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# Engineered liver tissue in vitro to mimic liver functions and its biomedical applications

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The liver is an important organ with metabolic functions, and plays the important role of deoxidizing, storing liver sugar, synthesizing secretory proteins and other functions inside the body. The prevalence of liver disease has been increasing in recent years. For acute liver failure, liver cancer and other serious liver diseases, the best treatment is liver transplantation. However, still there is shortage of organs in clinical practice. Under such a severe background, it is very important to cultivate liver tissue in vitro to make up for the shortage of organs and to test the hepatotoxicity of drugs. In this review, we will introduce the structure of liver and the function of liver cells, summarize several methods of liver tissue culture in vitro, and introduce the application of liver tissue culture in vitro. Finally, we forecast the development prospects of the liver tissue.

## 1. Introduction

The liver plays an important role in deoxidizing, storing liver sugar, and synthesizing secretory proteins. 1-3 The surface of the liver is covered by a thin capsule of dense connective tissue. The capsule penetrates the liver to form a reticular scaffold that separates the liver parenchyma into many hepatic lobules, which have similar morphology and the same function. The liver is made up of 500 000 to 1 million hexagonal lobules. There is a central vein in the center of the lobule, which is surrounded by a roughly radially arranged plate of liver cells

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(hepatic lamina). The liver is also composed of different cell types, including hepatocytes, sinusoidal endothelial cells, hepatic stellate cells, Kupffer cells, biliary epithelial cells, and dendritic cells (Fig. 1).4-8 Furthermore, it is important to maintain the polarity of hepatocytes under *in vitro* culture conditions.<sup>9,10</sup> Due to the structural arrangement and cellular heterogeneity of the liver, liver cells are exposed to a gradient of nutrients, hormones, and growth factors through the combined portal vein and hepatic artery blood supply. According to estimation, there are about 90 million people who are infected with HBV (hepatitis B virus) in China<sup>11</sup> and there are 7.6 million HCV patients (4.56 million chronic patients). 12 Every year, 330 000 people die of cirrhosis and primary liver cancer. In this severe situation, it is very urgent to understand the physiological function of liver cells and the related



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Fig. 1 The structure of the liver. (A) The structure of the liver and the distribution of major liver cells. (Reproduced with permission from ref. 4.) (B) The structure of the hepatic lobule. (Reproduced with permission from ref. 26.) (C) Structural diagram of hepatic sinuses and distribution of major cells (Reproduced with permission from ref. 27.) (D) The types of cells in the liver and their relationships. (Reproduced with permission from ref. 26.) (E) The interaction between cell and cell, and the interaction between cell and the extracellular matrix (ECM). (Reproduced with permission from ref. 26.)

disease mechanism, and to provide effective treatment. Unfortunately, traditional research models have many limitations. Primary human hepatocytes cannot be cultured in vitro for a long time to maintain cell morphology and function. 13,14 The genetic metabolic mechanisms of the animal models are different from those of humans, which prevents the results from being translated into clinical practice. Therefore, more closely resembling human hepatocytes are needed for experimental and therapeutic studies.

For drug screening, some drugs will still induce human toxicity and side effects after appearing on the market, and about 50% of them are due to the liver toxicity of the drug. It is particularly important to test drugs for hepatotoxicity.<sup>15</sup> Initially, researchers used live animal models to investigate treatments or drugs for liver diseases. However, the metabolic rules of the livers of mice, pigs and other animals used in the research are greatly different from those of human livers in many aspects, and the results may not be consistent with the actual effect. In addition, using live animals as models is very



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cruel to animals, which does not meet the humanitarian spirit. The liver models for drug screening are two-dimensional monolayer cultured cells. However, the two-dimensional monolayer culture model is limited by the distribution of receptors on the surface and cannot simulate the clustering state of the cells, which greatly limits the wide application of this technology. Therefore, the newly emerging liver tissue engineering has become a research hotspot in recent years because of its cell composition and similar three-dimensional structure, which is closed to human physiology. Compared with live animal models, a liver tissue constructed in vitro has almost the same mechanism as the human liver, possessing the genetic characteristics of the individual of origin, and avoids harm to animals. Compared with the two-dimensional monolayer tissue, liver tissue constructed in vitro can maintain cell polarity for a long time, and ensure cell surface protein transmission, avoiding the loss of liver tissue function. In recent years, with the continuous development of life science, material science and engineering science, researchers have made one by one breakthroughs in tissue construction and organ reconstruction in vitro. 16-18 Many tissue engineering products have been applied in clinical medicine, such as bone tissue and skin tissue. Liver tissue engineering is an important research field in tissue engineering, whose goal is to construct a transplantable liver tissue or organoid with the function of the human original liver, 19 so that patients who have liver injury can be treated and the deficiency of the clinical organ shortage can be made up.<sup>20–25</sup>

In this review, we will roughly divide the content into five parts. First, we will introduce the structure of the liver, and the types and functions of liver cells. Then, we will outline the cells and materials that are used in liver tissue engineering. With the above content as the basis, we will present eight methods to culture liver tissue in vitro, and list their applications in tissue reconstruction and drug preparation. Finally, the prospects of liver tissue culture will be prospected.

# Cells and materials for liver tissue engineering

## 2.1 Cells for liver tissue engineering

The selection of cells is very important in the process of liver tissue culture in vitro. There are several kinds of cells that can be used to grow liver tissue: bone marrow-derived stem cells, liver oval progenitor cells, liver cell lines and mature liver cells, embryonic stem cell (ESC), induced pluripotent stem cells (iPSC), bone marrow-derived very small embryonic-like stem cells (BM VSEL), and hepatic liver stem cells (HLSCs). 4,28

2.1.1 Liver progenitor cells. Progenitor cells refer to the group of cells that are derived from stem cells, which have lost the ability to self-renew, and can differentiate into a single cell under certain conditions. Hepatic progenitor cells are closely linked with adjacent hepatocytes and bile ducts at the end of differentiation in the liver, and belong to transitional bile ducts.<sup>29,30</sup> These cells cannot necessarily differentiate into mature liver cells, but have a variety of differentiation

options. 31 For example, liver progenitor cells can differentiate into other functional cells of the liver by different cellular signals. If the ideal differentiation conditions for the progenitor cells can be controlled to enable them to differentiate into the desired mature liver cells, the application of progenitor cells in liver tissue engineering will be very promising.

- 2.1.2 Mature liver cells. Mature liver cells can be obtained by the hepatic system. 32,33 The researchers have done much work to culture mature liver cells. It mainly focuses on two aspects: one is to use the extracellular matrix components of hepatocytes and the polarity of hepatocytes to promote the growth of hepatocyte colonies, and enhance the nutrition and oxygen supply in the colonies. Secondly, liver parenchymal cells were cultured with parenchymal cells. However, liver cells are terminal cells, so it is difficult to subculture. Furthermore, primary liver cells are mostly taken from animals such as mice and pigs, which are not suitable to directly use in the human body.
- 2.1.3 Liver cell lines. Liver cell lines mainly refer to tumorderived cell lines, including HepG2, C3A or SV40Tag, which can be cultured in large numbers for a long time. These cells have strong ability to proliferate, and have the function level of liver cells, such as albumin synthesis and cytochrome P450 activity, and others. They are the most promising type of cells to be applied in clinical trials of the artificial liver. 34-36
- 2.1.4 Embryonic stem cell (ESC). Embryonic stem cells (ESCs) are a kind of cells isolated from early embryos or primitive gonads. They have the characteristics of infinite proliferation, self-renewal and multidirectional differentiation in vitro culture. ESCs can be induced to differentiate into almost all cell types of the body, both in vitro and in vivo. Therefore, ESCs have the potential to differentiate into hepatocytes. However, due to ethical issues, the clinical use of these cells has certain limitations. In general, ESCs have high telomerase activity, which can proliferate strongly in vitro, when injected into mice, causing tumor formation and ultimately cancer. Therefore, it is a challenge to keep ESCs undifferentiated and differentiating them into liver cells after injection.
- 2.1.5 Induced pluripotent stem cells (iPSC). Induced pluripotent stem cells are produced by artificially inducing non-pluripotent cells to express a particular gene. These are similar to natural pluripotent stem cells in many aspects, such as the expression of certain stem cell genes and proteins, embryo body formation, the formation of different chimeras, and differentiation potential. In 2006, iPSCs were produced from mouse embryonic fibroblasts and adult mouse tail terminal fibroblasts, and in 2007 from human cells, which allowed researchers to harvest pluripotent stem cells for research and therapy. Because iPSCs are grown from a patient's own somatic cells, iPSCs treatment will not trigger an immune response. Differentiated cells (such as skin cells) can be converted to an undifferentiated pluripotent state by reprogramming factors. These reprogrammed cells, called induced pluripotent stem cells, can be manipulated outside the body to differentiate into a variety of cells. However, it has been suggested that phenotypic differences in iPSCs reprogramming and epigenetic

memory may lead to different differentiation pathways. In addition, iPSCs cells still have the possibility of tumor formation.

- 2.1.6 Mesenchymal stem cells (MSC). Mesenchymal stem cells (MSC) are pluripotent stem cells that have all the commonalities of stem cells, including the ability to self-renew and pluripotent differentiation. They can be obtained from different parts of the body, including adipose tissue, bone marrow, and the umbilical cord. They remain undifferentiated throughout their life cycle until the body needs them. MSC are ideal seed cells for liver tissue culture in vitro because of their multidirectional differentiation potential, stable genetic background, low antigenicity, and ease of culturing and amplification in clinic.
- 2.1.7 Bone marrow derived very small embryonic-like stem cells (BM VSEL). BM VSELs are pluripotent stem cells that can be recruited from the bone to an injured liver. Although they proliferate weakly, their proliferation increases when acute liver injury occurs, and researchers have shown that they may differentiate into liver cells.
- 2.1.8 Hepatic liver stem cells (HLSCs). A small number of adult stem cells in the liver can differentiate into liver cells. These adult stem cells can turn into progenitor cells, called ovoid cells. They have morphology and function similar to embryonic hepatoblasts, which can differentiate into bile duct cells and liver cells. HLSCs have the potential to differentiate into bone, lipids, endothelial cells, insulin-producing cells and hepatocytes, and can be differentiated into hepatocytes by the use of hepatocyte growth factor and fibroblast growth factor. In acute and chronic liver disease, these ovoid cells can partially repair the liver through a process similar to liver regeneration. However, the use of these cells is limited due to their complex isolation conditions.

