Association of Genomic Features with Integration

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Contents

1   Introduction

In this document, I examine the association of integration sites with various genomic features.

The data consist of both actual integration sites and sets of control sites, each set chosen to match the spacing (in bases) from the nearest restriction site (according to the direction in which the sequence was read) to an integration site. The numbers of insertion and matching sites for several data sets are shown below:

```
                    type
Origin.of.data.set    insertion match
  lenti-LN                  651  6490
  lenti-Sca1                639  6310
  lenti-marrow              130  1300
  lenti-pleural-fluid        55   550
  lenti-spleen              316  3160
  lenti-thymus              126  1260
  retro-LN                  334  3340
  retro-marrow              187  1870
  retro-spleen               91   910
  retro-thymus              146  1460
```

The advantage of choosing 'control' sites that match the spacing from the nearest restriction site is that biases due to location and density of restriction sites are eliminated by applying the classical multinomial logit model (reviewed in [2]). This model allows regression procedures to be applied to the study of integration intensity as a function of genomic features. The `clogit` function of the R `survival` library) implements estimation and fitting for such models along with the usual likelihood ratio and Wald tests.

The distribution of relative frequency of insertions across the chromosomes is given in this barplot:
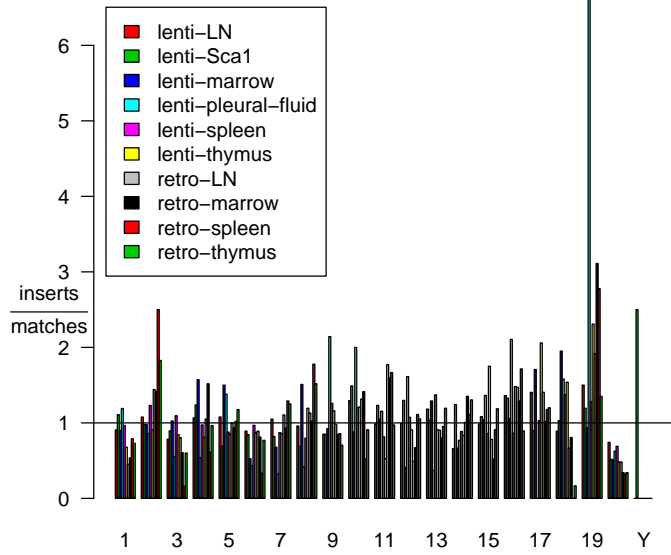
It seems evident that there are some chromosomes that are particularly favored for integration. This is reinforced by a test of statistical significance. The test performed used the likelihood ratio statistic for the multinomial logit model (reviewed in [2]) as implemented by the `clogit` function of the R `survival` library). The null hypothesis tested is that the ratio of true integration events to matched control sites is constant across all chromosomes. This test attains a p-value of $3.4105e - 13$.

2 Preference for Genes

2.1 refGene Genes

Here we examine the preference that integration events have for genes. In the following plot we show the relative frequency of integrations in genes according to the 'refGene' annotation. The bars grouped over the label "In Gene" give the relative frequency of integration events (compared to control sites) between bases located within refGene gene annotations, while the label "Not in Gene" give the relative frequency of integration events (compared to control sites) between bases not located within refGene gene annotations.
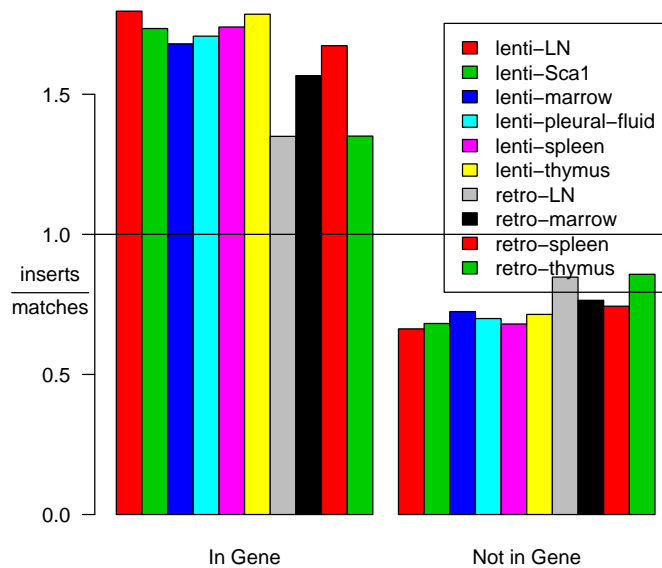
It seems evident that there is a strong tendency for insertions to occur in genes. A formal test of significance bears this out with a p-value of $< 2.22e - 16$. Also, it appears that the tendency of different viruses to integrate into genes varies, and a test for this hypothesis attains 0.0095238. Here is the table of coefficients of the log ratio of intensities for true insertion sites versus control insertion sites along with their standard errors, z statistics, and p-values for each data set:

```
                  coef     se     z         p
lenti-LN         1.000 0.0832 12.00 3.07e-33
lenti-Sca1       0.949 0.0845 11.20 3.12e-29
lenti-marrow     0.844 0.1860  4.53 5.82e-06
```

```
lenti-pleural-fluid 0.868 0.2820  3.07 2.12e-03
lenti-spleen        0.933 0.1190  7.84 4.44e-15
lenti-thymus        0.916 0.1890  4.84 1.31e-06
retro-LN            0.466 0.1180  3.96 7.37e-05
retro-marrow        0.716 0.1550  4.61 4.02e-06
retro-spleen        0.817 0.2240  3.65 2.64e-04
retro-thymus        0.452 0.1790  2.53 1.14e-02
```

As is evident, there are some differences in the coefficients. The largest coefficient is seen in the lenti-LN data set, while the smallest is seen in the retro-thymus data set.

In the following plot we show the relative frequency of insertions in exons according to the 'refGene' annotation. The bars grouped over the label "In Exon" give the relative frequency of integration events (compared to control sites) between bases located in exons according to the Acembly annotation, while the label "Not in Exon" give the relative frequency of integration events (compared to control sites) between bases not located in exons according to the Acembly gene annotation.
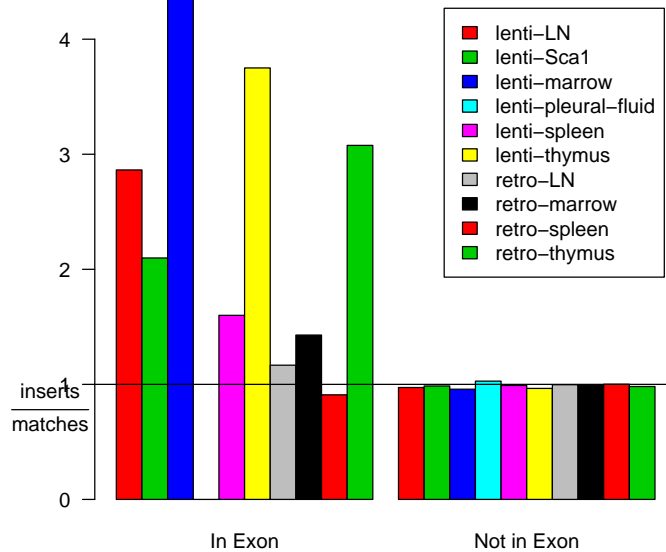

Here is the table of coefficients of the log ratio of intensities for true insertion sites versus control insertion sites along with their standard errors, z statistics, and p-values for each data set:

```
              coef       se       z       p
```

```
lenti-LN               0.5200    0.227   2.2900  0.022
lenti-Sca1             0.1830    0.273   0.6720  0.502
lenti-marrow           1.0200    0.473   2.1600  0.031
lenti-pleural-fluid  -13.9000  627.000  -0.0221  0.982
lenti-spleen          -0.1130    0.393  -0.2880  0.774
lenti-thymus           0.8140    0.496   1.6400  0.101
retro-LN              -0.1560    0.414  -0.3770  0.706
retro-marrow          -0.0938    0.544  -0.1720  0.863
retro-spleen          -0.7020    1.060  -0.6650  0.506
retro-thymus           0.8630    0.592   1.4600  0.145
```

The model on which these coefficients are based include terms for whether the site is in a gene or not. Thus, the effect shown as "In Exon" is net of that due to being in a gene. Note that in the barplot above the 'Not in Exon' bars include both the introns and intergenic regions, so the impression given by the table may differ from that for the barplot.

2.2 ensGenes

Here we examine the preference that insertions have for genes. In the following plot we show the relative frequency of insertions in genes according to the 'ensGene' annotation.
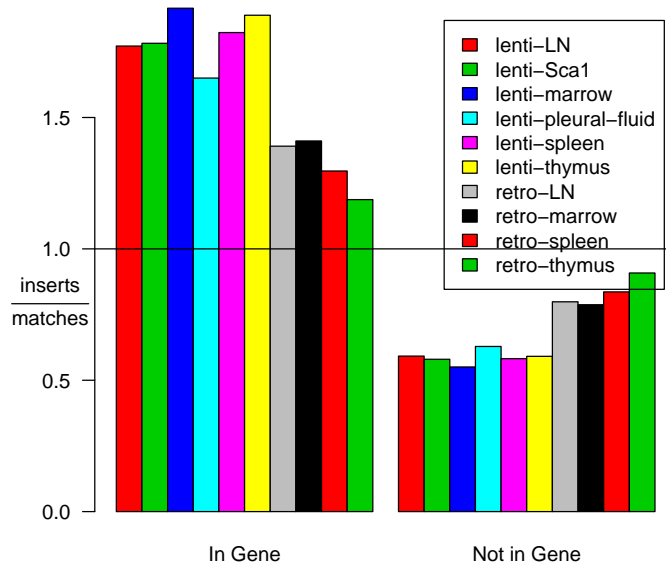
It seems evident that there is a strong tendency for insertions to occur in

genes. A formal test of significance bears this out with a p-value of $< 2.22e - 16$. Also, it appears that the tendency of different viruses to integrate into genes varies, and a test for this hypothesis attains $2.91e - 08$. Here is the table of coefficients of the log ratio of intensities for true insertion sites versus control insertion sites along with their standard errors, z statistics, and p-values for each data set:

```
                    coef     se     z        p
lenti-LN            1.100 0.0849 13.00 1.37e-38
lenti-Sca1          1.150 0.0871 13.20 1.06e-39
lenti-marrow        1.290 0.1950  6.60 4.01e-11
lenti-pleural-fluid 0.945 0.2870  3.29 1.01e-03
lenti-spleen        1.140 0.1220  9.35 8.58e-21
lenti-thymus        1.170 0.1930  6.07 1.30e-09
retro-LN            0.558 0.1160  4.82 1.46e-06
retro-marrow        0.576 0.1530  3.76 1.73e-04
retro-spleen        0.441 0.2220  1.99 4.71e-02
retro-thymus        0.268 0.1780  1.50 1.34e-01
```

As is evident, there are some differences in the coefficients. The largest coefficient is seen in the lenti-marrow data set, while the smallest is seen in the retro-thymus data set.

In the following plot we show the relative frequency of insertions in exons according to the 'ensGene' annotation.
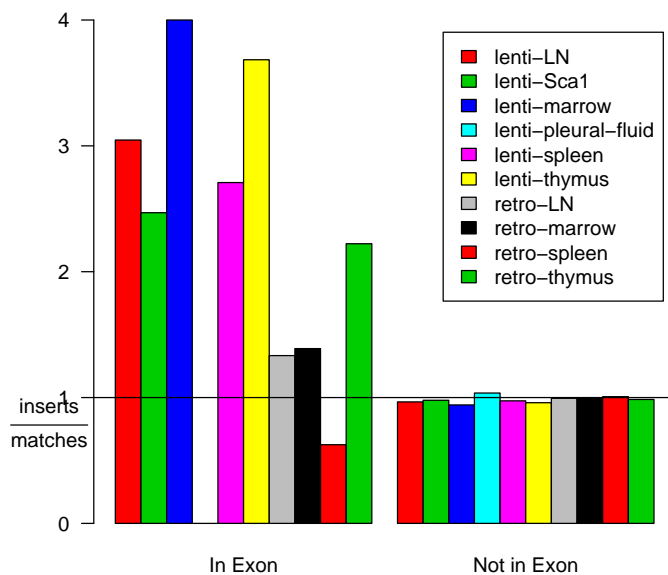
Here is the table of coefficients of the log ratio of intensities for true insertion sites versus control insertion sites along with their standard errors, z statistics, and p-values for each data set:

```
                       coef      se       z        p
lenti-LN             0.6180   0.208   2.970  0.00294
lenti-Sca1           0.3470   0.239   1.450  0.14700
lenti-marrow         0.7420   0.388   1.910  0.05570
lenti-pleural-fluid -13.9000 554.000 -0.025  0.98000
lenti-spleen         0.4050   0.324   1.250  0.21100
lenti-thymus         0.7260   0.460   1.580  0.11500
retro-LN            -0.0439   0.389  -0.113  0.91000
retro-marrow        -0.0170   0.487  -0.035  0.97200
retro-spleen        -0.7930   1.050  -0.758  0.44900
retro-thymus         0.6790   0.576   1.180  0.23800
```

The model on which these coefficients are based include terms for whether the site is in a gene or not. Thus, the effect shown as "In Exon" is net of that due to being in a gene.

2.3   genScan Genes

Here we examine the preference that insertions have for genes. In the following plot we show the relative frequency of insertions in genes according to the

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'genScan' annotation.

It seems evident that there is a strong tendency for insertions to occur in genes. A formal test of significance bears this out with a p-value of $3.5702e-10$. Also, it appears that the tendency of different viruses to integrate into genes varies, and a test for this hypothesis attains 0.00037706. Here is the table of coefficients of the log ratio of intensities for true insertion sites versus control insertion sites along with their standard errors, z statistics, and p-values for each data set:
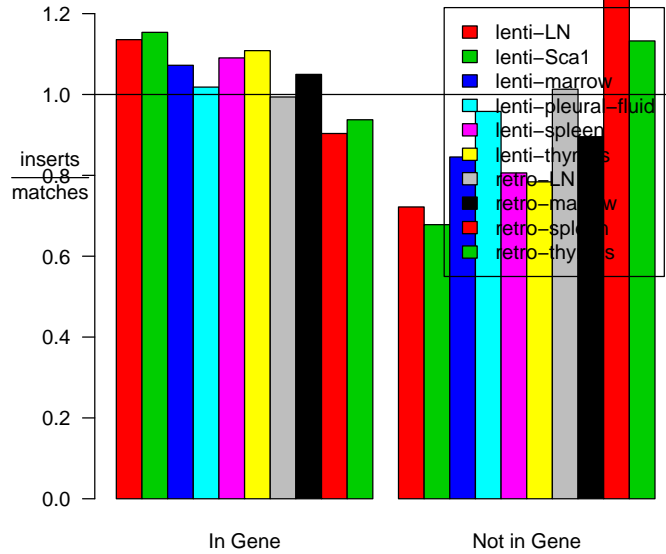
```
                     coef     se      z        p
lenti-LN            0.4660 0.0964  4.840 1.32e-06
lenti-Sca1          0.5370 0.0997  5.380 7.42e-08
lenti-marrow        0.2390 0.2070  1.160 2.48e-01
lenti-pleural-fluid 0.0606 0.3100  0.195 8.45e-01
lenti-spleen        0.3030 0.1350  2.250 2.43e-02
lenti-thymus        0.3430 0.2100  1.630 1.03e-01
retro-LN           -0.0193 0.1230 -0.157 8.75e-01
retro-marrow        0.1620 0.1700  0.950 3.42e-01
retro-spleen       -0.3130 0.2280 -1.370 1.71e-01
retro-thymus       -0.1880 0.1810 -1.040 2.98e-01
```

As is evident, there are some differences in the coefficients. The largest coefficient is seen in the lenti-Sca1 data set, while the smallest is seen in the

retro-spleen data set.

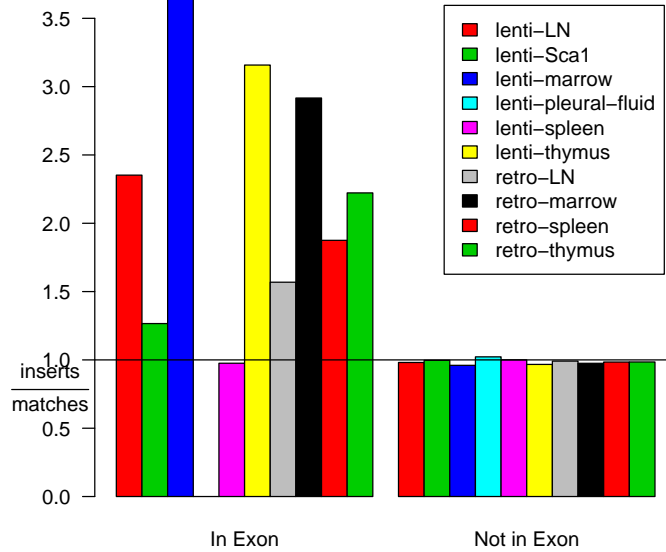In the following plot we show the relative frequency of insertions in exons according to the 'genScan' annotation.

Here is the table of coefficients of the log ratio of intensities for true insertion sites versus control insertion sites along with their standard errors, z statistics, and p-values for each data set:

```
                     coef      se       z       p
lenti-LN            0.741   0.247  3.0000 0.00271
lenti-Sca1          0.113   0.338  0.3330 0.73900
lenti-marrow        1.300   0.457  2.8400 0.00456
lenti-pleural-fluid -13.200 655.000 -0.0201 0.98400
lenti-spleen       -0.122   0.529 -0.2300 0.81800
lenti-thymus        1.100   0.481  2.2800 0.02270
retro-LN            0.483   0.393  1.2300 0.21900
retro-marrow        1.040   0.436  2.3800 0.01730
retro-spleen        0.744   0.635  1.1700 0.24100
retro-thymus        0.894   0.567  1.5800 0.11500
```

The model on which these coefficients are based include terms for whether the site is in a gene or not. Thus, the effect shown as "In Exon" is net of that due to being in a gene.

2.4 oncogenes

Here we examine the preference that insertions have for oncogenes. In the following plot we show the relative frequency of insertions with 50kb of an oncogene 5' end.

A formal test of oncogenic insertion returns p-value of $< 2.22e - 16$. The tendency of different viruses to integrate near oncogenes may vary, and a test for thi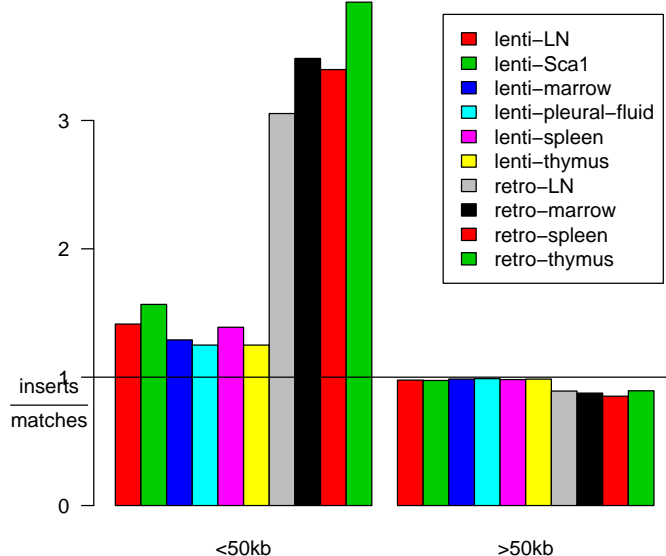s hypothesis attains $2.0046e - 06$. Here is the table of coefficients of the log ratio of intensities for true insertion sites versus control insertion sites along with their standard errors, z statistics, and p-values for each data set:

	coef	se	z	p
lenti-LN	-0.367	0.159	-2.310	2.10e-02
lenti-Sca1	-0.473	0.169	-2.800	5.09e-03
lenti-marrow	-0.267	0.386	-0.693	4.88e-01
lenti-pleural-fluid	-0.233	0.627	-0.371	7.10e-01
lenti-spleen	-0.346	0.246	-1.410	1.59e-01
lenti-thymus	-0.239	0.367	-0.651	5.15e-01
retro-LN	-1.250	0.175	-7.180	7.18e-13
retro-marrow	-1.420	0.231	-6.150	7.55e-10
retro-spleen	-1.440	0.310	-4.650	3.27e-06
retro-thymus	-1.480	0.281	-5.260	1.45e-07
lenti-LN	NA	0.000	NA	NA

```
lenti-Sca1            NA 0.000    NA      NA
lenti-marrow          NA 0.000    NA      NA
lenti-pleural-fluid   NA 0.000    NA      NA
lenti-spleen          NA 0.000    NA      NA
lenti-thymus          NA 0.000    NA      NA
retro-LN              NA 0.000    NA      NA
retro-marrow          NA 0.000    NA      NA
retro-spleen          NA 0.000    NA      NA
retro-thymus          NA 0.000    NA      NA
```

As is evident, there are some differences in the coefficients. The largest coefficient is seen in the lenti-pleural-fluid data set, while the smallest is seen in the retro-thymus data set.

3 CpG Island Neighborhoods

Here we study the effect of being in the neighborhood of CpG Islands. Following Wu et al [3], who found that the neighborhoods within ±1kb of CpG islands are enriched for MLV insertions, we study such neighborhoods.

3.1 1 kilobase neighborhoods

The following plot shows the effect of being in or within ±1kb of a CpG island:
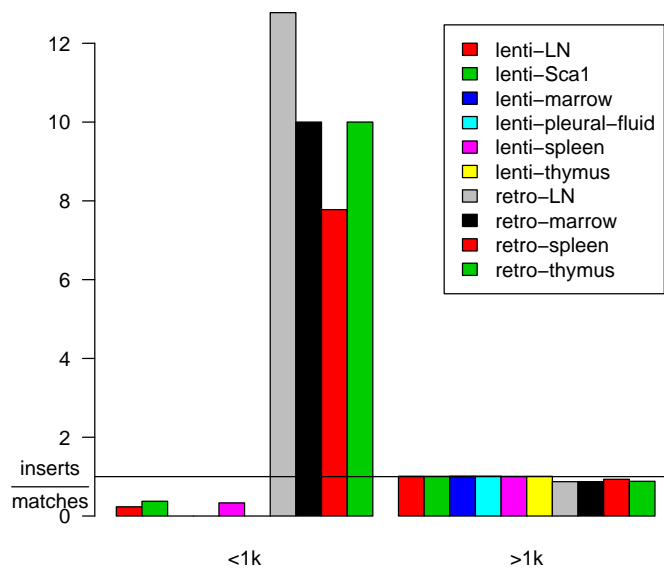
A formal test of significance comparing the difference attains a p-value of $< 2.22e - 16$. A test for differences between viruses attains $< 2.22e - 16$. Here is the table of coefficients of the log ratio of intensities for true insertion sites versus control insertion sites along with their standard errors, z statistics, and p-values for each data set:

```
                          coef       se        z         p
lenti-LN                 -1.46    0.716 -2.03000 4.21e-02
lenti-Sca1               -0.98    0.590 -1.66000 9.71e-02
lenti-marrow            -14.70 1130.000 -0.01300 9.90e-01
lenti-pleural-fluid     -14.60 1810.000 -0.00810 9.94e-01
lenti-spleen             -1.10    1.020 -1.08000 2.80e-01
lenti-thymus            -14.70 1520.000 -0.00965 9.92e-01
retro-LN                  2.67    0.236 11.30000 8.10e-30
```

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```
retro-marrow            2.50    0.305  8.22000 2.11e-16
retro-spleen            2.12    0.520  4.07000 4.69e-05
retro-thymus            2.37    0.335  7.08000 1.45e-12
```

The largest coefficient is seen in the retro-LN data set, while the smallest is seen in the lenti-thymus data set.

3.2   5 kilobase neighborhoods

The following plot shows the effect of being in or within ±5kb of a CpG island:
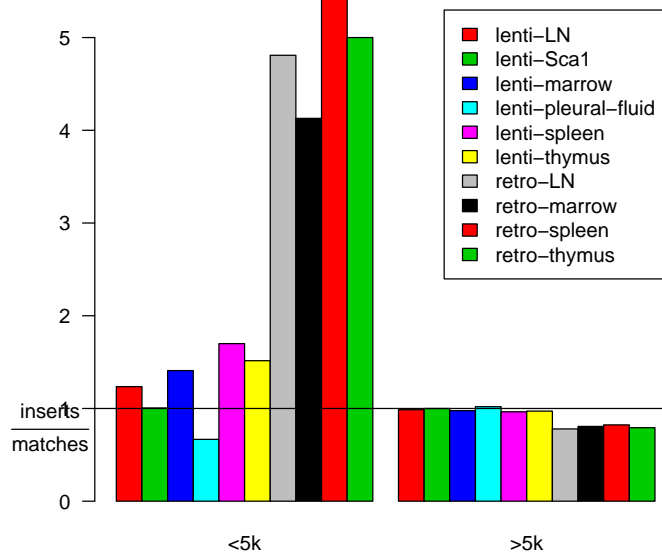
A formal test of significance comparing the difference attains a p-value of $< 2.22e - 16$. A test for differences between viruses attains $< 2.22e - 16$. Here is the table of coefficients of the log ratio of intensities for true insertion sites versus control insertion sites along with their standard errors, z statistics, and p-values for each data set:

