

# Protein\_Domain\_RNA\_Localization

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```
# BiocManager::install("biomaRt")
# install.packages("tidyverse")
```

Load Libraries

```
library(biomaRt)
```

```
## Warning: package 'biomaRt' was built under R version 4.1.1
```

```
library(tidyverse)
```

```
## -- Attaching packages ----- tidyverse 1.3.1 --
```

```
## v ggplot2 3.3.5      v purrr   0.3.4
```

```
## v tibble  3.1.6      v dplyr  1.0.8
```

```
## v tidyr   1.2.0      v stringr 1.4.0
```

```
## v readr   2.1.2      v forcats 0.5.1
```

```
## Warning: package 'tidyr' was built under R version 4.1.2
```

```
## Warning: package 'readr' was built under R version 4.1.2
```

```
## Warning: package 'dplyr' was built under R version 4.1.2
```

```
## -- Conflicts ----- tidyverse_conflicts() --
```

```
## x dplyr::filter() masks stats::filter()
```

```
## x dplyr::lag()     masks stats::lag()
```

```
## x dplyr::select() masks biomaRt::select()
```

```
library(openxlsx)
```

```
library(ComplexHeatmap)
```

```
## Loading required package: grid
```

```
## =====
```

```
## ComplexHeatmap version 2.8.0
```

```
## Bioconductor page: http://bioconductor.org/packages/ComplexHeatmap/
```

```
## Github page: https://github.com/jokergoo/ComplexHeatmap
```

```
## Documentation: http://jokergoo.github.io/ComplexHeatmap-reference
```

```
##
```

```
## If you use it in published research, please cite:
```

```
## Gu, Z. Complex heatmaps reveal patterns and correlations in multidimensional  
## genomic data. Bioinformatics 2016.
```

```
##
```

```
## The new InteractiveComplexHeatmap package can directly export static
```

```
## complex heatmaps into an interactive Shiny app with zero effort. Have a try!
```

```
##
```

```
## This message can be suppressed by:
```

```
## suppressPackageStartupMessages(library(ComplexHeatmap))
## =====
```

Download information from WormBase ParaSite BioMart.

Guide is located here: <https://parasite.wormbase.org/info/Tools/biomart.html>

```
paramart <- useMart("parasite_mart", dataset = "wbps_gene", host = "https://parasite.wormbase.org", port = 80)
```

```
protomain_df <- getBM(
  mart = paramart,
  filter = c("species_id_1010","biotype"),
  values = list(species_id_1010 = "caelegprjna13758", biotype = "protein_coding"),
  attributes = c("production_name_1010", "wormbase_gseq","wbps_gene_id", "wikigene_name", "interpro_id")
)
head(protomain_df)
```

```
##           production_name_1010 wormbase_gseq  wbps_gene_id wikigene_name
## 1 caenorhabditis_elegans_prjna13758    Y110A7A.10 WBGene000000001      aap-1
## 2 caenorhabditis_elegans_prjna13758    Y110A7A.10 WBGene000000001      aap-1
## 3 caenorhabditis_elegans_prjna13758    Y110A7A.10 WBGene000000001      aap-1
## 4 caenorhabditis_elegans_prjna13758    Y110A7A.10 WBGene000000001      aap-1
## 5 caenorhabditis_elegans_prjna13758    Y110A7A.10 WBGene000000001      aap-1
## 6 caenorhabditis_elegans_prjna13758    Y110A7A.10 WBGene000000001      aap-1
##   interpro_id interpro_short_description  interpro_description interpro_start
## 1   IPR036860          SH2_dom_sf SH2 domain superfamily           8
## 2   IPR036860          SH2_dom_sf SH2 domain superfamily          333
## 3   IPR036860          SH2_dom_sf SH2 domain superfamily           14
## 4   IPR036860          SH2_dom_sf SH2 domain superfamily          358
## 5   IPR000980              SH2          SH2 domain           360
## 6   IPR000980              SH2          SH2 domain            20
##   interpro_end
## 1           125
## 2           450
## 3           115
## 4           443
## 5           422
## 6            94
```

