{10}(\text{raw P-values}) \leq 0.001$  and  $-\log_{10}(\text{Permuted corrected P-values}) \leq 0.001$  for a 10000 permutation correction approach, 886 pairs of gene and probes were identified to be significantly anti-correlated. The result of this analysis is summarized in Figure 27. The heatmap is composed of other three heatmaps, one heatmap for the DNA methylation levels, one heatmap for the gene expression levels and one with the distance between the gene and the probe.

Table 15 – G-CIMP-high vs G-CIMP-low analysis: number of distal probes, significant differentially methylated probes, significant anti-correlated paired gene-probes and enriched motifs

Result	Value
Distal probes	135374
Significant differentially methylated probes	7540
Significant anti-correlated paired gene-probes	886
Enriched motifs	84

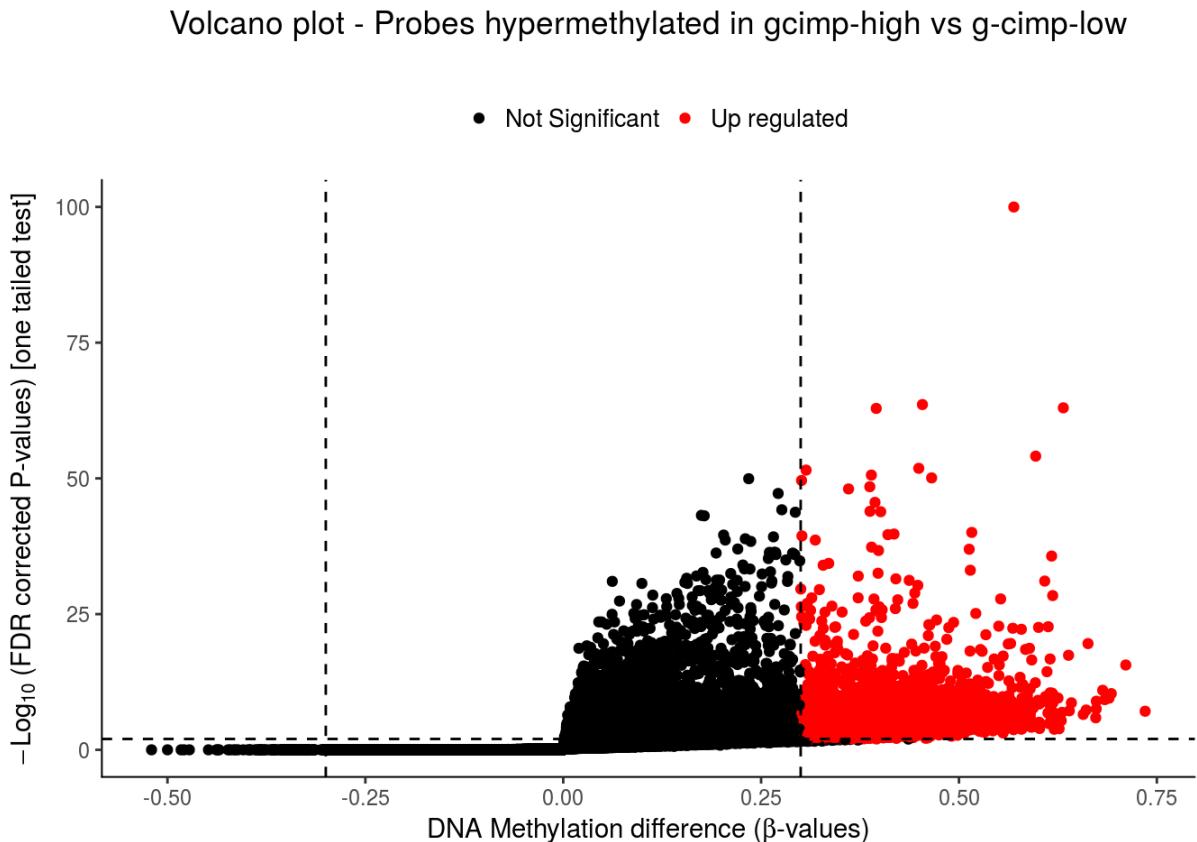


Figure 26 – G-CIMP analysis: DNA methylation Volcano plot. Difference of DNA methylation ( $\min \Delta \beta$ ) versus significance  $-\log_{10}(\text{FDR corrected P-values})$ .

