

Feature Classification Take 2

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Objective

Revise feature classifications to define situations that result in different performance expectation.

```
## Extracting a tidy dataframe with count values from MRexpiment objects
get_count_df <- function(mrobject){
  mrobject <- cumNorm(mrobject, p = 0.75)
  mrobject %>%
    # not sure whether or not to normalize counts prior to analysis
    MRcounts(norm = TRUE, log = FALSE, sl = 1000) %>%
    as.data.frame() %>%
    rownames_to_column(var = "feature_id") %>%
    gather("id", "count", -feature_id)
}

count_df <- mrexpr %>% map_df(get_count_df, .id = "pipe")

count_replicate_df <- pData(mrexpr$dada2) %>% right_join(count_df) %>%
  filter(biosample_id != "NTC") %>%
  mutate(detect = if_else(count > 0, 1, 0)) %>%
  group_by(pipe, biosample_id, titration, t_fctr, feature_id) %>%
  summarise(total_detect = sum(detect),
            n_replicates = n(),
            avg_non0_count = sum(count)/total_detect) %>%
  mutate(detect_prop = total_detect/n_replicates) %>%
  select(-total_detect)

rep_info <- count_replicate_df %>% ungroup() %>%
  mutate(t_fctr = paste0("T", t_fctr)) %>%
  select(pipe, biosample_id, feature_id, t_fctr, detect_prop)
```

Feature Categories

- Null - features not present in more than one PCR replicate for any sample of a biological replicate, and pipeline.
- Full - features present in at least one PCR replicate for all samples of a biological replicate, and pipeline.
- Mix - features only present in at least two PCR replicates for a mixed sample but not observed in any of the unmixed sample PCR replicates.
- Pre - present in three or more PCR replicates for unmixed pre-treatment samples, not observed in any PCR replicates of the unmixed post treatment samples, and present in at least 20 total PCR replicates.
- Post - present in three or more PCR replicates for the unmixed post-treatment samples, not observed in any PCR replicates of the unmixed pre-treatment samples, and present in at least 8 total PCR replicates.

```
prop_summary <- rep_info %>%
  group_by(pipe, biosample_id, feature_id) %>%
  summarise(prop_max = max(detect_prop),
```

```

        prop_min = min(detect_prop),
        prop_sum = sum(detect_prop))

unmix_prop <- rep_info %>%
  filter(t_fctr %in% c("T0","T20")) %>%
  spread(t_fctr, detect_prop)

feature_cat <- left_join(prop_summary, unmix_prop) %>%
  mutate(cat_null = if_else(prop_max < 0.5, 1, 0),
         cat_full = if_else(prop_min > 0, 1, 0),
         cat_mix  = if_else(prop_max >= 0.5 & T0 == 0 & T20 == 0, 1, 0),
         cat_pre  = if_else(T20 >= 0.75 & T0 == 0 & prop_sum > 5, 1, 0),
         cat_post = if_else(T0 >= 0.75 & T20 == 0 & prop_sum > 2, 1, 0),
         cat_none = if_else(cat_null + cat_full + cat_mix + cat_pre + cat_post == 0, 1, 0))

```

Category Sanity Check

```

cat_check <- feature_cat %>%
  select(pipe, biosample_id, feature_id,
         cat_null, cat_full, cat_mix, cat_pre, cat_post, cat_none) %>%
  gather(cat, value, -pipe, -biosample_id, -feature_id) %>%
  group_by(pipe, biosample_id, feature_id) %>%
  summarise(n_cat = sum(value))
cat_check %>% filter(n_cat != 1)

## Source: local data frame [0 x 4]
## Groups: pipe, biosample_id [0]
##
## # ... with 4 variables: pipe <chr>, biosample_id <chr>, feature_id <chr>,
## #   n_cat <dbl>

# cat_check <- feature_categories %>%
#   select(pipe, biosample_id, feature_id,
#          cat_null, cat_full, cat_mix, cat_pre, cat_post, cat_none) %>%
#   gather(cat, value, -pipe, -biosample_id, -feature_id) %>%
#   group_by(pipe, biosample_id, feature_id) %>%
#   mutate(n_cat = sum(value)) %>%
#   filter(n_cat != 1, value != 0)
# cat_check %>% arrange(feature_id)