```
                          coef    se      z        p
lenti-LN                0.2300 0.166   1.390 1.66e-01
lenti-Sca1              0.0188 0.185   0.102 9.19e-01
lenti-marrow            0.3610 0.348   1.040 3.00e-01
lenti-pleural-fluid -0.4280 0.747  -0.573 5.67e-01
lenti-spleen            0.5690 0.221   2.570 1.02e-02
lenti-thymus            0.4540 0.357   1.270 2.04e-01
```

```
retro-LN                    1.8300 0.148 12.400 4.39e-35
retro-marrow                1.6600 0.203  8.190 2.67e-16
retro-spleen                1.8500 0.309  5.990 2.14e-09
retro-thymus                1.8500 0.231  8.020 1.09e-15
```

The largest coefficient is seen in the retro-thymus data set, while the smallest is seen in the lenti-pleural-fluid data set.

3.3 10 kilobase neighborhoods

The following plot shows the effect of being in or within ± 10kb of a CpG island:
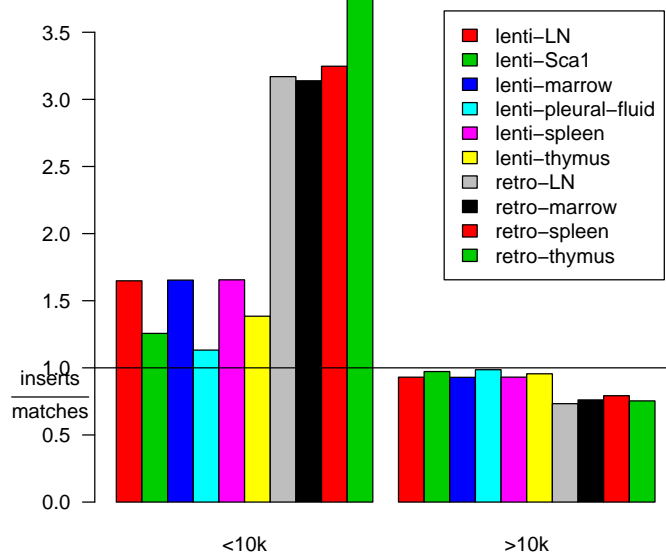
A formal test of significance comparing the difference attains a p-value of $< 2.22e - 16$. A test for differences between viruses attains $1.6562e - 15$. Here is the table of coefficients of the log ratio of intensities for true insertion sites versus control insertion sites along with their standard errors, z statistics, and p-values for each data set:

```
                        coef    se      z       p
lenti-LN                0.564 0.115   4.890 1.01e-06
lenti-Sca1              0.253 0.128   1.980 4.82e-02
lenti-marrow            0.569 0.254   2.240 2.51e-02
lenti-pleural-fluid 0.138 0.455   0.303 7.62e-01
lenti-spleen            0.582 0.167   3.490 4.81e-04
```

```
lenti-thymus          0.383 0.276  1.390 1.65e-01
retro-LN              1.470 0.129 11.400 2.74e-30
retro-marrow          1.450 0.180  8.060 7.86e-16
retro-spleen          1.390 0.262  5.310 1.12e-07
retro-thymus          1.600 0.205  7.810 5.70e-15
```

The largest coefficient is seen in the retro-thymus data set, while the smallest is seen in the lenti-pleural-fluid data set.

3.4   25 kilobase neighborhoods

The following plot shows the effect of being in or within ±25kb of a CpG island:
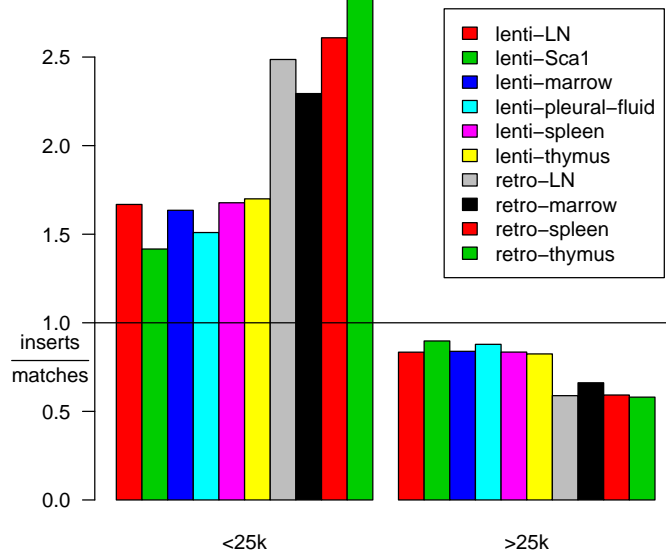
A formal test of significance comparing the difference attains a p-value of $< 2.22e - 16$. A test for differences between viruses attains $7.8411e - 14$. Here is the table of coefficients of the log ratio of intensities for true insertion sites versus control insertion sites along with their standard errors, z statistics, and p-values for each data set:

```
                   coef      se    z        p
lenti-LN          0.693 0.0893  7.76 8.31e-15
lenti-Sca1        0.462 0.0935  4.94 7.93e-07
lenti-marrow      0.657 0.1970  3.33 8.80e-04
lenti-pleural-fluid 0.550 0.3180 1.73 8.41e-02
```

```
lenti-spleen          0.692 0.1270  5.44 5.46e-08
lenti-thymus          0.749 0.2050  3.66 2.52e-04
retro-LN              1.440 0.1180 12.20 4.08e-34
retro-marrow          1.270 0.1600  7.90 2.78e-15
retro-spleen          1.510 0.2310  6.54 6.26e-11
retro-thymus          1.580 0.1790  8.81 1.20e-18
```

The largest coefficient is seen in the retro-thymus data set, while the smallest is seen in the lenti-Sca1 data set.

3.5   50 kilobase neighborhoods

The following plot shows the effect of being in or within ± 50kb of a CpG island:
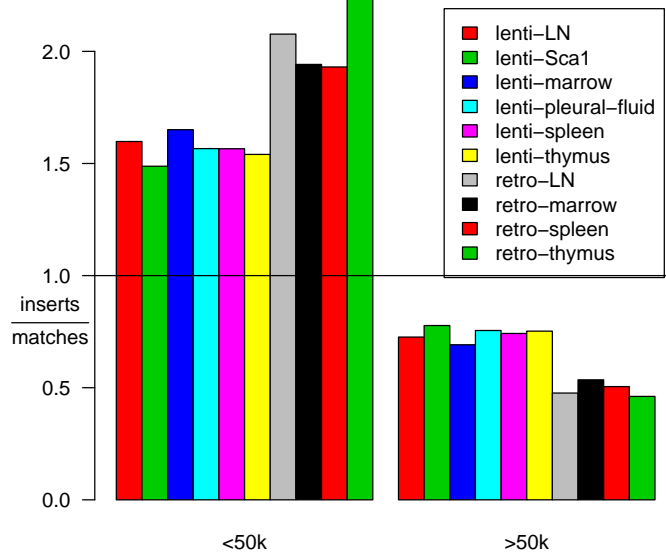
A formal test of significance comparing the difference attains a p-value of $< 2.22e - 16$. A test for differences between viruses attains $1.8251e - 09$. Here is the table of coefficients of the log ratio of intensities for true insertion sites versus control insertion sites along with their standard errors, z statistics, and p-values for each data set:

```
                      coef      se    z         p
lenti-LN              0.796 0.0834  9.54 1.40e-21
lenti-Sca1            0.662 0.0841  7.88 3.35e-15
lenti-marrow          0.857 0.1840  4.65 3.29e-06
```

```
lenti-pleural-fluid 0.760 0.2910  2.61 9.01e-03
lenti-spleen        0.746 0.1190  6.27 3.67e-10
lenti-thymus        0.734 0.1910  3.85 1.19e-04
retro-LN            1.470 0.1230 11.90 7.48e-33
retro-marrow        1.290 0.1610  8.01 1.18e-15
retro-spleen        1.360 0.2380  5.72 1.05e-08
retro-thymus        1.620 0.1910  8.51 1.77e-17
```

The largest coefficient is seen in the retro-thymus data set, while the smallest is seen in the lenti-Sca1 data set.

4 Gene Density, Expression 'Density', and CpG Island Density

In this section the association with gene density is examined. For expression analysis, the 'genes' that are counted are the genes represented on the microarray. In addition, we the number of such genes expressed at various levels. The levels are

low.ex Count genes whose expression is in the upper half and divide by number of bases

med.ex Count genes whose expression is in the upper $1/8^{th}$ and divide by number of bases

high.ex Count genes whose expression is in the upper $1/16^{th}$ and divide by number of bases

The bolded terms are used as abbreviations in what follows. The abbreviation **dens** is used to indicate gene density as number of genes per base.

4.1 25 kilobase Window

In the barplot that follows we examine the association of insertion sites with gene density in a 25 kilobase window surrounding each locus. More such plots will follow and the method of their construction is always to try to divide the data according to the deciles of density. However, it often happens that there is a very skewed distribution of density and even the 90th percentile is zero. In that case, the barplots simply show the sites for which the density is zero and those for which it is non-zero. If there are fewer than ten groups of bars, the groupings contain ten percent of the sites each except for the leftmost grouping which will contain all of the remaining sites.

Also note that the title of the plot contains clues as to its content; the prefix indicates the type of variable studied while the suffix indicates the window width in the number of bases. The p-value given is the result of fitting a cubic polynomial to the gene density values.

The following expression data and probe set were used for this report:

```
[1] "MEF-GSE3400-MGU74Av2"

[1] "MG_U74"

Density data too sparse for barplot
```

	coef	se	z	p
lenti-LN	0.702	0.119	5.900	3.74e-09
lenti-Sca1	0.643	0.123	5.240	1.58e-07
lenti-marrow	0.831	0.264	3.150	1.62e-03

```
lenti-pleural-fluid 0.117 0.498 0.235 8.14e-01
lenti-spleen        0.811 0.162 5.010 5.39e-07
lenti-thymus        0.733 0.278 2.630 8.44e-03
retro-LN            1.220 0.154 7.900 2.72e-15
retro-marrow        0.857 0.212 4.050 5.20e-05
retro-spleen        1.010 0.298 3.380 7.15e-04
retro-thymus        1.220 0.234 5.210 1.88e-07
```

Here are the results for expression density. First, we count just genes that are in the upper half.

```
Density data too sparse for barplot

                    coef    se     z        p
lenti-LN           0.965 0.141 6.870 6.62e-12
lenti-Sca1         0.757 0.153 4.940 7.80e-07
lenti-marrow       0.741 0.345 2.150 3.17e-02
lenti-pleural-fluid 0.233 0.627 0.371 7.10e-01
lenti-spleen       0.915 0.197 4.650 3.34e-06
lenti-thymus       0.882 0.336 2.630 8.61e-03
retro-LN           1.260 0.182 6.910 4.69e-12
retro-marrow       0.705 0.283 2.490 1.27e-02
retro-spleen       0.888 0.376 2.360 1.81e-02
retro-thymus       0.866 0.312 2.780 5.47e-03
```

Now we count genes in the upper $1/8^{th}$:

```
Density data too sparse for barplot
```

```
                   coef    se    z         p
lenti-LN           1.010 0.186 5.45 4.99e-08
lenti-Sca1         0.533 0.216 2.47 1.36e-02
lenti-marrow       0.539 0.456 1.18 2.37e-01
lenti-pleural-fluid 0.646 0.641 1.01 3.13e-01
lenti-spleen       0.589 0.287 2.06 3.98e-02
lenti-thymus       0.739 0.453 1.63 1.03e-01
retro-LN           1.160 0.241 4.83 1.38e-06
retro-marrow       0.707 0.370 1.91 5.59e-02
retro-spleen       0.823 0.474 1.74 8.24e-02
retro-thymus       1.120 0.384 2.93 3.43e-03
```

And here we count genes in the upper $1/16^{th}$:

```
Density data too sparse for barplot

                     coef    se     z        p
lenti-LN            0.982 0.238 4.130 3.65e-05
lenti-Sca1          0.558 0.293 1.900 5.69e-02
lenti-marrow        0.108 0.754 0.143 8.86e-01
lenti-pleural-fluid 3.000 1.220 2.450 1.44e-02
lenti-spleen        0.637 0.366 1.740 8.20e-02
lenti-thymus        0.856 0.556 1.540 1.24e-01
retro-LN            1.110 0.322 3.460 5.46e-04
retro-marrow        1.320 0.447 2.960 3.10e-03
retro-spleen        0.788 0.651 1.210 2.26e-01
retro-thymus        1.690 0.425 3.960 7.37e-05
```

Here the effect of density of CpG islands is studied:

rescale(cpg.dens.25k)
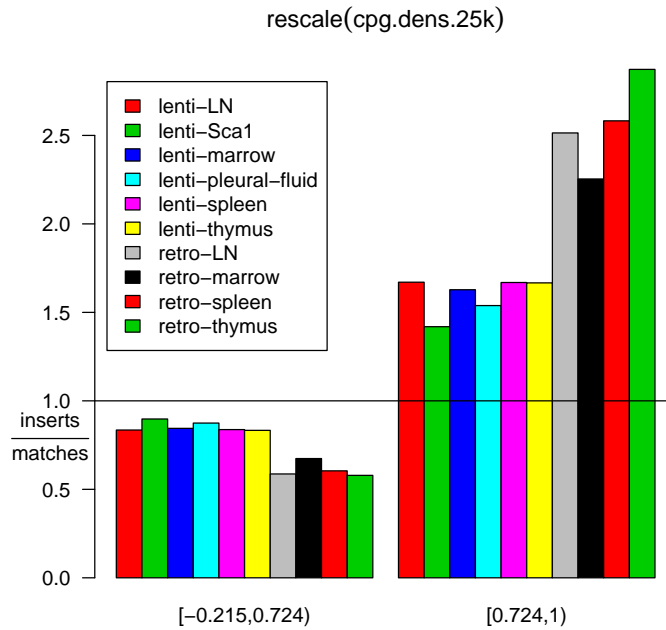
	coef	se	z	p
lenti-LN	0.695	0.0894	7.78	7.20e-15
lenti-Sca1	0.453	0.0938	4.83	1.37e-06
lenti-marrow	0.648	0.1990	3.26	1.13e-03
lenti-pleural-fluid	0.575	0.3190	1.80	7.13e-02
lenti-spleen	0.682	0.1280	5.34	9.16e-08
lenti-thymus	0.721	0.2060	3.49	4.76e-04
retro-LN	1.460	0.1180	12.30	9.72e-35
retro-marrow	1.230	0.1600	7.67	1.77e-14
retro-spleen	1.480	0.2300	6.42	1.38e-10
retro-thymus	1.600	0.1800	8.89	6.28e-19

4.2   50 kilobase Window

In the barplot that follows we examine the association of insertion sites with expression density in a 50 kilobase window surrounding each locus. First, we count just the number of genes represented on the chip.

rescale(dens.50k)
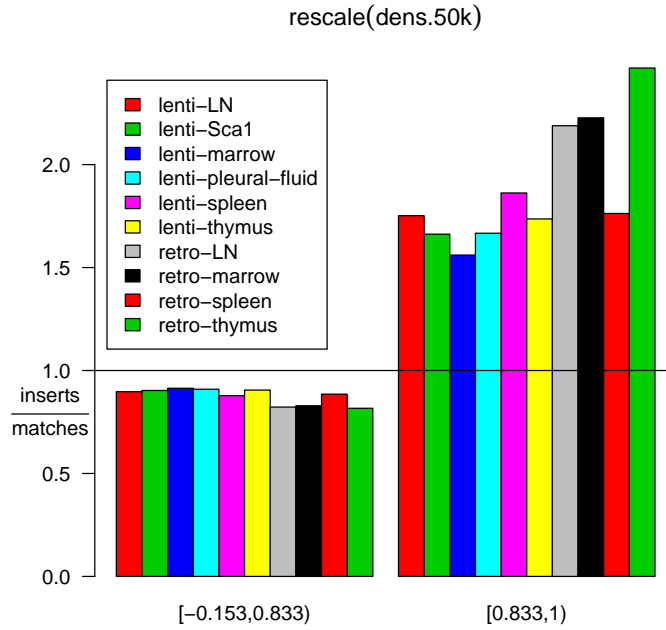
	coef	se	z	p
lenti-LN	0.703	0.0975	7.21	5.74e-13
lenti-Sca1	0.609	0.1000	6.09	1.15e-09
lenti-marrow	0.690	0.2150	3.22	1.30e-03
lenti-pleural-fluid	0.480	0.3620	1.33	1.84e-01
lenti-spleen	0.803	0.1360	5.88	4.03e-09
lenti-thymus	0.641	0.2290	2.80	5.14e-03
retro-LN	0.998	0.1290	7.72	1.14e-14
retro-marrow	1.010	0.1720	5.85	4.83e-09
retro-spleen	0.703	0.2540	2.77	5.60e-03
retro-thymus	1.100	0.2000	5.49	4.13e-08

Here are the results for expression density. First, we count just genes that are in the upper half.

```
Density data too sparse for barplot
```

```
                   coef    se    z        p
lenti-LN           0.880 0.111 7.92 2.39e-15
lenti-Sca1         0.704 0.118 5.99 2.06e-09
lenti-marrow       0.723 0.252 2.87 4.11e-03
lenti-pleural-fluid 0.663 0.410 1.62 1.06e-01
lenti-spleen       0.741 0.162 4.58 4.70e-06
lenti-thymus       0.715 0.276 2.59 9.55e-03
retro-LN           1.170 0.145 8.08 6.28e-16
retro-marrow       0.962 0.204 4.71 2.47e-06
retro-spleen       0.579 0.305 1.90 5.76e-02
retro-thymus       1.230 0.224 5.49 3.97e-08
```

Now we count genes in the upper $1/8^{th}$:

```
Density data too sparse for barplot
```

```
                       coef    se     z        p
lenti-LN              0.821 0.145 5.660 1.52e-08
lenti-Sca1            0.491 0.159 3.090 2.02e-03
lenti-marrow         0.384 0.337 1.140 2.54e-01
lenti-pleural-fluid  0.831 0.477 1.740 8.16e-02
lenti-spleen         0.619 0.214 2.890 3.83e-03
lenti-thymus         0.234 0.411 0.568 5.70e-01
retro-LN             1.220 0.176 6.910 4.99e-12
retro-marrow         1.050 0.249 4.210 2.58e-05
retro-spleen         0.748 0.353 2.120 3.41e-02
retro-thymus         1.270 0.273 4.650 3.32e-06
```

And here we count genes in the upper $1/16^{th}$:

```
Density data too sparse for barplot

                        coef    se     z        p
lenti-LN               0.890 0.185 4.810 1.48e-06
lenti-Sca1             0.235 0.235 1.000 3.17e-01
lenti-marrow           0.302 0.447 0.677 4.99e-01
lenti-pleural-fluid    1.320 0.677 1.950 5.09e-02
lenti-spleen           0.691 0.279 2.470 1.33e-02
lenti-thymus           0.935 0.463 2.020 4.34e-02
retro-LN               1.160 0.227 5.110 3.19e-07
retro-marrow           1.500 0.296 5.080 3.71e-07
retro-spleen           1.030 0.438 2.350 1.88e-02
retro-thymus           1.360 0.326 4.160 3.23e-05
```

Here the effect of density of CpG islands is studied:

rescale(cpg.dens.50k)
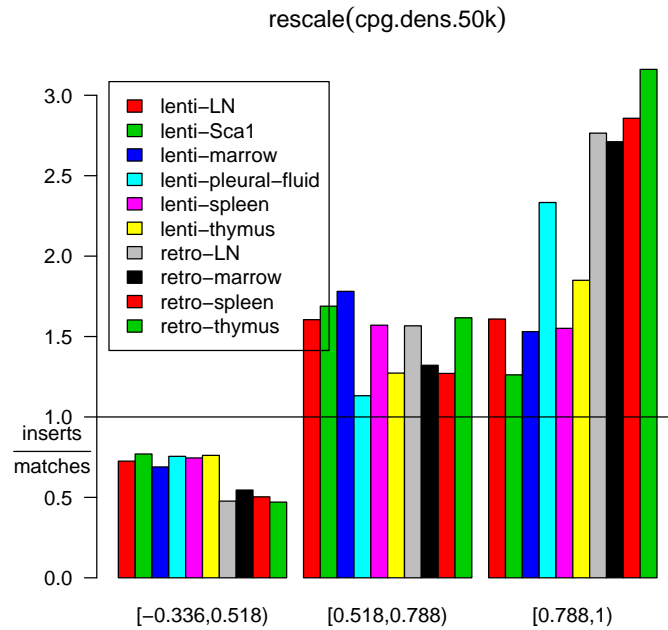
	coef	se	z	p
lenti-LN	0.802	0.0834	9.61	7.33e-22
lenti-Sca1	0.667	0.0840	7.93	2.16e-15
lenti-marrow	0.867	0.1840	4.71	2.52e-06
lenti-pleural-fluid	0.760	0.2910	2.61	9.01e-03
lenti-spleen	0.738	0.1190	6.21	5.40e-10
lenti-thymus	0.714	0.1910	3.74	1.87e-04
retro-LN	1.470	0.1230	12.00	4.53e-33
retro-marrow	1.270	0.1600	7.90	2.72e-15
retro-spleen	1.370	0.2380	5.76	8.27e-09
retro-thymus	1.590	0.1900	8.39	4.78e-17

4.3   100 kilobase Window

In the barplot that follows we examine the association of insertion sites with expression density in a 100 kilobase window surrounding each locus. First, we count just the number of genes represented on the chip.

rescale(dens.100k)
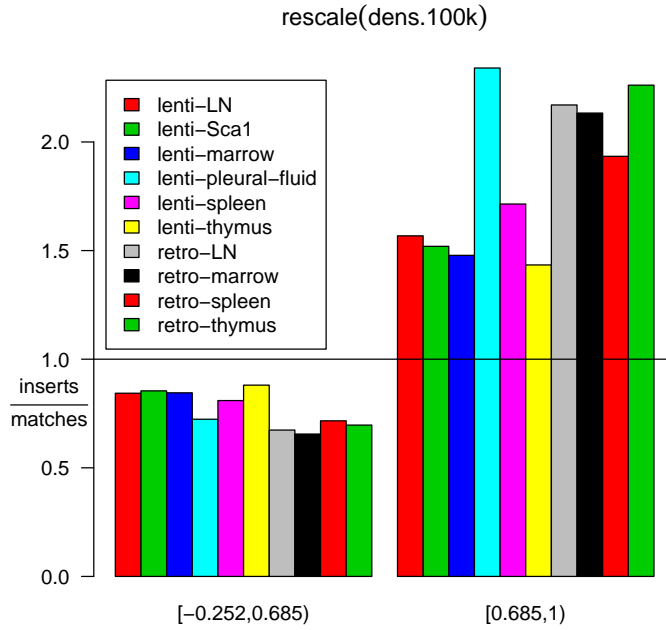
```
                     coef     se     z        p
lenti-LN            0.661 0.0861  7.67 1.73e-14
lenti-Sca1          0.562 0.0883  6.36 2.01e-10
lenti-marrow        0.564 0.1930  2.92 3.48e-03
lenti-pleural-fluid 1.100 0.2930  3.76 1.67e-04
lenti-spleen        0.744 0.1240  6.00 1.92e-09
lenti-thymus        0.510 0.1990  2.57 1.02e-02
retro-LN            1.220 0.1180 10.30 8.21e-25
retro-marrow        1.280 0.1580  8.10 5.62e-16
retro-spleen        1.190 0.2220  5.36 8.14e-08
retro-thymus        1.140 0.1780  6.40 1.58e-10
```

Here are the results for expression density. First, we count just genes that are in the upper half.

rescale(low.ex.100k)
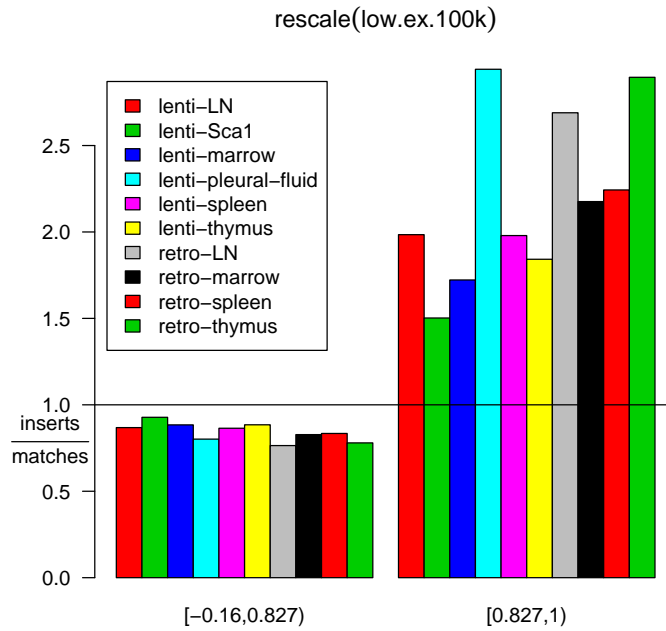
	coef	se	z	p
lenti-LN	0.880	0.0931	9.45	3.40e-21
lenti-Sca1	0.581	0.1000	5.81	6.32e-09
lenti-marrow	0.752	0.2060	3.64	2.70e-04
lenti-pleural-fluid	1.060	0.3210	3.30	9.55e-04
lenti-spleen	0.819	0.1360	6.03	1.64e-09
lenti-thymus	0.686	0.2170	3.17	1.53e-03
retro-LN	1.300	0.1240	10.50	1.03e-25
retro-marrow	1.080	0.1700	6.36	2.00e-10
retro-spleen	1.140	0.2380	4.78	1.77e-06
retro-thymus	1.280	0.1920	6.63	3.29e-11

Now we count genes in the upper $1/8^{th}$:

