Short INDELS: genetic markers for adaptive divergence

Original aspects of the short INDELs project

- Divergent natural selection vs neutral processes
- Species with high diversity
- Systems with imperfect genomes can still contain useful functional information

INDEL-SNP comparisons

- 1. Outlier sharing
- 2. Clustering of (different types) markers
- 3. Derived allele frequencies (in progress and for now simply minor)
- 4. Displacement of cline centres

1. Outlier sharing

Total number of Anja's SNP: 55106	Total number of SNP: 11225	Total number of INDEL: 1752
Proportion of SNP with significant clines.	Proportion of SNP with significant clines.	Proportion of INDEL with significant clines.
CZA left: 0.5122128	0.5317595	0.5296804
CZA right: 0.4238377	0.4457016	0.4549087
CZB left: 0.3201829	0.3277506	0.3413242
CZB right: 0.4114071	0.4244989	0.4092466
CZD left: 0.4393351	0.4473942	0.4737443
CZD right: 0.4732697	0.4823163	0.4834475
Proportions of SNP outliers that are shared.	Proportions of SNP outliers that are shared.	Proportions of INDEL outliers that are shared.
CZA left and right: 0.707804	0.6160714	0.7058824
CZB left and right: 0.5680581	0.5178571	0.4705882
CZD left and right: 0.6569873	0.6339286	0.6470588
CZA and CZB: 0.4650635	0.359375	0.3529412
CZA and CZD: 0.508167	0.4107143	0.4117647
CZB and CZD: 0.5426497	0.484375	0.4411765
Number of SNP outliers found in 1 hybrid zone(s): 602	142	24
Number of SNP outliers found in 2 hybrid zone(s): 258	66	7
Number of SNP outliers found in 3 hybrid zone(s): 137	29	7
Number of SNP outliers found in 4 hybrid zone(s): 117	27	5
Number of SNP outliers found in 5 hybrid zone(s): 95	25	1
Number of SNP outliers found in 6 hybrid zone(s): 139	13	3
Prop. of SNP outliers in inversions found in 1 zone(s): 0.648	0.556	0.625
Prop. of SNP outliers in inversions found in 2 zone(s): 0.698	0.636	0.57
Prop. of SNP outliers in inversions found in 3 zone(s): 0.912	0.862	0.86
Prop. of SNP outliers in inversions found in 4 zone(s): 0.923	0.889	1
Prop. of SNP outliers in inversions found in 5 zone(s): 0.979	0.92	1
Prop. of SNP outliers in inversions found in 6 zone(s): 1	1	1

Except this difference in the total number of SNPs, the proportions look quite similar but I have not run any statistical tests

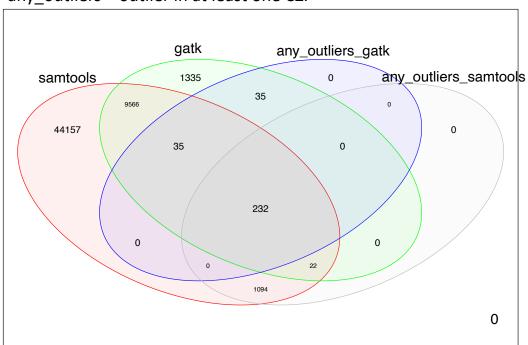
Given the difference between SAMtools and GATK in the total number of SNPs, I have run some diagnostics • Contigs after coverage filter (in progress)

Given the difference between SAMtools and GATK in the total number of SNPs, I have run some diagnostics SNPs after all the filters (in progress)

Given the difference between SAMtools and GATK in the total number of SNPs, I have run some diagnostics

SNPs after all the filters and cline/outlier analysis (i.e., clinal and non-clinal SNPs)

any outliers = outlier in at least one CZ.



all outliers = outlier shared by all six CZs (two per islands).