```

```

feature_categories <- feature_cat %>% select(pipe, biosample_id, feature_id,
      cat_null, cat_full, cat_mix, cat_pre, cat_post, cat_none) %>%
  gather(cat, value, -pipe, -biosample_id, -feature_id) %>%
  filter(value == 1) %>% select(-value)

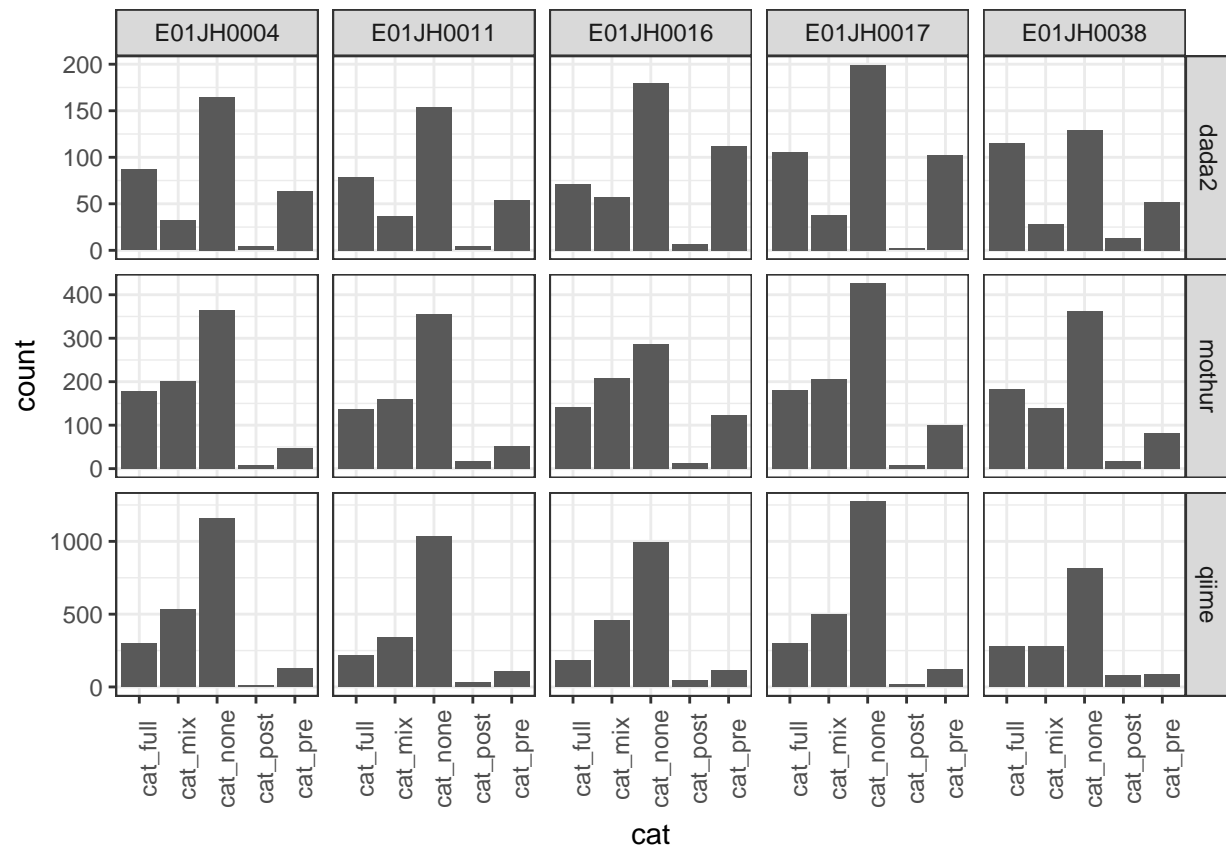
```

Save Data Frame as RDS

```
feature_categories %>% saveRDS("../data/feature_categories_df.rds")
```