```
Density data too sparse for barplot

                    coef    se    z       p
lenti-LN           0.792 0.115 6.90 5.31e-12
lenti-Sca1         0.283 0.131 2.16 3.11e-02
lenti-marrow       0.399 0.258 1.55 1.22e-01
lenti-pleural-fluid 1.260 0.371 3.41 6.59e-04
lenti-spleen       0.670 0.168 3.99 6.59e-05
lenti-thymus       0.508 0.282 1.80 7.20e-02
retro-LN           1.240 0.145 8.53 1.46e-17
retro-marrow       0.920 0.200 4.60 4.21e-06
retro-spleen       0.812 0.293 2.77 5.53e-03
retro-thymus       1.290 0.230 5.60 2.11e-08
```

And here we count genes in the upper $1/16^{th}$:

```
Density data too sparse for barplot
```

```
                         coef    se      z          p
lenti-LN               0.8250 0.145 5.690 1.27e-08
lenti-Sca1             0.0507 0.185 0.274 7.84e-01
lenti-marrow           0.3250 0.352 0.924 3.56e-01
lenti-pleural-fluid    1.1700 0.589 1.980 4.72e-02
lenti-spleen           0.7030 0.217 3.240 1.18e-03
lenti-thymus           0.8070 0.348 2.320 2.05e-02
retro-LN               1.0600 0.181 5.840 5.15e-09
retro-marrow           1.2300 0.234 5.240 1.59e-07
retro-spleen           1.2000 0.357 3.360 7.81e-04
retro-thymus           1.1800 0.278 4.260 2.08e-05
```

Here the effect of density of CpG islands is studied:

rescale(cpg.dens.100k)
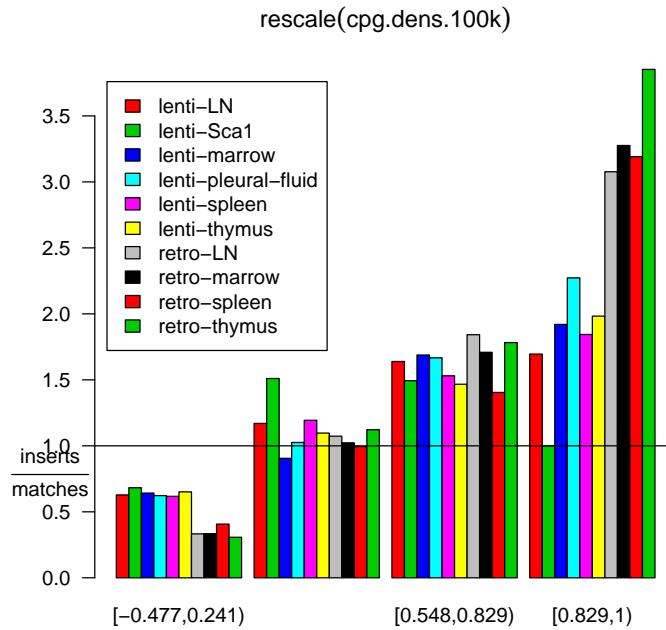
	coef	se	z	p
lenti-LN	0.844	0.0868	9.72	2.53e-22
lenti-Sca1	0.704	0.0855	8.24	1.77e-16
lenti-marrow	0.787	0.1920	4.10	4.09e-05
lenti-pleural-fluid	0.873	0.3000	2.91	3.57e-03
lenti-spleen	0.847	0.1240	6.85	7.41e-12
lenti-thymus	0.784	0.1950	4.03	5.58e-05
retro-LN	1.690	0.1480	11.40	2.60e-30
retro-marrow	1.650	0.1980	8.34	7.39e-17
retro-spleen	1.370	0.2670	5.15	2.60e-07
retro-thymus	1.830	0.2270	8.03	9.38e-16

4.4   250 kilobase Window

In the barplot that follows we examine the association of insertion sites with expression density in a 250 kilobase window surrounding each locus. First, we count just the number of genes represented on the chip.

rescale(dens.250k)
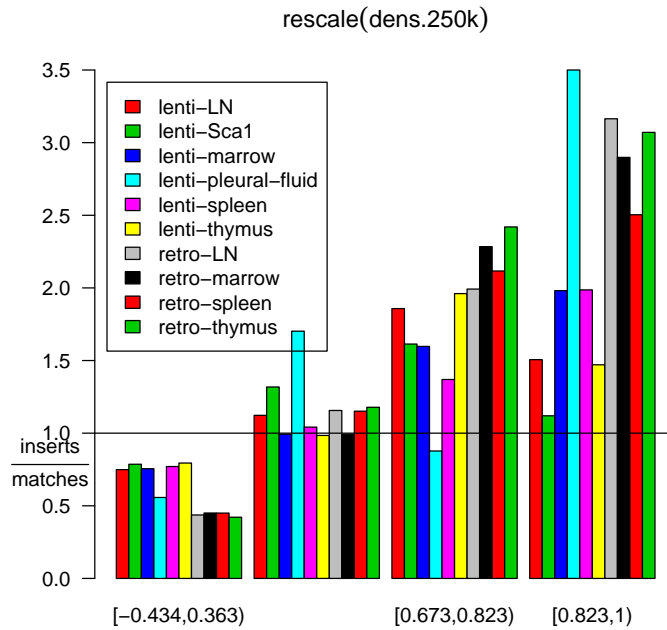
	coef	se	z	p
lenti-LN	0.644	0.0834	7.72	1.12e-14
lenti-Sca1	0.511	0.0837	6.11	1.03e-09
lenti-marrow	0.634	0.1860	3.41	6.55e-04
lenti-pleural-fluid	1.120	0.2930	3.82	1.35e-04
lenti-spleen	0.566	0.1190	4.76	1.90e-06
lenti-thymus	0.516	0.1880	2.75	6.02e-03
retro-LN	1.440	0.1320	10.90	6.90e-28
retro-marrow	1.400	0.1780	7.86	3.88e-15
retro-spleen	1.420	0.2590	5.49	4.04e-08
retro-thymus	1.510	0.2000	7.56	4.00e-14

Here are the results for expression density. First, we count just genes that are in the upper half.

rescale(low.ex.250k)
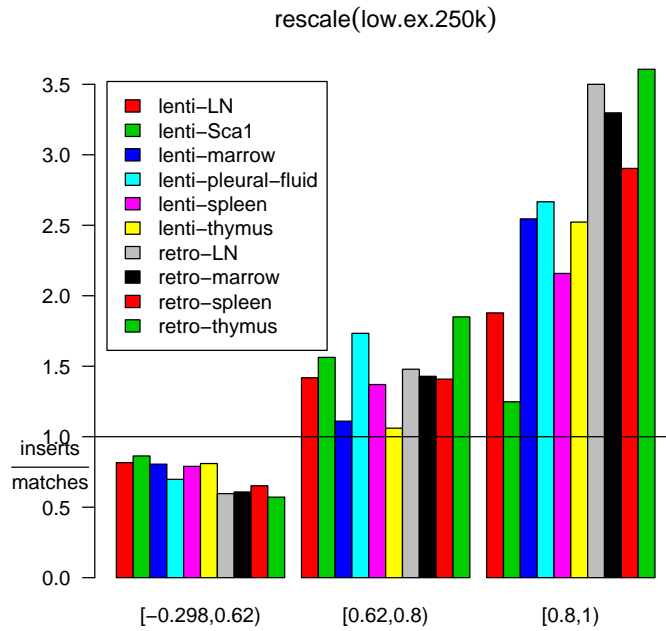
	coef	se	z	p
lenti-LN	0.701	0.0840	8.35	7.10e-17
lenti-Sca1	0.500	0.0869	5.75	8.68e-09
lenti-marrow	0.684	0.1870	3.66	2.50e-04
lenti-pleural-fluid	0.988	0.2860	3.45	5.55e-04
lenti-spleen	0.700	0.1210	5.81	6.43e-09
lenti-thymus	0.597	0.1910	3.12	1.79e-03
retro-LN	1.380	0.1200	11.50	1.35e-30
retro-marrow	1.300	0.1580	8.22	2.05e-16
retro-spleen	1.230	0.2280	5.38	7.45e-08
retro-thymus	1.540	0.1810	8.54	1.32e-17

Now we count genes in the upper $1/8^{th}$:

rescale(med.ex.250k)
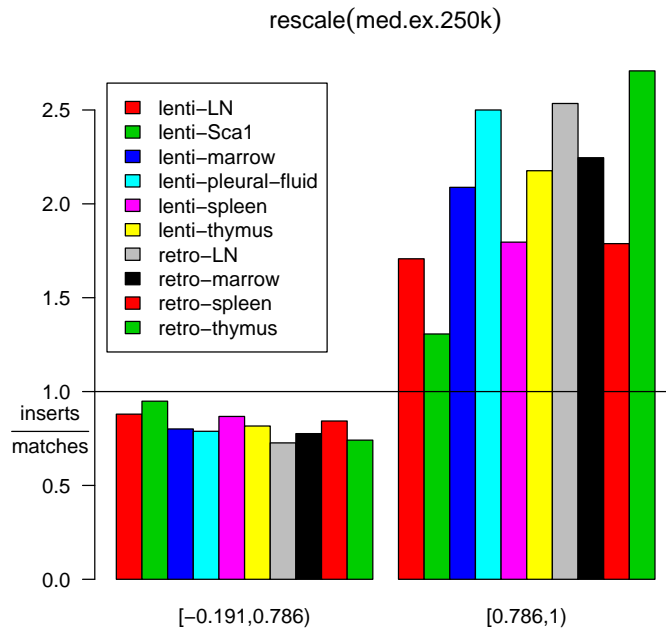
```
                        coef     se     z        p
lenti-LN               0.708  0.0921   7.69  1.45e-14
lenti-Sca1             0.269  0.1020   2.64  8.23e-03
lenti-marrow           0.721  0.1980   3.64  2.68e-04
lenti-pleural-fluid    0.953  0.3020   3.15  1.61e-03
lenti-spleen           0.663  0.1320   5.01  5.37e-07
lenti-thymus           0.853  0.2050   4.16  3.20e-05
retro-LN               1.240  0.1210  10.20  1.85e-24
retro-marrow           1.030  0.1590   6.47  1.01e-10
retro-spleen           0.650  0.2340   2.77  5.56e-03
retro-thymus           1.240  0.1880   6.57  4.92e-11
```

And here we count genes in the upper $1/16^{th}$:

rescale(high.ex.250k)
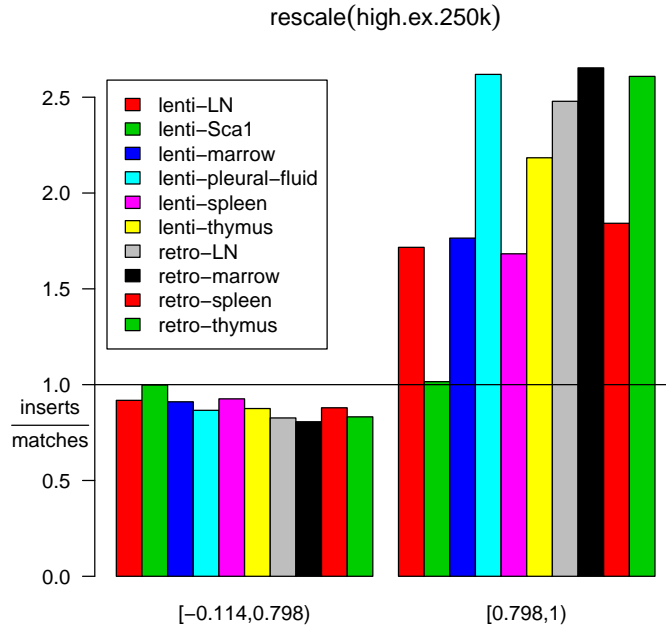
```
                         coef    se       z         p
lenti-LN              0.65300  0.109  5.9700  2.37e-09
lenti-Sca1            0.00165  0.135  0.0123  9.90e-01
lenti-marrow          0.71000  0.237  3.0000  2.70e-03
lenti-pleural-fluid   1.01000  0.363  2.7900  5.23e-03
lenti-spleen          0.64900  0.159  4.0700  4.61e-05
lenti-thymus          0.91700  0.233  3.9300  8.47e-05
retro-LN              1.12000  0.137  8.1400  4.08e-16
retro-marrow          1.16000  0.177  6.5600  5.56e-11
retro-spleen          0.71600  0.261  2.7500  6.01e-03
retro-thymus          1.14000  0.212  5.3800  7.30e-08
```

Here the effect of density of CpG islands is studied:

rescale(cpg.dens.250k)
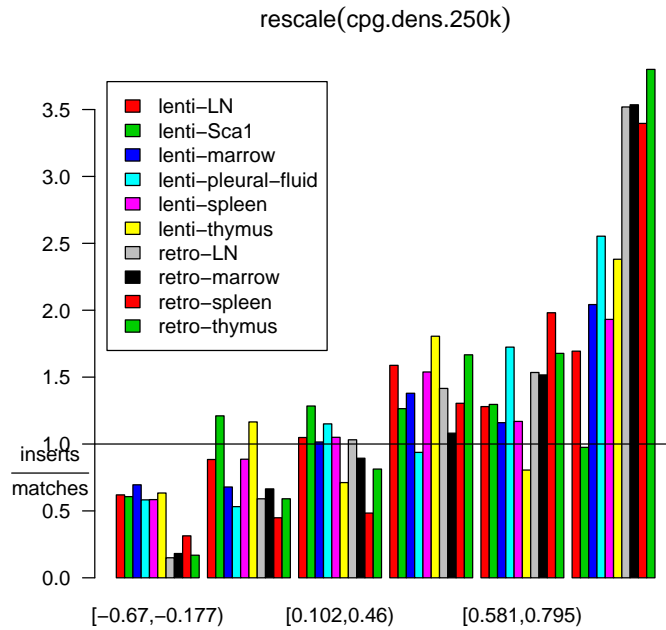
	coef	se	z	p
lenti-LN	0.621	0.0828	7.49	6.76e-14
lenti-Sca1	0.234	0.0843	2.77	5.56e-03
lenti-marrow	0.589	0.1830	3.21	1.31e-03
lenti-pleural-fluid	0.928	0.2870	3.24	1.22e-03
lenti-spleen	0.733	0.1190	6.14	8.46e-10
lenti-thymus	0.627	0.1890	3.32	8.99e-04
retro-LN	1.720	0.1350	12.70	6.68e-37
retro-marrow	1.500	0.1760	8.53	1.44e-17
retro-spleen	1.570	0.2560	6.16	7.38e-10
retro-thymus	2.010	0.2140	9.37	7.25e-21

4.5   500 kilobase Window

In the barplot that follows we examine the association of insertion sites with expression density in a 500 kilobase window surrounding each locus. First, we count just the number of genes represented on the chip.

rescale(dens.500k)
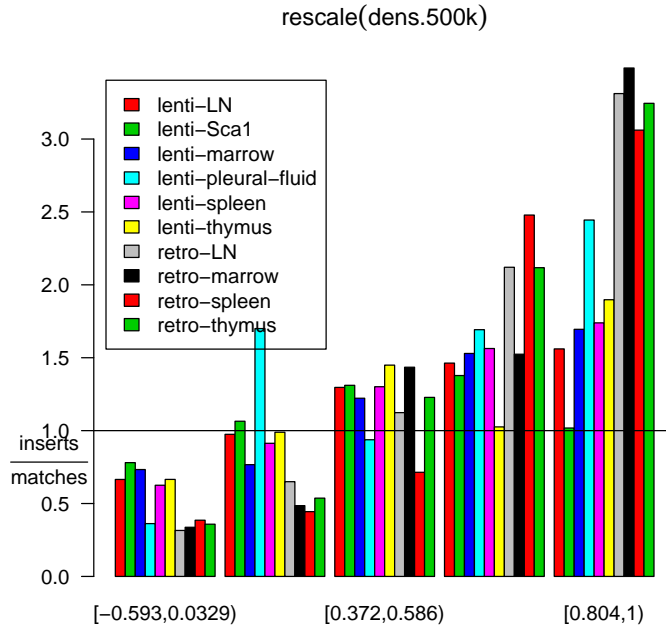
	coef	se	z	p
lenti-LN	0.648	0.0830	7.81	5.52e-15
lenti-Sca1	0.357	0.0841	4.24	2.19e-05
lenti-marrow	0.745	0.1860	4.02	5.92e-05
lenti-pleural-fluid	0.967	0.2920	3.31	9.31e-04
lenti-spleen	0.777	0.1190	6.53	6.75e-11
lenti-thymus	0.647	0.1880	3.44	5.82e-04
retro-LN	1.540	0.1290	11.90	9.06e-33
retro-marrow	1.590	0.1760	9.01	2.08e-19
retro-spleen	1.550	0.2530	6.13	8.93e-10
retro-thymus	1.590	0.1940	8.15	3.62e-16

Here are the results for expression density. First, we count just genes that are in the upper half.

rescale(low.ex.500k)
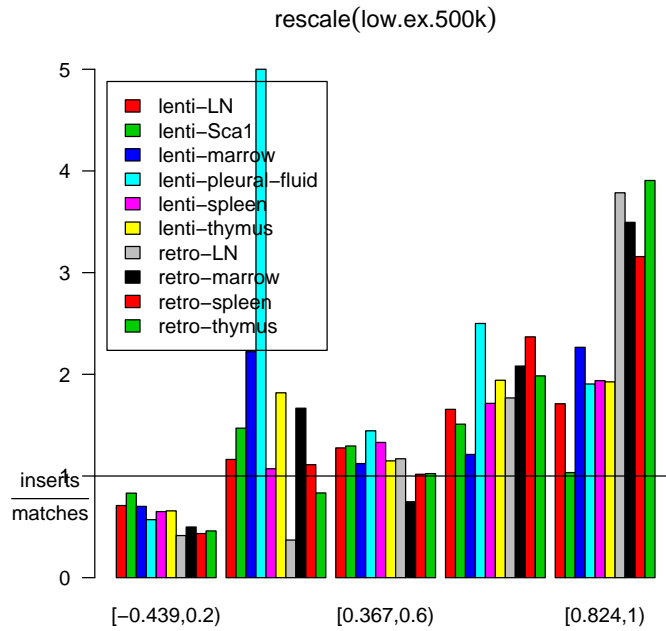
	coef	se	z	p
lenti-LN	0.777	0.0845	9.19	3.79e-20
lenti-Sca1	0.408	0.0837	4.88	1.07e-06
lenti-marrow	0.708	0.1900	3.73	1.90e-04
lenti-pleural-fluid	1.280	0.3030	4.23	2.35e-05
lenti-spleen	0.859	0.1220	7.05	1.73e-12
lenti-thymus	0.752	0.1950	3.86	1.14e-04
retro-LN	1.620	0.1390	11.60	2.94e-31
retro-marrow	1.430	0.1800	7.96	1.67e-15
retro-spleen	1.600	0.2660	6.00	1.96e-09
retro-thymus	1.480	0.1970	7.50	6.30e-14

Now we count genes in the upper $1/8^{th}$:

rescale(med.ex.500k)
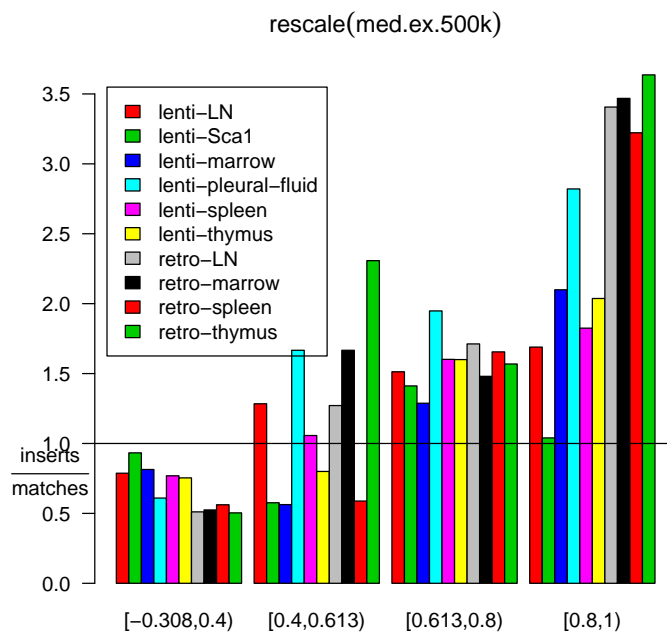
	coef	se	z	p
lenti-LN	0.659	0.0840	7.85	4.26e-15
lenti-Sca1	0.279	0.0882	3.16	1.59e-03
lenti-marrow	0.512	0.1850	2.77	5.55e-03
lenti-pleural-fluid	1.200	0.2830	4.24	2.25e-05
lenti-spleen	0.718	0.1200	5.98	2.26e-09
lenti-thymus	0.729	0.1910	3.81	1.38e-04
retro-LN	1.470	0.1210	12.10	1.07e-33
retro-marrow	1.410	0.1620	8.67	4.31e-18
retro-spleen	1.320	0.2300	5.71	1.10e-08
retro-thymus	1.550	0.1830	8.47	2.39e-17

And here we count genes in the upper $1/16^{th}$:

rescale(high.ex.500k)
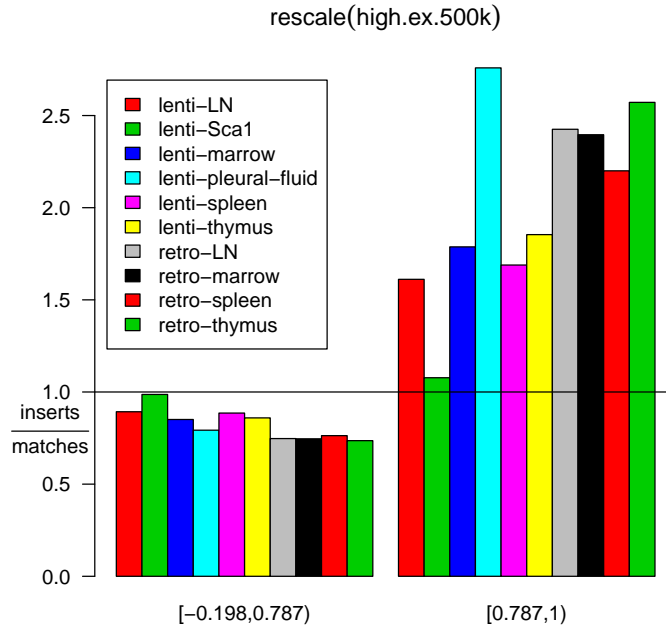
	coef	se	z	p
lenti-LN	0.5740	0.0928	6.180	6.39e-10
lenti-Sca1	0.0906	0.1050	0.866	3.87e-01
lenti-marrow	0.6620	0.2010	3.290	1.00e-03
lenti-pleural-fluid	1.2400	0.2880	4.300	1.68e-05
lenti-spleen	0.6720	0.1330	5.070	4.04e-07
lenti-thymus	0.8350	0.2040	4.100	4.11e-05
retro-LN	1.1600	0.1190	9.710	2.72e-22
retro-marrow	1.1800	0.1580	7.460	8.74e-14
retro-spleen	1.0900	0.2260	4.810	1.52e-06
retro-thymus	1.3000	0.1820	7.120	1.08e-12

Here the effect of density of CpG islands is studied:

rescale(cpg.dens.500k)
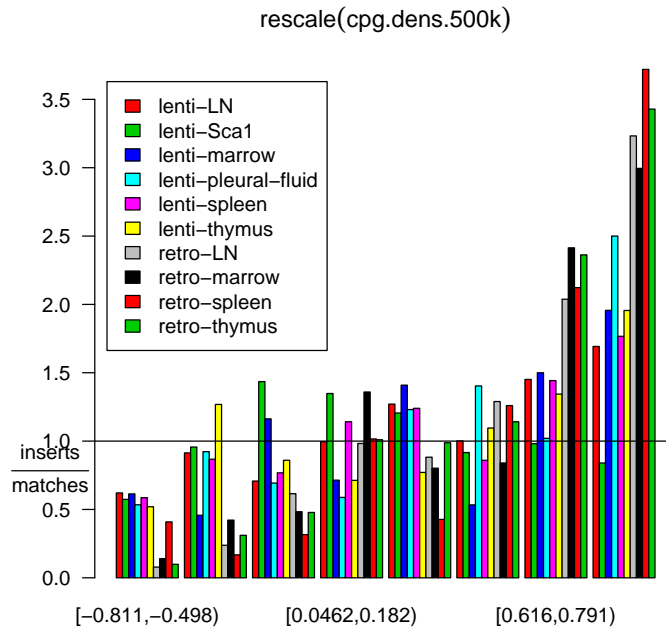
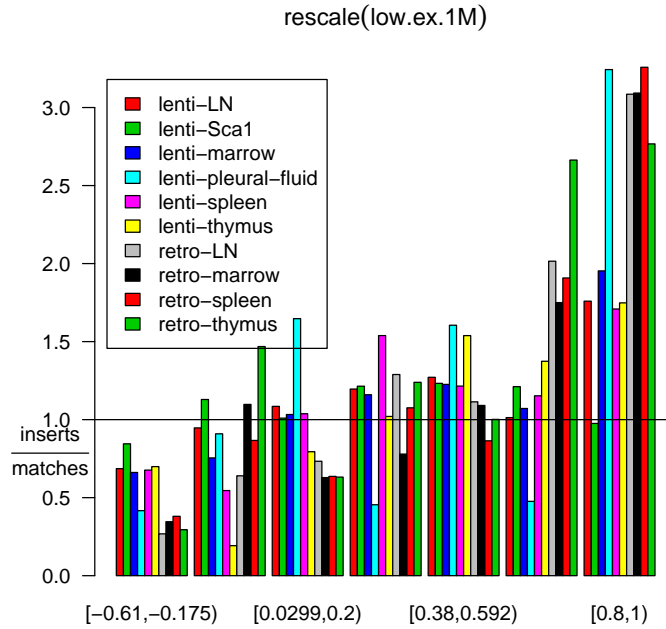|	coef	se	z	p
lenti-LN	0.5630	0.0831	6.770	1.25e-11
lenti-Sca1	-0.0163	0.0847	-0.192	8.47e-01
lenti-marrow	0.5700	0.1860	3.060	2.25e-03
lenti-pleural-fluid	0.8170	0.2890	2.830	4.69e-03
lenti-spleen	0.5260	0.1180	4.440	8.99e-06
lenti-thymus	0.4510	0.1870	2.420	1.57e-02
retro-LN	1.5100	0.1370	11.100	2.13e-28
retro-marrow	1.3400	0.1770	7.550	4.30e-14
retro-spleen	1.4800	0.2610	5.680	1.38e-08
retro-thymus	1.6800	0.2140	7.860	3.87e-15

4.6 1 megabase Window

In the barplot that follows we examine the association of insertion sites with expression density in a 1 megabase window surrounding each locus. First, we count just the number of genes represented on the chip.

rescale(dens.1M)
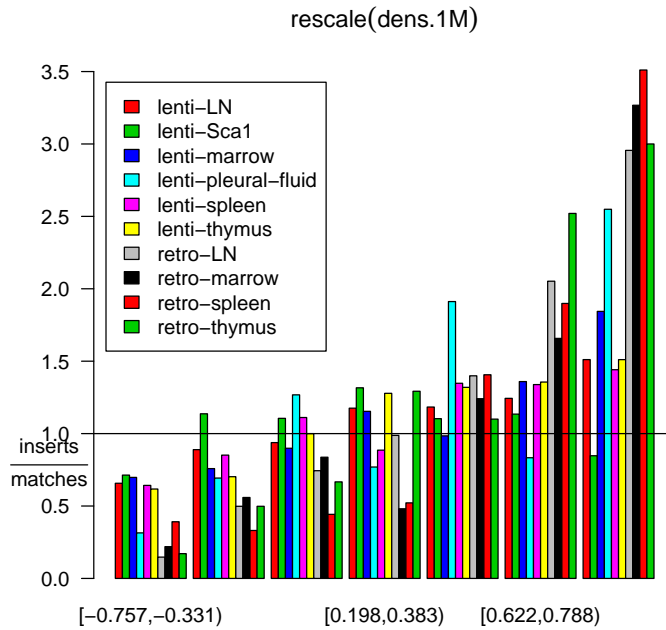
	coef	se	z	p
lenti-LN	0.437	0.0830	5.27	1.37e-07
lenti-Sca1	0.190	0.0836	2.27	2.34e-02
lenti-marrow	0.543	0.1870	2.90	3.70e-03
lenti-pleural-fluid	0.901	0.2930	3.07	2.12e-03
lenti-spleen	0.461	0.1180	3.91	9.41e-05
lenti-thymus	0.631	0.1940	3.26	1.13e-03
retro-LN	1.510	0.1390	10.90	1.36e-27
retro-marrow	1.240	0.1780	6.96	3.52e-12
retro-spleen	1.400	0.2670	5.26	1.47e-07
retro-thymus	1.610	0.2130	7.58	3.52e-14

Here are the results for expression density. First, we count just genes that are in the upper half.

rescale(low.ex.1M)

	coef	se	z	p
lenti-LN	0.479	0.0826	5.80	6.74e-09
lenti-Sca1	0.243	0.0842	2.89	3.90e-03
lenti-marrow	0.548	0.1850	2.97	3.00e-03
lenti-pleural-fluid	0.684	0.2840	2.41	1.61e-02
lenti-spleen	0.557	0.1180	4.71	2.42e-06
lenti-thymus	0.765	0.1910	4.01	6.11e-05
retro-LN	1.520	0.1320	11.60	5.33e-31
retro-marrow	1.310	0.1680	7.84	4.60e-15
retro-spleen	1.350	0.2470	5.47	4.55e-08
retro-thymus	1.420	0.1920	7.39	1.43e-13

Now we count genes in the upper $1/8^{th}$:

rescale(med.ex.1M)

