### The CGDS-R library

#### Anders Jacobsen

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#### Contents

I	Intr	oduction	1
2	The	CGDS R interface	1
	2.1	CGDS(): Create a CGDS connection object	1
	2.2	getCancerTypes(): Retrieve a set of available cancer types	2
	2.3	getGeneticProfiles(): Retrieve genetic data profiles for a spe-	
		cific cancer type	3
	2.4	getCaseLists(): Retrieve case lists for a specific cancer type .	3
	2.5	getProfileData(): Retrieve genomic profile data for genes and	
		genetic profiles	4
	2.6	${\tt getClinicalData(): Retrieve\ clinical\ data\ for\ a\ list\ of\ cases} \ \ .$	4
3	Exa	mples	5
	3.1	Example 1: Association of NF1 copy number alteration and mRNA	
		expression in glioblastoma	5
	3.2	Example 2: MDM2 and MDM4 mRNA expression levels in glioblas-	
		toma	7
	3.3	Example 3: Comparing expression of PTEN in primary and metastati	c
		prostate cancer tumors	9

### 1 Introduction

This package provides a basic set of R functions for querying the Cancer Genomic Data Server (CGDS) hosted by the Computational Biology Center (cBio) at the Memorial Sloan-Kettering Cancer Center (MSKCC). This service is a part of the cBio Cancer Genomics Portal, http://www.cbioportal.org/cgx/.

In summary, the library can issue the following types of queries:

- getCancerTypes(): What cancer types are hosted on the server? For example, TCGA glioblastoma or TCGA ovarian cancer.
- getGeneticProfiles(): What genetic profile types are available for cancer type X? For example, mRNA expression or copy number alterations.
- getCaseLists(): what case sets are available for cancer type X? For example, all samples or only samples corresponding to a given cancer subtype.

- getProfileData(): Retrieve slices of genomic data. For example, a client can retrieve all mutation data for PTEN and EGFR in TCGA glioblastoma
- getClinicalData(): Retrieve clinical data (e.g. patient survival time and age) for a given cancer type and list of cases.

Each of these functions will be briefly described in the following sections. The last part of this document includes some concrete examples of how to access and plot the data.

The purpose of this document is to give the reader a quick overview of the cgdsr package. Please refer to the corresponding R manual pages for a more detailed explanation of arguments and output for each function.

### 2 The CGDS R interface

#### 2.1 CGDS(): Create a CGDS connection object

Initially, we will establish a connection to the public CGDS server hosted by Memorial Sloan-Kettering Cancer Center. The function for creating a CGDS connection object requires the URL of the CGDS server service, in this case http://cbio.mskcc.org/cgds-public/, as an argument.

```
> library(cgdsr)
> mycgds = CGDS("http://cbio.mskcc.org/cgds-public/")
```

The variable mycgds is now a CGDS connection object pointing at the URL for the public CGDS server. This connection object must be included as an argument to all subsequent interface calls. Optionally, we can now perform a set of simple tests of the data returned from the CGDS connection object using the test function:

#### > test(mycgds)

```
getCancerTypes...
getCaseLists (1/2) ...
getCaseLists (2/2) ...
getGeneticProfiles (1/2) ...
getGeneticProfiles (2/2) ...
getClinicalData (1/4) ...
getClinicalData (2/4) ...
getClinicalData (3/4) ...
getClinicalData (4/4) ...
getProfileData (1/7) ...
getProfileData (2/7) ...
getProfileData (3/7) ...
getProfileData (4/7) ...
                          OK
getProfileData (5/7) ...
                          OK
getProfileData (6/7) ...
getProfileData (7/7) ...
```

### 2.2 getCancerTypes(): Retrieve a set of available cancer types

Having created a CGDS connection object, we can now retrieve a data frame with available cancer types using the getCancerTypes function:

> getCancerTypes(mycgds)[, c(1, 2)]

	cancer_type_id				name
1	pca	Pı	costate	Cancer	(MSKCC)
2	gbm		Gliob	lastoma	(TCGA)
3	ova	Serous	Ovarian	Cancer	(TCGA)
4	Sarc		Sarcoma	(MSKCC	/Broad)

Here we are only showing the first two columns, the cancer type ID and short name, of the result data frame. There is also a third column, a longer description of the cancer type. The cancer type ID must be used in subsequent interface calls to retrieve case lists and genetic data profiles (see below).