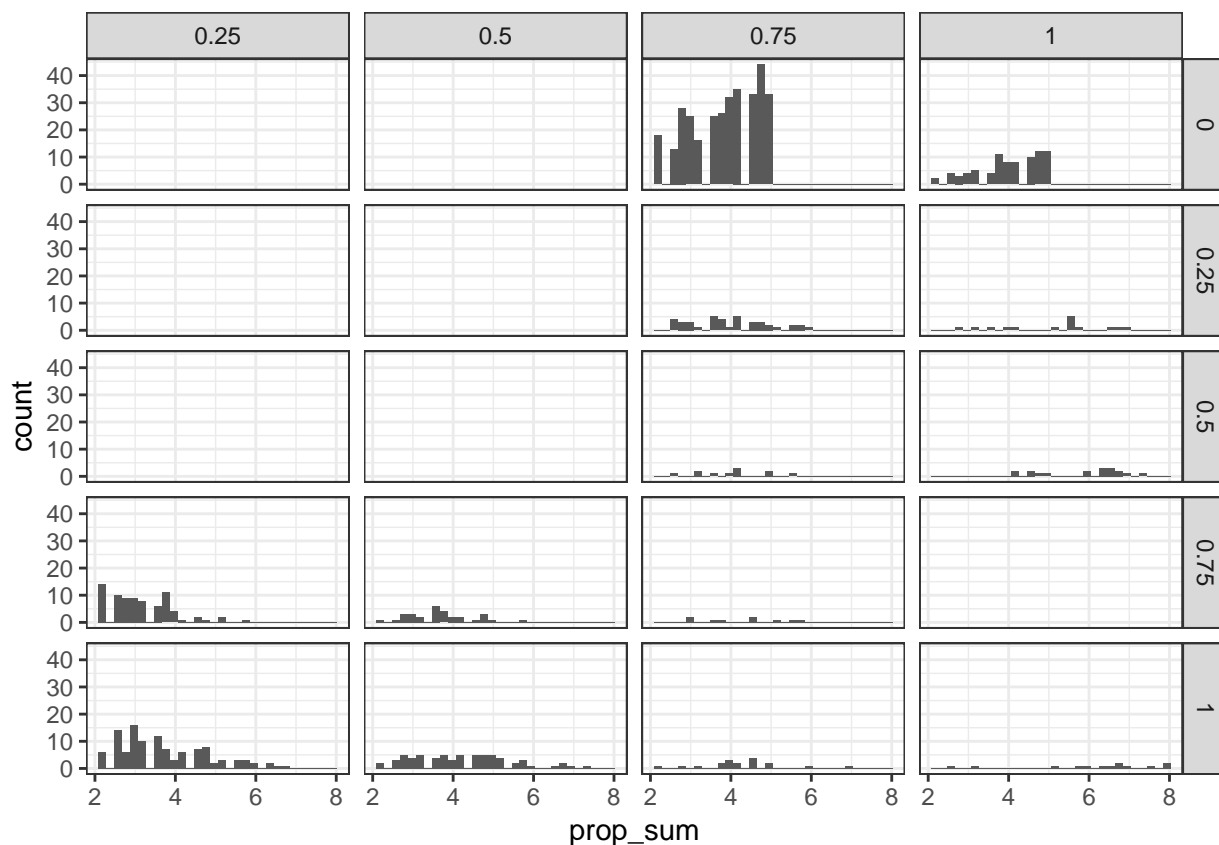
Summary Figures

```
feature_categories %>% filter(cat != "cat_null") %>%
  ggplot() + geom_bar(aes(x = cat)) +
  facet_grid(pipe ~ biosample_id, scales = "free_y") +
  theme_bw() + theme(axis.text.x = element_text(angle = 90))
```



While there are a large number of unclassified features, few are potentially informative. Ones that stand out are features detected in 3 of 4 T0 (pre-treatment features), and observed between 8 and 20 PCR replicates ($2 < \text{prop_sum} < 5$).

```
feature_cat %>% filter(cat_none == 1, prop_sum > 2, T0 > 0.5 | T20 > 0.5) %>%
  ggplot() + geom_histogram(aes(x = prop_sum)) + facet_grid(T0 ~ T20) + theme_bw()
```



