Methodology article

A reliable measure of similarity based on dependency for short time series: an application to gene expression networks

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Abstract

Background: Microarray techniques have become an important tool to the investigation of genetic relationships and the assignment of different phenotypes. Since microarrays are still very expensive, most of the experiments are performed with small samples. This paper introduces a method to quantify dependency between data series composed of few sample points. The method is used to construct gene co-expression subnetworks of highly significant edges.

Results: The results shown here are for an adapted subset of a Saccharomyces cerevisiae gene expression data set with low temporal resolution and poor statistics. The method reveals common transcription factors with a high confidence level and allows the construction of subnetworks with high biological relevance that reveals characteristic features of the processes driving the organism adaptations to specific environmental conditions.

Conclusion: Our method allows a reliable and sophisticated analysis of microarray data even under severe constraints. The utilization of systems biology improves the biologist’s ability to elucidate the mechanisms underlying cellular processes and to formulate new hypotheses.

Background

In recent years, the technology of DNA microarrays has been central for the knowledge assembly in molecular biology research. The possibility of measuring mRNA levels in a genomic scale in a comparative way enables the discovery of genes or clusters of genes which expressions are differentiated in a specific condition, thus providing important clues about gene functions and pathways. Statistical tools have been developed for the analysis of this great amount of biological data aiming to help biologists to make predictions and originate hypotheses to be experimentally tested.

In the past decade, systems biology approaches are emerging as a novel concept increasingly attractive to deal with this kind of data. The biological system is modeled as a graph and each biological unit (a gene, for instance) is represented as a node in this graph and if two nodes interact they are connected by an edge [1,2]. This approach produces a visually appealing structure generically called

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network where nodes with similar characteristics can be
grouped together and groups of tightly interconnected
odes are often associated with a specific cellular function [3,4].
Perhaps the most interesting feature of this application
is the possibility to analyze how these modules of
tightly interconnected nodes connect among themselves
or how different functions interact to produce the
observed phenotype. In fact, this kind of approach is prov-
ing to be a more robust tool for the extraction of relevant
information from the biological data and abstraction and
better comprehension of the biological system as a whole
[5].

In the specific case of networks inferred from gene expres-
sion data, an interaction between two genes can be deter-
mined by a measure of association between their
expression patterns. Thus, in order to get a reliable struc-
ture, one must first define a consistent measure to quan-
tify gene expression similarity. The most commonly ones
used are based on Pearson correlation (for linear associ-
ation) [6,7] or mutual information (that accounts for any
kind of association function) [8-10]. These are very robust
measures and are widely used in a number of applications
involving the inference about the degree of dependency
between biological signals.

However they are based on statistical assumptions that
depends on assymptotic supositions making them
strongly dependent on large samples sizes. This becomes
an important drawback in the context of molecular biol-
ogy applications. DNA microarrays are still a very expen-
sive technique and the design of experiments with a large
number of different conditions and/or time acquisition
demands such human and financial resources that could
make it unachievable in many research laboratories. Thus,
the usual procedure is to design experiments with a small
number of observations. In fact, according to the Gene
Expression Omnibus [11], about one third of microarray
studies involves experiments with 3–8 time points or
other types of non-temporal sequential data [12].

Limited sampling accentuates the difficulties related to
standard signal analyses. Among the most important
problems are the influence of noise that becomes even
more prominent with shorter series, enhancing the com-
plexity in distinguishing real from random patterns and
increasing the potential of misleading results [13]. The
efforts to overcome difficulties related to limited sampling
include strategies of simplification [14,15] and the incor-
poration of multi-source information [16]. In the first
case, the goal is to transform continuous data to discrete
representations prior to analysis, categorizing the gene
expression data into a set of different states trying to cap-
ture tendencies instead of absolute values. However, sim-
plication strategies are highly dependent on pre-
definitions about the a priori patterns of gene expression
in the discretization step, a process which is largely
dependent on the researchers’ expertise [13]. Additionaly,
valuable information is lost during this process. On the
other hand, incorporating multi-source information
includes prior knowledge or multi-scale and different lev-
eels of information from other sources to improve the
computational analysis of short time-series microarray
data. This approach faces the challenge of dealing with a
high heterogeneity of data which increases the difficulties
in extracting meaningful information. Furthermore, for
many organisms for which the state of research is still in
its infancy, there is no reasonable additional information
or any prior knowledge to rely on. Notwithstanding the
statistical difficulties inherent to this kind of analysis,
these data represent a rich source of information and the
design of a rational pipeline to better explore them is of
paramount importance.

