

Discovering interactions among BRCA1 and other candidate genes associated with sporadic breast cancer

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Analysis of a subset of case-control sporadic breast cancer data, [from the National Cancer Institute's Cancer Genetic Markers of Susceptibility (CGEMS) initiative], focusing on 18 breast cancer-related genes with 304 SNPs, indicates that there are many interesting interactions that form two- and three-way networks in which *BRCA1* plays a dominant and central role. The apparent interactions of *BRCA1* with many other genes suggests the conjecture that *BRCA1* serves as a protective gene and that some mutations in it or in related genes may prevent it from carrying out this protective function even if the patients are not carriers of known cancer-predisposing *BRCA1* mutations. The method of analysis features the evaluation of the effect of a gene by averaging the effects of the SNPs covered by that gene. Marginal methods that test one gene at a time fail to show any effect. That may be related to the fact that each of these 18 genes adds very little to the risk of cancer. Analysis that relates the ratio of interactions to the maximum of the first-order effects discovers significant gene pairs and triplets.

Breast cancer (MIM 114480) has complex causes. Known predisposition genes explain <15% of the breast cancer cases. It is generally believed that most sporadic breast cancers are triggered by unknown combined effects, possibly because of a large number of genes and other risk factors, each adding a small risk toward cancer etiology. Progress in seeking breast cancer genes other than *BRCA1* and *BRCA2* has been slow and limited because the individual risk due to each gene is small. This difficulty may be partly due to the fact that current methods rely largely on marginal information from genes studied one at a time and ignore potentially valuable information because of the interaction among multiple loci. Because each responsible gene may have a small marginal effect in causing disease, it is likely that such methods will fail to capture many responsible genes by studying a dataset where the disease may be due to a variety of different sources. The possible presence of many genes responsible for different subgroups of cancer patients may reduce the power of current methods to detect genes partly responsible for some forms of breast cancer. It is believed that methods effective in extracting interactive information from data should be developed.

What should be done when marginal effects are too weak to be detected? Our methods use interactive information from multiple sites as well as marginal information. They provide power to detect interactive genes. To test this claim and to demonstrate the practical value of these methods in real applications, we apply them to an important study: a subset of a large dataset collected from a case-control sporadic breast cancer study, focusing on gene-gene-based analysis. This partial dataset comprises 18 genes with 304 SNP markers. The application results in a number of scientific findings.

The message of this article is fourfold. First, if marginal methods fail, more powerful methods that take into account interactive information can be used effectively. We apply our proposed methods to this dataset to illustrate the detection of the interactions between genes. We point out that in our findings, none of the 18 selected genes show any detectable marginal

effects that are significantly higher than those generated by random fluctuations. In other words, all of the 18 genes would be missed if only marginal methods were used.

Second, we demonstrate how to carry out a gene-based analysis by treating each gene as a basic unit while incorporating relevant information from all SNPs within that gene. Two summary test scores are proposed to quantify the strength of interactions for each pair of genes. The pairwise interactions can be extended easily. We also provide results using third-order interactions.

Third, to establish statistical significance, we generate a large number of permutations of the dependent variable (case or control) to see how the measures of interaction for the real data compare with those from the many permutations.

Finally, when these procedures are applied to the data, they lead to a number of interesting findings. It is shown that there are a substantial number of significant interactions that form a network in which *BRCA1* plays a dominant role. The interactions of *BRCA1* with many of the other genes suggests the conjecture that *BRCA1* serves as a protective gene and that some mutations in it or in related genes may prevent it from carrying out the protective function.

Results

None of the 18 marginal effects are significantly higher than those generated by random permutations. This claim is made based on comparisons between the observed 18 V values and the values obtained from 1,000 permutations. The exact procedures carried out were described in steps 1 and 2 in *Methods*. Of the 18 genes, the most significant one is *BARD1* with a P value = 0.053, which is slightly short of being significant at 0.05. It is also noticed that *BRCA1* has a very large P value (0.944), meaning that the real effects of *BRCA1*, as well as those of the other genes in breast cancer, cannot be properly measured and reflected by its marginal effect alone. It is the interactive effect that reveals the roles of some of these genes in breast cancer. See [supporting information \(SI\) Fig. S1](#).

