3D cardiomyocyte-based biosensor with tissue engineering scaffold and microelectrode array

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Abstract:
Polylactic acid (PLA) and polycaprolactone (PCL) were selected as materials to fabricate tissue engineering scaffolds by 3D printing and electrospinning, which were used to culture cardiomyocytes of neonatal rats. Then the scaffolds with cardiomyocytes were coupled with microelectrode array (MEA) to form a 3D cell-based biosensor, which was used to detect the extracellular field potential (EFP) of cardiomyocytes. The experimental results demonstrated that cardiomyocytes adhered and grew well in scaffolds, and could drive fibers to produce combined beating due to the excitation-contraction coupling. After 48 hours, the beating rate of cardiomyocytes in the scaffolds tended to be stable. The detecting results demonstrated that scaffolds and MEA were coupled well to be a 3D cell-based biosensor system, which could detect the EFP of cardiomyocytes in scaffolds with stable and high-SNR signals, and the EFP amplitude and firing rate are both similar to the signals recorded from traditional 2D culturing method.

Key words: 3D cell-based biosensor; tissue engineering scaffold; microelectrode array; cardiomyocyte; extracellular field potential.

Introduction
Scaffolds, a very important part of tissue engineering, provide an ideal environment for cell attachment, migration and proliferation [1]. At present, there are many technologies for manufacturing tissue engineering scaffolds. Electrospinning, which can continuously produce nanometer or submicron Microfiber, has become the most important method of making tissue engineering scaffolds [2].

Microelectrode array (MEA) is one of the most commonly used cell-based biosensor, which can record extracellular potential signals of Cardiomyocytes or neurons [3]. Traditionally, to build a cell-based biosensor, cells are directly cultured on the sensor chip. However, this simplified method is difficult to reflect true response for lacking of the characteristics of in vivo 3D tissue [4].

In this study, we combined tissue engineering scaffolds and MEA chip to construct 3D cell-based biosensor, which can form a 3D cell sensing system to mimic the biological in vivo environment and detect extracellular field potential of cardiomyocytes.
Methods
Polyactic acid (PLA) and polycaprolactone (PCL) were selected as materials to fabricate tissue engineering scaffolds by 3D printing and electrospinning. Then the cardiomyocytes of neonatal rats were cultured in scaffolds. The scaffolds with cardiomyocytes were coupled with microelectrode array (MEA) to form a 3D cell-based biosensor, which was used to detect the extracellular field potential (EFP) of cardiomyocytes in scaffolds (see Fig. 1).

Results
The experimental results demonstrated that cardiomyocytes adhered and grew well in scaffolds, and could drive fibers to produce combined beating due to the excitation-contraction coupling. The beating rate of cardiomyocytes tended to be stable after 48 hours and reached (34.25±3.3) times/min after 72 hours. After coupling cardiomyocytes scaffold with MEA chip, 3D cardiomyocyte-based biosensor could record the different EFP patterns of cardiomyocytes in scaffolds with stable and high-SNR signals (see Fig. 2).

Discussion
EFP signal patterns were related to pacemaker and cell status of cardiomyocytes in scaffold. Before cardiomyocytes in scaffold fused fully to form synchronous beating, there may be multiple pacemaker points, so the sensor would record the EFP signals from several pacing points simultaneously. The more active the cardiomyocytes in scaffold behave, the larger EFP signal amplitude and the higher beating rate will be.

EFP signals of cardiomyocytes cultured in planar sensors and tissue engineering scaffolds are recorded respectively (see Fig. 3a). The amplitude and firing rate of EFP signal were statistically analyzed (see Fig. 3b-3c). The EFP signal amplitude of cardiomyocytes in 2D culture environment was (228.9±6.9) μV, and the firing rate was (34.2±4.6) times/min. Similarly, in 3D scaffold culture environment, EFP signal amplitude was (222.6±4.1) μV, and the firing rate was (33.9±8.4) times/min. Significant difference analysis was conducted with t test, and it was found that the EFP signals recorded in the two culture environments showed no significant difference in the amplitude (p>0.05, n=29) and firing rate (p>0.05, n=10).

In conclusion, this study provides a new method to construct 3D cell-based biosensor, which could detect the EFP of cardiomyocytes in scaffolds with stable and high-SNR signals. Cells in scaffolds can mimic the in vivo environment and has more physiological significances. This study is expected to be applied in the field of cardiac drug assessment and tissue engineering.

Fig.3. (a) EFP signals of cardiomyocytes cultured in planar sensors and scaffolds; (b) EFP firing rate in two modes of cultivation; (c) EFP amplitude in two modes of cultivation

References