The motif enrichment analysis results are summarized in Figure 28, which shows the Odds Ratio (OR) (x axis) for the enriched motifs, and in Table 16, which shows the candidate regulatory TFs whose expression anti-correlated with the DNA methylation level on the probes of each enriched motifs. From the most anti-correlated ones, ELMER uses the TFClass classification (WINGENDER *et al.*, 2015) to identifies which TFs are known to bind in those motifs. This classification has two levels, family and subfamily, which groups motifs with a similar signature. Depending on the motif, its family classification might have a very similar signature, otherwise, the subfamily classification is the most indicated.

For example, in Table 16 the motif HXD3 has as potential TF candidate the HOXD13 if we consider the family Transcription Factor (TF) classification and the HOXD3 TF candidate considering the subfamily TF classification. Figure 29 shows the motif signature for the HOX-related factors family from HOMO sapiens COmprehensive MOdel COllection (HOCOMOCO) database. The transcription factors Homeobox D13 (HOXD13) and Homeobox D3 (HOXD3) are in the same family (HOX-related factors) but in different subfamilies.

The analysis identified several candidate master regulator TF including PAX3, HMX3, FOXM1, HOXD13, HOXD11, HOXC10 and SOX11. Some of them have already been reported in previous research presented below.

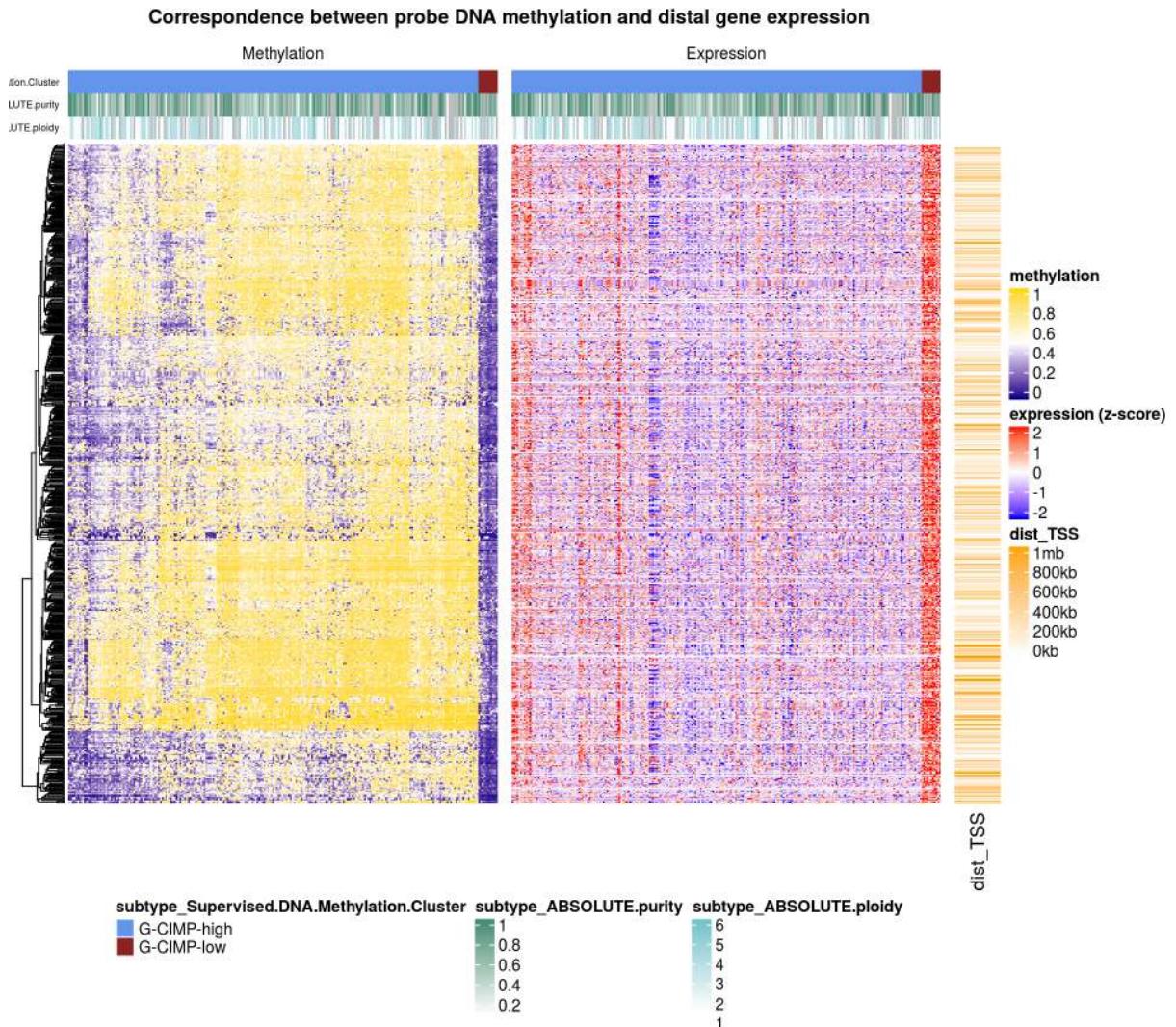


Figure 27 – G-CIMP analysis: Heatmap of paired probes and distal genes. The first heatmap (left one) shows DNA methylation  $\beta$  levels ranging from 0 (non-methylated) up to 1 (methylated probes). The second heatmap (middle one) shows z-scores for gene expression levels (standard deviations from each gene means). The last heatmap (right heatmap) shows the distance between the probe and gene anti-correlated. The top annotation in the heatmap was retrieved from Ceccarelli *et al.* (2016a)

MORALES-DELGADO *et al.* showed that HMX3 is a key player in the activation and maintenance of Gsh1 expression which resulted in subsequent downstream generation of GHRH (growth hormone-releasing hormone) expression in neurons (MORALES-DELGADO *et al.*, 2014). Furthermore, GHRH antagonists in glioblastoma cells showed efficacy of those drugs for experimental treatment of glioblastoma (POZSGAI *et al.*, 2010).

LEE *et al.* reported that HOXD13 is substantially up regulated in GBMs (Glioblastomas) and may contribute to malignancy. Also, PAX3 and FOXM1 were shown to be essential for gliomagenesis by XIA *et al.* and WANG *et al.*, respectively.

However, contrary to what was expected, ?? showed that SOX11 prevents gliomagenesis, which may indicate a false positive in our analyzes.

