Review

Review of Synthetic and Hybrid Scaffolds in Cartilage Tissue Engineering

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Abstract: Cartilage tissue is under extensive investigation in tissue engineering and regenerative medicine studies because of its limited regenerative potential. Currently, many scaffolds are undergoing scientific and clinical research. A key for appropriate scaffolding is the assurance of a temporary cellular environment that allows the cells to function as in native tissue. These scaffolds should meet the relevant requirements, including appropriate architecture and physicochemical and biological properties. This is necessary for proper cell growth, which is associated with the adequate regeneration of cartilage. This paper presents a review of the development of scaffolds from synthetic polymers and hybrid materials employed for the engineering of cartilage tissue and regenerative medicine. Initially, general information on articular cartilage and an overview of the clinical strategies for the treatment of cartilage defects are presented. Then, the requirements for scaffolds in regenerative medicine, materials intended for membranes, and methods for obtaining them are briefly described. We also describe the hybrid materials that combine the advantages of both synthetic and natural polymers, which provide better properties for the scaffold. The last part of the article is focused on scaffolds in cartilage tissue engineering that have been confirmed by undergoing preclinical and clinical tests.

Keywords: cartilage tissue engineering; articular cartilage; scaffolds; scaffold obtaining methods; materials for scaffolds; scaffold requirements; synthetic and hybrid scaffolds; chondrocytes; mesenchymal stem cells; tissue engineering; regenerative medicine

1. Introduction

Most human tissues and organs have a limited capacity to properly self-regenerate. Moreover, they are often exposed to damage as a result of injuries, accidents, and various diseases involving tissue dysfunction or devastating deficits [1,2]. Many surgical strategies have been developed to ameliorate these problems, including the transplantation of artificial substitutes, such as joint prostheses, heart valves, kidneys, or even tissues and organs [3–5]. Unfortunately, the main obstacles for organ transplantation are the deficit of donor organs and the necessity of lifelong immunosuppression. Nonbiological components can cause particular problems, such as a lack of biocompatibility, the development of serious infections, and limited durability [6,7]. Therefore, regenerative medicine including tissue engineering (TE) is a promising domain of research that can offer not only tissues and organs for transplantation but can also provide new perspectives for the treatment of many diseases [8]. This is due to the combination of biological sciences and material engineering methods enabling the development and acquisition of biological substitutes [2,9]. At present, regenerative medicine offers methods for treating various tissues, including the skin, musculoskeletal tissue, the liver, gastrointestinal tissue, nervous system tissue, and cardiovascular tissue [9–12] and can even treat diseases such as diabetes [13,14]. Currently, scaffolds are increasingly popular substitutes in TE. Scaffolds can be used for in vitro cultures of appropriate
cells, which can be implanted into the body as a bio-implant after a suitable amount of time and can also be transplanted directly into the organism as a medium for the colonization of host stem cells [15–18]. Scaffolds should be three-dimensional with a network of an interconnected pore structure and tunable sizes that depend on the kinds of cells. A scaffold needs to be biocompatible and provide appropriate mechanical stability and shape properties to resist stresses during cultivation and after being implanted into the body [19–21].

An example of a tissue with limited regenerative capacity is cartilage, due to its lack of vascularization and innervations [22]. Scientists and doctors are still looking for an effective method to regenerate cartilage, and the most promising method is to use scaffolds [22–25].

This review article presents a general description of articular cartilage, the problems faced by this organ, and the current methods for its treatment. Next, the requirements and materials for scaffolds in cartilage engineering are presented, along with general methods for their preparation. In this section, the requirements for scaffolds using chondrocytes and stem cells are also highlighted. The last section of the article presents scaffolds made of synthetic polymers and their combinations with natural materials (hybrid scaffolds). This work focuses on research from the last decade, taking into consideration scaffolds that are currently under development as well as those that have undergone or are undergoing clinical research.

2. Articular Cartilage and Clinical Strategies for Treatment

Cartilage is a skeletal connective tissue classified into three types, for which hyaline is necessary to enable proper movement [23,26]. Cartilage is still a still problem for regenerative medicine because there is no effective treatment for its reconstruction. Currently, supportive treatment methods are used. Articular cartilage and its general treatment methods are briefly discussed in this section.