In seeking pairwise interactions among the 18 genes, both Mean-ratio and Quantile-ratio methods were implemented. Although the Mean-ratio method using (M , R) is more conservative than the Quantile-ratio method using (M , Q), it still identified 16 of 153 pairs as significant gene pairs at the 0.05 level that are connected with breast cancer etiology with an estimated

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Table 1. Significance of the gene pairs identified by the Mean-ratio and Quantile-ratio methods with *P* values estimated by the curve and rank methods

Pair no.	Mean-ratio method			Quantile-ratio method		
	Gene pair	Curve <i>P</i>	Rank <i>P</i>	Gene pair	Curve <i>P</i>	Rank <i>P</i>
1	<i>ESR1 BRCA1</i>	0.017	≤0.001	<i>ESR1 BRCA1</i>	0.013	0.001
2	<i>BRCA1 PHB</i>	0.026	0.040	<i>BRCA1 PHB</i>	0.029	0.073
3	<i>KRAS2 BRCA1</i>	0.002	0.006	<i>KRAS2 BRCA1</i>	0.002	0.004
4	<i>SLC22A18 BRCA1</i>	0.032	0.072	<i>SLC22A18 BRCA1</i>	0.019	0.079
5	<i>RAD51 BRCA1</i>	0.052	0.090	<i>RAD51 BRCA1</i>	0.005	0.032
6	<i>RB1CC1 SLC22A18</i>	0.024	0.026	<i>ESR1 SLC22A18</i>	0.033	0.016
7	<i>CASP8 KRAS2</i>	0.043	0.038	<i>RB1CC1 SLC22A18</i>	0.009	0.008
8	<i>CASP8 SLC22A18</i>	0.042	0.048	<i>CASP8 KRAS2</i>	0.038	0.036
9	<i>PIK3CA BRCA1</i>	0.030	0.048	<i>CASP8 SLC22A18</i>	0.021	0.012
10	<i>PIK3CA ESR1</i>	0.047	0.032	<i>PIK3CA BRCA1</i>	0.014	0.049
11	<i>PIK3CA RB1CC1</i>	0.047	0.051	<i>PIK3CA ESR1</i>	0.021	0.005
12	<i>PIK3CA SLC22A18</i>	0.025	0.036	<i>PIK3CA RB1CC1</i>	0.044	0.053
13	<i>BRCA1 CHEK2</i>	0.016	0.031	<i>CASP8 PIK3CA</i>	0.007	0.009
14	<i>BARD1 BRCA1</i>	0.032	0.057	<i>BRCA1 CHEK2</i>	0.007	0.022
15	<i>BARD1 ESR1</i>	0.044	0.025	<i>BARD1 BRCA1</i>	0.003	0.015
16	<i>BARD1 TP53</i>	0.019	0.019	<i>BARD1 ESR1</i>	0.017	0.003
17				<i>BARD1 TP53</i>	0.015	0.010
18				<i>BARD1 SLC22A18</i>	0.056	0.063
	<i>CASP8 ESR1</i>	0.071	0.048	<i>CASP8 ESR1</i>	0.066	0.031
				<i>BARD1 KRAS2</i>	0.055	0.036
	<i>ESR1 KRAS2</i>	0.145	≤0.001	<i>ESR1 KRAS2</i>	0.103	≤0.001
	<i>ESR1 PPM1D</i>	0.252	0.021	<i>ESR1 PPM1D</i>	0.348	≤0.001

The gene pairs are listed in the same order as in Figs. 3 and 4.

false discovery rate (FDR) of 50%. Of these 16 pairs, three pairs, (*BRCA1, ESR1*), (*BARD1, ESR1*) and (*BRCA1, KRAS2*) are significant even under more stringent criteria. The estimated *P* values are 0.001, 0.003 and 0.004; see Table 1. Four pairs are significant at level 0.01, and the remaining pairs are significant at the 0.05 level by using the rank method. A pairwise network based on these 16 pairs is shown in Fig. 1. On the other hand, the Quantile-ratio method is more powerful. For example, 18 pairs of genes with an estimated FDR of 40% were identified. A pairwise network based on these 18 pairs is shown in Fig. 2. It is interesting to note that 15 of the 18 identified pairs were also found by the Mean-ratio method.

