Looking at genic data from SnpEff

Uploading data

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Converting information from SnpEff
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```
In[*]:= fullgenomeSNPEFF = Import[
          "/Users/otto/Documents/Students/AnnaBazzicalupo/MetalAdaptation/Genomics/
            snpEff genic haploid 19May2025.tsv"];
 In[*]:= header = fullgenomeSNPEFF[[1]]
Out[ • ]=
      {CHROM, chr, POS, REF, ALT, QUAL, FORMAT, gene, effect, mut_type, gene_code,
       base_change, aa_change, ANN, OLY077_S101, CdBM23_S1, CdBM25_S10, CdBM26_S18,
       CdBM29_S26, CdBM30_S34, CdBM32_S42, CdBM36_S50, CdBM37_S58, CdBM39_S2,
       CdBM42_S11, CdBM43_S19, CdBM44_S27, CdBM45_S35, CdBM46_S43, CdBM47_S51,
       CdBM48_S59, CoBM12_S4, CoBM14_S13, CoBM15_S21, CoBM16_S29, CoBM17_S37,
       CoBM18_S45, CoBM1_S3, CoBM20_S53, CoBM21_S61, CoBM2_S12, CoBM3_S20, CoBM4_S28,
       CoBM5_S36, CoBM6_S44, CoBM7_S52, CoBM8_S60, CuBM10_S54, CuBM11_S62, CuBM12_S6,
       CuBM13_S15, CuBM14_S23, CuBM15_S31, CuBM16_S39, CuBM17_S47, CuBM18_S55, CuBM3_S5,
       CuBM4_S14, CuBM6_S22, CuBM7_S30, CuBM8_S38, CuBM9_S46, MnBM12_S63, MnBM13_S7,
       MnBM14_S16, MnBM15_S24, MnBM16_S32, MnBM17_S40, MnBM18_S48, MnBM20_S56,
       MnBM21_S64, MnBM23_S8, MnBM24_S17, MnBM25_S25, MnBM27_S33, MnBM28_S41,
       MnBM29_S49, MnBM31_S57, MnBM32_S65, MnBM34_S66, MnBM38_S72, MnBM39_S78,
       MnBM42_S84, MnBM44_S90, NiBM11_S73, NiBM12_S79, NiBM14_S85, NiBM16_S91,
       NiBM17_S97, NiBM21_S103, NiBM22_S109, NiBM24_S68, NiBM25_S74, NiBM27_S80,
       NiBM28_S86, NiBM29_S92, NiBM30_S98, NiBM4_S96, NiBM6_S102, NiBM8_S108, NiBM9_S67,
       ZnBM11_S104, ZnBM12_S110, ZnBM15_S69, ZnBM16_S75, ZnBM17_S81, ZnBM19_S87,
       ZnBM22_S93, ZnBM23_S99, ZnBM25_S105, ZnBM28_S111, ZnBM29_S70, ZnBM31_S76,
       ZnBM34_S82, ZnBM37_S88, ZnBM38_S94, ZnBM39_S100, ZnBM41_S106, ZnBM42_S112,
       ZnBM43_S71, ZnBM44_S77, ZnBM45_S83, ZnBM46_S89, ZnBM47_S95, AC, AF, AN,
       BaseQRankSum, DP, FS, MLEAC, MLEAF, MQ, MQRankSum, QD, ReadPosRankSum, SOR, LOF}
 In[*]:= fullgenomeSNPEFF = Drop[fullgenomeSNPEFF, 1];
 In[*]:= Length[fullgenomeSNPEFF]
Out[ • ]=
      462
 In[*]:= Select[fullgenomeSNPEFF, #[Position[header, "effect"][1, 1]]] == "LOW" &] // Length
Out[ • ]=
      88
```

```
In[*]:= Select[fullgenomeSNPEFF, #[Position[header, "effect"][1, 1]] == "MODERATE" &] // Length
Out[ • ]=
       267
 In[*]:= Select[fullgenomeSNPEFF, #[Position[header, "effect"][1, 1]]] == "HIGH" &] // Length
Out[ • ]=
       107
      Filters (already applied in R) keep only those sites with depth ≥5 and that have some "1|1" genotypes
      when called as diploid (stripping out heterozygous only calls):
 In[*]:= droppos = -1 + Position[header, "OLY077_S101"] [[1, 1]];
       (*# columns to be dropped before first sample*)
 In[*]:= topos = Position[header, "ZnBM47_S95"][[1, 1]]; (*# last metal column*)
```

In[@]:= genelist = Sort[Union[fullgenomeSNPEFF[All, Position[header, "gene"][1, 1]]]] Out[•]=

{ABP1, ACB1, ACC1, ADE6, ADR1, AFT1, AGP1, AHK1, ALD5, ALY2, APC2, ARO3, ASN1, ATG11, ATG2, BDS1, BIR1, BLM10, BNA6, BNI1, BNI4, BRR2, BSC1, BSD2, BUD19, BUL1, BUL2, CAJ1, CCR4, CCW12, CDC25, CIC1, CLB3, CLN3, CNE1, COG1, COG3, COQ1, CPA1, CSM3, CUE3, CWC22, CWH41, CYK3, DAL2, DAN4, DBF20, DBP10, DDR48, DNF1, DOA4, DOT6, DSC2, DSE4, DUS4, EBS1, ECM14, ECM22, EFR3, ERB1, ERG1, ERG27, ERG7, ERG9, FAS2, FAU1, FCY2, FET4, FIG2, FIG4, FLC2, FLO1, FLO9, FMP52, FRE6, FSF1, FYV10, FYV6, FZF1, GAS4, GCD2, GPB1, GPB2, GPH1, GSC2, HAP1, HBT1, HEH2, HHF2, HIS4, HMF1, HO, HRT3, HSL1, HSP104, HUL5, HXT13, IDP1, IES3, ILV2, IMA1, INP2, IOC4, IPI3, IRC20, IRC8, IST2, ISU1, ITC1, KAR3, KGD1, KIN1, KIN82, KRE5, KSP1, KTR3, LAM1, LAM6, LOS1, LRG1, MAM3, MAS2, MCK1, MCM5, MCT1, MDH2, MDL2, MDM30, MET10, MET30, MEX67, MHP1, MKT1, MLF3, MLH3, MLS1, MMS4, MNN9, MOT1, MPA43, MRL1, MRPL36, MSC6, MSS11, MSS2, MTR4, MYO2, NAM8, NAR1, NFT1, NGG1, NIP100, NPL3, NSR1, NUC1, NUG1, OAF1, OCA4, OCA5, OPI1, OSH2, PBP2, PBS2, PCL5, PDA1, PDR1, PDR10, PDX1, PET111, PET127, PFK26, PHM7, PH023, PH081, PH084, PIB1, PIR3, PKP1, PLB2, PMA1, POL3, PPQ1, PRC1, PRI1, PRI2, PRP2, PRP8, PTK2, PYK2, QRI7, RAD16, RAD17, RAD26, RBK1, RCE1, RCK2, RCO1, REC102, RED1, RHO1, RIM11, RIM21, RKM3, RMP1, RNY1, ROG1, RPA135, RPI1, RPL22A, RPL2A, RPS15, RPS2, RPT5, RRB1, RRG7, RRP46, RRP6, RSC1, RSE1, RTC2, SAK1, SAP4, SBE2, SEC8, SET2, SFP1, SHP1, SIW14, SKN1, SMC2, SMY2, SNF7, SNT2, SPC105, SP077, SPS100, SRP40, SSE1, SSK2, SSQ1, SSZ1, STE5, STP3, STT3, SUB2, SUC2, SUM1, SWC5, SWS2, SWT1, SYF2, SYG1, TAF8, TAH11, TAO3, tC(GCA)B, TCO89, tE(UUC)G3, TFB1, TFG1, TIM54, tK(CUU)E2, TOM1, TOP1, TOP2, TOS3, TPS3, TRK2, TRL1, TRT2, tS(AGA)M, TUS1, UBC1, UBP5, UBR2, UBX7, UGA3, UGA4, UIP5, ULS1, UME6, URA2, URB1, USA1, UTP21, UTP8, VBA1, VCX1, VHR2, VID22, VPS13, VPS63, VPS74, VRP1, VTC1, VTC3, VTC4, VTC5, WAR1, YAP6, YBL109W, YBR134W, YBR242W, YBR292C, YCF1, YCK1, YCT1, YDL176W, YDL199C, YDR003W-A, YDR157W, YDR269C, YDR381C-A, YDR541C, YDR544C, YEF1, YER087C-A, YER156C, YFL021C-A, YFR036W-A, YGR126W, YGR130C, YGR266W, YHC1, YHC3, YHL008C, YHR028W-A, YHR071C-A, YIH1, YIL092W, YIR020W-A, YJL211C, YKR073C, YLL020C, YLL066W-B, YLR108C, YLR296W, YLR312C, YLR372W, YML119W, YMR027W, YMR317W, YNL109W, YOR029W, YOR1, YOR296W, YOR343C, YPK1, YPL025C, YPL114W, YPR078C, YPR089W, YPR117W, YRA1, YRB2, ZDS2, ZRT1, ZRT3}

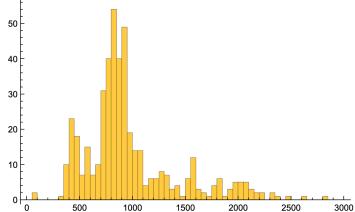
Plotting the depth of coverage at each site:

In[*]:= Max[fullgenomeSNPEFF[All, Position[header, "DP"][1, 1]]]]

Out[•]=

17792

```
In[*]:= fullgenedepth = Histogram[fullgenomeSNPEFF[All, Position[header, "DP"][1, 1]]],
         {50}, PlotRange → {{0, 3000}, Automatic}]
Out[ • ]=
```



There are two low outliers (BSC1 and HAP1, filtered out below for having too many "." genotypes):

```
In[*]:= Select[fullgenomeSNPEFF[All, Position[header, "DP"][1, 1]]], # < 300 &]</pre>
Out[ • ]=
       {91, 84}
```

There are three high outliers (all FLO1, filtered out below for having too many "." genotypes):

```
In[*]:= Select[fullgenomeSNPEFF[All, Position[header, "DP"][1, 1]]], # > 3000 &]
Out[ • ]=
       {15599, 15734, 17792}
```

Mean depth:

```
In[*]:= Mean[fullgenomeSNPEFF[All, 129]] // N
Out[ • ]=
       1072.34
```

Dropping tRNAs, checking overlapping coding regions (SnpEff), and dropping dubious genes

Dropping tRNAs, which we drop {"tC(GCA)B","tE(UUC)G3","tK(CUU)E2","tS(AGA)M"} and TRT2 (which is tT(CGU)K), but first checking that their annotations from SnpEff do not involve other genes (other than upstream or downstream modifiers):

```
In[*]:= dropgenes = {"tC(GCA)B", "tE(UUC)G3", "tK(CUU)E2", "tS(AGA)M", "TRT2"};
     Depth of coverage for these genes is pretty typical:
```

```
<code>In[⊕]:= Select[fullgenomeSNPEFF, MemberQ[dropgenes, #∏8∏] &] ∏All, {8, 129}∏</code>
       Mean[%[All, 2]] // N
Out[ • ]=
       {{tC(GCA)B, 1316}, {tK(CUU)E2, 1111}, {tE(UUC)G3, 398}, {TRT2, 2091}, {tS(AGA)M, 885}}
Out[ • ]=
       1160.2
      Dropping these five tRNAs
 In[*]:= Length[fullgenomeSNPEFF];
      fullgenomeSNPEFF = Select[fullgenomeSNPEFF, MemberQ[dropgenes, #[8]] == False &];
      %% - Length[fullgenomeSNPEFF]
Out[ • ]=
      5
      Looking over all genes whose most extreme SnpEff calls involve more than one protein-coding gene:
 In[*]:= catSNPEFF = {"LOW", "MODERATE", "HIGH"}; (*Not "MODIFIER"*)
 In[*]:= Clear[sumFULL]
      sumFULL = Table["ERROR", {i, 1, Length[fullgenomeSNPEFF]}];
      For[i = 1, i ≤ Length[fullgenomeSNPEFF], i++,
        temp = StringSplit[fullgenomeSNPEFF[i, 14], {"|"}];
        temp2 = Tally[temp];
        temp3 = Select[temp2, MemberQ[catSNPEFF, #[1]] &];
        sumFULL[[i]] = Total[temp3][[2]]
       1
 In[*]:= Position[sumFULL, _?(# > 1 &)]
Out[ • ]=
       \{\{84\}, \{120\}, \{227\}, \{229\}, \{296\}, \{317\}, \{356\}, \{380\}, \{439\}\}\}
 In[*]:= Position[sumFULL, _?(# > 2 &)]
Out[ • ]=
       {}
      #84: YDR269C is dubious, but CCC2 is verified (SGD) [Updated]
 In[*]:= fixme = 84;
 In[*]:= fullgenomeSNPEFF[[fixme, 8]]
Out[ • ]=
      YDR269C
       Manually swapping for the second gene:
 In[*]:= fullgenomeSNPEFF[[fixme]]
Out[ • ]=
       {chrIV, 4, 1005804, G, T, 283.29, GT:AD:DP:GQ:PL, YDR269C,
        MODERATE, missense_variant, YDR269C, c.185C>A, p.Thr62Asn,
```

```
T|missense_variant|MODERATE|YDR269C|YDR269C|transcript|YDR269C_mRNA|
  protein_coding|1/1|c.185C>A|p.Thr62Asn|185/324|185/324|62/107||,T|
  missense_variant|MODERATE|CCC2|YDR270W|transcript|YDR270W_mRNA|protein_coding|
  1/1|c.130G>T|p.Val44Leu|130/3015|130/3015|44/1004||,T|upstream_gene_variant|
  MODIFIER|HEL2|YDR266C|transcript|YDR266C_mRNA|protein_coding||c.-3781C>A|||||
  3781|,T|upstream_gene_variant|MODIFIER|CIA1|YDR267C|transcript|YDR267C_mRNA|
  protein_coding||c.-2302C>A|||||2302|,T|upstream_gene_variant|MODIFIER|GL02|
  YDR272W|transcript|YDR272W_mRNA|protein_coding||c.-3206G>T|||||3206|,T|
  upstream_gene_variant|MODIFIER|DON1|YDR273W|transcript|YDR273W_mRNA|
  protein_coding||c.-4372G>T|||||4372|,T|downstream_gene_variant|MODIFIER|MSW1|
  YDR268W|transcript|YDR268W_mRNA|protein_coding||c.*661G>T|||||661|,T|
  downstream_gene_variant|MODIFIER|YDR271C|YDR271C|transcript|YDR271C_mRNA|
  protein_coding||c.*2591C>A|||||2591|, 0:3,0:3:99:0,109, 0:5,0:5:99:0,135,
0:3,0:3:99:0,113, 0:4,0:4:99:0,104, 0:3,0:3:99:0,100, 0:4,0:4:99:0,115,
0:3,0:3:99:0,100, 0:3,0:3:99:0,100, 0:4,0:4:99:0,107, 0:4,0:4:99:0,100,
0:4,0:4:99:0,137, 0:3,0:3:99:0,100, 0:4,0:4:99:0,109, 0:7,0:7:99:0,125,
0:3,0:3:99:0,100,0:4,0:4:99:0,141,0:3,0:3:99:0,105,0:3,0:3:99:0,103,
0:3,0:3:99:0,109, 0:4,0:4:99:0,100, 0:3,0:3:99:0,100, 0:3,0:3:99:0,114,
0:4,0:4:99:0,141,0:3,0:3:99:0,100,0:5,0:5:99:0,128,0:3,0:3:99:0,122,
0:5,0:5:99:0,116, 1:0,11:11:99:304,0, 0:3,0:3:99:0,100, 0:3,0:3:99:0,100,
0:4,0:4:99:0,155, 0:3,0:3:99:0,118, 0:3,0:3:99:0,100, 0:4,0:4:99:0,135,
0:4,0:4:99:0,100, 0:3,0:3:99:0,99, 0:4,0:4:99:0,104, 0:3,0:3:99:0,100,
0:3,0:3:99:0,100, 0:8,0:8:99:0,148, 0:3,0:3:99:0,109, 0:4,0:4:99:0,154,
0:3,0:3:99:0,103,\ 0:3,0:3:99:0,100,\ 0:3,0:3:99:0,115,\ 0:4,0:4:99:0,132,
0:4,0:4:99:0,103, 0:4,0:4:99:0,131, 0:3,0:3:99:0,100, 0:3,0:3:99:0,122,
0:3,0:3:99:0,110,0:3,0:3:99:0,100,0:3,0:3:99:0,107,0:3,0:3:99:0,100,
0:3,0:3:99:0,100, 0:4,0:4:99:0,118, 0:3,0:3:99:0,100, 0:6,0:6:99:0,131,
0:3,0:3:99:0,100, 0:3,0:3:99:0,118, 0:5,0:5:99:0,122, 0:3,0:3:99:0,100,
0:4,0:4:99:0,132, 0:4,0:4:99:0,131, 0:3,0:3:99:0,100, 0:3,0:3:99:0,100,
0:3,0:3:99:0,113, 0:3,0:3:99:0,100, 0:3,0:3:99:0,100, 0:3,0:3:99:0,109,
0:3,0:3:99:0,100, 0:3,0:3:99:0,100, 0:3,0:3:99:0,112, 0:4,0:4:99:0,137,
0:3,0:3:99:0,118, 0:3,0:3:99:0,100, 0:5,0:5:99:0,159, 0:3,0:3:99:0,122,
0:3,0:3:99:0,109, 0:4,0:4:99:0,104, 0:3,0:3:99:0,114, 0:5,0:5:99:0,170,
0:5,0:5:99:0,144, 0:4,0:4:99:0,141, 0:3,0:3:99:0,122, 0:4,0:4:99:0,155,
0:10,0:10:99:0,153,0:3,0:3:99:0,100,0:3,0:3:99:0,122,0:3,0:3:99:0,100,
0:3,0:3:99:0,118, 0:4,0:4:99:0,119, 0:4,0:4:99:0,137, 0:5,0:5:99:0,141,
0:3,0:3:99:0,118, 0:4,0:4:99:0,126, 0:3,0:3:99:0,100, 0:3,0:3:99:0,100,
0:4,0:4:99:0,133, 0:9,0:9:99:0,157, 0:7,0:7:99:0,143, 0:4,0:4:99:0,100,
0:4,0:4:99:0,129, 0:3,0:3:99:0,100, 0:6,0:6:99:0,166, 0:6,0:6:99:0,180,
0:7,0:7:99:0,173, 0:4,0:4:99:0,104, 0:4,0:4:99:0,109, 0:3,0:3:99:0,114,
1, 0.009091, 110, Missing[NotAvailable], 426, 0., 1, 0.009091, 60.,
Missing[NotAvailable], 25.75, Missing[NotAvailable], 2.494, Missing[NotAvailable]}
```

In[*]:= fullgenomeSNPEFF[fixme] = {"chrIV", 4, 1005 804, "G", "T", 283.29`, "GT:AD:DP:GQ:PL", "CCC2", "MODERATE", "missense_variant", "YDR270W", "c.130G>T", "p.Val44Leu",

```
"T|missense_variant|MODERATE|YDR269C|YDR269C|transcript|YDR269C_mRNA|
  protein coding|1/1|c.185C>A|p.Thr62Asn|185/324|185/324|62/107||,T|
  missense_variant|MODERATE|CCC2|YDR270W|transcript|YDR270W_mRNA|
  protein_coding|1/1|c.130G>T|p.Val44Leu|130/3015|130/3015|44/1004||,T|
  upstream gene variant|MODIFIER|HEL2|YDR266C|transcript|YDR266C mRNA|
  protein coding||c.-3781C>A|||||3781|,T|upstream gene variant|MODIFIER|CIA1
  |YDR267C|transcript|YDR267C_mRNA|protein_coding||c.-2302C>A|||||2302|,T|
  upstream gene variant|MODIFIER|GLO2|YDR272W|transcript|YDR272W mRNA|
  protein_coding||c.-3206G>T|||||3206|,T|upstream_gene_variant|MODIFIER|DON1
  |YDR273W|transcript|YDR273W_mRNA|protein_coding||c.-4372G>T||||4372|,T|
  downstream gene variant|MODIFIER|MSW1|YDR268W|transcript|YDR268W mRNA|
  protein_coding||c.*661G>T|||||661|,T|downstream_gene_variant|MODIFIER|
  YDR271C|YDR271C|transcript|YDR271C mRNA|protein coding||c.*2591C>A|||||
  2591|", "0:3,0:3:99:0,109", "0:5,0:5:99:0,135",
"0:3,0:3:99:0,113", "0:4,0:4:99:0,104", "0:3,0:3:99:0,100", "0:4,0:4:99:0,115",
"0:3,0:3:99:0,100", "0:3,0:3:99:0,100", "0:4,0:4:99:0,107", "0:4,0:4:99:0,100",
"0:4,0:4:99:0,137", "0:3,0:3:99:0,100", "0:4,0:4:99:0,109", "0:7,0:7:99:0,125",
"0:3,0:3:99:0,100", "0:4,0:4:99:0,141", "0:3,0:3:99:0,105", "0:3,0:3:99:0,103",
"0:3,0:3:99:0,109", "0:4,0:4:99:0,100", "0:3,0:3:99:0,100", "0:3,0:3:99:0,114",
"0:4,0:4:99:0,141", "0:3,0:3:99:0,100", "0:5,0:5:99:0,128", "0:3,0:3:99:0,122",
"0:5,0:5:99:0,116", "1:0,11:11:99:304,0", "0:3,0:3:99:0,100",
"0:3,0:3:99:0,100", "0:4,0:4:99:0,155", "0:3,0:3:99:0,118", "0:3,0:3:99:0,100",
"0:4,0:4:99:0,135", "0:4,0:4:99:0,100", "0:3,0:3:99:0,99", "0:4,0:4:99:0,104",
"0:3,0:3:99:0,100", "0:3,0:3:99:0,100", "0:8,0:8:99:0,148", "0:3,0:3:99:0,109",
"0:4,0:4:99:0,154", "0:3,0:3:99:0,103", "0:3,0:3:99:0,100", "0:3,0:3:99:0,115",
"0:4,0:4:99:0,132", "0:4,0:4:99:0,103", "0:4,0:4:99:0,131", "0:3,0:3:99:0,100",
"0:3,0:3:99:0,122", "0:3,0:3:99:0,110", "0:3,0:3:99:0,100", "0:3,0:3:99:0,107",
"0:3,0:3:99:0,100", "0:3,0:3:99:0,100", "0:4,0:4:99:0,118", "0:3,0:3:99:0,100",
"0:6,0:6:99:0,131", "0:3,0:3:99:0,100", "0:3,0:3:99:0,118", "0:5,0:5:99:0,122",
"0:3,0:3:99:0,100", "0:4,0:4:99:0,132", "0:4,0:4:99:0,131", "0:3,0:3:99:0,100",
"0:3,0:3:99:0,100", "0:3,0:3:99:0,113", "0:3,0:3:99:0,100", "0:3,0:3:99:0,100",
"0:3,0:3:99:0,109", "0:3,0:3:99:0,100", "0:3,0:3:99:0,100", "0:3,0:3:99:0,112",
"0:4,0:4:99:0,137", "0:3,0:3:99:0,118", "0:3,0:3:99:0,100", "0:5,0:5:99:0,159",
"0:3,0:3:99:0,122", "0:3,0:3:99:0,109", "0:4,0:4:99:0,104", "0:3,0:3:99:0,114",
"0:5,0:5:99:0,170", "0:5,0:5:99:0,144", "0:4,0:4:99:0,141", "0:3,0:3:99:0,122",
"0:4,0:4:99:0,155", "0:10,0:10:99:0,153", "0:3,0:3:99:0,100",
"0:3,0:3:99:0,122", "0:3,0:3:99:0,100", "0:3,0:3:99:0,118", "0:4,0:4:99:0,119",
"0:4,0:4:99:0,137", "0:5,0:5:99:0,141", "0:3,0:3:99:0,118", "0:4,0:4:99:0,126",
"0:3,0:3:99:0,100", "0:3,0:3:99:0,100", "0:4,0:4:99:0,133", "0:9,0:9:99:0,157",
"0:7,0:7:99:0,143", "0:4,0:4:99:0,100", "0:4,0:4:99:0,129", "0:3,0:3:99:0,100",
"0:6,0:6:99:0,166", "0:6,0:6:99:0,180", "0:7,0:7:99:0,173",
"0:4,0:4:99:0,104", "0:4,0:4:99:0,109", "0:3,0:3:99:0,114", 1, 0.009091',
110, "NA", 426, 0.`, 1, 0.009091`, 60.`, "NA", 25.75`, "NA", 2.494`, "NA"};
```

#120: YFL021C-A is dubious, but GAT1 is verified (SGD) [Updated]

In[•]:= fixme = 120;

Manually swapping for the second gene:

In[*]:= fullgenomeSNPEFF[[fixme]]

Out[• 1=

```
{chrVI, 6, 96312, C, T, 839.29, GT:AD:DP:GQ:PL, YFL021C-A,
MODERATE, missense_variant, YFL021C-A, c.304G>A, p.Ala102Thr,
T|missense_variant|MODERATE|YFL021C-A|YFL021C-A|transcript|YFL021C-A_mRNA|
   protein_coding|1/1|c.304G>A|p.Ala102Thr|304/855|304/855|102/284||,T|
  missense_variant|MODERATE|GAT1|YFL021W|transcript|YFL021W_mRNA|protein_coding|
  1/1|c.347C>T|p.Ala116Val|347/1533|347/1533|116/510||,T|upstream_gene_variant|
  MODIFIER|FRS2|YFL022C|transcript|YFL022C_mRNA|protein_coding||c.-1302G>A|||||
   1302|,T|downstream_gene_variant|MODIFIER|BUD27|YFL023W|transcript|YFL023W_mRNA
   |protein_coding||c.*2936C>T|||||2936|,T|downstream_gene_variant|MODIFIER|PAU5|
  YFL020C|transcript|YFL020C_mRNA|protein_coding||c.*2919G>A|||||2919|,T|
  downstream_gene_variant|MODIFIER|YFL019C|YFL019C|transcript|YFL019C_mRNA|
  protein_coding||c.*3940G>A|||||3940|, 0:13,0:13:99:0,393,
0:12,0:12:99:0,329, 0:8,0:8:99:0,236, 0:20,0:20:99:0,484, 0:8,0:8:99:0,200,
0:14,0:14:99:0,452, 0:13,0:13:99:0,239, 0:24,0:24:99:0,545,
0:10,0:10:99:0,330, 0:11,0:11:99:0,348, 0:8,0:8:99:0,220, 0:10,0:10:99:0,267,
0:10,0:10:99:0,297, 0:16,0:16:99:0,469, 0:11,0:11:99:0,292, 0:14,0:14:99:0,399,
0:11,0:11:99:0,306, 0:9,0:9:99:0,278, 0:12,0:12:99:0,314, 0:11,0:11:99:0,401,
0:8,0:8:99:0,232, 0:10,0:10:99:0,294, 0:10,0:10:99:0,313, 0:16,0:16:99:0,464,
0:27,0:27:99:0,714,0:16,0:16:99:0,423,0:11,0:11:99:0,288,0:11,0:11:99:0,251,
0:6,0:6:99:0,172, 0:8,0:8:99:0,270, 0:19,0:19:99:0,514, 0:26,0:26:99:0,827,
0:23,0:23:99:0,744, 0:11,0:11:99:0,358, 0:21,0:21:99:0,582, 0:10,0:10:99:0,278,
0:3,0:3:99:0,109, 0:4,0:4:99:0,116, 0:7,0:7:99:0,121, 0:12,0:12:99:0,367,
0:17,0:17:99:0,344, 0:21,0:21:99:0,495, 0:10,0:10:99:0,270, 0:7,0:7:99:0,101,
0:11,0:11:99:0,279,0:12,0:12:99:0,331,0:6,0:6:99:0,178,0:9,0:9:99:0,274,
0:18,0:18:99:0,587, 0:7,0:7:99:0,221, 0:6,0:6:99:0,146, 0:6,0:6:99:0,124,
0:7,0:7:99:0,244, 0:12,0:12:99:0,384, 0:15,0:15:99:0,474, 0:10,0:10:99:0,129,
0:15,0:15:99:0,326,0:9,0:9:99:0,214,0:12,0:12:99:0,363,0:8,0:8:99:0,200,
0:5,0:5:99:0,140,0:7,0:7:99:0,200,0:11,0:11:99:0,377,0:16,0:16:99:0,323,
1:0,25:25:99:860,0,0:18,0:18:99:0,529,0:24,0:24:99:0,702,
0:17,0:17:99:0,510, 0:17,0:17:99:0,492, 0:15,0:15:99:0,504, 0:14,0:14:99:0,451,
0:10,0:10:99:0,308, 0:11,0:11:99:0,276, 0:9,0:9:99:0,247, 0:11,0:11:99:0,329,
0:14,0:14:99:0,421, 0:17,0:17:99:0,458, 0:28,0:28:99:0,800, 0:19,0:19:99:0,562,
0:12,0:12:99:0,359, 0:16,0:16:99:0,517, 0:9,0:9:99:0,266, 0:16,0:16:99:0,368,
0:8,0:8:99:0,268, 0:13,0:13:99:0,358, 0:15,0:15:99:0,285, 0:15,0:15:99:0,445,
0:13,0:13:99:0,452, 0:19,0:19:99:0,638, 0:7,0:7:99:0,203, 0:12,0:12:99:0,306,
0:6,0:6:99:0,100, 0:9,0:9:99:0,296, 0:6,0:6:99:0,200, 0:11,0:11:99:0,236,
0:12,0:12:99:0,364,0:14,0:14:99:0,396,0:18,0:18:99:0,550,0:19,0:19:99:0,485,
0:12,0:12:99:0,247, 0:9,0:9:99:0,306, 0:12,0:12:99:0,416, 0:5,0:5:99:0,100,
0:13,0:13:99:0,408, 0:9,0:9:99:0,306, 0:17,0:17:99:0,364, 0:9,0:9:99:0,180,
0:8,0:8:99:0,274, 0:12,0:12:99:0,302, 0:6,0:6:99:0,186, 1, 0.009091, 110,
Missing[NotAvailable], 1372, 0., 1, 0.009091, 60., Missing[NotAvailable],
33.57, Missing[NotAvailable], 2.584, Missing[NotAvailable]}
```

In[*]:= fullgenomeSNPEFF[fixme] = {"chrVI", 6, 96312, "C", "T", 839.29`, "GT:AD:DP:GQ:PL", "GAT1", "MODERATE", "missense_variant", "YFL021W", "c.347C>T", "p.Ala116Val",

```
"T|missense_variant|MODERATE|YFL021C-A|YFL021C-A|transcript|YFL021C-A mRNA|
  protein_coding|1/1|c.304G>A|p.Ala102Thr|304/855|304/855|102/284||,T|
  missense_variant|MODERATE|GAT1|YFL021W|transcript|YFL021W_mRNA|
  protein_coding|1/1|c.347C>T|p.Ala116Val|347/1533|347/1533|116/510||,T|
  upstream gene variant|MODIFIER|FRS2|YFL022C|transcript|YFL022C mRNA|
  protein_coding||c.-1302G>A|||||1302|,T|downstream_gene_variant|MODIFIER|
  BUD27|YFL023W|transcript|YFL023W_mRNA|protein_coding||c.*2936C>T|||||2936|
  ,T|downstream gene variant|MODIFIER|PAU5|YFL020C|transcript|YFL020C mRNA|
  protein_coding||c.*2919G>A|||||2919|,T|downstream_gene_variant|MODIFIER|
  YFL019C|YFL019C|transcript|YFL019C_mRNA|protein_coding||c.*3940G>A|||||
  3940|", "0:13,0:13:99:0,393", "0:12,0:12:99:0,329",
"0:8,0:8:99:0,236", "0:20,0:20:99:0,484", "0:8,0:8:99:0,200",
"0:14,0:14:99:0,452", "0:13,0:13:99:0,239", "0:24,0:24:99:0,545",
"0:10,0:10:99:0,330", "0:11,0:11:99:0,348", "0:8,0:8:99:0,220",
"0:10,0:10:99:0,267", "0:10,0:10:99:0,297", "0:16,0:16:99:0,469",
"0:11,0:11:99:0,292", "0:14,0:14:99:0,399", "0:11,0:11:99:0,306",
"0:9,0:9:99:0,278", "0:12,0:12:99:0,314", "0:11,0:11:99:0,401",
"0:8,0:8:99:0,232", "0:10,0:10:99:0,294", "0:10,0:10:99:0,313",
"0:16,0:16:99:0,464", "0:27,0:27:99:0,714", "0:16,0:16:99:0,423",
"0:11,0:11:99:0,288", "0:11,0:11:99:0,251", "0:6,0:6:99:0,172",
"0:8,0:8:99:0,270", "0:19,0:19:99:0,514", "0:26,0:26:99:0,827",
"0:23,0:23:99:0,744", "0:11,0:11:99:0,358", "0:21,0:21:99:0,582",
"0:10,0:10:99:0,278", "0:3,0:3:99:0,109", "0:4,0:4:99:0,116",
"0:7,0:7:99:0,121", "0:12,0:12:99:0,367", "0:17,0:17:99:0,344",
"0:21,0:21:99:0,495", "0:10,0:10:99:0,270", "0:7,0:7:99:0,101",
"0:11,0:11:99:0,279", "0:12,0:12:99:0,331", "0:6,0:6:99:0,178",
"0:9,0:9:99:0,274", "0:18,0:18:99:0,587", "0:7,0:7:99:0,221",
"0:6,0:6:99:0,146", "0:6,0:6:99:0,124", "0:7,0:7:99:0,244",
"0:12,0:12:99:0,384", "0:15,0:15:99:0,474", "0:10,0:10:99:0,129",
"0:15,0:15:99:0,326", "0:9,0:9:99:0,214", "0:12,0:12:99:0,363",
"0:8,0:8:99:0,200", "0:5,0:5:99:0,140", "0:7,0:7:99:0,200",
"0:11,0:11:99:0,377", "0:16,0:16:99:0,323", "1:0,25:25:99:860,0",
"0:18,0:18:99:0,529", "0:24,0:24:99:0,702", "0:17,0:17:99:0,510",
"0:17,0:17:99:0,492", "0:15,0:15:99:0,504", "0:14,0:14:99:0,451",
"0:10,0:10:99:0,308", "0:11,0:11:99:0,276", "0:9,0:9:99:0,247",
"0:11,0:11:99:0,329", "0:14,0:14:99:0,421", "0:17,0:17:99:0,458",
"0:28,0:28:99:0,800", "0:19,0:19:99:0,562", "0:12,0:12:99:0,359",
"0:16,0:16:99:0,517", "0:9,0:9:99:0,266", "0:16,0:16:99:0,368",
"0:8,0:8:99:0,268", "0:13,0:13:99:0,358", "0:15,0:15:99:0,285",
"0:15,0:15:99:0,445", "0:13,0:13:99:0,452", "0:19,0:19:99:0,638",
"0:7,0:7:99:0,203", "0:12,0:12:99:0,306", "0:6,0:6:99:0,100",
"0:9,0:9:99:0,296", "0:6,0:6:99:0,200", "0:11,0:11:99:0,236",
"0:12,0:12:99:0,364", "0:14,0:14:99:0,396", "0:18,0:18:99:0,550",
"0:19,0:19:99:0,485", "0:12,0:12:99:0,247", "0:9,0:9:99:0,306",
```

```
"0:12,0:12:99:0,416", "0:5,0:5:99:0,100", "0:13,0:13:99:0,408",
"0:9,0:9:99:0,306", "0:17,0:17:99:0,364", "0:9,0:9:99:0,180",
"0:8,0:8:99:0,274", "0:12,0:12:99:0,302", "0:6,0:6:99:0,186", 1, 0.009091`,
110, "NA", 1372, 0.`, 1, 0.009091`, 60.`, "NA", 33.57`, "NA", 2.584`, "NA"};
```

#227: YJL211C is dubious, but PEX2 is verified (SGD) [Updated]

```
In[ • ]:= fixme = 227;
```

Manually swapping for the second gene:

```
In[*]:= fullgenomeSNPEFF[[fixme]]
```

Out[•]=

```
{chrX, 10, 37037, C, T, 1239.29, GT:AD:DP:GQ:PL, YJL211C,
MODERATE, missense_variant, YJL211C, c.164G>A, p.Cys55Tyr,
T|missense variant|MODERATE|YJL211C|YJL211C|transcript|YJL211C mRNA|
   protein_coding|1/1|c.164G>A|p.Cys55Tyr|164/444|164/444|55/147||,T|
   missense_variant|MODERATE|PEX2|YJL210W|transcript|YJL210W_mRNA|protein_coding|
   1/1|c.119C>T|p.Ala40Val|119/816|119/816|40/271||,T|upstream_gene_variant|
   MODIFIER|OPT1|YJL212C|transcript|YJL212C_mRNA|protein_coding||c.-788G>A|||||
   788|,T|upstream_gene_variant|MODIFIER|CBP1|YJL209W|transcript|YJL209W_mRNA|
   protein_coding||c.-968C>T|||||968|,T|downstream_gene_variant|MODIFIER|YJL213W|
   YJL213W|transcript|YJL213W_mRNA|protein_coding||c.*3879C>T|||||3879|,T|
   downstream_gene_variant|MODIFIER|NUC1|YJL208C|transcript|YJL208C_mRNA|
   protein_coding||c.*3157G>A|||||3157|,T|downstream_gene_variant|MODIFIER|LAA1|
   YJL207C|transcript|YJL207C_mRNA|protein_coding||c.*4352G>A|||||4352|,
0:3,0:3:99:0,100, 0:8,0:8:99:0,129, 0:7,0:7:99:0,189, 0:3,0:3:99:0,109,
0:4,0:4:99:0,118, 0:3,0:3:99:0,111, 0:4,0:4:99:0,135, 0:3,0:3:99:0,100,
0:4,0:4:99:0,135, 0:3,0:3:99:0,109, 0:3,0:3:99:0,100, 0:4,0:4:99:0,119,
0:3,0:3:99:0,100, 0:4,0:4:99:0,135, 0:3,0:3:99:0,110, 0:3,0:3:99:0,100,
0:6,0:6:99:0,116, 0:3,0:3:99:0,100, 0:3,0:3:99:0,100, 0:3,0:3:99:0,104,
0:3,0:3:99:0,109, 0:4,0:4:99:0,119, 0:3,0:3:99:0,103, 0:4,0:4:99:0,114,
0:4,0:4:99:0,115, 0:4,0:4:99:0,119, 0:4,0:4:99:0,110, 0:4,0:4:99:0,141,
0:3,0:3:99:0,113, 0:3,0:3:99:0,100, 0:4,0:4:99:0,107, 0:3,0:3:99:0,100,
0:4,0:4:99:0,110, 0:4,0:4:99:0,130, 0:3,0:3:99:0,122, 0:3,0:3:99:0,100,
0:4,0:4:99:0,100, 0:4,0:4:99:0,115, 0:4,0:4:99:0,137, 0:3,0:3:99:0,109,
0:3,0:3:99:0,100, 0:3,0:3:99:0,105, 0:3,0:3:99:0,100, 0:4,0:4:99:0,151,
0:3,0:3:99:0,110,0:3,0:3:99:0,104,0:8,0:8:99:0,176,0:6,0:6:99:0,119,
0:3,0:3:99:0,100, 0:3,0:3:99:0,100, 0:3,0:3:99:0,113, 0:3,0:3:99:0,100,
0:3,0:3:99:0,100,0:4,0:4:99:0,100,0:3,0:3:99:0,100,0:3,0:3:99:0,103,
0:3,0:3:99:0,118, 0:3,0:3:99:0,100, 0:3,0:3:99:0,107, 0:3,0:3:99:0,107,
0:3,0:3:99:0,100, 0:7,0:7:99:0,227, 0:3,0:3:99:0,109, 0:3,0:3:99:0,113,
0:3,0:3:99:0,100, 0:4,0:4:99:0,139, 0:4,0:4:99:0,115, 0:4,0:4:99:0,114,
1:0,39:39:99:1260,0,0:5,0:5:99:0,132,0:4,0:4:99:0,133,
0:3,0:3:99:0,109, 0:4,0:4:99:0,110, 0:4,0:4:99:0,102, 0:3,0:3:99:0,100,
0:3,0:3:99:0,103, 0:5,0:5:99:0,135, 0:5,0:5:99:0,174, 0:3,0:3:99:0,100,
0:3,0:3:99:0,100, 0:3,0:3:99:0,107, 0:4,0:4:99:0,135, 0:3,0:3:99:0,104,
0:4,0:4:99:0,115, 0:4,0:4:99:0,128, 0:4,0:4:99:0,124, 0:5,0:5:99:0,181,
0:3,0:3:99:0,100, 0:4,0:4:99:0,123, 0:3,0:3:99:0,114, 0:4,0:4:99:0,132,
0:3,0:3:99:0,101, 0:3,0:3:99:0,100, 0:3,0:3:99:0,109, 0:3,0:3:99:0,100,
0:4,0:4:99:0,127, 0:3,0:3:99:0,117, 0:3,0:3:99:0,118, 0:3,0:3:99:0,100,
0:4,0:4:99:0,111, 0:4,0:4:99:0,114, 0:4,0:4:99:0,114, 0:3,0:3:99:0,100,
0:5,0:5:99:0,113, 0:4,0:4:99:0,135, 0:4,0:4:99:0,122, 0:3,0:3:99:0,114,
0:3,0:3:99:0,100, 0:3,0:3:99:0,100, 0:15,0:15:99:0,245, 1, 0.009091, 110,
Missing[NotAvailable], 452, 0., 1, 0.009091, 60., Missing[NotAvailable],
31.78, Missing[NotAvailable], 1.096, Missing[NotAvailable]}
```

