Comprehensive Transcriptional Analysis Reveals Gene-specific Transcriptional Variations in a Seed Plant, *Arabidopsis thaliana*

**Kohei Negishi**¹ and **Kengo Morohashi**¹,²

¹Department of Applied Biological Science, Faculty of Science and Technology, Tokyo University of Science, 2641 Yamazaki, Noda, Chiba 278-8510, Japan
²Department of Biochemistry and Molecular Biology, Michigan State University, Lansing, MI 48823, U.S.A.

Keywords: Systems Biology, Genomics, Gene Expression, Stochastic Variation.

Abstract: The multicellular biological organism comprises a number of cells connected, while each cell independently works. It seems to have a system to orchestrate a number of cells, like a parallel multi-agent intelligent system. In such a biological system, gene expression of even identical genes within homogeneous cell populations is varied due to a stochastic fluctuation of the transcriptional process. This gene expression variation (GEV) is observed in development, cell differentiation, and environmental responses. Although the GEV has been generally reported, a gene-specific GEV remains unclear. Using publicly available genome-wide gene expression data from a model plant, *Arabidopsis thaliana*, we successfully identified two groups of genes whose GEVs demonstrated consistently high and low. Analysis of 632 experimental conditions derived from more than 1,300 microarrays revealed that 65 and 296 genes had high and low GEVs, respectively. We named genes with the high GEV DOTABATA (DTBT), which means “romping about” in Japanese, and genes with the low GEV PISHIPASHI (PSPS), which means “over-discipline” in Japanese. Gene function enrichment analysis resulted that DTBT genes significantly enriched stress response genes. Our results suggest a gene-specific GEV, and the regulation of GEV would be involved in biological processes.

1 INTRODUCTION

The multicellular organism comprises a number of cells connected, while each cell independently works. The connection is often dynamic and appropriately responds to environmental stimuli. In particular, a plant adapts to environments where it grows. Since a plant does not have a central nervous system, plant cells should be autonomously organized by one another. This phenomenon inspires us to a parallel multi-agent intelligent system.

Gene expression is a fundamental process for various biological events such as development, homeostasis, and response to environmental stresses in biological organisms. The amount of gene expression products shows a variation due to a consequence of stochastic fluctuations occurred during a process of gene expression (Elowitz et al., 2002), which is called gene expression variation (GEV).

The GEV produces a phenotypic variation that benefits or hampers for fitness; therefore, living organisms seem to utilize the GEV for an adaptation (Fraser and Kaern, 2009; Raj and van Oudenaarden, 2008). The GEV is considered as a passive stochastic fluctuation, and the degree of the GEV is uniformly based on Gaussian distribution. However, regulation of gene expression is orchestrated by a gene regulatory network, which often plays a role as noise reduction or amplification (Chalancon et al., 2012). Since such modulation of the GEV by the gene regulatory network requires high energy cost (Lestas et al., 2010), a decisive role for the positive contribution of the GEV is postulated.

The GEV is widely observed from prokaryotes and eukaryotes and from unicellular to multicellular organisms. Recent studies have shown the plant's phenotypic variations, such as epidermal cell division timing in the sepal and phyllotactic patterning (Araújo et al., 2017; Besnard et al., 2014; Meyer et al., 2017). If the GEV gives a stochastic fluctuation to the entire system, the GEVs of entire genes would be equally distributed. However, the degree of GEV of each gene remains unclear.

We hypothesize that plants have a system to manage the GEV in each gene with different degrees.
To identify the genes with different degrees of the GEV, we applied a statistical approach. *Arabidopsis thaliana* is an excellent model plant that provides tremendous amounts of data, particularly omics data. More than 1,000 accessions that are adapted to their environments all over the world are available. *A. thaliana* is a valuable resource to integrate information from the molecular level to a worldwide scale. By using such publicly available expression data, we successfully identified the genes with high and low GEVs. Moreover, we found that variances of the gene expression correlated with the gene functions and some ranges of temperature shift within a day and a year.

2 MATERIALS AND METHODS

2.1 Screening of PISHIPASHI and DOTABATA Genes and Gene Functional Enrichment Analysis

The Arabidopsis Information Resource 10 transcriptome (TAIR10) annotation was used in this analysis. Genome-wide gene expression data of *Arabidopsis thaliana* was retrieved from ATTED (http://atted.jp/download.shtml). The data contain 22,591 genes and 631 experimental conditions and include the following accessions: Bay-0, C24, Col-0, Cvi-0, Est, Kin-0, Ler-2, Nd-1, Shahdara, and Van-0. Out of 631 conditions, 582 conditions involved greater equal than two replicates (maximum four replicates). We used genes which expression levels were higher than an average of entire genes, resulting in 2,008 genes. Firstly, a variance of a gene $g$ in a condition $c$, referred to $\text{conV}_{g,c}$, was calculated as followed.