Using the gene database from the National Center for Biotechnology Information (NCBI), we compare our results with published findings on these gene pairs. In Fig. 2, we have labeled with a red dot the interactions that were previously biologically identified.

There have been multiple reports on the biochemical interactions between *BRCA1* and *BARD1* (e.g., refs. 1 and 2). Meza *et al.* (1) studied, in detail, their interaction regarding the structure domains for the interaction, binding activities, etc. Further studies have shown that a subset of mutations at *BRCA1* can affect the *BRCA1–BARD1* interaction's relation with the susceptibility to breast cancer (3).

Functional interaction between *BRCA1* and *RAD51* has been identified and suggested to play a role during the S phase of the cell cycle, which is of critical importance for genome integrity (4). Levy-Lahad *et al.* (5) found evidence suggesting that a mutation at *RAD51* modified breast cancer risk among *BRCA2* carriers but not *BRCA1* carriers. It is interesting that we detected interaction between the *RAD51–BRCA1* genes among sporadic breast cancer patients, whereas neither gene demonstrated strong marginal effects.

CHEK2 is an important breast cancer candidate gene. Its association with breast cancer has been widely studied in multiple populations (e.g., refs. 6–8), and it has been shown to be a low-penetrance breast cancer-susceptibility gene (9) and suggested to have complex yet limited involvement in breast cancer etiology (7, 10). Interestingly, as noted in ref. 10, the *CHEK2**1100delC mutation was observed more frequently among cases with a first-degree-relative breast cancer patient. This may suggest that *CHEK2* may interact with important breast cancer-susceptibility genes as a risk modifier. The *BRCA1–CHEK2* interaction is known to be important for DNA damage response (10). Our findings suggested that the association between the *BRCA1–CHEK2* interaction and breast cancer susceptibility is much stronger than the marginal effects of these two

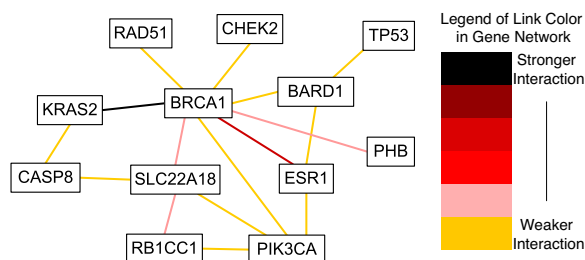


Fig. 1. Pairwise network based on 16 pairs of genes identified by the Mean-ratio method.

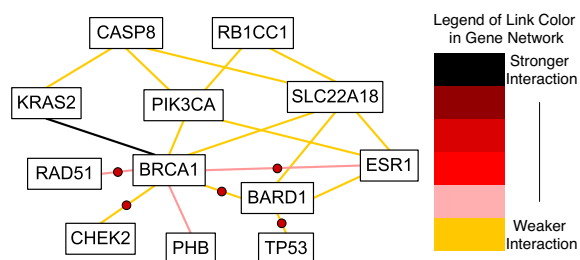


Fig. 2. Pairwise network based on 18 pairs of genes identified by the Quantile-ratio method.

The number of SNPs of the region that covers the corresponding gene varies from gene to gene, ranging from 2 to 78 SNPs. Because on average there are ≈3,000 bp between two consecutive SNPs, the largest region that covers the corresponding gene may spread over more than ≈100,000 bp. It is also noted that the genotypes of nearby SNPs are dependent on each other (with a noticeable linkage disequilibrium), and this dependence may stretch rather far along the regions. To take advantage of this dependence and to integrate the neighboring information, we treat each gene/region as a basic unit instead of treating each SNP as a unit. This illustrates our “gene-based approach,” in which information on SNPs is combined into one unit for that gene. Therefore, when we talk about the effect of a certain region or a certain gene, we simply mean the average effect of all individual effects due to all of the SNPs within that region/gene. Analogously, a two-way interaction of two genes, constitutes the average of all pairwise interactions of SNP pairs formed from the two regions/genes, one from each.