Here we present a method of inference of dependency
between series from a temporally poor data set and apply
this method to the construction of condition-specific sub-
networks from gene expression data. The proposed
method is an alternative to the standard measures of
dependency. It performs well with short series and
requires no a priori assumptions. In order to evaluate the
method, we show results for an adapted subset of a bench-
mark data set of *Saccharomyces cerevisiae* [17] aiming to
mimic the limiting conditions above described. We show
that the method is efficient in capturing dependencies
between the gene signals and that these dependencies can
be related to the existence of a common transcription reg-
ulatory factor with a high confidence level. Our results
strongly suggest that the method proposed allows the
application of systems biology to data sets obtained from
limited amount of experiments and that this method
offers the biologist a robust tool to analyze gene expres-
sion data.

**A similarity measure**
The output of a microarray experiment is given as the log
of the ratio between the amount of mRNA of an experi-
mental sample and a control sample. This ratio is com-
monly called log-ratio and gives the degree of modulation
of a gene relative to the control sample. A common
attempt in these experiments is to compare networks orig-
inated from different experimental conditions trying to
find either similarities or differences that explain the
observed phenotype. For example, one could create differ-
ent mutant strains inactivating key genes and then analyze
separately the expression profiles of each mutant in com-
parison to the expression profiles of the wild type. This
approach would generate different networks promptly
revealing the distinct adjustments the organism takes in
response to each experimental condition.
The present work aims to reconstruct the genes co-expression networks in a condition specific way. To make this reconstruction possible, we introduce a measure of association between gene profiles (log-ratios measurements). Consider, thus a data set consisting of log-ratios for $N$ genes in $\rho$ different experimental conditions. The parameter $\rho$ accounts for different series of experiments hence, different physiological conditions as well as different time points. Consider also that we are particularly interested in a specific subset of $\rho$ say, a specific series of experiments consisting of $\rho'$ sample points. The question addressed here is how to infer similarity between two gene profiles consisting of $\rho'$ sample points, considering usually $\rho' < 10$.

One can think of some points worth to consider when idealizing an efficient method:

1. The function that relates co-regulated genes is not necessarily linear. In fact it can assume very complex non-linear forms. Thus, an efficient method of inference of gene relations should be able to capture any association function that should arise naturally rather than being enforced a priori;

2. Co-regulated genes are expected to respond in a correlated way in different experimental conditions and even the lack of modulation in a given experimental condition is actually an useful information. Although interested in constructing condition-specific networks, we would like to use the valuable information contained in other possible series of experiments to determine the associated pairs. Thus, information about how a given pair of genes behaves in all known $\rho$ conditions could be taken into account when calculating a measure of similarity between pairs of gene expression profiles;

3. Microarray experiments involve several factors where chance fluctuations and random processes play a significant role. Thus, points around zero, or very small modulation rates are more likely to reflect experimental artifacts and lack biological meaning than higher values. It would thus be desirable if the method gave different weights to different levels of modulation;

Based on these leads, the similarity $S_{ij}$ between genes $i$ and $j$ concerning $\rho'$ conditions can be defined as:

$$S_{ij} = \sum_{r=1}^{\rho'} s(x^i_r; x^j_r),$$

(1)

where $x^i_r$ is the modulation rate of gene $i$ in condition $r$. Notice that the score $S$ can be obtained summing over any subset of $\rho$ allowing the researcher the choice of analysis of each perturbation individually or the entire data set depending on his/her interests.

Define now a random vector $(X; Y)$ whose elements are modulation rates of two genes to be observed in one experimental condition. Considering $(x; y)$ as a possible observation of $(X; Y)$, the function $s$ as used in this work is defined as follows:

$$s(x; y) = \log \frac{h(x; y)}{f(x)g(y)},$$

(2)

for $f$ and $g$ being, respectively, the marginal densities of $X$ and $Y$ and $h$ the joint probability density of $(X; Y)$. Equation 2 is known as a log-likelihood score in the context of testing independence of two random variables. We do not know the form of these densities a priori. In this work we use the data information contained in all $\rho$ experiments to estimate it. Note that if we have $h$ we simply derive the marginal densities $f$ and $g$. In the sequel we delineate the way $h$ is estimated.

