SUP MAT: Evolution of cross-tolerance to metals in yeast

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Data

Within the analysis for a particular metal (or for all metals combined), identical mutations (same exact SNP) were eliminated, as were mutations in MnBM14 and MnBM42. Rows do not sum to the "all" column because identical mutations observed in different metals were also removed when considering the dataset as a whole ("all"), but not from each metal considered in isolation. To assess parallelism across metals (but not within a metal), "difmetal" counts genes repeatedly hit ONLY in different metals (i.e., mutations in the same gene that occur within a metal are counted only once).

Out[•]//MatrixForm=

```
#hit all cd co cu mn ni zn difmetal
     186 43 63 18 47 17 20
                              200
 1
 2
     15
        2
            5
                   2
                      2
                        2
                               9
 3
     0
         1
            1
               0
                   1
                      1
                         2
                               1
 4
      3
         0
            1
               0
                   1
                      0
                               0
         0
            0
 6
         0
            1
                               0
      1
               0
                   0
                      0
 7
      0
         0
            0
               0
                   0
                      0
                        0
                               0
 8
      1
            0
                  0 0
 9
         0
            0
               0
                  0
                      0
                        0
                               0
         0 0
 10
               0
                   0
                      0 0
                               0
 11
         0 0
               0
                  0
                      0 0
                               0
 12
      0
         0
            0
               0
                      0
                               0
 13
            0
                               0
```

The number of independent mutations (leading to different SNPs) across all metals:

```
In[@]:= Drop[data[All, 2], 1].Drop[data[All, 1], 1]
Out[@]:=
270
```

The number of independent mutations (leading to different SNPs) across all metals, lumping together any mutations in the same gene that occur in the same metal (for testing parallelism across metals):

Out[•]=

```
In[@]:= Drop[data[All, 9], 1].Drop[data[All, 1], 1]
Out[ • ]=
       221
```

Analyses

Cadmium - significant parallelism

```
In[*]:= data[All, 3][[1]]
Out[ • ]=
       cd
 In[*]:= obsdata = Drop[data[All, 3], 1]
Out[ • ]=
       \{43, 2, 1, 0, 0, 0, 0, 0, 0, 0, 0, 0, 0\}
 In[*]:= nummut = obsdata.Drop[data[All, 1], 1]
Out[ • ]=
       50
       Of these, the number of multiply hit genes were:
 In[*]:= nummultiple = Total[Drop[data[All, 3], 2]]
Out[ • ]=
       3
       Randomizing the genes in which the mutations are observed:
 In[*]:= SeedRandom[129831]
```

Generating random draws of nummut mutations, each of which could occur in any one of 6607 possible genes (with an equal probability of mutating), repeating this 1000 times:

```
In[*]:= tab = Table[1/6607, {i, 1, 6607}];
     rantab = Table[BinCounts[
         RandomInteger[MultinomialDistribution[nummut, tab]], {0, 14, 1}], {i, 1, 1000}];
     The mean number of hits observed (j counts the number of mutations, accounting for genes hit 1, 2,
     ...13 times):
```

State hash: -4754988249305394038

Method: ExtendedCA

```
obsdata.Table[j (j), {j, 1, 13}]
 In[ • ]:=
        obsdata.Table[j, {j, 1, 13}]
Out[ • ]=
      1.2
```

RandomGeneratorState

Is higher than the mean numbers of all of 1000 randomizations:

```
meantab = Table \left[\frac{\text{rantab[i].Table[j}^2, \{j, 0, 13\}]}{\text{rantab[i].Table[j, <math>\{j, 0, 13\}]}}, \{i, 1, 1000\}\right] // N;
         Max[meantab]
Out[ • ]=
         1.12
        The 95% quantile for mean number of genes hit:
 In[a]:= {Quantile[meantab, 0.025], Quantile[meantab, 0.5], Quantile[meantab, 0.975]}
```

```
Out[ • ]=
        {1., 1., 1.04}
```

Thus the mean number of hits per mutated gene is significant (p<0.001).

Cobalt - significant parallelism

```
In[*]:= data[All, 4][1]
Out[ • ]=
       CO
 In[*]:= obsdata = Drop[data[All, 4], 1]
Out[ • ]=
        \{63, 5, 1, 1, 0, 1, 0, 0, 0, 0, 0, 0, 0, 0\}
 In[*]:= nummut = obsdata.Drop[data[All, 1], 1]
Out[ • ]=
       86
       Of these, the number of multiply hit genes were:
 In[*]:= nummultiple = Total[Drop[data[All, 4], 2]]
Out[ • ]=
       8
```

Randomizing the genes in which the mutations are observed:

```
In[*]:= SeedRandom[54219]
Out[ • ]=
                                  Method: ExtendedCA
       RandomGeneratorState
                                  State hash: 466 435 363 971759 394
```