```
                        coef     se     z        p
lenti-LN               0.447 0.0832  5.37 7.93e-08
lenti-Sca1             0.178 0.0835  2.13 3.34e-02
lenti-marrow           0.312 0.1840  1.69 9.05e-02
lenti-pleural-fluid    0.765 0.2900  2.63 8.48e-03
lenti-spleen           0.399 0.1180  3.38 7.34e-04
lenti-thymus           0.447 0.1900  2.36 1.85e-02
retro-LN               1.490 0.1390 10.80 4.52e-27
retro-marrow           1.320 0.1810  7.26 3.79e-13
retro-spleen           1.240 0.2520  4.92 8.65e-07
retro-thymus           1.550 0.2070  7.49 6.84e-14
```
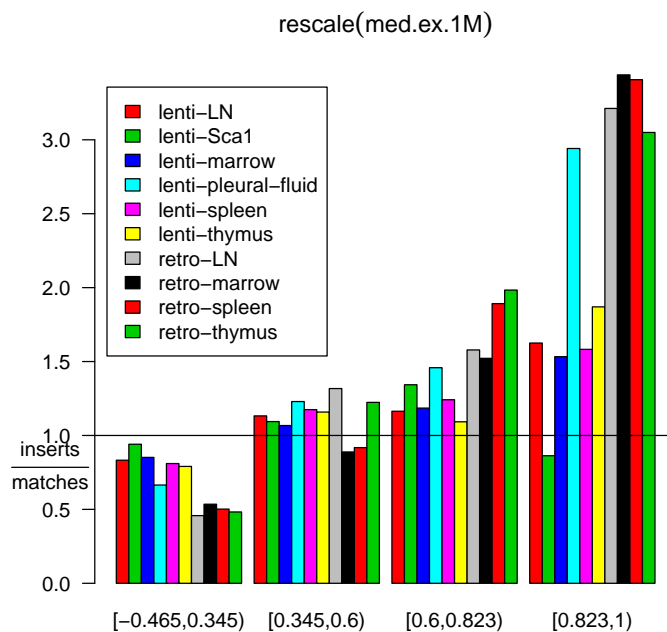
And here we count genes in the upper $1/16^{th}$:

rescale(high.ex.1M)
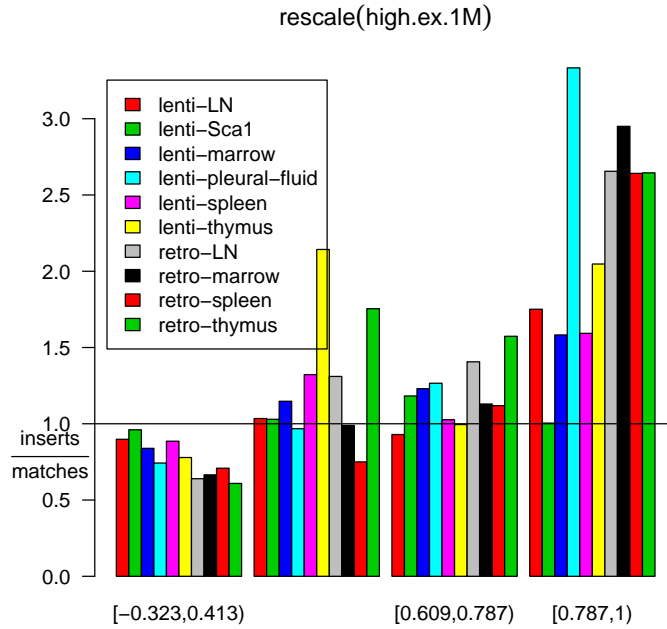
	coef	se	z	p
lenti-LN	0.312	0.0852	3.66	2.54e-04
lenti-Sca1	0.153	0.0883	1.73	8.33e-02
lenti-marrow	0.459	0.1870	2.45	1.42e-02
lenti-pleural-fluid	0.763	0.2840	2.69	7.18e-03
lenti-spleen	0.395	0.1210	3.25	1.13e-03
lenti-thymus	0.624	0.1900	3.28	1.04e-03
retro-LN	1.040	0.1170	8.90	5.78e-19
retro-marrow	0.941	0.1550	6.09	1.13e-09
retro-spleen	0.886	0.2220	3.99	6.59e-05
retro-thymus	1.110	0.1750	6.34	2.33e-10

Here the effect of density of CpG islands is studied:

rescale(cpg.dens.1M)
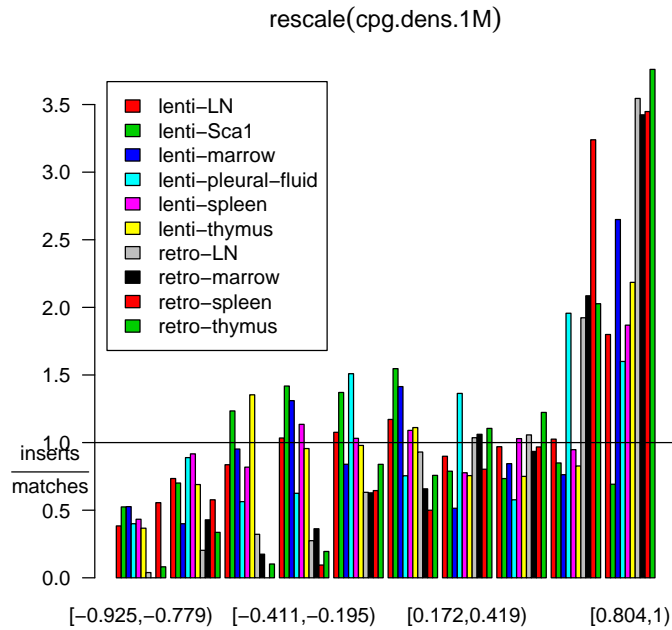
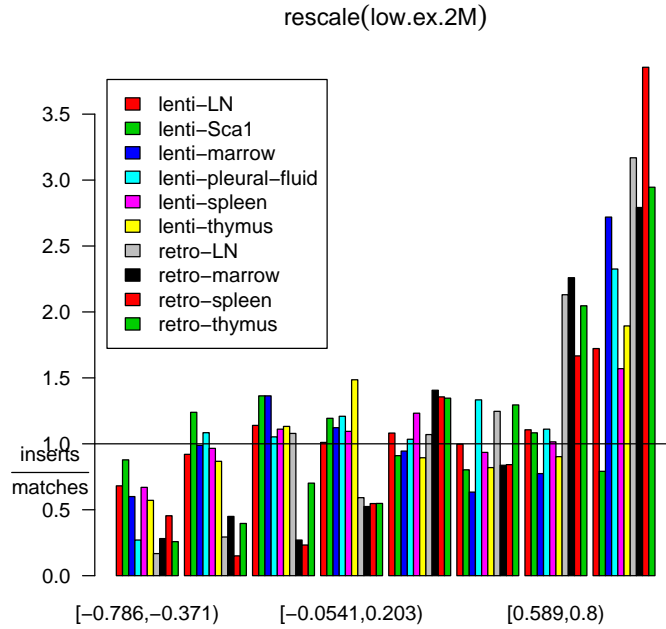|	coef	se	z	p
lenti-LN	0.2620	0.0823	3.18	1.48e-03
lenti-Sca1	-0.2570	0.0850	-3.02	2.52e-03
lenti-marrow	0.2230	0.1840	1.22	2.24e-01
lenti-pleural-fluid	0.6290	0.2830	2.22	2.65e-02
lenti-spleen	0.2220	0.1180	1.89	5.86e-02
lenti-thymus	0.0678	0.1880	0.36	7.19e-01
retro-LN	1.6200	0.1470	11.00	2.72e-28
retro-marrow	1.5600	0.1950	8.00	1.25e-15
retro-spleen	1.5700	0.2770	5.66	1.53e-08
retro-thymus	1.7800	0.2280	7.80	6.15e-15

4.7   2 megabase Window

In the barplot that follows we examine the association of insertion sites with expression density in a 2 megabase window surrounding each locus. First, we count just the number of genes represented on the chip.

rescale(dens.2M)
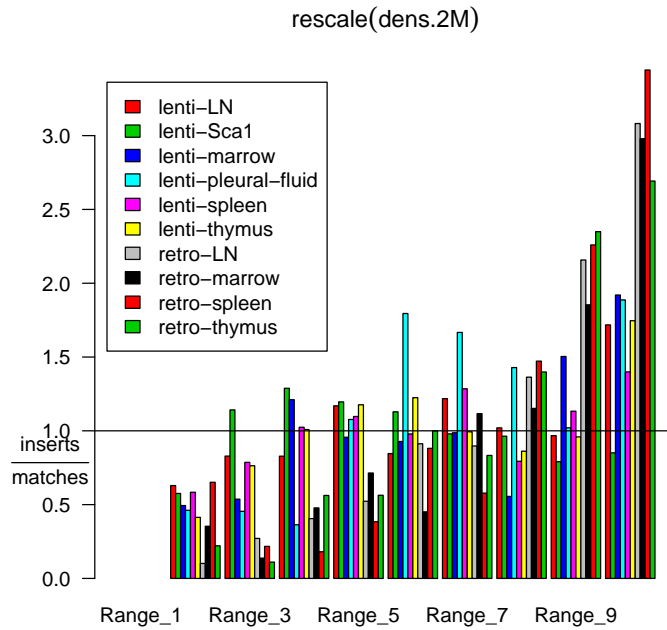
	coef	se	z	p
lenti-LN	0.367	0.0826	4.44	8.79e-06
lenti-Sca1	-0.170	0.0854	-1.99	4.65e-02
lenti-marrow	0.393	0.1850	2.13	3.35e-02
lenti-pleural-fluid	0.761	0.2880	2.65	8.13e-03
lenti-spleen	0.300	0.1180	2.55	1.07e-02
lenti-thymus	0.311	0.1880	1.65	9.81e-02
retro-LN	1.480	0.1370	10.80	4.37e-27
retro-marrow	1.460	0.1850	7.88	3.28e-15
retro-spleen	1.740	0.2870	6.06	1.37e-09
retro-thymus	1.490	0.2050	7.27	3.71e-13

Here are the results for expression density. First, we count just genes that are in the upper half.

rescale(low.ex.2M)

	coef	se	z	p
lenti-LN	0.331	0.0827	4.000	6.42e-05
lenti-Sca1	-0.132	0.0842	-1.570	1.17e-01
lenti-marrow	0.264	0.1840	1.430	1.51e-01
lenti-pleural-fluid	0.613	0.2860	2.140	3.22e-02
lenti-spleen	0.250	0.1180	2.120	3.41e-02
lenti-thymus	0.109	0.1880	0.581	5.61e-01
retro-LN	1.560	0.1460	10.700	1.23e-26
retro-marrow	1.570	0.1940	8.080	6.28e-16
retro-spleen	1.590	0.2790	5.710	1.12e-08
retro-thymus	1.500	0.2110	7.100	1.26e-12

Now we count genes in the upper $1/8^{th}$:

rescale(med.ex.2M)
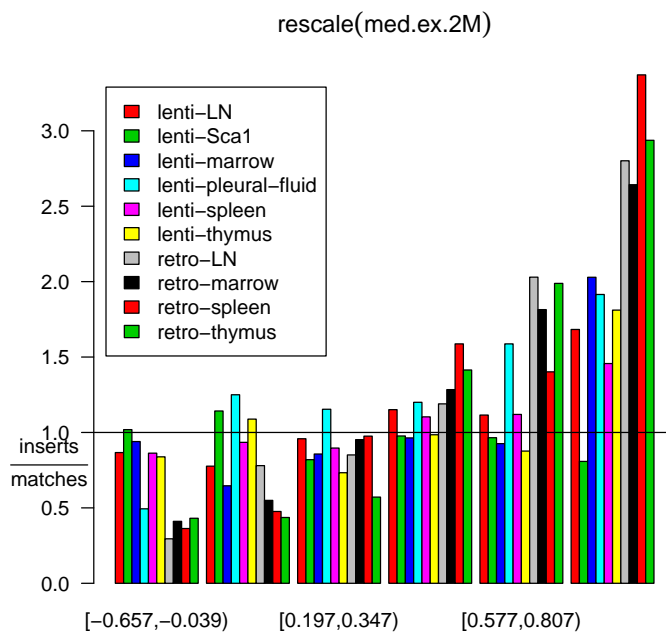
	coef	se	z	p
lenti-LN	0.372	0.0822	4.52	6.23e-06
lenti-Sca1	-0.124	0.0851	-1.45	1.46e-01
lenti-marrow	0.263	0.1830	1.44	1.51e-01
lenti-pleural-fluid	0.751	0.2900	2.59	9.63e-03
lenti-spleen	0.261	0.1180	2.21	2.70e-02
lenti-thymus	0.216	0.1880	1.15	2.50e-01
retro-LN	1.340	0.1320	10.20	3.17e-24
retro-marrow	1.400	0.1780	7.89	2.93e-15
retro-spleen	1.530	0.2590	5.92	3.30e-09
retro-thymus	1.410	0.1980	7.12	1.08e-12

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And here we count genes in the upper $1/16^{th}$:

rescale(high.ex.2M)
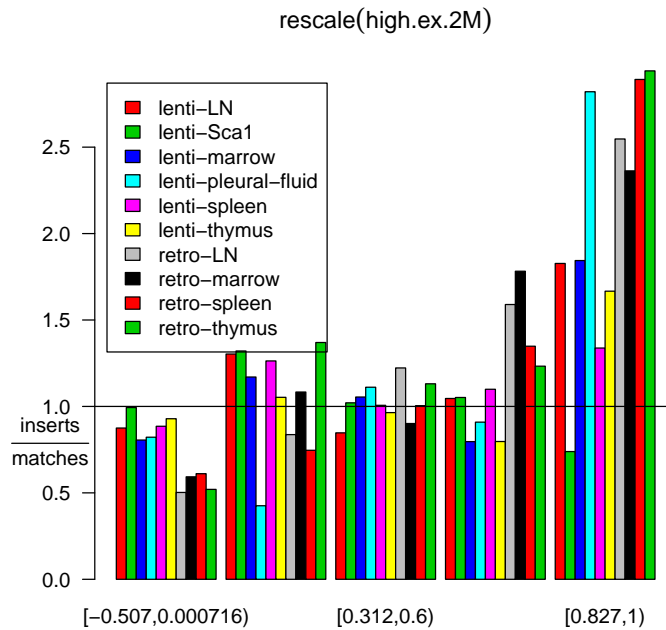
	coef	se	z	p
lenti-LN	0.2510	0.0827	3.030	2.46e-03
lenti-Sca1	0.0107	0.0833	0.129	8.98e-01
lenti-marrow	0.4110	0.1880	2.190	2.87e-02
lenti-pleural-fluid	0.3870	0.2880	1.340	1.79e-01
lenti-spleen	0.2350	0.1190	1.980	4.74e-02
lenti-thymus	0.1520	0.1870	0.813	4.16e-01
retro-LN	1.1200	0.1320	8.460	2.58e-17
retro-marrow	0.8560	0.1680	5.110	3.21e-07
retro-spleen	0.8150	0.2400	3.400	6.70e-04
retro-thymus	1.0900	0.1940	5.610	2.05e-08

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Here the effect of density of CpG islands is studied:

rescale(cpg.dens.2M)
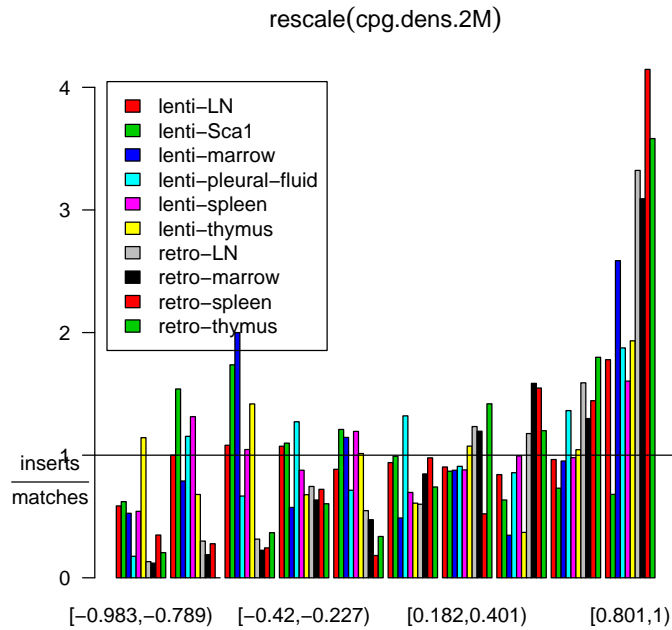
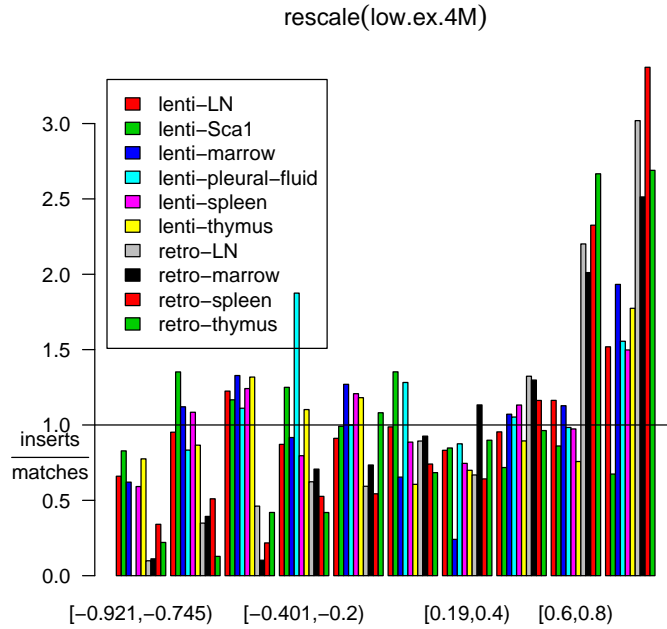|	coef	se	z	p
lenti-LN	0.1070	0.0823	1.300	1.95e-01
lenti-Sca1	-0.4570	0.0861	-5.300	1.13e-07
lenti-marrow	-0.0249	0.1850	-0.135	8.93e-01
lenti-pleural-fluid	0.4730	0.2840	1.670	9.51e-02
lenti-spleen	0.0178	0.1180	0.151	8.80e-01
lenti-thymus	-0.0771	0.1880	-0.410	6.82e-01
retro-LN	1.3600	0.1400	9.740	2.06e-22
retro-marrow	1.5600	0.1970	7.930	2.16e-15
retro-spleen	1.6300	0.2850	5.720	1.08e-08
retro-thymus	1.7500	0.2270	7.700	1.35e-14

4.8   4 megabase Window

In the barplot that follows we examine the association of insertion sites with
expression density in a 4 megabase window surrounding each locus. First, we
count just the number of genes represented on the chip.

rescale(dens.4M)
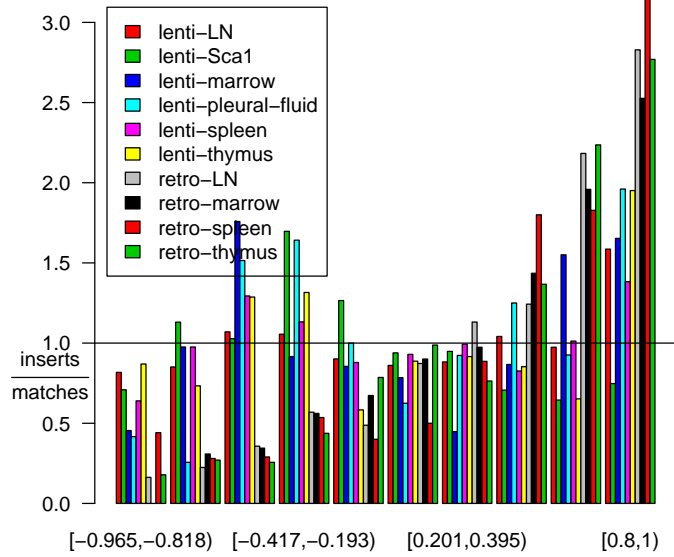
	coef	se	z	p
lenti-LN	0.1620	0.0823	1.960	4.96e-02
lenti-Sca1	-0.3930	0.0863	-4.560	5.21e-06
lenti-marrow	0.1720	0.1840	0.934	3.50e-01
lenti-pleural-fluid	0.1740	0.2820	0.616	5.38e-01
lenti-spleen	0.0465	0.1180	0.396	6.92e-01
lenti-thymus	0.1960	0.1870	1.050	2.93e-01
retro-LN	1.4900	0.1420	10.500	7.58e-26
retro-marrow	1.3100	0.1820	7.210	5.41e-13
retro-spleen	1.5600	0.2740	5.690	1.30e-08
retro-thymus	1.3600	0.2050	6.650	2.84e-11

Here are the results for expression density. First, we count just genes that are in the upper half.

rescale(low.ex.4M)

	coef	se	z	p
lenti-LN	0.14300	0.0821	1.7400	8.27e-02
lenti-Sca1	-0.27700	0.0854	-3.2500	1.17e-03
lenti-marrow	-0.00619	0.1850	-0.0336	9.73e-01
lenti-pleural-fluid	0.16600	0.2820	0.5900	5.55e-01
lenti-spleen	0.06730	0.1180	0.5700	5.69e-01
lenti-thymus	-0.01960	0.1900	-0.1030	9.18e-01
retro-LN	1.34000	0.1350	9.9100	3.61e-23
retro-marrow	1.35000	0.1810	7.4600	8.98e-14
retro-spleen	1.25000	0.2520	4.9400	7.93e-07
retro-thymus	1.28000	0.1980	6.4800	9.23e-11

Now we count genes in the upper $1/8^{th}$:

rescale(med.ex.4M)
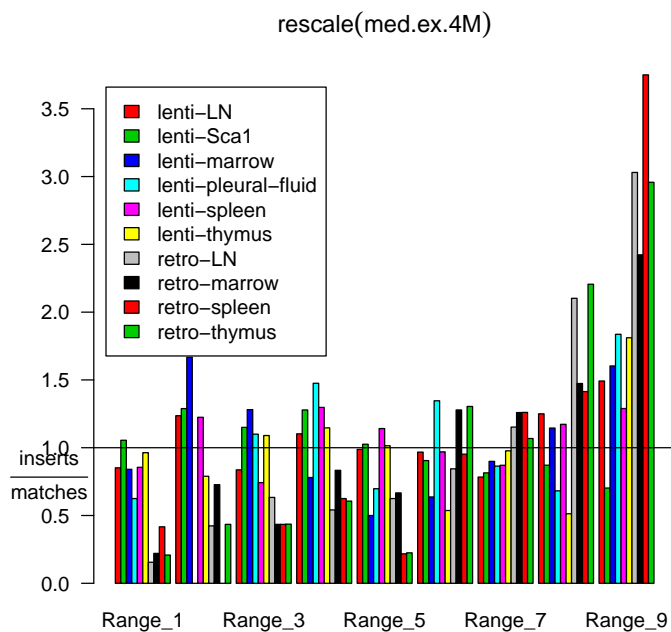