```
In[*]:= fullgenomeSNPEFF[fixme] = {"chrX", 10, 37037, "C", "T", 1239.29`, "GT:AD:DP:GQ:PL",
        "PEX2", "MODERATE", "missense_variant", "YJL210W", "c.119C>T", "p.Ala40Val",
        "T|missense_variant|MODERATE|YJL211C|YJL211C|transcript|YJL211C_mRNA|
          protein coding|1/1|c.164G>A|p.Cys55Tyr|164/444|164/444|55/147||,T|
          missense variant|MODERATE|PEX2|YJL210W|transcript|YJL210W mRNA|
          protein_coding|1/1|c.119C>T|p.Ala40Val|119/816|119/816|40/271||,T|
          upstream gene variant|MODIFIER|OPT1|YJL212C|transcript|YJL212C mRNA|
          protein_coding||c.-788G>A|||||788|,T|upstream_gene_variant|MODIFIER|CBP1|
          YJL209W|transcript|YJL209W_mRNA|protein_coding||c.-968C>T|||||968|,T|
          downstream gene variant|MODIFIER|YJL213W|YJL213W|transcript|YJL213W mRNA|
          protein_coding||c.*3879C>T|||||3879|,T|downstream_gene_variant|MODIFIER|
          NUC1|YJL208C|transcript|YJL208C_mRNA|protein_coding||c.*3157G>A|||||3157|,
          T|downstream gene variant|MODIFIER|LAA1|YJL207C|transcript|YJL207C mRNA|
          protein_coding||c.*4352G>A|||||4352|", "0:3,0:3:99:0,100",
        "0:8,0:8:99:0,129", "0:7,0:7:99:0,189", "0:3,0:3:99:0,109", "0:4,0:4:99:0,118",
        "0:3,0:3:99:0,111", "0:4,0:4:99:0,135", "0:3,0:3:99:0,100", "0:4,0:4:99:0,135",
        "0:3,0:3:99:0,109", "0:3,0:3:99:0,100", "0:4,0:4:99:0,119", "0:3,0:3:99:0,100",
        "0:4,0:4:99:0,135", "0:3,0:3:99:0,110", "0:3,0:3:99:0,100", "0:6,0:6:99:0,116",
        "0:3,0:3:99:0,100", "0:3,0:3:99:0,100", "0:3,0:3:99:0,104", "0:3,0:3:99:0,109",
        "0:4,0:4:99:0,119", "0:3,0:3:99:0,103", "0:4,0:4:99:0,114", "0:4,0:4:99:0,115",
        "0:4,0:4:99:0,119", "0:4,0:4:99:0,110", "0:4,0:4:99:0,141", "0:3,0:3:99:0,113",
        "0:3,0:3:99:0,100", "0:4,0:4:99:0,107", "0:3,0:3:99:0,100", "0:4,0:4:99:0,110",
        "0:4,0:4:99:0,130", "0:3,0:3:99:0,122", "0:3,0:3:99:0,100", "0:4,0:4:99:0,100",
        "0:4,0:4:99:0,115", "0:4,0:4:99:0,137", "0:3,0:3:99:0,109", "0:3,0:3:99:0,100",
        "0:3,0:3:99:0,105", "0:3,0:3:99:0,100", "0:4,0:4:99:0,151", "0:3,0:3:99:0,110",
        "0:3,0:3:99:0,104", "0:8,0:8:99:0,176", "0:6,0:6:99:0,119", "0:3,0:3:99:0,100",
        "0:3,0:3:99:0,100", "0:3,0:3:99:0,113", "0:3,0:3:99:0,100", "0:3,0:3:99:0,100",
        "0:4,0:4:99:0,100", "0:3,0:3:99:0,100", "0:3,0:3:99:0,103", "0:3,0:3:99:0,118",
        "0:3,0:3:99:0,100", "0:3,0:3:99:0,107", "0:3,0:3:99:0,107", "0:3,0:3:99:0,100",
        "0:7,0:7:99:0,227", "0:3,0:3:99:0,109", "0:3,0:3:99:0,113", "0:3,0:3:99:0,100",
        "0:4,0:4:99:0,139", "0:4,0:4:99:0,115", "0:4,0:4:99:0,114",
        "1:0,39:39:99:1260,0", "0:5,0:5:99:0,132", "0:4,0:4:99:0,133",
        "0:3,0:3:99:0,109", "0:4,0:4:99:0,110", "0:4,0:4:99:0,102", "0:3,0:3:99:0,100",
        "0:3,0:3:99:0,103", "0:5,0:5:99:0,135", "0:5,0:5:99:0,174", "0:3,0:3:99:0,100",
        "0:3,0:3:99:0,100", "0:3,0:3:99:0,107", "0:4,0:4:99:0,135", "0:3,0:3:99:0,104",
        "0:4,0:4:99:0,115", "0:4,0:4:99:0,128", "0:4,0:4:99:0,124", "0:5,0:5:99:0,181",
        "0:3,0:3:99:0,100", "0:4,0:4:99:0,123", "0:3,0:3:99:0,114", "0:4,0:4:99:0,132",
        "0:3,0:3:99:0,101", "0:3,0:3:99:0,100", "0:3,0:3:99:0,109", "0:3,0:3:99:0,100",
        "0:4,0:4:99:0,127", "0:3,0:3:99:0,117", "0:3,0:3:99:0,118", "0:3,0:3:99:0,100",
        "0:4,0:4:99:0,111", "0:4,0:4:99:0,114", "0:4,0:4:99:0,114", "0:3,0:3:99:0,100",
        "0:5,0:5:99:0,113", "0:4,0:4:99:0,135", "0:4,0:4:99:0,122", "0:3,0:3:99:0,114",
        "0:3,0:3:99:0,100", "0:3,0:3:99:0,100", "0:15,0:15:99:0,245", 1, 0.009091`,
        110, "NA", 452, 0.`, 1, 0.009091`, 60.`, "NA", 31.78`, "NA", 1.096`, "NA"};
```

#229: BUD19 is dubious [HIGH], but RPL39 is verified [LOW] (SGD) [Updated]

```
In[*]:= fixme = 229;
```

Manually swapping for the second gene:

In[*]:= fullgenomeSNPEFF[[fixme]]

Out[•]=

```
{chrX, 10, 76405, G, A, 839.29, GT:AD:DP:GQ:PL,
 BUD19, HIGH, stop_gained, YJL188C, c.106C>T, p.Gln36*,
 A|stop_gained|HIGH|BUD19|YJL188C|transcript|YJL188C_mRNA|protein_coding|1/1|c.106
     C>T|p.Gln36*|106/309|106/309|36/102||,A|synonymous_variant|LOW|RPL39|YJL189W|
     transcript | YJL189W\_mRNA | protein\_coding | 2/2 | c.87G>A | p. Leu29Leu | 87/156 | 87/156 | 29/156 | 29/156 | 29/156 | 29/156 | 29/156 | 29/156 | 29/156 | 29/156 | 29/156 | 29/156 | 29/156 | 29/156 | 29/156 | 29/156 | 29/156 | 29/156 | 29/156 | 29/156 | 29/156 | 29/156 | 29/156 | 29/156 | 29/156 | 29/156 | 29/156 | 29/156 | 29/156 | 29/156 | 29/156 | 29/156 | 29/156 | 29/156 | 29/156 | 29/156 | 29/156 | 29/156 | 29/156 | 29/156 | 29/156 | 29/156 | 29/156 | 29/156 | 29/156 | 29/156 | 29/156 | 29/156 | 29/156 | 29/156 | 29/156 | 29/156 | 29/156 | 29/156 | 29/156 | 29/156 | 29/156 | 29/156 | 29/156 | 29/156 | 29/156 | 29/156 | 29/156 | 29/156 | 29/156 | 29/156 | 29/156 | 29/156 | 29/156 | 29/156 | 29/156 | 29/156 | 29/156 | 29/156 | 29/156 | 29/156 | 29/156 | 29/156 | 29/156 | 29/156 | 29/156 | 29/156 | 29/156 | 29/156 | 29/156 | 29/156 | 29/156 | 29/156 | 29/156 | 29/156 | 29/156 | 29/156 | 29/156 | 29/156 | 29/156 | 29/156 | 29/156 | 29/156 | 29/156 | 29/156 | 29/156 | 29/156 | 29/156 | 29/156 | 29/156 | 29/156 | 29/156 | 29/156 | 29/156 | 29/156 | 29/156 | 29/156 | 29/156 | 29/156 | 29/156 | 29/156 | 29/156 | 29/156 | 29/156 | 29/156 | 29/156 | 29/156 | 29/156 | 29/156 | 29/156 | 29/156 | 29/156 | 29/156 | 29/156 | 29/156 | 29/156 | 29/156 | 29/156 | 29/156 | 29/156 | 29/156 | 29/156 | 29/156 | 29/156 | 29/156 | 29/156 | 29/156 | 29/156 | 29/156 | 29/156 | 29/156 | 29/156 | 29/156 | 29/156 | 29/156 | 29/156 | 29/156 | 29/156 | 29/156 | 29/156 | 29/156 | 29/156 | 29/156 | 29/156 | 29/156 | 29/156 | 29/156 | 29/156 | 29/156 | 29/156 | 29/156 | 29/156 | 29/156 | 29/156 | 29/156 | 29/156 | 29/156 | 29/156 | 29/156 | 29/156 | 29/156 | 29/156 | 29/156 | 29/156 | 29/156 | 29/156 | 29/156 | 29/156 | 29/156 | 29/156 | 29/156 | 29/156 | 29/156 | 29/156 | 29/156 | 29/156 | 29/156 | 29/156 | 29/156 | 29/156 | 29/156 | 29/156 | 29/156 | 29/156 | 29/156 | 29/156 | 29/156 | 29/156 | 29/156 | 29/156 | 29/156 | 29/156 | 29/156 | 29/156 | 29/156 | 29/156 | 29/156 | 29/156 | 29/156 | 29/156 | 29/156 | 29/156 | 29/156 | 29/156 
     /51||,A|upstream_gene_variant|MODIFIER|SOP4|YJL192C|transcript|YJL192C_mRNA|
     protein_coding||c.-2991C>T|||||2991|,A|upstream_gene_variant|MODIFIER|RPS22A|
     YJL190C|transcript|YJL190C_mRNA|protein_coding||c.-1103C>T||||1103|,A|
     upstream_gene_variant|MODIFIER|MNN5|YJL186W|transcript|YJL186W_mRNA|
     protein_coding||c.-3750G>A|||||3750|,A|downstream_gene_variant|MODIFIER|
     YJL193W|YJL193W|transcript|YJL193W_mRNA|protein_coding||c.*3831G>A|||||3831|,A
     |downstream_gene_variant|MODIFIER|RPS14B|YJL191W|transcript|YJL191W_mRNA|
     protein coding||c.*1794G>A|||||1794|,A|downstream gene variant|MODIFIER|SWE1|
     YJL187C|transcript|YJL187C_mRNA|protein_coding||c.*398C>T|||||398|,
 0:11,0:11:99:0,164, 0:9,0:9:99:0,270, 0:10,0:10:99:0,200, 0:8,0:8:99:0,103,
 0:3,0:3:99:0,118, 0:12,0:12:99:0,271, 0:4,0:4:99:0,135, 0:7,0:7:99:0,155,
 0:7,0:7:99:0,111, 0:14,0:14:99:0,458, 0:8,0:8:99:0,225, 0:6,0:6:99:0,139,
 0:7,0:7:99:0,152, 0:9,0:9:99:0,224, 0:3,0:3:99:0,110, 0:8,0:8:99:0,227,
 0:5,0:5:99:0,159, 0:6,0:6:99:0,153, 0:7,0:7:99:0,146, 0:11,0:11:99:0,277,
 0:7,0:7:99:0,223, 0:9,0:9:99:0,197, 0:3,0:3:99:0,118, 0:13,0:13:99:0,402,
 0:13,0:13:99:0,264, 0:13,0:13:99:0,107, 0:7,0:7:99:0,121, 0:10,0:10:99:0,165,
 0:8,0:8:99:0,217, 0:9,0:9:99:0,315, 0:7,0:7:99:0,135, 0:20,0:20:99:0,551,
 0:14,0:14:99:0,262, 0:12,0:12:99:0,209, 0:7,0:7:99:0,148, 0:8,0:8:99:0,140,
 0:3,0:3:99:0,100, 0:5,0:5:99:0,135, 0:4,0:4:99:0,116, 0:8,0:8:99:0,253,
 0:3,0:3:99:0,100, 0:11,0:11:99:0,314, 0:10,0:10:99:0,240, 0:4,0:4:99:0,103,
 0:5,0:5:99:0,159, 0:11,0:11:99:0,150, 0:3,0:3:99:0,118, 0:7,0:7:99:0,136,
 0:8,0:8:99:0,129, 0:9,0:9:99:0,180, 0:5,0:5:99:0,105, 0:7,0:7:99:0,127,
 0:3,0:3:99:0,109, 0:4,0:4:99:0,133, 0:5,0:5:99:0,115, 0:10,0:10:99:0,101,
 0:3,0:3:99:0,117, 0:3,0:3:99:0,100, 0:4,0:4:99:0,140, 0:3,0:3:99:0,107,
 0:6,0:6:99:0,218, 0:6,0:6:99:0,127, 0:8,0:8:99:0,169, 0:10,0:10:99:0,311,
 0:5,0:5:99:0,159, 0:5,0:5:99:0,118, 0:7,0:7:99:0,204, 0:9,0:9:99:0,250,
 0:15,0:15:99:0,182,0:7,0:7:99:0,172,0:13,0:13:99:0,197,0:7,0:7:99:0,198,
 0:9,0:9:99:0,177, 0:9,0:9:99:0,157, 0:4,0:4:99:0,127, 0:6,0:6:99:0,113,
 0:6,0:6:99:0,135, 0:20,0:20:99:0,348, 0:6,0:6:99:0,135, 0:5,0:5:99:0,173,
 0:12,0:12:99:0,338, 0:10,0:10:99:0,180, 0:15,0:15:99:0,143,
 0:12,0:12:99:0,355,0:4,0:4:99:0,128,0:14,0:14:99:0,174,0:12,0:12:99:0,139,
 0:14,0:14:99:0,157, 0:12,0:12:99:0,339, 0:9,0:9:99:0,124, 0:8,0:8:99:0,208,
```

```
0:5,0:5:99:0,102, 0:8,0:8:99:0,120, 0:5,0:5:99:0,135, 0:7,0:7:99:0,118,
      0:10,0:10:99:0,136, 0:12,0:12:99:0,194, 0:5,0:5:99:0,135, 0:9,0:9:99:0,178,
      0:8,0:8:99:0,245, 0:3,0:3:99:0,118, 0:4,0:4:99:0,122, 0:11,0:11:99:0,320,
      0:11,0:11:99:0,200, 0:6,0:6:99:0,124, 0:6,0:6:99:0,185, 1, 0.009091, 110,
      Missing[NotAvailable], 890, 0., 1, 0.009091, 60., Missing[NotAvailable],
      34.97, Missing[NotAvailable], 1.519, Missing[NotAvailable]}
In[*]:= fullgenomeSNPEFF[fixme] = {"chrX", 10, 76405, "G", "A", 839.29`, "GT:AD:DP:GQ:PL",
        "RPL39", "LOW", "synonymous_variant", "YJL189W", "c.87G>A", "p.Leu29Leu",
        "A|stop_gained|HIGH|BUD19|YJL188C|transcript|YJL188C_mRNA|protein_coding|1/1|c
          .106C>T|p.Gln36*|106/309|106/309|36/102||,A|synonymous_variant|LOW|RPL39|
          YJL189W|transcript|YJL189W_mRNA|protein_coding|2/2|c.87G>A|p.Leu29Leu|87/
          156|87/156|29/51||,A|upstream_gene_variant|MODIFIER|SOP4|YJL192C|
          transcript|YJL192C_mRNA|protein_coding||c.-2991C>T|||||2991|,A|
          upstream_gene_variant|MODIFIER|RPS22A|YJL190C|transcript|YJL190C_mRNA|
          protein_coding||c.-1103C>T|||||1103|,A|upstream_gene_variant|MODIFIER|MNN5
          |YJL186W|transcript|YJL186W_mRNA|protein_coding||c.-3750G>A||||3750|,A|
          downstream_gene_variant|MODIFIER|YJL193W|YJL193W|transcript|YJL193W_mRNA|
          protein coding||c.*3831G>A|||||3831|,A|downstream gene variant|MODIFIER|
          RPS14B|YJL191W|transcript|YJL191W_mRNA|protein_coding||c.*1794G>A|||||1794
          |,A|downstream_gene_variant|MODIFIER|SWE1|YJL187C|transcript|YJL187C_mRNA|
          protein coding||c.*398C>T|||||398|", "0:11,0:11:99:0,164",
        "0:9,0:9:99:0,270", "0:10,0:10:99:0,200", "0:8,0:8:99:0,103",
        "0:3,0:3:99:0,118", "0:12,0:12:99:0,271", "0:4,0:4:99:0,135",
        "0:7,0:7:99:0,155", "0:7,0:7:99:0,111", "0:14,0:14:99:0,458",
        "0:8,0:8:99:0,225", "0:6,0:6:99:0,139", "0:7,0:7:99:0,152", "0:9,0:9:99:0,224",
        "0:3,0:3:99:0,110", "0:8,0:8:99:0,227", "0:5,0:5:99:0,159", "0:6,0:6:99:0,153",
        "0:7,0:7:99:0,146", "0:11,0:11:99:0,277", "0:7,0:7:99:0,223",
        "0:9,0:9:99:0,197", "0:3,0:3:99:0,118", "0:13,0:13:99:0,402",
        "0:13,0:13:99:0,264", "0:13,0:13:99:0,107", "0:7,0:7:99:0,121",
        "0:10,0:10:99:0,165", "0:8,0:8:99:0,217", "0:9,0:9:99:0,315",
        "0:7,0:7:99:0,135", "0:20,0:20:99:0,551", "0:14,0:14:99:0,262",
        "0:12,0:12:99:0,209", "0:7,0:7:99:0,148", "0:8,0:8:99:0,140",
        "0:3,0:3:99:0,100", "0:5,0:5:99:0,135", "0:4,0:4:99:0,116", "0:8,0:8:99:0,253",
        "0:3,0:3:99:0,100", "0:11,0:11:99:0,314", "0:10,0:10:99:0,240",
        "0:4,0:4:99:0,103", "0:5,0:5:99:0,159", "0:11,0:11:99:0,150",
        "0:3,0:3:99:0,118", "0:7,0:7:99:0,136", "0:8,0:8:99:0,129", "0:9,0:9:99:0,180",
        "0:5,0:5:99:0,105", "0:7,0:7:99:0,127", "0:3,0:3:99:0,109", "0:4,0:4:99:0,133",
        "0:5,0:5:99:0,115", "0:10,0:10:99:0,101", "0:3,0:3:99:0,117",
        "0:3,0:3:99:0,100", "0:4,0:4:99:0,140", "0:3,0:3:99:0,107", "0:6,0:6:99:0,218",
        "0:6,0:6:99:0,127", "0:8,0:8:99:0,169", "0:10,0:10:99:0,311",
        "0:5,0:5:99:0,159", "0:5,0:5:99:0,118", "0:7,0:7:99:0,204",
        "0:9,0:9:99:0,250", "0:15,0:15:99:0,182", "0:7,0:7:99:0,172",
```

0:3,0:3:99:0,100, 0:7,0:7:99:0,184, 1:0,24:24:99:860,0, 0:8,0:8:99:0,196,

```
"0:13,0:13:99:0,197", "0:7,0:7:99:0,198", "0:9,0:9:99:0,177",
"0:9,0:9:99:0,157", "0:4,0:4:99:0,127", "0:6,0:6:99:0,113",
"0:6,0:6:99:0,135", "0:20,0:20:99:0,348", "0:6,0:6:99:0,135",
"0:5,0:5:99:0,173", "0:12,0:12:99:0,338", "0:10,0:10:99:0,180",
"0:15,0:15:99:0,143", "0:12,0:12:99:0,355", "0:4,0:4:99:0,128",
"0:14,0:14:99:0,174", "0:12,0:12:99:0,139", "0:14,0:14:99:0,157",
"0:12,0:12:99:0,339", "0:9,0:9:99:0,124", "0:8,0:8:99:0,208",
"0:3,0:3:99:0,100", "0:7,0:7:99:0,184", "1:0,24:24:99:860,0",
"0:8,0:8:99:0,196", "0:5,0:5:99:0,102", "0:8,0:8:99:0,120", "0:5,0:5:99:0,135",
"0:7,0:7:99:0,118", "0:10,0:10:99:0,136", "0:12,0:12:99:0,194",
"0:5,0:5:99:0,135", "0:9,0:9:99:0,178", "0:8,0:8:99:0,245",
"0:3,0:3:99:0,118", "0:4,0:4:99:0,122", "0:11,0:11:99:0,320",
"0:11,0:11:99:0,200", "0:6,0:6:99:0,124", "0:6,0:6:99:0,185", 1, 0.009091`,
110, "NA", 890, 0.`, 1, 0.009091`, 60.`, "NA", 34.97`, "NA", 1.519`, "NA"};
```

#296: VPS63 is uncharacterized, but YPT6 is verified (SGD) [Updated]

```
In[*]:= fixme = 296;
```

Manually swapping for the second gene:

```
In[*]:= fullgenomeSNPEFF[[fixme]]
```

```
Out[ • 1=
```

```
{chrXII, 12, 668352, C, G, 549.29, GT:AD:DP:GQ:PL, VPS63,
MODERATE, missense_variant, YLR261C, c.212G>C, p.Ser71Thr,
G|missense_variant|MODERATE|VPS63|YLR261C|transcript|YLR261C_mRNA|protein_coding|
   1/1|c.212G>C|p.Ser71Thr|212/327|212/327|71/108||,G|missense_variant|MODERATE|
  YPT6|YLR262C|transcript|YLR262C_mRNA|protein_coding|1/1|c.540G>C|p.Glu180Asp|
  540/648|540/648|180/215||,G|upstream_gene_variant|MODIFIER|HSP60|YLR259C|
  transcript|YLR259C_mRNA|protein_coding||c.-3350G>C|||||3350|,G|upstream_gene
   _variant|MODIFIER|RED1|YLR263W|transcript|YLR263W_mRNA|protein_coding||c.-1988
  C>G|||||1988|,G|upstream_gene_variant|MODIFIER|RPS28B|YLR264W|transcript|
  YLR264W_mRNA|protein_coding||c.-4779C>G|||||4779|,G|downstream_gene_variant|
  445|,G|downstream_gene_variant|MODIFIER|TMA7|YLR262C-A|transcript|YLR262C-
  A_mRNA|protein_coding||c.*1116G>C|||||1116|, 0:15,0:15:99:0,486,
0:33,0:33:99:0,1029, 0:20,0:20:99:0,656, 0:18,0:18:99:0,643,
0:9,0:9:99:0,253, 0:20,0:20:99:0,718, 0:19,0:19:99:0,660, 0:19,0:19:99:0,575,
0:7,0:7:99:0,217, 0:10,0:10:99:0,341, 0:11,0:11:99:0,278, 0:27,0:27:99:0,895,
0:14,0:14:99:0,489, 0:14,0:14:99:0,405, 0:17,0:17:99:0,519, 0:14,0:14:99:0,468,
0:12,0:12:99:0,423, 1:0,15:15:99:570,0, 0:9,0:9:99:0,270, 0:19,0:19:99:0,570,
0:17,0:17:99:0,572,0:11,0:11:99:0,360,0:14,0:14:99:0,436,0:16,0:16:99:0,530,
0:11,0:11:99:0,365, 0:24,0:24:99:0,847, 0:18,0:18:99:0,657, 0:7,0:7:99:0,238,
0:11,0:11:99:0,336, 0:14,0:14:99:0,341, 0:21,0:21:99:0,765, 0:22,0:22:99:0,790,
0:28,0:28:99:0,900,0:10,0:10:99:0,331,0:24,0:24:99:0,822,0:9,0:9:99:0,317,
0:3,0:3:99:0,100, 0:5,0:5:99:0,128, 0:8,0:8:99:0,239, 0:10,0:10:99:0,364,
0:11,0:11:99:0,292, 0:21,0:21:99:0,635, 0:19,0:19:99:0,558,
0:8,0:8:99:0,278, 0:11,0:11:99:0,403, 0:14,0:14:99:0,428, 0:18,0:18:99:0,628,
0:12,0:12:99:0,379, 0:13,0:13:99:0,474, 0:14,0:14:99:0,473, 0:13,0:13:99:0,423,
0:17,0:17:99:0,595, 0:4,0:4:99:0,129, 0:7,0:7:99:0,150, 0:8,0:8:99:0,274,
0:12,0:12:99:0,422, 0:19,0:19:99:0,619, 0:12,0:12:99:0,377, 0:14,0:14:99:0,341,
0:18,0:18:99:0,605, 0:8,0:8:99:0,277, 0:12,0:12:99:0,372, 0:19,0:19:99:0,592,
0:15,0:15:99:0,557, 0:12,0:12:99:0,437, 0:21,0:21:99:0,660, 0:17,0:17:99:0,560,
0:15,0:15:99:0,487, 0:28,0:28:99:0,905, 0:14,0:14:99:0,409, 0:20,0:20:99:0,685,
0:8,0:8:99:0,287, 0:19,0:19:99:0,698, 0:17,0:17:99:0,561, 0:24,0:24:99:0,811,
0:17,0:17:99:0,632, 0:10,0:10:99:0,370, 0:17,0:17:99:0,555, 0:20,0:20:99:0,668,
0:19,0:19:99:0,611, 0:24,0:24:99:0,812, 0:17,0:17:99:0,597, 0:20,0:20:99:0,706,
0:17,0:17:99:0,596, 0:15,0:15:99:0,505, 0:26,0:26:99:0,943, 0:16,0:16:99:0,539,
0:15,0:15:99:0,548, 0:16,0:16:99:0,534, 0:16,0:16:99:0,459, 0:18,0:18:99:0,602,
0:14,0:14:99:0,324,0:13,0:13:99:0,440,0:13,0:13:99:0,434,0:13,0:13:99:0,450,
0:11,0:11:99:0,365, 0:23,0:23:99:0,801, 0:13,0:13:99:0,413, 0:19,0:19:99:0,574,
0:13,0:13:99:0,393, 0:18,0:18:99:0,627, 0:9,0:9:99:0,281, 0:17,0:17:99:0,585,
0:15,0:15:99:0,540, 0:12,0:12:99:0,315, 0:18,0:18:99:0,563, 0:15,0:15:99:0,512,
0:15,0:15:99:0,563, 0:8,0:8:99:0,229, 0:7,0:7:99:0,237, 1, 0.009091, 110,
Missing[NotAvailable], 1668, 0., 1, 0.009091, 60., Missing[NotAvailable],
33.61, Missing[NotAvailable], 0.818, Missing[NotAvailable]}
```

In[*]:* fullgenomeSNPEFF[fixme] = {"chrXII", 12, 668352, "C", "G", 549.29`, "GT:AD:DP:GQ:PL", "YPT6", "MODERATE", "missense_variant", "YLR261C", "c.540G>C", "p.Glu180Asp",

```
"G|missense_variant|MODERATE|VPS63|YLR261C|transcript|YLR261C_mRNA|
  protein_coding|1/1|c.212G>C|p.Ser71Thr|212/327|212/327|71/108||,G|
  missense_variant|MODERATE|YPT6|YLR262C|transcript|YLR262C_mRNA|
  protein_coding|1/1|c.540G>C|p.Glu180Asp|540/648|540/648|180/215||,G|
  upstream gene variant|MODIFIER|HSP60|YLR259C|transcript|YLR259C mRNA|
  protein_coding||c.-3350G>C|||||3350|,G|upstream_gene_variant|MODIFIER|RED1
  |YLR263W|transcript|YLR263W_mRNA|protein_coding||c.-1988C>G|||||1988|,G|
  upstream gene variant|MODIFIER|RPS28B|YLR264W|transcript|YLR264W mRNA|
  protein_coding||c.-4779C>G|||||4779|,G|downstream_gene_variant|MODIFIER|
  LCB5|YLR260W|transcript|YLR260W_mRNA|protein_coding||c.*445C>G|||||445|,G|
  downstream_gene_variant|MODIFIER|TMA7|YLR262C-A|transcript|YLR262C-A mRNA|
  protein_coding||c.*1116G>C|||||1116|",
"0:15,0:15:99:0,486", "0:33,0:33:99:0,1029", "0:20,0:20:99:0,656",
"0:18,0:18:99:0,643", "0:9,0:9:99:0,253", "0:20,0:20:99:0,718",
"0:19,0:19:99:0,660", "0:19,0:19:99:0,575", "0:7,0:7:99:0,217",
"0:10,0:10:99:0,341", "0:11,0:11:99:0,278", "0:27,0:27:99:0,895",
"0:14,0:14:99:0,489", "0:14,0:14:99:0,405", "0:17,0:17:99:0,519",
"0:14,0:14:99:0,468", "0:12,0:12:99:0,423", "1:0,15:15:99:570,0",
"0:9,0:9:99:0,270", "0:19,0:19:99:0,570", "0:17,0:17:99:0,572",
"0:11,0:11:99:0,360", "0:14,0:14:99:0,436", "0:16,0:16:99:0,530",
"0:11,0:11:99:0,365", "0:24,0:24:99:0,847", "0:18,0:18:99:0,657",
"0:7,0:7:99:0,238", "0:11,0:11:99:0,336", "0:14,0:14:99:0,341",
"0:21,0:21:99:0,765", "0:22,0:22:99:0,790", "0:28,0:28:99:0,900",
"0:10,0:10:99:0,331", "0:24,0:24:99:0,822", "0:9,0:9:99:0,317",
"0:3,0:3:99:0,100", "0:5,0:5:99:0,128", "0:8,0:8:99:0,239",
"0:10,0:10:99:0,364", "0:11,0:11:99:0,292", "0:21,0:21:99:0,635",
"0:19,0:19:99:0,558", "0:8,0:8:99:0,278", "0:11,0:11:99:0,403",
"0:14,0:14:99:0,428", "0:18,0:18:99:0,628", "0:12,0:12:99:0,379",
"0:13,0:13:99:0,474", "0:14,0:14:99:0,473", "0:13,0:13:99:0,423",
"0:17,0:17:99:0,595", "0:4,0:4:99:0,129", "0:7,0:7:99:0,150",
"0:8,0:8:99:0,274", "0:12,0:12:99:0,422", "0:19,0:19:99:0,619",
"0:12,0:12:99:0,377", "0:14,0:14:99:0,341", "0:18,0:18:99:0,605",
"0:8,0:8:99:0,277", "0:12,0:12:99:0,372", "0:19,0:19:99:0,592",
"0:15,0:15:99:0,557", "0:12,0:12:99:0,437", "0:21,0:21:99:0,660",
"0:17,0:17:99:0,560", "0:15,0:15:99:0,487", "0:28,0:28:99:0,905",
"0:14,0:14:99:0,409", "0:20,0:20:99:0,685", "0:8,0:8:99:0,287",
"0:19,0:19:99:0,698", "0:17,0:17:99:0,561", "0:24,0:24:99:0,811",
"0:17,0:17:99:0,632", "0:10,0:10:99:0,370", "0:17,0:17:99:0,555",
"0:20,0:20:99:0,668", "0:19,0:19:99:0,611", "0:24,0:24:99:0,812",
"0:17,0:17:99:0,597", "0:20,0:20:99:0,706", "0:17,0:17:99:0,596",
"0:15,0:15:99:0,505", "0:26,0:26:99:0,943", "0:16,0:16:99:0,539",
"0:15,0:15:99:0,548", "0:16,0:16:99:0,534", "0:16,0:16:99:0,459",
"0:18,0:18:99:0,602", "0:14,0:14:99:0,324", "0:13,0:13:99:0,440",
"0:13,0:13:99:0,434", "0:13,0:13:99:0,450", "0:11,0:11:99:0,365",
```

```
"0:23,0:23:99:0,801", "0:13,0:13:99:0,413", "0:19,0:19:99:0,574",
"0:13,0:13:99:0,393", "0:18,0:18:99:0,627", "0:9,0:9:99:0,281",
"0:17,0:17:99:0,585", "0:15,0:15:99:0,540", "0:12,0:12:99:0,315",
"0:18,0:18:99:0,563", "0:15,0:15:99:0,512", "0:15,0:15:99:0,563",
"0:8,0:8:99:0,229", "0:7,0:7:99:0,237", 1, 0.009091`, 110, "NA",
1668, 0.', 1, 0.009091', 60.', "NA", 33.61', "NA", 0.818', "NA"};
```

#317: VRP1 is verified, but OPI9 is dubious (SGD) [Kept as is]

```
In[*]:= fixme = 317;
      Kept as is:
```

In[*]:= fullgenomeSNPEFF[[fixme]]

Out[•]=

```
{chrXII, 12, 804694, G, A, 355.29, GT:AD:DP:GQ:PL, VRP1,
MODERATE, missense_variant, YLR337C, c.413C>T, p.Ala138Val,
A | missense_variant | MODERATE | VRP1 | YLR337C | transcript | YLR337C mRNA | protein_coding | 1
   /1|c.413C>T|p.Ala138Val|413/2454|413/2454|138/817||,A|missense_variant|
   MODERATE | OPI9 | YLR338W | transcript | YLR338W_mRNA | protein_coding | 1/1 | c.349G>A | p.
   Ala117Thr|349/858|349/858|117/285||,A|upstream_gene_variant|MODIFIER|SGD1|
   YLR336C|transcript|YLR336C_mRNA|protein_coding||c.-2298C>T|||||2298|,A|
   upstream_gene_variant|MODIFIER|RPP0|YLR340W|transcript|YLR340W_mRNA|
   protein_coding||c.-1193G>A|||||1193|,A|upstream_gene_variant|MODIFIER|SP077|
   YLR341W|transcript|YLR341W_mRNA|protein_coding||c.-2691G>A|||||2691|,A|
   downstream_gene_variant|MODIFIER|YLR339C|YLR339C|transcript|YLR339C_mRNA|
   protein_coding||c.*1090C>T|||||1090|, 0:10,0:10:99:0,224,
0:18,0:18:99:0,224, 0:7,0:7:99:0,228, 0:8,0:8:99:0,201, 0:5,0:5:99:0,118,
0:13,0:13:99:0,378,0:10,0:10:99:0,180,0:11,0:11:99:0,244,0:7,0:7:99:0,228,
0:5,0:5:99:0,150, 0:5,0:5:99:0,135, 0:17,0:17:99:0,371, 0:4,0:4:99:0,128,
0:7,0:7:99:0,187, 0:8,0:8:99:0,246, 0:11,0:11:99:0,295, 0:5,0:5:99:0,115,
0:5,0:5:99:0,122,0:5,0:5:99:0,132,0:9,0:9:99:0,251,0:7,0:7:99:0,159,
0:5,0:5:99:0,180,0:11,0:11:99:0,338,0:13,0:13:99:0,357,0:3,0:3:99:0,107,
0:14,0:14:99:0,407, 0:11,0:11:99:0,306, 0:7,0:7:99:0,217, 0:4,0:4:99:0,99,
0:6,0:6:99:0,135, 0:14,0:14:99:0,407, 0:12,0:12:99:0,245, 0:16,0:16:99:0,568,
0:6,0:6:99:0,148,0:12,0:12:99:0,346,0:5,0:5:99:0,140,0:3,0:3:99:0,104,
0:5,0:5:99:0,122, 0:4,0:4:99:0,135, 0:4,0:4:99:0,113, 0:8,0:8:99:0,214,
0:12,0:12:99:0,392,0:6,0:6:99:0,218,0:8,0:8:99:0,161,0:4,0:4:99:0,119,
0:10,0:10:99:0,317,0:4,0:4:99:0,122,0:8,0:8:99:0,192,0:10,0:10:99:0,262,
0:10,0:10:99:0,197, 0:4,0:4:99:0,122, 0:6,0:6:99:0,204, 0:4,0:4:99:0,99,
0:4,0:4:99:0,154, 0:5,0:5:99:0,135, 0:8,0:8:99:0,162, 0:8,0:8:99:0,173,
0:7,0:7:99:0,191, 0:8,0:8:99:0,250, 0:6,0:6:99:0,190, 0:7,0:7:99:0,191,
0:7,0:7:99:0,228, 0:4,0:4:99:0,135, 0:10,0:10:99:0,263, 1:0,10:10:99:376,0,
0:9,0:9:99:0,225, 0:13,0:13:99:0,430, 0:9,0:9:99:0,336, 0:18,0:18:99:0,585,
0:11,0:11:99:0,283, 0:11,0:11:99:0,203, 0:8,0:8:99:0,103, 0:9,0:9:99:0,184,
0:9,0:9:99:0,240, 0:11,0:11:99:0,302, 0:10,0:10:99:0,351, 0:8,0:8:99:0,279,
0:16,0:16:99:0,521,0:12,0:12:99:0,345,0:8.99:0,241,0:14.0:14:99:0,509,
0:8,0:8:99:0,273, 0:12,0:12:99:0,435, 0:13,0:13:99:0,356, 0:7,0:7:99:0,209,
0:11,0:11:99:0,393, 0:7,0:7:99:0,225, 0:8,0:8:99:0,264, 0:9,0:9:99:0,232,
0:7,0:7:99:0,225, 0:9,0:9:99:0,296, 0:4,0:4:99:0,133, 0:7,0:7:99:0,219,
0:9,0:9:99:0,300, 0:8,0:8:99:0,221, 0:7,0:7:99:0,196, 0:10,0:10:99:0,276,
0:7,0:7:99:0,184, 0:12,0:12:99:0,358, 0:6,0:6:99:0,214, 0:10,0:10:99:0,316,
0:9,0:9:99:0,192,0:6,0:6:99:0,164,0:12,0:12:99:0,259,0:5,0:5:99:0,113,
0:9,0:9:99:0,162, 0:7,0:7:99:0,247, 0:9,0:9:99:0,210, 0:5,0:5:99:0,128,
0:4,0:4:99:0,107, 1, 0.009091, 110, Missing[NotAvailable], 924, 0., 1, 0.009091, 60.,
Missing[NotAvailable], 29.03, Missing[NotAvailable], 1.609, Missing[NotAvailable]}
```

#356: DDR48 is verified, but YMR173W-A is dubious (SGD) [Kept as is]