### 2.3 getGeneticProfiles(): Retrieve genetic data profiles for a specific cancer type

This function queries the CGDS API and returns the available genetic profiles, e.g. mutation or copy number profiles, stored about a specific cancer type. Below we list the current genetic profiles for the TCGA glioblastoma cancer type:

> getGeneticProfiles(mycgds, "gbm")[, c(1:2)]

```
genetic_profile_id
                                                 genetic_profile_name
1
                                                            Mutations
       gbm_mutations
2
  gbm_cna_consensus Putative copy-number alterations (GBM Pathways)
3
         gbm_cna_rae
                             Putative copy-number alterations (RAE)
4
            gbm_mrna
                                                      mRNA Expression
5
    gbm_mrna_zscores
                                             mRNA Expression z-Scores
```

Here we are only listing the first two columns, genetic profile ID and short name, of the resulting data frame. Please refer to the R manual pages for a more extended specification of the arguments and output.

## 2.4 getCaseLists(): Retrieve case lists for a specific cancer type

This function queries the CGDS API and returns available case lists for a specific cancer type. For example, within a particular study, only some cases may have sequence data, and another subset of cases may have been sequenced and treated with a specific therapeutic protocol. Multiple case lists may be associated with each cancer type, and this method enables you to retrieve meta-data regarding all of these case lists. Below we list the current case lists for the TCGA glioblastoma cancer type:

```
> getCaseLists(mycgds, "gbm")[, c(1:2)]
```

```
case_list_id
                                                   case_list_name
1
          gbm_3way_complete All Complete Tumors (seq, mRNA, CNA)
2
                    gbm_all
                                                       All Tumors
3
         gbm_expr_classical
                                     Expression Cluster Classical
4
                                   Expression Cluster Mesenchymal
       gbm_expr_mesenchymal
5
            gbm_expr_neural
                                        Expression Cluster Neural
6
         gbm_expr_proneural
                                     Expression Cluster Proneural
7
              gbm_seq_paper
                                 Sequenced Tumors, GBM Manuscript
                                      Sequenced, No Hypermutators
8
      gbm_sequenced_nohyper
9
                                           Sequenced, Not Treated
  gbm_sequenced_nottreated
10
      gbm_sequenced_treated
                                               Sequenced, Treated
```

Here we are only listing the first two columns, case list ID and short name, of the resulting data frame. Please refer to the R manual pages for a more extended specification of the arguments and output.

### 2.5 getProfileData(): Retrieve genomic profile data for genes and genetic profiles

The function queries the CGDS API and returns data based on gene(s), genetic profile(s), and a case list. The function only allows specifying a list of genes and a single genetic profile, or oppositely a single gene and a list of genetic profiles. Importantly, the format of the output data frame depends on if a single or a list of genes was specified in the arguments. Below we are retrieving mRNA expression and copy number alteration genetic profiles for the NF1 gene in all samples of the TCGA glioblastoma cancer type:

```
> getProfileData(mycgds, "NF1", c("gbm_cna_rae", "gbm_mrna"), "gbm_all")[c(1:5),
+ ]
```

```
gbm_cna_rae gbm_mrna
TCGA.02.0001 0 -0.296691953
TCGA.02.0003 0 -0.001066810
TCGA.02.0004 NaN -0.236265119
TCGA.02.0006 0 -0.169150677
TCGA.02.0007 0 -0.009593496
```

We are here only showing the first five rows of the data frame. In the next example, we are retrieving mRNA expression data for the MDM2 and MDM4 genes:

```
> getProfileData(mycgds, c("MDM2", "MDM4"), "gbm_mrna", "gbm_all")[c(1:5),
+ ]
```

```
MDM2 MDM4
TCGA.02.0001 -0.4232196 -0.48967339
TCGA.02.0003 -0.3428422 -0.10227876
TCGA.02.0004 -0.7600202 -0.47430488
TCGA.02.0006 2.3102925 0.02569512
TCGA.02.0007 -0.5988326 2.78569512
```

We are again only showing the first five rows of the data frame.