```
                        coef     se      z        p
lenti-LN              0.1400 0.0821   1.700 8.85e-02
lenti-Sca1           -0.3410 0.0858  -3.980 7.01e-05
lenti-marrow          0.0523 0.1840   0.284 7.76e-01
lenti-pleural-fluid   0.3170 0.2810   1.130 2.59e-01
lenti-spleen          0.0617 0.1180   0.525 6.00e-01
lenti-thymus         -0.0814 0.1900  -0.429 6.68e-01
retro-LN              1.2900 0.1340   9.660 4.32e-22
retro-marrow          1.2000 0.1760   6.830 8.45e-12
retro-spleen          1.3300 0.2550   5.220 1.78e-07
retro-thymus          1.4700 0.2090   7.010 2.40e-12
```

And here we count genes in the upper $1/16^{th}$:

rescale(high.ex.4M)
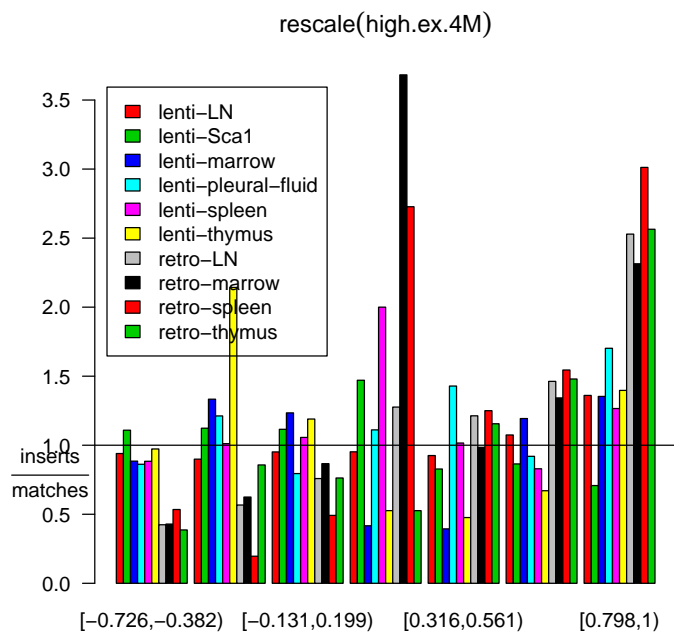
	coef	se	z	p
lenti-LN	0.1630	0.0825	1.970	4.84e-02
lenti-Sca1	-0.2030	0.0848	-2.400	1.65e-02
lenti-marrow	-0.1730	0.1850	-0.935	3.50e-01
lenti-pleural-fluid	0.2290	0.2800	0.819	4.13e-01
lenti-spleen	0.0641	0.1180	0.546	5.85e-01
lenti-thymus	-0.3230	0.1900	-1.700	9.00e-02
retro-LN	1.0300	0.1260	8.120	4.55e-16
retro-marrow	0.8350	0.1630	5.110	3.15e-07
retro-spleen	1.2900	0.2520	5.120	3.00e-07
retro-thymus	1.1100	0.1930	5.720	1.04e-08

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Here the effect of density of CpG islands is studied:

rescale(cpg.dens.4M)
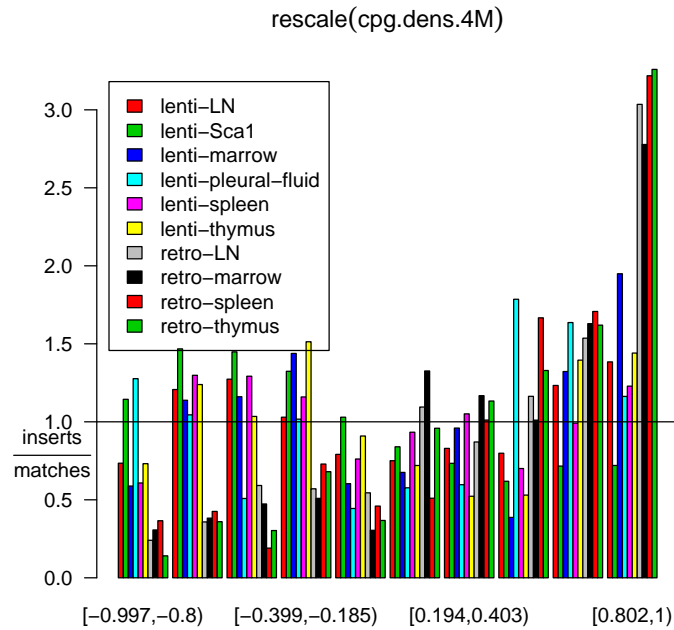
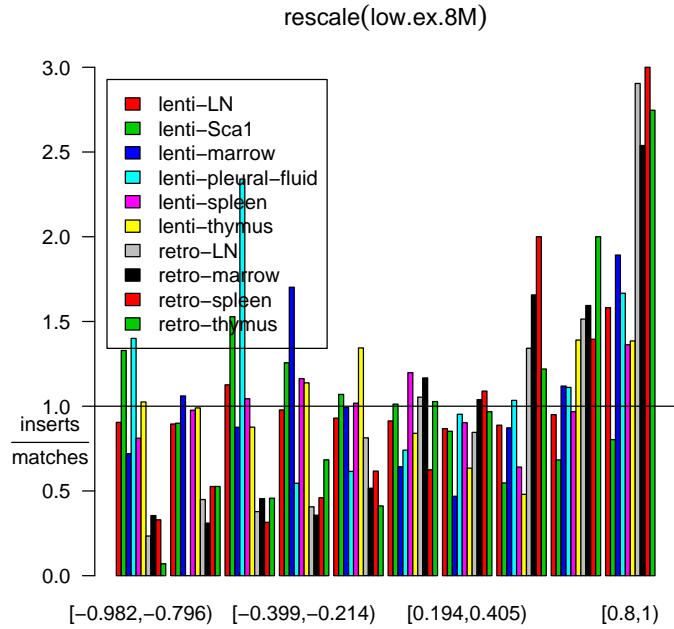|	coef	se	z	p
lenti-LN	-0.01310	0.0827	-0.1580	8.75e-01
lenti-Sca1	-0.55500	0.0874	-6.3500	2.21e-10
lenti-marrow	-0.00609	0.1830	-0.0333	9.73e-01
lenti-pleural-fluid	0.23100	0.2780	0.8290	4.07e-01
lenti-spleen	-0.08880	0.1170	-0.7560	4.50e-01
lenti-thymus	-0.27000	0.1880	-1.4400	1.51e-01
retro-LN	1.14000	0.1320	8.6500	5.03e-18
retro-marrow	1.34000	0.1830	7.2900	3.01e-13
retro-spleen	1.38000	0.2620	5.2500	1.48e-07
retro-thymus	1.40000	0.2130	6.6000	4.16e-11

4.9 8 megabase Window

In the barplot that follows we examine the association of insertion sites with
expression density in a 8 megabase window surrounding each locus. First, we
count just the number of genes represented on the chip.

rescale(dens.8M)
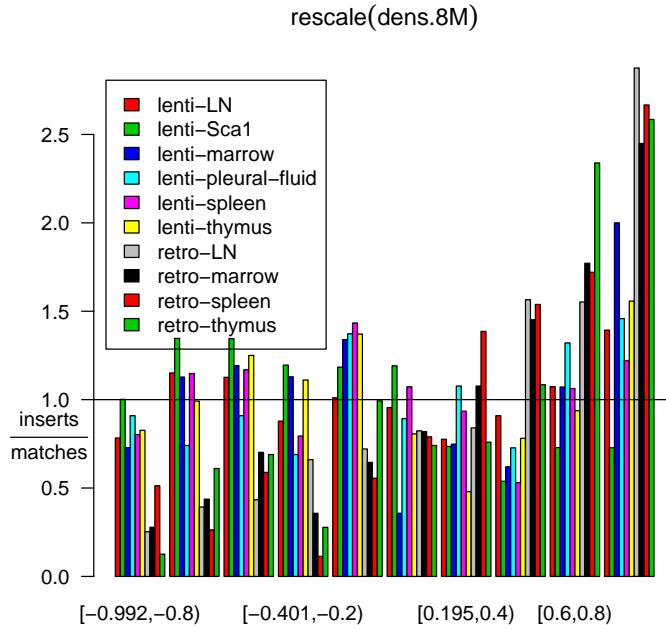
	coef	se	z	p
lenti-LN	0.0274	0.0828	0.331	7.40e-01
lenti-Sca1	-0.4440	0.0857	-5.180	2.20e-07
lenti-marrow	-0.1480	0.1840	-0.803	4.22e-01
lenti-pleural-fluid	0.1670	0.2830	0.589	5.56e-01
lenti-spleen	-0.1020	0.1180	-0.865	3.87e-01
lenti-thymus	-0.2110	0.1870	-1.130	2.58e-01
retro-LN	1.1200	0.1330	8.430	3.45e-17
retro-marrow	1.1300	0.1750	6.460	1.01e-10
retro-spleen	1.4300	0.2680	5.320	1.04e-07
retro-thymus	1.0100	0.1950	5.190	2.06e-07

Here are the results for expression density. First, we count just genes that are in the upper half.


rescale(low.ex.8M)

	coef	se	z	p
lenti-LN	0.0493	0.0828	0.595	5.52e-01
lenti-Sca1	-0.4560	0.0861	-5.300	1.18e-07
lenti-marrow	-0.0645	0.1840	-0.351	7.26e-01
lenti-pleural-fluid	0.1290	0.2810	0.459	6.46e-01
lenti-spleen	-0.0541	0.1180	-0.460	6.46e-01
lenti-thymus	-0.1200	0.1870	-0.644	5.19e-01
retro-LN	1.1600	0.1350	8.620	6.71e-18
retro-marrow	1.2800	0.1810	7.070	1.55e-12
retro-spleen	1.2900	0.2590	5.000	5.82e-07
retro-thymus	1.3000	0.2080	6.270	3.58e-10

Now we count genes in the upper $1/8^{th}$:

rescale(med.ex.8M)
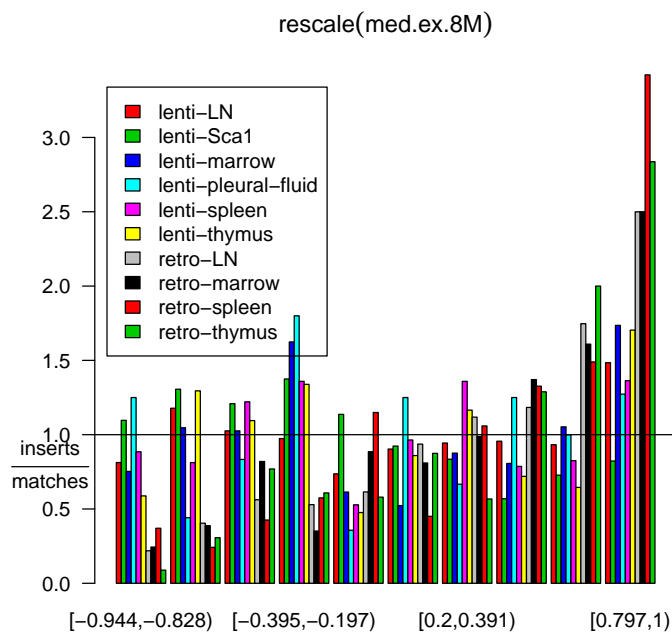
	coef	se	z	p
lenti-LN	0.0901	0.0824	1.090	2.74e-01
lenti-Sca1	-0.5100	0.0868	-5.880	4.22e-09
lenti-marrow	-0.0522	0.1840	-0.284	7.76e-01
lenti-pleural-fluid	0.2890	0.2830	1.020	3.08e-01
lenti-spleen	0.0806	0.1180	0.685	4.94e-01
lenti-thymus	0.0771	0.1880	0.410	6.81e-01
retro-LN	1.1100	0.1300	8.500	1.89e-17
retro-marrow	1.0200	0.1690	6.040	1.57e-09
retro-spleen	1.1000	0.2480	4.430	9.27e-06
retro-thymus	1.1200	0.1990	5.660	1.52e-08

And here we count genes in the upper $1/16^{th}$:

rescale(high.ex.8M)
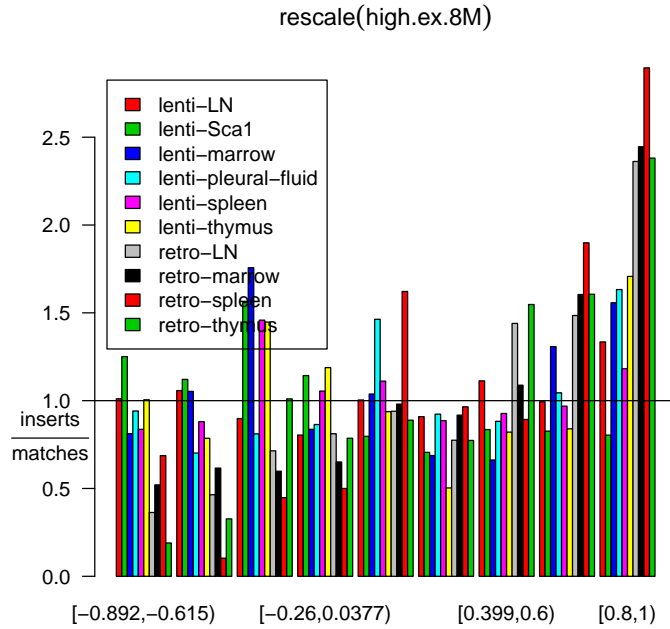
	coef	se	z	p
lenti-LN	0.1570	0.0827	1.900	5.71e-02
lenti-Sca1	-0.3650	0.0868	-4.200	2.61e-05
lenti-marrow	0.0576	0.1820	0.316	7.52e-01
lenti-pleural-fluid	0.2080	0.2810	0.739	4.60e-01
lenti-spleen	0.0230	0.1180	0.194	8.46e-01
lenti-thymus	-0.1820	0.1900	-0.958	3.38e-01
retro-LN	0.8430	0.1210	6.960	3.44e-12
retro-marrow	0.7950	0.1580	5.030	4.93e-07
retro-spleen	0.9160	0.2320	3.950	7.76e-05
retro-thymus	0.8990	0.1840	4.900	9.74e-07

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Here the effect of density of CpG islands is studied:

rescale(cpg.dens.8M)
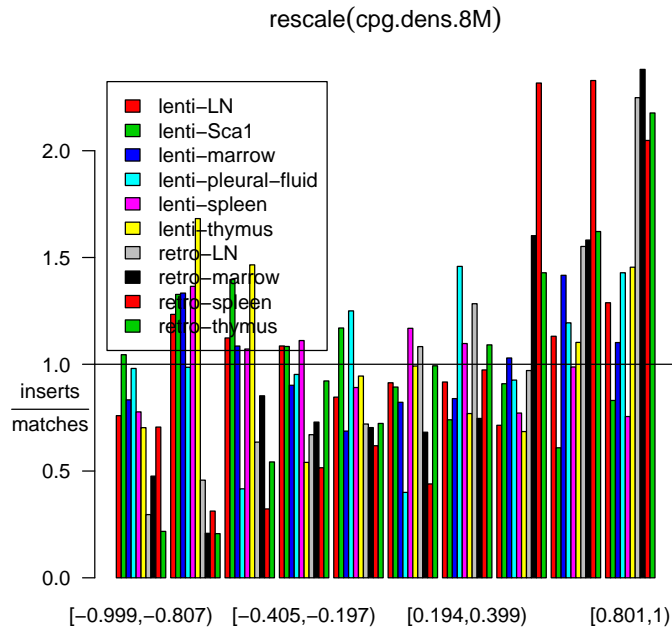