```
In[*]:= fixme = 356;
      Kept as is:
 In[*]:= fullgenomeSNPEFF[[fixme]]
Out[ • ]=
      {chrXIII, 13, 609388, A, G, 342.29, GT:AD:DP:GQ:PL, DDR48,
       MODERATE, missense_variant, YMR173W, c.700A>G, p.Asn234Asp,
       G|missense_variant|MODERATE|DDR48|YMR173W|transcript|YMR173W_mRNA|protein_coding|
         1/1|c.700A>G|p.Asn234Asp|700/1293|700/1293|234/430||,G|synonymous_variant|LOW|
         YMR173W-A|YMR173W-A|transcript|YMR173W-A_mRNA|protein\_coding|1/1|c.492A>G|p.
         Leu164Leu|492/1185|492/1185|164/394||,G|upstream_gene_variant|MODIFIER|EAR1|
         YMR171C|transcript|YMR171C_mRNA|protein_coding||c.-3868T>C|||||3868|,G|
         upstream_gene_variant|MODIFIER|YMR172C-A|YMR172C-A|transcript|YMR172C-A_mRNA|
         protein_coding||c.-1177T>C|||||1177|,G|upstream_gene_variant|MODIFIER|SIP18|
         YMR175W|transcript|YMR175W_mRNA|protein_coding||c.-1628A>G|||||1628|,G|
         upstream_gene_variant|MODIFIER|YMR175W-A|YMR175W-A|transcript|YMR175W-A_mRNA|
         protein_coding||c.-1926A>G|||||1926|,G|upstream_gene_variant|MODIFIER|ECM5|
         YMR176W|transcript|YMR176W_mRNA|protein_coding||c.-2352A>G|||||2352|,G|
         downstream_gene_variant|MODIFIER|HOT1|YMR172W|transcript|YMR172W_mRNA|
         protein_coding||c.*1248A>G|||||1248|,G|downstream_gene_variant|MODIFIER|PAI3|
         YMR174C|transcript|YMR174C_mRNA|protein_coding||c.*771T>C|||||771|,
       0:7,0:7:99:0,159, 0:11,0:11:99:0,111, 0:8,0:8:99:0,170, 0:7,0:7:99:0,246,
       0:5,0:5:99:0,103, 0:9,0:9:99:0,149, 0:9,0:9:99:0,168, 0:10,0:10:99:0,264,
       0:4,0:4:99:0,111, 0:6,0:6:99:0,104, 1:0,11:11:99:363,0, 0:6,0:6:99:0,142,
       0:4,0:4:99:0,128, 0:5,0:5:99:0,131, 0:5,0:5:99:0,163, 0:9,0:9:99:0,302,
       0:7,0:7:99:0,124, 0:7,0:7:99:0,111, 0:5,0:5:99:0,103, 0:8,0:8:99:0,202,
       0:5,0:5:99:0,155, 0:6,0:6:99:0,144, 0:11,0:11:99:0,116, 0:10,0:10:99:0,185,
       0:5,0:5:99:0,163, 0:9,0:9:99:0,251, 0:22,0:22:99:0,313, 0:5,0:5:99:0,140,
       0:3,0:3:99:0,105, 0:6,0:6:99:0,136, 0:12,0:12:99:0,208, 0:13,0:13:99:0,190,
       0:21,0:21:99:0,596, 0:7,0:7:99:0,123, 0:12,0:12:99:0,188, 0:5,0:5:99:0,166,
       0:4,0:4:99:0,109, 0:5,0:5:99:0,134, 0:4,0:4:99:0,104, 0:5,0:5:99:0,163,
       0:6,0:6:99:0,101,0:10,0:10:99:0,187,0:4,0:4:99:0,101,0:7,0:7:99:0,146,
       0:10,0:10:99:0,136, 0:8,0:8:99:0,270, 0:4,0:4:99:0,135, 0:5,0:5:99:0,125,
       0:7,0:7:99:0,110, 0:7,0:7:99:0,225, 0:5,0:5:99:0,103, 0:6,0:6:99:0,137,
       0:4,0:4:99:0,100, 0:4,0:4:99:0,107, 0:4,0:4:99:0,104, 0:8,0:8:99:0,235,
       0:6,0:6:99:0,159, 0:8,0:8:99:0,203, 0:6,0:6:99:0,139, 0:5,0:5:99:0,155,
       0:4,0:4:99:0,141,0:5,0:5:99:0,111,0:9,0:9:99:0,183,0:7,0:7:99:0,113,
       0:8,0:8:99:0,122, 0:13,0:13:99:0,399, 0:10,0:10:99:0,160, 0:9,0:9:99:0,183,
       0:11,0:11:99:0,342,0:7,0:7:99:0,174,0:9,0:9:99:0,202,0:19,0:19:99:0,297,
       0:19,0:19:99:0,632, 0:20,0:20:99:0,585, 0:10,0:10:99:0,157,
       0:8,0:8:99:0,194, 0:9,0:9:99:0,224, 0:9,0:9:99:0,105, 0:13,0:13:99:0,234,
```

0:8,0:8:99:0,263, 0:14,0:14:99:0,392, 0:19,0:19:99:0,151, 0:12,0:12:99:0,148,

Out[•]=

```
0:10,0:10:99:0,208,0:9,0:9:99:0,126,0:25,0:25:99:0,354,0:8,0:8:99:0,198,
0:7,0:7:99:0,114, 0:7,0:7:99:0,263, 0:6,0:6:99:0,180, 0:10,0:10:99:0,129,
0:3,0:3:99:0,104,0:6,0:6:99:0,218,0:7,0:7:99:0,137,0:7,0:7:99:0,191,
0:8,0:8:99:0,103, 0:11,0:11:99:0,334, 0:21,0:21:99:0,514, 0:12,0:12:99:0,128,
0:11,0:11:99:0,331,0:9,0:9:99:0,154,0:10,0:10:99:0,175,0:9,0:9:99:0,233,
0:25,0:25:99:0,281,0:5,0:5:99:0,141,0:22,0:22:99:0,455,0:7,0:7:99:0,195,
0:11,0:11:99:0,325, 0:3,0:3:99:0,114, 0:13,0:13:99:0,189, 1, 0.009091, 110,
Missing[NotAvailable], 972, 0., 1, 0.009091, 57.68, Missing[NotAvailable],
31.12, Missing[NotAvailable], 1.802, Missing[NotAvailable]}
```

#380: YNL109W is dubious, as is YNL108C (SGD) [To be dropped]

```
In[*]:= fixme = 380;
```

In[*]:= fullgenomeSNPEFF[[fixme]]

```
{chrXIV, 14, 419354, G, A, 446.29, GT:AD:DP:GQ:PL, YNL109W,
MODERATE, missense_variant, YNL109W, c.391G>A, p.Gly131Arg,
A|missense_variant|MODERATE|YNL109W|YNL109W|transcript|YNL109W_mRNA|
   protein_coding|1/1|c.391G>A|p.Gly131Arg|391/546|391/546|131/181||,A|
  missense_variant|MODERATE|YNL108C|YNL108C|transcript|YNL108C_mRNA|
  protein_coding|1/1|c.473C>T|p.Pro158Leu|473/813|473/813|158/270||,A|
  upstream_gene_variant|MODIFIER|CYB5|YNL111C|transcript|YNL111C_mRNA|
   protein_coding||c.-2052C>T|||||2052|,A|upstream_gene_variant|MODIFIER|NOP15|
  YNL110C|transcript|YNL110C_mRNA|protein_coding||c.-866C>T|||||866|,A|
  upstream_gene_variant|MODIFIER|YAF9|YNL107W|transcript|YNL107W_mRNA|
   protein_coding||c.-744G>A|||||744|,A|upstream_gene_variant|MODIFIER|RRT16|
  YNL105W|transcript|YNL105W_mRNA|protein_coding||c.-4801G>A||||4801|,A|
  downstream_gene_variant|MODIFIER|DBP2|YNL112W|transcript|YNL112W_mRNA|
   protein_coding||c.*3073G>A|||||3073|,A|downstream_gene_variant|MODIFIER|INP52|
  YNL106C|transcript|YNL106C_mRNA|protein_coding||c.*1590C>T|||||1590|,
0:10,0:10:99:0,151, 0:7,0:7:99:0,136, 0:5,0:5:99:0,159, 0:11,0:11:99:0,137,
0:4,0:4:99:0,109, 0:7,0:7:99:0,196, 0:8,0:8:99:0,105, 0:8,0:8:99:0,191,
0:5,0:5:99:0,113,0:7,0:7:99:0,99,0:7,0:7:99:0,123,0:8,0:8:99:0,237,
0:8,0:8:99:0,112, 0:7,0:7:99:0,125, 0:8,0:8:99:0,120, 0:6,0:6:99:0,125,
0:4,0:4:99:0,119,0:6,0:6:99:0,200,0:4,0:4:99:0,104,0:7,0:7:99:0,116,
0:6,0:6:99:0,150, 1:0,12:12:99:467,0, 0:13,0:13:99:0,106, 0:10,0:10:99:0,123,
0:12,0:12:99:0,151,0:10,0:10:99:0,130,0:9,0:9:99:0,177,0:7,0:7:99:0,114,
0:4,0:4:99:0,132, 0:9,0:9:99:0,101, 0:14,0:14:99:0,258, 0:12,0:12:99:0,172,
0:21,0:21:99:0,135,0:4,0:4:99:0,126,0:19,0:19:99:0,202,0:5,0:5:99:0,169,
0:4,0:4:99:0,135, 0:6,0:6:99:0,140, 0:4,0:4:99:0,135, 0:8,0:8:99:0,107,
0:13,0:13:99:0,138, 0:10,0:10:99:0,167, 0:7,0:7:99:0,99, 0:6,0:6:99:0,138,
0:6,0:6:99:0,130, 0:10,0:10:99:0,146, 0:7,0:7:99:0,223, 0:10,0:10:99:0,187,
0:9,0:9:99:0,259, 0:7,0:7:99:0,158, 0:7,0:7:99:0,142, 0:4,0:4:99:0,133,
0:4,0:4:99:0,113, 0:5,0:5:99:0,125, 0:4,0:4:99:0,103, 0:7,0:7:99:0,208,
```

```
0:9,0:9:99:0,103,0:3,0:3:99:0,100,0:6,0:6:99:0,138,0:7,0:7:99:0,123,
0:3,0:3:99:0,109, 0:6,0:6:99:0,105, 0:7,0:7:99:0,225, 0:9,0:9:99:0,283,
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0:9,0:9:99:0,166, 0:15,0:15:99:0,204, 0:13,0:13:99:0,184, 0:11,0:11:99:0,268,
0:21,0:21:99:0,130, 0:23,0:23:99:0,142, 0:17,0:17:99:0,198, 0:15,0:15:99:0,147,
0:12,0:12:99:0,178,0:13,0:13:99:0,121,0:22,0:22:99:0,125,0:11,0:11:99:0,342,
0:8,0:8:99:0,166, 0:19,0:19:99:0,102, 0:22,0:22:99:0,211, 0:11,0:11:99:0,249,
0:10,0:10:99:0,147,0:26,0:26:99:0,186,0:13,0:13:99:0,260,0:10,0:10:99:0,156,
0:11,0:11:99:0,237, 0:9,0:9:99:0,233, 0:8,0:8:99:0,109, 0:10,0:10:99:0,203,
0:4,0:4:99:0,114, 0:6,0:6:99:0,207, 0:8,0:8:99:0,137, 0:9,0:9:99:0,114,
0:8,0:8:99:0,177, 0:11,0:11:99:0,101, 0:10,0:10:99:0,173, 0:13,0:13:99:0,109,
0:6,0:6:99:0,138, 0:8,0:8:99:0,139, 0:7,0:7:99:0,119, 0:9,0:9:99:0,148,
0:11,0:11:99:0,138,0:6,0:6:99:0,112,0:9,0:9:99:0,101,0:10,0:10:99:0,114,
0:10,0:10:99:0,107,0:7,0:7:99:0,144,0:4,0:4:99:0,127,1,0.009091,110,
Missing[NotAvailable], 1025, 0., 1, 0.009091, 60., Missing[NotAvailable],
25.41, Missing[NotAvailable], 1.981, Missing[NotAvailable]}
```

Dropping this (will do in the next folder so it doesn't affect the numbering): fullgenomeSNPEFF=Select[fullgenomeSNPEFF,StringContainsQ[#[[8]],"YNL109W"]==False&];

#439: YPL114W is dubious, but YPL113C is verified (SGD) [Updated]

```
In[*]:= fixme = 439;
```

Out[•]=

Manually swapping for the second gene:

```
In[*]:= fullgenomeSNPEFF[[fixme]]
```

```
{chrXVI, 16, 336149, G, A, 896.29, GT:AD:DP:GQ:PL, YPL114W,
MODERATE, missense_variant, YPL114W, c.202G>A, p.Ala68Thr,
A|missense_variant|MODERATE|YPL114W|YPL114W|transcript|YPL114W_mRNA|
   protein_coding|1/1|c.202G>A|p.Ala68Thr|202/420|202/420|68/139||,A|
  missense_variant|MODERATE|YPL113C|YPL113C|transcript|YPL113C_mRNA|
  protein_coding|1/1|c.995C>T|p.Ala332Val|995/1191|995/1191|332/396||,A|
  upstream_gene_variant|MODIFIER|BEM3|YPL115C|transcript|YPL115C_mRNA|
   protein_coding||c.-663C>T|||||663|,A|upstream_gene_variant|MODIFIER|CAR1|
  YPL111W|transcript|YPL111W_mRNA|protein_coding||c.-3795G>A||||3795|,A|
  downstream_gene_variant|MODIFIER|HOS3|YPL116W|transcript|YPL116W_mRNA|
  protein_coding||c.*4450G>A|||||4450|,A|downstream_gene_variant|MODIFIER|PEX25|
  YPL112C|transcript|YPL112C_mRNA|protein_coding||c.*1287C>T|||||1287|,A|
  downstream_gene_variant|MODIFIER|IMT2|tM(CAU)P|transcript|tM(CAU)P_tRNA|
   protein_coding||c.*2699C>T|||||2699|WARNING_TRANSCRIPT_NO_START_CODON,A|
  downstream_gene_variant|MODIFIER|GDE1|YPL110C|transcript|YPL110C_mRNA|
   protein_coding||c.*4919C>T||||4919|, 0:10,0:10:99:0,307,
0:5,0:5:99:0,128,0:8,0:8:99:0,276,0:7,0:7:99:0,201,0:4,0:4:99:0,99,
0:7,0:7:99:0,185, 0:6,0:6:99:0,214, 0:6,0:6:99:0,194, 0:4,0:4:99:0,110,
```

```
0:5,0:5:99:0,163,0:6:99:0,169,0:5,0:5:99:0,174,0:6,0:6:99:0,180,
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0:4,0:4:99:0,132, 0:7,0:7:99:0,185, 0:10,0:10:99:0,162, 0:3,0:3:99:0,100,
0:9,0:9:99:0,275, 0:7,0:7:99:0,182, 1:0,28:28:99:917,0, 0:4,0:4:99:0,110,
0:5,0:5:99:0,169,0:10,0:10:99:0,321,0:10,0:10:99:0,360,0:19,0:19:99:0,577,
0:5,0:5:99:0,181, 0:7,0:7:99:0,259, 0:6,0:6:99:0,200, 0:3,0:3:99:0,100,
0:3,0:3:99:0,100, 0:3,0:3:99:0,100, 0:6,0:6:99:0,176, 0:7,0:7:99:0,180,
0:11,0:11:99:0,341,0:6,0:6:99:0,176,0:5,0:5:99:0,159,0:4,0:4:99:0,113,
0:9,0:9:99:0,244, 0:6,0:6:99:0,126, 0:7,0:7:99:0,137, 0:5,0:5:99:0,159,
0:8,0:8:99:0,258, 0:4,0:4:99:0,135, 0:5,0:5:99:0,150, 0:3,0:3:99:0,100,
0:4,0:4:99:0,100, 0:3,0:3:99:0,114, 0:9,0:9:99:0,280, 0:7,0:7:99:0,203,
0:5,0:5:99:0,173, 0:4,0:4:99:0,137, 0:6,0:6:99:0,204, 0:3,0:3:99:0,113,
0:7,0:7:99:0,190,0:8,0:8:99:0,254,0:5,0:5:99:0,144,0:7,0:7:99:0,229,
0:9,0:9:99:0,270, 0:12,0:12:99:0,380, 0:9,0:9:99:0,260, 0:9,0:9:99:0,256,
0:11,0:11:99:0,263, 0:9,0:9:99:0,278, 0:5,0:5:99:0,138, 0:9,0:9:99:0,146,
0:8,0:8:99:0,255, 0:8,0:8:99:0,270, 0:9,0:9:99:0,295, 0:8,0:8:99:0,260,
0:11,0:11:99:0,325,0:9,0:9:99:0,294,0:5,0:5:99:0,169,0:14,0:14:99:0,303,
0:6,0:6:99:0,222, 0:8,0:8:99:0,269, 0:8,0:8:99:0,249, 0:7,0:7:99:0,215,
0:8,0:8:99:0,219, 0:5,0:5:99:0,172, 0:9,0:9:99:0,273, 0:9,0:9:99:0,293,
0:5,0:5:99:0,159, 0:7,0:7:99:0,218, 0:4,0:4:99:0,113, 0:5,0:5:99:0,135,
0:7,0:7:99:0,198, 0:7,0:7:99:0,199, 0:6,0:6:99:0,173, 0:10,0:10:99:0,278,
0:8,0:8:99:0,257, 0:11,0:11:99:0,364, 0:8,0:8:99:0,214, 0:7,0:7:99:0,144,
0:7, 0:7:99:0, 113, \ 0:9, 0:9:99:0, 270, \ 0:11, 0:11:99:0, 360, \ 0:5, 0:5:99:0, 135, \\
0:6,0:6:99:0,132,0:6,0:6:99:0,180,0:9,0:9:99:0,246,0:4,0:4:99:0,100,
0:3,0:3:99:0,100, 1, 0.009091, 110, Missing[NotAvailable], 766, 0., 1, 0.009091, 60.,
Missing[NotAvailable], 32.01, Missing[NotAvailable], 0.997, Missing[NotAvailable]}
```

In[*]:= fullgenomeSNPEFF[[fixme]] = {"chrXVI", 16, 336149, "G", "A", 896.29`, "GT:AD:DP:GQ:PL", "YPL113C", "MODERATE", "missense_variant", "YPL113C", "c.995C>T", "p.Ala332Val", "A|missense_variant|MODERATE|YPL114W|YPL114W|transcript|YPL114W_mRNA| protein coding|1/1|c.202G>A|p.Ala68Thr|202/420|202/420|68/139||,A| missense_variant|MODERATE|YPL113C|YPL113C|transcript|YPL113C_mRNA| protein_coding|1/1|c.995C>T|p.Ala332Val|995/1191|995/1191|332/396||,A| upstream gene variant|MODIFIER|BEM3|YPL115C|transcript|YPL115C mRNA| protein_coding||c.-663C>T|||||663|,A|upstream_gene_variant|MODIFIER|CAR1| YPL111W|transcript|YPL111W_mRNA|protein_coding||c.-3795G>A|||||3795|,A| downstream gene variant|MODIFIER|HOS3|YPL116W|transcript|YPL116W mRNA| protein_coding||c.*4450G>A|||||4450|,A|downstream_gene_variant|MODIFIER| PEX25|YPL112C|transcript|YPL112C_mRNA|protein_coding||c.*1287C>T|||||1287| ,A|downstream gene variant|MODIFIER|IMT2|tM(CAU)P|transcript|tM(CAU)P tRNA |protein_coding||c.*2699C>T|||||2699|WARNING_TRANSCRIPT_NO_START_CODON,A| downstream_gene_variant|MODIFIER|GDE1|YPL110C|transcript|YPL110C_mRNA| protein_coding||c.*4919C>T|||||4919|", "0:10,0:10:99:0,307",

```
"0:5,0:5:99:0,128", "0:8,0:8:99:0,276", "0:7,0:7:99:0,201", "0:4,0:4:99:0,99",
"0:7,0:7:99:0,185", "0:6,0:6:99:0,214", "0:6,0:6:99:0,194", "0:4,0:4:99:0,110",
"0:5,0:5:99:0,163", "0:6,0:6:99:0,169", "0:5,0:5:99:0,174", "0:6,0:6:99:0,180",
"0:5,0:5:99:0,150", "0:4,0:4:99:0,123", "0:8,0:8:99:0,241", "0:4,0:4:99:0,112",
"0:4,0:4:99:0,119", "0:4,0:4:99:0,100", "0:7,0:7:99:0,178", "0:4,0:4:99:0,130",
"0:4,0:4:99:0,132", "0:7,0:7:99:0,185", "0:10,0:10:99:0,162",
"0:3,0:3:99:0,100", "0:9,0:9:99:0,275", "0:7,0:7:99:0,182",
"1:0,28:28:99:917,0", "0:4,0:4:99:0,110", "0:5,0:5:99:0,169",
"0:10,0:10:99:0,321", "0:10,0:10:99:0,360", "0:19,0:19:99:0,577",
"0:5,0:5:99:0,181", "0:7,0:7:99:0,259", "0:6,0:6:99:0,200", "0:3,0:3:99:0,100",
"0:3,0:3:99:0,100", "0:3,0:3:99:0,100", "0:6,0:6:99:0,176", "0:7,0:7:99:0,180",
"0:11,0:11:99:0,341", "0:6,0:6:99:0,176", "0:5,0:5:99:0,159",
"0:4,0:4:99:0,113", "0:9,0:9:99:0,244", "0:6,0:6:99:0,126", "0:7,0:7:99:0,137",
"0:5,0:5:99:0,159", "0:8,0:8:99:0,258", "0:4,0:4:99:0,135", "0:5,0:5:99:0,150",
"0:3,0:3:99:0,100", "0:4,0:4:99:0,100", "0:3,0:3:99:0,114", "0:9,0:9:99:0,280",
"0:7,0:7:99:0,203", "0:5,0:5:99:0,173", "0:4,0:4:99:0,137", "0:6,0:6:99:0,204",
"0:3,0:3:99:0,113", "0:7,0:7:99:0,190", "0:8,0:8:99:0,254", "0:5,0:5:99:0,144",
"0:7,0:7:99:0,229", "0:9,0:9:99:0,270", "0:12,0:12:99:0,380",
"0:9,0:9:99:0,260", "0:9,0:9:99:0,256", "0:11,0:11:99:0,263",
"0:9,0:9:99:0,278", "0:5,0:5:99:0,138", "0:9,0:9:99:0,146", "0:8,0:8:99:0,255",
"0:8,0:8:99:0,270", "0:9,0:9:99:0,295", "0:8,0:8:99:0,260",
"0:11,0:11:99:0,325", "0:9,0:9:99:0,294", "0:5,0:5:99:0,169",
"0:14,0:14:99:0,303", "0:6,0:6:99:0,222", "0:8,0:8:99:0,269",
"0:8,0:8:99:0,249", "0:7,0:7:99:0,215", "0:8,0:8:99:0,219", "0:5,0:5:99:0,172",
"0:9,0:9:99:0,273", "0:9,0:9:99:0,293", "0:5,0:5:99:0,159", "0:7,0:7:99:0,218",
"0:4,0:4:99:0,113", "0:5,0:5:99:0,135", "0:7,0:7:99:0,198", "0:7,0:7:99:0,199",
"0:6,0:6:99:0,173", "0:10,0:10:99:0,278", "0:8,0:8:99:0,257",
"0:11,0:11:99:0,364", "0:8,0:8:99:0,214", "0:7,0:7:99:0,144",
"0:7,0:7:99:0,113", "0:9,0:9:99:0,270", "0:11,0:11:99:0,360",
"0:5,0:5:99:0,135", "0:6,0:6:99:0,132", "0:6,0:6:99:0,180",
"0:9,0:9:99:0,246", "0:4,0:4:99:0,100", "0:3,0:3:99:0,100", 1, 0.009091`,
110, "NA", 766, 0.`, 1, 0.009091`, 60.`, "NA", 32.01`, "NA", 0.997`, "NA"};
```

Revised gene list

Dropping dubious YNL109W from above (done now to avoid affecting the numbering):

```
In[*]:= fullgenomeSNPEFF =
       Select[fullgenomeSNPEFF, StringContainsQ[#[8], "YNL109W"] == False &];
```

In[*]:= genelist = Sort[Union[fullgenomeSNPEFF[All, 8]]]]

Out[•]=

{ABP1, ACB1, ACC1, ADE6, ADR1, AFT1, AGP1, AHK1, ALD5, ALY2, APC2, ARO3, ASN1, ATG11, ATG2, BDS1, BIR1, BLM10, BNA6, BNI1, BNI4, BRR2, BSC1, BSD2, BUL1, BUL2, CAJ1, CCC2, CCR4, CCW12, CDC25, CIC1, CLB3, CLN3, CNE1, COG1, COG3, COQ1, CPA1, CSM3, CUE3, CWC22, CWH41, CYK3, DAL2, DAN4, DBF20, DBP10, DDR48, DNF1, DOA4, DOT6, DSC2, DSE4, DUS4, EBS1, ECM14, ECM22, EFR3, ERB1, ERG1, ERG27, ERG7, ERG9, FAS2, FAU1, FCY2, FET4, FIG2, FIG4, FLC2, FLO1, FLO9, FMP52, FRE6, FSF1, FYV10, FYV6, FZF1, GAS4, GAT1, GCD2, GPB1, GPB2, GPH1, GSC2, HAP1, HBT1, HEH2, HHF2, HIS4, HMF1, HO, HRT3, HSL1, HSP104, HUL5, HXT13, IDP1, IES3, ILV2, IMA1, INP2, IOC4, IPI3, IRC20, IRC8, IST2, ISU1, ITC1, KAR3, KGD1, KIN1, KIN82, KRE5, KSP1, KTR3, LAM1, LAM6, LOS1, LRG1, MAM3, MAS2, MCK1, MCM5, MCT1, MDH2, MDL2, MDM30, MET10, MET30, MEX67, MHP1, MKT1, MLF3, MLH3, MLS1, MMS4, MNN9, MOT1, MPA43, MRL1, MRPL36, MSC6, MSS11, MSS2, MTR4, MYO2, NAM8, NAR1, NFT1, NGG1, NIP100, NPL3, NSR1, NUC1, NUG1, OAF1, OCA4, OCA5, OPI1, OSH2, PBP2, PBS2, PCL5, PDA1, PDR1, PDR10, PDX1, PET111, PET127, PEX2, PFK26, PHM7, PH023, PH081, PH084, PIB1, PIR3, PKP1, PLB2, PMA1, POL3, PPQ1, PRC1, PRI1, PRI2, PRP2, PRP8, PTK2, PYK2, QRI7, RAD16, RAD17, RAD26, RBK1, RCE1, RCK2, RCO1, REC102, RED1, RHO1, RIM11, RIM21, RKM3, RMP1, RNY1, ROG1, RPA135, RPI1, RPL22A, RPL2A, RPL39, RPS15, RPS2, RPT5, RRB1, RRG7, RRP46, RRP6, RSC1, RSE1, RTC2, SAK1, SAP4, SBE2, SEC8, SET2, SFP1, SHP1, SIW14, SKN1, SMC2, SMY2, SNF7, SNT2, SPC105, SP077, SPS100, SRP40, SSE1, SSK2, SSQ1, SSZ1, STE5, STP3, STT3, SUB2, SUC2, SUM1, SWC5, SWS2, SWT1, SYF2, SYG1, TAF8, TAH11, TAO3, TC089, TFB1, TFG1, TIM54, TOM1, TOP1, TOP2, TOS3, TPS3, TRK2, TRL1, TUS1, UBC1, UBP5, UBR2, UBX7, UGA3, UGA4, UIP5, ULS1, UME6, URA2, URB1, USA1, UTP21, UTP8, VBA1, VCX1, VHR2, VID22, VPS13, VPS74, VRP1, VTC1, VTC3, VTC4, VTC5, WAR1, YAP6, YBL109W, YBR134W, YBR242W, YBR292C, YCF1, YCK1, YCT1, YDL176W, YDL199C, YDR003W-A, YDR157W, YDR381C-A, YDR541C, YDR544C, YEF1, YER087C-A, YER156C, YFR036W-A, YGR126W, YGR130C, YGR266W, YHC1, YHC3, YHL008C, YHR028W-A, YHR071C-A, YIH1, YIL092W, YIR020W-A, YKR073C, YLL020C, YLL066W-B, YLR108C, YLR296W, YLR312C, YLR372W, YML119W, YMR027W, YMR317W, YOR029W, YOR1, YOR296W, YOR343C, YPK1, YPL025C, YPL113C, YPR078C, YPR089W, YPR117W, YPT6, YRA1, YRB2, ZDS2, ZRT1, ZRT3}

Checking all of the systematic ("Y...") names:

```
In[*]:= checkme = {"YBL109W", "YBR134W", "YBR242W", "YBR292C", "YDL199C",
        "YDR003W-A", "YDR157W", "YDR541C", "YDR544C", "YER087C-A", "YFR036W-A",
        "YGR126W", "YGR130C", "YGR266W", "YHL008C", "YHR028W-A", "YHR071C-A",
        "YIL092W", "YIR020W-A", "YKR073C", "YLL020C", "YLL066W-B", "YLR108C",
        "YLR296W", "YML119W", "YMR027W", "YMR317W", "YOR029W", "YOR296W",
        "YOR343C", "YPL025C", "YPL113C", "YPL114W", "YPR078C", "YPR089W"};
```

Some now have standardized names (replacing these names):

```
In[@]:= Position[fullgenomeSNPEFF[All, 8], "YDL176W"][1, 1]
```

Out[•]=

```
In[*]:= fullgenomeSNPEFF[[%, 8]] = "GID12"
Out[ • ]=
       GID12
 In[*]:= Position[fullgenomeSNPEFF[All, 8], "YDR381C-A"][1, 1]
Out[ • ]=
       91
 In[*]:= fullgenomeSNPEFF[[%, 8]] = "COI1"
Out[ • ]=
       COI1
 In[@]:= Position[fullgenomeSNPEFF[All, 8]], "YER156C"] [[1, 1]]
Out[ • ]=
       113
 In[@]:= fullgenomeSNPEFF[[%, 8]] = "MYG1"
Out[ • ]=
       MYG1
 In[*]:= Position[fullgenomeSNPEFF[All, 8], "YLR312C"][1, 1]
Out[ • ]=
       315
 In[*]:= fullgenomeSNPEFF[[%, 8]] = "ATG39"
Out[ • ]=
       ATG39
 In[@]:= Position[fullgenomeSNPEFF[All, 8]], "YLR372W"] [[1, 1]]
Out[ • ]=
       321
 In[*]:= fullgenomeSNPEFF[[%, 8]] = "ELO3"
Out[ • ]=
       EL03
 In[*]:= Position[fullgenomeSNPEFF[All, 8], "YPR117W"][1, 1]
Out[ • ]=
       452
 In[*]:= fullgenomeSNPEFF[[%, 8]] = "HOB2"
Out[ • ]=
       HOB2
       Throwing out the dubious genes according to SGD [6 Nov 2024] (keeping uncharacterized):
 In[*]:= dropme = {"YBL109W", "YBR134W", "YDR544C", "YER087C-A", "YFR036W-A",
           "YHR028W-A", "YHR071C-A", "YIR020W-A", "YLL020C", "YPL025C"};
       Depth of coverage for these genes is pretty typical:
```

```
In[*]:= Select[fullgenomeSNPEFF, MemberQ[dropme, #[8]] &] [All, {8, 129}]
      Mean[%[All, 2]]] // N
Out[ • ]=
       {{YBL109W, 410}, {YBR134W, 976}, {YDR544C, 400},
        {YER087C-A, 814}, {YFR036W-A, 823}, {YHR028W-A, 691},
        {YHR071C-A, 968}, {YIR020W-A, 726}, {YLL020C, 1251}, {YPL025C, 846}}
Out[ • ]=
      790.5
      Dropping these ten dubious genes:
 In[*]:= Length[fullgenomeSNPEFF];
      fullgenomeSNPEFF = Select[fullgenomeSNPEFF, MemberQ[dropme, #[8]] == False &];
      %% - Length[fullgenomeSNPEFF]
Out[ • ]=
      10
```

In[*]:= genelist = Sort[Union[fullgenomeSNPEFF[All, 8]]]

{ABP1, ACB1, ACC1, ADE6, ADR1, AFT1, AGP1, AHK1, ALD5, ALY2, APC2, ARO3, ASN1, ATG11, ATG2, ATG39, BDS1, BIR1, BLM10, BNA6, BNI1, BNI4, BRR2, BSC1, BSD2, BUL1, BUL2, CAJ1, CCC2, CCR4, CCW12, CDC25, CIC1, CLB3, CLN3, CNE1, COG1, COG3, COI1, COQ1, CPA1, CSM3, CUE3, CWC22, CWH41, CYK3, DAL2, DAN4, DBF20, DBP10, DDR48, DNF1, DOA4, DOT6, DSC2, DSE4, DUS4, EBS1, ECM14, ECM22, EFR3, ELO3, ERB1, ERG1, ERG27, ERG7, ERG9, FAS2, FAU1, FCY2, FET4, FIG2, FIG4, FLC2, FLO1, FLO9, FMP52, FRE6, FSF1, FYV10, FYV6, FZF1, GAS4, GAT1, GCD2, GID12, GPB1, GPB2, GPH1, GSC2, HAP1, HBT1, HEH2, HHF2, HIS4, HMF1, HO, HOB2, HRT3, HSL1, HSP104, HUL5, HXT13, IDP1, IES3, ILV2, IMA1, INP2, IOC4, IPI3, IRC20, IRC8, IST2, ISU1, ITC1, KAR3, KGD1, KIN1, KIN82, KRE5, KSP1, KTR3, LAM1, LAM6, LOS1, LRG1, MAM3, MAS2, MCK1, MCM5, MCT1, MDH2, MDL2, MDM30, MET10, MET30, MEX67, MHP1, MKT1, MLF3, MLH3, MLS1, MMS4, MNN9, MOT1, MPA43, MRL1, MRPL36, MSC6, MSS11, MSS2, MTR4, MYG1, MYO2, NAM8, NAR1, NFT1, NGG1, NIP100, NPL3, NSR1, NUC1, NUG1, OAF1, OCA4, OCA5, OPI1, OSH2, PBP2, PBS2, PCL5, PDA1, PDR1, PDR10, PDX1, PET111, PET127, PEX2, PFK26, PHM7, PH023, PH081, PH084, PIB1, PIR3, PKP1, PLB2, PMA1, POL3, PPQ1, PRC1, PRI1, PRI2, PRP2, PRP8, PTK2, PYK2, QRI7, RAD16, RAD17, RAD26, RBK1, RCE1, RCK2, RCO1, REC102, RED1, RHO1, RIM11, RIM21, RKM3, RMP1, RNY1, ROG1, RPA135, RPI1, RPL22A, RPL2A, RPL39, RPS15, RPS2, RPT5, RRB1, RRG7, RRP46, RRP6, RSC1, RSE1, RTC2, SAK1, SAP4, SBE2, SEC8, SET2, SFP1, SHP1, SIW14, SKN1, SMC2, SMY2, SNF7, SNT2, SPC105, SP077, SPS100, SRP40, SSE1, SSK2, SSQ1, SSZ1, STE5, STP3, STT3, SUB2, SUC2, SUM1, SWC5, SWS2, SWT1, SYF2, SYG1, TAF8, TAH11, TAO3, TCO89, TFB1, TFG1, TIM54, TOM1, TOP1, TOP2, TOS3, TPS3, TRK2, TRL1, TUS1, UBC1, UBP5, UBR2, UBX7, UGA3, UGA4, UIP5, ULS1, UME6, URA2, URB1, USA1, UTP21, UTP8, VBA1, VCX1, VHR2, VID22, VPS13, VPS74, VRP1, VTC1, VTC3, VTC4, VTC5, WAR1, YAP6, YBR242W, YBR292C, YCF1, YCK1, YCT1, YDL199C, YDR003W-A, YDR157W, YDR541C, YEF1, YGR126W, YGR130C, YGR266W, YHC1, YHC3, YHL008C, YIH1, YIL092W, YKR073C, YLL066W-B, YLR108C, YLR296W, YML119W, YMR027W, YMR317W, YOR029W, YOR1, YOR296W, YOR343C, YPK1, YPL113C, YPR078C, YPR089W, YPT6, YRA1, YRB2, ZDS2, ZRT1, ZRT3}

In[*]:= Length[genelist]

Out[•]=

Out[•]=

342

Checking ExcessHet - No further filtering of data needed

In[*]:= fullgenomeDIP[[1]]

••• Part: Part specification fullgenomeDIP[1] is longer than depth of object. 0

Out[•]=

fullgenomeDIP[1]