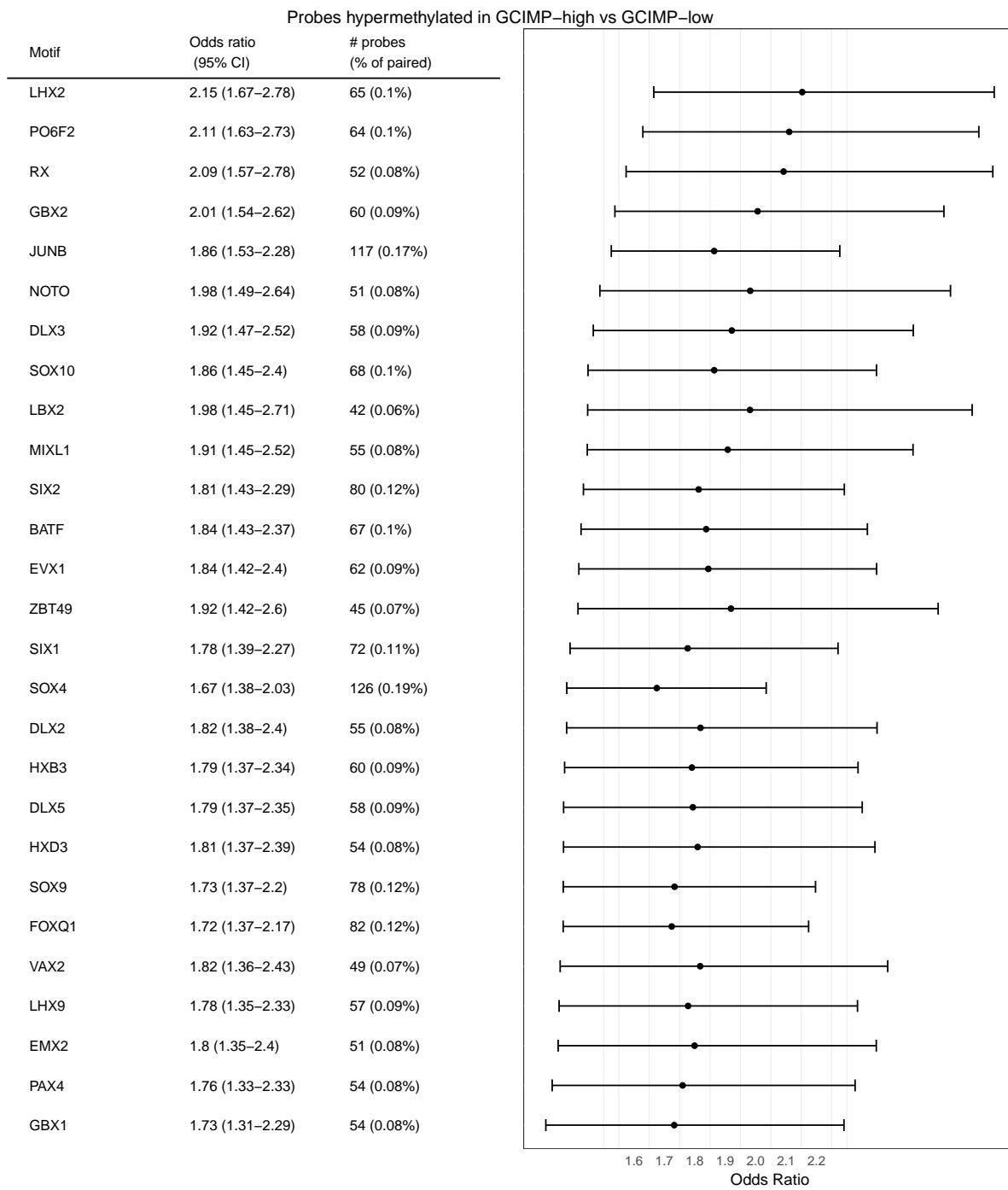


Figure 28 – Motif enrichment analysis: Odds Ratio (x-axis) for the selected motifs with lower OR above 1.3. The range shows the 95% confidence interval for each Odds Ratio.

To validate these findings, biological experiments are needed. By either knocking down these TFs or by regulating the DNA methylation levels of those binding regions will be able to verify if the downstream genes are being regulated.

Table 16 – TF ranking analysis: statistic For each enriched motif the anti-correlation level of all human TFs expression level with average DNA methylation level at sites with a given motif was access and ranked by the  $-\log_{10}(P_{value})$ , the most relevant one that belongs to the same family as the motif is shown in column *top.potential.TF.family* while the most relevant within the same sub-family classification is shown in column *top.potential.TF.subfamily*

