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[1]吴畏,徐海伟,屈娅,等.微流控电融合芯片的研制及其在体细胞重编程中的应用[J].第三军医大学学报,2013,35(01):29-33.

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Title: Development of microfluid cell electro-fusion chip and its application

in somatic cell reprogramming

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关键词: 电融合; 体细胞重编程; 微流控芯片; 胚胎干细胞

electro-fusion; somatic cells reprogramming; microfluid chips; embryonic stem Keywords:

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摘要: 开发高通量的细胞电融合平台,并通过此平台将小鼠胚胎干细胞 (mouse

> embryonic stem cells, mESCs) 与体细胞融合, 探讨融合细胞的多能性。 本实验室自主研发微流控芯片,将转染了绿色荧光的MESCs与转染了红色荧光的NIH3T3 细胞进行电融合,并计算其排队率与融合率。通过流式细胞仪筛选出表达两种荧光的融 合细胞,观察融合细胞的表型。RT-PCR检测NIH3T3、mESC以及融合细胞的多能性基因 (Nanog、OCT4、SOX2和LIN28) mRNA的表达水平。 结果 自主研发了微流控 芯片,构建了高通量的细胞电融合平台,通过此平台将mESCs与NIH3T3进行电融合,其 排队率为(44.35±10.99)%, 融合率为(59.88±20.03)%, 融合细胞可形成类似于 ESC样的克隆。RT-PCR法结果显示mESC和融合细胞均表达多能性基因Nanog、OCT4、 SOX2和LIN28 mRNA, 而NIH3T3不表达。 结论 通过自主研发的微流控芯片,可

实现mESCs与NIH3T3的高效融合,使体细胞重编程为多能性干细胞。

To establish a high throughput cell electro-fusion platform, by Abstract: Objective

> which to fuse mouse embryonic stem cells (mESCs) with somatic cells, and to explore the pluripotency of the fused cells. Methods With the microfluid chip developed by our lab, we fused the mESCs carrying GFP with

工具/TOOLS

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NIH3T3 cells carrying RFP, and explored the aligment efficiency and electrofusion efficiency. The fused cells were sorted by flow cytometery (FACS) and observed under a microscope. The mRNA levels of Nanog, Oct4, Sox2 and Lin28 in the three types of cells were detected by RT-PCR. Results A high throughput cell electro-fusion platform was developed. The aligment and electrofusion efficiencies of mESCs and NIH3T3 fused cells were $(44.35\pm10.99)\%$ and $(59.88\pm20.03)\%$, respectively. The fused cells can form embryonic stem cell-like clones. The RT-PCR results showed mESCs and fused cells rather than NIH3T3 cells could express Nanog, Oct4, Sox2 and Lin28. Conclusion Using the microfluid chip developed by our lab, the mESCs and NIH3T3 cells can be fused efficiently to reprogram somatic cells as pluripotent stem cells.

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吴畏,徐海伟,屈娅,等. 微流控电融合芯片的研制及其在体细胞重编程中的应用[J].第三军医大学学报,2013,35(1):29-33.

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