### 2.6 getClinicalData(): Retrieve clinical data for a list of cases

The function queries the CGDS API and returns available clinical data (e.g. patient survival time and age) for a given case list. Results are returned in a data frame with a row for each case and a column for each clinical attribute. The available clinical attributes are:

- overall\_survival\_months: Overall survival, in months.
- overall\_survival\_status: Overall survival status, usually indicated as "LIVING" or "DECEASED".
- disease\_free\_survival\_months: Disease free survival, in months.
- disease\_free\_survival\_status: Disease free survival status, usually indicated as "DiseaseFree" or "Recurred/Progressed".
- age\_at\_diagnosis: Age at diagnosis.

Below we retrieve clinical data for the TCGA ovarian cancer dataset (only first five cases/rows are shown):

> getClinicalData(mycgds, "ova\_all")[c(1:5), ]

	overall_survival_months	overa	all_survival_status
TCGA.04.1331	43.80		DECEASED
TCGA.04.1332	40.89		DECEASED
TCGA.04.1336	49.02		LIVING
TCGA.04.1337	2.03		DECEASED
TCGA.04.1338	46.49		LIVING
	disease_free_survival_mo	onths	${\tt disease\_free\_survival\_status}$
TCGA.04.1331	1	15.05	Recurred/Progressed
TCGA.04.1332	1	12.95	Recurred/Progressed
TCGA.04.1336	4	49.02	DiseaseFree
TCGA.04.1337		NA	Recurred/Progressed
TCGA.04.1338	1	12.46	Recurred/Progressed
	age_at_diagnosis		
TCGA.04.1331	79.04		
TCGA.04.1332	70.64		
TCGA.04.1336	55.53		
TCGA.04.1337	78.42		
TCGA.04.1338	78.87		

### 3 Examples

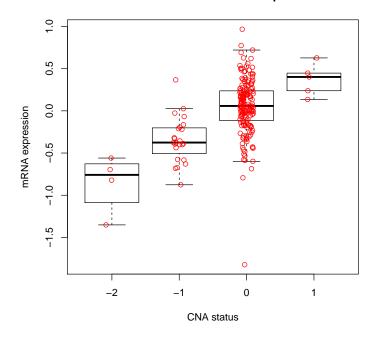
### 3.1 Example 1: Association of NF1 copy number alteration and mRNA expression in glioblastoma

As a simple example, we will generate a plot of the association between copy number alteration (CNA) status and mRNA expression change for the NF1 tumor suprpressor gene in glioblastoma. This plot is very similar to Figure 2b

in the TCGA research network paper on glioblastoma (McLendon et al. 2008). The mRNA expression of NF1 has been median adjusted on the gene level (by globally subtracting the median expression level of NF1 across all samples).

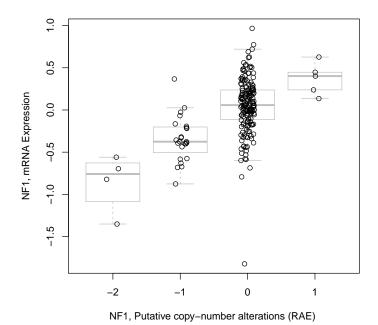
```
> df = getProfileData(mycgds, "NF1", c("gbm_cna_rae", "gbm_mrna"),
      "gbm_all")
> head(df)
             gbm_cna_rae
                              gbm_mrna
TCGA.02.0001
                       0 -0.296691953
TCGA.02.0003
                       0 -0.001066810
TCGA.02.0004
                     NaN -0.236265119
TCGA.02.0006
                       0 -0.169150677
TCGA.02.0007
                       0 -0.009593496
                       0 0.537073170
TCGA.02.0009
> boxplot(df[, 2] \tilde{} df[, 1], main = "NF1 : CNA status vs mRNA expression",
      xlab = "CNA status", ylab = "mRNA expression", outpch = NA)
> stripchart(df[, 2] ~ df[, 1], vertical = T, add = T, method = "jitter",
      pch = 1, col = "red")
```