Session information

```
s_info <- devtools::session_info()
print(s_info$platform)
```

```
## setting value
## version R version 3.3.3 (2017-03-06)
## system x86_64, darwin15.6.0
## ui unknown
## language (EN)
## collate en_US.UTF-8
## tz America/New_York
## date 2017-04-03
```

```
s_info$packages %>% filter(`*` == "*") %>% select(-`*`) %>%
  knitr::kable()
```

| package | version | date | source |
|--------------|---------|------------|-------------------------------|
| bbmle | 1.0.18 | 2016-02-11 | CRAN (R 3.3.2) |
| Biobase | 2.34.0 | 2016-11-07 | Bioconductor |
| BiocGenerics | 0.20.0 | 2016-11-07 | Bioconductor |
| BiocParallel | 1.8.1 | 2016-11-07 | Bioconductor |
| Biostrings | 2.42.1 | 2016-12-19 | Bioconductor |
| DESeq | 1.26.0 | 2016-11-28 | Bioconductor |
| DESeq2 | 1.15.28 | 2017-02-02 | bioc (readonly/DESeq2@125913) |

| package | version | date | source |
|----------------------|---------|------------|----------------|
| dplyr | 0.5.0 | 2016-06-24 | CRAN (R 3.3.2) |
| edgeR | 3.16.5 | 2017-02-02 | Bioconductor |
| forcats | 0.2.0 | 2017-01-23 | CRAN (R 3.3.2) |
| foreach | 1.4.3 | 2015-10-13 | CRAN (R 3.3.1) |
| GenomeInfoDb | 1.10.3 | 2017-03-28 | Bioconductor |
| GenomicAlignments | 1.10.1 | 2017-03-28 | Bioconductor |
| GenomicRanges | 1.26.4 | 2017-03-28 | Bioconductor |
| ggplot2 | 2.2.1 | 2016-12-30 | CRAN (R 3.3.2) |
| glmnet | 2.0-5 | 2016-03-17 | CRAN (R 3.3.1) |
| IRanges | 2.8.2 | 2017-03-28 | Bioconductor |
| knitr | 1.15.1 | 2016-11-22 | CRAN (R 3.3.2) |
| lattice | 0.20-34 | 2016-09-06 | CRAN (R 3.3.3) |
| limma | 3.30.13 | 2017-03-28 | Bioconductor |
| locfit | 1.5-9.1 | 2013-04-20 | CRAN (R 3.3.1) |
| Matrix | 1.2-8 | 2017-01-20 | CRAN (R 3.3.3) |
| metagenomeSeq | 1.16.0 | 2016-11-07 | Bioconductor |
| modelr | 0.1.0 | 2016-08-31 | cran (@0.1.0) |
| permute | 0.9-4 | 2016-09-09 | CRAN (R 3.3.1) |
| phyloseq | 1.19.1 | 2017-01-04 | Bioconductor |
| ProjectTemplate | 0.7 | 2016-08-11 | CRAN (R 3.3.1) |
| purrr | 0.2.2 | 2016-06-18 | CRAN (R 3.3.1) |
| RColorBrewer | 1.1-2 | 2014-12-07 | CRAN (R 3.3.1) |
| readr | 1.1.0 | 2017-03-22 | CRAN (R 3.3.2) |
| readxl | 0.1.1 | 2016-03-28 | cran (@0.1.1) |
| Rqc | 1.8.0 | 2016-11-07 | Bioconductor |
| Rsamtools | 1.26.1 | 2016-11-07 | Bioconductor |
| S4Vectors | 0.12.2 | 2017-03-28 | Bioconductor |
| sads | 0.3.1 | 2016-05-13 | CRAN (R 3.3.2) |
| savR | 1.12.0 | 2016-11-07 | Bioconductor |
| ShortRead | 1.32.1 | 2017-03-28 | Bioconductor |
| stringr | 1.2.0 | 2017-02-18 | CRAN (R 3.3.2) |
| SummarizedExperiment | 1.4.0 | 2016-11-07 | Bioconductor |
| tibble | 1.2 | 2016-08-26 | CRAN (R 3.3.1) |
| tidyr | 0.6.1 | 2017-01-10 | CRAN (R 3.3.2) |
| tidyverse | 1.1.1 | 2017-01-27 | CRAN (R 3.3.2) |
| vegan | 2.4-2 | 2017-01-17 | CRAN (R 3.3.2) |
| XVector | 0.14.1 | 2017-03-28 | Bioconductor |