```
In[•]:= For[i = 1; highEXHET = {}, i ≤ Length[fullgenomeDIP], i++,
        temp = StringSplit[fullgenomeDIP[i, 8], {";"}];
        temp2 = ToExpression[StringSplit[temp[6], {"="}]][2];
        highEXHET = Append[highEXHET, {fullgenomeDIP[i, 1], fullgenomeDIP[i, 2], temp2}]
       1
 IN[*]:= highEXHET = highEXHET /. {"ref|NC 001133|" → "chrI", "ref|NC 001134|" → "chrII",
           "ref|NC_001135|" → "chrIII", "ref|NC_001136|" → "chrIV",
           "ref|NC_001137|" → "chrV", "ref|NC_001138|" → "chrVI",
           "ref|NC_001139|" → "chrVII",
           "ref|NC_001140|" → "chrVIII", "ref|NC_001141|" → "chrVIX",
           "ref|NC_001142|" → "chrX",
           "ref|NC_001143|" → "chrXI", "ref|NC_001144|" → "chrXII",
           "ref|NC 001145|" \rightarrow "chrXIII", "ref|NC_001146|" \rightarrow "chrXIV", "ref|NC_001147|" \rightarrow
            "chrXV", "ref|NC_001148|" → "chrXVI", "ref|NC_001224|" → "chrmt"};
 In[*]:= highEXHET = Select[highEXHET, #[3] ≥ 50 &];
      Looking at the ExcessHet calls from the diploid lines above 50:
 In[@]:= {Min[highEXHET[All, 3]], Max[highEXHET[All, 3]]}
Out[ • ]=
       \{\infty, -\infty\}
      The only positions in common are:
 Intersection[highEXHET[All, 2], fullgenomeSNPEFF[All, 3]]
Out[ • ]=
       { }
      But this isn't on the same chromosome (position 25178) or is a site with more than one alternate allele
      and so filtered out (position 889748):
 In[@]:= Select[highEXHET, MemberQ[temppos, #[2]] &]
Out[ • ]=
       { }
 In[*]:= Select[fullgenomeSNPEFF[All, 1;; 3], MemberQ[temppos, #[3]] &]
Out[ • ]=
       { }
```

Checking MQ

The mapping quality is almost always 60:

In[@]:= Histogram[fullgenomeSNPEFF[[All, 133]]]

Out[•]=

> Many of the ones with lower mapping quality are the ones with missing genotype calls (filtered out below):

In[@]:= Select[fullgenomeSNPEFF[[All, {1, 3, 8, 133}]], #[[4]] < 60 &] // MatrixForm</pre>

Out[•]//MatrixForm= 25 487 FL09 58.61 chrI chrI 25 489 FL09 57.62 chrI 27 090 FL09 52.4 204 218 FL01 56.97 chrI chrI 206 360 FL01 54.5 chrI 206 363 FL01 54.41 FL01 chrI 206 375 53.78 chrV 22913 HXT13 51.91 chrVIII 93335 YHL008C 58.24 chrVIII 93350 YHL008C 58.24 chrVIII 93359 YHL008C 58.13 chrVIII 93361 YHL008C 58.11 chrVIII 93370 YHL008C 58.16 chrVIII 93375 YHL008C 58.16 chrVIII 93391 YHL008C 58.28 chrVIII 93577 YHL008C 51.63 chrVIII 93608 YHL008C 54.81 chrVIII 93624 YHL008C 56.2 chrVIII 93898 YHL008C 50.58 715 087 DAN4 58.93 chrX chrX 715 108 DAN4 57.88 chrX 715 114 DAN4 57.14 59.04 chrX 715 117 DAN4 chrX 715 141 DAN4 55.81 chrX 715 149 DAN4 59.1 chrXI 144 880 PIR3 55.87 chrXI 144883 PIR3 56.03 chrXI 578481 YKR073C 53.08 chrXI 578 485 YKR073C 50.17 chrXI 613636 SRP40 59.9 chrXII 5683 YLL066W-B 57.77 chrXII 650822 HAP1 59.76 chrXIII 588588 MSS11 59.85 chrXIII 609388 DDR48 57.68 chrXIII 908159 YMR317W 57.88 chrXIII 908168 YMR317W 57.87 chrXIII 908215 YMR317W 56.28 chrXIII 908221 YMR317W 55.95

```
In[*]:= Union[%[All, 3]]]
```

Out[•]=

{DAN4, DDR48, FL01, FL09, HAP1, HXT13, MSS11, PIR3, SRP40, YHL008C, YKR073C, YLL066W-B, YMR317W}

Mean mapping quality of all genes:

In[*]:= Mean[fullgenomeSNPEFF[All, 133]]]

Out[•]=

59.7012

Mean mapping quality of the genes dropped below with ≥5 "." calls:

58.8042

```
In[a]:= {"BSC1", "DAN4", "FLO1", "FLO9", "HAP1", "MSS11", "PIR3", "YHL008C", "YKR073C"};
       Select[fullgenomeSNPEFF, MemberQ[%, #[8]] &];
      Mean[%[All, 133]]]
Out[ • ]=
      56.6091
      Of the genes filtered out because of having 5 or more uncalled genotypes, all but BSC1 have MQ<60
       Mean mapping quality of the genes not dropped, but having some "." calls:
 In[*]:= {"SRP40", "YGR130C", "YLL066W-B", "YLR296W", "YMR317W"};
       Select[fullgenomeSNPEFF, MemberQ[%, #[8]] &];
      Mean[%[All, 133]]]
Out[ • ]=
```

Of the genes not filtered out (but of concern), three have MQ<60 {SRP40,YLL066W-B,YMR317W}.

Dropping genes with too many uncalled genotypes and dropping genotypic calls with <5 depth

Processing and dropping genes with two many poor quality genotypes

Sites that were hard to call (low depth or no genotype call) can indicate duplicated genes and alignment problems. Here we identify these:

```
In[@]:= droppos = -1 + Position[header, "OLY077_S101"] [[1, 1]];
       (*# columns to be dropped before first sample*)
 In[*]:* topos = Position[header, "ZnBM47_S95"] [[1, 1]]; (*# last sample column*)
       Extracting the ref vs alt call in genomeSNPEFF:
 In[*]:= genomeSNPEFF = Table[".", {i, 1, Length[fullgenomeSNPEFF]}, {j, 1, topos - droppos}];
 In[@]:= genomeSNPEFF[[1]] // Length
Out[ • ]=
      110
 In[⊕]:= For[i = 1, i ≤ Length[fullgenomeSNPEFF], i++,
        For [j = 1 + droppos, j \le topos, j++,
         temp = StringSplit[fullgenomeSNPEFF[i, j], {":"}];
         genomeSNPEFF[[i, j - droppos]] = temp[[1]];
        ]
       1
 In[*]:= genomeSNPEFF // Length
Out[ • ]=
      446
```

```
In[*]:= Tally[Select[genomeSNPEFF[[1]], # # "." &]]
Out[ • ]=
       \{\{0, 101\}, \{1, 9\}\}
       All mutant sites and the tally of mutations in them:
 In[*]:= summarytable = Table[{fullgenomeSNPEFF[[i, 1]],
            fullgenomeSNPEFF[i, 3], fullgenomeSNPEFF[i, 8], fullgenomeSNPEFF[i, 9],
            Sort[Tally[genomeSNPEFF[i]]]]}, {i, 1, Length[genomeSNPEFF]}];
       The set of sites where the number of "." calls was ≥5:
 ln[\ \circ\ ]:=\ dropthese = Select[summarytable, (#[[5, 1, 1]] == ".") \&\& (#[[5, 1, 2]] \ge 5) \&];
       dropthese // MatrixForm
Out[ • ]//MatrixForm=
          chrI
                   27090
                             FL09
                                         LOW
                                                 \{\{., 83\}, \{0, 2\}, \{1, 25\}\}
          chrI
                  204 218
                             FL01
                                     MODERATE \{\{., 71\}, \{0, 19\}, \{1, 20\}\}
           chrI
                  206 360
                             FL01
                                         LOW
                                                 \{\{., 80\}, \{0, 16\}, \{1, 14\}\}\
                  206 363
                             FL01
                                         LOW
                                                 \{\{., 70\}, \{0, 26\}, \{1, 14\}\}
          chrI
          chrI
                  206 375
                             FL01
                                         LOW
                                                 \{\{., 95\}, \{0, 1\}, \{1, 14\}\}
                                                 \{\{., 99\}, \{0, 7\}, \{1, 4\}\}
          chrIV
                  384 924
                             BSC1
                                        HIGH
         chrVIII 93335 YHL008C
                                         LOW
                                                 \{\{., 41\}, \{0, 1\}, \{1, 68\}\}
         chrVIII 93350
                           YHL008C
                                         LOW
                                                 \{\{., 39\}, \{0, 3\}, \{1, 68\}\}
        chrVIII 93359
                           YHL008C MODERATE {{., 29}, {0, 15}, {1, 66}}
         chrVIII 93375 YHL008C
                                        HIGH
                                                 \{\{., 30\}, \{0, 44\}, \{1, 36\}\}\
        chrVIII 93391 YHL008C MODERATE {{., 65}, {0, 9}, {1, 36}}
           chrX
                  715 087
                             DAN4
                                         LOW
                                                  \{\{., 6\}, \{0, 97\}, \{1, 7\}\}
          chrX
                  715 108
                             DAN4
                                         LOW
                                                 \{\{., 65\}, \{0, 2\}, \{1, 43\}\}
                  715 114
                             DAN4
                                         LOW
                                                 \{\{., 68\}, \{0, 5\}, \{1, 37\}\}\
          chrX
          chrX
                  715 141
                             DAN4
                                         LOW
                                                 \{\{., 61\}, \{0, 27\}, \{1, 22\}\}\
          chrXI 144880
                             PIR3
                                         LOW
                                                 \{\{., 76\}, \{0, 5\}, \{1, 29\}\}
          chrXI 144883
                                                 \{\{., 52\}, \{0, 29\}, \{1, 29\}\}
                             PIR3
                                         LOW
                  578 481 YKR073C
                                        HIGH
                                                 \{\{., 32\}, \{0, 77\}, \{1, 1\}\}
          chrXI
         chrXII 650822
                             HAP1
                                        HIGH
                                                 \{\{., 90\}, \{0, 14\}, \{1, 6\}\}
         chrXII 650830
                             HAP1
                                         LOW
                                                 \{\{., 98\}, \{0, 1\}, \{1, 11\}\}
         chrXIII 588588
                             MSS11 MODERATE {{., 8}, {0, 25}, {1, 77}}
 In[*]:= dropme = Union[dropthese[All, 3]]]
       % // Length
Out[ • ]=
       {BSC1, DAN4, FLO1, FLO9, HAP1, MSS11, PIR3, YHL008C, YKR073C}
Out[ • ]=
```

Depth of coverage for these genes is much higher on average, with both the high outliers (FLO1) and the two low outliers (BSC1 and HAP1):

```
In[*]:= Select[fullgenomeSNPEFF, MemberQ[dropme, #[8]] & [[All, {8, 129}]]
      Mean[%[All, 2]] // N
Out[ • ]=
       {{FL09, 1799}, {FL09, 1584}, {FL09, 2213}, {FL01, 348}, {FL01, 15599}, {FL01, 15734},
        {FL01, 17792}, {BSC1, 91}, {YHL008C, 2060}, {YHL008C, 2102}, {YHL008C, 2001},
        {YHL008C, 1983}, {YHL008C, 1921}, {YHL008C, 1941}, {YHL008C, 2062}, {YHL008C, 1441},
        {YHL008C, 1946}, {YHL008C, 1810}, {YHL008C, 2247}, {DAN4, 905}, {DAN4, 1522},
        {DAN4, 1559}, {DAN4, 1583}, {DAN4, 1310}, {DAN4, 628}, {PIR3, 2648}, {PIR3, 2800},
        {YKR073C, 1609}, {YKR073C, 1536}, {HAP1, 509}, {HAP1, 84}, {MSS11, 658}}
Out[ • ]=
      2938.28
      Mapping quality is also poorer on average, except for BSC1, with the mean at the bottom 3.5% tail:
 In[*]:= Select[fullgenomeSNPEFF, MemberQ[dropme, #[8]] &] [[All, {8, 133}]]
      Mean[%[All, 2]] // N
      Length[Select[fullgenomeSNPEFF[All, 133], # < % &]] /</pre>
          Length[fullgenomeSNPEFF[All, 133]] // N // PercentForm
Out[ • ]=
       {{FL09, 58.61}, {FL09, 57.62}, {FL09, 52.4}, {FL01, 56.97}, {FL01, 54.5},
        {FL01, 54.41}, {FL01, 53.78}, {BSC1, 60.}, {YHL008C, 58.24}, {YHL008C, 58.24},
        {YHL008C, 58.13}, {YHL008C, 58.11}, {YHL008C, 58.16}, {YHL008C, 58.16},
        {YHL008C, 58.28}, {YHL008C, 51.63}, {YHL008C, 54.81}, {YHL008C, 56.2},
        {YHL008C, 50.58}, {DAN4, 58.93}, {DAN4, 57.88}, {DAN4, 57.14},
        {DAN4, 59.04}, {DAN4, 55.81}, {DAN4, 59.1}, {PIR3, 55.87}, {PIR3, 56.03},
        {YKR073C, 53.08}, {YKR073C, 50.17}, {HAP1, 59.76}, {HAP1, 60.}, {MSS11, 59.85}}
Out[ • ]=
      56.6091
Out[ • ]//PercentForm=
      3.587%
      Dropping all sites (32) in the nine genes with ≥5 "." calls:
 In[*]:= Length[fullgenomeSNPEFF];
      fullgenomeSNPEFF = Select[fullgenomeSNPEFF, MemberQ[dropme, #[8]]] == False &];
      %% - Length[fullgenomeSNPEFF]
Out[ • ]=
      32
```

Considering the others with uncalled genotypes, first recalculating the summary table:

```
In[*]:= genomeSNPEFF = Table[".", {i, 1, Length[fullgenomeSNPEFF]}, {j, 1, topos - droppos}];
      For[i = 1, i ≤ Length[fullgenomeSNPEFF], i++,
        For [j = 1 + droppos, j \le topos, j++,
         temp = StringSplit[fullgenomeSNPEFF[[i, j]], {":"}];
         genomeSNPEFF[[i, j - droppos]] = temp[[1]];
        ]
       1
       summarytable = Table[{fullgenomeSNPEFF[[i, 1]],
            fullgenomeSNPEFF[i, 3], fullgenomeSNPEFF[i, 8], fullgenomeSNPEFF[i, 9],
            Sort[Tally[genomeSNPEFF[i]]]], {i, 1, Length[genomeSNPEFF]}];
 In[*]:= consider =
         Select[summarytable, (#[5, 1, 1] = ".") && (#[5, 1, 2] < 5) && (#[5, 1, 2] \ge 1) &];
       consider // MatrixForm
Out[ • ]//MatrixForm=
         chrVII 753296 YGR130C MODERATE {{., 1}, {0, 23}, {1, 86}}
         chrXI 613636
                           SRP40
                                      MODERATE {{., 1}, {0, 1}, {1, 108}}
         chrXII 5683 YLL066W-B
                                        HIGH
                                                 \{\{.,3\},\{0,99\},\{1,8\}\}
         chrXII 723168 YLR296W
                                                 \{\{., 2\}, \{0, 104\}, \{1, 4\}\}
                                        HIGH
        chrXIII 908215 YMR317W
                                         LOW
                                                 \{\{., 1\}, \{0, 17\}, \{1, 92\}\}
        chrXIII 908221 YMR317W
                                         LOW
                                                 \{\{., 2\}, \{0, 9\}, \{1, 99\}\}
 In[*]:= considergenes = Union[consider[All, 3]]
      % // Length
Out[ • ]=
       {SRP40, YGR130C, YLL066W-B, YLR296W, YMR317W}
Out[ • ]=
      Depth of coverage for these genes is somewhat higher on average:
 In[*]:= Select[fullgenomeSNPEFF, MemberQ[considergenes, #[8]] &] [[All, {8, 129}]]
       Mean[%[All, 2]] // N
Out[ • ]=
       {\( \text{YGR130C}, 2020 \)\), \( \text{YGR130C}, 2020 \)\), \( \text{YGR130C}, 1799 \)\), \( \text{YGR130C}, 1809 \)\,
        {SRP40, 1173}, {YLL066W-B, 607}, {YLR296W, 974}, {YLR296W, 977},
        {YMR317W, 1959}, {YMR317W, 1960}, {YMR317W, 1836}, {YMR317W, 1789}}
Out[ • ]=
      1576.92
```

Mapping quality is slightly smaller (in the bottom 1.7% tail):

```
In[0]:= Select[fullgenomeSNPEFF, MemberQ[considergenes, #[8]] &] [All, {8, 133}]
      Mean[%[All, 2]] // N
      Length[Select[fullgenomeSNPEFF[All, 133], # < % &]] /</pre>
          Length[fullgenomeSNPEFF[All, 133]] // N // PercentForm
Out[ • ]=
       {{YGR130C, 60.}, {YGR130C, 60.}, {YGR130C, 60.}, {YGR130C, 60.},
        {SRP40, 59.9}, {YLL066W-B, 57.77}, {YLR296W, 60.}, {YLR296W, 60.},
        {YMR317W, 57.88}, {YMR317W, 57.87}, {YMR317W, 56.28}, {YMR317W, 55.95}}
Out[ • ]=
      58.8042
Out[•]//PercentForm=
      1.691%
      These genes are kept but will be flagged as:
           {SRP40,YLL066W-B,YMR317W} have MQ<60
           {YGR130C,YMR317W} have depth ~ twice average depth (~1000)
```

Processing and dropping sites with <5 depth

We next scan for sites that are called with depth <5 (note that we do not filter on the depth for each allele, just total depth at that site in that sample).

An example of a site with low coverage:

```
In[*]:= Position[fullgenomeSNPEFF[All, 3], 723168][1, 1];
      fullgenomeSNPEFF[%, 106;; 110]
Out[ • ]=
      \{1:0,1:1:26:26,0,0:10,0:10:99:0,249,
       0:3,3:6:3:0,3,0:8,0:8:99:0,252,0:4,0:4:99:0,108
 In[•]:= For[i = 1; altered = {}, i ≤ Length[fullgenomeSNPEFF], i++,
       For [j = 1 + droppos, j \le topos, j++,
         temp = StringSplit[fullgenomeSNPEFF[[i, j]], {":"}];
         If[ToExpression[temp[3]]] < 5, temp[1] = ".";</pre>
          fullgenomeSNPEFF[[i, j]] = StringRiffle[temp, ":"];
          AppendTo[altered, {i, j}]];
       ]
      1
```

Out[•]=

 $\{\{0, 104\}, \{1, 1\}\}$

```
In[*]:= altered
Out[ • ]=
                                                       \{\{1,33\},\{1,51\},\{1,67\},\{1,75\},\{1,124\},\{2,33\},\{2,51\},\{2,67\},\{2,75\},\{2,124\},\{3,33\},
                                                            \{3, 51\}, \{3, 67\}, \{3, 75\}, \{4, 19\}, \{4, 36\}, \{4, 46\}, \{4, 51\}, \{4, 53\}, \{4, 55\}, \{4, 67\}, \{4, 68\}, \{4, 68\}, \{4, 68\}, \{4, 68\}, \{4, 68\}, \{4, 68\}, \{4, 68\}, \{4, 68\}, \{4, 68\}, \{4, 68\}, \{4, 68\}, \{4, 68\}, \{4, 68\}, \{4, 68\}, \{4, 68\}, \{4, 68\}, \{4, 68\}, \{4, 68\}, \{4, 68\}, \{4, 68\}, \{4, 68\}, \{4, 68\}, \{4, 68\}, \{4, 68\}, \{4, 68\}, \{4, 68\}, \{4, 68\}, \{4, 68\}, \{4, 68\}, \{4, 68\}, \{4, 68\}, \{4, 68\}, \{4, 68\}, \{4, 68\}, \{4, 68\}, \{4, 68\}, \{4, 68\}, \{4, 68\}, \{4, 68\}, \{4, 68\}, \{4, 68\}, \{4, 68\}, \{4, 68\}, \{4, 68\}, \{4, 68\}, \{4, 68\}, \{4, 68\}, \{4, 68\}, \{4, 68\}, \{4, 68\}, \{4, 68\}, \{4, 68\}, \{4, 68\}, \{4, 68\}, \{4, 68\}, \{4, 68\}, \{4, 68\}, \{4, 68\}, \{4, 68\}, \{4, 68\}, \{4, 68\}, \{4, 68\}, \{4, 68\}, \{4, 68\}, \{4, 68\}, \{4, 68\}, \{4, 68\}, \{4, 68\}, \{4, 68\}, \{4, 68\}, \{4, 68\}, \{4, 68\}, \{4, 68\}, \{4, 68\}, \{4, 68\}, \{4, 68\}, \{4, 68\}, \{4, 68\}, \{4, 68\}, \{4, 68\}, \{4, 68\}, \{4, 68\}, \{4, 68\}, \{4, 68\}, \{4, 68\}, \{4, 68\}, \{4, 68\}, \{4, 68\}, \{4, 68\}, \{4, 68\}, \{4, 68\}, \{4, 68\}, \{4, 68\}, \{4, 68\}, \{4, 68\}, \{4, 68\}, \{4, 68\}, \{4, 68\}, \{4, 68\}, \{4, 68\}, \{4, 68\}, \{4, 68\}, \{4, 68\}, \{4, 68\}, \{4, 68\}, \{4, 68\}, \{4, 68\}, \{4, 68\}, \{4, 68\}, \{4, 68\}, \{4, 68\}, \{4, 68\}, \{4, 68\}, \{4, 68\}, \{4, 68\}, \{4, 68\}, \{4, 68\}, \{4, 68\}, \{4, 68\}, \{4, 68\}, \{4, 68\}, \{4, 68\}, \{4, 68\}, \{4, 68\}, \{4, 68\}, \{4, 68\}, \{4, 68\}, \{4, 68\}, \{4, 68\}, \{4, 68\}, \{4, 68\}, \{4, 68\}, \{4, 68\}, \{4, 68\}, \{4, 68\}, \{4, 68\}, \{4, 68\}, \{4, 68\}, \{4, 68\}, \{4, 68\}, \{4, 68\}, \{4, 68\}, \{4, 68\}, \{4, 68\}, \{4, 68\}, \{4, 68\}, \{4, 68\}, \{4, 68\}, \{4, 68\}, \{4, 68\}, \{4, 68\}, \{4, 68\}, \{4, 68\}, \{4, 68\}, \{4, 68\}, \{4, 68\}, \{4, 68\}, \{4, 68\}, \{4, 68\}, \{4, 68\}, \{4, 68\}, \{4, 68\}, \{4, 68\}, \{4, 68\}, \{4, 68\}, \{4, 68\}, \{4, 68\}, \{4, 68\}, \{4, 68\}, \{4, 68\}, \{4, 68\}, \{4, 68\}, \{4, 68\}, \{4, 68\}, \{4, 68\}, \{4, 68\}, \{4, 68\}, \{4, 68\}, \{4, 68\}, \{4, 68\}, \{4, 68\}, \{4, 68\}, \{4, 68\}, \{4, 68\}, \{4, 68\}, \{4, 68\}, \{4, 68\}, \{4, 68\}, \{4, 68\}, \{4, 68\}, \{4, 68\}, \{4, 68\}, \{4, 68\}, \{4, 68\}, \{4, 68\}, \{4, 68\}, \{4, 68\}, \{4, 68\}, \{4, 68\}, \{4, 68\}, \{4, 68\}, \{4, 68\}, \{4, 68\}, \{4, 68\}, \{4, 68\}, \{4, 68\}, \{4, 68\}, \{4, 68\}, \{4, 68\}, \{4, 68\}, \{4, 68\}, \{4, 68\}, \{4, 68\}, \{4, 68\}, \{4, 68\}, \{4, 68\}, \{4, 68\}, \{
                                                           \{4, 69\}, \{4, 75\}, \{4, 76\}, \{4, 79\}, \{4, 116\}, \{4, 124\}, \{5, 19\}, \{5, 36\}, \{5, 46\}, \{5, 51\}, \{5, 53\},
                                                           \{5, 55\}, \{5, 67\}, \{5, 68\}, \{5, 69\}, \{5, 75\}, \{5, 76\}, \dots, \{413, 76\}, \{413, 77\}, \{413, 78\}, \{413, 78\}, \{413, 78\}, \{413, 78\}, \{413, 78\}, \{413, 78\}, \{413, 78\}, \{413, 78\}, \{413, 78\}, \{413, 78\}, \{413, 78\}, \{413, 78\}, \{413, 78\}, \{413, 78\}, \{413, 78\}, \{413, 78\}, \{413, 78\}, \{413, 78\}, \{413, 78\}, \{413, 78\}, \{413, 78\}, \{413, 78\}, \{413, 78\}, \{413, 78\}, \{413, 78\}, \{413, 78\}, \{413, 78\}, \{413, 78\}, \{413, 78\}, \{413, 78\}, \{413, 78\}, \{413, 78\}, \{413, 78\}, \{413, 78\}, \{413, 78\}, \{413, 78\}, \{413, 78\}, \{413, 78\}, \{413, 78\}, \{413, 78\}, \{413, 78\}, \{413, 78\}, \{413, 78\}, \{413, 78\}, \{413, 78\}, \{413, 78\}, \{413, 78\}, \{413, 78\}, \{413, 78\}, \{413, 78\}, \{413, 78\}, \{413, 78\}, \{413, 78\}, \{413, 78\}, \{413, 78\}, \{413, 78\}, \{413, 78\}, \{413, 78\}, \{413, 78\}, \{413, 78\}, \{413, 78\}, \{413, 78\}, \{413, 78\}, \{413, 78\}, \{413, 78\}, \{413, 78\}, \{413, 78\}, \{413, 78\}, \{413, 78\}, \{413, 78\}, \{413, 78\}, \{413, 78\}, \{413, 78\}, \{413, 78\}, \{413, 78\}, \{413, 78\}, \{413, 78\}, \{413, 78\}, \{413, 78\}, \{413, 78\}, \{413, 78\}, \{413, 78\}, \{413, 78\}, \{413, 78\}, \{413, 78\}, \{413, 78\}, \{413, 78\}, \{413, 78\}, \{413, 78\}, \{413, 78\}, \{413, 78\}, \{413, 78\}, \{413, 78\}, \{413, 78\}, \{413, 78\}, \{413, 78\}, \{413, 78\}, \{413, 78\}, \{413, 78\}, \{413, 78\}, \{413, 78\}, \{413, 78\}, \{413, 78\}, \{413, 78\}, \{413, 78\}, \{413, 78\}, \{413, 78\}, \{413, 78\}, \{413, 78\}, \{413, 78\}, \{413, 78\}, \{413, 78\}, \{413, 78\}, \{413, 78\}, \{413, 78\}, \{413, 78\}, \{413, 78\}, \{413, 78\}, \{413, 78\}, \{413, 78\}, \{413, 78\}, \{413, 78\}, \{413, 78\}, \{413, 78\}, \{413, 78\}, \{413, 78\}, \{413, 78\}, \{413, 78\}, \{413, 78\}, \{413, 78\}, \{413, 78\}, \{413, 78\}, \{413, 78\}, \{413, 78\}, \{413, 78\}, \{413, 78\}, \{413, 78\}, \{413, 78\}, \{413, 78\}, \{413, 78\}, \{413, 78\}, \{413, 78\}, \{413, 78\}, \{413, 78\}, \{413, 78\}, \{413, 78\}, \{413, 78\}, \{413, 78\}, \{413, 78\}, \{413, 78\}, \{413, 78\}, \{413, 78\}, \{413, 78\}, \{413, 78\}, \{413, 78\}, \{413, 78\}, \{413, 78\}, \{413, 78\}, \{413, 78\}, \{413, 78\}, \{413, 78\}, \{413, 78\}, \{413, 78\}, \{413, 78\}, \{413, 78\}, \{413, 78\}, \{413, 78\}, \{413, 78\}, \{413, 78\}, \{413, 78\}, \{413, 78\}, \{413, 78\}, \{413, 78\}, \{413, 78\}, \{413, 78\}, \{413, 78\}, \{413, 78\}, \{413, 78\}, \{413, 7
                                                           \{413,80\}, \{413,81\}, \{413,85\}, \{413,86\}, \{413,88\}, \{413,89\}, \{413,90\}, \{413,94\}, \{413,101\},
                                                           \{413, 104\}, \{413, 105\}, \{413, 106\}, \{413, 108\}, \{413, 109\}, \{413, 111\}, \{413, 115\}, \{413, 117\},
                                                            \{413, 118\}, \{413, 119\}, \{413, 123\}, \{413, 124\}, \{414, 19\}, \{414, 23\}, \{414, 31\}, \{414, 33\}, \{414, 43\}, \{414, 43\}, \{414, 414, 414\}, \{414, 414, 414\}, \{414, 414, 414\}, \{414, 414, 414\}, \{414, 414, 414\}, \{414, 414, 414\}, \{414, 414, 414\}, \{414, 414, 414\}, \{414, 414, 414\}, \{414, 414, 414\}, \{414, 414, 414\}, \{414, 414, 414\}, \{414, 414, 414\}, \{414, 414, 414\}, \{414, 414, 414\}, \{414, 414, 414\}, \{414, 414, 414\}, \{414, 414, 414\}, \{414, 414, 414\}, \{414, 414, 414\}, \{414, 414, 414\}, \{414, 414, 414\}, \{414, 414, 414\}, \{414, 414, 414\}, \{414, 414, 414\}, \{414, 414, 414\}, \{414, 414, 414\}, \{414, 414, 414\}, \{414, 414, 414\}, \{414, 414, 414\}, \{414, 414, 414\}, \{414, 414, 414\}, \{414, 414, 414\}, \{414, 414, 414\}, \{414, 414, 414\}, \{414, 414, 414\}, \{414, 414, 414\}, \{414, 414, 414\}, \{414, 414, 414\}, \{414, 414, 414\}, \{414, 414, 414\}, \{414, 414, 414\}, \{414, 414, 414\}, \{414, 414, 414\}, \{414, 414, 414\}, \{414, 414, 414\}, \{414, 414, 414\}, \{414, 414, 414\}, \{414, 414, 414\}, \{414, 414, 414\}, \{414, 414, 414\}, \{414, 414, 414\}, \{414, 414, 414\}, \{414, 414, 414\}, \{414, 414, 414\}, \{414, 414, 414\}, \{414, 414, 414\}, \{414, 414\}, \{414, 414\}, \{414, 414\}, \{414, 414\}, \{414, 414\}, \{414, 414\}, \{414, 414\}, \{414, 414\}, \{414, 414\}, \{414, 414\}, \{414, 414\}, \{414, 414\}, \{414, 414\}, \{414, 414\}, \{414, 414\}, \{414, 414\}, \{414, 414\}, \{414, 414\}, \{414, 414\}, \{414, 414\}, \{414, 414\}, \{414, 414\}, \{414, 414\}, \{414, 414\}, \{414, 414\}, \{414, 414\}, \{414, 414\}, \{414, 414\}, \{414, 414\}, \{414, 414\}, \{414, 414\}, \{414, 414\}, \{414, 414\}, \{414, 414\}, \{414, 414\}, \{414, 414\}, \{414, 414\}, \{414, 414\}, \{414, 414\}, \{414, 414\}, \{414, 414\}, \{414, 414\}, \{414, 414\}, \{414, 414\}, \{414, 414\}, \{414, 414\}, \{414, 414\}, \{414, 414\}, \{414, 414\}, \{414, 414\}, \{414, 414\}, \{414, 414\}, \{414, 414\}, \{414, 414\}, \{414, 414\}, \{414, 414\}, \{414, 414\}, \{414, 414\}, \{414, 414\}, \{414, 414\}, \{414, 414\}, \{414, 414\}, \{414, 414\}, \{414, 414\}, \{414, 414\}, \{414, 414\}, \{414, 414\}, \{414, 414\}, \{414, 414\}, \{414, 414\}, \{414, 414\}, \{414, 414\}, \{414, 414\}, \{414, 414\}, \{414, 414\}, \{414, 414\}, \{414, 414\}, \{414, 414\}, \{414, 414\}, \{414, 414\}, \{
                                                            \{414, 51\}, \{414, 53\}, \{414, 67\}, \{414, 68\}, \{414, 69\}, \{414, 71\}, \{414, 106\}, \{414, 123\}, \{414, 124\}\}
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                        £03
                                           The example is now corrected:
       In[@]:= Position[fullgenomeSNPEFF[All, 3], 723168][1, 1];
                                            fullgenomeSNPEFF[%, 106;; 110]
Out[ • ]=
                                             \{.:0,1:1:26:26,0,0:10,0:10:99:0,249,
                                                  0:3,3:6:3:0,3,0:8,0:8:99:0,252,.:4,0:4:99:0,108
                                            Extracting the ref vs alt call in genomeSNPEFF:
       in[*]:= genomeSNPEFF = Table[".", {i, 1, Length[fullgenomeSNPEFF]}, {j, 1, topos - droppos}];
       In[@]:= genomeSNPEFF[[1]] // Length
Out[ • ]=
                                            110
       In[*]:= For[i = 1, i ≤ Length[fullgenomeSNPEFF], i++,
                                                  For [j = 1 + droppos, j \le topos, j++,
                                                          temp = StringSplit[fullgenomeSNPEFF[i, j], {":"}];
                                                          genomeSNPEFF[[i, j - droppos]] = temp[[1]];
                                                 ]
                                            ]
       In[*]:= genomeSNPEFF // Length
Out[ • ]=
                                            414
       In[*]:= Tally[Select[genomeSNPEFF[[1]], # # "." &]]
```

All mutant sites and the tally of mutations in them (ignore the error, which comes from sorting so that "." is last):

```
In[*]:= summarytable =
```

```
Table[{fullgenomeSNPEFF[i, 1], fullgenomeSNPEFF[i, 3], fullgenomeSNPEFF[i, 8],
  fullgenomeSNPEFF[i, 9], SortBy[Tally[genomeSNPEFF[i]]], ToExpression[#] &]},
 {i, 1, Length[genomeSNPEFF]}];
```

```
ToExpression::sntx: Invalid syntax in or before ". ".
ToExpression::sntx: Invalid syntax in or before ". ".
ToExpression::sntx: Invalid syntax in or before ". ".
```

General::stop: Further output of ToExpression::sntx will be suppressed during this calculation.

This does not change the number of genes with mutations detected (i.e., all tallies with only two elements include the ref and alt allele):

In[⊕]:= Select[summarytable, Length[#[5]]] ≤ 2 &] // MatrixForm

Out[•]//MatrixForm=

```
chrII 634214 KTR3
                        HIGH
                               \{\{0, 108\}, \{1, 2\}\}
 chrII 682 726 SWC5 MODERATE {{0, 109}, {1, 1}}
 chrII 683 908 PBP2 MODERATE {{0, 109}, {1, 1}}
 chrXI 415 085 URB1 MODERATE {{0, 109}, {1, 1}}
 chrXI 529178 TRK2 MODERATE {{0, 109}, {1, 1}}
chrXII 672772 RED1
                               \{\{0, 109\}, \{1, 1\}\}
                        LOW
chrXIII 53 046 ZDS2
                        LOW
                               \{\{0, 109\}, \{1, 1\}\}
chrXIII 533 259 RRB1
                        LOW
                               \{\{0, 109\}, \{1, 1\}\}
chrXIII 547166 RIM11 MODERATE {{0, 109}, {1, 1}}
chrXIII 585173 INP2
                        HIGH {{0, 109}, {1, 1}}
chrXIII 914235 FET4
                        HIGH {{0, 109}, {1, 1}}
chrXIV 195289 ATG2
                        LOW {{0, 109}, {1, 1}}
```

Revised gene list

In[@]:= genelist = Sort[Union[fullgenomeSNPEFF[All, 8]]] Length[%]

Out[•]=

{ABP1, ACB1, ACC1, ADE6, ADR1, AFT1, AGP1, AHK1, ALD5, ALY2, APC2, ARO3, ASN1, ATG11, ATG2, ATG39, BDS1, BIR1, BLM10, BNA6, BNI1, BNI4, BRR2, BSD2, BUL1, BUL2, CAJ1, CCC2, CCR4, CCW12, CDC25, CIC1, CLB3, CLN3, CNE1, COG1, COG3, COI1, COQ1, CPA1, CSM3, CUE3, CWC22, CWH41, CYK3, DAL2, DBF20, DBP10, DDR48, DNF1, DOA4, DOT6, DSC2, DSE4, DUS4, EBS1, ECM14, ECM22, EFR3, ELO3, ERB1, ERG1, ERG27, ERG7, ERG9, FAS2, FAU1, FCY2, FET4, FIG2, FIG4, FLC2, FMP52, FRE6, FSF1, FYV10, FYV6, FZF1, GAS4, GAT1, GCD2, GID12, GPB1, GPB2, GPH1, GSC2, HBT1, HEH2, HHF2, HIS4, HMF1, HO, HOB2, HRT3, HSL1, HSP104, HUL5, HXT13, IDP1, IES3, ILV2, IMA1, INP2, IOC4, IPI3, IRC20, IRC8, IST2, ISU1, ITC1, KAR3, KGD1, KIN1, KIN82, KRE5, KSP1, KTR3, LAM1, LAM6, LOS1, LRG1, MAM3, MAS2, MCK1, MCM5, MCT1, MDH2, MDL2, MDM30, MET10, MET30, MEX67, MHP1, MKT1, MLF3, MLH3, MLS1, MMS4, MNN9, MOT1, MPA43, MRL1, MRPL36, MSC6, MSS2, MTR4, MYG1, MYO2, NAM8, NAR1, NFT1, NGG1, NIP100, NPL3, NSR1, NUC1, NUG1, OAF1, OCA4, OCA5, OPI1, OSH2, PBP2, PBS2, PCL5, PDA1, PDR1, PDR10, PDX1, PET111, PET127, PEX2, PFK26, PHM7, PH023, PH081, PH084, PIB1, PKP1, PLB2, PMA1, POL3, PPQ1, PRC1, PRI1, PRI2, PRP2, PRP8, PTK2, PYK2, QRI7, RAD16, RAD17, RAD26, RBK1, RCE1, RCK2, RCO1, REC102, RED1, RHO1, RIM11, RIM21, RKM3, RMP1, RNY1, ROG1, RPA135, RPI1, RPL22A, RPL2A, RPL39, RPS15, RPS2, RPT5, RRB1, RRG7, RRP46, RRP6, RSC1, RSE1, RTC2, SAK1, SAP4, SBE2, SEC8, SET2, SFP1, SHP1, SIW14, SKN1, SMC2, SMY2, SNF7, SNT2, SPC105, SP077, SPS100, SRP40, SSE1, SSK2, SSQ1, SSZ1, STE5, STP3, STT3, SUB2, SUC2, SUM1, SWC5, SWS2, SWT1, SYF2, SYG1, TAF8, TAH11, TAO3, TCO89, TFB1, TFG1, TIM54, TOM1, TOP1, TOP2, TOS3, TPS3, TRK2, TRL1, TUS1, UBC1, UBP5, UBR2, UBX7, UGA3, UGA4, UIP5, ULS1, UME6, URA2, URB1, USA1, UTP21, UTP8, VBA1, VCX1, VHR2, VID22, VPS13, VPS74, VRP1, VTC1, VTC3, VTC4, VTC5, WAR1, YAP6, YBR242W, YBR292C, YCF1, YCK1, YCT1, YDL199C, YDR003W-A, YDR157W, YDR541C, YEF1, YGR126W, YGR130C, YGR266W, YHC1, YHC3, YIH1, YIL092W, YLL066W-B, YLR108C, YLR296W, YML119W, YMR027W, YMR317W, YOR029W, YOR1, YOR296W, YOR343C, YPK1, YPL113C, YPR078C, YPR089W, YPT6, YRA1, YRB2, ZDS2, ZRT1, ZRT3}

Out[•]=

333

The systematic names:

In[@]:= genelistSYS = Sort[Union[fullgenomeSNPEFF[All, 11]]] Length[%]

Out[•]=