2.1. Articular Cartilage: Characteristics, Roles, Joint Diseases, and Traumatic Lesions

Articular cartilage (AC) is a type of hyaline cartilage. It is a hard and elastic tissue located between the bones. AC is composed of spheroid cells called chondrocytes and is 10 to 13 μm in diameter [23]. These cells constitute about 2% of the total volume of AC and produce an extracellular matrix (ECM) that is rich, among others, in collagen type II and proteoglycans. As a solid phase, AC is porous and permeable. The main component of the fluid phase of AC is water with inorganic ions such as sodium, chloride, and potassium. Cells are protected by the surrounding ECM from damaging forces. Cartilage is an avascular and aneural tissue, so it has no ability to transfer nutrients to cells (ECM helps transfer nutrients to chondrocytes via diffusion from the synovial fluid). This means that cartilage does not have a self-repair ability, which is why the role of doctors and scientists in cartilage tissue engineering/regenerative medicine is important. The principle functions of AC are, among others, to protect the ends of the bones from damage caused by movement (acting as a shock absorber). AC provides the mechanical ability to withstand loads and impacts and also provides a low-friction gliding surface. Trauma, an unhealthy lifestyle, traffic accidents, or various diseases (e.g., gene mutations and autoimmune disorders) can damage the cartilage, causing pain, movement limitations, stiffness, swelling, and even disability [23,26–29]. Examples of joint-damaging diseases include obesity and osteoporosis. This leads to an abrasion of the cartilage, which loses its elasticity and resistance to friction. Initially, this process is painless due to the lack of innervation and blood supply in the cartilage (it cannot be regenerated). Cartilage wear involves slow joint death, which is a consequence of aging and the accumulation of injuries from youth. Many people with knee ligament or meniscus damage have damaged cartilage after a few years [28,30]. Another type of cartilage damage occurs due to the disease osteochondritis dissecans, in which the bone dies and is secreted into the joint with the cartilage covering it. This bone and cartilage can then fracture and become loose [31]. Next, juvenile idiopathic arthritis (JIA) is the most common type of childhood arthritis. This autoimmune disease is a chronic inflammatory process that damages the articular cartilage, induces bone epiphysis, and is responsible for extra-articular symptoms and systemic complications. This disease can occur at any stage of
developmental age and its very wide symptomatology creates diagnostic problems, especially in the initial stages of the disease development [29,32]. The most common disease connected with articular cartilage defects is osteoarthritis (OA). This is the most common musculoskeletal disorder resulting from the degradation of cartilage and leads to a poor quality of life and disability. It can affect any joints in the body, including those in the knees, hips, spine, or fingers. Factors that can affect the development of this disease include genetic factors, obesity, inflammation, trauma, occupational factors, or metabolic syndrome. Moreover, OA progresses with age and mainly affects women. Without treatment, no recovery can be achieved [30,33–36].

2.2. Treatment Methods for Cartilage Regeneration

Despite much research on the matter, there are no effective treatments for OA. Current clinical methods focus mostly on pain treatment and are not satisfactory [37]. Many clinical techniques to repair/regenerate cartilage are known. Which method will be used depends on factors such as the area of damage, the depth, location, associated damage, chronicity, and age, as well as the physical activity of the patient. The depth or degree of cartilage damage is a key factor that determines the choice of treatment method. In classifying the degree of damage, many divisions are used to describe both the depth and the area of damage. The most widely used system is the Outerbridge classification, which takes into account size and depth (Table 1) [38,39].

| Grade of Damage | Description |
|-----------------|-------------|
| Grade 0         | Normal AC with a smooth surface |
| Grade I         | Soft and swollen cartilage with a reduced amount of proteoglycans and increased water content. |
| Grade II        | The surface is cracked up to half the thickness of the cartilage, a so-called “Blemish” of cartilage. Swelling or fraying is visible via Magnetic Resonance Imaging (MRI) imaging. The area of the damage does not exceed 1.25 cm² (less than 50%) of the surface. This corresponds to damage of an intermediate thickness. |
| Grade III       | The damage exceeds half the thickness of the cartilage and may reveal the subchondral bone; the surface of the damage exceeds 1.25 cm². The deep defect comprises more than 50%. |
| Grade IV        | Full thickness defect(s). Destruction with complete exposure of the subchondral bone. |

