metadata_script_for_lusha

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6/26/2023

packages

```
knitr::opts_chunk$set(echo = T, results = "show", fig.width=8, fig.height=6, tidy.opts=list(width.cutof
options(scipen = 0, digits = 3)  # controls base R output
# check if you have ISLR package, if not, install it
if(!require('pacman')) {install.packages('pacman')}
## Loading required package: pacman
```

Read in the data

```
setwd("~/OneDrive - Children's Hospital of Philadelphia/COG-pediatric-AML/metadata_v2")
wgs = read.csv("WGS.csv", header = T)  # point to path of wgs Cavatica CSV
rna = read.csv("RNA.csv", header = T)  # point to path of rna Cavatica CSV
new_pat.met = read.csv("NCTO1371981-D4-Dataset.csv")  # point to path of new metadata file
```

Change the name of rna case id -> usi

```
colnames(rna)[1] = "usi"
# head(rna)

colnames(wgs)[1] = "usi" # same for wgs
# head(wgs)
```

subset cavatica data

RNA

N individuals with WGS (normal) x RNA-seq tumor x Pat.Met -> 390

pacman::p_load(dplyr, readr, tidyverse, ggplot2, formatR)

```
rna_norm = filter(rna, sample_type == "Normal")
rna_tum = filter(rna, sample_type == "Tumor")

# x2 = filter(rna_tum, rna_tum$usi %in% wgs$usi & rna_tum$usi %in% regno$usi & !rna_tum %in% # rna_norm)

rna.wgs = merge(rna, wgs, by = "usi")
rna_tum.wgs = merge(rna_tum, wgs, by = "usi")
```

WGS

```
wgs_norm = filter(wgs, sample_type == "Normal")
wgs_tum = filter(wgs, sample_type == "Tumor")
```

Filter out Study Ineligible Patients

```
new_pat.met_eligible = new_pat.met %>%
filter(!grepl("02", consort_clsf))
```

Number of eligible patients with RNAseq data

```
length(intersect(rna$usi, new_pat.met_eligible$usi)) ## --> 1613
```

[1] 1040

creating table

row 1

WGS normal, !WGS tumor, !RNAseq normal, RNA-seq tumor, Patient Metadata

```
row1 = filter(new_pat.met_eligible, new_pat.met_eligible$usi %in% rna_tum$usi & new_pat.met_eligible$us
   wgs$usi & !new_pat.met_eligible$usi %in% rna_norm$usi)
nrow(row1)
```

[1] 390

row 2

!wgs_normal, !wgs_tumor, !rna_normal, rna_tumor, patient metadata

row2 = filter(new_pat.met_eligible, new_pat.met_eligible\$usi %in% rna_tum\$usi & !new_pat.met_eligible\$u
 wgs_norm\$usi & !new_pat.met_eligible\$usi %in% wgs_tum\$usi & !new_pat.met_eligible\$usi %in% rna_norm
 nrow(row2)

[1] 650

row 3

wgs_normal, !wgs_tumor, !rna_normal, rna_tumor, !patient metadata

row3 = filter(wgs, wgs\$usi %in% rna_tum\$usi & wgs\$usi %in% wgs_norm\$usi & !wgs\$usi %in% wgs_tum\$usi &
 !wgs\$usi %in% rna_norm\$usi & !wgs\$usi %in% new_pat.met_eligible\$usi)
nrow(row3)

[1] 0

row 4

wgs norm, !wgs tum, !rna norm, !rna tum, pat met

row4 = filter(wgs, wgs\$usi %in% wgs_norm\$usi & !wgs\$usi %in% wgs_tum\$usi & !wgs\$usi %in% rna_norm\$usi &
 !wgs\$usi %in% rna_tum\$usi & wgs\$usi %in% new_pat.met_eligible)
nrow(row4)

[1] 0

row 5

wgs_norm, !wgs_tum, !rna_norm, !rna_tum, !pat met

row5 = filter(wgs, wgs\$usi %in% wgs_norm\$usi & !wgs\$usi %in% wgs_tum\$usi & !wgs\$usi %in% rna_norm\$usi &
 !wgs\$usi %in% rna_tum\$usi & !wgs\$usi %in% new_pat.met_eligible)
nrow(row5)

[1] 18

row 6

!wgs_norm, !wgs_tum, rna_norm, !rna_tum, !pat met

row6 = filter(rna, !rna\$usi %in% wgs_norm\$usi & !rna\$usi %in% wgs_tum\$usi & rna\$usi %in% rna_norm\$usi &
 !rna\$usi %in% rna_tum\$usi & !rna\$usi %in% new_pat.met_eligible)
nrow(row6)

[1] 62

row 7

row7 = filter(rna, !rna\$usi %in% wgs_norm\$usi & !rna\$usi %in% wgs_tum\$usi & !rna\$usi %in% rna_norm\$usi e
rna\$usi %in% rna_tum\$usi & !rna\$usi %in% new_pat.met_eligible)
nrow(row7)

[1] 661