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Manipulating Fibroblast Environment to Study Specific Gene Expression

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Authors

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Abstract

We investigated a model system for cardiac fibrosis. Cardiac fibrosis is the thickening of the heart wall due to the inappropriate proliferation of cardiac fibrosis. Cardiac fibrosis is the thickening of the heart wall due to the inappropriate proliferation of cardiac fibrosis. To understand how the cells, respond to stress, we analyzed changes in gene expression. Our research imitated the stress conditions that the heart cells experience. We chose to analyze genes that have not previously been characterized under uniaxial, biaxial and stress-free environments to look at how gene expression varies under different conditions. We normalized all data to a validated housekeeping genes. This research will help people with various heart problems in repairing damaged tissue. We expect to increase the understanding of the cause of cardiac fibrosis and contribute to a solution. Our conclusions will compare gene expression during healthy conditions to damage repair conditions.

Introduction

Genes: CD44, CDH2, ECM1, EMILIN1, FN1, ITGA2, ITGA3, ITGA5, ITGAE, ITGAL, ITGAV, ITGB3, NCAM1, were observed under stress free conditions on tissue culture plastic, uniaxial stress and biaxial stress to analyze gene expression a 1 week, 2 week and 3 week period. Each conditions had three specimens. By observing which genes were expressed we can compare the expression in different conditions. At times cells responded to the different stresses by decreasing or increasing the level of the gene expressed. We are trying to imitate more accurate physiological relevance of fibroblasts.

Materials and methods

Materials used where C2C12 cells, which are mouse thigh muscle cells. Genes were watched for one, two and three weeks under different conditions: stress free on tissue culture plastic, uniaxial stress and biaxial stress.

Data of expressed genes was analyzed through Microsoft Excel.



Biaxial Stretch



Literature cited

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Cornell and Trevor Lujan.

Conclusions

From these results we can better understand how different genes are expressed under different stress conditions as a way to better identify what is happening in certain situations. We can see how a gene is expressed and try to pinpoint the way that gene is being manipulated by comparing it to the genes found from this experiment. We can also see how observing a gene on tissue culture plastic, which is traditionally how people look at cells, varies from how genes actually behave in cells when they are in ones body. With this data we can also conclude that with studying mutations in genes you can compare the level of expression to see how they correlate to the environment they were placed in.

Further information

- FN1 cell adhesion and migration
- ITGA2 cell and platelet adhesion
- ITGA3 cell surface adhesion molecules
- ITGAE adhesion
- ITGAL intercellular adhesion
- ITGA5 surface adhesion and signaling
- ITGB 3 cell adhesion and surface mediated signaling
- CDH2 development of nervous system, formation of cartilage and bone
- EMILIN1 development of elastic tissues
- CD44 cell-cell interactions, cell adhesion and migration
- ITGAV embryo implantations
- NCAM1 maintains skin integrity and homeostasis
- ECM1 maintains skin integrity and homeostasis