In grades I and II, conservative treatments, such as patient education, reduction of BMI, rehabilitation, or the application of pharmacological treatment and dieting, usually give good results. Surgical interventions are recommended for grades III or IV. The most commonly used surgical methods are the microfracture (MF) method, chondroplasty surgery, osteochondral transplantation, and mosaicplasty, as well as cell-based approaches, such as autologous chondrocyte implantation (ACI). The microfracture technique is used much more frequently than other techniques but is not satisfactory. In recent years, the progress of cartilage tissue engineering has provided great hope for the regeneration of damaged cartilage. In every case, the doctor must decide which method is appropriate to choose. Each method has its own indicators and limitations, as well as advantages and disadvantages. The most appropriate management should be implemented at every stage of cartilage damage. The more extensive and serious the damage is, the more difficult and complicated the therapy will be, and the lower the chance of a full recovery [22,24,25,40].

MF is a safe, minimally invasive, and cheap method for cartilage repair. MF is a subchondral bone marrow stimulation method where a blood clot fills the defect. This provides a suitable environment for tissue regeneration. Unfortunately, the MF technique promotes regeneration to fibrocartilage tissue with inferior biomechanical properties compared to hyaline cartilage [37,41].
More promising methods for the treatment of chondral lesion are cell-based approaches. These techniques enable the implantation of articular chondrocytes (ACs) from the patient in place of the defect. ACI with or without a scaffold is used in routine clinical practice. Three-dimensional scaffolds serve as a temporary matrix for chondrocytes isolated from a healthy non-load-bearing area of the patient’s cartilage. Generally, the therapeutic cells are cultured in vitro on scaffolds. Then, a bio-implant is transplanted into the tissue defect. The scaffold is gradually degraded along with cartilage formation (Figure 1) [22,25,40,42,43]. The schema in Figure 2 presents the ACI method with a scaffold, showing a version where ACs can be passaged to multiply them or placed directly in the scaffold for an in vitro culture.

Compared to MF, ACI allows the repair of larger cartilage defects. Moreover, studies indicate better results for ACI compared to MF [40,45]. Unfortunately, no current treatment for articular cartilage repair has recreated native hyaline cartilage. Current approaches reconstruct fibrocartilage, which is susceptible to further damage. However, combining different approaches, including advanced scaffolds, growth factors, or alternative cell types, such as mesenchymal stem cells (MSCs), provides an alternative for obtaining an effective cartilage treatment method. MSCs can be obtained from different sources, such as adipose tissue and bone marrow with the potential to differentiate into ACs. In addition, this approach can avoid the invasion of the joint for the initial harvesting of ACs [22,25,40,43,46–51].

Scaffolds can also serve as carriers of chondrogenic cells, MSCs, and the bioactive factors influencing chondrocyte growth and differentiation (growth factors) or their combinations. Recent treatments have indicated possibilities to regenerate articular cartilage using the surgical implantation of MSCs into articular cartilage lesions (Figure 2). Scaffolds for MSCs should meet the appropriate parameters described in Section 3.1.

Thanks to advances in medicine, it is now possible to collect stem cells not only from umbilical cord blood, endometrium, or bone marrow [46,47,49] but also from the adult tissues of each
organism, especially from adipose tissue (AT) [47,52,53]. The main advantage of AT is its availability and abundance. Studies show that there are many times more stem cells in AT than in bone marrow. MSCs obtained from their own AT have a significant advantage over bone marrow cells due to their availability and large number. The collection procedure itself is also less painful and invasive for the patient [47,52,54,55].

Figure 2. Cartilage repair methods via mesenchymal stem cell (MSC)-based therapies: (a) full-thickness cartilage injury; (b–d) therapies using MSCs and appropriate additives. The schema was modified from a previous article [53].

