

# Functional annotation of CNV breakpoints in Holstein-Friesian cows

Joanna Szyda, Magda Mielczarek, Magdalena Frąszczak

Riccardo Giannico, Giulietta Minozzi, Ezequiel Nicolazzi

Katarzyna Wojdak-Maksymiec



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# CNVs – Copy Number Variants

ATTCGACCGTAGCCGTAAGGTCTAGTTAACCCCCCGTAAGGTCTAGTTAACGACCGTA

ATTC

AGTTAACGACCGTA

ATTCGACCGTAGCCGTAAGGTCTAGTTAACCCCCCGTAAGGTCTGACCGTAGCCGTAAG  
GTCCTAGTTAACCCCCCGTAAGGTCTAGTTAACGACCGTA

# Why breakpoints ?

Are CNV breakpoints formed at  
random in the bovine genome ?

# Outline

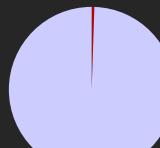
1. Material
2. CNV detection
3. Functional annotation of CNV breakpoints
  - SO terms
  - Gene functional category
  - All genes
4. Conclusions

# Whole genome sequence data

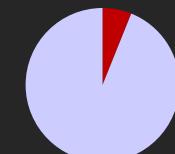
- 32 HF cows → 29 HF cows
- 16 paternal half-sibs
- WGS – Illumina HiSeq 2000
  - Paired end 100 bp
  - Raw reads 164,984,147 – 472,265,620
  - Coverage 5x – 17x



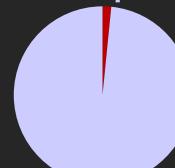
# Copy Number Variation

1. Alignment to UMD 3.1 – BWA.MEM
2. Read editing – SAMtools & Picard
3. CNV detection – CNVnator
  - Deletion      200 bp – 724,000 bp
  - Duplication      200 bp – 439,300 bp