```
                        coef     se       z        p
lenti-LN             -0.0161 0.0826 -0.1950 8.45e-01
lenti-Sca1           -0.4190 0.0855 -4.9100 9.29e-07
lenti-marrow          0.0756 0.1860  0.4060 6.85e-01
lenti-pleural-fluid   0.1590 0.2820  0.5630 5.73e-01
lenti-spleen         -0.1030 0.1180 -0.8760 3.81e-01
lenti-thymus         -0.0158 0.1860 -0.0847 9.33e-01
retro-LN              0.9050 0.1250  7.2400 4.50e-13
retro-marrow          0.8890 0.1670  5.3300 1.00e-07
retro-spleen          1.1500 0.2490  4.6100 3.94e-06
retro-thymus          1.0200 0.1950  5.2400 1.63e-07
```
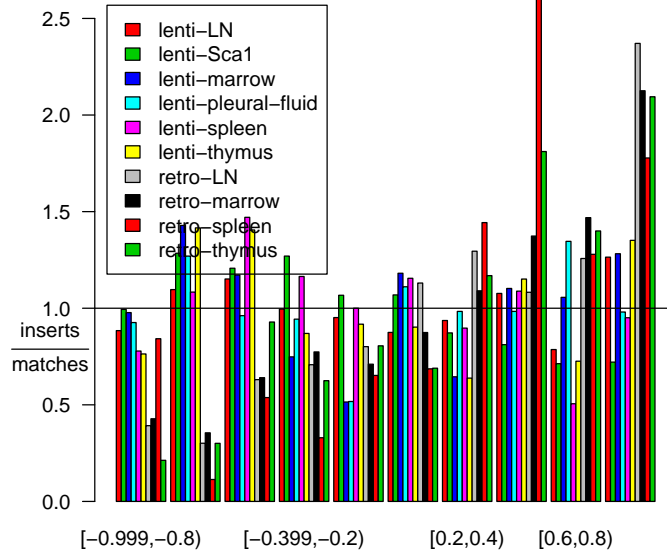
4.10   16 megabase Window

In the barplot that follows we examine the association of insertion sites with expression density in a 16 megabase window surrounding each locus. First, we count just the number of genes represented on the chip.

rescale(dens.16M)

	coef	se	z	p
lenti-LN	-0.0298	0.0827	-0.361	7.18e-01
lenti-Sca1	-0.3270	0.0845	-3.880	1.06e-04
lenti-marrow	0.1090	0.1850	0.589	5.56e-01
lenti-pleural-fluid	0.1500	0.2810	0.535	5.92e-01
lenti-spleen	-0.1920	0.1180	-1.620	1.05e-01
lenti-thymus	-0.1170	0.1870	-0.628	5.30e-01
retro-LN	0.9220	0.1270	7.280	3.29e-13
retro-marrow	0.8620	0.1690	5.100	3.36e-07
retro-spleen	1.1000	0.2500	4.390	1.14e-05
retro-thymus	0.8740	0.1900	4.590	4.43e-06

Here are the results for expression density. First, we count just genes that are in the upper half.

rescale(low.ex.16M)
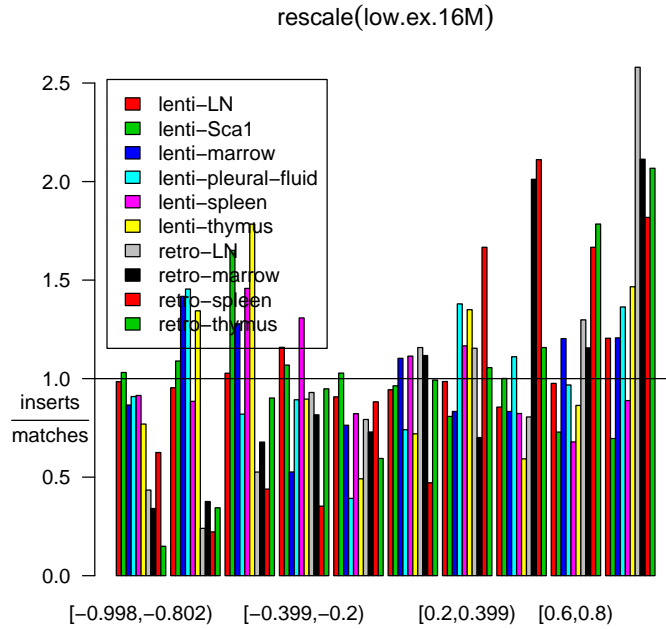
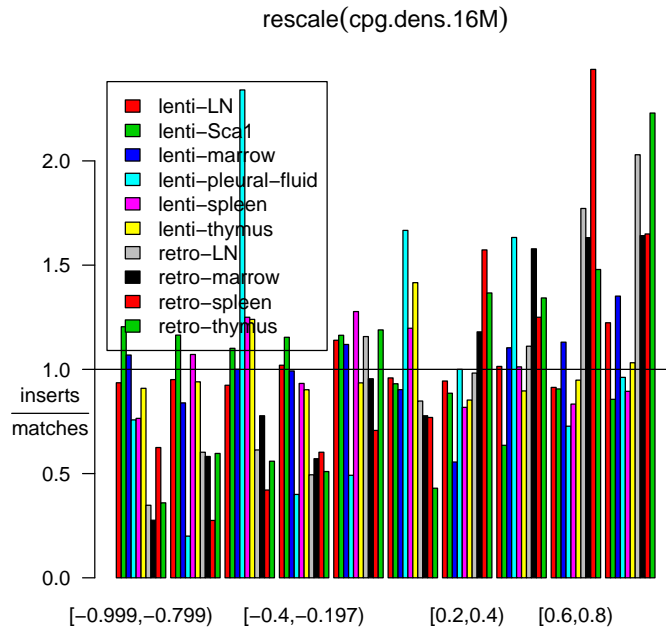|	coef	se	z	p
lenti-LN	-0.0112	0.0826	-0.135	8.92e-01
lenti-Sca1	-0.3500	0.0846	-4.130	3.60e-05
lenti-marrow	0.0715	0.1850	0.386	6.99e-01
lenti-pleural-fluid	0.2110	0.2830	0.743	4.57e-01
lenti-spleen	-0.1260	0.1180	-1.070	2.86e-01
lenti-thymus	-0.0825	0.1870	-0.442	6.59e-01
retro-LN	0.8670	0.1250	6.940	4.03e-12
retro-marrow	0.8630	0.1670	5.180	2.20e-07
retro-spleen	1.0900	0.2470	4.420	9.84e-06
retro-thymus	0.8900	0.1910	4.660	3.10e-06

Now we count genes in the upper $1/8^{th}$:

rescale(med.ex.16M)
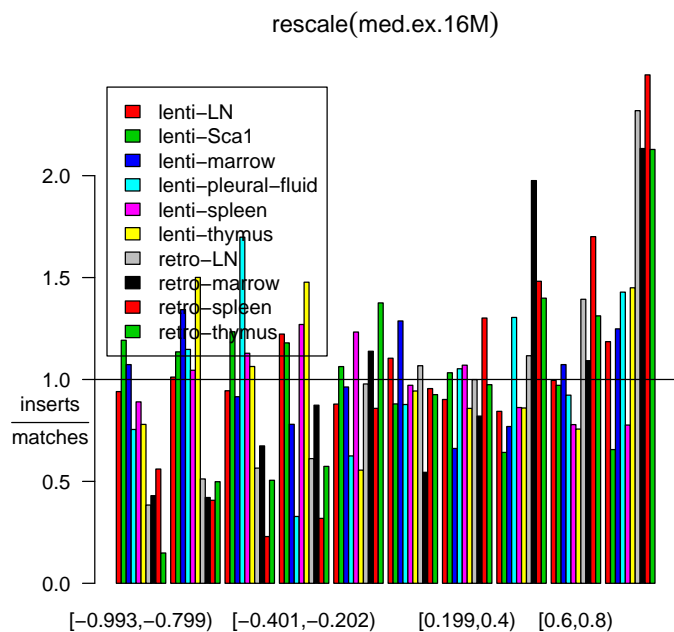
	coef	se	z	p
lenti-LN	-0.00062	0.0826	-0.0075	9.94e-01
lenti-Sca1	-0.32600	0.0846	-3.8600	1.14e-04
lenti-marrow	0.00934	0.1850	0.0505	9.60e-01
lenti-pleural-fluid	0.18800	0.2830	0.6660	5.06e-01
lenti-spleen	-0.20200	0.1180	-1.7100	8.76e-02
lenti-thymus	-0.05330	0.1860	-0.2870	7.74e-01
retro-LN	0.80700	0.1230	6.5500	5.75e-11
retro-marrow	0.59000	0.1590	3.7100	2.10e-04
retro-spleen	1.17000	0.2530	4.6000	4.18e-06
retro-thymus	0.78400	0.1880	4.1600	3.17e-05

And here we count genes in the upper $1/16^{th}$:

rescale(high.ex.16M)
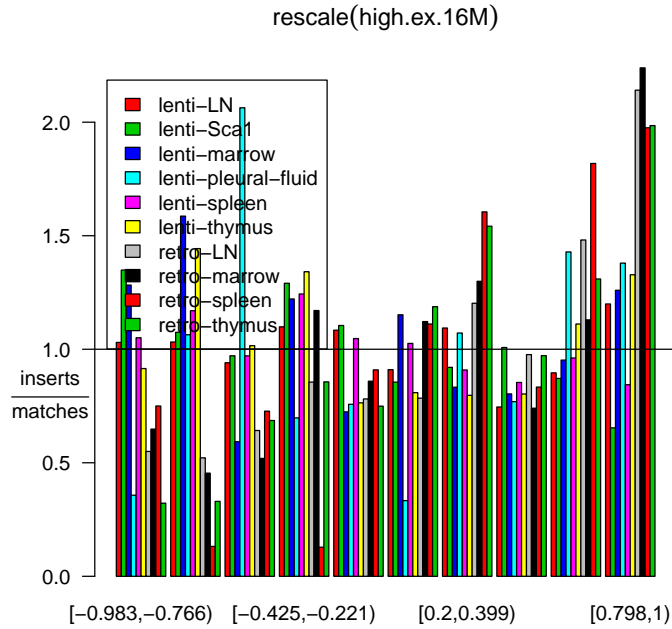
	coef	se	z	p
lenti-LN	-0.0463	0.0824	-0.5620	5.74e-01
lenti-Sca1	-0.2760	0.0847	-3.2600	1.12e-03
lenti-marrow	0.0155	0.1850	0.0840	9.33e-01
lenti-pleural-fluid	-0.0144	0.2820	-0.0512	9.59e-01
lenti-spleen	-0.1440	0.1180	-1.2200	2.23e-01
lenti-thymus	-0.0925	0.1870	-0.4940	6.21e-01
retro-LN	0.6910	0.1210	5.6900	1.23e-08
retro-marrow	0.6610	0.1600	4.1300	3.65e-05
retro-spleen	0.8810	0.2370	3.7100	2.04e-04
retro-thymus	0.8520	0.1920	4.4400	9.04e-06

Here the effect of density of CpG islands is studied:

rescale(cpg.dens.16M)

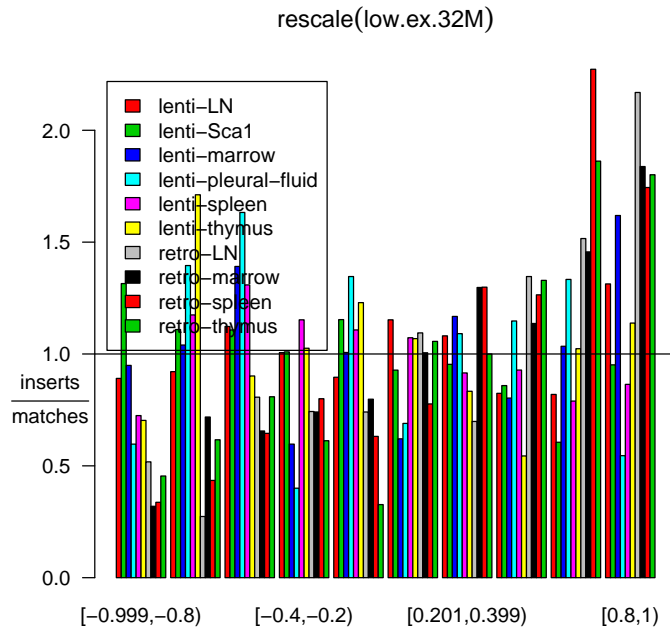|	coef	se	z	p
lenti-LN	0.01970	0.0823	0.2390	8.11e-01
lenti-Sca1	-0.33200	0.0850	-3.9100	9.33e-05
lenti-marrow	0.00618	0.1840	0.0335	9.73e-01
lenti-pleural-fluid	0.40300	0.2870	1.4000	1.61e-01
lenti-spleen	-0.08730	0.1180	-0.7390	4.60e-01
lenti-thymus	0.03420	0.1850	0.1850	8.53e-01
retro-LN	0.75200	0.1230	6.1400	8.49e-10
retro-marrow	0.77700	0.1640	4.7300	2.25e-06
retro-spleen	1.09000	0.2470	4.4200	9.81e-06
retro-thymus	0.78400	0.1880	4.1700	2.99e-05

4.11 32 megabase Window

In the barplot that follows we examine the association of insertion sites with expression density in a 32 megabase window surrounding each locus. First, we count just the number of genes represented on the chip.

rescale(dens.32M)
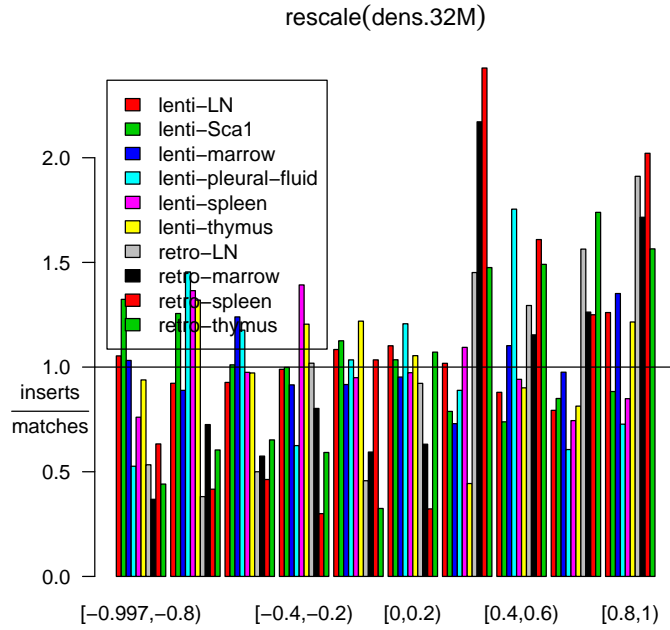
	coef	se	z	p
lenti-LN	0.0105	0.0823	0.1270	8.99e-01
lenti-Sca1	-0.2860	0.0846	-3.3700	7.39e-04
lenti-marrow	0.0154	0.1840	0.0836	9.33e-01
lenti-pleural-fluid	0.0655	0.2830	0.2310	8.17e-01
lenti-spleen	-0.1550	0.1180	-1.3100	1.91e-01
lenti-thymus	-0.2590	0.1870	-1.3900	1.66e-01
retro-LN	0.9110	0.1260	7.2400	4.40e-13
retro-marrow	0.8200	0.1660	4.9300	8.35e-07
retro-spleen	0.9700	0.2400	4.0400	5.30e-05
retro-thymus	1.0300	0.1950	5.2900	1.21e-07

Here are the results for expression density. First, we count just genes that are in the upper half.

rescale(low.ex.32M)

	coef	se	z	p
lenti-LN	0.0658	0.0823	0.800	4.24e-01
lenti-Sca1	-0.2730	0.0845	-3.230	1.23e-03
lenti-marrow	0.0337	0.1840	0.184	8.54e-01
lenti-pleural-fluid	-0.0513	0.2840	-0.181	8.57e-01
lenti-spleen	-0.1670	0.1190	-1.410	1.60e-01
lenti-thymus	-0.1880	0.1860	-1.010	3.12e-01
retro-LN	0.7920	0.1230	6.450	1.09e-10
retro-marrow	0.7190	0.1620	4.420	9.73e-06
retro-spleen	0.9110	0.2370	3.850	1.20e-04
retro-thymus	0.9310	0.1920	4.840	1.29e-06

Now we count genes in the upper $1/8^{th}$:

rescale(med.ex.32M)
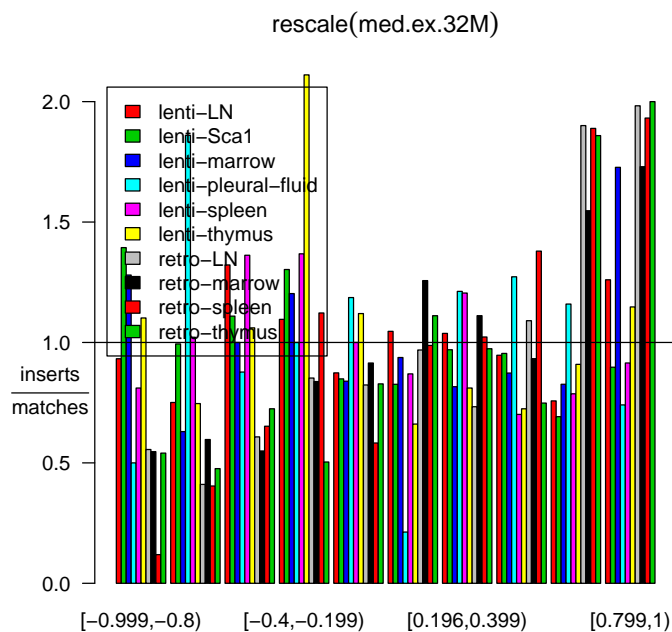
```
                          coef     se       z        p
lenti-LN              0.0166  0.0822   0.202  8.40e-01
lenti-Sca1           -0.2590  0.0845  -3.060  2.20e-03
lenti-marrow          0.0245  0.1840   0.134  8.94e-01
lenti-pleural-fluid  -0.0800  0.2830  -0.283  7.77e-01
lenti-spleen         -0.2120  0.1190  -1.780  7.50e-02
lenti-thymus         -0.3240  0.1890  -1.720  8.58e-02
retro-LN              0.7280  0.1220   5.960  2.58e-09
retro-marrow          0.6460  0.1620   4.000  6.40e-05
retro-spleen          0.8840  0.2340   3.770  1.61e-04
retro-thymus          0.7860  0.1880   4.170  3.01e-05
```

And here we count genes in the upper $1/16^{th}$:

rescale(high.ex.32M)
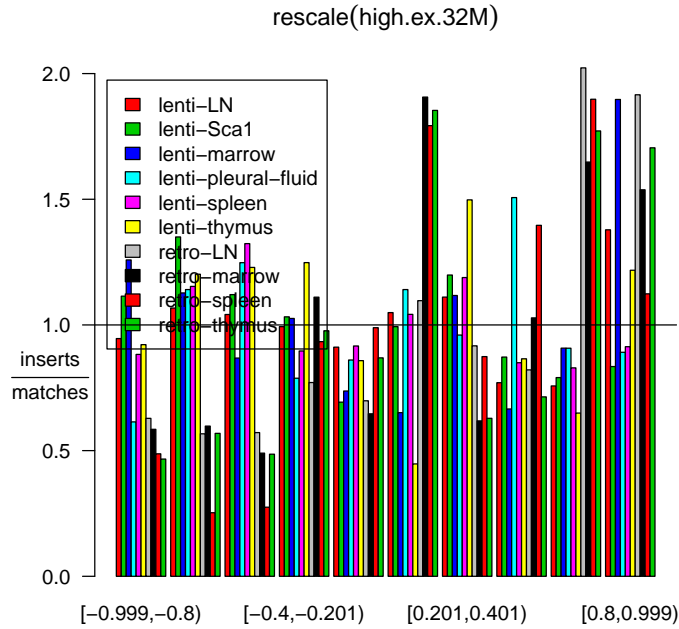
	coef	se	z	p
lenti-LN	0.0130	0.0824	0.157	8.75e-01
lenti-Sca1	-0.1160	0.0839	-1.380	1.66e-01
lenti-marrow	0.0185	0.1840	0.101	9.20e-01
lenti-pleural-fluid	0.1680	0.2920	0.578	5.63e-01
lenti-spleen	-0.0799	0.1180	-0.676	4.99e-01
lenti-thymus	-0.1660	0.1880	-0.885	3.76e-01
retro-LN	0.7320	0.1220	6.000	2.00e-09
retro-marrow	0.6620	0.1590	4.170	3.06e-05
retro-spleen	0.8420	0.2320	3.620	2.90e-04
retro-thymus	0.6780	0.1860	3.650	2.62e-04

Here the effect of density of CpG islands is studied:

rescale(cpg.dens.32M)
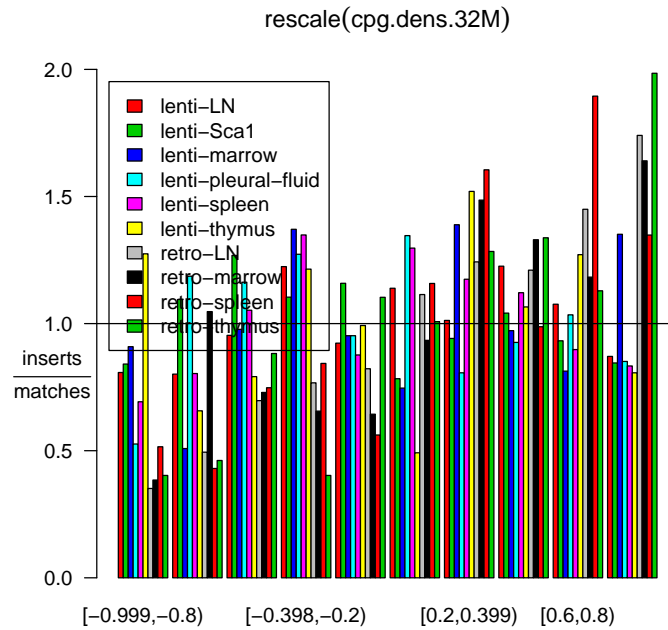