```
protomain_df %>% group_by(interpro_description) %>% count() %>% nrow
```

```
## [1] 8512
```

There are 8533 unique protein domain IDs

These are all the protein domains associated with “erm-1”, “frm-7”, and “imb-2”

```
# protomain_df %>% filter(wikigene_name == "erm-1")
# protomain_df %>% filter(wikigene_name == "frm-7")
# protomain_df %>% filter(interpro_short_description %in% c("PH_domain", "FERM_domain", "PH-like_dom_s

goi_domains <- protomain_df %>% filter(wikigene_name %in% c("erm-1")) %>% group_by(wikigene_name, interpro_short_description)
goi_domains
```

```
## # A tibble: 15 x 4
```

```
## # Groups:   wikigene_name, interpro_short_description, interpro_description
```

```
## #   [15]
```

```
##   wikigene_name interpro_short_description interpro_description      n
```

```

##      <chr>          <chr>          <chr>          <int>
## 1 erm-1      Band_41_domain      Band 4.1 domain      10
## 2 erm-1      ERM                  Ezrin/radixin/moesin      2
## 3 erm-1      ERM_C_dom            Ezrin/radixin/moesin, C-termi~      2
## 4 erm-1      ERM_FERM_C            ERM family, FERM domain C-lobe      2
## 5 erm-1      Ez/rad/moesin-like    Ezrin/radixin/moesin-like      16
## 6 erm-1      FERM_2                FERM superfamily, second doma~      2
## 7 erm-1      FERM_central          FERM central domain      4
## 8 erm-1      FERM_CS                FERM conserved site      4
## 9 erm-1      FERM_domain            FERM domain            2
## 10 erm-1     FERM_N                  FERM, N-terminal        2
## 11 erm-1     FERM_PH-like_C          FERM, C-terminal PH-like doma~      4
## 12 erm-1     FERM/acyl-CoA-bd_prot_sf FERM/acyl-CoA-binding protein~      2
## 13 erm-1     Moesin_tail_sf          Moesin tail domain superfamily      4
## 14 erm-1     PH-like_dom_sf          PH-like domain superfamily      2
## 15 erm-1     Ubiquitin-like_domsf     Ubiquitin-like domain superfa~      2

domain_hits <- protdomain_df %>% filter(interpro_short_description %in% goi_domains$interpro_short_desc
  !(interpro_short_description %in% c("Ubiquitin-like_domsf"))
) %>%
  select(wbps_gene_id, wikigene_name, interpro_description, interpro_short_description) %>%
  group_by(wbps_gene_id, wikigene_name, interpro_description, interpro_short_description) %>%
  count(name = "domain_count") %>%
  ungroup()
head(domain_hits)

## # A tibble: 6 x 5
##   wbps_gene_id  wikigene_name interpro_descripti~ interpro_short_~ domain_count
##   <chr>        <chr>          <chr>          <chr>          <int>
## 1 WBGene00000102 akt-1          PH-like domain sup~ PH-like_dom_sf      2
## 2 WBGene00000103 akt-2          PH-like domain sup~ PH-like_dom_sf      1
## 3 WBGene00000149 apl-1          PH-like domain sup~ PH-like_dom_sf      2
## 4 WBGene00000420 ced-6          PH-like domain sup~ PH-like_dom_sf      1
## 5 WBGene00000426 ced-12         PH-like domain sup~ PH-like_dom_sf      2
## 6 WBGene00000564 cnk-1          PH-like domain sup~ PH-like_dom_sf      1

length(unique(domain_hits$wbps_gene_id))

## [1] 144

domain_hits_totals <- protdomain_df %>% filter(interpro_short_description %in% goi_domains$interpro_sho
  !(interpro_short_description %in% c("Ubiquitin-like_domsf"))
) %>% group_by(interpro_short_description, interpro_description) %>% count(name = "domain_count") %>%
domain_hits_totals