### 2.2 Materials for liver tissue engineering

- 2.2.1 Natural biomaterials. Natural biomaterials usually refer to natural macromolecules, which are derived from animals, plants or human bodies. Natural biomaterials are widely used to repair or replace damaged tissues and organs in the body due to their good biocompatibility, biodegradability and remodeling ability. In addition, natural biomaterials have the function of promoting cell migration, proliferation, differentiation and adhesion. At present, natural biomaterials have been widely used in liver tissue engineering.
- 2.2.1.1 Alginate. Alginate is a natural polymer found in brown algae and often made into gels, which are similar to the structure of the extracellular matrix. Alginate is already widely used in wound healing, tissue engineering and drug delivery. High purity alginate has good biocompatibility and can be used in humans. Kinasiewicz et al. coated C3A (human immortalized liver cells) with alginate and cultured them statically for 7 days.<sup>37</sup> The cells then proliferated to form multicellular aggregates. The research demonstrated that alginate-coated C3A liver cancer cells could produce albumin, which is the main serum protein synthesized by the liver. The result proved that alginate microencapsulation could promote

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three-dimensional cell interaction and significantly improve cell metabolism. Fluidization allowed for better nutrient and oxygen supply to the intracellular cells, which increased the synthesis function. Lan et al. encapsulated HepG2 cells with alginate.<sup>38</sup> They demonstrated the possibility of HepG2 cell lines that could be encapsulated in a 3D alginate for drug screening. Liu et al. used calcium alginate in the study of simulating liver lobules in vitro. 39 They proposed a method of preparing calcium alginate cell sheets by electro-deposition, which were used to form liver lobules. This method provided a new "bottom-up" paradigm for constructing three-dimensional macroscopic liver tissue similar to that of a living organism. The assembled liver lobules were expected to serve as extracorporeal models of liver organs, and would promote new applications of electro-deposition methods in tissue engineering. Masumi Yamada et al. proposed an efficient method for the extracorporeal culture of a hepatic cord structure by using anisotropic alginate saline gel microfibers to generate complex hepatic microorgans, 40,41 which were composed of primary hepatocytes and parenchymal cells in vivo by microfluidic technique. The method provided an ideal microenvironment for hepatocytes, including cell and cell matrix interactions. Jeon et al. also used alginate to generate multi-layered three-dimensional structures of HepG2 cells by bioprinting. 41 After printing, the 3D alginate hydrogel was immersed in 1% calcium chloride solution for solid crosslinking. It turned out that 3D bioprinting could be used to create the liver with 3D structures, which contributed to the stable regeneration of HepG2 cells.

2.2.1.2 Collagen. Collagen is the most abundant protein in mammals, which is widely found in all tissues from the body surface of lower vertebrates to the body of mammals. Collagen mainly exists in the form of collagen fiber in the extracellular matrix such as tissue and bone, which has high tensile strength. It can maintain and stabilize the tissue structure of the body. Brown et al. used collagen to draw colloidal islands on polystyrene plates for tissue culture. 42 The liver cells selectively attached to colloidal islands, and parenchymal cells grew on the surrounding areas. Ukairo et al. also tested the bioactivity and toxicity of acetaminophen in rats by preparing the co-culture system of liver cell micrograph. 43 Okudaira et al. prepared a fibrous liver tissue by using the bottom-up method of the multilayer spheres.44 Collagen was attached to the spherical aggregates of hepatocytes, and then fibroblasts and human umbilical vein cells were attached to the co-culture with hepatocytes. Then, 3D liver tissue was constructed by high-density stacking cells. The results showed that cells could receive adequate oxygen supply, and could maintain cell viability and liver function for a long period of time. Moreover, this method does not require specialized or expensive materials and equipment. It can also build large-scale liver tissue.

2.2.1.3 Chitosan. Chitosan is the product of the deacetylation of chitin, which mainly exists in the shells of shrimps, crabs, insects and other crustaceans. It has good biocompatibility, and has been widely used in biomedical and pharmaceutical fields.

Feng et al. investigated the effects of nanofiber galactosylated chitosan (GC) scaffolds on the formation of primary hepatocyte aggregation and the maintenance of liver function in rats. 45 By using chitosan as the raw material, the galactose ligand was successfully combined into chitosan to improve the adhesion and synergistic function of liver cells. A nanofiber galactose chitosan scaffold was prepared by electrostatic spinning technology, and the primary liver cells were attached and cultured on the scaffold. The results of this research showed that the GC nanofiber scaffolds showed good cell activity and higher liver function, and their mechanical stability was better than that of three-dimensional spherical aggregates. Therefore, GC nanofiber scaffolds could be used as biodegradable natural scaffolds for primary liver cell culture in vivo or in vitro. Li et al. prepared microcapsules that encapsulated liver cells with alginate and chitosan as raw materials.46 Microencapsulation of hepatocytes can provide a new technology for the large-scale preparation of hepatocytes, which could also keep the high activity in vitro and long-term cryopreservation of hepatocytes.

2.2.1.4 Gelatin. Gelatin is a kind of macromolecule hydrophilic colloid, which is the product of the partial hydrolysis of collagen, and has been widely used in the medical field. In 3D bioprinting technology, the use of gelatin as a substrate can control the ink thickness and high printing fitness. Mazza et al. added gelatin to the alginate, squeezed a bio-ink mixed with pluripotent stem cell differentiated liver cells at low temperatures, and solidified it with calcium chloride to form liver tissue. <sup>47</sup> Nagamoto et al. used gelatin-coated scaffolds to harvest cell sheets, which were prepared by co-culture method. <sup>48</sup> The liver cells and fibroblasts were co-cultured for 14 days, placing the gelatin-coated scaffold on the cell sheet, and then the culture temperature was lowered to 20 °C for 60 min. Due to changes in the hydrophobicity of the temperature-sensitive surface, the cell sheet adhered to the gelatin-coated layer and was harvested.

2.2.1.5 Decellularized tissue matrix. Decellularized tissue matrix is a new kind of biological material that can be used to repair damaged tissues, and regenerates by decellularizing allogeneic or heterogeneic tissues. Decellularized tissue has good biocompatibility, good mechanical strength, and can induce and promote cell adhesion, proliferation, differentiation and tissue formation. Mazza et al. used high shear stress vibrationdecellularization technology to rapidly produce functional human liver scaffolds, and then cultured liver cells on scaffolds by perfusion method.49 Alejandro et al. removed all cells from isolated rat livers by combining the use of enzymes, detergents, and mechanical methods.<sup>50</sup> The decellularized liver is examined by morphological, biochemical, and immunolabeling techniques to preserve the primary matrix structure and components. To ensure maximum cell survival, they also evaluated three different methods of hepatocyte seeding: direct parenchymal infusion, multi-step infusion, and continuous infusion, and found that multi-step infusion produced the best results.

2.2.1.6 Hyaluronic acid. Hyaluronic acid is the main component of connective tissue such as intercellular stroma, ocular

vitreous and synovial fluid of joints. Hyaluronic acid can be used as a viscoelastic agent for ophthalmic intraocular lens implantation, as a filler for osteoarthritis and rheumatoid arthritis, and is widely used as a medium in eye drops. Turner et al. used hyaluronic acid to maintain the human hepatoblast phenotype. 51 The liver cells inoculated in the hyaluronic acid hydrogel maintained the stability of the early hepatocytes throughout the culture process. Although other culture conditions, such as plastic or embryonic collagen or embryonic feed, were also available, hyaluronic acid was the only culture condition that reliably promoted the survival, proliferation and maintenance of hepatoblasts.

2.2.1.7 Matrigel. The main components of matrix glue are laminin, type IV collagen, nestin, heparin sulfate glycoprotein, growth factors and matrix metalloproteinases. At room temperature, matrix glue polymerization can form a threedimensional matrix with biological activity, which simulates the structure, composition, physical properties and function of the cell basement membrane in vivo. It is beneficial to cell culture and differentiation in vitro, and can also be used for the study of the cell morphology, biochemical function, migration, infection and gene expression. Khetani et al. prepared the MPCC platform combined with the matrigel covering layer, which was useful to further mature iHeps toward a more adultlike PHH phenotype. 52 It also maintained the function of liver cells after several weeks of culture, so it could be used for chronic drug delivery research.

#### 2.2.2 Synthetic material

2.2.2.1 Polylactic acid-glycolic acid copolymer (PLGA). Polylactic acid-glycolic acid copolymer is a kind of biodegradable polymer organic compound, which is randomly polymerized by two monomers known as lactic acid and glycolic acid. It has good biocompatibility, non-toxicity and good performance of forming capsules and films. Now, PLGA has been widely used in the pharmaceutical, medical engineering materials and modern industry fields. Jessica H. Brown et al. developed a 3D nanofiber scaffold that was made of PLGA polymer by using the newly optimized wet electrostatic spinning technology.<sup>53</sup> It had a high pore structure and could accommodate primary human liver cells, simulating the ECM structure in vivo. Subsequently, it was proved that PLGA wet electrostatic spinning 3D nanofiber scaffolds modified with type I collagen enhanced the synthetic activity of primary human hepatocytes, which might be used as the starting material for the biosynthesis of primary hepatocytes in long-term culture.

2.2.2.2 Polycaprolactone (PCL). Polycaprolactone is a thermoplastic crystalline polyester obtained by ring-opening polymerization of caprolactone with dialcohol. Because PCL has good biodegradability, biocompatibility and nontoxicity, it has been widely used as medical biodegradable materials and drug control release system. Gao et al. prepared electrospun PCL mats by using an electrospinning system.<sup>54</sup> A HepG2 cell suspension was inoculated onto the PCL scaffold. Results have shown that the morphology of the fiber can significantly affect the behavior and growth of cells, and the fiber arrangement can affect cell elongation and orientation. Slivac et al. also manufactured electrospun PCL mats, and tested the effects of PCL mats and liver extracellular matrix scaffolds on the bioactivity of HepG2 cells. 55 Kim et al. promoted the regenerative treatment of the liver injury by stacking patient-specific progenitor cell sheets onto multiscale electrospun fibers.<sup>56</sup> The results showed that the liver patch was a technique for the generation of an artificial liver, which might have potential clinical application value in the future.