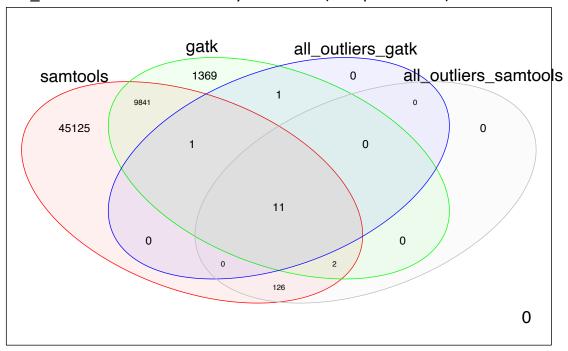
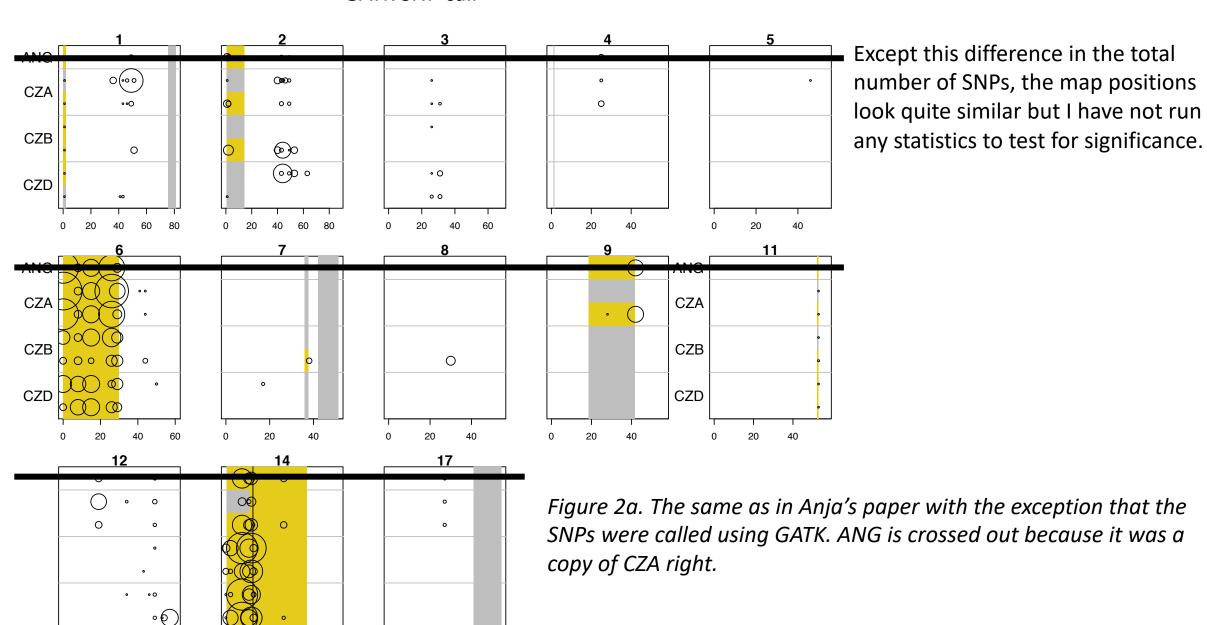


Figure 1. Venn diagrams of the number of SNPs after filtering and cline analysis. Left: SAMtools and GATK calls are intersected with the respective outliers that were at least present in one hybrid zone. Right: SAMtools and GATK calls are intersected with the respective outliers that were present in all six hybrid zones (two per islands).



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GATK INDEL call

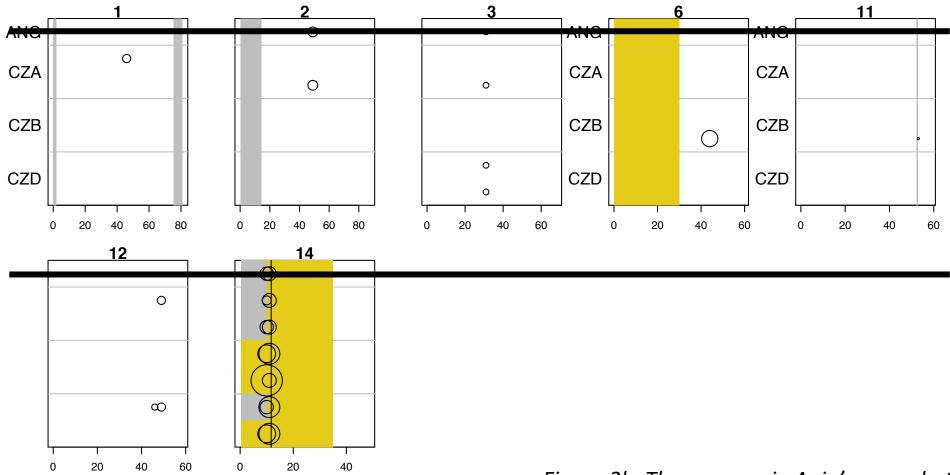


Figure 2b. The same as in Anja's paper but with INDELs. ANG is crossed out because it was a copy of CZA right.