In order to estimate $h$, we compile a list of co-occurrences for the $N$ genes in the $\rho$ experimental conditions. This list is obtained by taking all the possible pairwise combinations of modulation rates for each experimental condition. The number of pairs is $\rho^2 \frac{(N^2 - N)}{2}$. Note that the set of $N$ genes is previously filtered to comprise only the genes modulated in at least one experimental point. This step reduces largely the computational cost. With the co-occurrences list one can construct a bi-dimensional frequency matrix from which is estimated the joint probability density in equation 2. The rationale behind this is that, in an ideal situation all the genes modulated in a specific condition are somehow related to the perturbation(s) impinging to the system and, thus, directly or indirectly related to each other. Hence, the distribution of co-occurrences of these genes could be a good estimator of the function that describes the genes interdependencies. Of course, far from the ideal, the compilation of this list is also carrying a large amount of noise among the real signal. Nonetheless, we expect that the noise will contribute with a small amount or no information to the scoring function. Indeed, we will show that with this method we are able to identify and select the most significant links from the background and that these have a very relevant biological meaning.
Prior to the computation of the bi-dimensional frequency matrix, we make a discrete approximation of the values of expressions. The observed values of \( x_k \) were discretized into rectangular bins with size \( \delta \) defined according to Scott’s rule [18]:

\[
\delta = \frac{3.5\sigma}{N^{1/3}}
\]

where \( \sigma \) is the sample standard deviation and \( N \) is the number of samples in the data set. It is worth mentioning that the method is not sensitive to the choice of the size of the bins and the outcome of the analysis is quite conserved for a wide range of bin sizes and/or shapes (data not shown).

Figure 1 shows the form of the function \( s(x; y) \) for an artificial set of random Gaussian correlated variables. The \( x \)- and \( y \)-axis are the bins’ representative values. A high positive value of \( s \) (red colors) indicates that the pair \( (x; y) \) is likely to be strongly dependent on each other. The function highlights the variables lying in the diagonals (high correlation) and in the extremities of the distribution. On the other hand, log-ratios around zero contributes with zero information to the similarity score. However we would like to stress that a high measure of correlation (or dependency) between biological signals does not imply a causal relationship, hence any two given connected nodes in the networks generated from this approach are not guaranteed to be functionally related. In spite of this, our approach is a strong starting point for biological inferences.

We have proposed a method based on a standard log-likelihood score to infer association between gene expression profiles. The innovative in this work is how we calculate the \( h \) densities using different series of experiments altogether as a model of associated variables. Furthermore, although we use information on various sets of experiments to construct the scoring matrix, our method allows the analysis of each set individually since the similarity between pairs of genes can be assessed summing over the scores for any subset of the entire data set. Given the similarity matrix relating the \( N \) genes, one must set a cutoff

![Scoring matrix](image)
value $S_0$ above which the pairs of genes are considered connected in the network.

Removing edges originating from indirect interactions
Focusing on our objective of inferring subnetworks of strongly dependent genes, the process described above is subjected to an important limitation since genes separated by one or few intermediates and thus not directly interacting may achieve a high similarity value. In fact, this procedure originates very dense networks with a high occurrence of large cliques (groups of nodes completely interconnected) that does not agree with the expected structure of cellular networks [3]. To circumvent this drawback, we perform an additional step in our pipeline that removes potential indirect interactions using a known information theoretic property [19].

The procedure to remove edges originating from indirect interactions has already been used in gene co-expression networks [9,20] and was adapted here to our purposes. It states that if both genes $(i, j)$ and $(j, k)$ are directly interacting and $(i, k)$ are indirectly interacting through $j$, then $S_{ik} \leq S_{ij} \leq S_{jk}$. Thus we search for all the triangle loops in the network and discard the edges that satisfies the inequality. To account for inaccurate estimates of the difference between close values of $S$ we introduce a tolerance threshold:

$$S_{ik}^t = \begin{cases} 0 & \text{if } S_{ik} \leq S_{ij}(1-\epsilon) \text{ and } S_{ik} \leq S_{jk}(1-\epsilon); \\ S_{ik} & \text{otherwise.} \end{cases}$$

(4)

We acknowledge that the above restriction eliminates most of the indirect interactions at the expenses of eliminating also authentic direct interactions. Nevertheless, our main concern is to minimize the occurrence of false positives given the statistical constraints. The occurrence of false negatives, i.e., the absence of an interaction that actually exists in the co-regulated gene network reduces the potentiality of the method to infer new hypotheses about the system but, contrarily to the occurrence of false positives, it would hardly imply in unnecessary expenditures of time and resources to the involved laboratory.