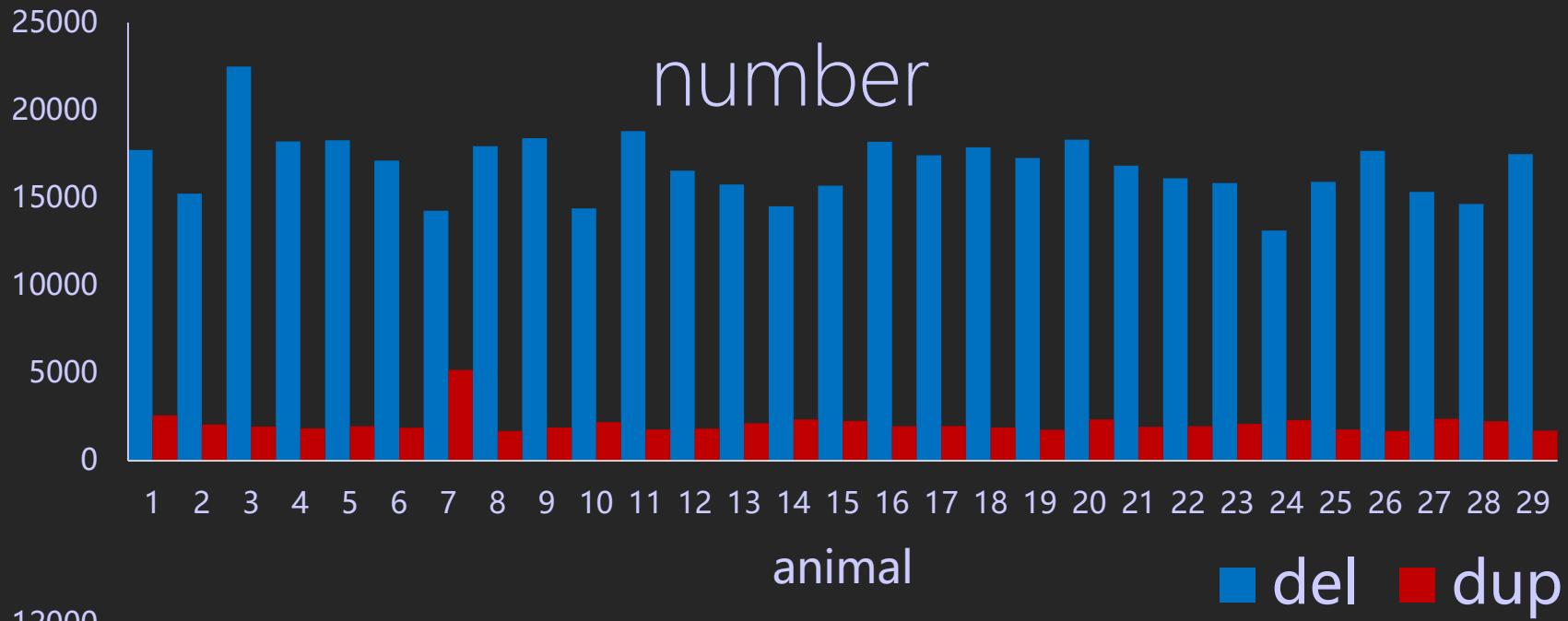
2.71 % – 6.09 %



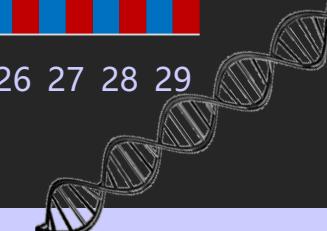
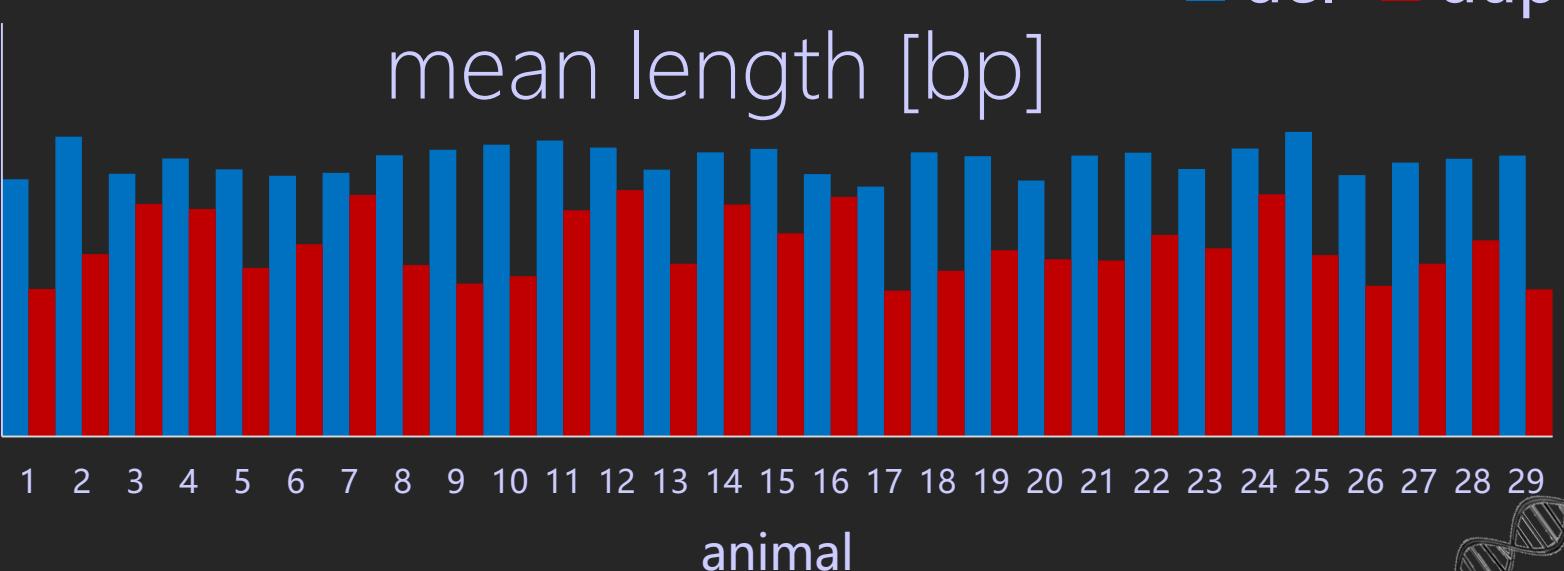
0.51 % – 1.58 %



# Copy Number Variation



mean length [bp]



# Functional annotation

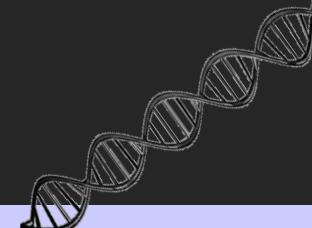
Genome assembly UMD 3.1 (release 2011)