$$\text{conV}_{g,c} = \frac{1}{r} \sum_{i=1}^{r} (x_i - \bar{x})^2$$

where $x$ is the normalized expression value of the gene $i$, and $r$ is the number of replicates in each condition $c$. In this study, the number of the genes and the conditions were 2,008 and 582, respectively. Average value of variances in each gene, referred to $\text{aveV}$, was calculated as followed,

$$\text{aveV}_{g} = \frac{1}{n} \sum_{i=1}^{n} \text{conV}_{g,i}$$

Then, an average of entire variances was calculated as followed.

$$\text{aveV}_{all} = \frac{1}{m} \sum_{j=1}^{m} \text{aveV}_{g}$$

where $m$ is the number of genes. By comparing $\text{aveV}$ with $\text{aveV}_{all}$, we figured out how the gene expression was fluctuated. $\text{VoV}_{g}$, which is the variance of $\text{conV}_{g}$, was calculated as

$$\text{VoV}_{g} = \frac{1}{n} \sum_{i=1}^{n} (\text{conV}_{g,i} - \text{conV})^2$$

To identify DTBT and PSPS genes, the following criteria were applied. In the case of DTBT, $\text{aveV}_{g}$ was lower than $\text{aveV}_{all}$, and $\text{VoV}_{g}$ was higher than one third of $\text{VoV}$.

Gene functional enrichment analysis was performed by using Cytoscape with BiNGO plugin. TAIR10 *Arabidopsis thaliana* gene annotation was used.

2.2 RNA Extraction and Quantitative Reverse-transcription PCR (qRT-PCR)

*A. thaliana* accessions (Col-0, Est, Kin, Cvi, and Van) used in this study were obtained from *Arabidopsis* Biological Resource Center (Ohio State University, Columbus, OH, USA). Seeds were sterilized in a solution of 50% commercial bleach (Kao, Singapore) containing 6% sodium hypochlorite for 6 min, followed by three washes with distilled water. Sterilized 0.1% agarose solution was added to the sterilized seeds, which were laid out on sterile filter paper or 50% Murashige and Skoog (MS) plant salt mixture (Wako Pure Chemical Industries, Osaka, Japan) with 1% sucrose (Wako Pure Chemical Industries) and 6% gellan gum (Wako Pure Chemical Industries). The extracted RNA was digested with DNase I (Sigma-Aldrich, St. Louis, Missouri) and reverse transcribed using ReverTra Ace® qPCR RT Kit (TOYOBO, Osaka, Japan). The synthesized cDNA was amplified by qRT-PCR using the THUNDERBIRD® SYBR® qPCR Mix (TOYOBO, Osaka, Japan). Transcripts were quantified using the $\Delta \Delta C_{t}$ method.
3 RESULTS

3.1 DOTABATA (DTBT) and PISHIPASHI (PSPS) Genes Constantly Show High and Low GEVs, Respectively

The dataset we used in this analysis consists of more than 1,388 publicly available microarray data derived from 632 conditions. Out of them, 582 conditions, each of which consists of replicates, were selected. A gene expression variation (GEV) in experimental replicates is supposed to come from biological and technical variations. Since the data were collected from diverse environmental conditions, the effect of technical variations was expected to be much less than those of biological variations (McCarthy et al., 2012). Therefore, GEVs in this analysis are most likely to be derived from the biological variations but not technical ones. A low expression gene tends to show high variation; therefore, before calculating the variances, we assessed a relationship between expression level and GEV to investigate whether low expression genes show high variances. Indeed, the plot of expression levels and variances of genes suggests that genes with low expression levels show high variances (Fig. 1). Therefore, to eliminate the variances due to expression level per se, we selected the genes with a higher level of expression than the average of overall genes, resulting in 2,008 genes for further analyses.

As a workflow shown in Fig 2A, we calculated the variance of each gene from a single condition, referred to $\text{conV}$. The $\text{conV}$ is likely to show a stochastic variation of gene expression from biological replicates. Then, we calculated the average variance of the gene, referred to $\text{aveV}$. The $\text{aveV}$ demonstrates how the gene expression was varied compared to entire genes. Finally, we calculated the variances of $\text{conV}$s, referred to $\text{VoV}$. The $\text{VoV}$ shows how the GEV was constant. When the $\text{VoV}$ is low, the GEV is likely to be constant, regardless the degree of $\text{conV}$. Fig 2B shows a scatter plot of logarithm $\text{aveV}$ and $\text{VoV}$. Those two factors tend to show a positive correlation. We attempted to find the genes with constant $\text{conV}$, instead of just high $\text{conV}$, caused by a particular condition such as environmental stimuli. To search the genes with constant $\text{conV}$, we set criteria combined of $\text{aveV}$ and $\text{VoV}$ as followed. As the first criterion, we tried to eliminate variations derived from specific conditions, we selected the genes with lower $\text{VoV}$ than average $\text{VoV}$ of entire genes (Fig 2B).

We divided those genes into two gene groups with high and low constant variances based on average of $\text{conV}$s (Fig 2B). As a result, we identified 65 and 279 genes with constantly high and low GEVs, respectively.
respectively (Fig 2B). We named the high variation genes as DOTABATA (DTBT) after romping about in Japanese and the low variation genes as PISHIPASHI (PSPS) after over-discipline.