Methods. Consider k SNPs in a case-control study. Because each SNP determines three genotypes, k SNPs define 3^k multilocus genotypes that form a partition Π of the dataset into corresponding subsets called partition elements. We shall use the following genotype statistics to measure the observed joint effect of these k SNPs on the disease status:

$$v = \sum_{s=1}^{3^k} \left(\frac{n_{D,s}}{n_D} - \frac{n_{U,s}}{n_U} \right)^2, \quad [1]$$

where $n_{D,s}$ and $n_{U,s}$ are counts of cases and controls in genotype (element) s , n_D , and n_U are the total number of cases and controls under study. Let $n_s = n_{D,s} + n_{U,s}$ be the number of observations in genotype s .

Let Y be the disease status, 1 for cases and 0 for controls. Then, for a genotype partition Π , the statistic v discussed in this article was used in our previous publications (32–35) and can be shown to be a multiple of a general version as follows,

$$I_{\Pi} = \sum_{s \in \Pi} n_s^2 (\bar{Y}_s - \bar{Y})^2. \quad [2]$$

The key statistics used in this article are all derived from v .

We offer the following view of how two genetic loci can act jointly to affect their disease outcome. If the joint effect exceeds their individual marginal effect, we regard this as evidence of “interaction.” Specifically, our concept of interaction of two loci at i and j is that v_{ij} is substantially greater than $\max(v_i, v_j)$. The implication is that such an effect, in a large population, suggests that there may be a causal mechanism of biological importance. In the rest of this article, the joint effect of two loci is referred to as a second-order effect, and the marginal effect of one locus is called a first-order effect.

First-order effects failed to indicate anything special about the 18 genes. Second-order effects for pairs of genes were also very disappointing until we compared the second-order effects with the first-order ones. The ratio of the improvement, due to second order over the first order, seemed to signal some connections. This indication was especially impressive when this ratio was compared with the maximum of the two first-order effects in a two-dimensional graph. Two alternative ratios, to be defined later, are considered to be used in conjunction with the maximum of the first-order effects. These will be called the Mean-ratio and Quantile-ratio.

To establish significance, we applied 1,000 permutations of the case-control outcomes, to determine the null distribution of the ratio and the maximum partially described above. We propose two methods of assigning significance levels. One will be called the curve method and the other, the rank method.

Summary of Procedures. 1. Suppose that there are m_i SNPs in gene i . For SNP d of gene i we calculate v_{id} (defined as in Eq. 1 for $k = 1$) and take the average over the m_i SNPs within the gene. This leads to 18 averages $V = \{V_1, V_2, \dots, V_{18}\}$, each representing the (marginal/main) effect of one of the genes. We also calculate the (pairwise/interaction) two-dimensional genotype statistics v_{idj_e} defined as in Eq. 1 for $k = 2$, for all $(304 \times 303)/2 = 46,056$ SNP pairs. Throughout the article, we will use the lowercase to denote the quantities at the SNP-level and uppercase for the gene-level. Also, although we confine ourselves here to SNPs on 18 genes, our methods would apply equally well to a set of SNPs that cover a region of interest that is not necessarily a gene.

2. To detect possible causal effects of genes on disease status, we should know what to expect if there is no relationship. To obtain this information, we destroy any relationship by means of a random permutation of the labels of case ($Y = 1$) and control ($Y = 0$) and carry out the step 1 above with the permuted data. For example, calculations based on the first part of step 1 generate 304 random effects (values) purely by chance, whereas the original

dependence among SNPs in each gene is retained. We repeat this procedure 1,000 times to produce 1,000 sets of 304 values denoted by $v^{(p)} = \{v_i^{(p)}, 1 \leq i \leq 304\}$, where $\{1 \leq p \leq 1,000\}$. Similarly, one can estimate the null distribution of v for a pair of SNPs when there is no relationship by using the permuted data 1,000 times following the second part of step 1. In this article we use 1,000 permutations for two-way interactions but only 100 for three-way interactions.