## 1. Sequence Ontology Terms

- define consequences of DNA sequence point mutations
  - e.g. transcript ablation, stop gained, frameshift variant, ...
- cumulated to 8 categories: coding, intron, splice region, non-coding transcript, UTR, up/down stream gene, intergenic

## 2. Gene functional categories

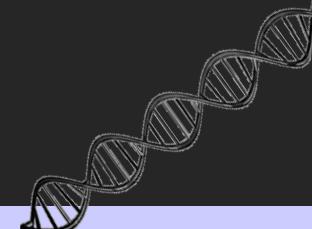
## 3. All genes



# Functional annotation

Genome assembly UMD 3.1 (release 2011)

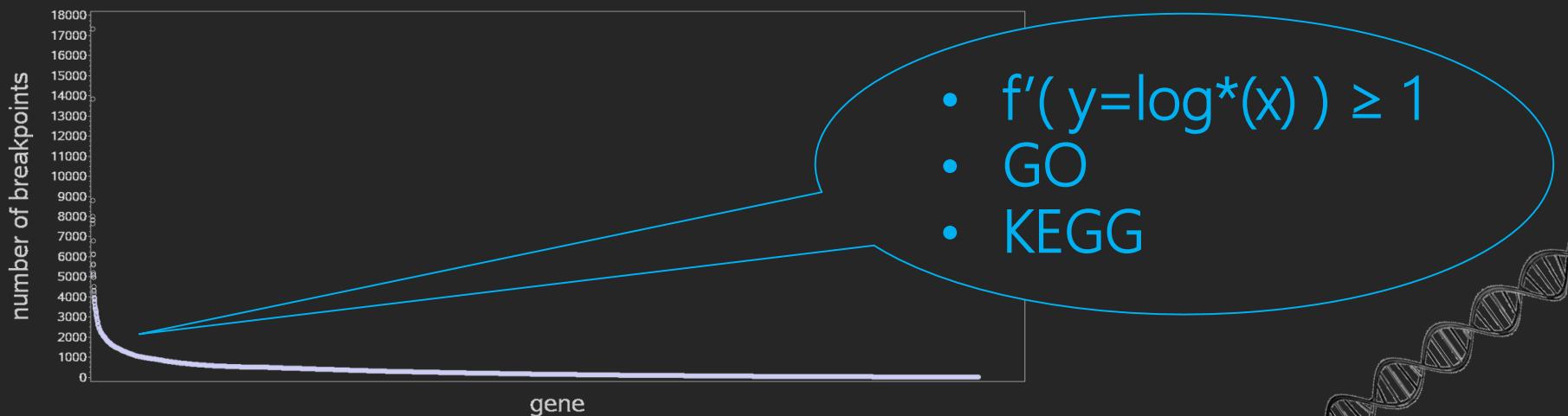
1. Sequence Ontology Terms
2. Gene functional categories
  - housekeeping, under strong selection, neutral
3. All genes



# Functional annotation

Genome assembly UMD 3.1 (release 2011)

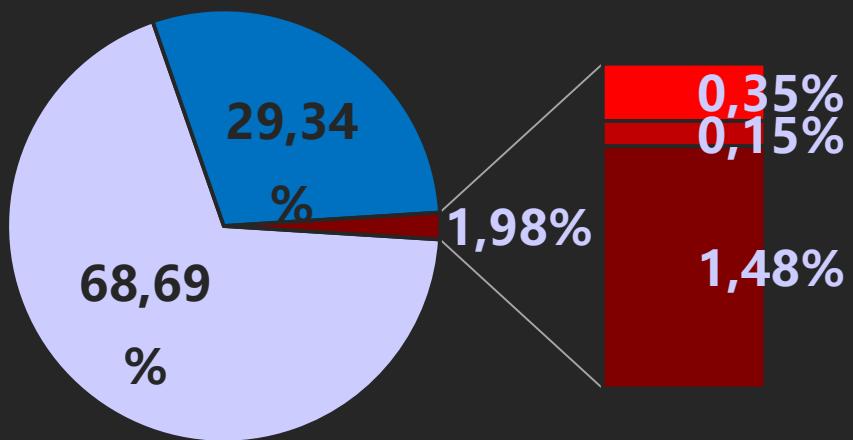
1. Sequence Ontology Terms
2. Gene functional categories
3. All genes
  - genes with a large number of breakpoints