#### NF1: CNA status vs mRNA expression



Alternatively, the generic cgdsr plot() function can be used to generate a similar plot:

```
> plot(mycgds, "gbm", "NF1", c("gbm_cna_rae", "gbm_mrna"), "gbm_all",
+ skin = "disc_cont")
```



# 3.2 Example 2: MDM2 and MDM4 mRNA expression levels in glioblastoma

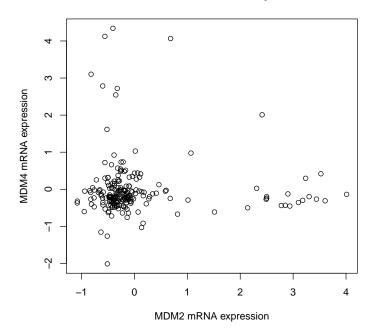
In this example, we evaluate the relationship of MDM2 and MDM4 expression levels in glioblastoma. mRNA expression levels of MDM2 and MDM4 have been median adjusted on the gene level (by globally subtracting the median expression level of the individual gene across all samples).

```
> df = getProfileData(mycgds, c("MDM2", "MDM4"), "gbm_mrna", "gbm_all")
> head(df)
```

```
MDM2 MDM4
TCGA.02.0001 -0.4232196 -0.48967339
TCGA.02.0003 -0.3428422 -0.10227876
TCGA.02.0004 -0.7600202 -0.47430488
TCGA.02.0006 2.3102925 0.02569512
TCGA.02.0007 -0.5988326 2.78569512
TCGA.02.0009 -0.4095651 0.02249049
```

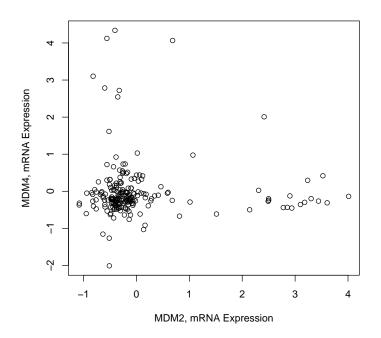
> plot(df, main = "MDM2 and MDM4 mRNA expression", xlab = "MDM2 mRNA expression",
+ ylab = "MDM4 mRNA expression")

### MDM2 and MDM4 mRNA expression



Alternatively, the generic  $\operatorname{\mathsf{cgdsr}}$   $\operatorname{\mathsf{plot}}()$  function can be used to generate a similar plot:

> plot(mycgds, "gbm", c("MDM2", "MDM4"), "gbm\_mrna", "gbm\_all")
[1] TRUE



## 3.3 Example 3: Comparing expression of PTEN in primary and metastatic prostate cancer tumors

In this example we plot the mRNA expression levels of PTEN in primary and metastatic prostate cancer tumors.

```
> df.pri = getProfileData(mycgds, "PTEN", "pca_mrna", "pca_primary")
> head(df.pri)
            PTEN
PCA0001 9.467183
PCA0002 9.041528
PCA0003 8.511305
PCA0004
             NaN
PCA0005 9.413217
PCA0006
             NaN
> df.met = getProfileData(mycgds, "PTEN", "pca_mrna", "pca_mets")
> head(df.met)
            PTEN
PCA0182 7.486938
PCA0183
             NaN
PCA0184 7.578755
PCA0185
             NaN
PCA0186
             NaN
PCA0187 8.756132
```

### PTEN expression in primary and metastatic tumors