In clinical methods, growth factors for ACs and MSCs are helpful. Studies have shown the effects of growth factors on chondrogenesis and the maintenance of the correct phenotypes of cells. These growth factors can be added to the medium or scaffold during cultivation. Polypeptide mediators, such as transforming growth factor β (TGF-β), insulin-like growth factor (IGF), and fibroblast growth factor (FGF), stimulate the proliferation of cartilage cells and stabilize their phenotypic expression and chondrogenesis. It has been shown that the therapeutic potential of growth factors in the process of cartilage regeneration is significant. Under the influence of these factors, tissue is formed, the histological structure and biochemical properties of which are similar to hyaline cartilage. Moreover, they hasten the healing of the defect and increase the content of type II collagen compared to I [46,53,56].
3. Scaffold for Articular Cartilage Repair: Requirements, Materials, and Method for Obtaining

The role of scaffolds in cartilage tissue engineering is to provide a suitable environment for cells and guarantee success in the tissue regeneration process; this is possible by providing an environment similar to native articular cartilage. Therefore, scaffolds must possess adequate parameters, such as correct architecture, biocompatibility, degradability, or specific chemical and physical properties. This can be achieved through the choice of appropriate materials, additives, such as pore precursors, and manufacturing methods [17,57].

This section presents the requirements for scaffolds in tissue cartilage engineering, the available materials, and techniques for obtaining said scaffolds.

3.1. Requirements for Scaffolds

Scaffolds for cartilage tissue engineering should provide an appropriate environment and enable cell adhesion, migration, and development by having an appropriate architecture, controlled degradability, adequate mechanical parameters, and good biocompatibility. Many structural features, including porosity, pore size, interconnectivity, and permeability, play a meaningful role in AC development and cartilage regeneration [15,17,51,57]. A three-dimensional design for scaffolds is necessary to prevent the dedifferentiation of chondrocytes into fibroblast-like cells or the chondrogenesis of MSCs [50,58–60]. Chondrocytes cultivated on flat surfaces lose their ability to produce particular proteins that are necessary to the formation of hyaline cartilage [61,62]. A highly porous membrane with an interconnected macro-pore network can improve cell seeding, cell migration, cell development, and tissue ingrowth [19,50]. Moreover, the membrane’s structure should be micro-porous to ensure the diffusion of oxygen, nutrients, and metabolism products. Regulated and controlled biodegradation are relevant to the formation of newly regenerating tissue cartilage and mainly depend on the materials used. The products released during degradation should be non-toxic to the body and easily removable. Scaffolds should maintain appropriate parameters in their stiffness, strength, and flexibility, conducive to integration and further tissue development. These parameters are important during cultivation and especially after implantation into the body due to the conditions in the knee [15,19,44,50,51,57–59,63–65]. Moreover, the parameters of scaffolds should be adapted to the types of cells. The sizes of the macro-pores must also be properly adjusted to the types of cells [15,17,19,64]. Pore sizes of about 150–250 μm are desirable for ACs, whereas large pore sizes of more than 300 μm are adequate for MSCs. Using the right pore sizes will support cell proliferation, the preservation of an appropriate phenotype, and chondrogenic differentiation (Figure 3) [17,20,64,66–70].

![Figure 3. General schematic demonstration of the scaffold properties for the appropriate growth of articular chondrocytes (ACs) and mesenchymal stem cells (MSCs).](image-url)
To obtain an appropriate scaffold structure depends on the relevant methods, materials, and pore precursors. One way to obtain suitable pores is the use of nonwovens produced by the electrospinning method. Depending on the nonwoven used, pores of 150 μm or greater can be produced [71,72]. To obtain information on the sizes of the pores in scaffolds, specialized programs can be used. One such program is MeMoExplorer™, an advanced membrane morphology software that analyzes SEM images. This software enables the contouring of pores and the measurement of their surfaces. These pores are partitioned into various size-classes, and measurements of the total areas (porosity coefficients) are provided [73,74]. Moreover, using pore precursors such as poly(vinyl pyrrolidone) or poly(ethylene glycol) can improve the hydrophilicity and mechanical properties of membranes. This is important due to the hydrophobic nature of most synthetic polymers used for scaffold manufacturing [67,75].

Thus, physical parameters such as stiffness, the structures of scaffolds (e.g., pore size, interconnection, and porosity), and culture conditions are important for the fate of the cells. The different conditions for ACs and MSCs are presented above.