## # A tibble: 14 x 3
##   interpro_short_description interpro_description      domain_count
##   <chr>                    <chr>                    <int>
## 1 PH-like_dom_sf          PH-like domain superfamily      282
## 2 Band_41_domain          Band 4.1 domain                101
## 3 FERM_central            FERM central domain            67
## 4 Ez/rad/moesin-like      Ezrin/radixin/moesin-like      47
## 5 FERM_domain             FERM domain                    42
## 6 FERM_PH-like_C          FERM, C-terminal PH-like domain      40
## 7 FERM_2                  FERM superfamily, second domain      37
## 8 FERM/acyl-CoA-bd_prot_sf FERM/acyl-CoA-binding protein superf~      36

```

## 9	FERM_CS	FERM conserved site	15
## 10	FERM_N	FERM, N-terminal	15
## 11	Moesin_tail_sf	Moesin tail domain superfamily	10
## 12	ERM_C_dom	Ezrin/radixin/moesin, C-terminal	6
## 13	ERM_FERM_C	ERM family, FERM domain C-lobe	3
## 14	ERM	Ezrin/radixin/moesin	2

```
present_sub <- read.xlsx(xlsxFile = "S1_Dataset_AB_P1_Transcriptome.xlsx",
  sheet = "present_subset",
  startRow = 2) %>% select(WBID)
AB_enr_sub <- read.xlsx(xlsxFile = "S1_Dataset_AB_P1_Transcriptome.xlsx",
  sheet = "AB-enriched_subset",
  startRow = 2) %>% select(WBID)
P1_enr_sub <- read.xlsx(xlsxFile = "S1_Dataset_AB_P1_Transcriptome.xlsx",
  sheet = "P1-enriched_subset",
  startRow = 2) %>% select(WBID)
symm_sub <- read.xlsx(xlsxFile = "S1_Dataset_AB_P1_Transcriptome.xlsx",
  sheet = "symm_subset",
  startRow = 2) %>% select(WBID)
c(nrow(present_sub), nrow(AB_enr_sub), nrow(P1_enr_sub), nrow(symm_sub))
```

```
## [1] 7945 80 201 3974
```

Add true/false for different AB/P1 category

```
twocell_domains <- domain_hits %>%
  mutate(present = case_when(wbps_gene_id %in% present_sub$WBID == TRUE ~ TRUE,
    wbps_gene_id %in% present_sub$WBID == FALSE ~ FALSE),
    AB_enriched = case_when(wbps_gene_id %in% AB_enr_sub$WBID == TRUE ~ TRUE,
    wbps_gene_id %in% AB_enr_sub$WBID == FALSE ~ FALSE),
    P1_enriched = case_when(wbps_gene_id %in% P1_enr_sub$WBID == TRUE ~ TRUE,
    wbps_gene_id %in% P1_enr_sub$WBID == FALSE ~ FALSE),
    symmetric = case_when(wbps_gene_id %in% symm_sub$WBID == TRUE ~ TRUE,
    wbps_gene_id %in% symm_sub$WBID == FALSE ~ FALSE),
  )
twocell_domains
```

```
## # A tibble: 273 x 9
```

##	wbps_gene_id	wikigene_name	interpro_descript~	interpro_short_~	domain_count
##	<chr>	<chr>	<chr>	<chr>	<int>
## 1	WBGene00000102	akt-1	PH-like domain su~	PH-like_dom_sf	2
## 2	WBGene00000103	akt-2	PH-like domain su~	PH-like_dom_sf	1
## 3	WBGene00000149	apl-1	PH-like domain su~	PH-like_dom_sf	2
## 4	WBGene00000420	ced-6	PH-like domain su~	PH-like_dom_sf	1
## 5	WBGene00000426	ced-12	PH-like domain su~	PH-like_dom_sf	2
## 6	WBGene00000564	cnk-1	PH-like domain su~	PH-like_dom_sf	1
## 7	WBGene00000565	cnt-1	PH-like domain su~	PH-like_dom_sf	3
## 8	WBGene00000894	dab-1	PH-like domain su~	PH-like_dom_sf	2
## 9	WBGene00001093	drp-1	PH-like domain su~	PH-like_dom_sf	2
## 10	WBGene00001116	dyc-1	PH-like domain su~	PH-like_dom_sf	2

## # ... with 263 more rows, and 4 more variables: present <lgl>,  
## # AB\_enriched <lgl>, P1\_enriched <lgl>, symmetric <lgl>

```
twocell_domain_genes <- twocell_domains %>% mutate(gene_type = case_when(
  present == TRUE & symmetric == FALSE & AB_enriched == FALSE ~ "no_sig_dif",
  present == TRUE & symmetric == TRUE ~ "symmetric",
```