For the benefit of nonstatisticians reading this paper, we digress to present a tutorial on significance levels because of the somewhat unusual applications we face in this presentation. The method of significance testing has had an important history in science. A scientist, who is interested in whether his data represent the presence of an effect of interest or are due to random fluctuations, formulates a null hypothesis that there is no effect. He calculates a statistic, T , based on the data. He finds the probability distribution of T , assuming the null hypothesis. He then calculates a P value that is the probability, under the null hypothesis, that T exceeds the observed value t of T . This probability is called the significance level or the P value. If the P value is small, that is regarded as a sign that there is an effect. What constitutes sufficiently small should depend on the background of the problem, although it is customary to use 0.05 or 0.01 as levels to determine statistical significance. If he chooses a statistic that is not sensitive to deviations from the null hypothesis, he is unlikely to get a low P value even when the hypothesis is false and there is an effect. So it pays for him to select a statistic and an experimental design that will be sensitive to those deviations from the null hypothesis in which he is interested.

Our applications have some unusual aspects. By regarding the question of whether a gene has an effect on breast cancer, we are effectively testing 18 distinct null hypotheses. In dealing with interactions, we are testing 153 null hypotheses simultaneously. Our reduced data for the latter problem consists of two statistics, R and M to be defined below. The methodology is ordinarily defined for only one test statistic. Our theory is inadequate, in our problem, to calculate the distribution under the null hypothesis, and thus we have to resort to randomized permutations. These aspects must be addressed here.

Returning to item 2, Fig. S1 presents, for the 18 genes, estimates of $P(V \geq t)$, as estimated by the permutations. The location of the observed value of V is indicated by a vertical line. The reader will note that none of the 18 P values represented is significant at the 0.05 level.

3. We seek pairwise interactions among the 18 genes. A definition of pairwise interactions between two genes is given first. There are a total of $18 \times 17/2 = 153$ gene pairs. We shall measure the amount of interaction between two genes, say g_i and g_j , as follows: For each pair of SNPs, (i_d, j_e) , one from gene i and another from gene j , define the SNP-wise interactions as the ratio of incremental interactions vs. the maximum of the two marginal effects. That is,

$$r(i_d, j_e) = \frac{v_{idj_e} - v_{id} \vee v_{j_e}}{v_{id} \vee v_{j_e}}, \quad [3]$$

where “ \vee ” stands for maximum of the two values. This ratio represents the relative amount of interactions of two SNPs with respect to their marginal effects. The amount of interactions between two genes i and j is then defined as the average of all SNP-wise ratios possibly formed from these two genes and will be denoted by

$$R_{ij} = \frac{\sum_{d=1}^{m_i} \sum_{e=1}^{m_j} r(i_d, j_e)}{m_i m_j}. \quad [4]$$

We term this as the “mean interaction ratio” between two genes, or simply, the Mean-ratio or R statistic. In step 6 below, we will introduce an alternative Quantile-ratio or Q statistic.

For each gene pair, we also define the “average maximum marginal v ” or “ M statistic” as

$$M_{ij} = \frac{\sum_{d=1}^{m_i} \sum_{e=1}^{m_j} (v_{id} \vee v_{j_e})}{m_i m_j} \quad [5]$$

4. From steps 1–3 above, we obtained a set of 153 points, $\{(M_{ij}, R_{ij}); 1 \leq i < j \leq 18\}$, corresponding to all possible gene pairs. We use a scatter plot of R vs. M in Fig. 3, to display these 153 points. The same procedure was repeated on 1,000 permuted datasets. This process will generate 1,000 sets of points, each set consisting of 153 points, $\{(M_{ij}^{(p)}, R_{ij}^{(p)}); 1 \leq i < j \leq 18\} = \Gamma^{(p)}$, where p stands for the p th permutation. It also generates 153 sets consisting of 1,000 points each.

5. Based on these 1,000 sets $\{\Gamma^{(p)}\}$, each having 153 points, we construct a 95% confidence curve on the (M, R) plane as shown in Fig. 3 by the dark blue curve, and as described below. Points above the dark blue curve will represent