2. Clustering of (different types) markers

- INDELs and SNPs after filtering and cline analysis.
- All six hybrid zones combined.

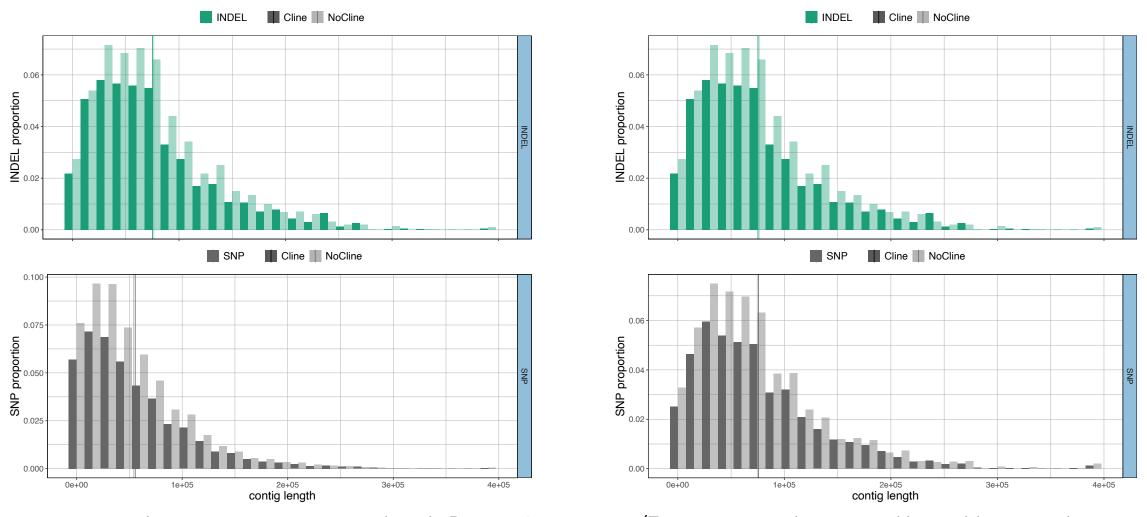


Figure 3a. Marker proportions over contig length. $Proportion = count/\Sigma$ count per marker type and bin width = 15000 base pairs. Clinal variants are dark coloured and non-clinal variants are light coloured. Left: SNP call using SAMtools and INDEL call using GATK. Right: both INDELs and SNPs were called with GATK.

2. Clustering of (different types) markers

All INDELs = clinal + non-clinal

$$n/N = \frac{n_i \ marker}{N \ marker}$$
 $n_i = \text{number of INDELs in contig } i$
 $N = \text{Total number of INDELs}$

Figure 3a. Proportions (left) and counts (right) of INDELs per contig.

... Clustering of (different types) markers

• All SNPs = clinal + non-clinal

$$n/N = \frac{n_i \ marker}{N \ marker}$$
 n_i number of SNPs in contig i
 $N = \text{Total number of SNPs}$

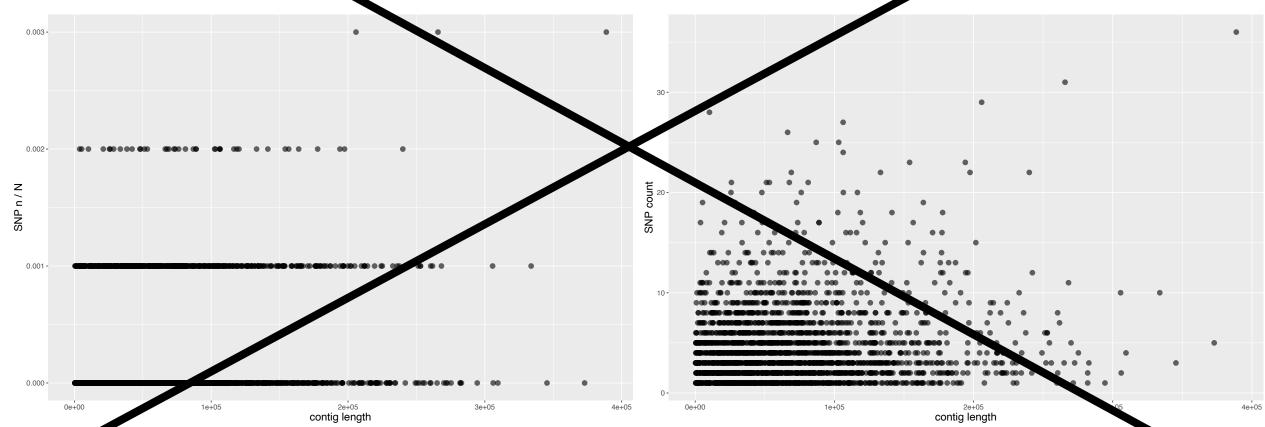


Figure 3b. Proportions (left) and counts (right) of SNPs per contig.