```
{YAL021C, YAL040C, YAL051W, YAL053W, YAL056W, YAL058W, YBL058W, YBR003W, YBR030W,
YBR086C, YBR098W, YBR114W, YBR122C, YBR147W, YBR172C, YBR205W, YBR231C, YBR233W,
YBR242W, YBR273C, YBR290W, YBR292C, YCL025C, YCL030C, YCR036W, YCR059C, YCR088W,
YCR089W, YCR091W, YCR095C, YDL019C, YDL031W, YDL066W, YDL073W, YDL084W, YDL102W,
YDL104C, YDL107W, YDL117W, YDL128W, YDL155W, YDL170W, YDL176W, YDL199C, YDL210W,
YDL223C, YDL227C, YDL240W, YDR003W-A, YDR035W, YDR069C, YDR089W, YDR103W,
YDR122W, YDR135C, YDR157W, YDR176W, YDR177W, YDR206W, YDR207C, YDR216W, YDR259C,
YDR270W, YDR310C, YDR311W, YDR313C, YDR351W, YDR372C, YDR381C-A, YDR381W,
YDR432W, YDR457W, YDR458C, YDR541C, YEL041W, YEL069C, YER004W, YER006W, YER048C,
YER056C, YER057C, YER064C, YER072W, YER073W, YER088C, YER129W, YER144C,
YER156C, YER157W, YER166W, YER172C, YER178W, YER183C, YFL007W, YFL021W,
YFR030W, YFR031C, YFR031C-A, YFR047C, YGL008C, YGL013C, YGL022W, YGL027C,
YGL071W, YGL093W, YGL110C, YGL123W, YGL131C, YGL133W, YGL141W, YGL144C,
YGL179C, YGL223C, YGL229C, YGL254W, YGL255W, YGR032W, YGR037C, YGR056W, YGR061C,
YGR083C, YGR095C, YGR126W, YGR128C, YGR129W, YGR130C, YGR143W, YGR159C,
YGR175C, YGR186W, YGR193C, YGR233C, YGR266W, YGR278W, YGR281W, YGR287C, YHL020C,
YHL029C, YHR024C, YHR052W, YHR064C, YHR071W, YHR072W, YHR082C, YHR086W, YHR132C,
YHR135C, YHR139C, YHR155W, YHR165C, YHR190W, YIL042C, YIL046W, YIL047C, YIL063C,
YIL092W, YIL097W, YIL107C, YIL119C, YIL125W, YIL129C, YIL162W, YIR008C, YIR029W,
YJL012C, YJL042W, YJL050W, YJL051W, YJL054W, YJL059W, YJL084C, YJL087C, YJL128C,
YJL130C, YJL168C, YJL189W, YJL208C, YJL210W, YJR035W, YJR046W, YJR059W, YJR089W,
YKL014C, YKL045W, YKL101W, YKL126W, YKL175W, YKL205W, YKR044W, YKR050W, YKR092C,
YKR103W, YLL026W, YLL040C, YLL051C, YLL055W, YLL066W-B, YLR024C, YLR025W,
YLR052W, YLR061W, YLR072W, YLR097C, YLR100W, YLR108C, YLR110C, YLR127C, YLR145W,
YLR228C, YLR247C, YLR248W, YLR261C, YLR263W, YLR274W, YLR296W, YLR298C, YLR310C,
YLR312C, YLR329W, YLR337C, YLR341W, YLR368W, YLR369W, YLR372W, YLR373C, YLR375W,
YLR403W, YLR405W, YLR409C, YLR425W, YML029W, YML049C, YML076C, YML109W, YML111W,
YML114C, YML119W, YML123C, YMR006C, YMR027W, YMR044W, YMR048W, YMR049C, YMR075W,
YMR088C, YMR108W, YMR131C, YMR139W, YMR163C, YMR173W, YMR212C, YMR257C, YMR261C,
YMR274C, YMR275C, YMR297W, YMR317W, YMR319C, YNL030W, YNL032W, YNL074C, YNL081C,
YNL085W, YNL088W, YNL097C, YNL117W, YNL133C, YNL182C, YNL233W, YNL240C, YNL242W,
YNL249C, YNL271C, YNL294C, YNL307C, YNL325C, YNR011C, YNR016C, YNR031C, YNR067C,
YOL006C, YOL040C, YOL060C, YOL073C, YOL084W, YOL126C, YOL132W, YOL164W, YOR001W,
YOR017W, YOR029W, YOR117W, YOR166C, YOR191W, YOR221C, YOR271C, YOR296W, YOR303W,
YOR305W, YOR326W, YOR328W, YOR336W, YOR343C, YOR347C, YOR354C, YOR368W, YOR371C,
YPL019C, YPL050C, YPL082C, YPL106C, YPL113C, YPL123C, YPL135W, YPL164C, YPL169C,
YPL174C, YPL179W, YPL180W, YPL231W, YPL270W, YPR010C, YPR049C, YPR055W, YPR078C,
YPR079W, YPR089W, YPR111W, YPR117W, YPR141C, YPR145W, YPR160W, YPR165W}
```

```
Plotting the depth of coverage at that site:
```

```
In[*]:= Mean[fullgenomeSNPEFF[All, 129]] // N
Out[ • ]=
       933.978
 In[*]:= Max[fullgenomeSNPEFF[[All, 129]]]
Out[ • ]=
       2469
 In[*]:= Histogram[fullgenomeSNPEFF[All, 129], {50}, PlotRange → {{0, 3000}, Automatic}]
Out[ • ]=
       50
       40
       30
       20
       10
        0 |-
                  500
```

Relating mutations to phenotypes (metal resistance, phosphorous metabolism, DNA repair)

Metal Resistance:

The list of genes affecting metal resistance was downloaded from SGD (https://yeastgenome.org/observable/APO:0000090) on 23 May 2025:

```
In[*]:= metallist = Import[
          "/Users/otto/Documents/Students/AnnaBazzicalupo/MetalAdaptation/Genomics/
            metal_resistance_annotations_23May2025.txt", "tsv"];
 In[*]:= metalhead = metallist[[2]]
Out[ • ]=
      {Gene, Gene Systematic Name, Phenotype, Experiment Type, Experiment Type Category,
       Mutant Information, Strain Background, Chemical, Details, Reference}
 In[*]:= metallist = Drop[metallist, 2];
      Here we will use the Gene Systematic Name for comparison:
 In[@]:= metalgenenames = Union[metallist[All, 2]];
      Length[%]
Out[ • ]=
      1791
```

Of the 333 genes bearing mutations, 121 (36.34%) are annotated as having metal tolerance:

```
In[*]:= Intersection[genelistSYS, metalgenenames]
      {Length[genelistSYS], Length[%], Length[%] / Length[genelistSYS] // N // PercentForm}
Out[ • ]=
      {YAL021C, YAL040C, YAL051W, YAL053W, YAL056W, YBL058W, YBR098W, YBR114W, YBR122C,
       YBR172C, YBR231C, YBR233W, YBR273C, YBR290W, YCL030C, YCR036W, YCR088W,
       YDL066W, YDL073W, YDL128W, YDL155W, YDL176W, YDR069C, YDR089W, YDR103W,
       YDR122W, YDR135C, YDR176W, YDR207C, YDR216W, YDR270W, YDR310C, YDR351W,
       YDR372C, YDR432W, YDR457W, YDR458C, YER072W, YER129W, YER156C, YER178W,
       YFL007W, YGL027C, YGL071W, YGL141W, YGL223C, YGL254W, YGL255W, YGR037C,
       YGR056W, YHR064C, YHR082C, YHR086W, YHR135C, YIL047C, YIL097W, YIL125W,
       YJL012C, YJL042W, YJL128C, YJL189W, YJL208C, YJR059W, YKL101W, YKL126W,
       YKL175W, YKR050W, YKR092C, YLL040C, YLL051C, YLR024C, YLR025W, YLR061W,
       YLR072W, YLR228C, YLR248W, YLR261C, YLR337C, YLR341W, YLR368W, YLR369W,
       YLR372W, YLR373C, YLR375W, YLR403W, YLR425W, YML111W, YML123C, YMR075W,
       YMR108W, YMR139W, YMR173W, YMR261C, YMR275C, YMR319C, YNL032W, YNL074C,
       YNL097C, YNL233W, YNL242W, YNL271C, YNL294C, YNL307C, YNL325C, YNR031C,
       YOL006C, YOL060C, YOR221C, YOR296W, YOR326W, YOR347C, YPL019C, YPL050C,
       YPL106C, YPL123C, YPL135W, YPL179W, YPL180W, YPR049C, YPR141C, YPR160W}
Out[ • ]=
      {333, 121, 36.34%}
      Appending "metal gene resistance" phenotype information to the data:
 In[•]:= For[i = 1; newcolumn = {}, i ≤ Length[fullgenomeSNPEFF], i++,
       nameSYS = fullgenomeSNPEFF[[i, 11]];
       If[MemberQ[metalgenenames, nameSYS],
        AppendTo[newcolumn, "Metal"], AppendTo[newcolumn, "No"]]
 In[*]:= fullgenomeSNPEFF = Transpose[Append[Transpose[fullgenomeSNPEFF], newcolumn]];
      Phosphorus metabolic process:
      The list of genes affecting phosphorous metabolism was downloaded from Amigo 2 (https://amigo.ge-
      neontology.org/amigo/term/GO:0006793) on 17 June2025:
 In[*]:= pholist = Import[
          "/Users/otto/Documents/Students/AnnaBazzicalupo/MetalAdaptation/Genomics/
            phosphate metabolism annotations 17Jun2025.tsv", "tsv"];
 In[*]:= phohead = pholist[[1]]
Out[ • ]=
      {gene, gene_id, gene_name, go_id, evidence}
 In[*]:= pholist = Drop[pholist, 1];
```

The gene name but not the systematic name is listed:

```
In[*]:= phogenenames = pholist[All, 1];
      Length[%]
Out[ • ]=
      480
      Of the 333 genes bearing mutations, 27 (8.1%) are annotated as affecting phosphorous metabolism:
 In[*]:= Intersection[genelist, phogenenames]
       {Length[genelist], Length[%], Length[%] / Length[genelist] // N // PercentForm}
Out[ • ]=
       {ACC1, ADE6, AFT1, BNA6, ERG9, FIG4, IDP1, KRE5, MPA43, OPI1, PDA1, PDX1, PFK26, PHO81,
       PH084, PLB2, PYK2, RBK1, SIW14, URA2, VTC1, VTC3, VTC4, VTC5, YBR242W, YEF1, YPK1}
Out[ • ]=
      {333, 27, 8.108%}
      Appending phosphorous metabolism phenotype information to the data:
 In[•]:= For[i = 1; newcolumn = {}, i ≤ Length[fullgenomeSNPEFF], i++,
        name = fullgenomeSNPEFF[i, 8];
       If[MemberQ[phogenenames, name],
         AppendTo[newcolumn, "PHO"], AppendTo[newcolumn, "No"]]
      ]
 In[*]:= fullgenomeSNPEFF = Transpose[Append[Transpose[fullgenomeSNPEFF], newcolumn]];
      DNA Repair:
      The list of genes affecting DNA repair was downloaded from Amigo 2 (https://amigo.geneontolo-
      gy.org/amigo/term/GO:0006281) on 17 June 2025:
 In[*]:= repairlist = Import[
          "/Users/otto/Documents/Students/AnnaBazzicalupo/MetalAdaptation/Genomics/
            dna_repair_amigo_17Jun2025.tsv", "tsv"];
 In[*]:= repairhead = repairlist[[1]]
Out[ • ]=
       {gene, gene_id, gene_name, go_id, evidence}
 In[*]:= repairlist = Drop[repairlist, 1];
 In[*]:= repairlist[2]
Out[ • ]=
       {RFC2, SGD:S000003829,
       Subunit of heteropentameric Replication factor C (RF-C), GO:0006298, TAS}
      The gene name but not the systematic name is listed:
 In[@]:= repairgenenames = Union[repairlist[All, 1]]];
      Length[%]
Out[ • ]=
      322
```

Of the 333 genes bearing mutations, 23 (6.9%) are annotated as affecting DNA repair:

```
In[*]:= Intersection[genelist, repairgenenames]
      {Length[genelist], Length[%], Length[%] / Length[genelist] // N // PercentForm}
Out[ • ]=
      {BLM10, CSM3, DDR48, FYV6, IRC20, MCK1, MCM5, MLH3, MMS4, PDR10, POL3, PRI2,
       RAD16, RAD17, RAD26, RSC1, SET2, SUB2, TAH11, TFB1, ULS1, VID22, YRA1}
Out[ • ]=
      \{333, 23, 6.907\%\}
      Appending DNA repair phenotype information to the data:
 In[*]:= For[i = 1; newcolumn = {}, i ≤ Length[fullgenomeSNPEFF], i++,
       name = fullgenomeSNPEFF[[i, 8]];
       If[MemberQ[repairgenenames, name],
        AppendTo[newcolumn, "Repair"], AppendTo[newcolumn, "No"]]
      1
 In[a]:= fullgenomeSNPEFF = Transpose[Append[Transpose[fullgenomeSNPEFF], newcolumn]];
      Exporting
      For printing, we want a shorter header table:
 in[*]:= newheader = Table[StringSplit[header[i]], "_"][1], {i, 1, Length[header]}];
      newheader = Flatten[AppendTo[newheader, {"Metal resistance gene (SGD)",
           "Phosphorous metabolism (Amigo 2)", "DNA repair (SGD)"}]]
Out[ • ]=
      {CHROM, chr, POS, REF, ALT, QUAL, FORMAT, gene, effect, mut, gene, base, aa, ANN,
       OLY077, CdBM23, CdBM25, CdBM26, CdBM29, CdBM30, CdBM32, CdBM36, CdBM37, CdBM39,
       CdBM42, CdBM43, CdBM44, CdBM45, CdBM46, CdBM47, CdBM48, CoBM12, CoBM14, CoBM15,
       COBM16, COBM17, COBM18, COBM1, COBM20, COBM21, COBM2, COBM3, COBM4, COBM5, COBM6,
       CoBM7, CoBM8, CuBM10, CuBM11, CuBM12, CuBM13, CuBM14, CuBM15, CuBM16, CuBM17,
       CuBM18, CuBM3, CuBM4, CuBM6, CuBM7, CuBM8, CuBM9, MnBM12, MnBM13, MnBM14, MnBM15,
       MnBM16, MnBM17, MnBM18, MnBM20, MnBM21, MnBM23, MnBM24, MnBM25, MnBM27, MnBM28,
       MnBM29, MnBM31, MnBM32, MnBM34, MnBM38, MnBM39, MnBM42, MnBM44, NiBM11, NiBM12,
       NiBM14, NiBM16, NiBM17, NiBM21, NiBM22, NiBM24, NiBM25, NiBM27, NiBM28, NiBM29,
       NiBM30, NiBM4, NiBM6, NiBM8, NiBM9, ZnBM11, ZnBM12, ZnBM15, ZnBM16, ZnBM17,
       ZnBM19, ZnBM22, ZnBM23, ZnBM25, ZnBM28, ZnBM29, ZnBM31, ZnBM34, ZnBM37, ZnBM38,
       ZnBM39, ZnBM41, ZnBM42, ZnBM43, ZnBM44, ZnBM45, ZnBM46, ZnBM47, AC, AF, AN,
       BaseQRankSum, DP, FS, MLEAC, MLEAF, MQ, MQRankSum, QD, ReadPosRankSum, SOR, LOF,
       Metal resistance gene (SGD), Phosphorous metabolism (Amigo 2), DNA repair (SGD)}
```

First recalculating the summary table:

```
In[*]:= genomeSNPEFF = Table[-1, {i, 1, Length[fullgenomeSNPEFF]}, {j, 1, topos - droppos}];
       For[i = 1, i ≤ Length[fullgenomeSNPEFF], i++,
        For [j = 1 + droppos, j \le topos, j++,
         temp = StringSplit[fullgenomeSNPEFF[i, j], {":"}];
         temp2 = temp[[1]];
         genomeSNPEFF[[i, j - droppos]] = temp2;
        ]
       ]
       The following sites are "1"s in OLY077 and so we swap 1's for 0's and call these revertants
 In[@]:= Position[genomeSNPEFF, _ ? (#[1]] == "1" &) ] [All, 1]
       ••• Part: Part specification List[1] is longer than depth of object. 1
       ••• Part: Part specification List[1] is longer than depth of object. 1
       Part: Part specification O[1] is longer than depth of object.
       😶 General: Further output of Part::partd will be suppressed during this calculation. 🕡
Out[ • ]=
       {155, 234, 322, 323, 324, 325}
       Swapping 0<->1 and calling "." 0 (like OLY077):
 In[\bullet]:= For[i = 1, i \leq Length[%], i++,
        genomeSNPEFF[%[i]] =
         Mod[ToExpression[StringReplace[genomeSNPEFF[%[i]]], "." → "1"]] + 1, 2]
       1
       Swapped variants positions:
 In[*]:= fullgenomeSNPEFF[[%%, {1, 3, 8}]]
Out[ • ]=
       {{chrVII, 753 296, YGR130C}, {chrXI, 613 636, SRP40}, {chrXIII, 908 159, YMR317W},
        {chrXIII, 908168, YMR317W}, {chrXIII, 908215, YMR317W}, {chrXIII, 908221, YMR317W}}
       All remaining "." are treated as identical to OLY077 (not mutant) and 0/1 treated as numbers (only in
       genomeSNPEFF, summarytable and fullgenomeSNPEFF remain the same):
 In[*]:= genomeSNPEFF = genomeSNPEFF /. "." → "0";
       genomeSNPEFF = ToExpression /@ genomeSNPEFF;
       Number of unique mutations:
 In[*]:= Length[genomeSNPEFF]
Out[ • ]=
       414
       Number of mutational hits:
 In[*]:= Total[Total[genomeSNPEFF]]
Out[ • ]=
       506
```

As expected (once revertants were recoded), OLY077 has no mutations

```
In[*]:= Total[genomeSNPEFF[All, 1]]]
Out[ • ]=
       Number of lines per unique mutation:
 In[@]:= Sort[Tally[Sum[genomeSNPEFF[All, i], {i, 1, 110}]]]
Out[ • ]=
       \{\{1, 384\}, \{2, 18\}, \{3, 3\}, \{4, 1\}, \{5, 1\}, \{6, 3\}, \{8, 1\}, \{9, 1\}, \{10, 1\}, \{23, 1\}\}\}
       Genes with excessive SNP mutations (≥5) include:
 In[•]:= toopoly = Position[genomeSNPEFF, _? (Total[#] ≥ 5 &)] [All, 1]
Out[ • ]=
       {68, 155, 159, 236, 322, 323, 324, 325}
      The genotype call for OLY077 at these sites:
 In[*]:* fullgenomeSNPEFF[toopoly, Position[newheader, "OLY077"][1, 1]]] // MatrixForm
Out[ • ]//MatrixForm=
         .:3,0:3:99:0,100
        1:0,21:21:99:945,0
         0:8,0:8:99:0,175
        0:13,0:13:99:0,226
        1:0,10:10:99:450,0
        1:0,10:10:99:450,0
         1:0,9:9:99:406,0
         1:0,9:9:99:406,0
 In[*]:= consider = summarytable[[toopoly]];
       consider // MatrixForm
Out[ • ]//MatrixForm=
         chrIV 815 428
                             NGG1
                                      MODERATE \{\{0, 38\}, \{1, 10\}, \{., 62\}\}
                                      MODERATE {{0, 23}, {1, 81}, {., 6}}
         chrVII 753296 YGR130C
         chrVII 869872
                             TFG1
                                      MODERATE {{0, 73}, {1, 6}, {., 31}}
         chrXII
                   5683 YLL066W-B
                                         HIGH
                                                  \{\{0, 46\}, \{1, 8\}, \{., 56\}\}\
                                                 \{\{0, 5\}, \{1, 80\}, \{., 25\}\}
        chrXIII 908 159 YMR317W
                                         HIGH
        chrXIII 908168 YMR317W
                                         HIGH
                                                  \{\{0, 6\}, \{1, 63\}, \{., 41\}\}
        chrXIII 908215
                           YMR317W
                                         LOW
                                                  \{\{0, 9\}, \{1, 60\}, \{., 41\}\}
        chrXIII 908221 YMR317W
                                         LOW
                                                  \{\{0,6\},\{1,68\},\{.,36\}\}
       Depth of coverage for these genes is higher on average:
 In[*]:= fullgenomeSNPEFF[toopoly, {8, 129}]
       Mean[%[All, 2]] // N
Out[ • ]=
       {{NGG1, 642}, {YGR130C, 1809}, {TFG1, 798}, {YLL066W-B, 607},
        {YMR317W, 1959}, {YMR317W, 1960}, {YMR317W, 1836}, {YMR317W, 1789}}
Out[ • ]=
       1425.
```

```
Mapping quality is also poorer on average:
 In[*]:= fullgenomeSNPEFF[toopoly, {8, 133}]
      Mean[%[All, 2]] // N
      Length[Select[fullgenomeSNPEFF[All, 133], # < % &]] /</pre>
          Length[fullgenomeSNPEFF[[All, 133]] // N // PercentForm
Out[ • ]=
      {{NGG1, 60.}, {YGR130C, 60.}, {TFG1, 60.}, {YLL066W-B, 57.77},
       {YMR317W, 57.88}, {YMR317W, 57.87}, {YMR317W, 56.28}, {YMR317W, 55.95}}
Out[ • ]=
      58.2188
Out[ • ]//PercentForm=
      1.691%
 In[*]:= considergenes = Union[consider[All, 3]]
      % // Length
Out[ • ]=
      {NGG1, TFG1, YGR130C, YLL066W-B, YMR317W}
Out[•]=
 In[*]:= Position[newheader, "Metal resistance gene (SGD)"] [1, 1]
Out[ • ]=
      139
 In[*]:= toprintmultiple =
         Sort[Table[{fullgenomeSNPEFF[[toopoly[i]], 8], fullgenomeSNPEFF[[toopoly[i]], 11]],
            StringJoin[fullgenomeSNPEFF[toopoly[i]], 1]], ".",
             ToString[fullgenomeSNPEFF[[toopoly[i]], 3]]],
            fullgenomeSNPEFF[toopoly[i], 12], fullgenomeSNPEFF[toopoly[i], 13],
            fullgenomeSNPEFF[[toopoly[i]], 9]], fullgenomeSNPEFF[[toopoly[i]],
             Position[newheader, "Metal resistance gene (SGD)"][[1, 1]]], fullgenomeSNPEFF[
             toopoly[i], Position[newheader, "Phosphorous metabolism (Amigo 2)"][1, 1]],
            fullgenomeSNPEFF[toopoly[i]], Position[newheader, "DNA repair (SGD)"][1, 1]]],
            summarytable[toopoly[i]], 5], genomeSNPEFF[toopoly[i]]].
             newheader[[1 + droppos ;; topos]]}, {i, 1, Length[toopoly]}]];
      MatrixForm[%]
Out[ • ]//MatrixForm=
                               chrIV.815428
          NGG1
                    YDR176W
                                                                c.977G>A
                                                                                               р.
          TFG1
                    YGR186W
                              chrVII.869872
                                                               c.2099T>C
                                                                                               р.
         YGR130C
                    YGR130C chrVII.753296
                                                          c.543_548delATCGTC
                                                                                           p.Serl
        YLL066W-B YLL066W-B
                              chrXII.5683
                                                    c.85_103delCACACCCACACCCACACAC
                                                                                                p
                   YMR317W chrXIII.908159
                                                         c.797_804delCGGCAACG
         YMR317W
                                                                                                p.
         YMR317W
                    YMR317W
                              chrXIII.908168 c.806_833delCTAGCGTAATTAGTTCAGAAGCTTCATG
                                                                                                p.
         YMR317W
                    YMR317W
                              chrXIII.908215
                                                                c.852G>A
                                                                                               р.
         YMR317W
                    YMR317W
                              chrXIII.908221
                                                                c.858G>A
                                                                                               р.
```

```
In[*]:= Export["/Users/otto/Documents/Students/AnnaBazzicalupo/MetalAdaptation/Genomics/
           toopolymorphic_17June2025.tsv",
         Join[{{"Gene", "Systematic name", "chrom.pos", "base_change",
            "aa_change", "effect", "Metal resistance gene (SGD)",
            "Phosphorous metabolism (Amigo 2)", "DNA repair (SGD)", "# lines",
            "Lines carrying variant not seen in W303 ancestor (OLY077)"}},
          toprintmultiple]];
      The remaining sites:
 In[*]:* rest = Complement[Table[i, {i, 1, Length[fullgenomeSNPEFF]}], toopoly];
 In[*]:= Length[rest] + Length[toopoly] - Length[fullgenomeSNPEFF]
Out[ • ]=
      0
      Including NGG1 and TFG1:
 In[@]:= fullgenomeSNPEFF[[68, 8]]
Out[ • ]=
      NGG1
 In[*]:= fullgenomeSNPEFF[[159, 8]]
Out[ • ]=
      TFG1
 In[*]:= rest = Sort[Join[rest, {68, 159}]];
```

```
In[*]:= toprint = Table[{fullgenomeSNPEFF[rest[i]], 8],
          fullgenomeSNPEFF[[rest[[i]], 11]], StringJoin[fullgenomeSNPEFF[[rest[[i]], 1]], ".",
            ToString[fullgenomeSNPEFF[rest[i]], 3]]], fullgenomeSNPEFF[rest[i]], 12],
           fullgenomeSNPEFF[[rest[[i]], 13]], fullgenomeSNPEFF[[rest[[i]], 9]],
           fullgenomeSNPEFF[rest[i]],
           Position[newheader, "Metal resistance gene (SGD)"][1, 1]], fullgenomeSNPEFF[
            rest[i], Position[newheader, "Phosphorous metabolism (Amigo 2)"][1, 1]],
           fullgenomeSNPEFF[rest[i], Position[newheader, "DNA repair (SGD)"][1, 1]],
          genomeSNPEFF[[rest[i]]].newheader[[1 + droppos;; topos]]}, {i, 1, Length[rest]}];
      MatrixForm[%[1;; 12]]
      (*The first 12 lines shows the nature of parallel SNPs in different lines*)
Out[ • ]//MatrixForm=
       CNE1 YAL058W chrI.37834
                                   c.371C>T p.Thr124Met MODERATE
                                                                                     MnBM14
                                                                           No No
       CNE1 YAL058W chrI.38041
                                  c.578C>T p.Ser193Leu MODERATE
                                                                           No No
                                                                                     MnBM14
       GPB2 YAL056W chrI.40593 c.1335A>C p.Leu445Phe MODERATE Metal No No
                                                                                     CoBM17
       FLC2 YAL053W
                      chrI.48206 c.2308A>G p.Asn770Asp MODERATE Metal No No
                                                                                     MnBM14
       OAF1 YALO51W chrI.48640 c.77C>T p.Ala26Val MODERATE Metal No No
                                                                                     MnBM14
       CLN3 YAL040C chrI.66785 c.736T>G p.Leu246Val MODERATE Metal No No
                                                                                     CoBM3
       CCR4 YAL021C chrI.112637 c.723C>G p.Asp241Glu MODERATE Metal No No
                                                                                     MnBM42
       CCR4 YAL021C chrI.113278 c.82C>T p.Leu28Leu
                                                             LOW
                                                                    Metal No No
                                                                                     MnBM42
       SHP1 YBL058W chrII.112395 c.959A>T p.Glu320Val MODERATE Metal No No
                                                                                     CoBM6
       COQ1 YBR003W chrII.243326 c.518C>A p.Pro173His MODERATE
                                                                                     CuBM15
                                                                           No No
       RKM3 YBR030W chrII.299270 c.979G>C p.Gly327Arg MODERATE
                                                                      No
                                                                           No No CoBM4 + NiBM2
       IST2 YBR086C chrII.421076 c.1966G>T p.Ala656Ser MODERATE
                                                                                     NiBM12
                                                                      No
                                                                           No No
 In[*]:= Length[toprint]
Out[ • ]=
      408
      For each SNP that occurs multiple times, make a separate row:
 In[*]:= namepos = 10; (*Line names*)
      For[i = 1; toappend = {}, i ≤ Length[toprint], i++,
       splits = Length[toprint[i, namepos]];
       If[splits > 1,
        temp = toprint[i, namepos];
        toprint[i, namepos] = toprint[i, namepos][1];
        For [j = 2, j \le splits, j++,
         toadd = toprint[i];
         toadd[namepos] = temp[j];
         toappend = Append[toappend, toadd]
        ]
       ]
      ]
      Sorting then putting the metal first:
```

```
In[@]:= Sort[Join[toprint, toappend]];
      mat =
        Transpose[Join[{Transpose[%[All, namepos]]}}, Transpose[%[All, 1;; namepos - 1]]]]];
      MatrixForm[mat[1;; 12]]
Out[ • ]//MatrixForm=
       CuBM18 ABP1 YCR088W chrIII.265691 c.624A>C p.Leu208Phe MODERATE Metal No
       MnBM42 ACB1 YGR037C chrVII.559830 c.165G>A
                                                       p.Lys55Lys
                                                                      LOW
                                                                             Metal No
                                                                                        No
       ZnBM12 ACC1 YNR016C chrXIV.660428 c.947G>T
                                                       p.Gly316Val MODERATE
                                                                               No
                                                                                    PHO No
       ZnBM15 ACC1 YNR016C chrXIV.660428 c.947G>T p.Gly316Val MODERATE
                                                                                    PHO No
                                                                               No
       ZnBM23 ADE6 YGR061C chrVII.613265 c.2701G>C p.Val901Leu MODERATE
                                                                               No
                                                                                   PHO No
       MnBM23 ADR1 YDR216W chrIV.896208 c.1174C>T p.His392Tyr MODERATE Metal No No
       CoBM17 AFT1 YGL071W chrVII.372520 c.509C>T p.Ser170Phe MODERATE Metal PHO No
       MnBM31 AGP1 YCL025C chrIII.76154 c.1765delA p.Ile589fs
                                                                      HIGH
                                                                               No
                                                                                    No
                                                                                        No
       MnBM31 AGP1 YCL025C chrIII.76159 c.1761G>T p.Leu587Leu
                                                                      LOW
                                                                               No
                                                                                    No No
       CdBM25 AHK1 YDL073W chrIV.329143 c.2531G>A p.Gly844Glu MODERATE Metal No No
       MnBM18 ALD5 YER073W
                             chrV.304100
                                             c.71C>A
                                                       p.Ser24Tyr MODERATE
                                                                               No
                                                                                    No
                                                                                        No
       CoBM16 ALY2 YJL084C chrX.277922
                                            c.80delT
                                                        p.Leu27fs
                                                                     HIGH
                                                                               No
                                                                                    No No
      The length matches that expected from the tally of multiply hit genes:
 In[*]:= Length[mat]
Out[ • ]=
      449
 In[o]:= Sort[Tally[Sum[genomeSNPEFF[All, i], {i, 1, 110}]]][1;; 4]
      %[All, 1].%[All, 2] + (6 + 10) (*Adding NGG1 and TFG1*)
Out[ • ]=
      \{\{1, 384\}, \{2, 18\}, \{3, 3\}, \{4, 1\}\}
Out[ • ]=
      449
 In[*]:= Export[
        "/Users/otto/Documents/Students/AnnaBazzicalupo/MetalAdaptation/Genomics/SNPs 17
          June2025.tsv",
        Join[{{"Line", "Gene", "Systematic name", "chrom.pos", "base_change",
            "aa change", "effect", "Metal resistance gene (SGD)",
```

"Phosphorous metabolism (Amigo 2)", "DNA repair (SGD)"}}, mat]];

Tally by line:

109

Out[•]=

In[@]:= newheader[[16;; 124]] // Length

```
In[*]:= bymetal = Tally[Sort[Join[mat[All, 1]], newheader[[16;; 124]]]]];
      bymetal[All, 2] = bymetal[All, 2] - 1;
      bymetal
Out[ • ]=
      {{CdBM23, 4}, {CdBM25, 2}, {CdBM26, 5}, {CdBM29, 5}, {CdBM30, 9}, {CdBM32, 7},
        {CdBM36, 0}, {CdBM37, 4}, {CdBM39, 6}, {CdBM42, 3}, {CdBM43, 3}, {CdBM44, 3},
        {CdBM45, 1}, {CdBM46, 3}, {CdBM47, 2}, {CdBM48, 1}, {CoBM1, 4}, {CoBM12, 6},
        {COBM14, 1}, {COBM15, 8}, {COBM16, 5}, {COBM17, 6}, {COBM18, 7}, {COBM2, 6},
        {COBM20, 2}, {COBM21, 5}, {COBM3, 8}, {COBM4, 4}, {COBM5, 6}, {COBM6, 8}, {COBM7, 5},
        {CoBM8, 5}, {CuBM10, 2}, {CuBM11, 2}, {CuBM12, 2}, {CuBM13, 2}, {CuBM14, 5},
        {CuBM15, 1}, {CuBM16, 0}, {CuBM17, 3}, {CuBM18, 5}, {CuBM3, 1}, {CuBM4, 3},
        {CuBM6, 2}, {CuBM7, 3}, {CuBM8, 0}, {CuBM9, 4}, {MnBM12, 5}, {MnBM13, 5},
        {MnBM14, 36}, {MnBM15, 1}, {MnBM16, 5}, {MnBM17, 4}, {MnBM18, 3}, {MnBM20, 2},
        {MnBM21, 1}, {MnBM23, 5}, {MnBM24, 4}, {MnBM25, 2}, {MnBM27, 3}, {MnBM28, 1},
        {MnBM29, 3}, {MnBM31, 4}, {MnBM32, 13}, {MnBM34, 0}, {MnBM38, 5}, {MnBM39, 4},
        {MnBM42, 102}, {MnBM44, 1}, {NiBM11, 2}, {NiBM12, 1}, {NiBM14, 0}, {NiBM16, 1},
        {NiBM17, 1}, {NiBM21, 2}, {NiBM22, 0}, {NiBM24, 3}, {NiBM25, 0}, {NiBM27, 1},
        {NiBM28, 2}, {NiBM29, 0}, {NiBM30, 2}, {NiBM4, 4}, {NiBM6, 4}, {NiBM8, 0},
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In[*]:= Sort[Union[mat[All, 2]]] Length[%]

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Out[•]=

331

Gene list with all lines:

In[*]:= Sort[Union[mat[All, 2]]] Length[%]

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Out[•]=

In[*]:= Sort[Union[mat[All, 2]]] Length[%]

Out[•]=

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Out[•]=

331

Gene list without the mutator lines MnBM14 and MnBM42:

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       COI1, COQ1, CSM3, CYK3, DBF20, DBP10, DDR48, DNF1, DOA4, DSE4, DUS4, EBS1, ECM14,
       EFR3, EL03, ERB1, ERG1, ERG27, ERG9, FCY2, FET4, FIG4, FRE6, FSF1, FYV10, FZF1,
       GAT1, GCD2, GPB1, GPB2, HBT1, HHF2, HIS4, HMF1, HO, HSL1, HSP104, HUL5, HXT13,
       IDP1, ILV2, INP2, IOC4, IST2, ISU1, KAR3, KIN1, KIN82, KRE5, KSP1, KTR3, LOS1,
       LRG1, MAM3, MAS2, MCK1, MCT1, MDL2, MET10, MET30, MLS1, MMS4, MPA43, MSC6, MSS2,
       MYG1, MYO2, NAM8, NFT1, NGG1, NPL3, NSR1, NUG1, OCA4, OCA5, PBP2, PBS2, PDR1,
       PDR10, PDX1, PET127, PFK26, PHM7, PH023, PH081, PH084, PKP1, PLB2, PMA1, POL3,
       PPQ1, PRI2, PRP2, PRP8, PTK2, PYK2, RAD16, RBK1, RCE1, RCK2, RCO1, RED1, RHO1,
       RIM11, RIM21, RKM3, RMP1, ROG1, RPL39, RPS15, RPS2, RPT5, RRG7, RRP46, RRP6,
       RSC1, RSE1, RTC2, SAK1, SAP4, SBE2, SET2, SFP1, SHP1, SIW14, SMC2, SMY2, SNF7,
       SNT2, SP077, SPS100, SRP40, SSK2, SSQ1, STE5, STP3, SUB2, SWT1, SYG1, TAF8,
       TC089, TFB1, TFG1, TOM1, TPS3, TRK2, TRL1, TUS1, UBC1, UBP5, UBR2, UBX7, UGA4,
       ULS1, UME6, URB1, VCX1, VHR2, VPS74, VRP1, VTC1, VTC4, VTC5, YAP6, YBR242W,
       YBR292C, YCT1, YDL199C, YDR003W-A, YDR157W, YEF1, YHC1, YIL092W, YLR108C,
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Out[ • ]=
      210
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      s_4Nov2024.tsv):
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 In[*]:= Export["/Users/otto/Documents/Students/AnnaBazzicalupo/MetalAdaptation/Genomics/
           fullgenomeSNPEFF_Afterfiltering_17June2025.tsv", toprint2];
      Correlation of phenotypes with cross-tolerance scores
 In[*]:= crosstolerance = Import[
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Below wincluding NGG1 and TFG1:

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       CoBM8, CuBM10, CuBM11, CuBM12, CuBM13, CuBM14, CuBM15, CuBM17, CuBM18, CuBM3,
       CuBM4, CuBM6, CuBM7, CuBM9, MnBM12, MnBM13, MnBM14, MnBM15, MnBM16, MnBM17,
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       ZnBM16, ZnBM17, ZnBM19, ZnBM22, ZnBM23, ZnBM28, ZnBM29, ZnBM31, ZnBM37,
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        part2 = If[
           MemberQ[Union[Select[mat, #[1] == linenames[i] &] [All, 9]], "PHO"], "PHO", "NO"];
        part3 = If[MemberQ[Union[Select[mat, #[1]] == linenames[i]] &] [All, 10]], "Repair"],
           "Repair", "NO"];
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        AppendTo[phenolist, {part1, part2, part3, part4}]
       ];
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       {CoBM15, Metal, PHO, NO, 0.0458346}, {CoBM16, Metal, PHO, NO, 0.0357453},
       {CoBM17, Metal, PHO, Repair, 0.0597343}, {CoBM18, Metal, PHO, Repair, 0.0413294},
       {CoBM2, Metal, PH0, N0, 0.0706202}, {CoBM20, Metal, PH0, N0, 0.0606192},
       {CoBM21, Metal, PHO, NO, 0.0100054}, {CoBM3, Metal, PHO, NO, 0.0703313},
       {CoBM4, Metal, PH0, N0, 0.0305671}, {CoBM5, Metal, PH0, N0, 0.0741003},
       {CoBM6, Metal, PH0, NO, 0.0712429}, {CoBM7, Metal, NO, Repair, 0.0970299},
       {CoBM8, Metal, PH0, N0, 0.0994528}, {CuBM10, Metal, N0, N0, 0.0342128},
       {CuBM11, Metal, NO, NO, 0.0346486}, {CuBM12, Metal, NO, NO, 0.0424993},
```