2.2.2.3 Polydimethylsiloxane (PDMS). Polydimethylsiloxane is a kind of hydrophobic silicone material. Yu Du et al. introduced the process of manufacturing microfluidic chips by using PDMs as raw materials when introducing the microfluidic chips that simulate the structure and function of liver sine.39,57

Liu et al. used electrodeposition technology to prepare alginate gel cell sheets to assemble multilayer hepatic lobule models.<sup>39</sup> After the cell sheets were collected, they were transferred to PDMS molds for assembly. Khetani et al. used PDMS templates made by photolithography to draw an extracellular matrix or collagen on tissue-cultured polystyrene. 52 Hepatocytes only attached to the ECM domain and the parenchymal cells were seeded, and attached to areas not occupied by hepatocytes. Pang et al. used PDMS-based cellular microarray culture plates to construct an aggregate of endothelialized rat liver cells.<sup>58</sup> Endothelialized rat hepatocytes were aggregated in the microporous apparatus.

2.2.2.4 Polyvinylidene fluoride (PVDF). Polyvinylidene fluoride mainly refers to the partial fluorine ethylene homopolymer or partial fluorine ethylene, and other small amounts of fluoride copolymer of vinyl monomers. PVDF membranes bind proteins and can separate small fragments of proteins. Its stability and corrosion resistance make it ideal for protein sequencing, and it is still in use today. Chu et al. developed a three-dimensional liver model to test sodium nitrite and acrylamide-induced hepatotoxicity,<sup>59</sup> which forms cell cylinders in PVDF hollow fibers to mimic the microenvironment of liver tissue.

2.2.2.5 1-Polylactic acid (PLLA). PLLA is an important biodegradable polymer material, which is characterized by nontoxicity, nonirritating property, biodegradable absorption, strength, good plasticity and easy processing. Its degradation cycle is 2-12 months. The degradation cycle can also be changed according to the addition of modifiers. PLLA is decomposed by enzymes in organisms. It eventually forms carbon dioxide and water, which has good biological compatibility. PLLA can be used to prepare scaffolds in the biological field. Jiang et al. cultured infant liver cells with a 3D PLLA scaffold.60 The results showed that the PLLA scaffold is a good supporting material for infant liver cells, which can be used for in vitro culture of infant liver cells. Bierwolf et al. used PLLA as a raw material to prepare 3D nanofibers for toxicology and medical research.<sup>61</sup> The data showed that PALL nanofiber scaffolds provided a good microenvironment for the survival

and new tissue formation of primary rat hepatocytes, and retained specific enzyme functions and morphology.

# 3. Methods of liver tissue culture

### Bottom-up tissue engineering method

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**3.1.1** Micromode coculture method. In the past few years, various liver models have been developed and used to investigate the basic function and metabolism of liver cells. However, most models have some limitations in reproducing human liver biology. To overcome the above disadvantages, Bhatia et al. subsequently proposed a micromode coculture method (MPCC) to culture the liver parenchymal cells with liver nonparenchymal cells in mice (Fig. 2A). The development of MPCCs was inspired by earlier studies of physical homomorphic (hepatocellularhepatocellular) and heteromorphic (hepatocellular-stromal) interactions regulating hepatocyte function in vitro. 62-65 Previous researchers found that some functions could be temporarily induced when epithelial cell types from the liver were cocultured with primary human liver cells. However, in these studies, the two types of cells were randomly distributed. It was not possible to explore the effect of controlled cell-cell interactions on liver phenotypes in the absence of confounding variables of cell seeding density. To overcome this limitation, Bhatia et al. used a photolithography method in the semiconductor industry to locate two-dimensional islands of primary rat hepatocytes surrounded by supportive 3T3-J2 mouse embryonic fibroblasts on the attached collagen. 48,66 This co-culture system is robust, reproducible, and miniaturized in a standard porous plate format compatible with an automated workflow environment, sustaining the hepatocyte culture for 4-6 weeks. The size and longevity of the micropatterned hepatocyte colonies were significantly increased by contact co-culture with stromal cells. 61 In particular, the functional screening showed that the 3T3 mouse embryonic fibroblasts were the best choice for induction of hepatocytes from multiple species (rats and humans). Another

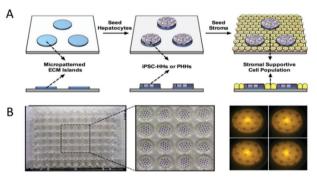


Fig. 2 Coculture of hepatocytes and nonparenchymal cells. (A) Polydimethylsiloxane (PDMS) templates are made by photolithography. Round collagen islands are planted on the template. Hepatocytes can only attach to collagen islands, and the hepatocytes on the outside of the collagen islands are washed away. The next day, the parenchymal cells are seeded and attach to areas not occupied by hepatocytes. (B) The polydimethylsiloxane (PDMS) template is made using the photolithography method. (Reproduced with permission from ref. 68.)

notable advance made by Khetani and Bhatia was the development of soft lithography, which allowed for the rapid creation of MPCCs in miniaturized formats with higher throughput screening (Fig. 2B). Currently, "the best practice" cultures consist of 500 µm islands (approximately 200 hepatocytes per island), and the remaining spaces are seeded with mouse 3T3-J2 fibroblasts. Cells on this model have good growth and differentiation characteristics. MPCCs have been successfully used to study infection and drug response in patients with HBV and HCV. 12,67 Brown et al. constructed human livers in vitro by micromode coculture method. 42 The design characteristics and validation data of this model were presented, which greatly advanced our understanding of liver function and injury. This helped reduce the risk of drug-induced organotoxicity to patients. Marchdengren et al. used MPCCs to obtain a stable and robust in vitro primary human hepatocyte model consisting of primary human hepatocytes and fibroblasts. 64 The model was used to demonstrate how to summarize the liver life cycle of hepatitis B and C viruses, and plasmodium falciparum and Plasmodium vivax pathogens in vitro. This revealed all aspects of host-pathogen interactions, which has the potential for drug and vaccine development.

3.1.2 Spherical aggregate culture method. The culture of primary hepatocytes has been used in various pharmacological and toxicological studies for decades. At present, the method of growing hepatocytes into a self-assembled three-dimensional spherical aggregate has been widely used on non-adhesive or highly compliant surfaces. 69,70 In this method, hepatocytes cannot fully diffuse. They have high cell density, and maintain the key ECM components in and around the aggregates. 71,72 In previous studies, Tatsuya Okudaira et al. constructed threedimensional liver tissue by high-density stacking these spherical aggregates. Researchers evaluated the cell viability and liver-specific function of the liver tissue. Overall, the spherical aggregate culture of hepatocytes has been shown to improve hepatocyte function in several types. This is due to the establishment of homotypic cell-cell contact, and the presence of key ECM components in and around aggregates.73 The cultured cells have good growth state and differentiation characteristics. They do not need a special carrier, or adipic function of the liver tissue. The results of many studies showed that the culture of spherical aggregates of HUVEC-covered liver cells improved the liver-specific function and cell survival rate. In addition to human umbilical vein endothelial cells, NIH/3T3 cells were subsequently covered on the surface of spherical hepatocyte aggregates. These were known to promote the vascularization of spherical endothelial cells. Okudaira et al. covered hepatocytes spherical aggregates with endothelial cells (Ecs) and NIH/3T3 cells, and then these spherical aggregates were stacked in fibers. Finally, the liver tissue was constructed by stacking these spherical aggregates. (Fig. 3A-a and b). 44,74 The practicability of this bottom-up approach was demonstrated by evaluating the liver specific function and oxygen supply of the cells in the liver tissue. Moreover, liver tissue could be collected from the hollow fibers, 75-78 which demonstrated that this structure was suitable for liver transplantation.

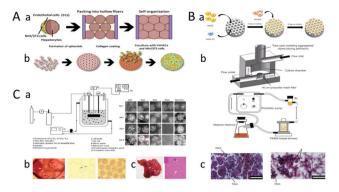


Fig. 3 (A) Heap hepatocytes covered by the HUVECs-3T3 spherical aggregate to form liver tissue. (a) The spherical aggregates form liver tissue through the bottom-up accumulation. (b) The fabrication process of the hepatocytes spherical aggregates coated with HUVECs-3T3. (Reproduced with permission from ref. 44.) (B) The spherical aggregates are assembled into liver tissue using bioreactor perfusion. (a) Endothelialized hepatocyte aggregates are prepared by PDMS. (b) Schematic diagram of the perfusion function of a bioreactor. (c) Culture of simple HepG2 aggregates (left). Culture of aggregates of endothelialized hepatocytes (right). (Reproduced with permission from ref. 61.) (C) Liver tissue is cultured by microcapsules. (a) The installation diagram of a bioreactor. Fabrication of chitosan-coated microcapsules of HepG2 cells. The microscopy images of microencapsulated HepG2 cells at an initial cell density of  $2 \times 10^6$  cells per ml for each microcapsule during 10 days cultivation in the stirred bioreactor (right). (b) Treat acute liver failure by transplant alginate microencapsulated hepatocytes. Distribution of hepatocytes in alginate microspheres. (c) 7 days after transplantation, microbeads spread throughout the abdominal cavity (black arrow). The liver of the surviving mice shows normal lobular structures after 7 days transplantation. (Reproduced with permission from ref. 83.)

A type of model is the formation of hepatocyte aggregates on a gel substrate or non-adherent plate. However, the key challenge is the inconsistent size distribution of the spherical aggregates due to diffusion limitations of key nutrients and oxygen. To overcome these challenges, scaffolds and channels are used to guide the assembly of the spheres. He et al. obtained HUVECs by co-culturing HepG2 cells, Swiss 3T3 cells and human umbilical vein endothelial cells with a microspace cell culture plate made of permeable PDMS (Fig. 3B-a). The aggregates were then placed in a bioreactor made of PDMS for 10 days (Fig. 3B-b). The liquid enters the culture chamber from the upper flow port, and is discharged from the lower flow port. A 40 µm polyester mesh filter is placed at the bottom of the culture chamber to capture the cultures. Pipe jacking is designed to seed fibers, which is clamped during perfusion. Endothelialized hepatocyte aggregates are mixed with PLLA fibers in a 1:1 ratio, and loaded into a bioreactor. In these processes, biodegradable PLLA fibers were used as scaffolds that aggregated with aggregates to form a loosely packed mode (Fig. 3B-c). These aggregates exhibited higher levels of cell proliferation and liver-specific function compared with hepatocyte aggregates alone. This study verified the important role of fibroblasts in liver cell co-culture.