In the following sections, we present results for a benchmark data set obtained from Spellman [17]. This set comprises data of DNA microarrays from yeast cultures synchronized by four independent methods, which are referred to as perturbations: alpha-factor arrest, elutriation and arrest of a $cdc15$ and a $cdc28$ temperature-sensitive mutants. Since our main objective is to evaluate a new method of exploring microarray data, a long discussion of biological results is beyond our purposes. Thus, we restrict our discussions to the results obtained with the alpha-factor and the $cdc15$ sets. Similar results were observed for elutriation and $cdc28$ temperature-sensitive mutant sets.

Results and Discussion

*S. cerevisiae* is one of the most studied organisms to date and there is a great amount of data about gene-to-gene and protein-to-protein interactions already validated on the web (for a review, see Saccharomyces Genome Database – http://www.yeastgenome.org). The Spellman data set [17,21] is being widely used with the aim of validating novel gene expression data exploratory methodologies. It consists of time samples taken in four different ways of synchronizing the cell cycle. After releasing the cultures from the stimuli, samples were taken over time totaling 73 samples: alpha (with 18 samples collected every 7 minutes), $cdc15$ (24 samples collected every 10 or 20 minutes), $cdc28$ (17 samples collected every 10 minutes) and elutriation (14 samples every 30 minutes). In order to recreate the conditions met in the experiments targeted by this study, we selected 12 samples – 3 time points from each experiment – according to Table 1 and all the calculations were performed over this adapted set.

The joint probability density distribution and the distribution of scores $s$ obtained for the yeast data set are shown in Figures 2A and 2B, respectively. Intensities are represented in colors in both figures where red colors represent higher intensities. The scale is given in the colorbar on the right of each figure.

The function $S$ defined in Equation 1 was applied separately to each of the four perturbation sets and similarity matrices were obtained for each one. The distribution of $S$ values obtained for the alpha set and the $cdc15$ set are

| Experiments                  | alpha        | cdc15        | cdc28        | elutriation |
|------------------------------|--------------|--------------|--------------|-------------|
| Points sampled:              | 28, 63 and 98 min | 30, 150 and 270 min | 30, 90 and 150 min | 120, 240 and 360 min |
| Original dataset:            | 0 to 119 min (2 cycles) | 10 to 290 min (3 cycles) | 0 to 160 min (2 cycles) | 0 to 390 min (1 cycle) |

First row are the samples chosen for the construction of the Saccharomyces bi-dimensional frequency matrix. Second and last rows correspond to the time range comprised by the original dataset and the number of cycles comprised by the time intervals as given by the author.
Figure 2
**Probability density distribution and scoring matrix for the yeast data.** The joint probability density distribution $h(x; y)$ of the yeast data set (panel A) and the respective distribution of log-scores $s$ (panel B). x- and y-axes are log-ratios. The intensities of the bivariate functions are given in colors. Hot colors represent higher values as scaled in the colorbars on the right. In this study, $N = 4461$ and $\rho = 12$. Genes were considered modulated if $x_k > |0.5|$ in at least one experimental condition.

Figure 3
**Distribution of scores – real data vs. shuffled ensemble.** Distribution of similarity values $S$ for the alpha and cdc15 data sets (shaded histogram) and their respective shuffled set (red filled histogram) representing the null model. alpha set – panel A; cdc15 set – panel B. The alpha set comprises 734 genes and the cdc15 set comprises 2517 genes modulated in at least one of $\rho'$ = 3 experimental conditions.
shown in Figures 3A and 3B, respectively. The distribution of scores S for a set of correlated variables presents a wide range of variation and the definition of a suitable cut-off above which two nodes are said to be significantly similar is quite arbitrary. Even for uncorrelated random variables, the distribution of S can achieve high positive values and thus, one must pay extra care in order to define a suitable cut-off value $S_0$. To deal with this, we generated an ensemble of uncorrelated signals by randomly shuffling the data set and averaged the obtained S distribution for 1000 repetitions. We then set $S_0$ according to a value of $p < 10^{-4}$. The networks were compiled assigning an undirected edge to each pair of nodes with $S > S_0$. The distributions of S for the shuffled sets are shown superimposed to the distributions of the real data in Figure 3. One can notice the remarkable difference between the real and shuffled distributions implying that the method is capturing some biological phenomena driving the expression behavior of the genes.