```
                          coef     se       z        p
lenti-LN               0.1310 0.0824   1.5900 1.11e-01
lenti-Sca1            -0.1860 0.0841  -2.2100 2.69e-02
lenti-marrow           0.1090 0.1860   0.5890 5.56e-01
lenti-pleural-fluid   -0.0145 0.2830  -0.0514 9.59e-01
lenti-spleen           0.1140 0.1180   0.9640 3.35e-01
lenti-thymus           0.0589 0.1850   0.3190 7.50e-01
retro-LN               0.7890 0.1240   6.3600 2.05e-10
retro-marrow           0.6380 0.1590   4.0200 5.87e-05
retro-spleen           0.7610 0.2300   3.3100 9.38e-04
retro-thymus           0.7170 0.1850   3.8700 1.08e-04
```

5 Juxtaposition with Gene Start and End Positions

5.1 Refseq Annotations

In this section we study the effect of juxtaposition in terms of gene start and end positions. The first barplot shows the effect of gene width for those insertions that are located within an RefSeq gene. The table following the barplot shows the p-values for a test of the hypothesis that the proportions in each of the categories that define the bars are equal in the insertions and their matches. This p-value is obtained from the $5 \times 2 \times k$ table of counts defined by gene width category, insertion/match status, and stratum (consisting of an insertion and its matched sites) using a likelihood ratio test for the hypothesis of no association between gene width category and insertion/match status. The test used compared the log-linear model [1] with all two-way configurations to that with no gene width category and insertion/match status configuration.

refSeq gene.width
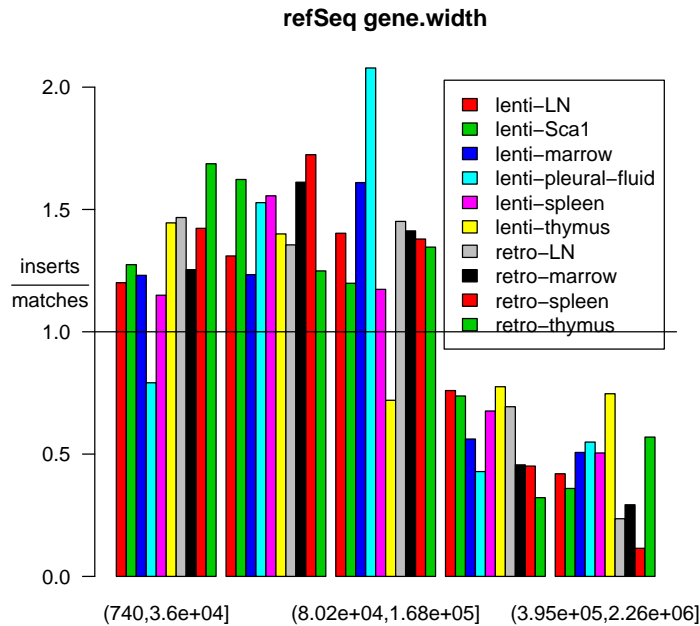
lenti-LN	lenti-Sca1	lenti-marrow	lenti-pleural-fluid
1.08e-11	9.32e-13	7.65e-03	1.94e-02
lenti-spleen	lenti-thymus	retro-LN	retro-marrow
3.51e-06	2.11e-01	3.65e-09	1.47e-05
retro-spleen	retro-thymus		
1.73e-04	7.00e-03		

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The next plot uses the width of a non-gene region for insertions that fall into such regions.

refSeq other.width
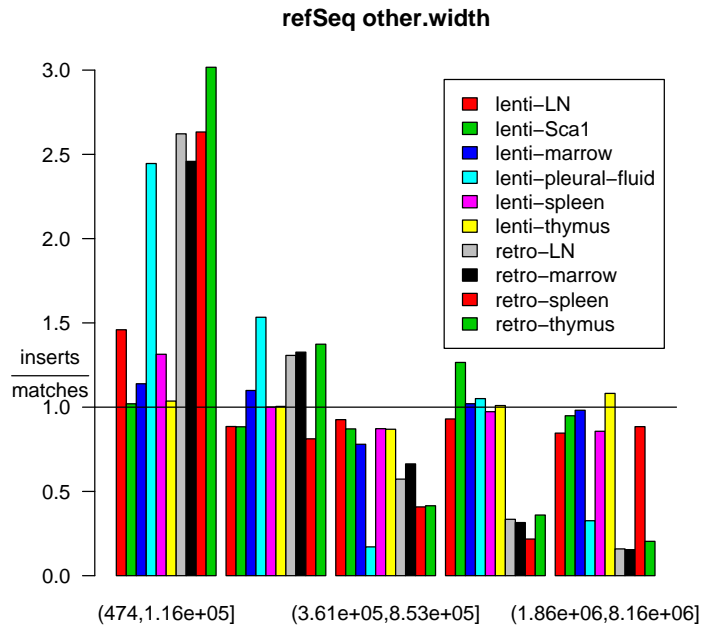
lenti-LN	lenti-Sca1	lenti-marrow	lenti-pleural-fluid
1.36e-02	1.94e-01	8.20e-01	2.12e-03
lenti-spleen	lenti-thymus	retro-LN	retro-marrow
5.31e-01	8.46e-01	2.17e-29	1.73e-14
retro-spleen	retro-thymus		
9.63e-07	1.07e-15		

The next plot studies the distance to the nearest boundary between a gene and a non-gene region. The distance is expressed as a fraction of the length of the region. Thus, '0.25' refers to one quarter of the distance from the site to nearest boundary divided by the total width of the region.

refSeq boundary.dist
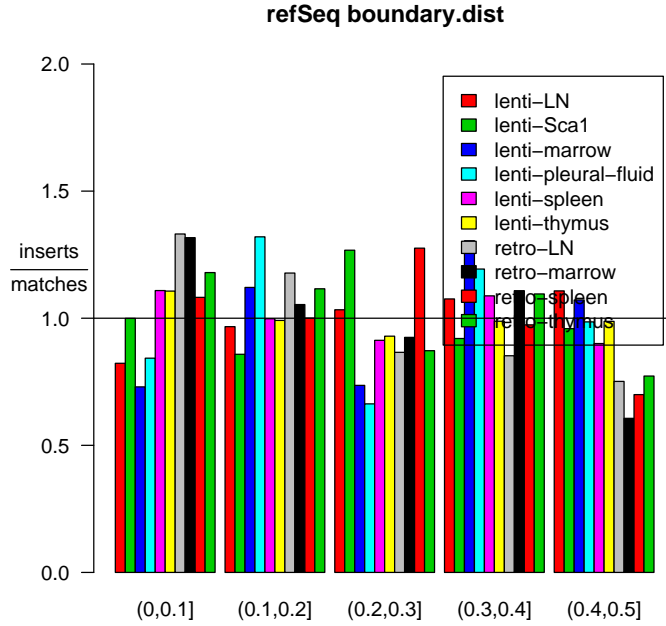
	lenti-LN	lenti-Sca1	lenti-marrow	lenti-pleural-fluid
	0.13900	0.01840	0.18300	0.56300
lenti-spleen	lenti-thymus	retro-LN	retro-marrow
	0.67700	0.98200	0.00282	0.03220
retro-spleen	retro-thymus		
	0.53400	0.45500		

This plot studies the effect of nearness to the beginning of a transcript. For sites in genes, it is the distance to the start of the gene divided by the width of the gene. For other sites it is the distance from the site to the nearer gene if that gene boundary is also a transcription starting point. Locations near '0' are relatively near the beginning of transcription, while those near '1' are near the termination of the transcript.
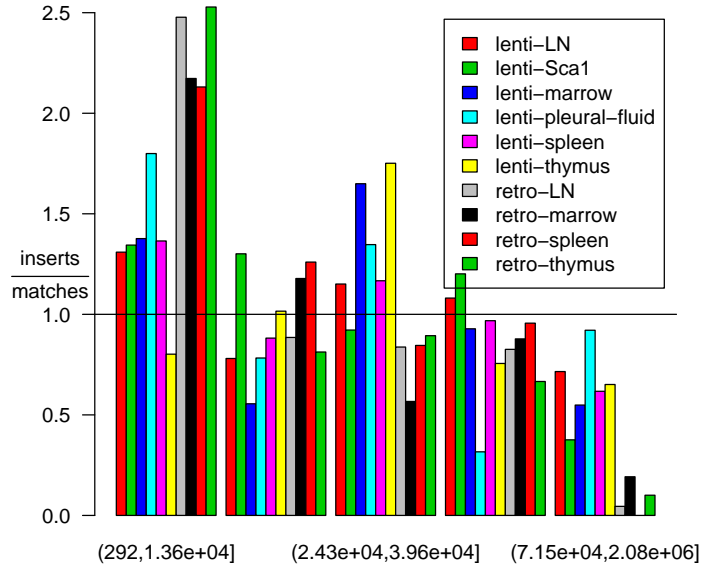
refSeq start.dist
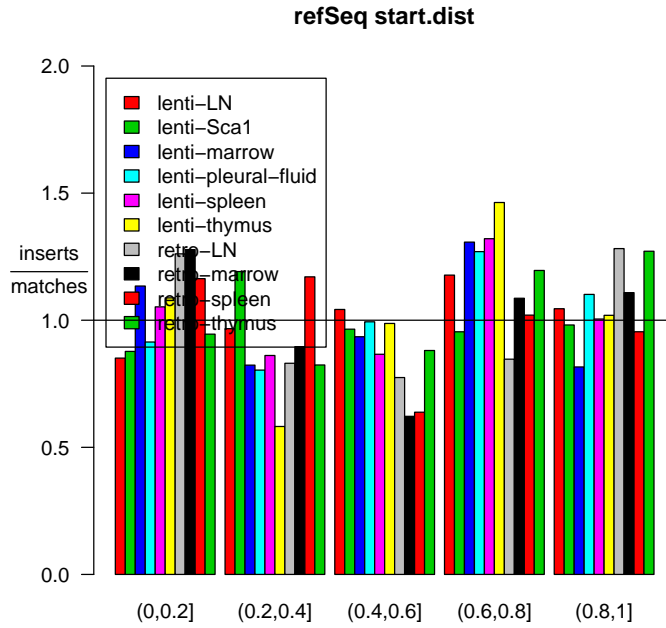
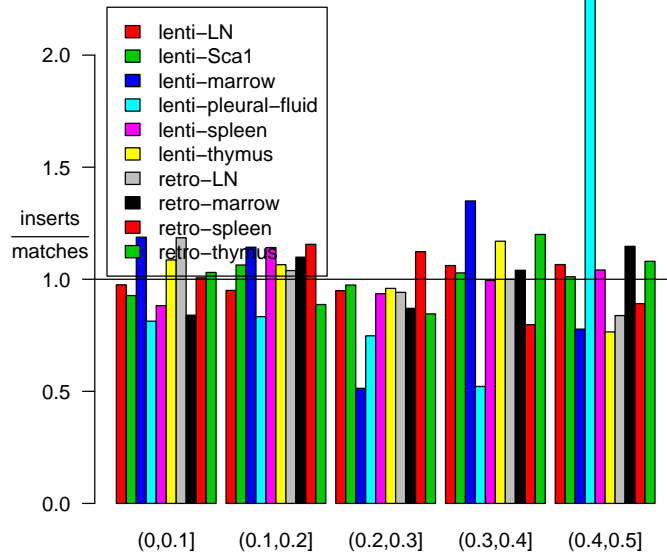|	lenti-LN	lenti-Sca1	lenti-marrow	lenti-pleural-fluid
	0.1130	0.1320	0.6990	0.9320
	lenti-spleen	lenti-thymus	retro-LN	retro-marrow
	0.1750	0.1240	0.0041	0.0706
	retro-spleen	retro-thymus		
	0.4720	0.5430		

5.2   genScan Annotations

genScan gene.width
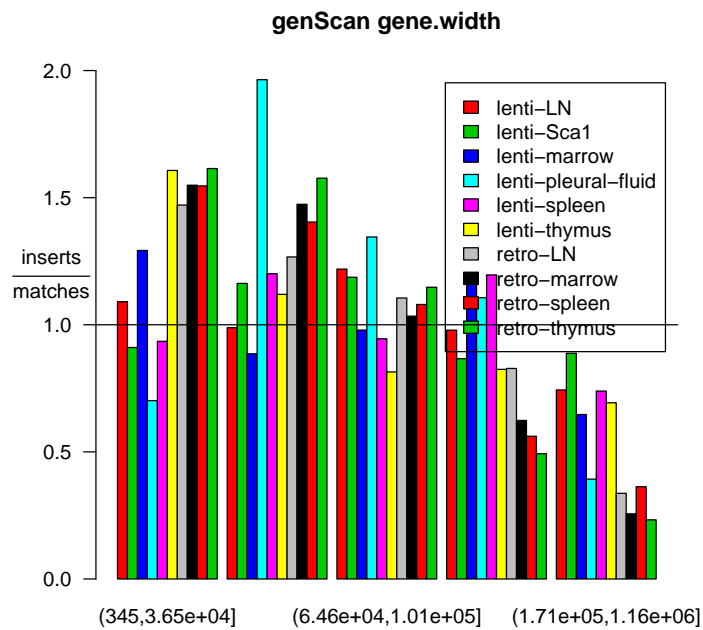
lenti-LN	lenti-Sca1	lenti-marrow	lenti-pleural-fluid
3.05e-02	7.71e-02	3.73e-01	7.07e-02
lenti-spleen	lenti-thymus	retro-LN	retro-marrow
9.22e-02	6.39e-02	3.47e-08	4.15e-07
retro-spleen	retro-thymus		
7.83e-03	1.28e-06		

genScan other.width

	lenti-LN	lenti-Sca1	lenti-marrow	lenti-pleural-fluid
	4.97e-03	1.13e-05	3.57e-02	2.39e-01
	lenti-spleen	lenti-thymus	retro-LN	retro-marrow
	1.38e-01	5.03e-02	8.03e-13	5.80e-07
	retro-spleen	retro-thymus		
	2.10e-04	5.37e-07		

genScan boundary.dist

lenti-LN	lenti-Sca1	lenti-marrow	lenti-pleural-fluid
0.78600	0.84800	0.01020	0.00247
lenti-spleen	lenti-thymus	retro-LN	retro-marrow
0.63300	0.63000	0.37200	0.59900
retro-spleen	retro-thymus		
0.79300	0.64400		

genScan start.dist
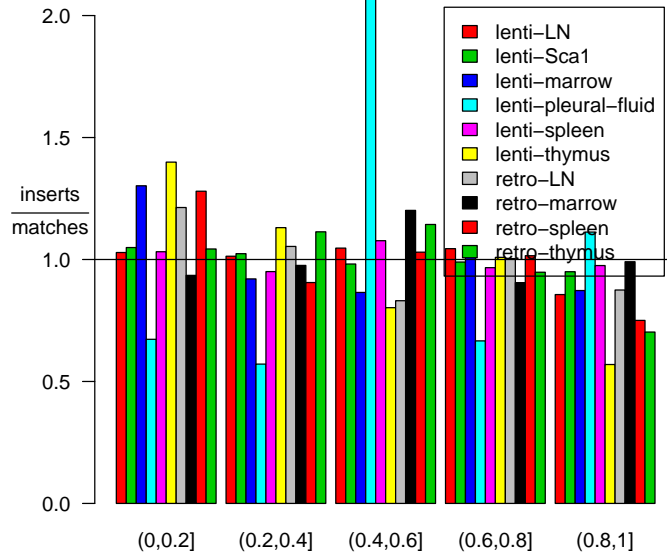
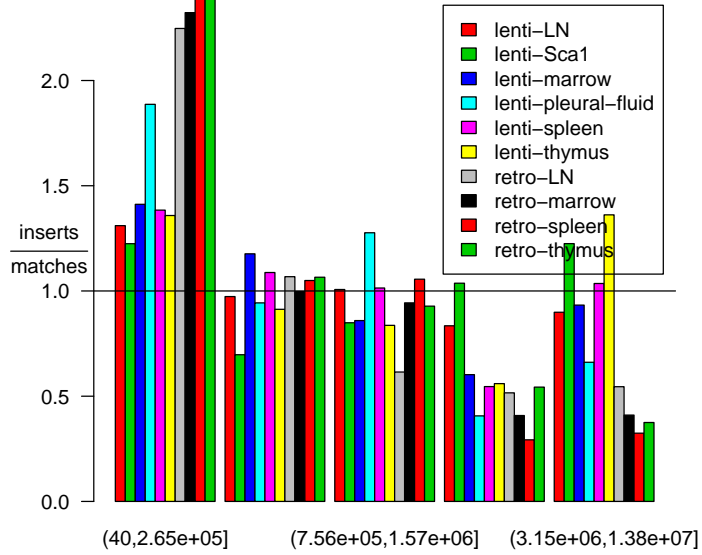| lenti-LN	lenti-Sca1	lenti-marrow	lenti-pleural-fluid
0.5290	0.9500	0.4860	0.0165
lenti-spleen	lenti-thymus	retro-LN	retro-marrow
0.9430	0.0522	0.1790	0.7820
retro-spleen	retro-thymus		
0.7460	0.5070		

6 Oncogenes

This plot studies the effect of nearness to the 5' end of an oncogene transcript. Positive values represent distances in which the integration site is upstream of the nearest oncogene 5' end, negative downstream.

onco.dist
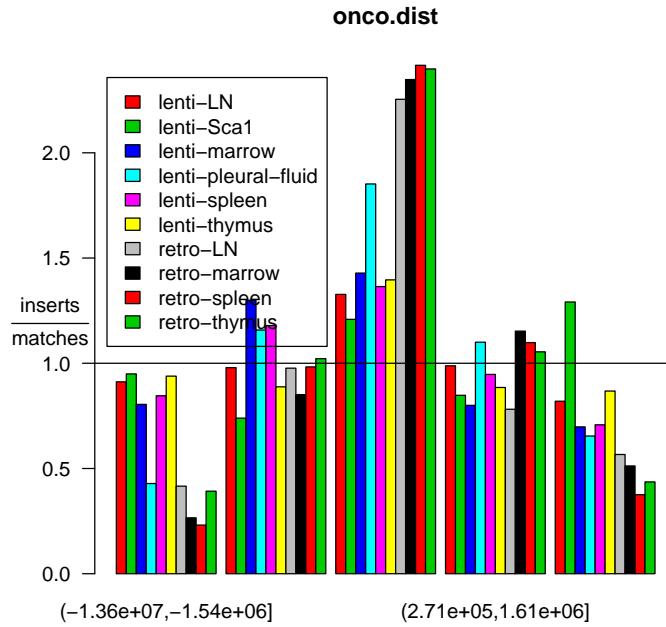
	lenti-LN	lenti-Sca1	lenti-marrow	lenti-pleural-fluid
	1.38e-03	1.81e-05	2.66e-02	9.04e-03
	lenti-spleen	lenti-thymus	retro-LN	retro-marrow
	2.05e-03	2.99e-01	6.41e-26	6.15e-19
	retro-spleen	retro-thymus		
	1.14e-10	9.43e-13		

Here is the same plot using absolute distance

abs abs.onco.dist

	lenti-LN	lenti-Sca1	lenti-marrow	lenti-pleural-fluid
	2.94e-03	1.72e-05	4.37e-02	6.78e-03
	lenti-spleen	lenti-thymus	retro-LN	retro-marrow
	4.83e-05	1.29e-02	8.82e-26	8.81e-17
	retro-spleen	retro-thymus		
	1.12e-10	4.37e-12		

7 GC content

Here we study the effect of GC content on insertion. The GC content is taken from the Mouse Genome Draft at GoldenPath from the table
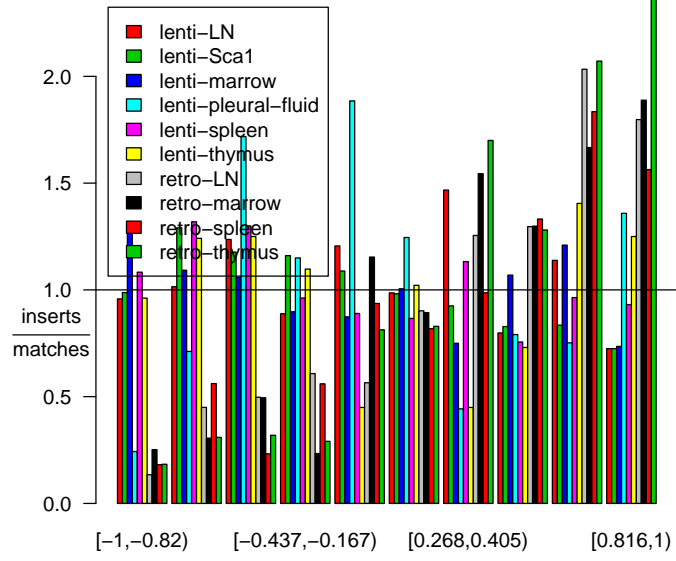
Following the plot is a table of fitted coefficients based on splitting the GC percent data at the median.

rescale(gc20)
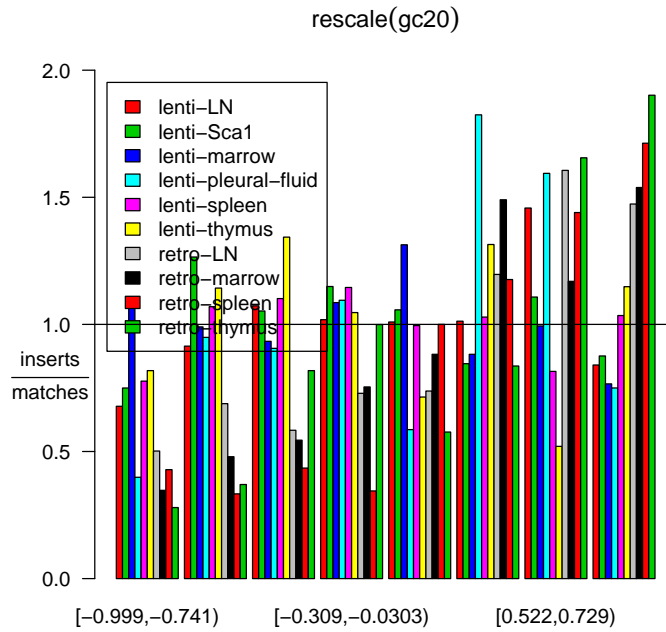
```
                          coef      se       z         p
lenti-LN               0.11100  0.0830   1.3400  1.79e-01
lenti-Sca1            -0.12300  0.0856  -1.4400  1.51e-01
lenti-marrow         -0.24300  0.1900  -1.2800  2.00e-01
lenti-pleural-fluid   0.48300  0.2840   1.7000  8.90e-02
lenti-spleen         -0.06170  0.1210  -0.5100  6.10e-01
lenti-thymus          0.00985  0.1900   0.0518  9.59e-01
retro-LN              0.77500  0.1200   6.4800  8.92e-11
retro-marrow          0.78900  0.1610   4.8900  1.03e-06
retro-spleen          0.98100  0.2420   4.0500  5.17e-05
retro-thymus          0.82100  0.1820   4.5100  6.43e-06

                          coef      se       z         p
lenti-LN              -0.04140  0.0831  -0.4980  6.19e-01
lenti-Sca1            -0.31100  0.0859  -3.6300  2.87e-04
lenti-marrow         -0.09020  0.1850  -0.4870  6.27e-01
lenti-pleural-fluid  -0.00655  0.2830  -0.0232  9.82e-01
lenti-spleen         -0.13600  0.1200  -1.1300  2.60e-01
lenti-thymus          0.01300  0.1890   0.0687  9.45e-01
retro-LN              0.96500  0.1260   7.6600  1.89e-14
retro-marrow          0.93500  0.1690   5.5400  3.03e-08
retro-spleen          0.88200  0.2470   3.5700  3.63e-04
retro-thymus          1.21000  0.1960   6.1600  7.48e-10
```

rescale(gc50)

Legend:
- lenti–LN
- lenti–Sca1
- lenti–marrow
- lenti–pleural–fluid
- lenti–spleen
- lenti–thymus
- retro–LN
- retro–marrow
- retro–spleen
- retro–thymus

x-axis: [−1,−0.82) [−0.437,−0.167) [0.268,0.405) [0.816,1)

y-axis: $\frac{\text{inserts}}{\text{matches}}$

