Data article

Expression of biomarker genes of differentiation in D3 mouse embryonic stem cells after exposure to different embryotoxicant and non-embryotoxicant model chemicals

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A B S T R A C T

There is a necessity to develop in vitro methods for testing embryotoxicity (Romero et al., 2015) [1]. We studied the progress of D3 mouse embryonic stem cells differentiation exposed to model embryotoxicants and non-embryotoxicants chemicals through the expression of biomarker genes. We studied a set of 16 different genes biomarkers of general cellular processes (Cdk1, Myc, Jun, Mixl, Cer and Wnt3), ectoderm formation (Nrcam, Nes, Shh and Pnpla6), mesoderm formation (Mesp1, Vegfa, Myo1e and Hdac7) and endoderm formation (Flk1 and Afp). We offer dose response in order to derive the concentration causing either 50% or 200% of expression of the biomarker gene. These records revealed to be a valuable end-point to predict in vitro the embryotoxicity of chemicals (Romero et al., 2015) [1].

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Specifications Table

Subject area Toxicology, embryotoxicity, developmental toxicity

ARTICLE INFO

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There is a necessity to develop in vitro methods for testing embryotoxicity (Romero et al., 2015) [1]. We studied the progress of D3 mouse embryonic stem cells differentiation exposed to model embryotoxicants and non-embryotoxicants chemicals through the expression of biomarker genes. We studied a set of 16 different genes biomarkers of general cellular processes (Cdk1, Myc, Jun, Mixl, Cer and Wnt3), ectoderm formation (Nrcam, Nes, Shh and Pnpla6), mesoderm formation (Mesp1, Vegfa, Myo1e and Hdac7) and endoderm formation (Flk1 and Afp). We offer dose response in order to derive the concentration causing either 50% or 200% of expression of the biomarker gene. These records revealed to be a valuable end-point to predict in vitro the embryotoxicity of chemicals (Romero et al., 2015) [1].

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More specific subject area: Alternative testing methods, embryonic stem cell, cell differentiation

Type of data: 2 Tables, 1 Scheme and 7 figures

How data was acquired: Quantitative real time PCR (StepOnePlus Real-Time PCR System (Applied Biosystems) equipment)

Data format: Analyzed and plotted data

Experimental factors: D3 mouse embryonic stem cells under spontaneous differentiation were exposed during 5 days to several concentrations of the model embryotoxic chemicals.

Experimental features: After exposure cells were lysed, mRNA was extracted and the expression of the biomarker genes analyzed

Data source location: Elche, Alicante (Spain)

Data accessibility: Data is provided in the article

Value of the data:

- We offer to the readers examples of widely used chemicals with different in vivo embryotoxicity potency.
- We offer to the readers the primer sequence and their respective annealing temperatures to assay using Power SYBR Green methodology the quantitative real time PCR the expression of 6 different genes (5 biomarkers of differentiation plus a house-keeping).
- We show as the treatments do not affect the expression of the house-keeping gene, which is an unavoidable requirement for validating the quantification of gene expression.
- We show doses–responses of model chemicals that allow deriving the concentrations causing either 50% or 200% of expression of the biomarker genes.

1. Data

We needed to select model chemicals with different embryotoxicity in order to develop a cellular method for testing embryotoxicity based on the alterations of the differentiation of D3 mouse embryonic stem cells. We finally selected our model chemicals (Table 1) among those that were previously used in the pre-validation or validation study of an embryonic stem cell method sponsored by the European Union Reference Laboratory for Alternatives to Animal Testing and by other papers dealing with the development of in vitro methods for testing embryotoxicity [2–4].

We needed to assay the effect of the selected chemicals (Table 1) on the alterations of D3 cells monitoring changes in biomarker genes. For that, we used quantitative PCR with Power SYBR Green methodology for 5 biomarker genes (plus in house-keeping gene). We designed for this purpose the primers shown in Table 2. Table 2 is also displaying annealing temperatures of such primers.

In order to check if the chemicals alter the expression of the house-keeping gene (β-actin) we determined that there were no statistical significant differences among the number of thermal cycles of control samples and samples exposed to all the tested concentrations of all model chemicals listed in Table 1 (Scheme 1). These findings are needed in order to validate further results with the biomarker genes.

We determined the effect of gene expression of biomarker genes of all the selected model embryotoxicants (Figs. 1–7). The dose–response plots were used to derive ECD50 or ECD200, which
were used as end-points for enhancing the performance of embryonic stem cell methods for testing embryotoxicity [1].

2. Experimental design, materials and methods

D3 cells cultured on monolayer under spontaneous differentiation were exposed to several concentrations of the strong embryotoxicants 5-fluorouracil (Fig. 1) and retinoic acid (Fig. 2); of the weak embryotoxicants 5,5-diphenylhydantoin (Fig. 3), valproic acid (Fig. 4) and LiCl (Fig. 5); and of the non-embryotoxicants saccharin (Fig. 6) and penicillin G (Fig. 7) for 5 days. At the end of exposure, cells were lysed, RNA was extracted and retrotranscribed to cDNA, and each gene was amplified and quantified by quantitative real time PCR as previously described [1,5,6] and using the 2^−ΔΔCt calculations [7] and β-actin as a house-keeping control gene. The expression of each gene was normalized against the expression of this same gene in the control (non-exposed) cells. The mean ± s.d. of three independent biological replicates run in the experiment is shown. (* = statistically different form control for at least p < 0.05 in Dunnett test; ** = statistically different form control for at least p < 0.01 in Dunnett test.)
Scheme 1. CTs recorded during actin expression. It is displayed number of number of thermal cycles needed to reach the threshold of fluorescence previously set during quantitative real time PCR experiments. It is displayed for each treatment the mean ± s.d. for three biological replicates.
Fig. 1. Effect of 5FU on the expression of biomarker genes.
Fig. 2. Effect of RA on the expression of biomarker genes.
Fig. 3. Effect of DPH on the expression of biomarker genes.
**Fig. 4.** Effect of VA on the expression of biomarker genes.
**Fig. 5.** Effect of LiCl on the expression of biomarker genes.
Fig. 6. Effect of saccharin on the expression of biomarker genes.
Fig. 7. Effect of PG on the expression of biomarker genes.
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