```
{CuBM13, Metal, NO, NO, 0.0659556}, {CuBM14, Metal, PHO, NO, 0.0276601},
      {CuBM15, NO, NO, NO, 0.0233375}, {CuBM17, Metal, NO, NO, 0.0318022},
      {CuBM18, Metal, NO, NO, 0.0338427}, {CuBM3, NO, NO, NO, 0.0420565},
      {CuBM4, Metal, NO, Repair, 0.0399191}, {CuBM6, Metal, NO, NO, 0.0296793},
      {CuBM7, Metal, NO, Repair, 0.0409789}, {CuBM9, Metal, NO, Repair, 0.0379313},
      {MnBM12, Metal, NO, Repair, 0.0169373}, {MnBM13, Metal, NO, NO, 0.00492092},
      {MnBM14, Metal, PHO, Repair, 0.0327898}, {MnBM15, NO, NO, NO, 0.0113796},
      {MnBM16, Metal, NO, Repair, 0.00232012}, {MnBM17, Metal, PHO, NO, 0.0149786},
      {MnBM18, NO, NO, NO, 0.0129248}, {MnBM20, NO, NO, NO, -0.00458364},
      {MnBM21, NO, NO, NO, -0.000900768}, {MnBM23, Metal, NO, NO, 0.0301072},
      {MnBM24, NO, NO, Repair, 0.011251}, {MnBM25, NO, NO, Repair, 0.0389341},
      {MnBM27, NO, NO, NO, -0.00624868}, {MnBM28, Metal, NO, NO, -0.0036978},
      {MnBM29, Metal, NO, NO, 0.0114235}, {MnBM31, Metal, NO, NO, 0.0238223},
      {MnBM32, Metal, NO, NO, 0.024686}, {MnBM38, Metal, PHO, NO, 0.0131807},
      \{MnBM39, Metal, NO, NO, 0.038027\}, \{MnBM42, Metal, PHO, Repair, 0.0167083\}, \}
      {MnBM44, NO, NO, NO, 0.0406336}, {NiBM11, Metal, PHO, NO, 0.0559507},
      {NiBM12, NO, NO, NO, 0.025158}, {NiBM16, NO, NO, NO, 0.0568948},
      {NiBM17, NO, NO, NO, 0.0653254}, {NiBM21, Metal, PHO, NO, 0.00717952},
      {NiBM24, Metal, NO, NO, 0.04489}, {NiBM27, Metal, NO, NO, -0.002682},
      {NiBM28, Metal, PHO, NO, 0.00404369}, {NiBM30, Metal, NO, NO, 0.03299},
      {NiBM4, Metal, PH0, N0, 0.0295935}, {NiBM6, Metal, PH0, N0, 0.0074981},
      {NiBM9, Metal, PHO, NO, 0.0379949}, {ZnBM11, Metal, PHO, NO, 0.0557065},
      {ZnBM12, NO, PHO, NO, 0.0316056}, {ZnBM15, NO, PHO, NO, 0.0529235},
      {ZnBM16, Metal, NO, NO, 0.0137405}, {ZnBM17, NO, NO, NO, 0.0505517},
      {ZnBM19, Metal, NO, NO, 0.0309429}, {ZnBM22, Metal, NO, NO, 0.00630274},
      {ZnBM23, Metal, PHO, NO, 0.111893}, {ZnBM28, Metal, NO, NO, 0.0390185},
      {ZnBM29, NO, NO, NO, 0.0610565}, {ZnBM31, Metal, NO, NO, 0.0299658},
      {ZnBM37, NO, NO, NO, 0.0234296}, {ZnBM38, Metal, PHO, NO, 0.0640663},
      {ZnBM39, Metal, NO, NO, 0.0516385}, {ZnBM41, NO, NO, NO, 0.0529478},
      {ZnBM42, Metal, PHO, NO, 0.0394398}, {ZnBM43, Metal, NO, NO, 0.0325736},
      {ZnBM44, Metal, NO, NO, 0.0223613}, {ZnBM45, Metal, NO, NO, 0.0207541},
      {ZnBM46, Metal, NO, NO, 0.0593916}, {ZnBM47, Metal, NO, NO, 0.0202203}}
In[*]:= Export["/Users/otto/Documents/Students/AnnaBazzicalupo/MetalAdaptation/Genomics/
         lines_by_phenotype_17June2025.tsv",
       Join[{{"Line", "Metal resistance gene (SGD)", "Phosphorous metabolism (Amigo 2)",
           "DNA repair (SGD)", "Broad cross tolerance"}}, linephenotypes]];
```

Looking at intergenic data from SnpEff

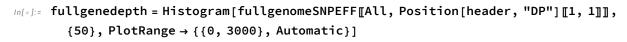
According to Shih and Fay (2021):

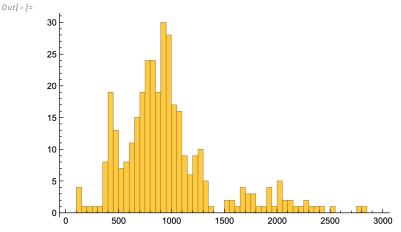
"In yeast, cis-acting variants most likely occur within the small (~500 bp) intergenic region upstream of a gene, but could also occur within the coding or 30 region of a gene."

Uploading data

Converting information from SnpEff

```
In[*]:= fullgenomeSNPEFF = Import[
          "/Users/otto/Documents/Students/AnnaBazzicalupo/MetalAdaptation/Genomics/
            snpEff_intergenic_haploid_19May2025.tsv"];
 In[*]:= header = fullgenomeSNPEFF[[1]]
Out[ • ]=
      {CHROM, chr, POS, REF, ALT, QUAL, FORMAT, gene, effect, mut_type, gene_code,
       base_change, aa_change, ANN, OLY077_S101, CdBM23_S1, CdBM25_S10, CdBM26_S18,
       CdBM29 S26, CdBM30 S34, CdBM32 S42, CdBM36 S50, CdBM37 S58, CdBM39 S2,
       CdBM42_S11, CdBM43_S19, CdBM44_S27, CdBM45_S35, CdBM46_S43, CdBM47_S51,
       CdBM48_S59, CoBM12_S4, CoBM14_S13, CoBM15_S21, CoBM16_S29, CoBM17_S37,
       COBM18_S45, COBM1_S3, COBM20_S53, COBM21_S61, COBM2_S12, COBM3_S20, COBM4_S28,
       CoBM5_S36, CoBM6_S44, CoBM7_S52, CoBM8_S60, CuBM10_S54, CuBM11_S62, CuBM12_S6,
       CuBM13_S15, CuBM14_S23, CuBM15_S31, CuBM16_S39, CuBM17_S47, CuBM18_S55, CuBM3_S5,
       CuBM4_S14, CuBM6_S22, CuBM7_S30, CuBM8_S38, CuBM9_S46, MnBM12_S63, MnBM13_S7,
       MnBM14_S16, MnBM15_S24, MnBM16_S32, MnBM17_S40, MnBM18_S48, MnBM20_S56,
       MnBM21_S64, MnBM23_S8, MnBM24_S17, MnBM25_S25, MnBM27_S33, MnBM28_S41,
       MnBM29_S49, MnBM31_S57, MnBM32_S65, MnBM34_S66, MnBM38_S72, MnBM39_S78,
       MnBM42_S84, MnBM44_S90, NiBM11_S73, NiBM12_S79, NiBM14_S85, NiBM16_S91,
       NiBM17_S97, NiBM21_S103, NiBM22_S109, NiBM24_S68, NiBM25_S74, NiBM27_S80,
       NiBM28_S86, NiBM29_S92, NiBM30_S98, NiBM4_S96, NiBM6_S102, NiBM8_S108, NiBM9_S67,
       ZnBM11_S104, ZnBM12_S110, ZnBM15_S69, ZnBM16_S75, ZnBM17_S81, ZnBM19_S87,
       ZnBM22_S93, ZnBM23_S99, ZnBM25_S105, ZnBM28_S111, ZnBM29_S70, ZnBM31_S76,
       ZnBM34_S82, ZnBM37_S88, ZnBM38_S94, ZnBM39_S100, ZnBM41_S106, ZnBM42_S112,
       ZnBM43_S71, ZnBM44_S77, ZnBM45_S83, ZnBM46_S89, ZnBM47_S95, AC, AF, AN,
       BaseQRankSum, DP, FS, MLEAC, MLEAF, MQ, MQRankSum, QD, ReadPosRankSum, SOR, LOF}
 In[*]:= fullgenomeSNPEFF = Drop[fullgenomeSNPEFF, 1];
 In[*]:= Length[fullgenomeSNPEFF]
Out[ • ]=
      348
 In[*]:= Select[fullgenomeSNPEFF, #[Position[header, "effect"][[1, 1]]] == "MODIFIER" &] // Length
Out[ • ]=
      348
      Plotting the depth of coverage at each site:
 In[*]:= Max[fullgenomeSNPEFF[All, Position[header, "DP"][1, 1]]]]
Out[ • ]=
      4828
```





There are a few low outliers:

There is one high outlier:

Mean depth:

```
In[*]:= Mean[fullgenomeSNPEFF[[All, 129]]] // N
Out[ • ]=
```

Below, we will also compare genes bearing mutations to those with upstream variants, so copying from above:

The list of 333 genes that bear a mutation from the previous section:

```
In[*]:= genelist = {"ABP1", "ACB1", "ACC1", "ADE6", "ADR1", "AFT1", "AGP1", "AHK1", "ALD5",
        "ALY2", "APC2", "ARO3", "ASN1", "ATG11", "ATG2", "ATG39", "BDS1", "BIR1",
        "BLM10", "BNA6", "BNI1", "BNI4", "BRR2", "BSD2", "BUL1", "BUL2", "CAJ1", "CCC2",
        "CCR4", "CCW12", "CDC25", "CIC1", "CLB3", "CLN3", "CNE1", "COG1", "COG3",
        "COI1", "COQ1", "CPA1", "CSM3", "CUE3", "CWC22", "CWH41", "CYK3", "DAL2",
        "DBF20", "DBP10", "DDR48", "DNF1", "DOA4", "DOT6", "DSC2", "DSE4", "DUS4",
        "EBS1", "ECM14", "ECM22", "EFR3", "ELO3", "ERB1", "ERG1", "ERG27", "ERG7",
        "ERG9", "FAS2", "FAU1", "FCY2", "FET4", "FIG2", "FIG4", "FLC2", "FMP52", "FRE6",
        "FSF1", "FYV10", "FYV6", "FZF1", "GAS4", "GAT1", "GCD2", "GID12", "GPB1",
        "GPB2", "GPH1", "GSC2", "HBT1", "HEH2", "HHF2", "HIS4", "HMF1", "HO", "HOB2",
        "HRT3", "HSL1", "HSP104", "HUL5", "HXT13", "IDP1", "IES3", "ILV2", "IMA1",
        "INP2", "IOC4", "IPI3", "IRC20", "IRC8", "IST2", "ISU1", "ITC1", "KAR3",
        "KGD1", "KIN1", "KIN82", "KRE5", "KSP1", "KTR3", "LAM1", "LAM6", "LOS1",
        "LRG1", "MAM3", "MAS2", "MCK1", "MCM5", "MCT1", "MDH2", "MDL2", "MDM30",
        "MET10", "MET30", "MEX67", "MHP1", "MKT1", "MLF3", "MLH3", "MLS1", "MMS4",
        "MNN9", "MOT1", "MPA43", "MRL1", "MRPL36", "MSC6", "MSS2", "MTR4", "MYG1",
        "MY02", "NAM8", "NAR1", "NFT1", "NGG1", "NIP100", "NPL3", "NSR1", "NUC1",
        "NUG1", "OAF1", "OCA4", "OCA5", "OPI1", "OSH2", "PBP2", "PBS2", "PCL5", "PDA1",
        "PDR1", "PDR10", "PDX1", "PET111", "PET127", "PEX2", "PFK26", "PHM7", "PHO23",
        "PH081", "PH084", "PIB1", "PKP1", "PLB2", "PMA1", "POL3", "PPQ1", "PRC1",
        "PRI1", "PRI2", "PRP2", "PRP8", "PTK2", "PYK2", "QRI7", "RAD16", "RAD17",
        "RAD26", "RBK1", "RCE1", "RCK2", "RCO1", "REC102", "RED1", "RH01", "RIM11",
        "RIM21", "RKM3", "RMP1", "RNY1", "ROG1", "RPA135", "RPI1", "RPL22A", "RPL2A",
        "RPL39", "RPS15", "RPS2", "RPT5", "RRB1", "RRG7", "RRP46", "RRP6", "RSC1",
        "RSE1", "RTC2", "SAK1", "SAP4", "SBE2", "SEC8", "SET2", "SFP1", "SHP1", "SIW14",
        "SKN1", "SMC2", "SMY2", "SNF7", "SNT2", "SPC105", "SPO77", "SPS100", "SRP40",
        "SSE1", "SSK2", "SSQ1", "SSZ1", "STE5", "STP3", "STT3", "SUB2", "SUC2", "SUM1",
        "SWC5", "SWS2", "SWT1", "SYF2", "SYG1", "TAF8", "TAH11", "TAO3", "TC089",
        "TFB1", "TFG1", "TIM54", "TOM1", "TOP1", "TOP2", "TOS3", "TPS3", "TRK2",
        "TRL1", "TUS1", "UBC1", "UBP5", "UBR2", "UBX7", "UGA3", "UGA4", "UIP5", "ULS1",
        "UME6", "URA2", "URB1", "USA1", "UTP21", "UTP8", "VBA1", "VCX1", "VHR2",
        "VID22", "VPS13", "VPS74", "VRP1", "VTC1", "VTC3", "VTC4", "VTC5", "WAR1",
        "YAP6", "YBR242W", "YBR292C", "YCF1", "YCK1", "YCT1", "YDL199C", "YDR003W-A",
        "YDR157W", "YDR541C", "YEF1", "YGR126W", "YGR130C", "YGR266W", "YHC1", "YHC3",
        "YIH1", "YIL092W", "YLL066W-B", "YLR108C", "YLR296W", "YML119W", "YMR027W",
        "YMR317W", "YOR029W", "YOR1", "YOR296W", "YOR343C", "YPK1", "YPL113C",
        "YPR078C", "YPR089W", "YPT6", "YRA1", "YRB2", "ZDS2", "ZRT1", "ZRT3"};
In[*]:= genelistSYS = {"YAL021C", "YAL040C", "YAL051W", "YAL053W", "YAL056W", "YAL058W",
        "YBL058W", "YBR003W", "YBR030W", "YBR086C", "YBR098W", "YBR114W", "YBR122C",
        "YBR147W", "YBR172C", "YBR205W", "YBR231C", "YBR233W", "YBR242W", "YBR273C",
        "YBR290W", "YBR292C", "YCL025C", "YCL030C", "YCR036W", "YCR059C", "YCR088W",
        "YCR089W", "YCR091W", "YCR095C", "YDL019C", "YDL031W", "YDL066W", "YDL073W",
        "YDL1084W", "YDL102W", "YDL104C", "YDL107W", "YDL117W", "YDL128W", "YDL155W",
```

```
"YDL170W", "YDL176W", "YDL199C", "YDL210W", "YDL223C", "YDL227C", "YDL240W",
"YDR003W-A", "YDR035W", "YDR069C", "YDR089W", "YDR103W", "YDR122W", "YDR135C",
"YDR157W", "YDR176W", "YDR177W", "YDR206W", "YDR207C", "YDR216W", "YDR259C",
"YDR270W", "YDR310C", "YDR311W", "YDR313C", "YDR351W", "YDR372C", "YDR381C-A",
"YDR381W", "YDR432W", "YDR457W", "YDR458C", "YDR541C", "YEL041W", "YEL069C",
"YER004W", "YER006W", "YER048C", "YER056C", "YER057C", "YER064C", "YER072W",
"YER073W", "YER088C", "YER129W", "YER144C", "YER156C", "YER157W", "YER166W",
"YER172C", "YER178W", "YER183C", "YFL007W", "YFL021W", "YFR030W", "YFR031C",
"YFR031C-A", "YFR047C", "YGL008C", "YGL013C", "YGL022W", "YGL027C",
"YGL071W", "YGL093W", "YGL110C", "YGL123W", "YGL131C", "YGL133W", "YGL141W",
"YGL144C", "YGL179C", "YGL223C", "YGL229C", "YGL254W", "YGL255W", "YGR032W",
"YGR037C", "YGR056W", "YGR061C", "YGR083C", "YGR095C", "YGR126W", "YGR128C",
"YGR129W", "YGR130C", "YGR143W", "YGR159C", "YGR175C", "YGR186W", "YGR193C",
"YGR233C", "YGR266W", "YGR278W", "YGR281W", "YGR287C", "YHL020C", "YHL029C",
"YHR024C", "YHR052W", "YHR064C", "YHR071W", "YHR072W", "YHR082C", "YHR086W".
"YHR132C", "YHR135C", "YHR139C", "YHR155W", "YHR165C", "YHR190W", "YIL042C",
"YIL046W", "YIL047C", "YIL063C", "YIL092W", "YIL097W", "YIL1119C",
"YIL125W", "YIL129C", "YIL162W", "YIR008C", "YIR029W", "YJL012C", "YJL042W",
"YJL050W", "YJL051W", "YJL054W", "YJL059W", "YJL084C", "YJL087C", "YJL128C",
"YJL130C", "YJL168C", "YJL189W", "YJL208C", "YJL210W", "YJR035W", "YJR046W",
"YJR059W", "YJR089W", "YKL014C", "YKL045W", "YKL101W", "YKL126W", "YKL175W",
"YKL205W", "YKR044W", "YKR050W", "YKR092C", "YKR103W", "YLL026W", "YLL040C",
"YLL051C", "YLL055W", "YLL066W-B", "YLR024C", "YLR025W", "YLR052W",
"YLR061W", "YLR072W", "YLR097C", "YLR100W", "YLR108C", "YLR110C", "YLR127C",
"YLR145W", "YLR228C", "YLR247C", "YLR248W", "YLR261C", "YLR263W", "YLR274W",
"YLR296W", "YLR298C", "YLR310C", "YLR312C", "YLR329W", "YLR337C", "YLR341W",
"YLR368W", "YLR369W", "YLR372W", "YLR373C", "YLR375W", "YLR403W", "YLR405W",
"YLR409C", "YLR425W", "YML029W", "YML049C", "YML076C", "YML111W",
"YML114C", "YML119W", "YML123C", "YMR006C", "YMR027W", "YMR044W", "YMR048W",
"YMR049C", "YMR075W", "YMR088C", "YMR108W", "YMR131C", "YMR139W", "YMR163C",
"YMR173W", "YMR212C", "YMR257C", "YMR261C", "YMR274C", "YMR275C", "YMR297W",
"YMR317W", "YMR319C", "YNL030W", "YNL032W", "YNL074C", "YNL081C", "YNL085W",
"YNL088W", "YNL097C", "YNL117W", "YNL133C", "YNL182C", "YNL233W", "YNL240C",
"YNL242W", "YNL249C", "YNL271C", "YNL294C", "YNL307C", "YNL325C", "YNR011C",
"YNR016C", "YNR031C", "YNR067C", "YOL006C", "YOL040C", "YOL060C", "YOL073C",
"YOL084W", "YOL126C", "YOL132W", "YOL164W", "YOR001W", "YOR017W", "YOR029W",
"YOR117W", "YOR166C", "YOR191W", "YOR221C", "YOR271C", "YOR296W", "YOR303W",
"YOR305W", "YOR326W", "YOR328W", "YOR336W", "YOR343C", "YOR347C", "YOR354C",
"YOR368W", "YOR371C", "YPL019C", "YPL050C", "YPL082C", "YPL106C", "YPL113C",
"YPL123C", "YPL135W", "YPL164C", "YPL169C", "YPL174C", "YPL179W", "YPL180W",
"YPL231W", "YPL270W", "YPR010C", "YPR049C", "YPR055W", "YPR078C", "YPR079W",
"YPR089W", "YPR111W", "YPR117W", "YPR141C", "YPR145W", "YPR160W", "YPR165W"};
```

```
Out[ • ]=
        {333, 333}
```

Dropping genes with too many uncalled genotypes and dropping genotypic calls with <5 depth

Processing and dropping genes with two many poor quality genotypes

Sites that were hard to call (low depth or no genotype call) can indicate duplicated genes and alignment problems. Here we identify these:

```
In[@]:= droppos = -1 + Position[header, "OLY077_S101"] [[1, 1]];
       (*# columns to be dropped before first sample*)
 In[*]:* topos = Position[header, "ZnBM47_S95"] [1, 1]; (*# last sample column*)
       Extracting the ref vs alt call in genomeSNPEFF:
 In[*]:= genomeSNPEFF = Table[".", {i, 1, Length[fullgenomeSNPEFF]}, {j, 1, topos - droppos}];
 In[*]:= genomeSNPEFF[[1]] // Length
Out[ • ]=
       110
 In[*]:= For[i = 1, i ≤ Length[fullgenomeSNPEFF], i++,
        For [j = 1 + droppos, j \le topos, j++,
         temp = StringSplit[fullgenomeSNPEFF[i, j], {":"}];
         genomeSNPEFF[[i, j - droppos]] = temp[[1]];
        ]
       1
 In[*]:= genomeSNPEFF // Length
Out[ • ]=
 In[*]:= Tally[Select[genomeSNPEFF[[1]], # # "." &]]
Out[ • ]=
       \{\{1, 36\}, \{0, 73\}\}
      All mutant sites and the tally of mutations in them:
 In[@]:= summarytable = Table[{fullgenomeSNPEFF[[i, 1]],
            fullgenomeSNPEFF[i, 3], fullgenomeSNPEFF[i, 8], fullgenomeSNPEFF[i, 9],
            Sort[Tally[genomeSNPEFF[i]]]], {i, 1, Length[genomeSNPEFF]}];
      The set of sites where the number of "." calls was ≥5:
```

ln[a]:= dropthese = Select[summarytable, (#[5, 1, 1] == ".") && (#[5, 1, 2] \geq 5) &]; dropthese // MatrixForm dropthese // Length

```
Out[ • ]//MatrixForm=
         chrII
                  258 784
                               FLR1
                                       MODIFIER
                                                 \{\{., 53\}, \{0, 49\}, \{1, 8\}\}
         chrII
                  643 488
                            DUR1%2C2
                                       MODIFIER
                                                  \{\{., 96\}, \{0, 1\}, \{1, 13\}\}\
         chrII
                  643 489
                            DUR1%2C2
                                       MODIFIER
                                                  \{\{., 104\}, \{0, 1\}, \{1, 5\}\}
                                                 \{\{., 35\}, \{0, 72\}, \{1, 3\}\}
         chrIII
                  295 234
                              HMRA2
                                       MODIFIER
         chrIV
                  273 629
                              PH02
                                       MODIFIER {{., 11}, {0, 74}, {1, 25}}
         chrIV
                  336 206
                              CBS1
                                       MODIFIER
                                                 \{\{.,5\},\{0,91\},\{1,14\}\}
                                       MODIFIER {{., 17}, {0, 31}, {1, 62}}
         chrIV
                  384 223
                              NAT1
                                       MODIFIER {{., 38}, {0, 10}, {1, 62}}
         chrIV
                  384 226
                              NAT1
         chrIV
                  384 262
                              NAT1
                                       MODIFIER {{., 24}, {0, 25}, {1, 61}}
                                       MODIFIER {{., 11}, {0, 70}, {1, 29}}
         chrIV
                  384 290
                              NAT1
                                                  \{\{.,5\},\{0,98\},\{1,7\}\}
         chrIV
                 1341641
                              DOT1
                                       MODIFIER
          chrV
                  116118
                              URA3
                                       MODIFIER \{\{., 7\}, \{0, 81\}, \{1, 22\}\}
         chrVII
                  278 779
                              MON1
                                       MODIFIER
                                                  \{\{.,5\},\{0,96\},\{1,9\}\}
                                       MODIFIER {{., 24}, {0, 76}, {1, 10}}
         chrVII
                  323 365
                              TOS8
         chrVII
                  701 187
                              SRB5
                                       MODIFIER {{., 43}, {0, 61}, {1, 6}}
         chrVII
                  735 813
                               COG2
                                       MODIFIER {{., 13}, {0, 95}, {1, 2}}
         chrVII
                  735 814
                                       MODIFIER {{., 13}, {0, 95}, {1, 2}}
                              COG2
         chrVII
                  811448
                                       MODIFIER {{., 92}, {0, 13}, {1, 5}}
                              NSR1
        chrVIII
                   2292
                               COS8
                                       MODIFIER {{., 10}, {0, 85}, {1, 15}}
        chrVIII
                   85 388
                               PRS3
                                       MODIFIER
                                                 \{\{., 23\}, \{0, 85\}, \{1, 2\}\}
        chrVIII
                              PRS3
                                       MODIFIER {{., 23}, {0, 85}, {1, 2}}
                   85 390
          chrX
                  745 669
                               COS5
                                       MODIFIER {{., 27}, {0, 39}, {1, 44}}
         chrXI
                  196211
                             01-0ct
                                       MODIFIER \{\{., 59\}, \{0, 44\}, \{1, 7\}\}
                                                  \{\{., 60\}, \{0, 49\}, \{1, 1\}\}
         chrXI
                  379 671
                           tV(AAC)K2 MODIFIER
         chrXI
                  379672 tV(AAC)K2 MODIFIER
                                                  \{\{., 28\}, \{0, 81\}, \{1, 1\}\}
         chrXI
                  517 454
                               KAE1
                                       MODIFIER
                                                  \{\{., 107\}, \{0, 1\}, \{1, 2\}\}
                                                  \{\{., 105\}, \{0, 1\}, \{1, 4\}\}
         chrXI
                  517 455
                               KAE1
                                       MODIFIER
         chrXII
                  251 194
                              FCF2
                                       MODIFIER {{., 10}, {0, 97}, {1, 3}}
         chrXII
                  421 496
                              TIS11
                                       MODIFIER
                                                  \{\{.,9\},\{0,93\},\{1,8\}\}
                                                  \{\{., 95\}, \{0, 1\}, \{1, 14\}\}
         chrXII
                  593 143
                             YLR225C
                                       MODIFIER
                                                  \{\{., 12\}, \{0, 97\}, \{1, 1\}\}
         chrXII 1065029
                            YLR460C
                                       MODIFIER
                  850633 YPR158W-A MODIFIER {{., 91}, {0, 14}, {1, 5}}
         chrXVI
```

Out[•]= 32

In[*]:= dropme = Union[dropthese[All, 3]]] % // Length

Out[•]= {01-Oct, CBS1, COG2, COS5, COS8, DOT1, DUR1%2C2, FCF2, FLR1, HMRA2, KAE1, MON1, NAT1, NSR1, PHO2, PRS3, SRB5, TIS11, TOS8, tV(AAC)K2, URA3, YLR225C, YLR460C, YPR158W-A} Out[•]=

24

We'll remove these specific sites by position too (not by gene, as these sites may be upstream of many genes, which hasn't been processed yet):

```
In[*]:= dropmeposition = Union[dropthese[All, 2]];
      Depth of coverage for these genes is similar:
 In[@]:= Select[fullgenomeSNPEFF, MemberQ[dropme, #[8]] &] [All, {8, 129}]
      Mean[%[All, 2]] // N
Out[ • ]=
       {{FLR1, 303}, {DUR1%2C2, 292}, {DUR1%2C2, 158}, {HMRA2, 1986}, {PHO2, 702},
        {CBS1, 709}, {NAT1, 1327}, {NAT1, 1253}, {NAT1, 1231}, {NAT1, 987},
        {NAT1, 838}, {DOT1, 978}, {URA3, 1248}, {MON1, 821}, {TOS8, 502}, {TOS8, 2018},
        {SRB5, 1355}, {SRB5, 1281}, {COG2, 860}, {COG2, 859}, {NSR1, 130}, {COS8, 773},
        {PRS3, 1684}, {PRS3, 1522}, {COS5, 657}, {01-Oct, 2022}, {tV(AAC)K2, 1747},
        {tV(AAC)K2, 143}, {tV(AAC)K2, 205}, {KAE1, 129}, {KAE1, 124}, {FCF2, 928},
        {TIS11, 721}, {YLR225C, 450}, {YLR460C, 1124}, {YPR158W-A, 355}}
Out[ • ]=
      900.611
      Mapping quality is also poorer on average, with the mean at the bottom 8.9% tail:
 In[*]:= Select[fullgenomeSNPEFF, MemberQ[dropme, #[8]] &] [[All, {8, 133}]]
      Mean[%[All, 2]] // N
      Length[Select[fullgenomeSNPEFF[All, 133], # < % &]] /</pre>
          Length[fullgenomeSNPEFF[All, 133]] // N // PercentForm
Out[ • ]=
       {{FLR1, 55.54}, {DUR1%2C2, 59.98}, {DUR1%2C2, 60.}, {HMRA2, 56.3}, {PHO2, 60.},
        {CBS1, 60.}, {NAT1, 58.99}, {NAT1, 58.98}, {NAT1, 57.38}, {NAT1, 53.14},
        {NAT1, 52.7}, {DOT1, 60.}, {URA3, 60.}, {MON1, 57.64}, {TOS8, 53.69}, {TOS8, 60.},
        {SRB5, 59.95}, {SRB5, 59.94}, {COG2, 58.17}, {COG2, 58.1}, {NSR1, 60.},
        {COS8, 59.57}, {PRS3, 52.77}, {PRS3, 52.02}, {COS5, 57.36}, {01-0ct, 60.},
        {tV(AAC)K2, 56.25}, {tV(AAC)K2, 51.59}, {tV(AAC)K2, 50.94}, {KAE1, 60.}, {KAE1, 60.},
        {FCF2, 60.}, {TIS11, 60.}, {YLR225C, 60.}, {YLR460C, 54.43}, {YPR158W-A, 59.56}}
Out[ • ]=
      57.6386
Out[ • ]//PercentForm=
      8.908%
      Dropping all sites (32) with ≥5 "." calls:
 In[*]:= Length[fullgenomeSNPEFF];
      fullgenomeSNPEFF = Select[fullgenomeSNPEFF,
          (MemberQ[dropme, #[8]]] == False) || (MemberQ[dropmeposition, #[3]] == False) &];
      %% - Length[fullgenomeSNPEFF]
Out[ • ]=
      32
```

Processing and dropping sites with <5 depth

We next scan for sites that are called with depth <5 (note that we do not filter on the depth for each allele, just total depth at that site in that sample).

An example of a site with low coverage:

Out[•]=

110

```
In[@]:= Position[fullgenomeSNPEFF[All, 3], 415 764][1, 1];
        fullgenomeSNPEFF[[%, 49;; 53]]
Out[ • ]=
        \{0:10,0:10:99:0,301,1:1,1:2:2:2,0,
         0:3,0:3:99:0,100,0:5,0:5:99:0,99,0:2,1:3:21:0,21
 In[•]:= For[i = 1; altered = {}, i ≤ Length[fullgenomeSNPEFF], i++,
         For [j = 1 + droppos, j \le topos, j++,
          temp = StringSplit[fullgenomeSNPEFF[i, j], {":"}];
          If[ToExpression[temp[3]]] < 5, temp[1] = ".";</pre>
            fullgenomeSNPEFF[[i, j]] = StringRiffle[temp, ":"];
            AppendTo[altered, {i, j}]];
         ]
        1
 In[•]:= altered
Out[ • ]=
          \big\{\{2,33\},\{2,51\},\{2,67\},\{2,75\},\{2,106\},\{3,36\},\{3,46\},\{3,55\},\{3,67\},\{3,68\},\{3,69\},
          \{3, 76\}, \{3, 79\}, \{3, 116\}, \{3, 124\}, \{4, 19\}, \{4, 36\}, \{4, 46\}, \{4, 51\}, \{4, 55\}, \{4, 67\}, \{4, 68\},
          \{4, 69\}, \{4, 75\}, \{4, 76\}, \{4, 79\}, \{4, 116\}, \{4, 124\}, \{5, 19\}, \{5, 36\}, \{5, 46\}, \{5, 51\}, \{5, 53\},
          \{5,55\}, \{5,67\}, \{5,68\}, \{5,69\}, \{5,75\}, \{5,76\}, \{5,79\}, \{5,116\}, \{5,124\}, \{6,19\},
           9255 \dots, 313, 106, 313, 108, 313, 109, 313, 111, 313, 115, 313, 117, 313, 118,
          \{313, 119\}, \{313, 123\}, \{313, 124\}, \{314, 19\}, \{314, 23\}, \{314, 31\}, \{314, 33\}, \{314, 43\}, \{314, 51\},
          {314, 53}, {314, 67}, {314, 68}, {314, 69}, {314, 71}, {314, 106}, {314, 123}, {314, 124},
           \{315, 19\}, \{315, 33\}, \{315, 51\}, \{315, 53\}, \{315, 65\}, \{315, 67\}, \{315, 68\}, \{315, 69\}, \{315, 124\},
           \{316, 19\}, \{316, 33\}, \{316, 51\}, \{316, 53\}, \{316, 65\}, \{316, 67\}, \{316, 68\}, \{316, 69\}, \{316, 124\}\}
         Full expression not available (original memory size: 0.9 MB)
       The example is now corrected:
 In[@]:= Position[fullgenomeSNPEFF[All, 3], 415 764][1, 1];
        fullgenomeSNPEFF[%, 49;; 53]
Out[ • ]=
        \{0:10,0:10:99:0,301, .:1,1:2:2:2,0,
         .:3,0:3:99:0,100,0:5,0:5:99:0,99,.:2,1:3:21:0,21
        Extracting the ref vs alt call in genomeSNPEFF:
 In[*]:= genomeSNPEFF = Table[".", {i, 1, Length[fullgenomeSNPEFF]}, {j, 1, topos - droppos}];
       genomeSNPEFF[[1]] // Length
 In[ • ]:=
```

```
In[*]:= For[i = 1, i ≤ Length[fullgenomeSNPEFF], i++,
        For [j = 1 + droppos, j \le topos, j++,
         temp = StringSplit[fullgenomeSNPEFF[i, j], {":"}];
         genomeSNPEFF[[i, j - droppos]] = temp[[1]];
        ]
       1
 In[*]:= genomeSNPEFF // Length
Out[ • ]=
 In[*]:= Tally[Select[genomeSNPEFF[[1]], # # "." &]]
Out[ • ]=
       \{\{1, 36\}, \{0, 73\}\}
       All mutant sites and the tally of mutations in them (ignore the error, which comes from sorting so that
       "." is last):
 In[*]:= summarytable =
         Table[{fullgenomeSNPEFF[i, 1], fullgenomeSNPEFF[i, 3], fullgenomeSNPEFF[i, 8],
            fullgenomeSNPEFF[i, 9], SortBy[Tally[genomeSNPEFF[i]]], ToExpression[#] &]},
           {i, 1, Length[genomeSNPEFF]}];
       ToExpression::sntx: Invalid syntax in or before ". ".
       ToExpression::sntx: Invalid syntax in or before ". ".
       ToExpression::sntx: Invalid syntax in or before ". ".
       General::stop: Further output of ToExpression::sntx will be suppressed during this calculation.
       This does not change the number of genes with mutations detected (i.e., all tallies with only two
       elements include the ref and alt allele):
```

In[⊕]:= Select[summarytable, Length[#[5]]] ≤ 2 &] // MatrixForm

```
Out[•]//MatrixForm=
       chrII 684840 0M14
                               MODIFIER {{0, 2}, {1, 108}}
       chrIII 309030 YCR102C MODIFIER {{0, 1}, {1, 109}}
        chrV 406772
                     GL03
                               MODIFIER {{0, 109}, {1, 1}}
        chrV 407063 GL03
                               MODIFIER {{0, 1}, {1, 109}}
       chrVI 193508 tS(GCU)F MODIFIER \{\{0, 1\}, \{1, 109\}\}
       chrVII 649 059 GCD2
                               MODIFIER {{0, 109}, {1, 1}}
       chrXI 146956
                       PIR1
                               MODIFIER {{0, 109}, {1, 1}}
       chrXI 379623 tV(AAC)K2 MODIFIER {{0, 1}, {1, 109}}
                               MODIFIER {{0, 109}, {1, 1}}
       chrXI 489841 DBP7
       chrXII 185314
                       YEH2
                               MODIFIER {{0, 1}, {1, 109}}
       chrXII 881947
                       SEC61
                               MODIFIER {{0, 1}, {1, 109}}
```

Stats for the remaining SNPs

Plotting the depth of coverage at each site:

```
Mean[fullgenomeSNPEFF[All, 129]] // N
 In[ • ]:=
Out[ • ]=
       983.203
 In[*]:= Max[fullgenomeSNPEFF[[All, 129]]]
Out[ • ]=
       4828
 In[*]:= Histogram[fullgenomeSNPEFF[All, 129]], {50}, PlotRange → {{0, 3000}, Automatic}]
Out[ • ]=
       30 |
       25
       20
       15
       10
        5
        0
                  500
                           1000
                                   1500
                                            2000
                                                     2500
```

Processing SnpEff

The gene list provided by SnpEff just picks the first of the genes, not necessarily the closest:

```
In[*]:= fullgenomeSNPEFF[[1;; 4, Position[header, "gene"][[1, 1]]]
Out[ • ]=
       {YAL065C, FL09, YAL056C-A, ACS1}
```

in[*]:= annotations = fullgenomeSNPEFF[[All, Position[header, "ANN"][[1, 1]]]; Looking through the SnpEff annotations and finding which genes are nearest, with the SNP in the upstream region:

```
In[ • ]:= For [ i = 1;
       regpotential = {};
       morethanone = {};
       closeregpotential = {}, i ≤ Length[annotations], i++,
       temp = StringSplit[
          StringSplit[fullgenomeSNPEFF[i, Position[header, "ANN"][1, 1]], ","], "|"];
       temp2 = Select[temp, #[2] == "upstream_gene_variant" &];
       temp2 =
         temp2 /. "ADE5%2C7" → "ADE5,7" /. "WARNING_TRANSCRIPT_NO_START_CODON" → 10<sup>6</sup> /.
           "WARNING_TRANSCRIPT_MULTIPLE_STOP_CODONS" → 10<sup>6</sup> /.
          "WARNING_TRANSCRIPT_INCOMPLETE" → 10<sup>6</sup>;
       If[Length[temp2] == 0, AppendTo[regpotential, {"", "", "", ""}],
         For[j = 1;
          dist = {};
          closegene = {};
          systname = \{\}, j \le Length[temp2], j++,
          AppendTo[dist, ToExpression[Last[temp2[j]]]];
          AppendTo[closegene, temp2[j, 4]];
          AppendTo[systname, temp2[j, 5]]
         ];
         closest = PositionSmallest[ToExpression[dist]][1];
         AppendTo[regpotential,
          {temp2[closest, 4], temp2[closest, 5], dist[closest], temp2[closest, 10]}];
         If[Length[numclose = Select[Transpose[{dist, closegene}], #[1] ≤ 500 &]] > 1,
          AppendTo[morethanone, {i, numclose}]];
       ]
 regpotential[1;; 4]
Out[ • ]=
      {{TDA8, YAL064C-A, 1849, c.-1849C>G}, {GDH3, YAL062W, 771, c.-771G>A},
       {FLC2, YAL053W, 3064, c.-3064C>T}, {FLC2, YAL053W, 412, c.-412C>T}}
      Appending the nearest gene (SNP must be upstream) and distance to fullgenomeSNPEFF and updating
      the header:
 In[*]:= fullgenomeSNPEFF = Table[Flatten[{fullgenomeSNPEFF[i]], regpotential[i]]}] /.
           Missing["NotAvailable"] → "", {i, 1, Length[fullgenomeSNPEFF]}];
```

In[*]:= newheader = Join[Table[StringSplit[header[i]], "_"][1], {i, 1, Length[header]}], {"Upstream of nearest gene", "Systematic name of nearest gene", "Distance", "Effect on closest"}]

Out[•]=

{CHROM, chr, POS, REF, ALT, QUAL, FORMAT, gene, effect, mut, gene, base, aa, ANN, OLY077, CdBM23, CdBM25, CdBM26, CdBM29, CdBM30, CdBM32, CdBM36, CdBM37, CdBM39, CdBM42, CdBM43, CdBM44, CdBM45, CdBM46, CdBM47, CdBM48, CoBM12, CoBM14, COBM15, COBM16, COBM17, COBM18, COBM1, COBM20, COBM21, COBM2, COBM3, COBM4, CoBM5, CoBM6, CoBM7, CoBM8, CuBM10, CuBM11, CuBM12, CuBM13, CuBM14, CuBM15, CuBM16, CuBM17, CuBM18, CuBM3, CuBM4, CuBM6, CuBM7, CuBM8, CuBM9, MnBM12, MnBM13, MnBM14, MnBM15, MnBM16, MnBM17, MnBM18, MnBM20, MnBM21, MnBM23, MnBM24, MnBM25, MnBM27, MnBM28, MnBM29, MnBM31, MnBM32, MnBM34, MnBM38, MnBM39, MnBM42, MnBM44, NiBM11, NiBM12, NiBM14, NiBM16, NiBM17, NiBM21, NiBM22, NiBM24, NiBM25, NiBM27, NiBM28, NiBM29, NiBM30, NiBM4, NiBM6, NiBM8, NiBM9, ZnBM11, ZnBM12, ZnBM15, ZnBM16, ZnBM17, ZnBM19, ZnBM22, ZnBM23, ZnBM25, ZnBM28, ZnBM29, ZnBM31, ZnBM34, ZnBM37, ZnBM38, ZnBM39, ZnBM41, ZnBM42, ZnBM43, ZnBM44, ZnBM45, ZnBM46, ZnBM47, AC, AF, AN, BaseQRankSum, DP, FS, MLEAC, MLEAF, MQ, MQRankSum, QD, ReadPosRankSum, SOR, LOF, Upstream of nearest gene, Systematic name of nearest gene, Distance, Effect on closest}

In[*]:= intergenicCLOSE = Sort[Select[regpotential, #[3]] ≤ 500 &]][All, {1, 3}] Length[%]

Out[•]=