Microcapsule technology is also one of the methods used to obtain spherical aggregates. It has been widely used in the fields of immobilization of cells and enzymes, large-scale culture of microbial and animal and plant cells,<sup>79</sup> controlling

the release of drugs, screening of anticancer drugs,80 and separation of proteins. Cell microcapsule technology is a promising technology that provides a three-dimensional microenvironment for cells to maintain cell vitality and function through the exchange of nutrients and wastes. 61,81,82 Khodabakhshaghdam et al. designed a small stirred bioreactor to study the effects of different stirring rates on the proliferation and liver specific functions of HepG2 cells cultured with alginate chitosan core-shell microcapsules (Fig. 3C-a). In this study, precooled type I collagen solution was mixed with alginate solution to make alginate collagen solution, and the microcapsules were cleaned twice. Then, the microcapsules were suspended in 0.3% (wt.) chitosan solution (pH = 6.5-6.6) for 5 min to obtain chitosan-coated microcapsules. The spherical aggregates were assembled into the liver tissue by agitation of flat blades. When the stirring rate increased, a large number of smaller and denser aggregates were formed. Finally, rat liver cell microspheres were transplanted into the abdominal cavity of rats with acute liver injury. Then, they found that the microspheres spread throughout the abdomen 7 days after transplantation (black arrow shows the cell high density culture is realized). These results showed that microencapsulated hydrogels loaded with liver cells were cultured in a small stirred bioreactor at a low mixing rate and had great potential for mass production of liver cells, while maintaining liver specific function. At present, although spherical aggregates culture technology has been used for many years, it is mainly used in bioreactors or artificial liver devices.

3.1.3 Cell sheet culture method. With the development of tissue engineering, researchers have developed a method to separate cell sheets from thermosensitive surfaces.84-86 Cells are attached to the culture surface via membrane receptors and cellular adhesion proteins. With the development of the liver tissue engineering, cell sheets have been widely used to grow liver tissue in vitro. 86 Subcutaneous hepatocyte sheet implantation is an attractive option for the treatment of various liver diseases. Traditional hepatocyte sheets are made up of primary hepatocytes and fibroblasts. Thin slices of tissue are successfully transplanted into the spleen subcutaneously (Fig. 4A). 44,48

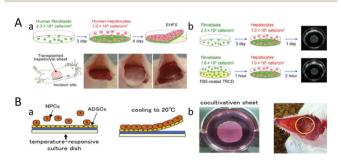


Fig. 4 The liver tissue was cultured with thin cell sheets. (A) (a) Liver tissue is cultured from human liver parenchyma cells and human fibroblasts. The cell sheets are transplanted into mice. (b) Rapid tissue culture can be achieved by using high density fibroblast cells. (Reproduced with permission from ref. 44.) (B) (a) Liver tissue is cultured from hepatic non-parenchymal cells (NPCs) and adipose-derived stem cells (ADSCs). (b) The cell sheets are transplanted into mice. (Reproduced with permission from ref. 87.)

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In another research study, the researchers constructed liver tissue from thin slices of tissue co-cultured with adipogenic stem cells (ADSCs) that produced angiogenic factors under the skin of rats (Fig. 4B).87

3.1.4 Microfluidic culture method. Microfluidic chip technology integrates the basic operation units of biological, chemical and medical analysis, such as sample preparation, reaction, separation and detection, into micrometer-scale chips,88 automatically completing the whole process of analysis. At present, various liver tissues such as hepatic sinuses, hepatic lamina and hepatic lobule can be cultured in vitro by microfluidic technology. The hepatic lobule is the structural and functional unit of the liver, which is centered on the central vein. The hepatic plate and hepatic sinuses are radially distributed around the central vein. 89-92 In the process of liver lobule culture by microfluidic technique, two layers of PDMS form tissue cavities that allow for blood flow between the layers to mimic the central vein of the liver lobules (Fig. 5A). 57,88 Fluid flows through the feed network that mimics porous endothelial cells in the microenvironment of liver tissue, allowing nutrients and applied drugs to diffuse into the tissue lumens. The biorelated geometry of the device can form an oxygen gradient perfusion and metabolic zones. Sinusoidal microfluidic

structures that simulate the central vein can radially arrange the cells in each lobule cavity, and ensure that the diffusion network can cover the liver tissue to the maximum extent. Hepatic sinusoidal endothelial cells, hepatic stellate cells, hepatic cells and Kupffer cells are distributed in three-dimensional layers in hepatic sinusoids. Sinusoidal endothelial cells are introduced to the top of the membrane to form a discrete layer of top cells, 93,94 and liver cells are placed on a collagen pre-coated PDMS substrate to form a basal cell layer. The upper channel of the device is connected at one end to the injection pump and at the other end to the media bracket to assemble all of the components together (Fig. 5B).<sup>57</sup> Human immortalized hepatocyte sodium alginate cell suspension, human umbilical vein cell sodium alginate cell suspension, buffer solution and gelatin solution are injected in an orderly manner into the microfluidic chip by microinjection pump. Laminar flow occurs when the four liquids meet at the main channel (Fig. 5C). Yajima et al. fabricated microfibers embedded with HepG2 cells using microfluidic technology. The microfibers were then recycled with rollers and bundled into bundles. The obtained bundles were placed into the perfusion chamber, and the cell culture medium was introduced for perfusion culture. Finally, the tissue needed was obtained (Fig. 5D). In addition, Knut Rennert et al. constructed a liver-like organ

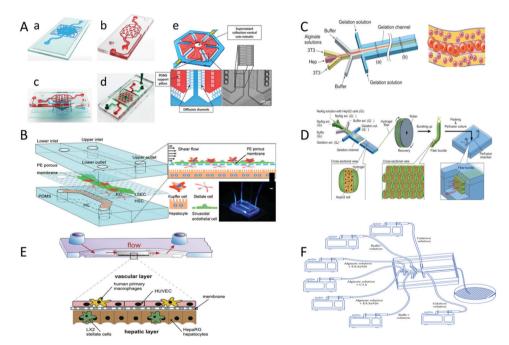


Fig. 5 The liver tissue is prepared through microfluidic method. (A) Hepatic lobules are fabricated by microfluidic technique. (a) The lower layer is the culture chamber, and (b) the upper layer provides nutrients to the cells. (c) is the side view of the microfluidic device. (d) is the top view of the device. Red and green food dyes (d) are injected into the tissue lumen and surrounding feed channels. A single lobule binds to the bottom layer of the cell compartment (Red), surrounding flow channels (Blue) and PDMS walls 2 µm (White) (e) diffusion channels. (Reproduced with permission from ref. 88.) (B) Hepatic sinusoids are fabricated through the microfluidic method. The sinusoidal chip consists of two PDMS cavities, which are combined by plasma treatment with the upper and lower channels and membranes. (Reproduced with permission from ref. 57.) (C) The liver plate was fabricated through microfluidic technology. Liver cell fluid, endothelial cell fluid, buffer fluid and gelatin fluid flow through the chip from the inside to the outside. (Reproduced with permission from ref. 99.) (D) The hepatic lobules were fabricated through microfluidic method and perfusion. Fabricated microfibers embedded with HepG2 cells by using microfluidic technology. The fibers are then bundled into bundles by using rollers. (Reproduced with permission from ref. 100.) (E) The liver-like organ were fabricated through the microfluidic method. (F) The hepatic cords were fabricated through microfluidic technology. The sodium alginate solution containing hepatocytes and 3T3 cells is continuously introduced into the microfluidic channel to generate cellloaded sodium alginate hydrogel microfibers. (Reproduced with permission from ref. 96.)

using microfluidic technology. Liver parenchymal cells were integrated between the vascular layer composed of endothelial cells and tissue macrophages, and a liver layer composed of stellate cells and liver cells (Fig. 5E).95 The three-dimensional hepatoid organ was embedded into a microfluid-infused biochip, which could provide sufficient nutrient supply for the hepatoid organ. It was similar to the shape of human hepatic sinuses. It used a suspension membrane to simulate the space of the sinus cavity shape gap. Sensors on the chip could detect increased oxygen consumption of the cells and increased expression of various proteins in the liver-like organ. The results indicated that perfusion liver organoids have similar morphological and functional characteristics with the human liver, and proved the feasibility of microfluidic technology in vitro culture of liver organoids. The hepatic cord refers to the uneven and roughly radial liver plate. The adjacent liver plate is anastomosed and connected to form a labyrinth-like structure. 96-98 In the process of hepatic cord culture by microfluidic technique, a sodium alginate solution containing liver cells and fibroblasts is continuously injected into microfluidic channels to generate anisotropic sodium alginate hydrogel microfibers. Under high oxygen tension conditions, the hydrogel fiber culture can form heteromorphic microorgans of 1 mm in length and 50 mm in diameter to mimic the hepatic cord structure found in the liver. High hepatocyte viability can be maintained for more than one month. This technique can be used to construct heterotypic micro-organs with precisely arranged multiple cell types, which will also contribute to the development of a new method of liver tissue engineering (Fig. 5F).

#### 3.2 Top-down tissue engineering method

3.2.1 3D bioprinting culture method. Zhong et al. fabricated a 3D hydrogel scaffold using a bioprinter. L02 cells (HL-7702, human liver cell line) were cultured and deposited, and then transplanted into the liver of mice with partial hepatectomy or radiation-induced liver injury (RILD) (Fig. 6A). The levels of enzymes and proteins in the liver of mice in each group were compared. The results showed that the 3D printing method could ensure cell viability and the content of the enzymes. Jeon et al. used a 3D bioprinting system to construct an alginate 3D printed liver structure (Fig. 6B). The results showed that the cells grew well on the alginate scaffold and the expression of liver-specific genes increased. Compared with the 2D culture, the 3D cultured cells showed better liver properties. For the construction of the liver lobule, Caddeo et al. used the sequential 3D bioprinting of the human hiPSC-derived liver cells in a hexagonal lobule structure with supporting cells from the endothelial and mesenchymal original cells, filling the lining of the lobules (Fig. 6C). 101,102 Firstly, they mixed the liver parenchymal cells that were induced from human pluripotent stem cells with photocurable hydrogel solution. HiPSC-derived hepatocytes were graphically mapped with a digital mask of the desired geometry, and then the sertoli cells were graphically mapped with the second digital mask (Fig. 6C-a). Sertoli cells and HiPSC-derived hepatocytes were fluorescently labeled with

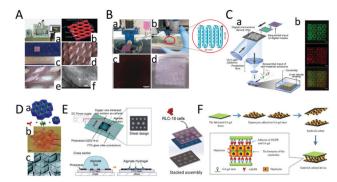


Fig. 6 Creation of liver tissue using 3D bioprinting technology. (A) Creation of mini livers using 3D printing technology. (a) The 3D printing equipment. (b) The 3D-printed structure containing pores. (c) A view of the structure obtained through 3D bioprinting technology. (d and e) The picture of the pores. (f) The single triangle refers to the cells, and the double triangles refer to the hydrogel. (Reproduced from ref. 102.) (B) (a) The 3D printing equipment. (b) An alginate containing mCherry-HepG2 cells is extruded utilizing nozzle pressure. (c) The fluorescence micrograph of mCherry-HepG2 cells. (d) Multilayer mCherry-HepG2 cells solidified successively to form tissue. (Reproduced with permission from ref. 104.) (C) (a) Liver lobules are prepared through 3D bioprinting technology. (b) Sertoli cells and HiPSCderived hepatocytes are fluorescently labeled with red and green dye. (Reproduced with permission from ref. 101.) (D) Multiple layers are printed to form liver tissue. (Reproduced with permission from ref. 103.) (E) Multiple layers stack to form hepatic lobules. (Reproduced with permission from ref. 39.) (F) Liver cell coagulation tablets were prepared using 3D bioprinting technology. Hepatocytes were seeded and adhered to a gel sheet. (Reproduced with permission from ref. 105.)

red and green dye (Fig. 6C-b). After photopolymerization, structures resembling healthy liver tissue were obtained.