Generating random draws of nummut mutations, each of which could occur in any one of 6607 possible genes (with an equal probability of mutating), repeating this 1000 times:

```
In[*]:= tab = Table[1/6607, {i, 1, 6607}];
     rantab = Table[BinCounts[
        RandomInteger[MultinomialDistribution[nummut, tab]], {0, 14, 1}], {i, 1, 1000}];
```

The mean number of hits observed (j counts the number of mutations, accounting for genes hit 1, 2, ...13 times):

Is higher than the mean numbers of all of 1000 randomizations:

$$In[*]:= meantab = Table \left[\frac{rantab[i].Table[j^2, \{j, 0, 13\}]}{rantab[i].Table[j, \{j, 0, 13\}]}, \{i, 1, 1000\} \right] // N;$$

Max[meantab]

Out[*]= 1.09302

The 95% quantile for mean number of genes hit:

```
In[@]:= {Quantile[meantab, 0.025], Quantile[meantab, 0.5], Quantile[meantab, 0.975]}
Out[@]:= {1., 1., 1.04651}
```

Thus the mean number of hits per mutated gene is significant (p<0.001).

Copper - no parallelism

Manganese - significant parallelism

For manganese, we drop the putative mutators (MnBM14 and MnBM42)

```
In[*]:= data[All, 6][1]
Out[ • ]=
       mn
 In[*]:= obsdata = Drop[data[All, 6], 1]
Out[ • ]=
       \{47, 2, 1, 1, 0, 0, 0, 0, 0, 0, 0, 0, 1\}
 In[*]:= nummut = obsdata.Drop[data[All, 1], 1]
Out[ • ]=
       71
       Of these, the number of multiply hit genes were:
```

```
In[*]:= nummultiple = Total[Drop[data[All, 6], 2]]
Out[ • ]=
       5
```

Randomizing the genes in which the mutations are observed:

```
In[*]:= SeedRandom[77 127]
Out[ • ]=
```

RandomGeneratorState Method: ExtendedCA State hash: 8 628 400 527 046 742 293

Generating random draws of nummut mutations, each of which could occur in any one of 6607 possible genes (with an equal probability of mutating), repeating this 1000 times:

```
In[*]:= tab = Table[1/6607, {i, 1, 6607}];
     rantab = Table[BinCounts[
        RandomInteger[MultinomialDistribution[nummut, tab]], {0, 14, 1}], {i, 1, 1000}];
```

The mean number of hits observed (j counts the number of mutations, accounting for genes hit 1, 2, ...13 times):

$$\frac{\textit{ln[o]:=}}{\textit{obsdata.Table[j (j), {j, 1, 13}]}} // N$$

$$\textit{Out[o]=}$$

3.50704

Is higher than the mean numbers of all of 1000 randomizations:

The 95% quantile for mean number of genes hit:

```
In[a]:= {Quantile[meantab, 0.025], Quantile[meantab, 0.5], Quantile[meantab, 0.975]}
Out[ • ]=
       {1., 1., 1.05634}
```

Even if we drop the gene hit 13 times in manganese (CDC25), the result is highly significant and outside the range of all 1000 randomizations:

Thus the mean number of hits per mutated gene is significant (p<0.001).

Nickle - significant parallelism

```
In[*]:= data[All, 7][1]
Out[ • ]=
       ni
 In[*]:= obsdata = Drop[data[All, 7], 1]
Out[ • ]=
        \{17, 2, 1, 0, 0, 0, 0, 0, 0, 0, 0, 0, 0, 0\}
 In[*]:= nummut = obsdata.Drop[data[All, 1], 1]
Out[ • ]=
       24
       Of these, the number of multiply hit genes were:
 In[@]:= nummultiple = Total[Drop[data[All, 7], 2]]
Out[ • ]=
       Randomizing the genes in which the mutations are observed:
 In[*]:= SeedRandom[32412]
Out[ • ]=
                                   Method: ExtendedCA
State hash: -8692798250509348668
       RandomGeneratorState
```

Generating random draws of nummut mutations, each of which could occur in any one of 6607 possible genes (with an equal probability of mutating), repeating this 1000 times:

```
In[*]:= tab = Table[1/6607, {i, 1, 6607}];
     rantab = Table[BinCounts[
        RandomInteger[MultinomialDistribution[nummut, tab]], {0, 14, 1}], {i, 1, 1000}];
```