motif	top.potential.TF.family	top.potential.TF.subfamily
LHX2_HUMAN.H11MO.0.A	NA	NA
PO6F2_HUMAN.H11MO.0.D	NA	NA
RX_HUMAN.H11MO.0.D	NA	NA
GBX2_HUMAN.H11MO.0.D	HOXD13	NA
JUNB_HUMAN.H11MO.0.A	NA	NA
NOTO_HUMAN.H11MO.0.D	HMX3	NA
DLX3_HUMAN.H11MO.0.C	HMX3	NA
SOX10_HUMAN.H11MO.0.B	SOX11	NA
LBX2_HUMAN.H11MO.0.D	HMX3	NA
MIXL1_HUMAN.H11MO.0.D	NA	NA
SIX2_HUMAN.H11MO.0.A	NA	NA
BATF_HUMAN.H11MO.1.A	NA	NA
EVX1_HUMAN.H11MO.0.D	HOXD13	NA
ZBT49_HUMAN.H11MO.0.D	ZSCAN16	ZSCAN16
SIX1_HUMAN.H11MO.0.A	NA	NA
SOX4_HUMAN.H11MO.0.B	SOX11	SOX11
DLX2_HUMAN.H11MO.0.D	HMX3	NA
HXB3_HUMAN.H11MO.0.D	HOXD13	HOXD3
DLX5_HUMAN.H11MO.0.D	HMX3	NA
HXD3_HUMAN.H11MO.0.D	HOXD13	HOXD3
SOX9_HUMAN.H11MO.0.B	SOX11	NA
FOXQ1_HUMAN.H11MO.0.C	FOXM1	NA
VAX2_HUMAN.H11MO.0.D	HMX3	NA
LHX9_HUMAN.H11MO.0.D	NA	NA
EMX2_HUMAN.H11MO.0.D	HMX3	NA
PAX4_HUMAN.H11MO.0.D	PAX3	NA
GBX1_HUMAN.H11MO.0.D	HOXD13	NA