Hepatic lobule structures could also be obtained by sequential 3D bioprinting of various materials (Fig. 6D). 103 Liu et al. used a multilayer photopatterning platform to embed cells in complex hydrogels to make liver tissue. First, a closed chamber was filled with a cell-loaded prepolymer solution. This was exposed to light through a mask, and then photocrosslinked to form patterns with microscale characteristics. The height of the chambers was increased to repeat this process to form a multi-layer cell-rich liver tissue structure with microscale characteristics.

Another way to construct liver lobules using 3D bioprinting is with micro-patterned electrode devices. 103,104 Liu et al. developed a micropatterned electrode device using lithography technology. Two kinds of hepatic lobule patterns with different diameters (1.5 mm and 2.0 mm) were designed. A DC voltage was applied to the device to trigger the electrodeposition process. An alginate solution formed the alginate gel structure according to the principle of electrodeposition. Finally, the prepared cell-loaded alginate sheets were further separated and stacked to form multilayer liver lobule tissue (Fig. 6E).<sup>39</sup>

In addition to mixing cells with bio-ink to form a cellprinting solution, Lee et al. printed gel sheets using a 3D bioprinter. Liver cells can then be seeded and attached to the gel sheets to obtain liver tissue. The researchers synthesized a polymer of galactosylated alginate (GA) to produce the gel sheet because it contained a sufficient number of galactosylated chains that bind easily to the salivation glycoprotein receptor

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(ASGPR) of liver cells. For a comparative study, liver cells were individually inoculated on alginate gel sheets and GA-gel sheets. The results showed that the hepatocytes inoculated on the GA-gel sheets showed good adhesion characteristics, but the adhesion of the liver cells inoculated on the alginate gel sheet was poor (Fig. 6F). 105

3.2.2 Decellularization-cellularization culture method. "Decellularization-cellularization culture method" has developed into important technologies for constructing vital organs (liver), which opens up new ideas for the treatment of end-stage liver diseases. Livers from humans, mice and pigs are decellularized, then inoculated with human liver cells and transplanted into the body to treat many kinds of liver diseases (Fig. 7A). ECM is a complex network of macromolecules, 103 which not only provides an extracellular scaffold for cells, but also plays an important role in cell function regulation. 104 Fully decellular human liver scaffolds obtained by the decellularization of untransplanted human livers can solve the problems of biocompatibility and with different types of human liver cells 106 (Fig. 7B). High shear stress oscillatory decellularization is one of the best methods. The process involves removing the immunogenic cellular material, while retaining ECM proteins by progressively increasing the gravity strength (Fig. 7C).107 The macroscopic appearance and histology of the scaffold were analyzed after decellularization.

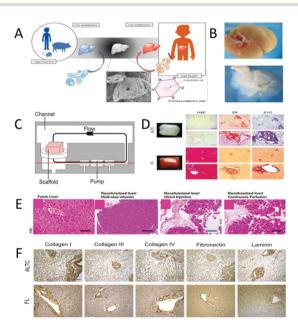


Fig. 7 Decellularization-cellularization technology is used to produce liver tissue. (A) Livers from humans, mice and pigs are decellularized, and then inoculated with human liver cells and transplanted into the body. (Reproduced with permission from ref. 108.) (B) Fresh harvested liver (top) and the decellularized liver (bottom). (C) A reperfusion device for regeneration. (D) Rapid production of liver tissue by high shear stress decellularizationrecellularization technique. (E) Reaggregation of the acellular liver matrix with adult hepatocytes. Three different cell seeding techniques are used for recellularized liver transplantation. (Reproduced with permission from ref. 50.) (F) Comparison of the expression and distribution of type I collagen, type III collagen, type IV collagen, fibronectin and laminin. The results show that the acellular sample (top) is consistent with the fresh sample (bottom). (Reproduced with permission from ref. 107.)

Staining results showed that the nucleus disappeared (blue, H&E), and the cellular material (yellow, SR), collagen (red, SR) and elastin (blue/black, EVG) were preserved (above). This was different from the macroscopic appearance and histological image of fresh human liver (below) (Fig. 7D). After obtaining the decellular biological scaffold, 108,109 the hepatocytes need to be replanted on this scaffold. Researchers used the three different cell seeding techniques for recellularized liver transplantation and evaluated these models individually. After extensive evaluation of the integrity, attachment, function, and distribution of the transplanted cells, the results of the multi-step perfusion technique were found to be the most appropriate technology (Fig. 7E). Park et al. obtained pig iPSC-Heps, which exhibits a variety of liver functions. 106 IPSC-Heps was re-implanted into decellular liver scaffolds, and the recellularized liver was then cultured using a continuous perfusion system. Finally, the recellularized liver was transplanted into rats and the grafts expressed hepatocyte markers without rupture. These results lay the foundation for the development of bioengineered livers using stem cells and decellular scaffolds (Fig. 7F).

3.2.3 Nanofiber scaffold culture method. Scaffolds fabricated using the biomolecule and biodegradable polymer electrospun nanofibers could allow cells to attach to and grow inwards to eventually replace damaged tissues and organs in living organisms. 96,110,111 At present, the tensile method, template synthesis method, phase separation method, selfassembly method and electrostatic spinning method can be used to prepare nanofibers. Among these methods, the electrostatic spinning method is widely used due to its simple operation, wide application range and relatively high production efficiency. Electrospun nanofibers are biomimetic materials of natural fibrous extracellular matrix (ECM), 56,112,113 and have been successfully used as scaffold materials for liver tissue engineering. In this review, five scaffolds commonly used in liver tissue engineering are introduced and proved to be suitable: GC nanofibers, 45,114,115 SAPNF self-assembled peptide nanofibers, 116 PCL scaffolds, 54,55 PLGA scaffolds,<sup>53</sup> and chitosan nanofibers scaffolds.<sup>113</sup>

Gao et al. investigated the response of liver cells to the surface characteristics of electrospinning fibers. PCL scaffolds with large surface depression (2 μm), small surface depression (0.37 µm) and no surface depression (NSD) were prepared. HepG2 was inoculated onto the PCL scaffold for a maximum of 14 days. The comparison and analysis of the cell activity and DNA content in each group showed that the scaffolds could promote the gene expression of albumin, and it was found that the HepG2 cells may prefer small depressions to smooth and large depressions (Fig. 8A).<sup>54</sup> In addition, Slivac et al. compared a synthetic polyester PCL with a natural ECM isolated from pig liver<sup>55</sup> (Fig. 8B-a). Cells are transfected with a plasmid carrying the red fluorescent protein (RFP) gene. Green cells are living cells, red and orange cells are successfully transfected cells. There is no significant difference in the number of transfections between the two scaffolds. It was found that the cell diffusion rates and the percentage of transfected cells on the two scaffolds were quite similar (Fig. 8B-b). Mechanical and structural analyses showed that the tensile stress resistance of PCL increased

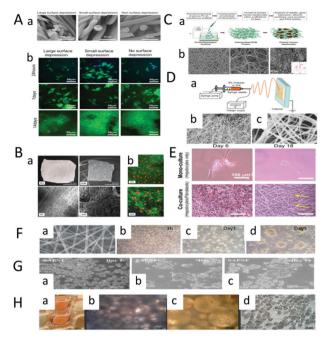


Fig. 8 Preparation of liver tissue based on nanofiber scaffolds. (A) Fabrication of the liver tissue based on PCL scaffolds. (a) Scanning electron microscopy (SEM) images of large, small and non-pitted fibers. (b) Seed liver cells on fibrous scaffolds with large, small and no surface depressions. The distribution of cells is observed under a microscope at 24 h, 7 d and 14 d. The nucleus is blue. (Reproduced with a permission from ref. 54.) (B) Comparison of the synthetic polyester PCL scaffold with natural protein-based ECM. (a) Physical and electron microscopic views of the electrospun PCL felt and pig liver ECM. (b) Cells are transfected with a plasmid carrying the red fluorescent protein (RFP) gene. Green cells are living cells, red and orange cells are successfully transfected cells. (Reproduced with permission from ref. 60.) (C) Fabrication of liver tissue based on the PLGA scaffolds. (a) Porous PLGA nanofiber scaffolds are prepared by wet electrospinning. Seed human primary hepatocytes on the scaffolds and culture for 14 days. (b) Polylactic acid scaffolds with different thicknesses (top). Human primary hepatocytes implanted on scaffolds (bottom). (Reproduced with permission from ref. 53.) (D) (a) Diagram of the electrospinning device. (b and c) SEM images of the chitosan nanofiber scaffolds at low and high magnification. (E) Morphology of the single and co-cultured hepatocytes on the surface of the chitosan nanofiber scaffolds. (Reproduced with permission from ref. 115.) (F) Fabrication of liver tissue based on galactosylated chitosan (GC) nanofiber scaffolds. (a) Fabrication galactosylated chitosan (GC) nanofibers by electrostatic spinning. (b) Distribution of human liver cells at different times after inoculation into scaffolds. (Reproduced with permission from ref. 45.) (G) Fabrication of liver tissue based on SAPNF self-assembled peptide nanofiber scaffolds. (a-c) At 24 h after inoculation, they aggregated into twins and triplets. (Reproduced with permission from ref. 116.) (H) Fabrication of liver tissue based on three-dimensional nanofiber PLLA scaffolds. (a and b) Schematic diagram of the scaffold after the implantation of primary rat hepatocytes by centrifugal seeding technique. (c) After 7 days of 3D culture, the nanofiber scaffold is observed by transmission light microscope. (d) Transmission light images of 2D cells culture in a 96-well Petri dish for 7 days. (Reproduced with permission from ref. 60.)

by about 12 times compared with the natural ECM. These results indicated that PCL electrospinning pads can be used to simulate natural scaffolds.