3.2 No Significant Difference in the Gene Structures of DTBT and PSPS Genes

We looked into a difference between DTBT and PSPS besides variances, and we concluded that gene structures in the genome do not contribute GEVs. Loci of both DTBT and PSPS genes are evenly located throughout the *A. thaliana* genome, and we did not find any significant difference as to location in the genome. To see whether or not GEV might be affected the length of the genic region, we compared the gene length of DTBT, PSPS, and other genes based on TAIR10 gene model. As a result, the gene length did not show significant correlations with variances (Fig 3).

Since gene expression is often affected by a position in the genome, we measured a gene density of around where DTBT, PSPS, and other genes are located. The results show no significant difference (Fig 4). Also, when we compared the length of untranslated regions (5’UTR and 3’UTR) among DTBT, PSPS, and other genes, we did not find any significant difference (Fig 5). Taken together, the genome structure around DTBT and PSPS genes is unlikely to contribute to the characteristics of DTBT and PSPS genes.

3.3 GO Enrichment Analysis Demonstrates Stress Response and Housekeeping Genes in DTBT and PSPS Clusters, Respectively

To assess gene functions of DTBT and PSPS genes, we performed GO analysis, for which we calculated enrichments of gene functions in DTBT and PSPS genes. As a result, stress response genes were statistically enriched in DTBT genes, whereas housekeeping genes were enriched in PSPS genes ($P<0.001$). In DTBT, the most enriched gene function was a stress response. Other highly enriched gene functions were also involved in response to abiotic stress, such as temperature. On the other hand, PSPS gene group enriched functions as a cellular metabolic process such as proton transport, which functions to maintain metabolic state in a cell (Fig 6).
The result that DTBT genes enriched stress genes brought us to wonder if our analysis might not work since we attempted to exclude specific responses, mainly from stress. We experimentally validated DTBT and PSPS gene expressions. Six seedlings were grown on a plate under a mild condition, followed by RNA extraction from an individual plant and measurement of the representative of DTBT and PSPS gene expression. Figure 7 clearly show that the DTBT genes were varied even in a homogenous condition, whereas the GEVs of the PSPS genes were constant, suggesting that while DTBT genes enriched stress-related genes, expression of DTBT genes are constantly varied without certain stimuli.

4 DISCUSSION

This report successfully found the gene-specific GEV even under a homogenous condition. We considered the GEV as variations of transcript level in a genome-wide manner. Living organisms need to gain fitness in their surrounding environments. In particular, unicellular organisms often appear phenotypic variations. Since phenotype originated from how genes involved in the phenotype are expressed, phenotypic variations are partly related to the extent of the number of transcripts derived from those genes. Therefore, even though thermodynamic fluctuations cause GEV, living organisms might utilize such stochastic fluctuations in their fitness and eventually their adaptation to their surrounding environments. This hypothesis is not applied to only unicellular organisms. In particular, a plant must adapt to its environment, which is required more than animals that can move. We hypothesized that plants have a system to manage fluctuations of gene expression.

One can argue that the variances we calculated were derived mainly from technical noises. However, it is unlikely to be the case for the following reasons. We chose substantial experimental conditions (582 conditions). These data set were derived from unbiased research groups, and experimental conditions were also unbiased. Therefore, differences of values among replicates are likely to be from biological ones, not technical ones. Gene expression and variances are indeed relatively correlated, particularly in the case of low gene expression. Therefore, we used genes which expression levels were higher than the average of entire genes. Nevertheless, we observed that $\text{aveV}$ and $\text{VoV}$ were positively correlated, suggesting a need for further work to eliminate the effects of expression-variance relationships.

If a gene responds to some stimulus, its expression should be changed. Thus, when we compared gene expression levels between unstimulated and stimulated conditions, we see the difference, suggesting the variances higher than non-responding genes. Since we considered a constant GEV, we ignored those genes. To do it, we calculated the variance of variances. The $\text{conV}$ from a single condition indicates to what extent the gene expression fluctuated. The variance of the $\text{conV}$, $\text{VoV}$, shows how the fluctuations of gene expression are constant. The high $\text{VoV}$ might indicate stimulus-responding genes since the gene expression variation differed in experimental conditions. We tried to find the genes that expression variation is constant; we defined a cutoff $\text{VoV}$ as the average $\text{VoV}$ from entire genes. Even...
the genes show stable GEVs; they can be classified based on a degree of variations. One is constantly high variation, and the other is constantly low. DTBT gene is former; PSPS is the latter.

5 CONCLUSIONS

In conclusion, our computational analysis using publicly available large data sets explored that the GEVs were observed in a gene-specific manner. This study suggests that plants would manage a stochastic fluctuation for their adaptations. In future work, we plan to elucidate a mechanism of DTBT and PSPS gene regulations. These findings would contribute to the biological field, such as a phenotypic variation, and the artificial intelligence field, such as a super distributed and multi-agent intelligent system.

ACKNOWLEDGEMENTS

We thank S. Yasui (Tokyo University of Science) for providing useful suggestions. We also thank S. Okamoto for providing invaluable assistance in conducting the experiments.

REFERENCES