Model	LOGO	Transcription factor	Quality	TF family	TF subfamily
			D	HOX-related	
HXD3_HUMAN.H11MO.0.D		HOXD3 (GeneCards)	D	HOX-related factors{3.1.1}	HOX3{3.1.1.3}
HXD4_HUMAN.H11MO.0.D		HOXD4 (GeneCards)	D	HOX-related factors{3.1.1}	HOX4{3.1.1.4}
HXA5_HUMAN.H11MO.0.D		HOXA5 (GeneCards)	D	HOX-related factors{3.1.1}	HOX5{3.1.1.5}
HXA7_HUMAN.H11MO.0.D		HOXA7 (GeneCards)	D	HOX-related factors{3.1.1}	HOX6-7{3.1.1.6}
HXB6_HUMAN.H11MO.0.D		HOXB6 (GeneCards)	D	HOX-related factors{3.1.1}	HOX6-7{3.1.1.6}
HXC6_HUMAN.H11MO.0.D		HOXC6 (GeneCards)	D	HOX-related factors{3.1.1}	HOX6-7{3.1.1.6}
HXC8_HUMAN.H11MO.0.D		HOXC8 (GeneCards)	D	HOX-related factors{3.1.1}	HOX8{3.1.1.7}
HXD8_HUMAN.H11MO.0.D		HOXD8 (GeneCards)	D	HOX-related factors{3.1.1}	HOX8{3.1.1.7}
HXA11_HUMAN.H11MO.0.D		HOXA11 (GeneCards)	D	HOX-related factors{3.1.1}	HOX9-13{3.1.1.8}
HXC10_HUMAN.H11MO.0.D		HOXC10 (GeneCards)	D	HOX-related factors{3.1.1}	HOX9-13{3.1.1.8}
HXC11_HUMAN.H11MO.0.D		HOXC11 (GeneCards)	D	HOX-related factors{3.1.1}	HOX9-13{3.1.1.8}
HXC12_HUMAN.H11MO.0.D		HOXC12 (GeneCards)	D	HOX-related factors{3.1.1}	HOX9-13{3.1.1.8}
HXC13_HUMAN.H11MO.0.D		HOXC13 (GeneCards)	D	HOX-related factors{3.1.1}	HOX9-13{3.1.1.8}
HXD10_HUMAN.H11MO.0.D		HOXD10 (GeneCards)	D	HOX-related factors{3.1.1}	HOX9-13{3.1.1.8}
HXD11_HUMAN.H11MO.0.D		HOXD11 (GeneCards)	D	HOX-related factors{3.1.1}	HOX9-13{3.1.1.8}
HXD12_HUMAN.H11MO.0.D		HOXD12 (GeneCards)	D	HOX-related factors{3.1.1}	HOX9-13{3.1.1.8}
HXD13_HUMAN.H11MO.0.D		HOXD13 (GeneCards)	D	HOX-related factors{3.1.1}	HOX9-13{3.1.1.8}

Figure 29 – HOCOMOCO V11: HOX-related factors family. Transcription factors HOXD13 and HOXD3 are in the same family (HOX-related factors) but in different subfamilies.



# CONCLUSION

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## 5.1 Conclusions

The main goal of this project has been to develop tools and workflows to perform integrative analysis as well as their application in the analysis of public cancer data. The main motivation was that the integration of data from different databases, which are stored in different formats and with different biological meanings, is both a complex process from the computational point of view, because dealing with a large amount of data requires the optimization of the use of computational resources, as well as from the biological point of view, because it is a biological process still under study, often unknown, or even without a scientific consensus.

## 5.2 Conclusions and future studies

In this section, we list the conclusions and future studies for the results reported in the previous chapters.

### 5.2.1 *Conclusions and future works of TCGAbiolinks*

This work was an effort towards the search, download, and processing of data from the NCI's Genomic Data Commons (GDC) data portal for downstream analysis. It offered several single-dataset exploratory analysis such as clustering methods, box plots, differential gene expression analysis, differential methylated CpGs (DMCs) analysis, survival analysis, and some integrative analysis such as the correlation of the regions differently methylated with levels of expression of the nearest gene to identify those affected by the DNA methylation, and gene set enrichment analysis. As future works, the improvement of the integrative analysis performed by the starburst plot is suggested. This method which integrates the results from two different analysis DEA (differential expression analysis) and DMR (differential methylated regions), should compare the gene expression and DNA methylation of the same sample instead of using the average levels of all samples in a group.