Brown et al. developed a three-dimensional nanofiber scaffold based on the PLGA polymer and used the wet electrostatic spinning technology to create a porous structure. ECM proteins (type I collagen or fibronectin) were chemically linked to

electrospun PLGA via amine-coupling to maintain hepatocyte function in vitro (Fig. 8C).53 Rajendran et al. used chitosan nanofiber scaffolds to construct 3D liver models<sup>115</sup> (Fig. 8D). The chitosan nanofiber scaffolds can be obtained by applying a DC voltage of 20-25 kV between the syringe needle and the square metal collector. By comparing the single and co-cultured hepatocytes, it was found that the co-cultured hepatocytes formed colonies, and maintained their morphology and function for a long time (Fig. 8E). The 3D liver tissue model established in this study provides a useful tool for drug screening and tissue engineering development of the engineered liver tissue. Feng et al. developed a novel natural nanofiber scaffold with a surface galactose ligand to enhance the biological activity and mechanical stability of the cultured primary hepatocytes. The nanofiber scaffolds were prepared using natural material GC by electrostatic spinning.45

Based on the disintegration assessment and Young's modulus test, GC nanofiber scaffolds were found to degrade slowly and have suitable mechanical properties as ECM. Morphological characterization, dual staining fluorescence analysis and functional analysis showed that the liver cells cultured on the GC nanofiber scaffolds formed stable three-dimensional aggregates, which were similar to human liver in terms of the albumin secretion. The result indicated that this GC-based nanofiber scaffold can be used in a variety of applications, such as the construction of bioartificial liver (Fig. 8F). Alvarez et al. compared the effects of SAPNF on hepatocyte metabolism and secretory activity after hepatocyte isolation (Fig. 8G). 116 The isolated porcine liver cells were cultured in SAPNF and type I collagen, respectively, and a morphological evaluation was conducted by scanning electron microscopy at different time points. It was found that the liver cells cultured with SAPNF showed a three-dimensional spherical structure and maintained cell differentiation during 2 weeks of culture, which indicates that SAPNF plays an important role in maintaining the differentiation function of porcine liver cells. Bierwolf et al. cultured primary rat liver cells on a 3D nanofiber PLLA scaffold for 7 days.<sup>60</sup> After 7 days of 3D culture, the nanofiber scaffold was observed by transmission light microscope, and the pores were randomly full of hepatocytes and no morphological changes were noticed. T cell viability and hepatocellular specific functions (albumin secretion, glycogen storage capacity) showed that the primary rat hepatocytes cultured on nanofiber scaffolds had high viability and well-preserved glycogen storage. Albumin was secreted during the whole culture process, and hepatocytes were highly differentiated and preserved. These phenomena indicate the applicability of three-dimensional nanofiber PLLA scaffolds in liver tissue construction (Fig. 8H).

Furthermore, the various methods for liver tissue construction in vitro are summarized as shown in Table 1.

# 4. Applications of liver tissue

#### 4.1 Tissue reconstruction

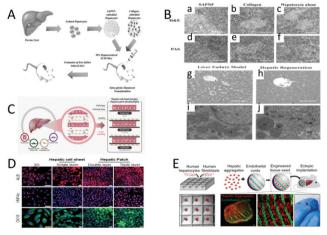
Although liver transplantation has become a standard treatment, the mortality rate from liver disease remains high. In the

Table 1 Advantages and disadvantages of various methods for liver tissue culture in vitro

Methods	Advantages	Disadvantages	Ref.
Micromode coculture	- The cells had good growth status and differentiation	– Cell separation technology is high;	48, 61–64, 66 and 68
method	characteristics;	eel 10 0 100 101	
Curls and and a manuscripts	- High cell density.	- The cultivation conditions are high.	44 64 70 75 00
Spherical aggregate culture method	<ul> <li>The cells had good growth status and differentiation characteristics;</li> </ul>	i – High purity cens are needed;	44, 61, 70, 75–83 and 117–122
culture illetilou	- Simple equipment and low cost	- High level of operation.	anu 117–122
	- High cell density;	riigii level of operation.	
	- No specific carrier is required.		
Cell sheets culture method	- Simple operation;	- Cells cannot be grown on a large scale;	39, 44, 48, 86, 87 and 105
	- The cells had good growth status and differentiation characteristics in a good microenvironment.	- Liver cells can easily lose differentiation phenotype.	
Microfluidic culture method	- The number of cells required is small and the test dose is small:		57, 88, 96, 99, 100 and 123–126
	- The cells had good growth status and differentiation	- Cell growth is slow and susceptibility is	
	characteristics;	low;	
	- Cells can maintain a specific phenotype.	- Cells cannot be grown on a large scale.	
3D bioprinting culture method	- Provides cells with an <i>in vivo</i> 3D environment;	<ul> <li>The material is required to be photosensitive.</li> </ul>	16, 39, 79, 83, 101– 105 and 127–134
	- Cells adhere easily.	•	
Decellularized-recellu- larized culture method		- Problems such as shear forces and nutrient supply may affect cell growth.	47, 50, 95, 106–109, 135 and 136
Nanofiber scaffold culture method	- Large and homogeneous space for cultivation;	- Cells easily lose normal morphology and differentiation function;	45, 53–56, 58, 60, 71, 96, 110–116 and 137–142
	– Cells adhere easily and grow well.	– Problems such as shear forces and nutrient supply may affect cell growth.	

research phase of liver cell transplantation, Sakai et al. treated acute liver failure in mice with xenogeneic hepatocyte transplantation (Fig. 9A). First, porcine liver cells embedded in SAPNF or type I collagen were injected directly into 90% of hepatectomy mice with acute liver failure. After transplantation, the researchers followed up with these mice to assess their survival. H&E was used to stain the nucleus and cytoplasm. On the seventh day after transplantation, a considerable number of porcine liver cells were detected in the spleens of mice injected with self-assembled peptide nanofibers, which had trabecular arrangement, well-developed bile ducts and tight intercellular spaces. Moreover, the glycogen storage capacity was detected after staining with PAS (Fig. 9B-a, d). Small clusters of porcine liver cells were observed in the spleen of collagen-injected mice. However, the result of PAS staining was weak, which indicated poor glycogen storage capacity (Fig. 10B-b, e). Few hepatocytes were detected when transplanted alone (Fig. 10B-c, f).

Kim et al. tried to explore the effect of the number of nanofiber layers on liver tissue repair (Fig. 9C). 143 The regenerative treatments of liver injury are promoted by stacking patient-specific progenitor cells on electrospun nanofibers. HCdH-Heps cells are seeded onto fiber sheets to form a hepatic cell sheet (single layer). Human umbilical vein cells and liver cells are seeded on the thin sheet of electrospun fibers to form a hepatic cell patch (double layer and triple layer). ALB, CK18, and HNF4 are differentiation products, so the effects of hepatic sheets and hepatic patches on tissue reconstruction can be determined by the detection of ALB (red, top), CK18 (green, bottom), and HNF4 (red, middle). Functional analysis revealed that the hepatic patch synthesized using hCdHs recapitulated a liver tissue-like structure. Therefore, patient-specific hCdHs



Liver reconstruction. (A) Treatment of acute liver failure in mice by xenogeneic hepatocyte transplantation. (B) Histological findings of mice after hepatocyte transplantation. H&E stains the nucleus and cytoplasm. PAS staining shows glycogen storage capacity. (a and d) Pig liver cells are seeded on the SAPNF scaffold, and then transplanted into the spleen of mice. (b and e) Pig liver cells are seeded on the collagen, and then transplanted into the spleen of mice. (c and f) The liver cells are individually injected into the spleens of the mice. (g and i) Liver injury model. (h and j) Liver repair. Intercellular interstitial junction (Gj) and tight junction (Tj), well-developed bile duct (Bc), and glycogen wreath (Gly). (Reproduced with permission from ref. 143.) (C) Promoting regenerative treatment of the liver injury by stacking the patient-specific progenitor cells on electrospun nanofibers. (D) HCdhsheps differentiate on single, double, or three-layer electrostatic spun fiber stacked sheets express mature hepatocyte specific markers. (Reproduced with permission from ref. 56.) (E) A treatment that uses autologous (or allogeneic) adult cells (or stem cells) from patients to repair tissues or organs. (Reproduced with permission from ref. 144.)

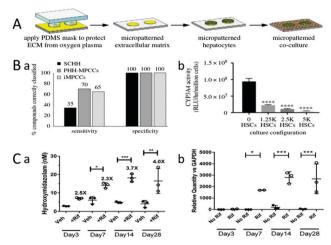


Fig. 10 Drug efficacy testing. (A) Building a liver model. Micromode coculture (MPCCs) allows the regulation of homotypic interactions between the hepatocytes and heterotypic interfaces between the hepatocytes and fibroblasts, while maintaining a constant cell count/ratio in various mode configurations. (B) (b) Tissue cultured by MPCCs can maintain a high level of CYP450 activity for several weeks, which can greatly reduce the influence of culture time on CYP450 activity. (Reproduced with permission from ref. 132.)

may be a novel source to develop hepatic cell sheets or hepatic cell patches for application in personalized tissue engineering and cell sheet-based regenerative medicine. It is also obvious that the hepatic cell patch is a better choice to promote tissue regeneration than the hepatic cell sheet (Fig. 9D). Kim *et al.* investigated the importance of the relationship between liver cells, endothelial cells, and stromal cells in tissue regeneration. After transplantation of liver seeds into mice with liver injury, large amounts of Ki67 protein and cytokeratin-18 were observed (left, white arrow), and mitosis of cells was also observed (right, white arrow), which demonstrated that ectopic liver seed transplantation can repair damaged liver tissue (Fig. 9E).<sup>144</sup> Furthermore, the relationship between hepatocytes, endothelial cells and stromal cells proved to be important in tissue regeneration.