3.2. Materials Intended for Scaffolds

Materials for scaffolds should be biocompatible, exhibit adequate mechanical parameters, and be biodegradable into non-toxic and non-inflammatory components in the host organism. These materials should also be resistant to the conditions in the body, such as pH and body temperature. Therefore, appropriate materials for the production of scaffolds should be selected. Such materials can be made of synthetic or natural polymers or a combination of both (i.e., hybrid materials (hybrid)) [17,25,51,57,64,76]. Natural materials such as collagen [64,69,77], hyaluronic acid (HA) [78,79], chitosan (CH) [80,81], chondroitin sulfate (CS) [82,83], and fibrin [25,84,85] are widely used in the production of scaffolds for cartilage regeneration. These materials are characterized by their high biocompatibility and bioactivity. Due to their origins, these materials have properties similar to those of native tissues, and most of such materials naturally occur in the human body. These materials support cell attachment and stimulate the production of the ECM. Unfortunately, natural materials have disadvantages. Because of their rapid hydrolysis, natural materials quickly lose their properties suitable for the scaffold structure. Their low mechanical stability is also not adequate to support cells, and their products are thus insufficient for the regeneration of tissue. Moreover, the methods for obtaining such materials are limited due to the low resistance of natural polymers to changes in process parameters, such as high temperatures [25,50,51,57,59,64,76,86]. Synthetic polymers, such as poly(ethylene glycol) (PEG) [87,88], polycaprolactone (PCL) [89,90], polyactic acid (PLA) [87,91,92], polyurethane [93,94], poly(glycolic acid) (PGA) [87,95], polyethersulfone (PES) [96–99], and polysulfone[100,101], are more diverse and promising. Some of these materials have been approved by the FDA for clinical human use [49,51,57,102–105]. Unfortunately, decisions of the FDA may be overturned. This change is associated with a new validation request, which is a long and difficult process. Unlike natural materials, synthetic polymers can be used to produce various shapes of membranes via many techniques and provide cell attachment, as well as good mechanical, physical, and chemical properties that can be modified to improve the parameters of the material. Most of these polymers degrade into components that are metabolized in the body. Moreover, the mechanical properties and degradation time can be controlled by combining these polymers (as copolymers or blends) [20,59,64,74,76,86,102,106–113].

Synthetic materials, like natural materials, have some disadvantages. One of them is an unexpected degradation time, which can cause brittleness of the scaffolds, even during culturing [59]. One example is PLA, which can be influenced by the use of a PLA copolymer with PCL or PEG and will affect the quality of the material [107,110,114]. Synthetic materials also lack desirable biological properties [115]. Moreover, the degradation products can result in side-effects for the host organism. These side-effects mostly involve acids that can be toxic to cells during cultivation or even elicit an inflammatory response in the host organism [59,86,103,116].

To date, studies have been conducted to obtain hybrid materials. Hybrid scaffolds combine the advantages of both synthetic and natural materials, allowing one to obtain membranes with
defined mechanical properties featuring the retained bio-functionality and tunable degradation necessary for the regeneration of cartilage [51,57,58,115].

3.3. Methods for Obtaining Scaffolds

The desired architecture, mechanical parameters, and forms of a scaffold can be obtained by selecting appropriate scaffold production methods. Scaffolds can be formed into 3D membranes (sponges), hydrogels, nonwovens (nanofibers) (Figure 4), or combinations thereof.

![Figure 4](image.png)

**Figure 4.** Schematic illustration of the main forms of scaffolds for cartilage tissue engineering: (A,B) hydrogels; (C,D) sponges; (E,F) nonwoven (nanofibers). Scale bars: D—300 μm; E—1000 nm; F—300 μm.

The literature describes many methods for producing a network of connected pores offering control over the mechanical properties and the time of scaffold degradation. This section will present the most popular techniques for obtaining synthetic and hybrid scaffolds for cartilage tissue engineering [17,19,21,40,59,86,117,118].