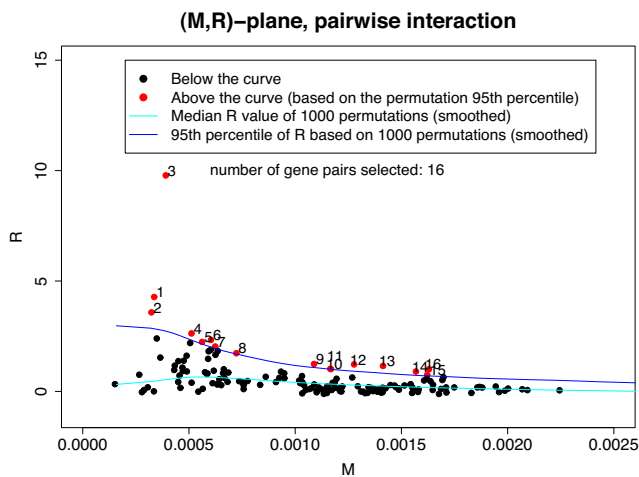


Fig. 3. (M,R)-plane: observed data and permutation quantiles. 1-*ESR1* *BRCA1*, 2-*BRCA1* *PHB*, 3-*KRAS2* *BRCA1*, 4-*SLC22A18* *BRCA1*, 5-*RAD51* *BRCA1*, 6-*RB1CC1* *SLC22A18*, 7-*CASP8* *KRAS2*, 8-*CASP8* *SLC22A18*, 9-*PIK3CA* *BRCA1*, 10-*PIK3CA* *ESR1*, 11-*PIK3CA* *RB1CC1*, 12-*PIK3CA* *SLC22A18*, 13-*BRCA1* *CHEK2*, 14-*BARD1* *BRCA1*, 15-*BARD1* *ESR1*, 16-*BARD1* *TP53*

potentially significant gene pairs. This method will be called the Mean-ratio curve method.

We place all points (total $153 \times 1,000 = 153,000$ points) obtained from these 1,000 permutations on the (M, R) plane. The values of the M coordinate are binned into 100 bins, each with 1,530 points. Each bin is represented by a point with coordinates (M^*, R^*) , where M^* is the mid value of the bin, and R^* is the 95th percentile of those R scores falling into the bin. A smoothing fit, using a spline with 15 knots, is made to the curve formed by these 100 points (M^*, R^*) , and the resulting dark blue curve is shown in Fig. 3. We also introduce the following curve method of assigning significance levels. For each gene pair (i, j) , we shall determine its nominal P value by $S_{ij}/1,530$ when the pair's R value is the S_{ij} highest among all 1,530 R values obtained from 1,000 permutations that fall into the same bin. We use the term "nominal" because this method does not satisfy the standard criterion of applying the null distribution of a one-dimensional test statistic. The Rank method to be described later does satisfy criterion.

6. The ratio measure of interaction is sensitive to the possibility that a relatively low interaction for two SNPs has a high ratio when both marginal effects are very small. The resulting high ratio may have a large effect on the overall mean R_{ij} . Instead of using $R = R_{ij}$, we moderate this problem by introducing an alternative measure of interaction between two genes i and j as the top 90th quantile of the SNP-wise ratios formed by the $m_j m_i$ SNP pairs. This will be denoted by $Q = Q_{ij}$ and will be called the Quantile statistic. With R_{ij} replaced by Q_{ij} , we carried out steps 4 and 5 and obtained 153 points $\{(M_{ij}, Q_{ij}); 1 \leq i < j \leq 18\}$. The same procedure was applied to 1,000 permuted datasets generating 153,000 points. We apply the smoothing techniques described in step 5 to these points, leading to a dark blue curve shown in Fig. 4. We will call this the Quantile-ratio method.

7. With mild modifications and adequate computational power, the six steps described above can be generalized to detect third- or higher-order interactions. For example, in three-way interactions, consider three SNPs as a triplet, (i_d, j_e, k_f) , each selected from the respective genes i, j , and k . The three-way-SNP-wise interaction is now defined as the ratio of incremental interactions vs. the maximum of the three marginal effects. That is,

$$r(i_d, j_e, k_f) = \frac{v_{i_d j_e k_f} - v_{i_d j_e} \vee v_{i_d k_f} \vee v_{j_e k_f}}{v_{i_d j_e} \vee v_{i_d k_f} \vee v_{j_e k_f}} \quad [6]$$

At the gene level, we define the three-way interaction with genes i, j , and k as the average of all three-way SNP-wise ratios formed from these three genes, to be denoted by

1. Meza JE, Brzovic PS, King MC, Kleivt RE (1999) Mapping the functional domains of BRCA1 interaction of the ring finger domains of BRCA1 and BARD1. *J Biol Chem* 274:5659–5665.