```
                         coef     se      z       p
lenti-LN              0.00894 0.0838  0.107 9.15e-01
lenti-Sca1           -0.23700 0.0856 -2.770 5.69e-03
lenti-marrow         -0.18500 0.1880 -0.983 3.26e-01
lenti-pleural-fluid   0.08250 0.2870  0.288 7.74e-01
lenti-spleen         -0.21500 0.1220 -1.760 7.84e-02
lenti-thymus          0.04960 0.1910  0.260 7.95e-01
retro-LN              1.22000 0.1340  9.100 9.38e-20
retro-marrow          1.06000 0.1750  6.090 1.15e-09
retro-spleen          0.91200 0.2520  3.620 2.96e-04
retro-thymus          1.24000 0.1980  6.260 3.86e-10
```


rescale(gc100)
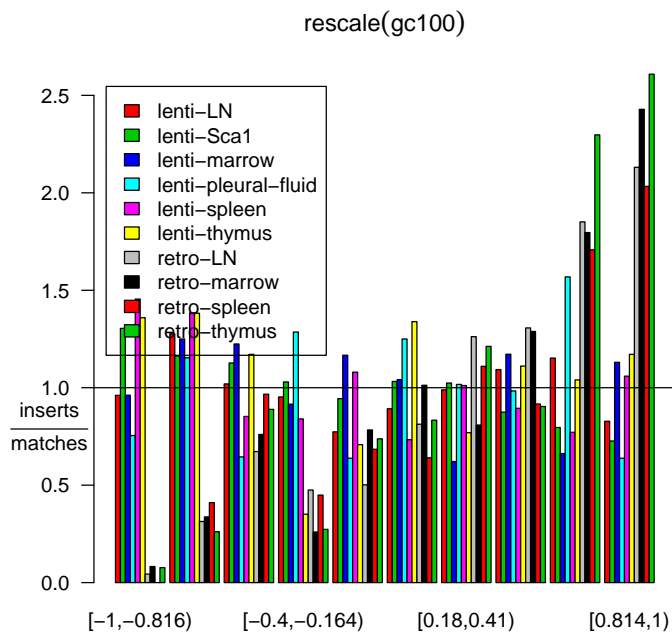
```
                      coef      se       z        p
lenti-LN           -0.0769  0.0834  -0.922  3.57e-01
lenti-Sca1         -0.3260  0.0859  -3.800  1.47e-04
lenti-marrow       -0.2180  0.1880  -1.160  2.48e-01
lenti-pleural-fluid -0.1040  0.2870  -0.363  7.17e-01
lenti-spleen       -0.2610  0.1210  -2.150  3.13e-02
lenti-thymus       -0.2790  0.1930  -1.450  1.47e-01
retro-LN            1.5400  0.1450  10.600  2.89e-26
retro-marrow        1.4200  0.1920   7.400  1.39e-13
retro-spleen        1.2500  0.2640   4.730  2.27e-06
retro-thymus        1.4200  0.2080   6.820  9.12e-12
```

rescale(gc250)

90
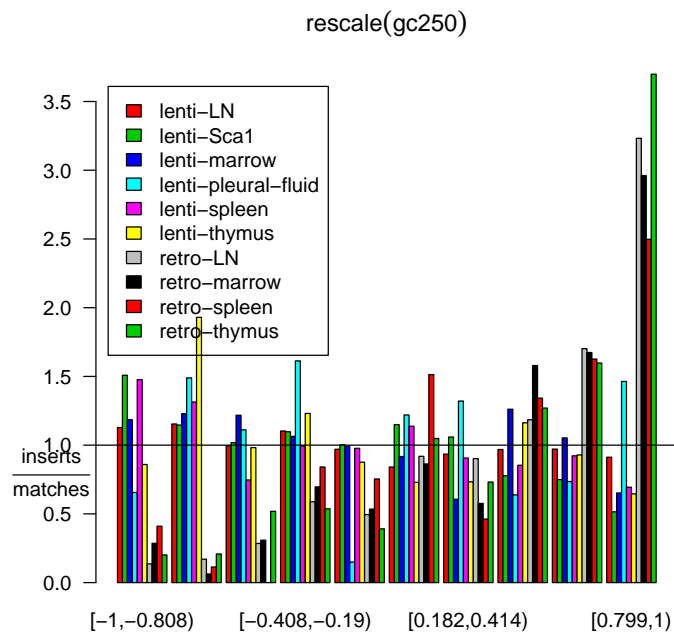
```
                        coef      se       z        p
lenti-LN             -0.1470  0.0832  -1.770  7.61e-02
lenti-Sca1           -0.5380  0.0866  -6.210  5.25e-10
lenti-marrow         -0.0343  0.1850  -0.185  8.53e-01
lenti-pleural-fluid  -0.3790  0.2920  -1.300  1.95e-01
lenti-spleen         -0.1650  0.1190  -1.390  1.66e-01
lenti-thymus         -0.1670  0.1920  -0.870  3.84e-01
retro-LN              1.6300  0.1550  10.600  4.30e-26
retro-marrow          1.8200  0.2200   8.260  1.47e-16
retro-spleen          1.4300  0.2830   5.050  4.41e-07
retro-thymus          1.8400  0.2410   7.610  2.66e-14
```

rescale(gc500)

91
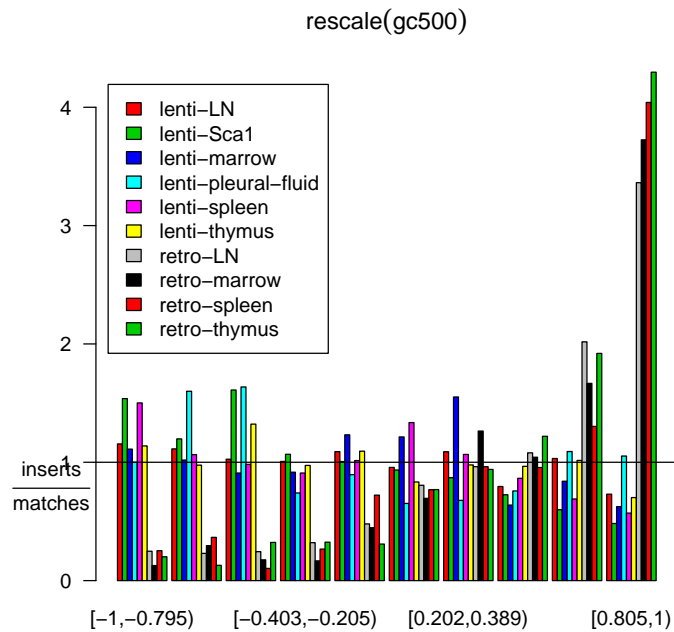
```
                     coef     se       z        p
lenti-LN            -0.476  0.0858  -5.550  2.83e-08
lenti-Sca1          -0.787  0.0893  -8.820  1.17e-18
lenti-marrow        -0.615  0.1950  -3.150  1.64e-03
lenti-pleural-fluid -0.289  0.2920  -0.991  3.22e-01
lenti-spleen        -0.464  0.1220  -3.800  1.47e-04
lenti-thymus        -0.452  0.1940  -2.330  1.98e-02
retro-LN             1.630  0.1550  10.500  8.14e-26
retro-marrow         1.920  0.2290   8.400  4.39e-17
retro-spleen         1.500  0.2940   5.100  3.44e-07
retro-thymus         1.730  0.2300   7.530  4.91e-14
```


rescale(gc1000)
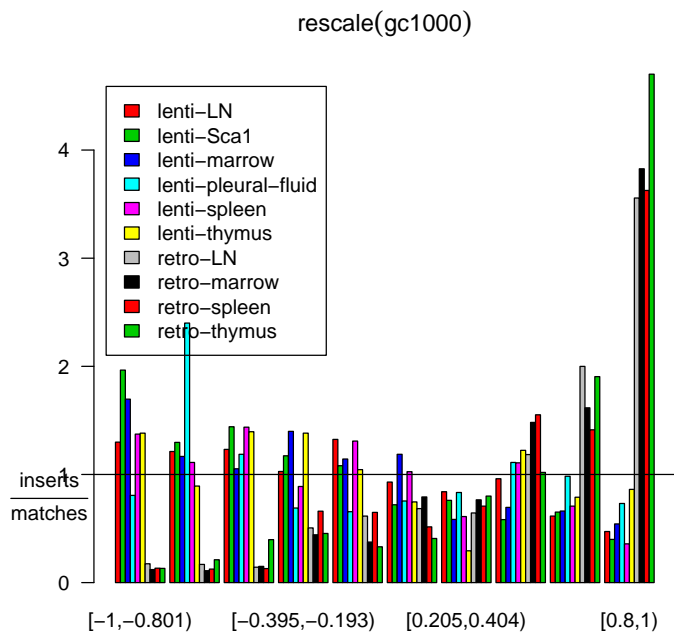
```
                     coef      se       z         p
lenti-LN            -0.502  0.0861   -5.83  5.48e-09
lenti-Sca1          -1.000  0.0934  -10.80  5.61e-27
lenti-marrow        -0.706  0.1960   -3.61  3.08e-04
lenti-pleural-fluid -0.423  0.2950   -1.43  1.52e-01
lenti-spleen        -0.471  0.1220   -3.86  1.14e-04
lenti-thymus        -0.458  0.1930   -2.37  1.79e-02
retro-LN             1.760  0.1630   10.80  2.88e-27
retro-marrow         2.210  0.2540    8.68  4.06e-18
retro-spleen         1.740  0.3170    5.47  4.42e-08
retro-thymus         1.990  0.2540    7.84  4.62e-15
```


rescale(gc2000)
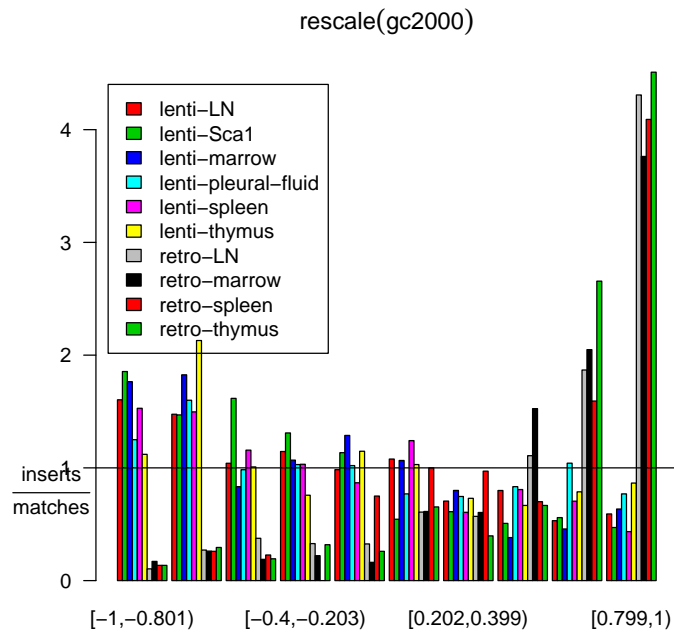
```
                     coef     se      z        p
lenti-LN           -0.429  0.0853  -5.030  4.93e-07
lenti-Sca1         -0.855  0.0915  -9.350  8.52e-21
lenti-marrow       -0.745  0.1950  -3.820  1.34e-04
lenti-pleural-fluid -0.103 0.2850  -0.361  7.18e-01
lenti-spleen       -0.479  0.1230  -3.910  9.10e-05
lenti-thymus       -0.317  0.1920  -1.660  9.77e-02
retro-LN            2.010  0.1780  11.200  2.58e-29
retro-marrow        2.200  0.2530   8.690  3.62e-18
retro-spleen        1.770  0.3180   5.580  2.34e-08
retro-thymus        2.150  0.2700   7.940  1.98e-15
```

rescale(gc5000)
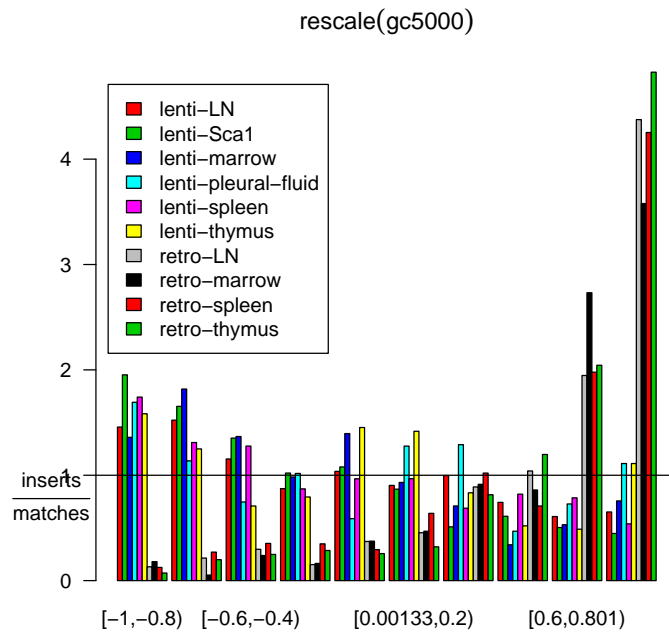
```
                       coef     se       z        p
lenti-LN              -0.446  0.0853  -5.230  1.69e-07
lenti-Sca1            -0.711  0.0887  -8.010  1.13e-15
lenti-marrow          -0.731  0.1950  -3.760  1.73e-04
lenti-pleural-fluid   -0.110  0.2850  -0.387  6.99e-01
lenti-spleen          -0.490  0.1220  -4.020  5.84e-05
lenti-thymus          -0.478  0.1920  -2.490  1.29e-02
retro-LN               2.170  0.1880  11.500  8.34e-31
retro-marrow           1.920  0.2290   8.380  5.11e-17
retro-spleen           1.780  0.3180   5.610  1.99e-08
retro-thymus           2.420  0.2990   8.090  5.86e-16
```


rescale(gc10000)
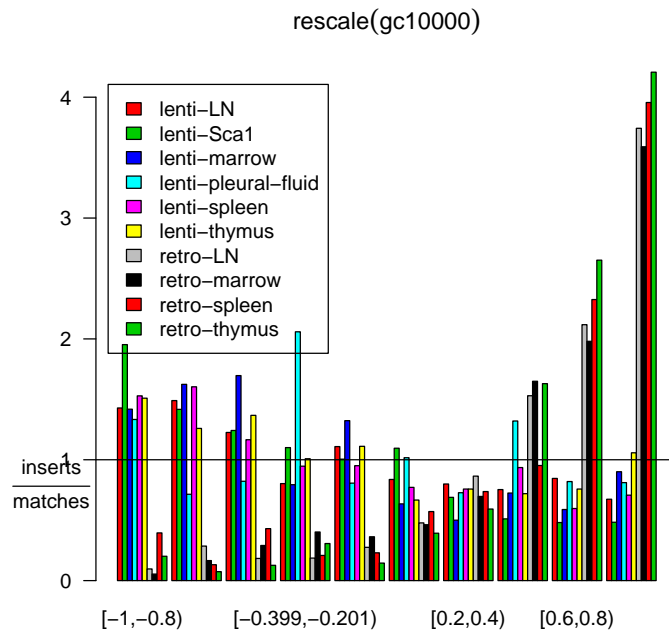
```
                      coef      se       z         p
lenti-LN            -0.342  0.0839  -4.080  4.51e-05
lenti-Sca1          -0.691  0.0885  -7.810  5.55e-15
lenti-marrow        -0.545  0.1890  -2.880  3.98e-03
lenti-pleural-fluid  0.102  0.2830   0.361  7.18e-01
lenti-spleen        -0.394  0.1210  -3.270  1.09e-03
lenti-thymus        -0.383  0.1900  -2.020  4.35e-02
retro-LN             2.110  0.1830  11.500  8.90e-31
retro-marrow         1.890  0.2250   8.400  4.35e-17
retro-spleen         1.710  0.3060   5.580  2.42e-08
retro-thymus         2.230  0.2720   8.180  2.95e-16
```

rescale(gc25000)
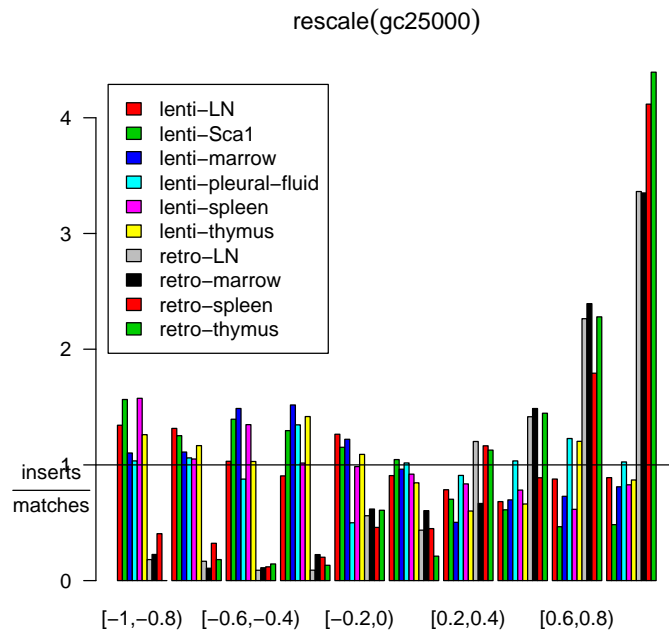
```
                     coef      se       z        p
lenti-LN           -0.2810  0.0838  -3.360  7.93e-04
lenti-Sca1         -0.6580  0.0880  -7.480  7.59e-14
lenti-marrow       -0.4350  0.1870  -2.330  1.99e-02
lenti-pleural-fluid 0.0862  0.2810   0.307  7.59e-01
lenti-spleen       -0.3140  0.1200  -2.630  8.61e-03
lenti-thymus       -0.3440  0.1890  -1.820  6.90e-02
retro-LN            2.1200  0.1850  11.500  1.55e-30
retro-marrow        1.7500  0.2130   8.190  2.67e-16
retro-spleen        1.7200  0.3080   5.590  2.23e-08
retro-thymus        2.3600  0.2890   8.160  3.23e-16
```

rescale(gc50000)
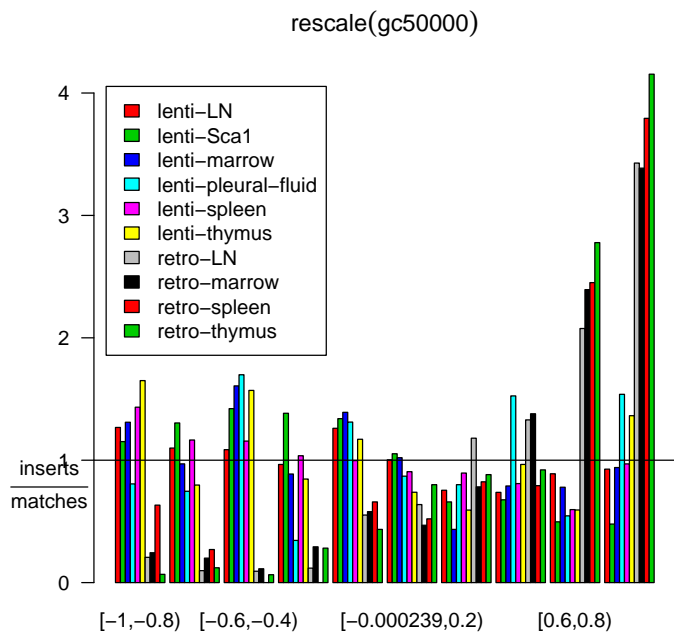
```
                     coef     se       z        p
lenti-LN            -0.170  0.0831  -2.050  4.03e-02
lenti-Sca1          -0.489  0.0860  -5.690  1.26e-08
lenti-marrow        -0.194  0.1840  -1.050  2.93e-01
lenti-pleural-fluid  0.187  0.2810   0.667  5.05e-01
lenti-spleen        -0.148  0.1180  -1.250  2.10e-01
lenti-thymus        -0.196  0.1880  -1.040  2.98e-01
retro-LN             2.060  0.1800  11.400  2.55e-30
retro-marrow         1.810  0.2170   8.320  8.60e-17
retro-spleen         1.720  0.3070   5.590  2.27e-08
retro-thymus         2.160  0.2650   8.170  2.97e-16
```

rescale(gc100000)
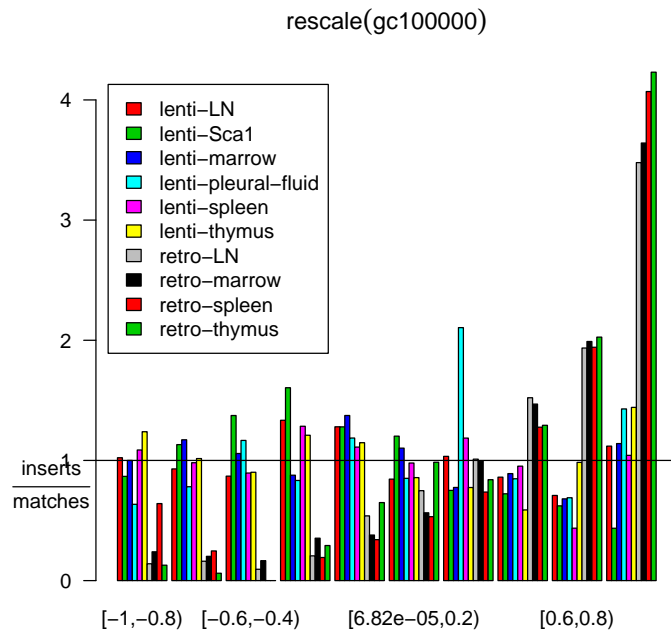
```
                          coef       se       z         p
lenti-LN                0.0532   0.0825   0.645  5.19e-01
lenti-Sca1             -0.3920   0.0851  -4.610  4.07e-06
lenti-marrow            0.2400   0.1850   1.300  1.94e-01
lenti-pleural-fluid     0.4740   0.2830   1.680  9.38e-02
lenti-spleen            0.0487   0.1170   0.415  6.78e-01
lenti-thymus            0.1340   0.1870   0.714  4.75e-01
retro-LN                2.0600   0.1780  11.600  4.14e-31
retro-marrow            1.7900   0.2160   8.290  1.10e-16
retro-spleen            1.8100   0.3160   5.740  9.22e-09
retro-thymus            2.3700   0.2880   8.220  2.02e-16
```


rescale(gc250000)
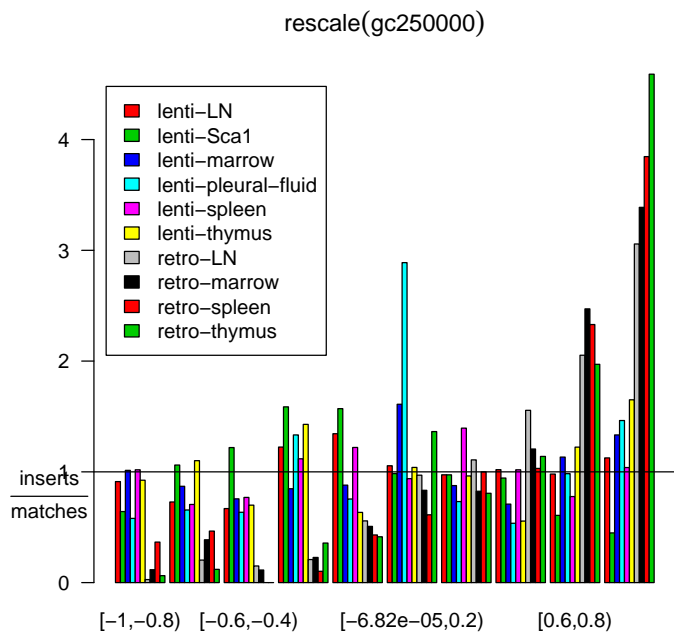
```
                     coef     se      z         p
lenti-LN            0.130  0.0824   1.570  1.16e-01
lenti-Sca1         -0.303  0.0842  -3.600  3.19e-04
lenti-marrow        0.229  0.1850   1.230  2.17e-01
lenti-pleural-fluid 0.570  0.2850   2.000  4.53e-02
lenti-spleen        0.153  0.1180   1.300  1.93e-01
lenti-thymus        0.149  0.1890   0.787  4.31e-01
retro-LN            1.810  0.1630  11.100  1.15e-28
retro-marrow       1.640  0.2070   7.940  1.95e-15
retro-spleen        1.760  0.3060   5.740  9.53e-09
retro-thymus        2.090  0.2590   8.070  6.95e-16
```


rescale(gc500000)

100
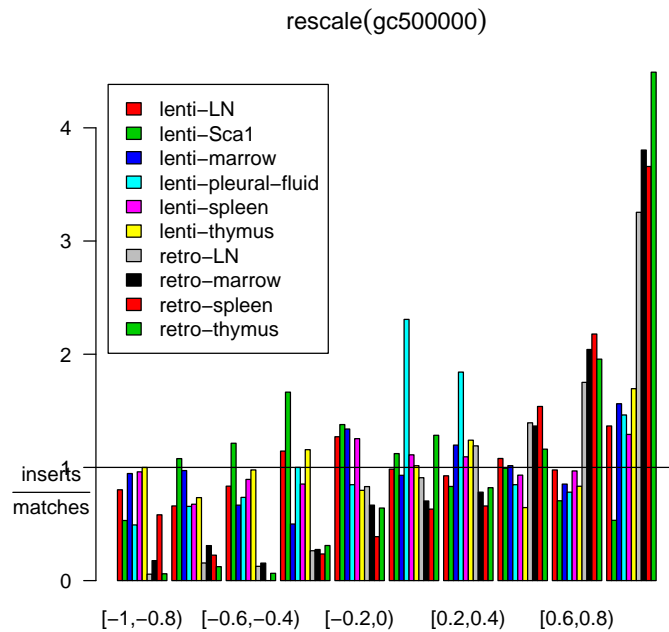
```
                      coef     se     z         p
lenti-LN             0.141 0.0825  1.71 8.73e-02
lenti-Sca1          -0.328 0.0847 -3.88 1.06e-04
lenti-marrow         0.272 0.1860  1.47 1.42e-01
lenti-pleural-fluid  0.602 0.2870  2.10 3.59e-02
lenti-spleen         0.139 0.1170  1.18 2.37e-01
lenti-thymus         0.238 0.1900  1.25 2.11e-01
retro-LN             1.910 0.1690 11.30 1.42e-29
retro-marrow         1.570 0.2010  7.81 5.52e-15
retro-spleen         1.780 0.3070  5.80 6.60e-09
retro-thymus         2.030 0.2540  7.96 1.66e-15
```


rescale(gc1000000)
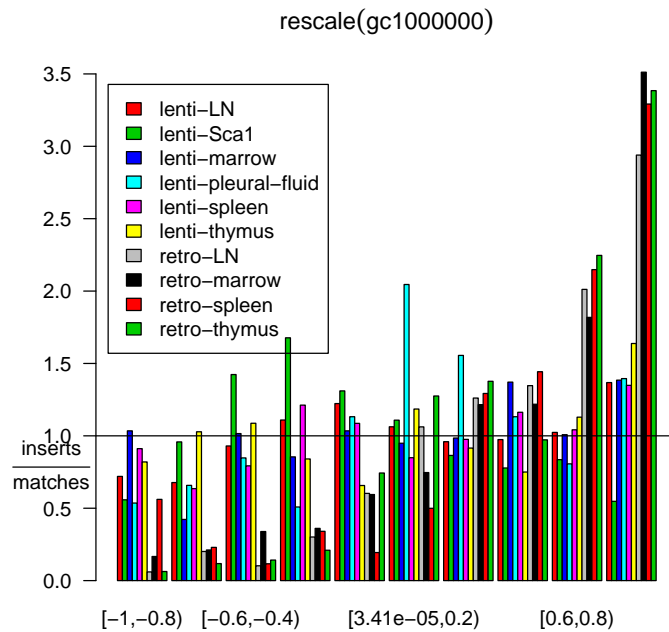
	coef	se	z	p
lenti-LN	0.04070	0.0824	0.4940	6.21e-01
lenti-Sca1	-0.53100	0.0865	-6.1300	8.62e-10
lenti-marrow	-0.04660	0.1850	-0.2520	8.01e-01
lenti-pleural-fluid	0.30700	0.2820	1.0900	2.76e-01
lenti-spleen	-0.00877	0.1170	-0.0747	9.40e-01
lenti-thymus	-0.15400	0.1880	-0.8180	4.13e-01
retro-LN	1.43000	0.1440	9.9100	3.82e-23
retro-marrow	1.36000	0.1890	7.2200	5.13e-13
retro-spleen	1.28000	0.2600	4.9200	8.63e-07
retro-thymus	1.51000	0.2180	6.9100	4.73e-12

rescale(gc5000000)
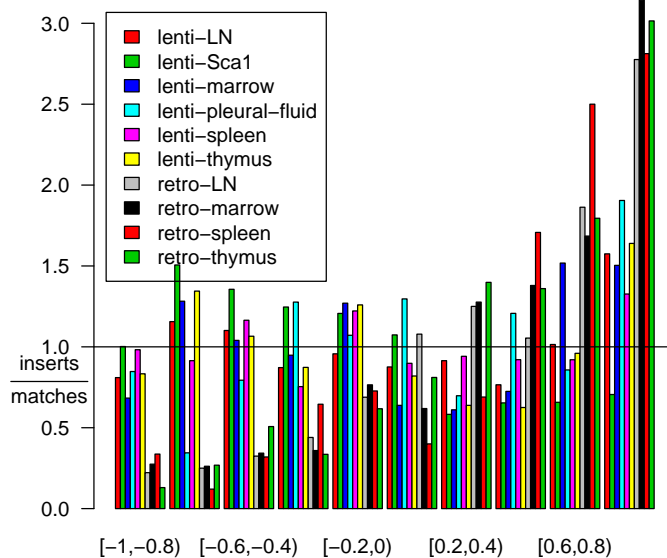
```
                      coef      se       z        p
lenti-LN           -0.0161  0.0825  -0.195  8.45e-01
lenti-Sca1         -0.4830  0.0862  -5.600  2.10e-08
lenti-marrow       -0.0432  0.1840  -0.234  8.15e-01
lenti-pleural-fluid  0.1500  0.2800   0.534  5.93e-01
lenti-spleen       -0.0585  0.1170  -0.500  6.17e-01
lenti-thymus       -0.1200  0.1870  -0.644  5.19e-01
retro-LN            1.1300  0.1330   8.480  2.25e-17
retro-marrow        1.3200  0.1870   7.060  1.70e-12
retro-spleen        1.2900  0.2580   4.990  6.17e-07
retro-thymus        1.0500  0.1980   5.320  1.02e-07
```
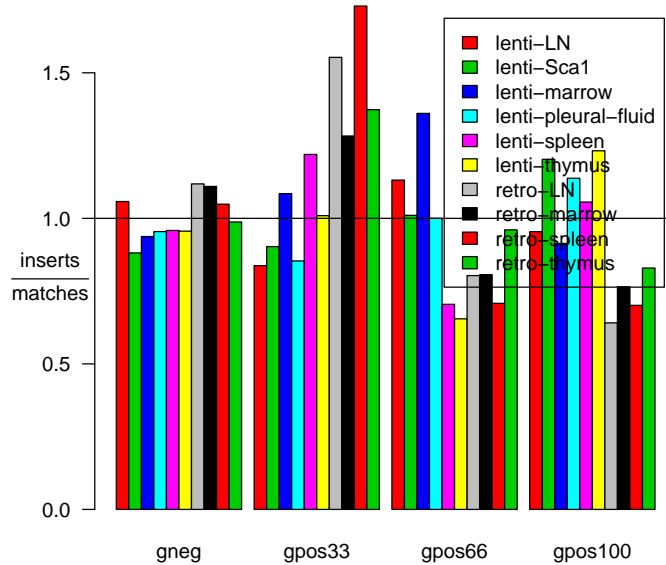
rescale(gc10000000)
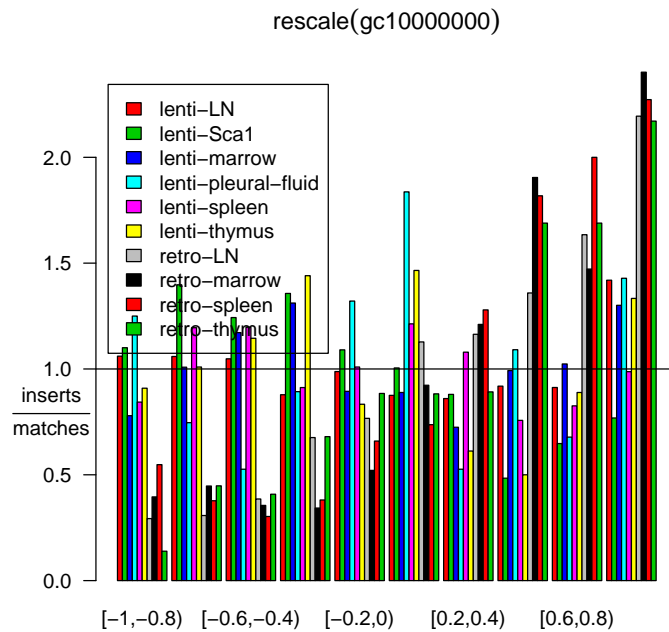
8 Cytobands

Here we study the association of cytoband with insertion intensity. The data are obtained from
`http://genome.ucsc.edu/goldenPath/hg17/database/cytoBand.txt.gz`.

A formal test of significance attains a p-value of 0.026514. Here is the table of coefficients of the log ratio of intensities for true insertion sites versus control insertion sites (comparing each category of Giemsa staining to 'gneg') along with their standard errors, z statistics, and p-values:

```
                   coef     se      z      p
cyto.typegpos100 -0.0284 0.0488 -0.582 0.560
cyto.typegpos33   0.0933 0.0579  1.610 0.107
cyto.typegpos66  -0.0524 0.0676 -0.774 0.439
```

References

[1] Yvonne M.M. Bishop, Stephen E. Fienberg, and Paul W. Holland. *Discrete multivariate analyses: Theory and practice* (MIT Press, 1975).

[2] P. McCullagh and John A. Nelder. *Generalized linear models.* (Chapman & Hall ltd, 1999).

[3] Xiaolin Wu, Yuan Li, Bruce Crise, Shawn M. Burgess "Transcription Start Regions in the Human Genome Are Favored Targets for MLV Integration," *Science,* **300**(5626), (June 2003): 1749-1751.