The mean number of hits observed (j counts the number of mutations, accounting for genes hit 1, 2,

```
...13 times):
        \frac{\text{obsdata.Table[j (j), {j, 1, 13}]}}{\text{obsdata.Table[j, {j, 1, 13}]}} \; // \; N
Out[ • ]=
        1.41667
        Is higher than the mean numbers of all of 1000 randomizations:
       meantab = Table \left[ \frac{rantab[i].Table[j^2, \{j, 0, 13\}]}{rantab[i].Table[j, \{j, 0, 13\}]}, \{i, 1, 1000\} \right] // N;
        Max[meantab]
Out[ • ]=
        1.08333
        The 95% quantile for mean number of genes hit:
 In[*]:= {Quantile[meantab, 0.025], Quantile[meantab, 0.5], Quantile[meantab, 0.975]}
Out[ • ]=
        {1., 1., 1.08333}
        Thus the mean number of hits per mutated gene is significant (p<0.001).
        Zinc - significant parallelism
 In[*]:= data[All, 8][1]
Out[ • ]=
 In[*]:= obsdata = Drop[data[All, 8], 1]
Out[ • ]=
        {20, 2, 2, 0, 1, 0, 0, 0, 0, 0, 0, 0, 0}
 In[*]:= nummut = obsdata.Drop[data[All, 1], 1]
Out[ • ]=
        35
        Of these, the number of multiply hit genes were:
       nummultiple = Total[Drop[data[All, 8], 2]]
 In[ • ]:=
Out[ • ]=
        5
        Randomizing the genes in which the mutations are observed:
 In[*]:= SeedRandom[82712]
Out[ • ]=
        RandomGeneratorState Method: ExtendedCA State hash: -1951316623039499653
```

Generating random draws of nummut mutations, each of which could occur in any one of 6607 possible genes (with an equal probability of mutating), repeating this 1000 times:

```
In[*]:= tab = Table[1/6607, {i, 1, 6607}];
     rantab = Table[BinCounts[
        RandomInteger[MultinomialDistribution[nummut, tab]], {0, 14, 1}], {i, 1, 1000}];
```

The mean number of hits observed (j counts the number of mutations, accounting for genes hit 1, 2, ...13 times):

Is higher than the mean numbers of all of 1000 randomizations:

In[*]:= meantab = Table
$$\left[\frac{\text{rantab}[i].Table[j^2, {j, 0, 13}]}{\text{rantab}[i].Table[j, {j, 0, 13}]}, {i, 1, 1000}\right] // N;$$
Max[meantab]

Out[•]= 1.11429

The 95% quantile for mean number of genes hit:

```
In[a]:= {Quantile[meantab, 0.025], Quantile[meantab, 0.5], Quantile[meantab, 0.975]}
Out[ • ]=
       {1., 1., 1.05714}
```

Thus the mean number of hits per mutated gene is significant (p<0.001).

All metals together - significant parallelism

```
In[*]:= data[All, 2][[1]]
Out[ • ]=
       all
 In[*]:= obsdata = Drop[data[All, 2], 1]
Out[ • ]=
        \{186, 15, 0, 3, 3, 1, 0, 1, 0, 0, 0, 0, 1\}
 In[*]:= nummut = obsdata.Drop[data[All, 1], 1]
Out[ • ]=
       270
```

Of these, the number of multiply hit genes were:

```
In[*]:= nummultiple = Total[Drop[data[All, 2], 2]]
Out[ • ]=
       24
```

Randomizing the genes in which the mutations are observed:

```
In[*]:= SeedRandom[2120]
```

Out[•]=

RandomGeneratorState Method: ExtendedCA State hash: 1510 253 275 902 383 272

Generating random draws of nummut mutations, each of which could occur in any one of 6607 possible genes (with an equal probability of mutating), repeating this 1000 times:

```
In[*]:= tab = Table[1/6607, {i, 1, 6607}];
     rantab = Table[BinCounts[
        RandomInteger[MultinomialDistribution[nummut, tab]], {0, 14, 1}], {i, 1, 1000}];
```

The mean number of hits observed (j counts the number of mutations, accounting for genes hit 1, 2, ...13 times):

2.36296

Is higher than the mean numbers of all of 1000 randomizations:

In[*]:= meantab = Table
$$\left[\frac{\text{rantab[i].Table[j}^2, \{j, 0, 13\}]}{\text{rantab[i].Table[j, \{j, 0, 13\}]}}, \{i, 1, 1000\}\right] // N;$$

Max[meantab]

Out[•]= 1.11111

The 95% quantile for mean number of genes hit:

```
In[a]:= {Quantile[meantab, 0.025], Quantile[meantab, 0.5], Quantile[meantab, 0.975]}
Out[ • ]=
       {1.00741, 1.03704, 1.07407}
```

Even if we drop the gene hit 13 times in manganese (CDC25), the result is highly significant and outside the range of all 1000 randomizations:

1.8249

Out[•]=

Double hits: The 95% quantile for expected number of double hit genes is 1-10 (median of 5), whereas

15 were observed:

```
In[*]:= {Quantile[rantab[All, 3], 0.025],
       Quantile[rantab[All, 3], 0.5], Quantile[rantab[All, 3], 0.975]}
Out[ • ]=
       \{1, 5, 10\}
```