```
{{AAC3, 218}, {ADE5,7, 256}, {ADH4, 312}, {ADH7, 39}, {ALE1, 354}, {ALT2, 425},
  {ARC15, 145}, {ARG4, 124}, {ARO7, 178}, {ATH1, 121}, {AVO2, 46}, {BMT5, 125},
  {BNA6, 418}, {BUB2, 75}, {CBP4, 483}, {CCA1, 302}, {CLB5, 393}, {COG5, 74},
  {COQ8, 159}, {CRG1, 388}, {CSE1, 136}, {CWC27, 135}, {DAD3, 136}, {DAM1, 69},
  {DAT1, 100}, {DNF3, 48}, {EAP1, 25}, {EHT1, 298}, {EMI2, 199}, {EP01, 366},
  {ERG26, 228}, {ESF1, 107}, {FCY21, 35}, {FLC2, 412}, {FLC3, 74}, {FRA1, 58},
  {GCN3, 187}, {GEX1, 250}, {GLC8, 209}, {GPI13, 119}, {GPR1, 90}, {GSC2, 193},
  {HFA1, 78}, {HMG2, 500}, {HOR7, 468}, {HOT1, 113}, {HOT13, 40}, {HRD1, 403},
  {HSK3, 21}, {HSP104, 430}, {HYM1, 408}, {IML2, 414}, {IRC3, 120}, {ITC1, 286},
  {KTI11, 12}, {LDB16, 121}, {LIA1, 178}, {MEK1, 44}, {MMS1, 77}, {MMT1, 208},
  {MNN5, 182}, {MRP49, 334}, {MRPL9, 124}, {MSF1, 181}, {MSO1, 121}, {MST1, 193},
  {NAM2, 59}, {NAR1, 246}, {NBP1, 199}, {NDD1, 145}, {NOC2, 97}, {NOP4, 161},
  \{NOP56, 163\}, \{NOP9, 144\}, \{OAF1, 31\}, \{OCA5, 132\}, \{OM45, 339\}, \{ORC4, 68\}, \{ORC4, 68\},
  {PAH1, 156}, {PDA1, 13}, {PDC6, 459}, {PEP4, 56}, {PHO80, 184}, {PHO91, 146},
  {PIL1, 171}, {PMT6, 334}, {POM34, 146}, {POP2, 388}, {PPT2, 130}, {PPZ1, 319},
  {PRM7, 223}, {PR02, 222}, {PRP11, 96}, {PUF4, 329}, {RAD14, 159}, {RAS1, 314},
  {RBD2, 106}, {RFC3, 128}, {RPA12, 109}, {RPH1, 21}, {RPL11B, 152},
  {RPL17B, 151}, {RPL18A, 212}, {RPL22B, 182}, {RPL3, 187}, {RPL34B, 418},
  {RP041, 393}, {RPS0A, 457}, {RPS22A, 287}, {RRM3, 167}, {RSA1, 210},
  {RSE1, 3}, {SAM50, 271}, {SCM4, 171}, {SDA1, 245}, {SDD2, 171}, {SDD4, 156},
  {SD01, 389}, {SEC12, 92}, {SEC31, 86}, {SET5, 10}, {SET5, 11}, {SMA1, 137},
  {SMI1, 441}, {SNR61, 94}, {SNR63, 207}, {SOD2, 365}, {SPC42, 490}, {SPO21, 157},
  {SSA1, 331}, {SWI6, 47}, {SYC1, 137}, {SYG1, 48}, {TCD2, 270}, {THI2, 165},
  {TIF5, 50}, {TOK1, 222}, {TOM7, 312}, {TOP1, 244}, {TRE1, 71}, {UBP13, 153},
  {UBR2, 104}, {UBX4, 81}, {VHR1, 365}, {VMA13, 103}, {VMA13, 302}, {VPH1, 27},
  {VPS54, 170}, {VPS68, 482}, {VTA1, 125}, {VTC5, 232}, {YBR300C, 372},
  {YBR300C, 388}, {YCL012C, 89}, {YCR006C, 476}, {YDL180W, 439}, {YDR169C-A, 75},
  {YEL043W, 82}, {YGL063C-A, 134}, {YGL063C-A, 135}, {YGR125W, 167},
  {YGR293C, 385}, {YHR139C-A, 41}, {YIL169C, 437}, {YJL107C, 203}, {YJL193W, 394},
  {YKE2, 264}, {YKR005C, 142}, {YLL054C, 175}, {YME2, 393}, {YMR001C-A, 116},
  {YNL190W, 91}, {YOP1, 328}, {YOR121C, 106}, {YOR121C, 112}, {YPK9, 170},
  {YPL068C, 134}, {YPL222C-A, 342}, {YPQ2, 188}, {YPR078C, 262}, {ZWF1, 373}}
```

IntergenicFURTHER = Sort[Select[regpotential, #[3] > 500 &]][All, {1, 3}] Length[%]

```
Out[ • ]=
      {{AGA1, 549}, {ALE1, 3743}, {AMF1, 1098}, {ATG18, 1303}, {BET4, 823}, {BI02, 2017},
       {BRX1, 2745}, {BUD3, 3536}, {BUL2, 881}, {CNL1, 2327}, {COF1, 810}, {CUP9, 1728},
       {DDR2, 560}, {DLD3, 1630}, {DNF2, 609}, {ERB1, 2135}, {ERR2, 2305}, {ETT1, 520},
       {FAP7, 902}, {FLC2, 3064}, {FL01, 4814}, {FL01, 4821}, {FL010, 1151}, {FMN1, 1812},
       {FRE8, 2238}, {GDH3, 771}, {GLC3, 554}, {GLO3, 2420}, {GLO3, 2711}, {GLO3, 2788},
       {GUA1, 3110}, {HXT7, 530}, {KGD1, 599}, {LGE1, 1771}, {LYS9, 1323}, {MDN1, 2506},
       {MIX17, 3488}, {MLP1, 2247}, {MNN4, 755}, {MRH1, 1209}, {MSC6, 709}, {MSS4, 3233},
       {NCE102, 738}, {NGG1, 1285}, {NSR1, 843}, {OCA6, 1410}, {PAD1, 696}, {PAD1, 850},
       {PAU10, 928}, {PAU17, 1332}, {PDS5, 919}, {PEX2, 3572}, {PGA1, 1965},
       {PMT4, 1133}, {PRE10, 843}, {PSA1, 2688}, {PSA1, 3022}, {PSP2, 1835}, {RCK1, 691},
       {RPC82, 3823}, {RPL4B, 1275}, {RPL8B, 3391}, {RP026, 1539}, {SAM1, 2118},
       {SDD4, 509}, {SDD4, 686}, {SEC27, 1488}, {SMC6, 3340}, {SNR52, 2977}, {SPT21, 519},
       {SQT1, 1213}, {SSF2, 523}, {TDA8, 1849}, {TIM10, 3772}, {TMS1, 2180}, {TOS8, 1751},
       {TRM11, 2882}, {URA2, 1149}, {USO1, 1660}, {VOA1, 1401}, {VPS1, 4066},
       {WTM1, 908}, {YAE1, 1312}, {YAL042C-A, 4154}, {YAP1802, 1319}, {YBL070C, 2626},
       {YBL070C, 2681}, {YBL111C, 1148}, {YBL111C, 1149}, {YBL111C, 1166},
       {YBL111C, 1173}, {YBL111C, 1182}, {YDL022C-A, 832}, {YDL186W, 1823},
       {YDL240C-A, 3782}, {YDR034C-D, 544}, {YDR537C, 2496}, {YER076C, 970},
       {YER137C, 1594}, {YER137C, 1595}, {YER137C, 1597}, {YFL021C-A, 2539},
       {YGL014C-A, 2488}, {YGR269W, 1642}, {YHL005C, 1286}, {YHR140W, 629},
       {YIL154C, 2257}, {YIR018C-A, 680}, {YIR018C-A, 3557}, {YJL045W, 1390},
       {YJL107C, 514}, {YJL169W, 2579}, {YJL169W, 2593}, {YKL031W, 2448}, {YKL162C-A, 882},
       {YLL066W-A, 632}, {YLL066W-A, 635}, {YLR302C, 2362}, {YML018C, 701},
       {YMR182W-A, 3578}, {YMR242W-A, 1338}, {YNR062C, 3447}, {YOL150C, 909},
       {YOR343W-A, 4558}, {YOR376W, 1775}, {YOR392W, 1075}, {YPL257W-A, 647},
       {YPR014C, 2379}, {YPR142C, 822}, {YR02, 660}, {ZRT1, 961}, {ZRT1, 1577}}
Out[ • ]=
```

Some sites have more than one gene within 500bp, but these were not analysed (only the closest gene):

```
In[a]:= morethanone(*row, closest gene, distance to all genes within 500bp*)
      Length[%]
Out[ • ]=
      {{4, {{465, ACS1}, {412, FLC2}}}},
       {7, {{331, SSA1}, {412, EFB1}}}, {15, {{410, YBL071C-B}, {12, KTI11}}},
        {29, {{188, SNR43}, {121, LDB16}}}, {82, {{302, CCA1}, {398, RPH1}}},
        {86, {{182, RPL22B}, {363, MIL1}}}, {94, {{430, MT01}, {256, ADE5,7}}},
        {99, {{346, PRP43}, {159, COQ8}}}, {101, {{134, YGL063C-A}, {411, PYC1}}},
        {102, {{135, YGL063C-A}, {410, PYC1}}}, {103, {{377, YGL015C}, {329, PUF4}}},
        {130, {{167, RRM3}, {207, ERC1}}}, {132, {{41, YHR139C-A}, {112, YHR140W}}},
        {139, {{492, TMA108}, {339, 0M45}}}, {145, {{48, SYG1}, {438, YIL046W-A}}},
        {151, {{287, RPS22A}, {344, RPL39}}}, {159, {{222, TOK1}, {389, SRS2}}},
        {163, {{144, NOP9}, {149, YJL009W}}}, {164, {{237, NTA1}, {109, RPA12}}},
        {175, {{40, HOT13}, {452, YKL083W}}}, {200, {{125, VTA1}, {145, SWI6}}},
        {201, {{223, VTA1}, {47, SWI6}}}, {207, {{59, NAM2}, {478, SMC6}}},
        {209, {{199, NBP1}, {364, GAB1}}}, {210, {{370, TAF8}, {100, DAT1}}},
        {212, {{3, RSE1}, {118, GSF2}}}, {217, {{116, YMR001C-A}, {500, MIX17}}},
        {222, {{171, SDD2}, {338, RC01}}}, {227, {{348, EAR1}, {113, HOT1}}},
        {231, {{159, RAD14}, {334, ERG2}}}, {238, {{275, MID1}, {128, RFC3}}},
        {240, {{246, NAR1}, {346, LAP3}}}, {247, {{92, SEC12}, {142, BUD17}}},
        {268, {{354, ALE1}, {373, HEM15}}}, {271, {{181, GEP3}, {97, NOC2}}},
        {280, {{145, NDD1}, {218, NUD1}}}, {286, {{106, RBD2}, {183, YPL245W}}},
        {290, {{71, TRE1}, {196, SPT14}}}, {292, {{130, PPT2}, {310, PXA1}}},
        {294, {{135, CWC27}, {188, TIM50}}}, {309, {{262, YPR078C}, {342, MRL1}}}}
Out[ • ]=
      41
      These were the genes with more than 1 intergenic mutation within 500 basepairs
 In[@]:= Select[Tally[intergenicCLOSE[All, 1]]], #[2] > 1 &]
Out[ • ]=
      {{SET5, 2}, {VMA13, 2}, {YBR300C, 2}, {YGL063C-A, 2}, {YOR121C, 2}}
      Of these, the last three are dubious open-reading frames, leaving
        "SET5" 2 \
       \"VMA13" 2
```

Only VMA13 mutants are known to affect metal resistance (SGD).

These are the set of genes with both intergenic and genic SNPS in the dataset:

```
In[@]:= Intersection[genelist, intergenicCLOSE[[All, 1]]]
      Length[%]
Out[ • ]=
       {BNA6, FLC2, GSC2, HSP104, ITC1, NAR1, OAF1,
        OCA5, PDA1, RSE1, SYG1, TOP1, UBR2, VTC5, YPR078C}
Out[ • ]=
      15
```

Relating mutations to phenotypes (metal resistance, phosphorous metabolism, DNA repair)

Metal Resistance: The list of genes affecting metal resistance was downloaded from SGD (https://yeastgenome.org/observable/APO:0000090) on 23 May 2025:

```
In[*]:= metallist = Import[
          "/Users/otto/Documents/Students/AnnaBazzicalupo/MetalAdaptation/Genomics/
            metal_resistance_annotations_23May2025.txt", "tsv"];
 In[*]:= metalhead = metallist[[2]]
Out[ • ]=
      {Gene, Gene Systematic Name, Phenotype, Experiment Type, Experiment Type Category,
       Mutant Information, Strain Background, Chemical, Details, Reference}
 In[*]:= metallist = Drop[metallist, 2];
      Here we will use the Gene Systematic Name for comparison:
 In[@]:= metalgenenames = Union[metallist[All, 2]]];
      Length[%]
Out[ • ]=
      1791
```

Of the SNPs that are within 500bp of gene (upstream position), 176 genes are involved and 31.82% are listed as having metal tolerance phenotypes:

Out[•]=

15

```
m[*]:= genelistCLOSESYS = Union[Sort[Select[regpotential, #¶3] ≤ 500 &]][All, 2]];
      Intersection[genelistCLOSESYS, metalgenenames]
       {Length[genelistCLOSESYS], Length[%],
       Length[%] / Length[genelistCLOSESYS] // N // PercentForm}
Out[ • ]=
       {YAL051W, YAL053W, YBL071W-A, YBR085W, YCL073C, YDR027C, YDR089W, YEL043W,
       YER169W, YER178W, YGL234W, YGR086C, YGR214W, YGR229C, YHR008C, YIL047C,
       YIL052C, YIL136W, YJL177W, YJL190C, YKL204W, YLL029W, YLR024C, YLR181C,
       YLR182W, YLR200W, YLR382C, YLR450W, YML016C, YML113W, YMR172W, YMR177W,
       YMR201C, YNL051W, YNL241C, YNR013C, YNR026C, YNR049C, YNR052C, YOL001W,
       YOL006C, YOL013C, YOL129W, YOR101W, YOR206W, YOR270C, YOR291W, YOR323C,
       YOR372C, YPL148C, YPL154C, YPL176C, YPL193W, YPR036W, YPR060C, YPR120C}
Out[ • ]=
       {176, 56, 31.82%}
      By contrast, for the 333 genic SNPs, 121 (36.34%) were annotated as having metal tolerance, which is
      slightly higher by 14.2% than the close intergenic SNPs.
      By comparison, of the SNPs that are not within 500bp of gene (upstream position), 115 genes are
      involved and only 20.87% are listed as having metal tolerance phenotypes:
 In[a]:= genelistFURTHERSYS = Union[Sort[Select[regpotential, #[3] > 500 &]][All, 2]];
      Intersection[genelistFURTHERSYS, metalgenenames]
       {Length[genelistFURTHERSYS], Length[%],
       Length[%] / Length[genelistFURTHERSYS] // N // PercentForm}
Out[ • ]=
       {YAL053W, YDR067C, YDR105C, YDR176W, YFR021W, YGL255W, YGR106C, YIL125W,
       YIL154C, YKL201C, YKR001C, YLL045C, YLR047C, YLR180W, YML017W, YML018C,
       YML111W, YMR179W, YMR242W-A, YOL124C, YOR376W, YPL055C, YPL177C, YPR190C}
Out[ • ]=
      \{115, 24, 20.87\%\}
      By contrast, for the 333 genic SNPs, 121 (36.34%) were annotated as having metal tolerance, which is
      substantially higher (by 74%) than for the further intergenic SNPs.
      These are the set of genes with both close intergenic and genic SNPS in the dataset:
 In[*]:= bothgenicintergenic = Intersection[genelistSYS, genelistCLOSESYS]
      Length[%]
Out[ • ]=
       {YAL051W, YAL053W, YDR089W, YER178W, YFR047C, YGL133W, YGR032W,
       YHL029C, YIL047C, YLL026W, YLR024C, YML049C, YNL240C, YOL006C, YPR078C}
```

7 out of these 15 have metal tolerance effects noted in SGD:

```
In[*]:* whichones = Intersection[bothgenicintergenic, metalgenenames]
      {Length[bothgenicintergenic], Length[%],
       Length[%] / Length[bothgenicintergenic] // N // PercentForm}
Out[ • ]=
      {YAL051W, YAL053W, YDR089W, YER178W, YIL047C, YLR024C, YOL006C}
Out[ • ]=
      {15, 7, 46.67%}
 In[*]:= Select[regpotential, MemberQ[whichones, #[2]] &] // Sort // MatrixForm
        FLC2 YAL053W 412
                              c.-412C>T
       FLC2 YAL053W 3064
                              c.-3064C>T
                             c.-31C>T
       OAF1 YAL051W 31
       PDA1 YER178W 13
                               c.-13C>A
       SYG1 YIL047C 48 c.-48_-47insT
                             c.-244delA
       TOP1 YOL006C 244
       UBR2 YLR024C 104
                              c.-104delT
       VTC5 YDR089W 232 c.-232_-231delAA
      Appending metal resistance phenotype information to the data:
 In[*]:= For[i = 1; newcolumn = {}, i ≤ Length[fullgenomeSNPEFF], i++,
       nameSYS = fullgenomeSNPEFF[[i, 11]];
       If[MemberQ[metalgenenames, nameSYS],
        AppendTo[newcolumn, "METAL"], AppendTo[newcolumn, "No"]]
      1
 In[*]:= fullgenomeSNPEFF = Transpose[Append[Transpose[fullgenomeSNPEFF], newcolumn]];
      Phosphorus metabolic process:
      The list of genes affecting phosphorous metabolism was downloaded from Amigo 2 (https://amigo.ge-
      neontology.org/amigo/term/GO:0006793) on 17 June2025:
 In[*]:= pholist = Import[
          "/Users/otto/Documents/Students/AnnaBazzicalupo/MetalAdaptation/Genomics/
            phosphate_metabolism_annotations_17Jun2025.tsv", "tsv"];
 In[*]:= phohead = pholist[[1]]
Out[ • ]=
      {gene, gene_id, gene_name, go_id, evidence}
 In[*]:= pholist = Drop[pholist, 1];
      The gene name but not the systematic name is listed:
 In[*]:= phogenenames = pholist[All, 1];
      Length[%]
Out[ • ]=
      480
```

Of the 287 genes with upstream intergenic mutations, 20 (7.0%) are annotated as affecting phosphorous metabolism:

1

```
In[@]:= intergenelist = Union[Sort[regpotential] [All, 1]]
      intergenelist = Drop[intergenelist, 1];
Out[ • ]=
      {, AAC3, ADE5,7, ADH4, ADH7, AGA1, ALE1, ALT2, AMF1, ARC15, ARG4, ARO7, ATG18, ATH1,
       AVO2, BET4, BIO2, BMT5, BNA6, BRX1, BUB2, BUD3, BUL2, CBP4, CCA1, CLB5, CNL1,
       COF1, COG5, COQ8, CRG1, CSE1, CUP9, CWC27, DAD3, DAM1, DAT1, DDR2, DLD3, DNF2,
       DNF3, EAP1, EHT1, EMI2, EP01, ERB1, ERG26, ERR2, ESF1, ETT1, FAP7, FCY21, FLC2,
       FLC3, FLO1, FLO10, FMN1, FRA1, FRE8, GCN3, GDH3, GEX1, GLC3, GLC8, GLO3, GPI13,
       GPR1, GSC2, GUA1, HFA1, HMG2, HOR7, HOT1, HOT13, HRD1, HSK3, HSP104, HXT7, HYM1,
       IML2, IRC3, ITC1, KGD1, KTI11, LDB16, LGE1, LIA1, LYS9, MDN1, MEK1, MIX17, MLP1,
       MMS1, MMT1, MNN4, MNN5, MRH1, MRP49, MRPL9, MSC6, MSF1, MSO1, MSS4, MST1, NAM2,
       NAR1, NBP1, NCE102, NDD1, NGG1, NOC2, NOP4, NOP56, NOP9, NSR1, OAF1, OCA5, OCA6,
       OM45, ORC4, PAD1, PAH1, PAU10, PAU17, PDA1, PDC6, PDS5, PEP4, PEX2, PGA1, PHO80,
       PH091, PIL1, PMT4, PMT6, POM34, POP2, PPT2, PPZ1, PRE10, PRM7, PRO2, PRP11, PSA1,
       PSP2, PUF4, RAD14, RAS1, RBD2, RCK1, RFC3, RPA12, RPC82, RPH1, RPL11B, RPL17B,
       RPL18A, RPL22B, RPL3, RPL34B, RPL4B, RPL8B, RP026, RP041, RPS0A, RPS22A, RRM3,
       RSA1, RSE1, SAM1, SAM50, SCM4, SDA1, SDD2, SDD4, SD01, SEC12, SEC27, SEC31, SET5,
       SMA1, SMC6, SMI1, SNR52, SNR61, SNR63, SOD2, SPC42, SPO21, SPT21, SQT1, SSA1,
       SSF2, SWI6, SYC1, SYG1, TCD2, TDA8, THI2, TIF5, TIM10, TMS1, TOK1, TOM7, TOP1,
       TOS8, TRE1, TRM11, UBP13, UBR2, UBX4, URA2, USO1, VHR1, VMA13, VOA1, VPH1, VPS1,
       VPS54, VPS68, VTA1, VTC5, WTM1, YAE1, YAL042C-A, YAP1802, YBL070C, YBL111C,
       YBR300C, YCL012C, YCR006C, YDL022C-A, YDL180W, YDL186W, YDL240C-A, YDR034C-D,
       YDR169C-A, YDR537C, YEL043W, YER076C, YER137C, YFL021C-A, YGL014C-A, YGL063C-A,
       YGR125W, YGR269W, YGR293C, YHL005C, YHR139C-A, YHR140W, YIL154C, YIL169C,
       YIR018C-A, YJL045W, YJL107C, YJL169W, YJL193W, YKE2, YKL031W, YKL162C-A, YKR005C,
       YLL054C, YLL066W-A, YLR302C, YME2, YML018C, YMR001C-A, YMR182W-A, YMR242W-A,
       YNL190W, YNR062C, YOL150C, YOP1, YOR121C, YOR343W-A, YOR376W, YOR392W, YPK9,
       YPL068C, YPL222C-A, YPL257W-A, YPQ2, YPR014C, YPR078C, YPR142C, YR02, ZRT1, ZWF1}
 In[*]:= Intersection[intergenelist, phogenenames]
      {Length[intergenelist], Length[%],
       Length[%] / Length[intergenelist] // N // PercentForm}
Out[ • ]=
      {ALE1, BNA6, EMI2, ERR2, FAP7, FMN1, GPI13, GUA1, HFA1,
       HMG2, MSS4, PAH1, PDA1, PGA1, PH091, PSA1, URA2, VPH1, VTC5, ZWF1}
Out[•]=
      {287, 20, 6.969%}
      Appending phosphorous metabolism phenotype information to the data:
 In[*]:= For[i = 1; newcolumn = {}, i ≤ Length[fullgenomeSNPEFF], i++,
       name = fullgenomeSNPEFF[[i, 8]];
       If[MemberQ[phogenenames, name],
        AppendTo[newcolumn, "PHO"], AppendTo[newcolumn, "No"]]
```

```
In[*]:= fullgenomeSNPEFF = Transpose[Append[Transpose[fullgenomeSNPEFF], newcolumn]];
      DNA Repair:
      The list of genes affecting DNA repair was downloaded from Amigo 2 (https://amigo.geneontolo-
      gy.org/amigo/term/GO:0006281) on 17 June 2025:
 In[*]:= repairlist = Import[
          "/Users/otto/Documents/Students/AnnaBazzicalupo/MetalAdaptation/Genomics/
            dna_repair_amigo_17Jun2025.tsv", "tsv"];
 In[*]:= repairhead = repairlist[[1]]
Out[ • ]=
       {gene, gene_id, gene_name, go_id, evidence}
 In[*]:= repairlist = Drop[repairlist, 1];
 In[*]:= repairlist[2]
Out[ • ]=
       {RFC2, SGD:S000003829,
       Subunit of heteropentameric Replication factor C (RF-C), GO:0006298, TAS}
      Here we will use the Gene Systematic Name for comparison:
 In[@]:= repairgenenames = Union[repairlist[All, 2]];
      Length[%]
Out[ • ]=
      322
      Of the 287 genes with upstream intergenic mutations, 6 (2.1%) are annotated as affecting DNA repair:
 Intersection[Union[genelistCLOSESYS, genelistFURTHERSYS], repairgenenames]
       {Length[Union[genelistCLOSESYS, genelistFURTHERSYS]], Length[%],
       Length[%] / Length[Union[genelistCLOSESYS, genelistFURTHERSYS]] // N // PercentForm}
Out[•]=
       { }
Out[ • ]=
      {287, 0, 0%}
      Appending DNA repair phenotype information to the data:
 In[•]:= For[i = 1; newcolumn = {}, i ≤ Length[fullgenomeSNPEFF], i++,
       nameSYS = fullgenomeSNPEFF[[i, 11]];
       If[MemberQ[repairgenenames, nameSYS],
         AppendTo[newcolumn, "Repair"], AppendTo[newcolumn, "No"]]
      1
 In[*]:= fullgenomeSNPEFF = Transpose[Append[Transpose[fullgenomeSNPEFF], newcolumn]];
```

Exporting

If the entire SNPEff and VCF information is needed:

Swapped variants positions:

```
In[*]:= newheader = Flatten[AppendTo[newheader, {"Metal resistance gene (SGD)",
             "Phosphorous metabolism (Amigo 2)", "DNA repair (SGD)"}]];
 In[@]:= toprint = Join[{newheader}, fullgenomeSNPEFF];
 In[*]:= Export["/Users/otto/Documents/Students/AnnaBazzicalupo/MetalAdaptation/Genomics/
            intergenicfullgenomeSNPEFF_Afterfiltering_17June2025.tsv", toprint];
      To print the data by line, we first recalculate the summary table:
 In[*]:= genomeSNPEFF = Table[-1, {i, 1, Length[fullgenomeSNPEFF]}, {j, 1, topos - droppos}];
       For[i = 1, i ≤ Length[fullgenomeSNPEFF], i++,
        For [j = 1 + droppos, j \le topos, j++,
         temp = StringSplit[fullgenomeSNPEFF[i, j], {":"}];
         temp2 = temp[1];
         genomeSNPEFF[[i, j - droppos]] = temp2;
        ]
       ]
      The following sites are "1"s in OLY077 and so we swap 1's for 0's and call these revertants
 In[@]:= Position[genomeSNPEFF, _ ? (#[1]] == "1" &) ] [All, 1]
       ••• Part: Part specification List[1] is longer than depth of object. 0
       ••• Part: Part specification List[1] is longer than depth of object. 🕡
       Part: Part specification 1[1] is longer than depth of object. 1
       😶 General: Further output of Part::partd will be suppressed during this calculation. 🕖
Out[ • ]=
       {1, 8, 9, 22, 31, 32, 36, 38, 41, 43, 44, 76, 88, 105, 114, 128,
        131, 138, 141, 144, 145, 177, 189, 194, 195, 206, 213, 253, 263, 275}
       Swapping 0<->1 and calling "." 0 (like OLY077):
 In[*]:= For[i = 1, i ≤ Length[%], i++,
        genomeSNPEFF[%[i]] =
         Mod[ToExpression[StringReplace[genomeSNPEFF[%[i]]], "." → "1"]] + 1, 2]
       ]
```

```
m[\cdot]:= fullgenomeSNPEFF[%%, {1, 3, Position[newheader, "Upstream of nearest gene"][1, 1]]]
Out[ • ]=
       {{chrI, 15592, TDA8}, {chrI, 198582, FL01}, {chrI, 198589, FL01},
        {chrII, 684 840, DAD3}, {chrIII, 309 030, ADH7}, {chrIV, 26 389, YDL240C-A},
        {chrIV, 164 943, FAP7}, {chrIV, 344 004, US01}, {chrIV, 376 575, PRP11},
        {chrIV, 392146, GPR1}, {chrIV, 410887, YDL022C-A}, {chrV, 407063, GL03},
        {chrVI, 193508, ATG18}, {chrVII, 496729, ERG26}, {chrVII, 808498, NSR1},
        {chrVIII, 123 954, SOD2}, {chrVIII, 379 942, YHR140W}, {chrIX, 57 277, YIL154C},
        {chrIX, 183 251, BMT5}, {chrIX, 257 480, RPL34B}, {chrIX, 267 870, SYG1},
        {chrXI, 379623, YKL031W}, {chrXII, 52019, RPL8B}, {chrXII, 179751, P0M34},
        {chrXII, 185314, SD01}, {chrXII, 881947, SMC6}, {chrXIII, 234752, PSP2},
        {chrXV, 45692, YOL150C}, {chrXV, 426603, ETT1}, {chrXV, 923126, PRO2}}
      All remaining "." are treated as identical to OLY077 (not mutant) and 0/1 treated as numbers (only in
       genomeSNPEFF, summarytable and fullgenomeSNPEFF remain the same):
 In[*]:= genomeSNPEFF = genomeSNPEFF /. "." → "0";
       genomeSNPEFF = ToExpression /@ genomeSNPEFF;
      Number of unique mutations:
 In[*]:= Length[genomeSNPEFF]
Out[ • ]=
      316
      Number of mutational hits:
 In[*]:= Total[Total[genomeSNPEFF]]
Out[ • ]=
      592
      As expected (once revertants were recoded), OLY077 has no mutations
 In[@]:= Total[genomeSNPEFF[All, 1]]]
Out[ • ]=
      Number of lines per unique mutation:
 In[@]:= Sort[Tally[Sum[genomeSNPEFF[All, i], {i, 1, 110}]]]
Out[ • ]=
       \{\{1, 252\}, \{2, 35\}, \{3, 13\}, \{4, 3\}, \{5, 2\}, \{7, 2\},
        \{10, 2\}, \{11, 1\}, \{12, 2\}, \{13, 1\}, \{14, 1\}, \{40, 1\}, \{73, 1\}\}
      Genes with excessive SNP mutations (≥5) include:
 In[*]:= toopoly = Position[genomeSNPEFF, ? (Total[#] ≥ 5 &)][All, 1]
Out[ • ]=
       \{1, 9, 42, 50, 54, 105, 111, 145, 184, 185, 186, 243, 300\}
      The genotype call for OLY077 at these sites:
```

n[∘]:= fullgenomeSNPEFF¶toopoly, Position[newheader, "OLY077"]¶1, 1∏∏ // MatrixForm

Out[•]//MatrixForm= 1:17,19:36:23:23,0 1:6,9:15:99:153,0 0:16,0:16:99:0,135 .:0,3:3:99:99,0 .:3,0:3:99:0,100 1:1,12:13:99:209,0 0:14,0:14:99:0,242 1:2,6:8:99:106,0 .:4,0:4:99:0,104 0:13,0:13:99:0,226 0:28,0:28:91:0,91 0:9,0:9:99:0,253 0:5,0:5:99:0,112

> To list the remaining mutant sites and the tally of mutations in them, we first update the summary table to carry the names and distances to the nearest gene, as well as the phenotypes:

```
In[*]:= summarytable[[1]]
Out[ • ]=
      {chrI, 15592, YAL065C, MODIFIER, {{0, 73}, {1, 36}, {., 1}}}
 In[*]:= summarytable[[1]]
Out[ • ]=
      {chrI, 15592, YAL065C, MODIFIER, {{0, 73}, {1, 36}, {., 1}}}
 In[*]:= summarytableTEMP = summarytable;
 In[*]:= For[i = 1, i ≤ Length[summarytable], i++,
       hold = summarytable[i, 5];
       summarytable[i, 3] =
        fullgenomeSNPEFF[i, Position[newheader, "Upstream of nearest gene"][1, 1]]];
       summarytable[i, 4] = fullgenomeSNPEFF[i,
          Position[newheader, "Systematic name of nearest gene"] [1, 1]];
       summarytable[i, 5] = fullgenomeSNPEFF[i, Position[newheader, "Distance"][1, 1]];
       AppendTo[summarytable[i],
        fullgenomeSNPEFF∏i, Position[newheader, "Effect on closest"]∏1, 1∏∏];
       AppendTo[summarytable[i],
         fullgenomeSNPEFF[i, Position[newheader, "Metal resistance gene (SGD)"][[1, 1]]]];
       AppendTo[summarytable[i]], fullgenomeSNPEFF[i,
          Position[newheader, "Phosphorous metabolism (Amigo 2)"][1, 1]]];
       AppendTo[summarytable[i], fullgenomeSNPEFF[i,
          Position[newheader, "DNA repair (SGD)"][1, 1]]]];
       AppendTo[summarytable[i], hold]
      1
```

In[*]:= consider = summarytable [toopoly]; consider // MatrixForm Out[•]//MatrixForm= chrI 15 592 TDA8 YAL064C-A 1849 c.-1849C>GchrI 198589 FL01 YAR050W 4814 c.-4814A>GchrIV 384303 PRM7 YDL039C 223 c.-250_-223delCGTAGTCAGTTCAAGTTCAGCTGATCTA chrIV 621879 VTC5 YDR089W 232 c.-232_-231delAA chrIV 794798 YDR169C-A YDR169C-A 75 c.-75A>C chrVII 496729 ERG26 YGL001C 228 c.-228_-227insT chrVII 701186 VOA1 YGR106C 1401 c.-1401_-1400insACA chrIX 267870 SYG1 YIL047C 48 c.-48_-47insT chrXI 666614 chrXII 5851 YLL066W-A YLL066W-A 635 c.-635C>T chrXII 5854 YLL066W-A YLL066W-A 632 c.-632G>T chrXIV 493 053 TOM7 YNL070W 312 c.-312delA chrXVI 607820 SDD4 YPR022C 509 c.-509delA Depth of coverage for these genes is higher on average: In[*]:= fullgenomeSNPEFF[[toopoly, {Position[newheader, "Upstream of nearest gene"][1, 1], 129}] Mean[%[All, 2]] // N Out[•]= {{TDA8, 4828}, {FL01, 1738}, {PRM7, 838}, {VTC5, 696}, {YDR169C-A, 657}, {ERG26, 1141}, {VOA1, 1355}, {SYG1, 901}, {, 606}, {YLL066W-A, 1016}, {YLL066W-A, 1073}, {TOM7, 779}, {SDD4, 676}} Out[•]= 1254.15 Mapping quality is also slightly poorer on average: In[*]:= fullgenomeSNPEFF[toopoly, {Position[newheader, "Upstream of nearest gene"][1, 1], 133}] Mean[%[All, 2]] // N Length[Select[fullgenomeSNPEFF[All, 133], # < % &]] /</pre> Length[fullgenomeSNPEFF[All, 133]] // N // PercentForm Out[•]= {{TDA8, 55.02}, {FL01, 56.74}, {PRM7, 52.7}, {VTC5, 59.97}, {YDR169C-A, 60.}, {ERG26, 60.}, {VOA1, 59.95}, {SYG1, 60.}, {, 58.61}, {YLL066W-A, 55.18}, {YLL066W-A, 54.84}, {TOM7, 59.05}, {SDD4, 60.}} Out[•]= 57.8508

The set of highly polymorphic (and likely wrong) genes:

Out[•]//PercentForm=

6.329%

```
In[*]:= toprintmultiple =
        Sort[Table[{summarytable[toopoly[i], 3], summarytable[toopoly[i], 4],
            StringJoin[fullgenomeSNPEFF[toopoly[i]], 1]], ".",
            ToString[fullgenomeSNPEFF[toopoly[i]], 3]]], summarytable[toopoly[i]], 5],
            summarytable[toopoly[i], 6], summarytable[toopoly[i], 7],
            summarytable[toopoly[i], 8], summarytable[toopoly[i], 9],
            summarytable[toopoly[i]], 10], genomeSNPEFF[toopoly[i]].
             newheader[[1 + droppos;; topos]]}, {i, 1, Length[toopoly]}]];
      MatrixForm[%]
Out[ • ]//MatrixForm=
                             chrXI.666614
                   YGL001C chrVII.496729 228
         ERG26
                                                               c.-228_-227insT
          FL01
                   YAR050W
                             chrI.198589 4814
                                                                 c.-4814A>G
          PRM7
                   YDL039C chrIV.384303 223 c.-250_-223delCGTAGTCAAGTTCAAGTTCAGCTGATCTA
                  YPR022C chrXVI.607820 509
          SDD4
                                                                 c.-509delA
          SYG1
                  YIL047C chrIX.267870
                                            48
                                                                c.-48_-47insT
          TDA8
                  YAL064C-A chrI.15592 1849
                                                                 c.-1849C>G
          TOM7
                   YNL070W chrXIV.493053 312
                                                                 c.-312delA
          VOA1
                   YGR106C chrVII.701186 1401
                                                           c.-1401_-1400insACA
          VTC5
                  YDR089W chrIV.621879 232
                                                              c.-232_-231delAA
       YDR169C-A YDR169C-A chrIV.794798
                                            75
                                                                  c.-75A>C
       YLL066W-A YLL066W-A chrXII.5851
                                            635
                                                                  c.-635C>T
       YLL066W-A YLL066W-A chrXII.5854
                                            632
                                                                  c.-632G>T
 In[*]:= Export["/Users/otto/Documents/Students/AnnaBazzicalupo/MetalAdaptation/Genomics/
          intergenictoopolymorphic_17June2025.tsv", Join[
         {{"Gene", "Systematic name", "chrom.pos", "Distance from gene (upstream only)",
            "SNP effect", "Metal resistance gene (SGD)",
           "Phosphorous metabolism (Amigo 2)", "DNA repair (SGD)", "# lines",
            "Lines carrying variant not seen in ancestor (OLY077)"}}, toprintmultiple]];
      The remaining sites:
 In[*]:= rest = Complement[Table[i, {i, 1, Length[fullgenomeSNPEFF]}], toopoly];
      Length[rest]
Out[ • ]=
      303
 In[*]:= Length[rest] + Length[toopoly] - Length[fullgenomeSNPEFF]
Out[ • ]=
 In[*]:= summarytable[1]
Out[ • ]=
      {chrI, 15592, TDA8, YAL064C-A, 1849, c.-1849C>G, No, No, No, {{0, 73}, {1, 36}, {., 1}}}
```