One of the most frequently used methods is phase inversion. Depending on the factor that induces the phase separation of the polymer solution, phase inversion can be carried out in two ways. One such method involves the temperature. This method is called thermal-induced phase separation (TIPS) and can be performed in liquid–liquid and liquid–solid systems, where the temperature of the process is appropriately selected. This method can obtain large and small pores with different membrane porosities [59,66,119–121]. In the second case, the phase inversion factor is non-solvent. This is the so-called non-solvent induced phase separation (NIPS) method. Here, the properly formed polymer solution is immersed into a non-solvent of the polymer. The phase inversion then produces a membrane. In these cases, like with the TIPS method, membranes with different porosities and pore sizes can be obtained [122–124]. A variant of this method involves adding a pore precursor to a previously prepared polymer solution or during the formation of a membrane. This approach promotes the formation of larger pore sizes and higher porosity. Pore precursors are ultimately removed from the scaffold by an appropriate solvent (porogen-leaching) [71,97,117,118,125–127]. Another similar method is solvent-casting particulate leaching (SCPL). This method involves dispersing salt particles in a biocompatible polymer solution. The solvent used to dissolve the polymer is then evaporated to obtain a polymer/salt composite membrane. Then, the salt is leached out by dipping the membrane into water or other salt solvents (not polymer solvents). The obtained membrane is dried to produce a porous compatible membrane. The most commonly used pore precursors are sodium bicarbonate, sodium chloride, and a sodium acetate preparation of polycaprolactone [128].
The freeze-drying technique is a method that uses the sublimation process. In the first step, the polymer is dissolved in a suitable solvent, and then the polymer solution is cooled to its freezing point. In this way, by means of sublimation, the solid solvent is evaporated to obtain a scaffold with multiple pores. With this technique, the dissolved substances can be separated in the ice phase and a small porous structure can be obtained. The final scaffolds are then formed after the final drying. The advantage of this method is its application in biomedical contexts due to the use of water and ice crystals instead of organic solvents in the preparation of the scaffolds. One is also able to control the sizes of the pores by changing the freezing method. The disadvantages of this method are its high energy consumption and a long preparation process [21,59,117,118].

Another common method used to obtain scaffolds is electrospinning. This is a simple and effective technique that can obtain nonwovens from both natural and synthetic polymers. In this method, electrostatic forces are used to produce fibers or spheres with different morphologies and sizes at the micrometer and nanometer scales. Electrospun scaffolds can be characterized by their high porosity, good mechanical properties, and flexibility [117,129–132]. Moreover, the fibers can be modified using this method, e.g., by functionalizing the fiber surface via enzyme immobilization [133,134]. Therefore, the fibers obtained by the electrospinning technique have wide biomedical applications in addition to their use as scaffolds in tissue engineering [129–131,135].

The aforementioned methods are considered conventional. A group of more advanced methods, known as rapid prototyping (RP) techniques, enable the production of three-dimensional objects with precise spatial control over the polymer structure. In this way, scaffolds can be obtained gradually, layer by layer, according to computerized data, such as computer-aided design (CAD) or computed tomography (CT) data [5,59,136]. The most common and popular RP techniques involve 3D printing (3DP). As mentioned above, 3DP consists of creating tools and prototype functions directly from computer models. This technique is carried out by applying powdered material in layers and selectively fusing the powder through “inkjet printing” the adhesive. Then, after the layers are deposited, the unbound powder is removed and a 3D object is obtained. Three-dimensional printing can be used to precisely control the structures of the scaffolds at the micron level; however, it requires close monitoring of the tissue structure and the mechanical properties of the scaffold. The 3DP technique also includes bioprinting [5,137–140]. Another RP method is selective laser sintering (SLS), where a laser is the power source used to sinter the powdered material. The advantage of this technique is the excellent control it offers over the microstructures of the obtained scaffolds by adjusting the process parameters, e.g., the percentage composition of the mixed polymer/composite powder blend. Moreover, this method can use ultra-high molecular weights of polyethylene. Unfortunately, in this process, an additional procedure is required to remove the injected powder, which, in addition to its high operating temperature, is the main disadvantage of SLS [5,141]. In addition to RP, relevant methods include stereolithography (SLA) [142–144] and fused deposition modeling (FDM) for creating an object through the controlled deposition of molten material [5,145,146].

