

عنوان مقاله:

New Strategy of Decellularization and Production of Human Kidney Scaffolds

محل انتشار:

سومین جشنواره ملی و کنگره بین المللی علوم و فناوری های سلول های بنیادی و پزشکی بازساختی (سال: 1397)

تعداد صفحات اصل مقاله: 1

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خلاصه مقاله:

Background and Aim: Incidence of end-stage renal disease (ESRD) is greatly increasing. Renal transplantation is one of the goal treatments for ESRD. The development of tissue engineering and regenerative medicine have catalyzed due to a gap between limited organ supply and increasing demands. Natural scaffolds prepared from an extracellular matrix (ECM), have emerged as an ideal tissue microenvironment for this goal. An important technique in regenerative medicine to prepare an acellular ECM is the decellularization of native tissues. Therefore, the aim of this study was to determine the effective method for decellularization of human kidney and producing the natural human kidney scaffold. Methods: After Nephrectomy, human kidneys that could not be transplanted were used in this study. Adipose tissue and capsule around the kidney were removed. We cut kidneys into transverse sections (approximately $10 \times 10 \times 2$ mm³ pieces) using a scalpel. Then Cortex-medulla kidney sections were washed twice with phosphate buffered saline (PBS), followed by decellularization in a solution of either 1% Triton X-100 or sodium dodecyl sulfate 1% (SDS). The sample was decellularized at 4 using shaker (200 rpm). Decellularization solution was changed 4 hours after initial tissue harvesting and then every 24 hours until tissues were transparent (for 14 days). In order to confirm decellularization, hematoxylin-eosin (H&E) staining was performed on days 2, 5, 10, and 14. Results: Comparison of H&E staining of the decellularized and native kidney tissue revealed successful elimination of cell nucleus in SDS and Triton-treated sections. Also, H&E staining revealed that in the Triton-treated sections the native ECM architecture, integration of renal vascular and glomerular structures was more preserved than the SDS-treated sections. Conclusion: We have developed an effective decellularization method for the preparation of human renal ECM scaffold. Additionally, it may be possible to use the scaffolds that prepared with 1% Triton X-100 for kidney regeneration. These results also indicate that discarded human kidneys are a suitable source of renal scaffolds and their use for tissue engineering applications may be more clinically applicable and beneficial than kidneys derived from animals.

کلمات کلیدی:

Human kidney; Regenerative medicine; Scaffold; Decellularization; 1% Triton X-100