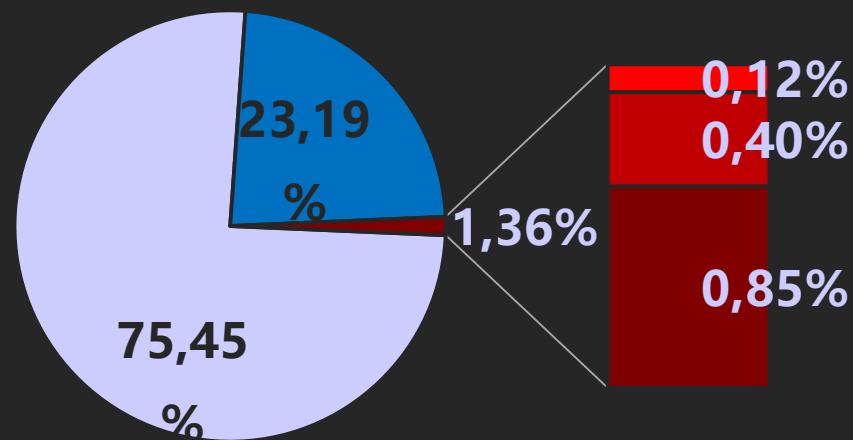
# Functional annotation – results

## Annotation to SO terms

Deletions (487 746)



Duplications (61 789)



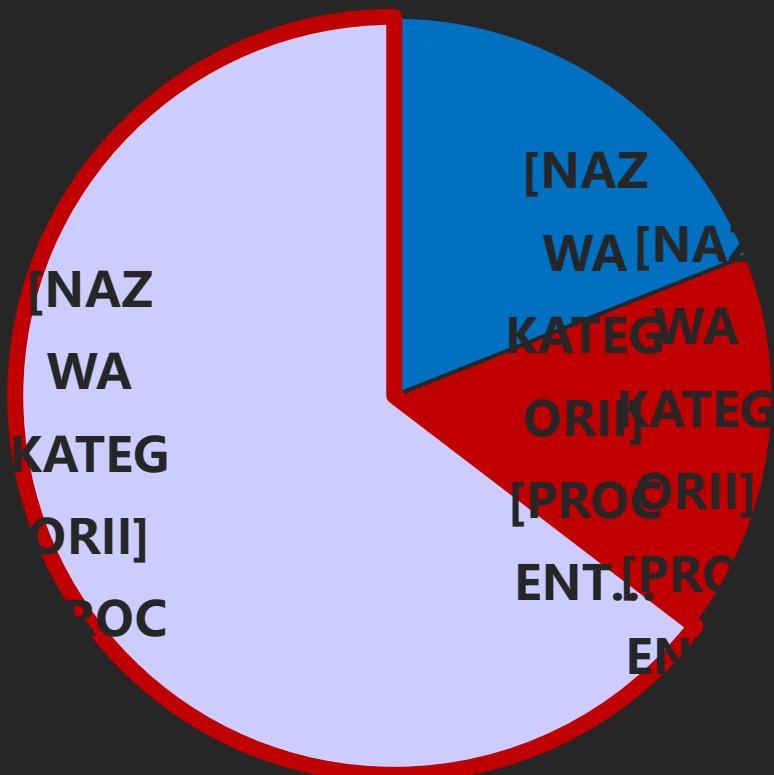
- intergenic
- intron
- non-coding regions of transcripts
- splice regions
- coding regions

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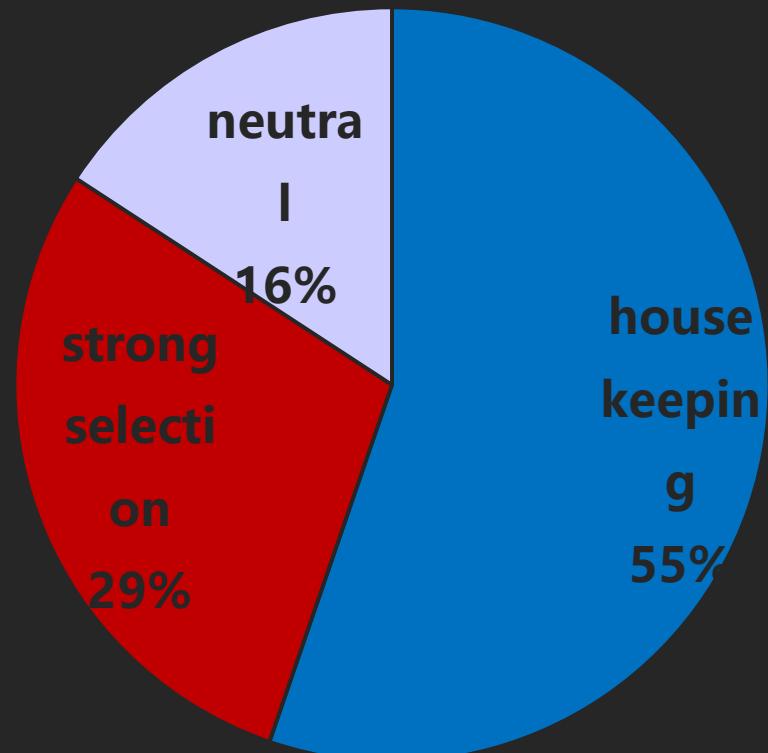
# Functional annotation – results