### 5.2.2 ***Conclusions and future works of TCGAbiolinksGUI***

This work created a Graphical User Interface (GUI) to our command-line tools to help users without programming knowledge to perform deeper downstream analysis. Possible improvements in future work include improving the import of external user data and improving the tool modularization, so that the addition of new menus and packages are facilitated.

### 5.2.3 ***Conclusions and future works of ELMER***

This work was an effort towards an integrative analysis using RNA-seq, DNA methylation, and histone marks to identify a candidate regulatory network. ELMER identifies distal regions with a difference in DNA methylation and correlates them with the gene expression levels of upstream and downstream genes. A motif enrichment analysis is performed on the regulatory regions that affect the expression of distant genes to identify potential regulatory TF candidates. As future work, we suggest expanding the algorithm to consider promoter regions, accepting whole-genome bisulfite sequencing (WGBS) data in DNA methylation analysis, and using mutation information instead of DNA methylation to identify regulatory regions mutated that might have affected the regulation of the upstream/downstream genes.

## 5.3 Publications, presentations, and software of the Doctorate Period

The work produced during the Doctorate period in form of scientific articles, softwares and conference presentations is shown in the next subsections. The published scientific articles were divided into two groups, 1) the first authorship, and 2) co-authorship. The published manuscripts are organized into: "First-authored papers", "Co-authored papers", "First-authored softwares" and "Co-authored softwares". The subsection "workshops and workflows" contains all the material created to help users to use the developed tools. Finally, the subsection "Conferences & presentations" list all oral and poster presentations in international and national conferences.

### 5.3.1 ***First-authored papers***

- COLAPRICO, A.; SILVA, T. C.; OLSEN, C.; GAROFANO, L.; CAVA, C.; GAROLINI, D.; SABEDOT, T. S.; MALTA, T. M.; PAGNOTTA, S. M.; CASTIGLIONI, I.; CECCARELLI, M.; BONTEMPI, G.; NOUSHMEHR, H. TCGAbiolinks: an r/bioconductor package for integrative analysis of tcga data. **Nucleic Acids Research**, v. 44, n. 8, p. e71, 2016. Available at: <http://nar.oxfordjournals.org/content/44/8/e71.abstract>.

- The main contributions to the work were the structuring of the package according to the standards of the Bioconductor project, the creation of all data functions (query,

download and prepare) and some function for analysis and visualization (survival analysis, DNA methylation analysis and plots and integration of DNA methylation and gene expression and visualization in a starburst plot function), the creation of unitary tests, the creation of the documentation for the functions listed above, and the creation of two use case.

- SILVA, T.; COLAPRICO, A.; OLSEN, C.; D'ANGELO, F.; BONTEMPI, G.; CECCARELLI, M.; NOUSHMEHR, H. TCGA workflow: Analyze cancer genomics and epigenomics data using bioconductor packages [version 2; referees: 1 approved, 1 approved with reservations]. **F1000Research**, v. 5, n. 1542, 2016.
  - The main contributions to the work were the development and testing of the sections "Experimental data", "DNA methylation analysis", "Motif analysis" and "Integrative analysis", the creation and maintenance of the workflow version available in the Bioconductor website.
- SILVA, T. C.; COLAPRICO, A.; OLSEN, C.; BONTEMPI, G.; CECCARELLI, M.; BERMAN, B. P.; NOUSHMEHR, H. Tcgabiolinksgui: A graphical user interface to analyze gdc cancer molecular and clinical data. **bioRxiv**, Cold Spring Harbor Labs Journals, 2017. Available at: <http://www.biorxiv.org/content/early/2017/08/17/147496>.
  - The main contributions to the work were the creation of the graphical user interface structure and all the menus except the "Transcriptomic analysis - Network of inference and differential expression analysis" menus, the creation of the documentation, docker image, and tutorials (PDF and youtube videos).
- SILVA, T. C.; COETZEE, S. G.; YAO, L.; HAZELETT, D. J.; NOUSHMEHR, H.; BERMAN, B. P. Enhancer linking by methylation/expression relationships with the r package elmer version 2. **bioRxiv**, Cold Spring Harbor Labs Journals, p. 148726, 2017.
  - The main contributions to the work were the code refactoring to provide greater stability, performance, and ease of use, the alteration of the main data structure to a standard Bioconductor data structure, the integration of the ELMER and TCGAbiolinks tools to import data from GDC, the creation of a graphical user interface and interactive HTML reports, the expansion of the algorithm to consider a supervised analysis mode, the creation of the case of study in the article, the restructuring of all the documentation by changing from a PDF format to an HTML and the addition of unit tests to the tool.