Upon the removal of the indirect interactions ($\varepsilon = 0.1$ in Equation 4), the resulting edges were investigated for validity according to validated biological data.

Validation of results
To validate the obtained results, each interaction found for each network (the undirected edges) is searched for biological meaning in the BioGRID (The Biological General Repository for Interaction Datasets) and the YeastRACT (Yeast Search for Transcriptional Regulators And Consensus Tracking) databases [22,23]. The former is a repository of the protein and genetic interactions reported to date for some model organisms including the baker yeast. The YeastRACT is a curated repository of regulatory associations between transcription factors and target genes in S. cerevisiae based on bibliographic references. We compared the obtained edges in two ways. First, we searched for the existence of a catalogued physical or genetic interaction in the BioGRID data bank. Second, we searched in the YeastRACT data bank for the existence of a common transcription-
tion factor regulating both genes in each resulting edge. Our results show a good agreement with the literature description. In the following section we show the results for the alpha-factor and cdc15 networks.

The networks
The resulting edges were investigated for validity both in the BioGRID and the Yeastact data banks and results are shown in the pie charts in Figure 4. This Figure presents the percentage of connections (edges) that have either been identified in the BioGRID data bank or that are regulated by the same transcription factor as given by the Yeastact data bank. Connections involving genes not documented in the cited databases counts as unknown genes.

Connections not found in either data banks are labeled as mis-assignments and are subdivided into two groups: those that involve dubious or uncharacterized ORFs and the rest of it. In order to evaluate the results a null model has been tested by randomly assigning edges among the modulated genes while keeping the same connectivity distribution of the original network. Averaged results for this random network are shown in shaded areas.

According to these results, one can expect that in average 35% of two randomly chosen genes modulated in the alpha perturbation are regulated by the same transcription factor in contrast to 72% obtained with our method ($p < 10^{-5}$). Similarly, in average 30% of two randomly chosen genes modulated in the cdc15 perturbation share a common transcription factor in contrast to 71% obtained with our method. One could argue that the rate of correct assignments for the random model is overestimated. However, we want to emphasize that the set of genes analyzed here and subjected to the described procedures comprises genes with a rate of modulation above a given threshold under the considered experimental treatment. Thus, we expect that they are all somehow related. Nonetheless, the efficiency of the method is highly above the expected by chance.

It is worth mentioning that the results shown in Figure 4 are mostly due to common documented transcription factors regulating the genes connected by an edge, i.e., due to agreement with the Yeastact data bank solely. The percentage of interactions found in the BioGRID data bank is around 10% for the alpha network and 3% for the cdc15 network ($p < 10^{-5}$). Although this result is statistically significant, we stress that the proposed procedure is best suited for elucidation of possible common regulation pathways rather than physical or genetic interactions as given by the BioGRID bank.

These results together reinforce the common assumption that the connections found with methods based on similarity between gene profiles are only conceptual. They mean solely that the involved pair presents patterns of expressions that are highly dependent under the statistical assumptions. Furthermore, the edges participating in the networks are selected under a rank of high statistical significance and some of them can be replaced if the parameters are changed. In spite of this, the kernel structure is usually maintained, i.e., central nodes and modular organization are usually kept under different parameter choices and can provide unparalleled valuable informations concerning genes functions and cellular states.

In order to evaluate the robustness of the method, we applied the same procedure to the complete data set (73 time points). We compared the results obtained with the complete sets of alpha and cdc15 (18 and 24 time points, respectively) to the results previously obtained (with only three time points). We observed a massive overlap between the nodes from both networks – 88% of the nodes in the alpha network and 97% of the nodes in the cdc15 network. Concerning the edges, we observed that about 10% are present in both networks. It is worth stressing that this is not a trivial result since the genes selected as nodes in the networks comprehend only a fraction of the total number of modulated genes.