Clustering of (different types) markers All INDEEs and SNPs in the same contigs

Figure 3c. Relationship between SNPs and INDELs with respect to their proportions (left) and their counts (right) per continues of SNPs and INDELs are the same as in Fig. 1-2.

INDEL count

0.003 INDEL n / N

0.004

- 3. Derived allele frequencies (in progress; for now minor allele frequencies Fig. 4-5)
- Ancestral allele was inferred from called genotypes:
 - 1. Reference allele = ancestral allele compressa is homo for the reference allele (0)
 - 2. Alternative allele = ancestral allele compressa is homo for the alternative allele (2)
 - 3. Unknown ancestry compressa is het (1)

3. Minor allele frequencies (GATK call)

- INDELs and SNPs after filtering but before cline analysis.
- All six hybrid zones combined.

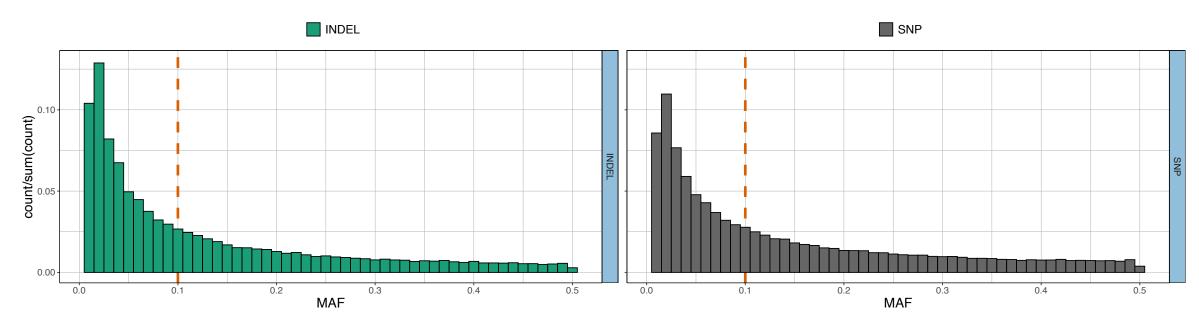


Figure 4a. Proportions of minor allele frequencies of INDELs (left) and SNPs (right) after filtering but before cline analysis. Bin width is 0.01 and orange dashed line marks the maf filter in the cline analysis (0.1).

3. Minor allele frequencies (GATK call)

For the joint AFS between INDELs and SNPs, see (attached) file short_indels.nb.html

- INDELs and SNPs after filtering and cline analysis.
- All six hybrid zones combined.

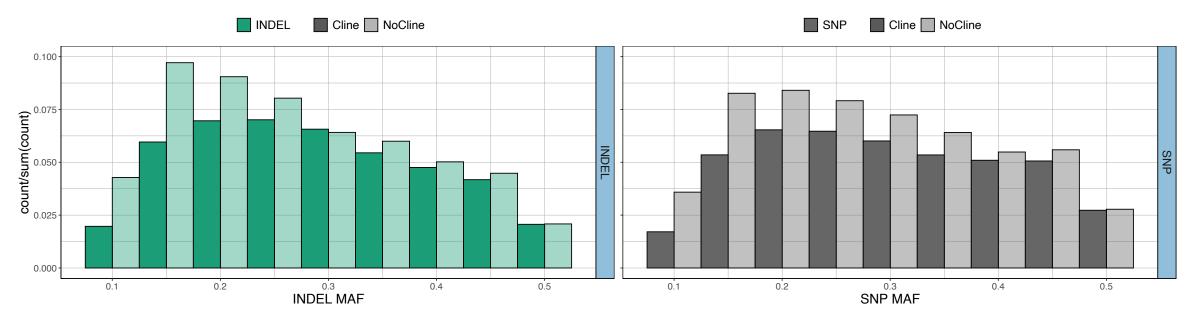


Figure 5a. Proportions of minor allele frequencies of INDELs (left) and SNPs (right) after filtering and cline analysis. Bin width is 0.05. Clinal variants are dark coloured and non-clinal variants are light coloured.