### **5.3.2 Co-authored papers**

- LIN, D.-C.; DINH, H. Q.; XIE, J.-J.; MAYAKONDA, A.; SILVA, T. C.; JIANG, Y.-Y.; DING, L.-W.; HE, J.-Z.; XU, X.-E.; HAO, J.-J.; WANG, M.-R.; LI, C.; XU, L.-Y.; LI,

E.-M.; BERMAN, B. P.; KOEFFLER, H. P. Identification of distinct mutational patterns and new driver genes in oesophageal squamous cell carcinomas and adenocarcinomas. **Gut**, BMJ Publishing Group, 2017. ISSN 0017-5749. Available at: <http://gut.bmj.com/content/early/2017/09/02/gutjnl-2017-314607>.

- The main contribution to the work was to perform the integrative analysis of esophageal cancer using the ELMER package.
- CECCARELLI, M.; BARTHEL, F.; MALTA, T.; SABEDOT, T.; SALAMA, S.; MURRAY, B.; AL., O. M. et. Molecular profiling reveals biologically discrete subsets and pathways of progression in diffuse glioma. **Cell**, v. 164, n. 3, p. 550 – 563, 2016. ISSN 0092-8674. Available at: <http://www.sciencedirect.com/science/article/pii/S009286741501692X>.
  - The main contributions to the work were the validation of the results found by reanalyzing all DNA methylation data and clinical data. In addition, the analysis methods used in this article have been made available in the TCGAbiolinks package.
- MALTA, T. M.; SOUZA, C. F.; SABEDOT, T. S.; SILVA, T. C.; MOSELLA, M. Q.; KALKANIS, S. N.; SNYDER, J.; CASTRO, A. V. B.; NOUSHMEHR, H. Glioma cpg island methylator phenotype (g-cimp): Biological and clinical implications. **bioRxiv**, Cold Spring Harbor Labs Journals, p. 169680, 2017.
  - The main contribution to the work was to help in writing about bioinformatics tools and analysis.
- CAVA, C.; COLAPRICO, A.; BERTOLI, G.; GRAUDENZI, A.; SILVA, T. C.; OLSEN, C.; NOUSHMEHR, H.; BONTEMPI, G.; MAURI, G.; CASTIGLIONI, I. Spidermir: An r/bioconductor package for integrative analysis with mirna data. **International journal of molecular sciences**, Multidisciplinary Digital Publishing Institute, v. 18, n. 2, p. 274, 2017.
  - The main contributions to the project were the integration of the tools TCGAbiolinks and SpidermiR, the structuring of the code as a package and help in the creation of the documentation.
- GOOD, E. E.; MCCORMACK, E. P.; SILVA, T. C.; PARONETT, E. M.; NOUSHMER, H.; LEE, N. H.; MAYNARD, T. M.; LAMANTIA, A. S.; SHERMAN, J. H. Foxj1, a potential biomarker for glioma. Manuscript in prep.
  - The main contributions to the study were differential gene expression analysis for samples that had high expression of the FOXJ1 gene compared to those with low expression and the integration of the results with copy number alteration and mutation data using TCGAbiolinks.

### 5.3.3 First-authored software

**TCGAbiolinks (version 2.0)** An R/Bioconductor package for integrative analysis of TCGA data. Published in Bioconductor <http://bioconductor.org/packages/TCGAbiolinks/>. Source code available in GitHub <https://github.com/BioinformaticsFMRP/TCGAbiolinks>.

**TCGAbiolinksGUI** A Graphical User Interface to analyze cancer genomics and epigenomics data. Published in GitHub <https://github.com/BioinformaticsFMRP/TCGAbiolinksGUI>.

**ELMER (version 2.0)** Enhancer Linking by Methylation/Expression Relationship (ELMER) is a package to identify tumor-specific changes in DNA methylation within distal enhancers, and link these enhancers to downstream target genes. Published in Bioconductor <http://bioconductor.org/packages/ELMER/>. Source code available in GitHub <https://github.com/tiagochst/ELMER>.

### 5.3.4 Co-authored software

**SpidermiR: An R/Bioconductor package for integrative network analysis with miRNA data**

Published in Bioconductor <http://bioconductor.org/packages/SpidermiR/>. Source code available in GitHub <https://github.com/claudiacava/SpidermiR>.