The alpha-factor network
The alpha network is depicted in Figure 5. The two most connected genes CS T1 and MFA1 encode endochitinase (required for cell separation after mitosis, activated during late GAP (G) and early mitosis (M) cell cycle phases) and Mating pheromone a-factor (which interacts with alpha cells to induce cell cycle arrest and other responses leading to mating), respectively. A list with all the genes and its annotations according to the GEO is given as Supplementary file.

Since interactions are only conceptual we suggest that instead of analyzing individual genes and edges in an isolated way, one could extract more valuable information from the analysis of groups of densely connected genes. The symbolic representation of a network in groups or modules can help one to better comprehend its structure. In Figure 5 one can nearly distinguish four groups of nodes. To facilitate the evaluation of the modules, we applied a well-known community detection procedure based on edge betweenness over the network [24,25]. The edge betweenness community algorithm splits apart modules of nodes densely connected by successively removing the most central edges. This method produces a dendrogram displaying the nodes in the x-axis and the distance among them is proportional to the length of the y-axis.
The dendrogram relative to the alpha-factor network is given in Figure 6.

Spellman experiment's main objective was to investigate cell-cycle regulated genes [17,21]. With this aim, the authors synchronized yeast cultures in different ways analyzing transcript levels as a function of time after the release of the cultures. Each different method used to synchronize cells introduces characteristic artifacts that can persist for some time even after the stimulus is removed. Specifically, this network is originated from the use of the \( \alpha \) pheromone to arrest MAT\( a \) cells in G\( 1 \). Yeast have two mating types, \( a \) and \( \alpha \) (genotypes MAT\( a \) and MAT\( \alpha \), respectively) that can fuse to form a diploid MAT\( a \)/MAT\( \alpha \). Once a MAT\( a \) cell is exposed to \( \alpha \)-factor (the pheromone purified from the opposite mating type), the cell undergo a reversible process of differentiation of vegetatively growing cells to cells with characteristics of gametes. The cell ceases dividing and starts elongating towards the highest concentration of pheromone, forming a structure termed mating projection [26]. Yeast cells are non-motile; they have rigid cell wall and can't form filopodia like some protozoans. Thus, this chemotropic morphogenesis involves a series of cell wall modifications. Proteins involved in signaling, polarization, cell adhesion and fusion are localized to the mating projection [27,28]. On the other hand, in the yeast response pathway negative feedback loops operate at many levels to promote desensitization/adaptation and recovery. Among those negative feedback mechanisms, phosphorylation and dephosphorylation play crucial role in the modulation of signal intensity. Phosphatases and kinases operate at every level of the pathway.

The analysis of the alpha network reveals very interesting characteristics of the process going on with the studied organism. Firstly, the great majority of the genes are related to cell cycle in some way, as expected. Notwithstanding, one can discern distinct functions in different modules and most of these are directly related to the response to the pheromone, e.g. mating and cell wall remodeling. It is remarkable the presence of the clique composed by the proteins PHO5, PHO11 and PHO12. This module is formed by cell-cycle regulated transcripts peaking in the transition M/G1 of the cell cycle (Mitosis to the GAP1 phase of interphase). The acid phosphatases are
involved in the response to phosphate starvation that occurs late in the cell cycle [29] due to the consumption of inorganic phosphate for the synthesis of nucleotid and phospholipids during the metaphase. The MCM2 and MS A1 are both involved in G1/S (GAP1/DNA Synthesis) transition progression of the cell cycle [17].

The histones module comprehends mainly transcripts peaking at the early stages of cell cycle such as the MFA1, S VS 1 and S PC98 peaking in G1 and the histones themselves peaking in S phase. The histones play a crucial role in DNA replication and are thus highly synthesized during S. The MFA1 is the pheromone produced by a cells in response to mating stimulus. In our sorted time samples, it is expressed only in the very first point of the signal, maybe due to the recent stimulation with alpha-factor.