```
In[@]:= toprint = Table[Flatten[{summarytable[rest[i]], 3]],
           summarytable[rest[i]], 4]], StringJoin[fullgenomeSNPEFF[rest[i]], 1], ".",
            ToString[fullgenomeSNPEFF[[rest[[i]], 3]]]], summarytable[[rest[[i]], 5;; 9]],
           genomeSNPEFF[[rest[i]]].newheader[[1 + droppos;; topos]]}], {i, 1, Length[rest]}];
      MatrixForm[%[1;; 12]]
      (*The first 12 lines shows the nature of parallel SNPs in different lines*)
Out[ • ]//MatrixForm=
         GDH3
                  YAL062W
                            chrI.30796
                                        771
                                                       c.-771G>A
                                                                            No
                                                                                 No
                                                                                     No
         FLC2
                  YAL053W
                           chrI.42835 3064
                                                      c.-3064C>T
                                                                            No
                                                                                 No
                                                                                     No
                  YAL053W chrI.45487 412
         FLC2
                                                       c.-412C>T
                                                                            No
                                                                                 PHO No
         OAF1
                  YAL051W chrI.48533
                                         31
                                                       c.-31C>T
                                                                            No
                                                                                 PHO No
       YAL042C-A YAL042C-A chrI.65761 4154
                                                     c.-4154delA
                                                                            No
                                                                                     No
                                                                                 No
                  YAL005C chrI.141762 331
         SSA1
                                                      c.-331C>G
                                                                          METAL
                                                                                 No
                                                                                     No
         FL01
                  YAR050W chrI.198582 4821
                                                      c.-4821G>A
                                                                            No
                                                                                 No
                                                                                     No Cd
        YBL111C
                  YBL111C chrII.6157 1148
                                                      c.-1148A>G
                                                                                 No No
                                                                            No
                  YBL111C
        YBL111C
                           chrII.6158 1149
                                                      c.-1149T>C
                                                                            No
                                                                                 No
                                                                                     No
                  YBL111C chrII.6175 1166
        YBL111C
                                                      c.-1166A>G
                                                                            No
                                                                                 No
                                                                                     No
        YBL111C
                  YBL111C chrII.6181 1173 c.-1173_-1172insAGTGGTGGTGGT
                                                                            No
                                                                                 No No
                            chrII.6191 1182
        YBL111C
                  YBL111C
                                                      c.-1182G>A
                                                                            No
                                                                                 No No
```

For each SNP that occurs multiple times, make a separate row:

```
In[*]:= namepos = 9; (*Line names*)
     For[i = 1; toappend = {}, i ≤ Length[toprint], i++,
      splits = Length[toprint[i, namepos]];
      If[splits > 1,
       temp = toprint[i, namepos];
        toprint[i, namepos] = toprint[i, namepos][1];
        For [j = 2, j \le splits, j++,
         toadd = toprint[i];
         toadd[namepos] = temp[j];
         toappend = Append[toappend, toadd]
        ]
      ]
     1
```

Sorting then putting the metal first:

```
In[*]:= Sort[Join[toprint, toappend]];
      mat = Transpose[
          Join[{Transpose[%[All, namepos]]]}, Transpose[%[All, 1;; (namepos - 1)]]]]];
      MatrixForm[mat[1;; 12]]
Out[•]//MatrixForm=
        CdBM45
                                  chrV.21321
                                                                       No
                                                                                No
                                                                             No
       MnBM24
                                chrVIII.543677
                                                                       No
                                                                             No
                                                                                 No
       CdBM43
                AAC3 YBR085W
                                 chrII.415764
                                                                     METAL
                                                 218
                                                       c.-218delT
                                                                             No
                                                                                 No
       CdBM26 ADE5,7 YGL234W
                                 chrVII.56225
                                                 256
                                                        c.-256delT
                                                                       No
                                                                             No
                                                                                 No
       CdBM37
                ADH4
                       YGL256W
                                 chrVII.14846
                                                 312
                                                        c.-312delA
                                                                       No
                                                                             No
                                                                                 No
                ADH7 YCR105W
       CdBM23
                                chrIII.309030
                                                 39 c.-40_-39insA METAL No
                                                                                No
       MnBM17
               AGA1 YNR044W chrXIV.703150
                                                 549
                                                        c.-549C>G
                                                                       No
                                                                             No
                                                                                No
       CdBM29
                ALE1
                       YOR175C
                                chrXV.662028
                                                 354
                                                        c.-354A>T
                                                                       No
                                                                            PHO No
       CdBM42 ALE1 YOR175C
                                chrXV.665416
                                                3743 c.-3743delT
                                                                            PHO No
                                                                       No
       MnBM27 ALT2 YDR111C
                                chrIV.680189
                                                 425
                                                        c.-425T>C
                                                                       No
                                                                            PHO No
       CdBM46
                AMF1
                       YOR378W
                                 chrXV.1048412
                                                 1098
                                                       c.-1098delT
                                                                       No
                                                                             No
                                                                                 No
       MnBM42 ARC15 YIL062C
                                 chrIX.244607
                                                                            PHO No
                                                 145
                                                        c.-145C>T
                                                                       No
      The length matches that expected from the tally of multiply hit genes:
 In[*]:= Length[mat]
Out[ • ]=
      373
 In[*]:= Sort[Tally[Sum[genomeSNPEFF[All, i], {i, 1, 110}]]]
      %[All, 1]].%[All, 2]
Out[ • ]=
      \{\{1, 252\}, \{2, 35\}, \{3, 13\}, \{4, 3\}, \{5, 2\}, \{7, 2\},
       \{10, 2\}, \{11, 1\}, \{12, 2\}, \{13, 1\}, \{14, 1\}, \{40, 1\}, \{73, 1\}\}
Out[ • ]=
      592
 In[*]:= Export["/Users/otto/Documents/Students/AnnaBazzicalupo/MetalAdaptation/Genomics/
           intergenicSNPs_17June2025.tsv", Join[
          {{"Line", "Gene", "Gene_Code", "chrom.pos", "Distance from gene (upstream only)",
            "Mutation", "Metal resistance gene (SGD)",
            "Phosphorous metabolism (Amigo 2)", "DNA repair (SGD)"}}, mat]];
      Tally by line:
 In[*]:= newheader[[16;; 124]] // Length
Out[ • ]=
      109
```

```
In[*]:= bymetal = Tally[Sort[Join[mat[All, 1]], newheader[[16;; 124]]]]];
      bymetal[All, 2] = bymetal[All, 2] - 1;
      bymetal
Out[ • ]=
      {{CdBM23, 16}, {CdBM25, 2}, {CdBM26, 17}, {CdBM29, 15}, {CdBM30, 12}, {CdBM32, 18},
        {CdBM36, 0}, {CdBM37, 23}, {CdBM39, 18}, {CdBM42, 18}, {CdBM43, 21}, {CdBM44, 7},
        {CdBM45, 3}, {CdBM46, 3}, {CdBM47, 3}, {CdBM48, 7}, {CoBM1, 4}, {CoBM12, 4},
        {COBM14, 2}, {COBM15, 5}, {COBM16, 0}, {COBM17, 0}, {COBM18, 5}, {COBM2, 0},
        {COBM20, 1}, {COBM21, 2}, {COBM3, 0}, {COBM4, 0}, {COBM5, 4}, {COBM6, 4}, {COBM7, 0},
        {CoBM8, 6}, {CuBM10, 1}, {CuBM11, 2}, {CuBM12, 1}, {CuBM13, 4}, {CuBM14, 0},
        {CuBM15, 3}, {CuBM16, 1}, {CuBM17, 0}, {CuBM18, 1}, {CuBM3, 0}, {CuBM4, 0},
        {CuBM6, 0}, {CuBM7, 1}, {CuBM8, 0}, {CuBM9, 4}, {MnBM12, 0}, {MnBM13, 0},
        {MnBM14, 12}, {MnBM15, 2}, {MnBM16, 2}, {MnBM17, 6}, {MnBM18, 3}, {MnBM20, 0},
        {MnBM21, 0}, {MnBM23, 0}, {MnBM24, 4}, {MnBM25, 2}, {MnBM27, 2}, {MnBM28, 2},
        {MnBM29, 2}, {MnBM31, 1}, {MnBM32, 5}, {MnBM34, 1}, {MnBM38, 3}, {MnBM39, 1},
        {MnBM42, 37}, {MnBM44, 0}, {NiBM11, 0}, {NiBM12, 2}, {NiBM14, 1}, {NiBM16, 2},
        {NiBM17, 1}, {NiBM21, 2}, {NiBM22, 2}, {NiBM24, 1}, {NiBM25, 0}, {NiBM27, 2},
        {NiBM28, 0}, {NiBM29, 1}, {NiBM30, 0}, {NiBM4, 1}, {NiBM6, 0}, {NiBM8, 0},
        {NiBM9, 0}, {ZnBM11, 1}, {ZnBM12, 3}, {ZnBM15, 1}, {ZnBM16, 2}, {ZnBM17, 1},
        {ZnBM19, 2}, {ZnBM22, 1}, {ZnBM23, 0}, {ZnBM25, 5}, {ZnBM28, 2}, {ZnBM29, 1},
        {ZnBM31, 1}, {ZnBM34, 1}, {ZnBM37, 0}, {ZnBM38, 0}, {ZnBM39, 0}, {ZnBM41, 1},
        {ZnBM42, 1}, {ZnBM43, 0}, {ZnBM44, 6}, {ZnBM45, 2}, {ZnBM46, 3}, {ZnBM47, 1}}
 In[*]:= Total[bymetal[All, 2]]]
Out[ • ]=
      373
```

Gene list with all lines:

In[*]:= Sort[Union[mat[All, 2]]] Length[%]

Out[•]=

{, AAC3, ADE5,7, ADH4, ADH7, AGA1, ALE1, ALT2, AMF1, ARC15, ARG4, ARO7, ATG18, ATH1, AVO2, BET4, BIO2, BMT5, BNA6, BRX1, BUB2, BUD3, BUL2, CBP4, CCA1, CLB5, CNL1, COF1, COG5, COQ8, CRG1, CSE1, CUP9, CWC27, DAD3, DAM1, DAT1, DDR2, DLD3, DNF2, DNF3, EAP1, EHT1, EMI2, EP01, ERB1, ERR2, ESF1, ETT1, FAP7, FCY21, FLC2, FLC3, FLO1, FLO10, FMN1, FRA1, FRE8, GCN3, GDH3, GEX1, GLC3, GLC8, GLO3, GPI13, GPR1, GSC2, GUA1, HFA1, HMG2, HOR7, HOT1, HOT13, HRD1, HSK3, HSP104, HXT7, HYM1, IML2, IRC3, ITC1, KGD1, KTI11, LDB16, LGE1, LIA1, LYS9, MDN1, MEK1, MIX17, MLP1, MMS1, MMT1, MNN4, MNN5, MRH1, MRP49, MRPL9, MSC6, MSF1, MSO1, MSS4, MST1, NAM2, NAR1, NBP1, NCE102, NDD1, NGG1, NOC2, NOP4, NOP56, NOP9, NSR1, OAF1, OCA5, OCA6, OM45, ORC4, PAD1, PAH1, PAU10, PAU17, PDA1, PDC6, PDS5, PEP4, PEX2, PGA1, PH080, PH091, PIL1, PMT4, PMT6, P0M34, P0P2, PPT2, PPZ1, PRE10, PR02, PRP11, PSA1, PSP2, PUF4, RAD14, RAS1, RBD2, RCK1, RFC3, RPA12, RPC82, RPH1, RPL11B, RPL17B, RPL18A, RPL22B, RPL3, RPL34B, RPL4B, RPL8B, RP026, RP041, RPS0A, RPS22A, RRM3, RSA1, RSE1, SAM1, SAM50, SCM4, SDA1, SDD2, SDD4, SDO1, SEC12, SEC27, SEC31, SET5, SMA1, SMC6, SMI1, SNR52, SNR61, SNR63, SOD2, SPC42, SP021, SPT21, SQT1, SSA1, SSF2, SWI6, SYC1, TCD2, THI2, TIF5, TIM10, TMS1, TOK1, TOP1, TOS8, TRE1, TRM11, UBP13, UBR2, UBX4, URA2, USO1, VHR1, VMA13, VPH1, VPS1, VPS54, VPS68, VTA1, WTM1, YAE1, YAL042C-A, YAP1802, YBL070C, YBL111C, YBR300C, YCL012C, YCR006C, YDL022C-A, YDL180W, YDL186W, YDL240C-A, YDR034C-D, YDR537C, YEL043W, YER076C, YER137C, YFL021C-A, YGL014C-A, YGL063C-A, YGR125W, YGR269W, YGR293C, YHL005C, YHR139C-A, YHR140W, YIL154C, YIL169C, YIR018C-A, YJL045W, YJL107C, YJL169W, YJL193W, YKE2, YKL031W, YKL162C-A, YKR005C, YLL054C, YLR302C, YME2, YML018C, YMR001C-A, YMR182W-A, YMR242W-A, YNL190W, YNR062C, YOL150C, YOP1, YOR121C, YOR343W-A, YOR376W, YOR392W, YPK9, YPL068C, YPL222C-A, YPL257W-A, YPQ2, YPR014C, YPR078C, YPR142C, YR02, ZRT1, ZWF1}

Out[•]=

279

Gene list without the mutator lines MnBM14 and MnBM42:

In[*]:= Union[Sort[Select[mat, MemberQ[{"MnBM14", "MnBM42"}, #[1]] == False &] [All, 2]]] Length[%]

Out[•]=

{, AAC3, ADE5,7, ADH4, ADH7, AGA1, ALE1, ALT2, AMF1, ARG4, ARO7, ATG18, ATH1, AVO2, BET4, BIO2, BMT5, BNA6, BRX1, BUB2, BUD3, CCA1, CLB5, CNL1, COF1, COG5, CRG1, CSE1, CUP9, CWC27, DAD3, DAM1, DAT1, DDR2, DLD3, DNF2, DNF3, EHT1, EMI2, EP01, ERB1, ERR2, ESF1, ETT1, FAP7, FCY21, FLC3, FLO1, FLO10, FRA1, FRE8, GCN3, GEX1, GLC3, GLO3, GPI13, GPR1, GSC2, GUA1, HMG2, HOR7, HOT1, HOT13, HXT7, HYM1, IML2, IRC3, ITC1, KGD1, LDB16, LGE1, LIA1, LYS9, MDN1, MEK1, MIX17, MLP1, MNN4, MNN5, MRH1, MRP49, MRPL9, MSC6, MSF1, MSO1, MSS4, MST1, NBP1, NCE102, NDD1, NGG1, NOP4, NOP56, NOP9, NSR1, OCA5, ORC4, PAU17, PDA1, PDS5, PEP4, PEX2, PGA1, PHO80, PH091, PMT4, PMT6, POM34, POP2, PPT2, PPZ1, PRE10, PRO2, PRP11, PSP2, PUF4, RAD14, RAS1, RBD2, RCK1, RFC3, RPA12, RPC82, RPL18A, RPL22B, RPL3, RPL34B, RPL4B, RPL8B, RP026, RP041, RPS0A, RPS22A, RRM3, RSA1, RSE1, SAM1, SAM50, SCM4, SDA1, SDD2, SDD4, SDO1, SEC12, SEC31, SET5, SMA1, SMC6, SMI1, SNR52, SNR63, SOD2, SPC42, SPO21, SPT21, SQT1, SSA1, SWI6, SYC1, TCD2, THI2, TIF5, TIM10, TMS1, TOK1, TOP1, TOS8, UBP13, UBR2, UBX4, URA2, USO1, VHR1, VMA13, VPH1, VPS1, VPS54, VPS68, VTA1, WTM1, YAE1, YAL042C-A, YAP1802, YBL070C, YBL111C, YBR300C, YCL012C, YCR006C, YDL022C-A, YDL180W, YDL186W, YDL240C-A, YDR034C-D, YDR537C, YEL043W, YER137C, YFL021C-A, YGL014C-A, YGL063C-A, YGR125W, YGR269W, YGR293C, YHL005C, YHR140W, YIL154C, YIL169C, YIR018C-A, YJL045W, YJL169W, YJL193W, YKE2, YKL031W, YKL162C-A, YKR005C, YLL054C, YLR302C, YME2, YML018C, YMR001C-A, YMR182W-A, YMR242W-A, YNL190W, YNR062C, YOL150C, YOP1, YOR343W-A, YOR376W, YOR392W, YPK9, YPL068C, YPL222C-A, YPQ2, YPR014C, YPR078C, YPR142C, YR02, ZRT1}

Out[•]=

237

Statistical analyses

Fraction of genic vs intergenic mutations

```
ln[ \circ ] := vec = \{414, 316\};
     exp = \{0.7, 0.3\};
Out[ • ]=
     61.3764
```

Out[•]=

0.0408403

```
In[*]:= 1 - CDF[ChiSquareDistribution[1], %]
Out[ • ]=
       \textbf{4.66294} \times \textbf{10}^{-15}
       Fraction of metal-associated close vs further intergenic mutations
 In[*]:= tab = Join[{{"Gene", "Intergenic Close", "Intergenic Far"}}, Transpose[
             {{333, 121, 36.34%}, {176, 56, 31.82%}, {115, 24, 20.87%}}]] // Transpose;
       tab // TableForm
Out[•]//TableForm=
       Gene
                              333
                                      121
                                              0.363363
       Intergenic Close
                              176
                                      56
                                              0.318182
                                              0.208696
       Intergenic Far
                              115
                                      24
 ln[*]:= tabCF = {tab[[2][[2]; 3] - {tab[[2][[3], 0}, tab[[3][[2]; 3]] - {tab[[3][[3], 0}};
       tabCF // TableForm
Out[ • ]//TableForm=
              56
       120
               24
       91
       The row proportions:
 In[*]:= row = Total[Transpose[tabCF]] / Total[Total[Transpose[tabCF]]] // N
Out[ • ]=
       {0.604811, 0.395189}
       The column proportions:
 In[@]:= col = Total[tabCF] / Total[Total[tabCF]] // N
Out[ • ]=
       {0.725086, 0.274914}
       The expected matrix with the same row and column sums:
 In[@]:= expCF = Total[Total[tabCF]] x Transpose[{row}].{col}
Out[ • ]=
       \{\{127.615, 48.3849\}, \{83.3849, 31.6151\}\}
 In[•]:= Total [Total [ \frac{(tabCF - expCF)^2}{expCF} ]]
Out[ • ]=
       4.18263
 In[*]:= 1 - CDF[ChiSquareDistribution[1], %]
```

Fraction of metal-associated close vs genic mutations

```
In[*]:= tab = Join[{{"Gene", "Intergenic Close", "Intergenic Far"}}, Transpose[
             {{333, 121, 36.34%}, {176, 56, 31.82%}, {115, 24, 20.87%}}]] // Transpose;
      tab // TableForm
Out[•]//TableForm=
                              333
                                      121
                                             0.363363
                              176
       Intergenic Close
                                      56
                                             0.318182
                                             0.208696
       Intergenic Far
                              115
                                      24
 In[*]:= tabCF = {tab[[2][[2]; 3] - {tab[[2][[3]], 0}, tab[[1][[2]; 3]] - {tab[[1][[3]], 0}};
       tabCF // TableForm
Out[ • ]//TableForm=
              56
       120
       212
              121
      The row proportions:
 In[*]:= row = Total[Transpose[tabCF]] / Total[Total[Transpose[tabCF]]] // N
Out[ • ]=
       \{0.345776, 0.654224\}
      The column proportions:
 In[@]:= col = Total[tabCF] / Total[Total[tabCF]] // N
Out[ • ]=
       {0.652259, 0.347741}
      The expected matrix with the same row and column sums:
 In[@]:= expCF = Total[Total[tabCF]] x Transpose[{row}].{col}
Out[ • ]=
       \{\{114.798, 61.2024\}, \{217.202, 115.798\}\}
 In[*]:= Total \left[ \text{Total} \left[ \frac{(\text{tabCF - expCF})^2}{\text{expCF}} \right] \right]
Out[ • ]=
       1.0363
 In[*]:= 1 - CDF[ChiSquareDistribution[1], %]
Out[ • ]=
       0.308684
       Cross-tolerance by phenotypes
 Infepers = { ("CdBM23", "NO", "NO", "NO", 0.0408102148645912) },
           {"CdBM25", "Metal", "NO", "NO", 0.0390631227075351`},
           {"CdBM26", "Metal", "NO", "NO", 0.0737050353872304`},
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{"CdBM43", "NO", "NO", "NO", 0.0316342610972205`},
{"CdBM44", "Metal", "PHO", "NO", 0.0282256599435741`},
{"CdBM45", "Metal", "NO", "NO", 0.0473659618242343`},
{"CdBM46", "Metal", "NO", "NO", 0.0444138767868399`},
{"CdBM47", "Metal", "NO", "NO", 0.0701281571352517`},
{"CdBM48", "NO", "NO", "NO", 0.0492242624666969`},
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{"CoBM15", "Metal", "PHO", "NO", 0.0458346221316833`},
{"CoBM16", "Metal", "PHO", "NO", 0.0357453244509767`},
{"CoBM17", "Metal", "PHO", "Repair", 0.0597342559378414`},
{"CoBM18", "Metal", "PHO", "Repair", 0.0413293636583137`},
{"CoBM2", "Metal", "PHO", "NO", 0.0706202022286716`},
{"CoBM20", "Metal", "PHO", "NO", 0.0606191536353199`},
{"CoBM21", "Metal", "PHO", "NO", 0.0100054482758219`},
{"CoBM3", "Metal", "PHO", "NO", 0.0703312657166637`},
{"CoBM4", "Metal", "PHO", "NO", 0.0305670689864274`},
{"CoBM5", "Metal", "PHO", "NO", 0.0741003153178232`},
{"CoBM6", "Metal", "PHO", "NO", 0.0712429398379751`},
{"CoBM7", "Metal", "NO", "Repair", 0.0970299424029689`},
{"CoBM8", "Metal", "PHO", "NO", 0.0994528207296415`},
{"CuBM10", "Metal", "NO", "NO", 0.0342127642207387`},
{"CuBM11", "Metal", "NO", "NO", 0.034648617736055`},
{"CuBM12", "Metal", "NO", "NO", 0.0424992612246359`},
{"CuBM13", "Metal", "NO", "NO", 0.0659556115059866`},
{"CuBM14", "Metal", "PHO", "NO", 0.0276601268729923`},
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{"CuBM17", "Metal", "NO", "NO", 0.0318022493818922`},
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{"CuBM4", "Metal", "NO", "Repair", 0.0399190942736623`},
{"CuBM6", "Metal", "NO", "NO", 0.0296793452947361`},
{"CuBM7", "Metal", "NO", "Repair", 0.0409788621512595`},
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{"MnBM12", "Metal", "NO", "Repair", 0.0169372808934513`},
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{"MnBM14", "Metal", "PHO", "Repair", 0.0327898094910443`},
{"MnBM15", "NO", "NO", "NO", 0.011379586406605`},
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{"MnBM38", "Metal", "PHO", "NO", 0.0131807268686594`},
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{"MnBM42", "Metal", "PHO", "Repair", 0.01670830341897`},
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{"ZnBM15", "NO", "PHO", "NO", 0.0529235430658802`},
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{"ZnBM19", "Metal", "NO", "NO", 0.0309429263721979`},
{"ZnBM22", "Metal", "NO", "NO", 0.00630274392454666`},
{"ZnBM23", "Metal", "PHO", "NO", 0.111893288460111`},
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{"ZnBM38", "Metal", "PHO", "NO", 0.0640662934781045`},
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Out[•]=

Out[•]=

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         {"ZnBM46", "Metal", "NO", "NO", 0.0593915710521536`},
         {"ZnBM47", "Metal", "NO", "NO", 0.0202202824239883`}};
     Metal Resistance: Having or not having a mutation in a metal resistance gene is not a significant
     predictor of cross-tolerance:
     [Not expected]
in[*]:= tabMETAL = Select[linephenotypes, #[2] == "Metal" &] [All, 5];
In[*]:= tabNoMETAL = Select[linephenotypes, #[2] == "NO" &] [All, 5];
     p-value of t-test:
In[@]:= TTest[{tabMETAL, tabNoMETAL}]
     ••• TTest: At least one of the p−values in {0.0178719, 0.676747}, resulting from a test for normality, is below
          0.025`. The tests in {T} require that the data is normally distributed.
     0.321505
     Violin plot with mean and SE:
In[ • ]:= Show[
      DistributionChart[{tabMETAL, tabNoMETAL},
        ChartLegends → {"Metal", "No"}, ChartStyle → {Pink, LightBlue}],
       ErrorListPlot[
        {{1, Mean[tabMETAL], StandardDeviation[tabMETAL]/Sqrt[Length[tabMETAL]]}},
         {2, Mean[tabNoMETAL], StandardDeviation[tabNoMETAL] / Sqrt[Length[tabNoMETAL]]}},
        PlotStyle → Black],
      ListPlot[{{0.5, Mean[Join[tabMETAL, tabNoMETAL]]}},
         {2.5, Mean[Join[tabMETAL, tabNoMETAL]]}},
       PlotStyle → {Black, Dashed}, Joined → True]
     ]
     0.10
                                                            Metal
     0.05
                                                            ■ No
                                           Ŧ
     0.00
```

Phosphorous metabolism: Having or not having a mutation in a phosphorous metabolism gene does predict cross-tolerance:

```
in[*]:= tabPHO = Select[linephenotypes, #[3] == "PHO" &] [All, 5];
 In[@]:= tabNoPHO = Select[linephenotypes, #[3] == "NO" &] [All, 5];
 In[*]:= {Mean[tabPH0], Mean[tabNoPH0]}
Out[ • ]=
       {0.0430593, 0.0328959}
 In[@]:= {StandardDeviation[tabPH0] / Sqrt[Length[tabPH0]],
       StandardDeviation[tabNoPH0] / Sqrt[Length[tabNoPH0]]}
Out[ • ]=
       {0.00464377, 0.00263299}
      p-value of t-test:
 In[*]:= TTest[{tabPH0, tabNoPH0}]
Out[ • ]=
      0.0430228
      Violin plot with mean and SE:
 In[ • ]:= Show[
       DistributionChart[{tabPH0, tabNoPH0},
         ChartLegends → {"PHO", "No"}, ChartStyle → {Pink, LightBlue}],
        ErrorListPlot[
         {{1, Mean[tabPH0], StandardDeviation[tabPH0]/Sqrt[Length[tabPH0]]},
          {2, Mean[tabNoPH0], StandardDeviation[tabNoPH0]/Sqrt[Length[tabNoPH0]]}},
         PlotStyle → Black],
       ListPlot[{{0.5, Mean[Join[tabPHO, tabNoPHO]]}},
          {2.5, Mean[Join[tabPH0, tabNoPH0]]}}, PlotStyle → {Black, Dashed}, Joined → True]
      1
Out[ • ]=
      0.10
                                                           PHO
      0.05
                                                           ■ No
      0.00
```

Repair: Having or not having a mutation in a repair gene does predict cross-tolerance:

```
[Not expected]
 In[*]:= tabREPAIR = Select[linephenotypes, #[4] == "Repair" &] [All, 5];
 In[*]:= tabNoREPAIR = Select[linephenotypes, #[4] == "NO" &] [All, 5];
      p-value of t-test:
 In[@]:= TTest[{tabREPAIR, tabNoREPAIR}]
Out[ • ]=
      0.841256
      Violin plot with mean and SE:
 In[ • ]:= Show[
       DistributionChart[{tabREPAIR, tabNoREPAIR},
         ChartLegends → {"Repair", "No"}, ChartStyle → {Pink, LightBlue}],
       ErrorListPlot[
         {{1, Mean[tabREPAIR], StandardDeviation[tabREPAIR]/Sqrt[Length[tabREPAIR]]},
          {2, Mean[tabNoREPAIR], StandardDeviation[tabNoREPAIR] /
            Sqrt[Length[tabNoREPAIR]]}}, PlotStyle → Black],
       ListPlot[{{0.5, Mean[Join[tabREPAIR, tabNoREPAIR]]}},
          {2.5, Mean[Join[tabREPAIR, tabNoREPAIR]]}},
         PlotStyle → {Black, Dashed}, Joined → True]
      ]
Out[ • ]=
      0.10
                                                           Repair
      0.05
                                                           ■ No
      0.00
 In[*]:= temp = Union[mat[All, {2, 8, 9, 10}]]]
Out[ • ]=
      {{ABP1, Metal, No, No}, {ACB1, Metal, No, No}, {ACC1, No, PHO, No},
        {ADE6, No, PH0, No}, {ADR1, Metal, No, No}, {AFT1, Metal, PH0, No},
        {AGP1, No, No, No}, {AHK1, Metal, No, No}, {ALD5, No, No, No}, {ALY2, No, No, No},
        {APC2, No, No, No}, {ARO3, No, No, No}, {ASN1, No, No, No}, {ATG11, Metal, No, No},
        {ATG2, Metal, No, No}, {ATG39, No, No, No}, {BDS1, No, No, No}, {BIR1, No, No, No},
        {BLM10, Metal, No, Repair}, {BNA6, No, PHO, No}, {BNI1, Metal, No, No},
        {BNI4, Metal, No, No}, {BRR2, No, No, No}, {BSD2, Metal, No, No},
```

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{BUL1, Metal, No, No}, {BUL2, Metal, No, No}, {CAJ1, No, No, No},
{CCC2, Metal, No, No}, {CCR4, Metal, No, No}, {CCW12, No, No, No}, {CDC25, No, No, No},
{CIC1, No, No, No}, {CLB3, Metal, No, No}, {CLN3, Metal, No, No}, {CNE1, No, No, No},
{COG1, Metal, No, No}, {COG3, No, No, No}, {COI1, No, No, No}, {COQ1, No, No, No},
{CPA1, No, No, No}, {CSM3, No, No, Repair}, {CUE3, No, No, No}, {CWC22, No, No, No},
{CWH41, Metal, No, No}, {CYK3, No, No, No}, {DAL2, No, No, No}, {DBF20, No, No, No},
{DBP10, No, No, No}, {DDR48, Metal, No, Repair}, {DNF1, No, No, No},
{DOA4, Metal, No, No}, {DOT6, No, No, No}, {DSC2, No, No, No}, {DSE4, No, No, No},
{DUS4, No, No, No}, {EBS1, No, No, No}, {ECM14, No, No, No}, {ECM22, Metal, No, No},
{EFR3, No, No, No}, {ELO3, Metal, No, No}, {ERB1, No, No, No}, {ERG1, No, No, No},
{ERG27, No, No, No}, {ERG7, No, No, No}, {ERG9, No, PHO, No}, {FAS2, No, No, No},
{FAU1, No, No, No}, {FCY2, No, No, No}, {FET4, Metal, No, No}, {FIG2, No, No, No},
{FIG4, Metal, PH0, No}, {FLC2, Metal, No, No}, {FMP52, No, No, No},
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{RSE1, No, No, No}, {RTC2, No, No, No}, {SAK1, Metal, No, No}, {SAP4, No, No, No},
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                     No No
        HUL5 Metal No No
        MAM3 Metal No No
        PHO84 Metal PHO No
        PMA1
               No No No
        PTK2 Metal No No
        SFP1
              Metal No No
        SIW14 Metal PHO No
        VCX1 Metal No No
        VTC4
              Metal PHO No
        VTC5 Metal PHO No
       YGR130C No No No
        YOR1
                No
                     No No
```

Cross-tolerance by genes mutated in multiple metals

For the list of genes that were bore mutations in different metals, we might predict that they would be more cross tolerant. Here we explore the relationship between these lines and the phenotypes of interest.

```
In[*]:= multiplemetalgenelist =
         {"DNF1", "FYV10", "KSP1", "MAM3", "PDR1", "PH084", "PMA1", "PTK2", "SFP1",
          "VTC5", "YLR296W", "HUL5", "PPQ1", "SNT2", "TOM1", "YDL199C", "YOR1"};
 In[ • ]:= mat[1]
Out[ • ]=
      {CuBM18, ABP1, YCR088W, chrIII.265691, c.624A>C, p.Leu208Phe, MODERATE, Metal, No, No}
 In[@]:= Select[mat, MemberQ[multiplemetalgenelist, #[2]] &];
 In[*]:= linewithmultiplemetalhits = Union[%[All, 1]]
Out[ • ]=
      {CdBM25, CdBM29, CdBM30, CdBM32, CdBM39, CdBM45, CdBM46, CdBM47,
       CoBM1, CoBM12, CoBM15, CoBM18, CoBM21, CoBM3, CoBM4, CoBM7, CoBM8,
       CuBM17, CuBM3, CuBM6, CuBM9, MnBM13, MnBM14, MnBM16, MnBM17, MnBM31,
       MnBM39, MnBM42, NiBM11, NiBM16, NiBM4, NiBM9, ZnBM11, ZnBM17, ZnBM19,
       ZnBM29, ZnBM37, ZnBM38, ZnBM41, ZnBM42, ZnBM43, ZnBM45, ZnBM46, ZnBM47}
      This gets the phenotypes for the lines with and without hits in these genes:
 In[*]:= withmultmetalhits =
         Select[linephenotypes, MemberQ[linewithmultiplemetalhits, #[1]] &];
      withoutmultmetalhits =
         Select[linephenotypes, MemberQ[linewithmultiplemetalhits, #[1]] == False &];
      For example:
```

```
In[*]:= withmultmetalhits[[1]]
Out[ • ]=
       {CdBM25, Metal, NO, NO, 0.0390631}
      Cross tolerance - No significant difference in broad cross tolerance:
 In[*]:= {Mean[withmultmetalhits[All, 5]]}, Mean[withoutmultmetalhits[All, 5]]}
Out[ • ]=
       {0.0395263, 0.0337043}
 In[@]:= {StandardDeviation[withmultmetalhits[All, 5]]] /
         Sqrt[Length[withmultmetalhits[All, 5]]],
        StandardDeviation[withoutmultmetalhits[All, 5]] /
         Sqrt[Length[withoutmultmetalhits[All, 5]]]}
Out[•]=
       {0.00332774, 0.00334899}
 In[@]:= TTest[{withmultmetalhits[[All, 5]], withoutmultmetalhits[[All, 5]]}]
Out[ • ]=
      0.225657
      Metal resistance - There is significant association with metal resistance using a chi-squared goodness
      of fit test with df=1:
      [Code from https://mathematica.stackexchange.com/questions/5271/doing-a-chi-square-indepen-
```

dence-test-in-mathematica by whuber] In[*]:= {Sort[Tally[withmultmetalhits[All, 2]]]], Sort[Tally[withoutmultmetalhits[All, 2]]]}

```
Out[ • ]=
       {{{Metal, 38}, {NO, 6}}}, {{Metal, 37}, {NO, 17}}}
```

```
I_{n[a]} := data = \{ \%[1, 1, 2], \%[1, 2, 2] \}, \{ \%[2, 1, 2], \%[2, 2, 2] \} \};
      rc = {{"with", "without"}, {"yes", "no"}};
      TableForm[data, TableHeadings → rc]
      fit = Outer[Times, Total[data, {2}], Total[data]] / Total[data, 2];
       (*TableForm[fit//N,TableHeadings→rc]*)
      residual = data - fit;
       (*TableForm[residual//N,TableHeadings→rc]*)
      χ2array = residual^2/fit;
       (*TableForm[χ2array//N,TableHeadings→rc]*)
      \chi2 = Total[\chi2array, 2];
      \chi 2 // N
      df = (Times @@ # - Total[#] + 1) &[Dimensions[data]];
      pvalue = 1 - CDF[ChiSquareDistribution[df], \chi2] // N;
      Print["df=", df, ", p=", pvalue]
Out[ • ]//TableForm=
                  yes
                          no
      with
                          6
                   38
      without 37
                          17
Out[ • ]=
      4.29855
      df=1, p=0.0381448
```

PHO metabolism - There is no significant association with phosphorous metabolism using a chisquared goodness of fit test with df=1:

[Code from https://mathematica.stackexchange.com/questions/5271/doing-a-chi-square-independence-test-in-mathematica by whuber]

```
In[*]:= {Sort[Tally[withmultmetalhits[All, 3]]]}, Sort[Tally[withoutmultmetalhits[All, 3]]]}
Out[ • ]=
      {{{NO, 26}, {PHO, 18}}, {{NO, 39}, {PHO, 15}}}
```

```
I_{n[a]} := data = \{ \%[1, 1, 2], \%[1, 2, 2] \}, \{ \%[2, 1, 2], \%[2, 2, 2] \} \};
      rc = {{"with", "without"}, {"yes", "no"}};
      TableForm[data, TableHeadings → rc]
      fit = Outer[Times, Total[data, {2}], Total[data]] / Total[data, 2];
       (*TableForm[fit//N,TableHeadings→rc]*)
       residual = data - fit;
       (*TableForm[residual//N,TableHeadings→rc]*)
      χ2array = residual^2/fit;
       (*TableForm[χ2array//N,TableHeadings→rc]*)
      \chi2 = Total[\chi2array, 2];
      \chi 2 // N
      df = (Times @@ # - Total[#] + 1) &[Dimensions[data]];
      pvalue = 1 - CDF[ChiSquareDistribution[df], \chi2] // N;
      Print["df=", df, ", p=", pvalue]
Out[ • ]//TableForm=
                  yes
                          no
      with
                   26
                          18
      without 39
                          15
Out[ • ]=
      1.87181
      df=1, p=0.171268
```

Repair - There is no significant association with phosphorous metabolism using a chi-squared goodness of fit test with df=1:

[Code from https://mathematica.stackexchange.com/questions/5271/doing-a-chi-square-independence-test-in-mathematica by whuber]

```
In[*]:= {Sort[Tally[withmultmetalhits[All, 4]]]}, Sort[Tally[withoutmultmetalhits[All, 4]]]}
Out[ • ]=
      {{{NO, 37}, {Repair, 7}}, {{NO, 47}, {Repair, 7}}}
```

```
ln[\circ]:= data = \{ \{ [1, 1, 2], [1, 2, 2] \}, \{ [2, 1, 2], [2, 2, 2] \} \};
      rc = {{"with", "without"}, {"yes", "no"}};
      TableForm[data, TableHeadings → rc]
      fit = Outer[Times, Total[data, {2}], Total[data]] / Total[data, 2];
      (*TableForm[fit//N,TableHeadings→rc]*)
      residual = data - fit;
      (*TableForm[residual//N,TableHeadings→rc]*)
      χ2array = residual^2/fit;
      (*TableForm[χ2array//N,TableHeadings→rc]*)
      \chi2 = Total[\chi2array, 2];
      \chi 2 // N
      df = (Times @@ # - Total[#] + 1) &[Dimensions[data]];
      pvalue = 1 - CDF[ChiSquareDistribution[df], \chi2] // N;
      Print["df=", df, ", p=", pvalue]
Out[ • ]//TableForm=
                  yes
                         no
                          7
      with
                  37
      without 47
                          7
Out[ • ]=
      0.171857
```

df=1, p=0.678466