## Annotation to gene categories

Deletions (199)



Duplications (38)



# Functional annotation – results

Annotation to genes with many breakpoints -

KEGG

## Deletions

- olfactory transduction
- $P=0.11$
- breakpoints depleted
- $Pr(0.001)$  vs  $Pr(0.010)$



## Duplications

- no significance



# Functional annotation – results

Annotation to genes with many breakpoints –  
GO term clusters

Deletions

highest enrichment



regulation of cell  
adhesion

Duplications

highest enrichment



binding of  
macromolecules

# Conclusions

1. Dynamic landscape of CNVs
2. Variability across
3. CNV formation

# Conclusions

1. Dynamic landscape of CNVs

2. Variability across

- Individuals
- Chromosomes
- Functional genome elements

3. CNV formation

# Conclusions

1. Dynamic landscape of CNVs
2. Variability across
3. CNV formation
  - Supressed in coding regions
  - Deletions → suppressed in housekeeping genes and genes under strong selection
  - Duplications → most common in housekeeping genes
  - More duplications → molecule binding → immune response

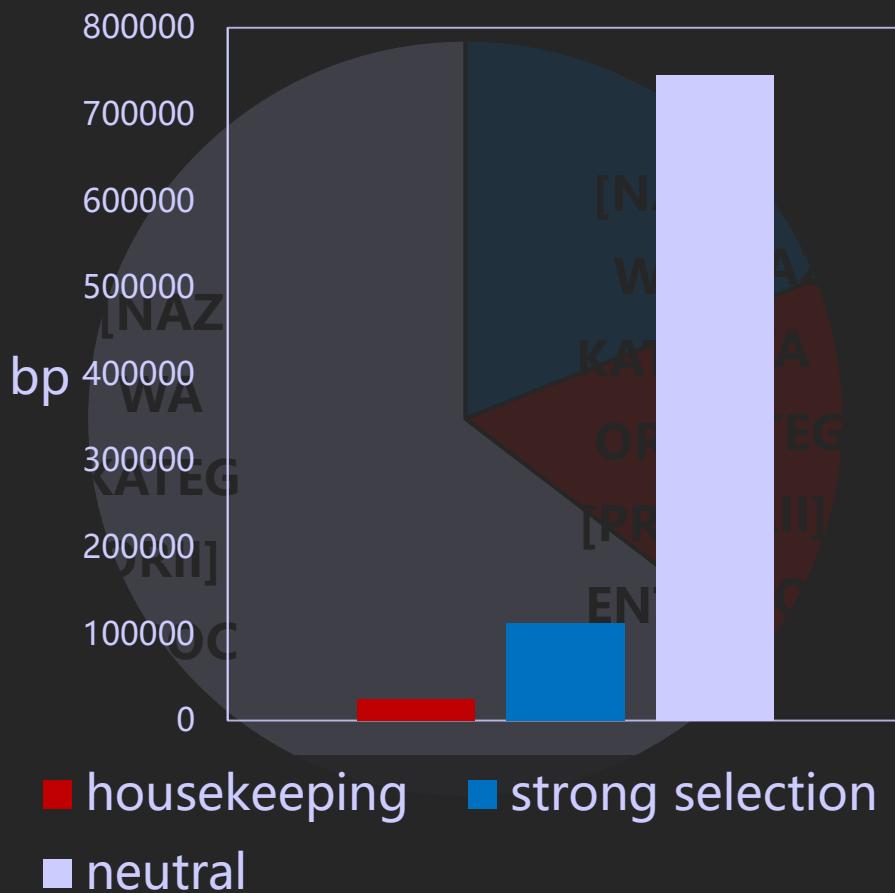
# Thank you for attention



# Functional annotation of CNV breakpoints

## Annotation to gene categories

### Del gene length



### l transcript length