The group of genes displayed in the leftmost part of Figure 4 presents an enrichment of genes involved directly or indirectly with the cell’s rearrangements in response to mating. Most of these genes peak in M or in the transition M/G1. It’s worth stressing that such proteins are not present in any of the other studied networks. This module is connected to the other modules in the network through an endochitinase CTS 1 that is strongly cell-cycle regulated and plays a role in cytokinesis. It is also interesting to notice that this central node connects all modules through at least one transcript related to cell wall assembly or remodeling.

These results show a high level of coherence in face of the known biological processes. The subnetwork of most dependent genes modulated at the alpha-factor stimulated cell-cycle arrest shows groups of more interdependent transcripts that whether participate in a common well-defined process or happens to be expressed in the same stages of the cell-cycle. The method reveals clusters of genes cell-cycle-related or function-related and corroborate the results found in the literature.

The cdc15 network

The cdc15 subnetwork is depicted in Figure 7. In contrast to the alpha network, the cdc15 net is much denser and modules are more interconnected. The top connected vertex is NCE102 – a protein of unknown function, involved in secretion of proteins that lack classical secretory signal sequences; component of the detergent-insoluble glycolipid-enriched complexes (DIGs). According to the dendro-
gram in Figure 8, one can distinguish 6 modules in the network.

In this experiment, the authors used a temperature-sensitive mutant strain. Temperature sensitive mutants are able to grow and develop normally under a range of temperature called *permissive* but if the temperature is shifted to a so-called *restrictive* range, the organism develops the mutant phenotype and triggers a series of modifications (driven by the stress response pathways) among which is a transient cell cycle arrest. In this experiment, the authors took advantage of this genetic trait to synchronize cultures growing cells under the restrictive temperature. This procedure, however can trigger some other effects such as the stress responses induced by heat. Heat damages cells in a number of ways, perhaps most critically by disrupting the integrity of membranes and by causing proteins to denature and aggregate [30]. In response, cells induce a number of changes involving membrane fluidity and structure, an increased turnover of several plasma membrane proteins and induction of sphingolipid biosynthesis. Another characteristic feature is the strong induction of a small number of heat-shock proteins (HSPs). Their function range from the synthesis of the dissacharide trehalose, which acts as a thermoprotectant, to protein chaperones involved in protein folding, and the machinery involved in protein degradation, in particular ubiquitin [27,31,32]. Among these, the Hsp104 protein plays the most important role in yeast recovery after heat shock exposure and a number of studies shows evidence of a connection between Hsp104 content and mitochondrial activity [33]. Mitochondria are the main source of energy in cell. They are necessary not only for growth and development, but also for the repair of heat shock induced injury. Apart from the function of the mitochondrion in energy supply, this organelle seems to regulate the expression of the HSP104 gene and probably the expression of other heat shock-regulated genes in *S. cerevisiae* [33]. Evidence suggest that the hyperpolarization of the inner mitochondrial membrane by a mild heat shock is one of several signals triggering the chain of reactions that culminates in the responses described above. In the *cdc15* network one can notice the occurrence of several heat shock proteins including the Hsp104. In Figures 6 and 7 one can also notice that this protein is inserted in a group of proteins related to mitochondrial functions or mitochondrial integral membrane proteins. Another interesting feature of this network is the number of plasma membrane proteins, in particular proton ATPases, permeases and transporters. Certain inducers of stress responses such as heat

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**Figure 7**

*Cdc15 network*. The *cdc15* network. The identification of the genes and their description are given in Additional file 1.
shock permeabilizes membranes, thereby causing pronounced disturbance to transmembrane ion gradients [34]. The maintenance of the electrochemical gradient across the plasma membrane is of vital importance for the cellular functioning. Thus, a pronounced turnover of the enzymes and carrier systems associated with the membrane is expected in response to stress. This feature is being revealed in this system’s depiction.

Like in the alpha network, here we can also discern modules of genes related to cell cycle progression such as the module centered around TOS4 and its neighbor centered around AMN1 that are mostly composed by genes that peak in the G1 and S phases. The striking predominance of cell-cycle related genes in these modules suggest a similar role to the neighboring uncharacterized proteins.