More than two hits: The 95% quantile for expected number of triple-plus hit genes is 0-1 (median of 0), whereas 9 were observed:

```
In[@]:= {Quantile[Sum[rantab[All, i], {i, 4, 10}], 0.025],
       Quantile[Sum[rantab[All, i], {i, 4, 10}], 0.5],
       Quantile[Sum[rantab[All, i], {i, 4, 10}], 0.975]}
Out[ • ]=
      {0,0,1}
```

Number of hits per gene and count of the maximum time that # of hits was observed:

```
In[@]:= Table[{i - 1, Max[rantab[All, i]]}, {i, 1, 10}]
Out[ • ]=
        \{\{0, 6351\}, \{1, 270\}, \{2, 12\}, \{3, 2\}, \{4, 1\}, \{5, 0\}, \{6, 0\}, \{7, 0\}, \{8, 0\}, \{9, 0\}\}\}
```

Only 6.7% of simulations had any genes hit more than twice, whereas 9 were observed:

```
In[@]:= Total[Sum[rantab[All, i]], {i, 4, 10}]] / 1000.
Out[ • ]=
       0.067
```

Thus both the mean number of hits per gene (the main test) is significant (p<0.001), as is the number of genes hit twice or more than twice.

All metals together (dropping repeated hits in the same gene) - significant parallelism

```
In[*]:= data[All, 9][1]
Out[ • ]=
       difmetal
 In[*]:= obsdata = Drop[data[All, 9], 1]
Out[ • ]=
       {200, 9, 1, 0, 0, 0, 0, 0, 0, 0, 0, 0, 0}
 In[*]:= nummut = obsdata.Drop[data[All, 1], 1]
Out[ • ]=
       221
```

Of these, the number of multiply hit genes were:

```
In[@]:= nummultiple = Total[Drop[data[All, 9], 2]]
Out[ • ]=
       10
```

Randomizing the genes in which the mutations are observed:

```
In[*]:= SeedRandom[8329]
```

Out[•]=

RandomGeneratorState Method: ExtendedCA
State hash: -4294397200794781781

Generating random draws of nummut mutations, each of which could occur in any one of 6607 possible genes (with an equal probability of mutating), repeating this 1000 times:

```
In[*]:= tab = Table[1/6607, {i, 1, 6607}];
     rantab = Table[BinCounts[
        RandomInteger[MultinomialDistribution[nummut, tab]], {0, 14, 1}], {i, 1, 1000}];
```

The mean number of hits observed (j counts the number of mutations, accounting for genes hit 1, 2, ...13 times):

1.1086

Is above the highest of all 1000 randomizations:

In[*]:= meantab = Table
$$\left[\frac{\text{rantab[i].Table[j}^2, \{j, 0, 13\}]}{\text{rantab[i].Table[j, \{j, 0, 13\}]}}, \{i, 1, 1000\}\right] // N;$$

Max[meantab]

Out[•]= 1.0905

The 95% quantile for mean number of genes hit:

```
In[a]:= {Quantile[meantab, 0.025], Quantile[meantab, 0.5], Quantile[meantab, 0.975]}
Out[ • ]=
       \{1., 1.02715, 1.0724\}
```

Double hits: The 95% quantile for expected number of double hit genes is 0-7 (median of 3) is less than the 9 observed:

```
In[*]:= {Quantile[rantab[All, 3], 0.025],
        Quantile[rantab[All, 3], 0.5], Quantile[rantab[All, 3], 0.975]}
Out[ • ]=
       \{0, 3, 7\}
```

More than two hits: The 95% quantile for expected number of triple-plus hit genes is 0-1 (median of

0), whereas 1 was observed:

```
In[@]:= {Quantile[Sum[rantab[All, i], {i, 4, 10}], 0.025],
       Quantile[Sum[rantab[All, i], {i, 4, 10}], 0.5],
       Quantile[Sum[rantab[All, i], {i, 4, 10}], 0.975]}
Out[ • ]=
      {0,0,1}
```

Number of hits per gene and count of the maximum time that # of hits was observed:

```
In[@]:= Table[{i - 1, Max[rantab[All, i]]]}, {i, 1, 10}]
Out[ • ]=
        \{\{0, 6396\}, \{1, 221\}, \{2, 10\}, \{3, 1\}, \{4, 0\}, \{5, 0\}, \{6, 0\}, \{7, 0\}, \{8, 0\}, \{9, 0\}\}\}
```

Only 3.3% of simulations had any genes hit more than twice, whereas one was observed:

```
In[@]:= Total[Sum[rantab[All, i], {i, 4, 10}]] / 1000.
Out[ • ]=
       0.033
```

Thus both the mean number of hits per gene (the main test) is significant (p<0.001), as is the number of genes hit more than twice (PMA1)