```

present == TRUE & AB_enriched == TRUE ~ "AB_enriched",
present == FALSE ~ "not_detected",
)) %>% select(wbps_gene_id:domain_count, gene_type)
twocell_domain_genes

```

```

## # A tibble: 273 x 6
##   wbps_gene_id wikigene_name interpro_descript~ interpro_short_~ domain_count
##   <chr>         <chr>         <chr>         <chr>         <int>
## 1 WBGene00000102 akt-1          PH-like domain su~ PH-like_dom_sf      2
## 2 WBGene00000103 akt-2          PH-like domain su~ PH-like_dom_sf      1
## 3 WBGene00000149 apl-1          PH-like domain su~ PH-like_dom_sf      2
## 4 WBGene00000420 ced-6          PH-like domain su~ PH-like_dom_sf      1
## 5 WBGene00000426 ced-12         PH-like domain su~ PH-like_dom_sf      2
## 6 WBGene00000564 cnk-1          PH-like domain su~ PH-like_dom_sf      1
## 7 WBGene00000565 cnt-1          PH-like domain su~ PH-like_dom_sf      3
## 8 WBGene00000894 dab-1          PH-like domain su~ PH-like_dom_sf      2
## 9 WBGene00001093 drp-1          PH-like domain su~ PH-like_dom_sf      2
## 10 WBGene00001116 dyc-1          PH-like domain su~ PH-like_dom_sf      2
## # ... with 263 more rows, and 1 more variable: gene_type <chr>

```

```

# Number of unique genes in dataset
# Make sure the numbers match the plot above
table((twocell_domain_genes %>% distinct(wikigene_name, .keep_all = TRUE))$gene_type)

```

```

##
## AB_enriched no_sig_dif not_detected symmetric
##           6           57           20           61

```

Number of protein domain types in each two cell embryo gene category

```
table(twocell_domain_genes$gene_type)
```

```

##
## AB_enriched no_sig_dif not_detected symmetric
##           25           106           40           102

```

Get the names of AB enriched genes

```
unique((twocell_domain_genes %>% filter(gene_type == "AB_enriched"))$wikigene_name)
```

```
## [1] "akt-1" "erm-1" "frm-7" "sma-1" "unc-73" "F22G12.5"
```

Get the names of symmetric enriched genes

```
unique((twocell_domain_genes %>% filter(gene_type == "symmetric"))$wikigene_name)
```

```

## [1] "akt-2" "cnk-1" "cnt-1" "dyc-1" "ech-4" "exc-5"
## [7] "feh-1" "frm-10" "grp-1" "hmg-3" "hum-4" "icln-1"
## [13] "ist-1" "kin-32" "max-1" "mtm-1" "mtm-6" "mtm-9"
## [19] "nfm-1" "npp-16" "pdk-1" "plc-1" "ptp-1" "ran-5"
## [25] "soc-1" "sos-1" "stn-1" "tag-52" "unc-31" "unc-34"
## [31] "unc-104" "vav-1" "wsp-1" "F07C6.4" "vps-36" "snx-17"
## [37] "F31D4.5" "F38B7.3" "T10B10.3" "acbp-5" "dkf-2" "dkf-1"
## [43] "Y37D8A.25" "Y41E3.7" "ani-3" "obr-1" "exoc-8" "ZK632.12"
## [49] "obr-3" "obr-4" "shc-1" "ani-2" "K10B4.3" "bris-1"
## [55] "M03A8.3" "gtf-2H1" "shc-2" "dcap-1" "prhg-1" "ZC328.3"
## [61] "tbc-2"

```

Total number of genes

```
twocell_domain_genes %>% distinct(wbps_gene_id) %>% nrow()
```

```
## [1] 144
```

Output the list of protein domains

```
write.xlsx(twocell_domain_genes %>% select(wbps_gene_id:domain_count), file = "Protein_Domains_2-Cell_E
```

Output the list of genes with 2 cell data annotation

```
write.xlsx(twocell_domain_genes %>% select(wbps_gene_id, wikigene_name, AB_vs_P1 = gene_type) %>% disti
```