**Conclusion**

Temporal gene expression data are hard to obtain and thus the design of experiments with a large set of time samples is frequently unfeasible. Instead, the majority of experiments are performed with as few as three or four time samples usually taken for as many experimental conditions. Notwithstanding the statistical poorness of the data, researchers are used to extract important clues about the biological systems by simply applying classical clustering techniques [35,36]. The availability of a more sophisticated approach could allow one to further explore the data saving time and resources in additional experimental procedures and generating more robust hypotheses to be tested. The depiction of biological systems as graphs is becoming increasingly present in the literature. The possibility of eliminating indirect interactions and to evaluate connections among groups of genes makes this a very powerful tool. Furthermore, it has been shown that cen-

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**Figure 8**

*Cdc15 dendrogram*. The dendrogram resulting from the community detection applied to the cdc15 network. The colours of the leaf nodes correspond to the described protein functions: **Red**: Response to stress – Proteins involved in stress response; **Yellow**: Cell Wall – Proteins related to cell wall remodeling (cell-cycle regulated); **Violet**: Membrane – Membrane integral proteins; **Green**: Mitochondria – Mitochondrial proteins; **Blue**: Chromatin – Proteins required for chromatin assembly and chromosome function; **Light Green**: Cell Cycle – Proteins involved in regulation of cell cycle progression; **Brown**: Other – several unshared biological functions; **Grey**: Unknown – Uncharacterized ORFs or proteins with unknown biological function.
trality measures, for instance the most connected nodes, can identify gene products that play crucial roles in the analyzed processes and that these measures does not correlate with the mean expression level [37], thus rendering unparalleled informations.

Our objective in this work is to propose a metric that makes viable more robust analyses of data where standard well-established techniques are not appropriate due to the statistical constraints. Log-likelihood scores are a common tool in the inference of correlation between time series [38] that needs no prior assumptions about the functions governing it and is very well suited for the analysis of systems where little is known about it. Here we use the information contained in different experiments about how the modulated genes behave relative to each other to derive our hypothesis of association among variables.

One point to argue in this paper is the disregard of temporal correlations. Indeed, microarray time series are known to be autoregressive and good results have been obtained with linear regression models in the context of gene networks [39-41], originating directed structures where a causal relationship is implied. However, the computational cost for this kind of approach can be restrictively high and some prior assumptions have to be made concerning the regression function and the number of variables to fit the model. In this work we are proposing a more simple yet sophisticated approach. Given the nature of our data and our purposes, we chose to disregard time correlations.

Despite the simplification, our results show a very good support from biological meanings. We stress that although our method is not based on the presence or absence of oscillatory patterns, it is applicable in cases when oscillatory patterns can be assumed negligible due to the difference in time scale with the known rhythms or asynchrony of the cell culture, for instance.

We showed results for the yeast *S. cerevisiae*, a well-known biological system and discussed two very diverse cellular processes based on the networks results. The proposed $S$ score allowed us to infer subnetworks of strongly dependent genes that show great biological relevance. Although the data set used here has been constructed from a large temporal series with a good time resolution, we have designed it in a way to disassemble time correlations and to impoverish statistics to an extreme condition. Even so, we were able to explain, in some depth, non-trivial biochemical processes related to the synchronization procedures. We have also applied the same algorithm to the original complete data set and the results are stunningly robust. The networks originated from the adapted subsets seem to be samplings of the complete ones with only edges rearranged (although around 10% of the edges are conserved).

Our results showed that the co-expression networks constructed from this kind of poorly temporally resolved microarray data are not appropriate to reveal physical interactions. However, it can provide a reliable tool to infer pairs of genes regulated by common transcription factors and thus extract valuable clues about how the system works. Finally, we would like to stress that this method can be applied to arbitrarily complex organisms and does not rely on prior knowledge. Likewise, it is also well suited for the analysis of data series extracted from any complex system such as social relationships or economics.

**Authors’ contributions**

MGC: Conducted research, designed the study, designed the algorithm, performed literature validation, wrote manuscript. FMS and IM: Participated in design of the study, participated in literature validation. OK: Participated in the theoretical discussions. CABB: Statistical discussions and revisions, wrote manuscript. GHG: Supervised research, wrote manuscript. All authors read and approved the final manuscript.

**Additional material**

**Additional file 1**

Annotations. A list with the genes’ names and functions obtained from the SGD (Saccharomyces Genomic Database) for all the genes presented in the networks described in the text. Click here for file [http://www.biomedcentral.com/content/supplementary/1471-2105-10-270-S1.xls](http://www.biomedcentral.com/content/supplementary/1471-2105-10-270-S1.xls)

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