### 5.3.5 Workshops and workflows

**Workshop** Integrative analysis workshop with TCGAbiolinks and ELMER. Dana Farber Cancer Institute, Boston, MA. Link to workshop: <https://bioinformaticsfmrp.github.io/Bioc2017.TCGAbiolinks.ELMER/index.html>. Source code available in GitHub <https://github.com/BioinformaticsFMRP/Bioc2017.TCGAbiolinks.ELMER>.

**Workflow** TCGA Workflow: Analyze cancer genomics and epigenomics data using Bioconductor packages. Available at <https://www.bioconductor.org/help/workflows/TCGAWorkflow/>. Source code available in GitHub <https://github.com/BioinformaticsFMRP/TCGAWorkflow>.

### 5.3.6 Conferences & presentations

**SNOLA 2016 - Update on Neuro-Oncology - Oral presentation - 04/19/2016** "EPIGENOMIC AND TRANSCRIPTOMIC ANALYSIS OF ADULT GLIOMA REVEALS CANDIDATE DRIVER TRANSCRIPTION FACTORS INVOLVED IN GLIOMA PROGRESSION." Windsor Barra Hotel, Rio de Janeiro - Brazil

**Chromatin and Epigenetics - Poster presentation - 5/5/2017** TCGAbiolinksGUI: A Graphical User Interface to analyze cancer genomics and epigenomics data. EMBL, Heidelberg, Germany

**Omics Seminar - Oral presentation - 06/06/2017** Enhancer Linking by Methylation/Expression Relationship: a case study using Breast Cancer. Cedars-Sinai Medical Center, Los Angeles, California.

**Bioc2017 - Oral presentation - 07/28/2017** Workshop: Integrative analysis workshop with TCGAbiolinks and ELMER. Dana Farber Cancer Institute, Boston, MA. Link to workshop: <https://bioinformaticsfmrp.github.io/Bioc2017.TCGAbiolinks.ELMER/index.html>

**From Single to Multiomics - Poster presentation - 11/13/2017** Enhancer Linking by Methylation/Expression Relationships with the R package (ELMER) version 2. EMBL, Heidelberg, Germany

## BIBLIOGRAPHY

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## APPENDIX

A

## ETHICS COMMITTEE



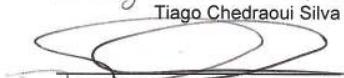
Faculdade de Medicina de Ribeirão Preto  
Universidade de São Paulo

Ilma. Senhora Profa. Dra. Marcia Guimarães Villanova,  
(Coordenadora do Comitê de Ética em Pesquisa do HCFMRP-USP)

Venho, por meio desta, solicitar dispensa de apreciação ética do projeto de pesquisa de Doutorado intitulado "Bioinformatic tool to integrate and understand aberrant epigenomic and genomic changes associated with cancer:Methods, development and analysis" ao Comitê de Ética em Pesquisa do Hospital das Clínicas da Faculdade de Medicina de Ribeirão Preto, Universidade de São Paulo (HCFMRP-USP). A solicitação de dispensa justifica-se uma vez que esta pesquisa consiste no desenvolvimento de uma ferramenta de bioinformática para a análise dados genômicos e epigenômicos e utiliza somente dados genômicos de amostras humanas que estão publicamente disponíveis à comunidade científica no banco de dados internacional *The Cancer Genome Atlas* ou TCGA (disponível em <https://tcga-data.nci.nih.gov/tcga/>), com livre acesso a todos os usuários para baixar e analisar os dados sem necessidade de cadastro ou identificação prévios. Este projeto de Doutorado está sendo desenvolvido no Departamento de Genética da FMRP-USP pelo aluno de Pós-Graduação Tiago Chedraoui Silva e sob orientação do Professor da Faculdade de Medicina de Ribeirão Preto Dr. Houtan Noushmehr. As amostras estão de acordo com a ética e a legislação norte-americana vigente, tendo em vista que o consórcio TCGA, responsável pelo banco de dados, mantém em sigilo todas as informações que possam levar à identificação dos pacientes.

Ribeirão Preto, 11 de maio de 2016.



  
Tiago Chedraoui Silva  


Prof. Dr. Houtan Noushmehr

*Não há necessidade de  
submissões ao Comitê de Ética  
em Pesquisa 23/05/16*

*Marcia Villanova*  
DRA MARCIA GUIMARÃES VILLANOVA  
Coordenadora do Comitê de Ética em Pesquisa  
do HCFMRP-USP e da FMRP-USP