(M,Q)-plane, pairwise interaction

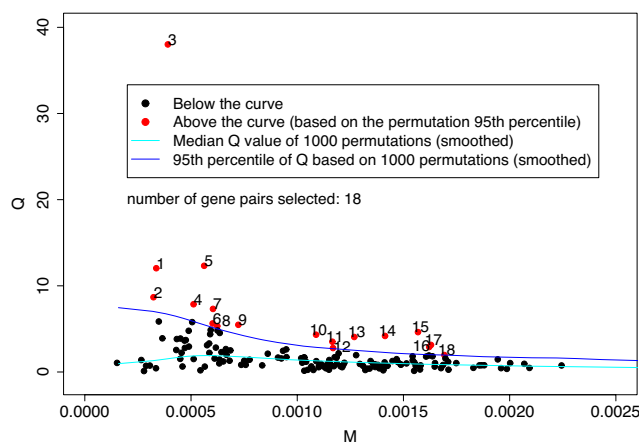


Fig. 4. (M,Q)-plane: observed data and permutation quantiles. 1-*ESR1* *BRCA1*, 2-*BRCA1* *PHB*, 3-*KRAS2* *BRCA1*, 4-*SLC22A18* *BRCA1*, 5-*RAD51* *BRCA1*, 6-*ESR1* *SLC22A18*, 7-*RB1CC1* *SLC22A18*, 8-*CASP8* *KRAS2*, 9-*CASP8* *SLC22A18*, 10-*PIK3CA* *BRCA1*, 11-*PIK3CA* *ESR1*, 12-*PIK3CA* *RB1CC1*, 13-*CASP8* *PIK3CA*, 14-*BRCA1* *CHEK2*, 15-*BARD1* *BRCA1*, 16-*BARD1* *ESR1*, 17-*BARD1* *TP53*, 18-*BARD1* *SLC22A18*

$$R_{ijk} = \frac{\sum r(i_d, j_e, k_f)}{m_j m_i m_k} \quad [7]$$

Analogously, one can mimic the quantile statistic to calculate Q_{ijk} for the three genes i, j , and k .

For each gene triplets, we also define

$$M_{ijk} = \frac{\sum (v_{i_d j_e} \vee v_{i_d k_f} \vee v_{j_e k_f})}{m_j m_i m_k} \quad [8]$$

With the scatter plots of (M, R) and (M, Q) based on a number of permuted datasets, we can derive the threshold curves to set various significant levels. Every point of these sets corresponds to a triplet with three genes involved. For the current data, there are $18 \times 17 \times 16/6 = 816$ such triplets. Instead of the 95-percentile level, we use the 99.5% level to determine R^* values in each bin and the threshold curves (Fig. S4 and Fig. S5).

8. The methods used in steps 5–7, to determine which pairs of genes yield a significant signal, depend for their power on an implicit assumption that the 1,000 points for one pair of genes gives rise to a similar cloud as for another pair of genes. Although an informal view supports this assumption, an alternative rank procedure is more reliable and more directly in keeping with the principle of using a single test statistic to evaluate the P value for a given hypothesis. This method uses the bins of step 5. We have called the first method the curve method, and this alternative, the rank method.

Consider the gene pair (i, j) and the corresponding statistics (M_{ij}, R_{ij}) . From the permutations we have 1,530 points (M, R) in each of 100 bins. The values of R in each bin are ranked from 1 to 1,530. If R_{ij} falls between the permutation derived points with ranks $s - 1$ and s in the bin for (i, j) , the gene pair is given the value $T = s$. For this gene pair, there are 1,000 values of $T^{(p)}$ derived from the permutations, and the proportion of $T^{(p)} \geq T$ is the P value for the gene pair. In effect, the rank method uses the bins for constructing the original method of finding pairs that are significant at the 5% level as the source of a one-dimensional statistic from which a traditional P value can be constructed. The same approach applies to (M, Q) . Table 1 displays the respective P values obtained from the curve and rank procedures for those gene pairs above the curve in Figs. 3 and 4.

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