Liver cell transplantation is one of the most effective ways to treat liver failure caused by alcohol consumption, hepatotoxic drugs or viral infection. However, the use of human liver cells has been limited by the shortage of donors, the limited number of cells suitable for transplantation, and the low efficiency of transplantation. In addition, human liver cells may lose their reproductive potential during in vitro culture, making it difficult to grow and reproduce in vitro. These above problems have been largely resolved with the use of HCdHs with proliferative and differentiated potential and cell sheets. The discovery of HCdHs solved the problem of a shortage of donors and a limited number of cells suitable for transplantation. Tissue engineering based on cell sheets solves the problem of low transplantation efficiency. Previous studies have shown that the function and therapeutic effect of transplanted liver cell sheets are significantly higher than that of direct cell perfusion. The liver sheets made up of HCdHs and HUVECs not only have the function of the original human liver, but also have remarkable therapeutic effect on acute liver injury. It is expected that replacing HUVECs with hepatic sinusoidal endothelial cells and hepatic stellate cells will have a better effect.

However, growing a tissue with a fully functional vascular system in vitro has still been a major challenge. A developed vascular system allows the implanted tissue to overcome diffuse limitations, ensuring the long-term survival of the implanted cells. In liver tissue engineering, due to the high metabolic activity of liver cells requiring adequate oxygen supply, the generation of a vascular system is more important. So far, no one has successfully developed a perfused microvascular system. A preliminary approach is to develop vascularized organoids by the co-culture of liver tissue directly with endothelial cells. Inamori et al. filled hollow fibers for plasma separation with hepatocyte spheres covered by human umbilical vein endothelial cells. The dense spheres then connected to each other to form a large tissue with regular distribution of HUVECs. On the ninth day, HUVECs invaded the hepatocytes spheres and formed a dense vascular network. 145 LeCluyse et al. also reconstructed the multicellular structure and hemodynamic characteristics of the liver using this method.<sup>74</sup> Another approach that seemed to work was to coat liver cells, endothelial cells and pericytes in a hydrogel. The results showed that these cells remained active for a month, and formed a network of blood vessels with the hydrogel. By using PCL as a skeleton material, Lee et al. successfully induced the formation of vascular networks and hepatocyte growth by injecting collagen bioinks containing three different cell types into the channels of the PCL framework. 146 Jeon et al. also experimentally verified the feasibility of this approach. 104 In addition to the above two methods, the decellularized matrix shows good effects in the construction of liver tissue vascularization in vitro. In Soto-Gutierrez's experiment, the resulting hepatic matrix preserved the entire vascular bed and bile duct after the decellularization treatment.<sup>50</sup> Researchers are currently developing many similar methods to build vascular networks, but there is still room for improvement.

### 4.2 Drug screening

At present, drug-induced liver injury (DILI) remains one of the major causes of preclinical and clinical drug consumption, and the emergence of marketed drugs withdrawal from the market. Since animal drug screening cannot accurately reflect the effects of drugs on the human body, the field of *in vitro* human tissue engineering is becoming increasingly important. 12,42,52,64,127,132,147–152

Rifampicin has a strong antibacterial effect on tuberculosis bacilli, and also has a curative effect on Gram-positive or negative bacteria and viruses. Rifampicin can activate the CYP enzyme, which is known as cytochrome oxidase P450, and is a group of isoenzymes encoded by superfamily genes related by structure and function. They are monooxygenases mainly existing in the liver and intestine, and mostly located in the endoplasmic reticulum. Furthermore, they are terminal enzymes of mixed functional oxidase system on the endoplasmic reticulum. It is involved in the metabolism of endogenous substances (such as fatty acids, vitamins and cholic acids),

metabolites.

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detoxification of exogenous substances (such as drugs), activation of pre-carcinogens (such as aromatic substances), and plays an important role in drug metabolism. CYP3A4 and CYP2D6 account for a large proportion of CYP enzymes, and their activities can also be activated by rifampicin. The activity of CYP3A4 can be assessed by metabolizing testosterone into 4-hydroxymidazolam, <sup>132</sup> and the activity of CYP2D6 can be assessed by metabolism of dextromethorphan into cathane. Brown et al. constructed a bioengineered liver model by the co-culture of 3T3-J2 fibroblasts with primary hepatocytes. The ECM, such as collagen, was evenly coated on polystyrene (or glass) and protected with a PDMS stamp. The exposed ECM region was ablated under oxygen plasma, leaving tiny ECM islands that matched the geometry of the PDMS stamp. Hepatocytes selectively attached to the ECM island, and parenchymal cells (NPCs) filled the surrounding area. So, a bioengineered liver model can be obtained (Fig. 10A). Tissue cultured by MPCCs can maintain a high level of CYP2D6 (Cytochrome P450 isozymes) activity for several weeks, which can greatly reduce the influence of the culture time on CYP2D6 activity, ensuring that changes in the CYP2D6 activity were caused by rifampicin (Fig. 10B). Then, the activity of CYP2D6

(Cytochrome P450 isozymes) can be assessed by the metabolism of dextromethorphan into coffee alkanes. Finally, the efficacy of

rifampicin can be evaluated by testing the content of the

Malaria is an insect-borne infection caused by the bite of anopheles mosquitoes or the transmission of plasmodium into the blood of people carrying the parasite. At present, there are many kinds of drugs used to treat malaria, such as artemisinin, aminoquinoline and quinoline alcohol. Primaquine is mainly used to control the recurrence of malaria. Its mechanism of action is mainly by interfering with the reduction of coenzyme II, so that the plasmodium cannot carry out the metabolic oxidation of sugar. The gametophyte of the plasmodium in the infrared phase is then quickly killed, so as to prevent the recurrence and spread of malaria. Primaquine should be screened before it is used for clinical treatment. First, Sandra March et al. made micropatterned collagen islands in the 24-well or 96-well plates. 64 Then, MPCCs were formed by seeding primary human hepatocytes continuously onto the collagen island and seeding 3T3-J2 fibroblasts onto the spaces around the island (Fig. 11A). Under different experimental interferences, infected MPCCs by Plasmodium falciparum sporozoa were cryopreserved to detect their sensitivity to primaquine (Fig. 11B-a and b). They reproduced the liver life cycles of the plasmodium pathogens Plasmodium falciparum and Plasmodium vivax in vitro. The results showed that MPCCs had the ability to distinguish live spores from attenuated spores (Fig. 11B-c). The MPCC platform can be used to uncover various aspects of host-pathogen interactions, and has the potential for drug and vaccine development.

In addition to drug efficacy and hepatotoxicity detection, drug metabolism detection is very necessary. 127 Chang et al. developed three-dimensional liver micro-organs as an in vitro drug metabolism model. They obtained the liver tissue by 3D bioprinting, and tested its ability to metabolize drugs through

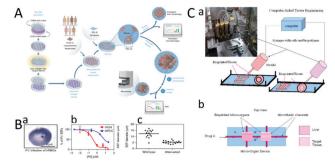
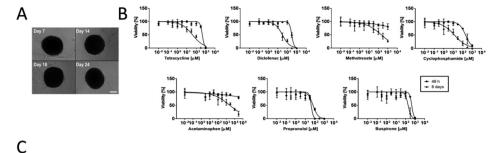


Fig. 11 Application of MPCC in the infection of hepatophilic pathogens in vitro. (A) The preparation, sowing and cultivation of MPCC. (B) (a) MPCC is infected with cryopreserved Plasmodium falciparum sporozoan, and its sensitivity to primaguine is tested. (b) Detection of the susceptibility of virus-infected MPCC to primaquine. (c) MPCC has the ability to distinguish between live and attenuated spores. Model of drug metabolism in vitro. (Reproduced from ref. 64.) (C) (a) Cells write directly. (b) Diagram of a device used to detect drug metabolism. (Reproduced with permission from ref. 127.)

an in vitro dynamic drug infusion (Fig. 11C-a).52 Finally, the drug EFC (7-ethoxy-4-trifluoromethyl Coumarin) was injected into the tissue system, and the metabolic degree of the drug was determined by observing the fluorescently-labeled metabolite HFC (7-hydroxy-4-trifluoromethyl Coumarin) (Fig. 11Cb). This research confirmed the feasibility of the model in drug metabolism.

Tetracycline, diclofenac, methotrexate, cyclophosphamide and acetaminophen are known as five hepatotoxic drugs, while propranolol and buspirone are two non-drug liver injury compounds. Spherical aggregates composed of hepatocyte were exposed to the above drugs individually for a long term. Starting from the 7th day after sowing, the shape of the sphere is shown in Fig. 12A. The spheres were treated with a single dose once within 48 h (circle) or repeated every 2 days within 8 d (triangle) to observe the coordinate diagram of the effect of the drug content on the hepatocyte activity 153 (Fig. 12B). In both cases, the ATP levels are used as a marker of viability. The IC50 is the half-inhibitory concentration, which indicates the concentration required for a drug to inhibit cell growth, viral replication, or other properties, at 50% of the required concentration. After a single dose of methotrexate and acetaminophen, the cell viability decreased slightly and IC50 values were higher. In contrast, the IC50 value of re-administration was lower and consistent with the previously reported  $C_{\text{max}}$  values in mice. Roproxolol and buspirone produced fairly stable doseresponse curves with similar IC50 values for single or repeated dosing. The IC50 values of tetracycline, diclofenac and cyclophosphamide were 10 times lower after repeated treatment than after a single treatment. The corresponding concentrations are correlated with the maximum plasma concentration  $(C_{\text{max}})$ , suggesting that spherical aggregates may be suitable for studying hepatotoxicity in vivo (Fig. 12C). From the substances tested, tetracycline, diclofenac and cyclophosphamide were labeled as hepatotoxic after 48 hours, while methotrexate and acetaminophen showed hepatotoxicity only after repeated exposure. As expected, both propranolol and buspirone were



IC<sub>50</sub> values of DILI and non-DILI compounds in 3D liver spheroids as well as reported 3D in vitro IC50 [µM] Tetracyclin Diclofenac 294.0 29.7 Methotrexate 389'98 206. 197.1 Cyclophosphamid 13.29 644'301'371 2750 2216 Acetaminopher Propranolo 65.84 41.3

Fig. 12 Drug screening for DILI. (A) Spherical aggregates of hepatocytes. (B) Long-term exposure tests were performed on globular cells composed of parenchymal cells using the known hepatotoxic drugs tetracycline, diclofenac, methotrexate, cyclophosphamide, acetaminophen, and the nonpharmaceutical liver injury compounds propranolol and buspirone. From the 7th day after seeding, the balls are treated with a single dose once within 48 h (round) or repeated every 2 days within 8 d (triangle). (C) IC<sub>50</sub> values of DILI and non-DILI compounds in 3D liver spheroids, as well as reported mouse  $C_{\text{max}}$  values. (Reproduced with permission from ref. 153.)

correctly classified as non-pharmacological liver injury compounds in both short-and long-term exposure trials. The research indicated the applicability of liver tissue engineering in the testing of drug hepatotoxicity.

