Prediction of therapeutic effects of human cardiomyocytes in myocardial infarction using non-human primates model

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Cardiovascular disease (CVD) is a critical disease that accounts for 34% of all deaths and consumes almost $300 billion annually in the United States (1). Since myocardial infarction (MI) is the most severe and serious CVD with high lethality (2), attempts to overcome the disease are highly demanded. Although the development of pharmacological and intervention therapy enhances the survival rates, there are clear limitations to preventing or reversing the heart failure due to the massive muscle loss after MI. Currently, heart transplantation is considered to be the most fundamental treatment, but it also has many issues to resolve such as, complex post-operative management, very expensive strategies, and limited number of organ donors. Indeed, development of additional cardiovascular therapy is essential (3).

In spite of numerous therapeutic approaches that have been attempted using stem cells to treat CVD, few positive results and obvious limitations are reported in heart regeneration. Thus, there are crucial checkpoints to be confirmed before treating CVD through cell therapy. First, the transplanted cells must be differentiated into cardiac muscles. Second, the myotube formation with the cell graft and implanted heart, and eventually the electromechanical maturation of the graft with host myocardium must be confirmed, resulting in the transplantation of external cardiomyocytes which could restore the infarcted heart (3). When these prerequisites are confirmed, human pluripotent stem cells (hPSCs)-derived cardiac cells are ready to be engrafted into the injured heart and are expected to have a spontaneous action potential from the cell therapy (4).

In this study, Liu et al. demonstrated that transplantation of millions of cryopreserved human embryonic stem cell-derived cardiomyocytes (hESC-CMs) successfully enhances cardiac function in macaque monkeys with large MI. The primate animal is the most suitable model for predictions in a clinical study due to being evolutionary adjacent and physiological similar to humans (5). After the hESC-CMs injection in the macaque heart disease model, the authors confirmed the functional recovery and described it in two steps: (I) mechanical maturation without teratoma formation, and (II) lower burden of ventricular arrhythmia. The authors induced larger infarcts in macaque hearts and showed the restoration effect of the transplanted hESC-CMs by cardiac magnetic resonance imaging (MRI) exclusively developed for macaque monkeys, which can provide the information related to left ventricular ejection fraction (LVEF), LV mass, the myocardial motion, as well as general information, such as heart structure (6). The results showed that hESC-CM transplantation improved ~10% and ~22% in LVEF and wall thickening, respectively. The enhancement of LVEF indicates an increased amount of blood released in each contraction and the enhanced efficiency of the heartbeat for systemic circulation (7). In addition, heart wall thickening could be interpreted as the...
loss of muscle due to the infarction being partially covered. However, to explain the therapeutic effect of transplanted cells in infarcted hearts, additional aspects of confirmation in histology are needed: (I) successful transplantation in infarcted region; (II) decreased scar formation; (III) sufficient maturation to function in macaque infarction model (8).

MI leads to massive loss of cardiomyocytes in the ischemic region and fibrotic scar formation, resulting in re-expression of heart failure (9). In regard to the success of transplantation aspect, the authors observed large grafts of myocardium (5.7–15.6% of the infarcted region) in the central infarcted region, using cardiac troponin 1, and prevention of the wall thinning after 4 weeks of transplantation in the hESC-CM-treated group. Subsequently, the reduction of scar formation (from 25.2% to 19.9% of the LV in the hESC-CM-treated group compared to from 25.9% to 23.6% in the sham group) was histologically confirmed. Since the electromechanical maturation of the graft cells is very important to keep its beating regular and synchronized with the recipient’s cardiac tissue (10), the authors confirmed the formation of mature N-cadherin+ adherens junctions and Connexin 43+ gap junctions with the host heart. However, a minor limitation in the distribution of Connexin 43 remained. In addition, the authors could show vascularization with host cells by the graft using CD31+ in blood vessels, as well as proliferation of the graft cells using pericentriolar material-1 (PCM-1) and Ki67. Altogether, this data provided strong evidence that the implantation of hESC-CM reduced the fibrotic scar, filled up the muscle loss caused by the fibrosis process and could recover structural stability of the left ventricular wall. Meaningfully, the injected cardiomyocytes successfully formed a considerable maturation with the macaque infarcted heart.

The authors then investigated a new perspective on the cause of arrhythmias. An arrhythmia is an abnormal condition in which the heart contracts irregularly, and its severity presumes the degree of myocardial dysfunction. It has been known as a main limitation of cell-based therapy in heart disease because it has occurred when the external cells were implanted in an infarcted heart and has more frequently occurred in large rather than small animals (11,12). Although its mechanism is unclear, differences of electrophysiologic maturity, gap junction isoatypes, and wave propagation between the host heart and the graft are thought to be potential causes of arrhythmia (4,5). Almeida et al. raised the possibility that the newly formed pacemaker from transplanted cells could be the cause of arrhythmia formation (13). From this point of view, Liu et al. performed an electronic mapping and determined that an ectopic activation source generated in transplanted hESC-CMs improved the pulse generated by ectopic pacemaker cells. As mentioned above, since the stability is considered as the first priority in stem cell-mediated therapy (14), the authors excluded the chance of teratoma formation derived from undifferentiated pluripotent cells. In order to do that, the authors used cardiomyocytes fully differentiated by well-known methods and ultimately confirmed 86–99% of differentiation based on flow cytometry for cardiac troponin T and then confirmed the occurrence of teratoma by performing full necropsy on all animals with an experienced pathologist. As a result, teratoma was not found in any of the five hESC-CMs treated animals, except one macaque that had an extremely small carcinoma, which originated not from the grafted human cardiomyocytes but from the host macaque. It is suggested that the treatment of hESC-CM for infarction may not induce teratoma or cancer by using almost fully differentiated cardiomyocytes.

In conclusion, the authors optimized the protocol on the cell type and dose, and the application time for hESC-CMs and injected the cells into the macaque infarction model in order to introduce stem cells for cell therapy in further preclinical experiments and human clinical trials. Additionally, this study confirmed the therapeutic effects of hESC-CMs transplantation through electromechanical maturation and safety by showing the result without teratoma formation of transplanted cells. However, there is still a lot of room for improvement of cell therapy in heart disease. If the cell therapy taken in this study is tested as the first priority in stem cell-mediated therapy (14), there is still a lot of room for improvement of cell therapy in heart disease. If the cell therapy taken in this study is tested in the five hESC-CMs treated animals, except one macaque that had an extremely small carcinoma, which originated not from the grafted human cardiomyocytes but from the host macaque. It is suggested that the treatment of hESC-CM for infarction may not induce teratoma or cancer by using almost fully differentiated cardiomyocytes.
cardiomyocytes and to construct a better microenvironment in the MI region. There are still unsolved obstacles in the use of stem cell therapy in MI, but this study provided a good possibility in clinical therapy with a well-applied protocol utilizing hESC-CMs.

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**Footnote**

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**References**