Although drug screening based on liver engineering has been widely used, its development process is the joint efforts of many researchers. At an early stage, Elaut et al. found that the cell survival, phenotype and liver specific function were rapidly lost when primary isolated hepatocytes were cultured in a 2D model with simple construction. These phenotypic changes are mainly due to changes in gene expression caused by reduced transcription of relevant liver-specific genes, which also can be called dedifferentiation of isolated hepatocytes. 154 Rowe et al. also demonstrated that when liver cells were isolated and cultured by conventional methods, they developed dedifferentiation as matured. 155 Then, they quantitatively compared the proteome of human fetal liver, adult hepatocytes, and HepG2 cell lines. They also demonstrated the proteomic changes typical of hepatocyte dedifferentiation in the ECM interlayer culture, a new culture method that can better preserve the phenotype and function of hepatocytes. Nudischer et al. measured cell viability in single and co-culture cells by assessing intracellular ATP levels. It was found that the ATP level of the co-cultured cells was about 40% higher than that of the single cultured cells on the seventh day.<sup>153</sup> Nguyen et al. evaluated the toxicity of terovasfloxacin in 2D and 3D hepatocyte cultures. First, both models were administered continuously for 7 days, and the cell activity and albumin production were assessed. It was found that after seven days of administration, the albumin and ATP levels produced by the 3D model were significantly reduced. Meanwhile, the 2D model under the same treatment conditions showed only the highest dose of terovasfloxacin toxicity. 132 Chu et al. conducted toxicity tests of sodium nitrite and acrylamide in 3D and monolayer liver models,

respectively, to study the sensitivity of the 3D model to toxicity. As expected, HepG2 cells in the 3D model reacted with a lower concentration and shorter exposure time than cells in the monolayer model.59

The results of the above studies indicated that liver tissue cultured by traditional methods lacked the complexity of natural tissue in drug toxicity screening and drug metabolism identification, so the predictive ability was limited and the sensitivity was about 30-50%. However, the liver tissue model cultured by the co-culture method can significantly improve the detection of toxicity and metabolites, and increase the sensitivity to 70-75%. The function of liver tissue cultured in vitro by this method can be maintained for several weeks. There is evidence that the coculture of different liver cells results in better tissue stability and function than culture alone. 127,152,153 Therefore, 3D liver tissues constructed in vitro are more suitable for the accurate screening of drugs than traditional 2D tissues.

### 4.3 Disease modeling

In recent years, the incidence of fatty liver, hepatitis, and alcoholic liver has increased sharply. 156 Various disease models have been produced in laboratory animals by various methods. However, the main problem is that there are considerable differences in the types of lesions between animals and humans.157 For example, cirrhosis induced in animals is rarely accompanied by the decompensated features of human disease, namely portal hypertension, ascites, encephalopathy and esophageal varicose veins. Therefore, it is necessary to construct the model of human liver disease in vitro, and then study the drug research of the disease.<sup>52</sup> In the establishment of the hepatitis C model, a normal liver tissue model was obtained by co-culture of liver cells and fibroblasts. The model was then infected with hepatitis C virus to obtain a hepatitis C model. The model was stably infected in vitro for several weeks.

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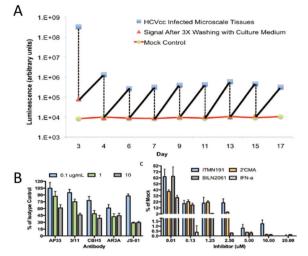


Fig. 13 (A) A disease model of infectious disease HCV was obtained. The zigzag pattern is the result of medium change every 2 days, viral replication, and luciferase secretion from the supernatant. MPCCs are treated with antibody or inhibitor to prevent the virus from entering cells. (B) Establishment of a cirrhosis model. Application of DMNA to the liver tissue to induce liver cirrhosis. The surface of the liver capsule is irregular and the reticular bands intersect with the liver parenchyma. Early nodular formation can be seen at the lower left. Hepatocytes vary considerably in size with the presence of several necrotic hepatocytes (marked with arrows). (Reproduced with permission from ref. 52.)

The zigzag pattern was the result of medium change every 2 days, viral replication, and luciferase secretion in the supernatant (Fig. 13A). The MPCCs model can then be treated with interferon or various antiviral protein drugs, and the effect of these drugs on hepatitis C can be observed (Fig. 13B). In addition, the development of therapeutics and vaccines for human liver disease requires the study of models of hostpathogen interactions. However, in vitro models of liver infection usually use physiologically abnormal hepatocellular carcinoma cell lines. However, this model will rapidly lose liver characteristics under in vitro culture conditions. To achieve a stable and robust in vitro model of human primary hepatocytes, March et al. developed MPCCs, which consisted of primary human hepatocytes and fibroblasts. This system can be used to establish hepatitis B and C and malaria models in vitro. The MPCC platform can simulate multiple liver disease models in vitro, which can be used to uncover host-pathogen interactions and has the potential for drug and vaccine development.<sup>64</sup>

In conclusion, the results of the studies suggest that the mechanisms associated with diseases such as HCV and HBV can be reproduced in a liver tissue model constructed in vitro, which can remain active and functional for several weeks. However, the current disease model also faces an important challenge, which is the low pathogen infection rate (for example, HCV infection efficiency is 5%), which will limit the application of disease models. To overcome this challenge, the researchers studied interactions between the host and virus in greater depth. One interesting example is the use of broad-spectrum Janus kinase (JAK) inhibitors to attenuate the immune response, making HCV and HBV pathogens more susceptible to infection.

We believe that with the continuous development of technology, liver tissue models constructed in vitro will be widely used in clinical medicine in the future.

## 5. Conclusion

The liver is an organ of metabolism in the body, which plays an important role in our body. Construction of liver tissue in vitro provides an effective approach for liver drug screening and transplantation. Nowadays, many techniques involving spherical aggregates of liver cells, nanofiber scaffolds, 3D bioprinting and microfluidic technology, have emerged to construct 3D liver tissue. As a multifunctional model at the intersection of in vivo and in vitro experiments, liver tissue in vitro has a good application prospect in disease model construction, drug screening, and cell therapy. Liver tissue can partially reproduce the process of liver development in vitro, maintaining the genetic characteristics of the source individual for a long time. Liver tissue reconstructed in vitro can perfectly simulate the cell-to-cell and cell-to-matrix contact environment in vivo, which helps to reconstruct the polarity of liver cells and improve the growth activity of liver cells. In vitro cultured liver tissue can survive for several months. After transplantation into the body, there is no immune rejection. It can rapidly develop into mature liver cells.

In terms of drug screening, the sensitivity of the 3D liver tissue model constructed in vitro is more than twice that of the 2D liver tissue model cultured by traditional methods. It can effectively detect the hepatotoxicity and metabolic characteristics of drugs, reduce the cost of drug research and development, and reduce the adverse reactions of drugs to the human body. In terms of liver cell transplantation, the use of thin sheets of HCdHs with proliferative and differentiated potential can largely solve the problems of donor shortage, low transplantation efficiency and loss of function. Previous studies have shown that the function and therapeutic effect of transplanted thin sheets are significantly higher than that of direct cell perfusion. The liver sheets composed of HCdHs and HUVECs not only has the function of the original human liver, but also has a significant therapeutic effect on acute liver injury. In terms of tissue models, liver models will provide pharmaceutical companies with more predictability, lower drug development costs, and reduce drug toxicity to humans. These models will also help identify new molecular targets affecting liver disease, which could lead to the development of treatments with fewer side effects.

However, the construction of liver tissue still faces many challenges. First, there is no single chamber for the collection of secreted bile. Bile secreted by liver cells in the body flows slowly through the hepatic tubules, usually in the opposite direction to the blood flow in the hepatic sinuses, and flows out of the bile duct. Bile is a complex liquid that contains bile acids, cholesterol and bilirubin, which is a toxic breakdown product of hemoglobin. Due to the lack of a functional clearance mechanism for bile, these toxic products currently accumulate and mix with the base culture medium in vitro, potentially

causing significant damage to cultured liver cells over time. One current solution is to inoculate liver cells in microchannels spaced 50 µm apart, limiting the tissue to two neatly arranged rows of liver cells. The liver cells will then be exposed to the shear stream by layers of collagen and monolayer endothelial cells. This device can clear the secreted bile from the liver cells below, which can also be used to study the reaction of bile acids to various stimuli. In addition, bile ducts can be obtained by incubating hepatic progenitor cells or bile duct cells with hydrogel. Tanimizu et al. explored the influence of matrix composition, matrix composition, and growth factors on bile duct formation. 158 Katsuda et al. co-cultured ABECs and FLCs to form functional bile duct structures. Experimental results showed that ABECs could help to establish a bile duct network. 159 However, the integration of the bile duct network and its application in 3D implantable structures remain to be explored. Second, liver organs have not yet included all the cell types in the liver. Naturally, it cannot fully realize the functions of the mature liver, as well as the vascularization, immunization and nerve innervation of the organ. To overcome this problem, Inamori et al. filled hollow fibers with hepatocyte spheres covered by human umbilical vein endothelial cells to form a dense vascular network. 145 LeCluyse et al. also reconstructed the multicellular structure and hemodynamic characteristics of the liver using this method.<sup>74</sup> In addition, Lee et al. successfully induced the formation of vascular networks by injecting collagen bioinks containing three different cell types into the channels of the PCL framework. 146 Researchers are developing many similar methods to build vascular networks, but there is still room for improvement. Third, due to the limited source of liver cells, xenogeneic liver cells have to be introduced. However, xenogeneic liver cells will face immune rejection and infection. Fourth, the functional expression of cultured liver tissue requires the co-culture of various functional cells and the interaction of various growth factors. Although some problems still exist, the results of liver tissue engineering research have been remarkable. Although implantable complete livers cannot be constructed at present, it is believed that with the continuous development of science and technology, implantable complete livers will be manufactured in the next few years, which will become a good experimental tool and clinical treatment model.

## Conflicts of interest

There are no conflicts to declare.

# Acknowledgements

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