The scaffold material, biological factors, and even cells can be used in RP methods. These elements make it possible to obtain a construct with a precise, controllable, and complex internal structure featuring appropriate mechanical properties [59,118,136,137,147,148]. A combination of the above methods can obtain hybrid membranes, which are important due to the sensitivity of some materials to technological conditions, such as temperature. This approach can obtain scaffolds with appropriate mechanical properties as well as appropriate biological parameters. For example, it is possible to combine electrospinning techniques with 3D printing [118,149–151].

Cross-linking should also be mentioned as a common method used for the preparation of hybrid scaffolds. Cross-linking can be done in essentially two ways: (1) the formation of a multifunctional molecule with a low molecular weight, resulting in higher molecular weight branched structures and, ultimately, continuous cross-linked structures, and (2) obtaining a networked structure by bonding long linear polymer molecules. Cross-linked polymers have properties that give them numerous applications, including their (1) resistance to solvents, (2) common high softening and heat-distortion temperatures, and (3) excellent dimensional stability. In freeze-drying
and cross-linking techniques, it is possible to cross-link polymers during freeze-drying fabrication. Cross-linking polymerization is commonly used to produce hydrogels using hydrophilic monomers with cross-linkers. The components can be of both natural and synthetic origin [155]. Table 2 presents the advantages and disadvantages of the methods for obtaining scaffolds.

Table 2. Rapid prototyping (RP) and conventional methods for obtaining scaffolds for tissue engineering cartilage.

| Technique                        | Advantages                                                                 | Disadvantages                                                  |
|----------------------------------|-----------------------------------------------------------------------------|----------------------------------------------------------------|
| 3D printing (3DP)                | • Possibility of using hydrogels and cells                                  | • Low precision                                                |
|                                  |                                                                             | • Long-standing process                                         |
|                                  |                                                                             | • Poor mechanical properties                                   |
| Selective laser sintering (SLS)  | • Smart process                                                             | • High temperature                                             |
|                                  | • High precision                                                            | • Rough surface                                                |
|                                  | • No need for support                                                       |                                                                |
|                                  | • Construction                                                              |                                                                |
| Stereolithography (SLA)          | • High precision                                                            | • Risk of high process temperature                             |
|                                  | • Smart process                                                             | • Untreated                                                    |
|                                  | • Soft surface                                                              | • Material may be cytotoxic                                    |
|                                  |                                                                             | • high costs                                                   |
| Fused deposition modeling (FDM)  | • Good mechanical properties                                                | • Poor precision                                               |
|                                  |                                                                             | • High temperature                                             |
|                                  |                                                                             | • Narrow range of parameters                                   |
|                                  |                                                                             | • Limits in application to biodegradable polymers               |
| Bioprinting                      | • High precision                                                            | • Toxicity of using solvents                                   |
|                                  | • Low costs                                                                 | • Depends on the cell’s existence                              |
|                                  | • High speed of printing                                                    | • Depends on many factors                                      |
|                                  | • Possibility of supporting high cell viability                             | • Obtaining 3D structure or/and adequate pore sizes for biomedical applications can be problematic |
| Electrospinning                  | • Standard technique for obtaining nanofibrous scaffolds                    | • Toxicity when using solvents                                 |
|                                  | • Capability of controlling the pore size                                   | • High energy consumption                                       |
|                                  | • Possibility of obtaining high temperatures                                | • Irregular obtained size pores                                 |
| Freeze-drying                    | • Used for multiple purposes                                                |                                                                |
| Thermal-induced phase separation (TIPS) | • Possibility of using a low temperature                               | • Used only for thermoplastics                                 |
|                                  | • Very high porosity surface-to-volume ratio                               |                                                                |
|                                  | • Scaffolds obtained from a thermoplastic crystalline polymer               |                                                                |
| Solvent-casting particulate leaching (SCPL) | • High porosity                                                | • High toxicity when using solvents                           |
|                                  | • Low costs                                                                 | • Time consuming for thin membranes                           |
|                                  | • Can be used for fabricating thin membranes of thin-wall 3D specimens      |                                                                |
4. Scaffolds for Cartilage Treatment

Scaffolds for cartilage regeneration can be made of synthetic or natural polymers or a combination of both (a hybrid scaffold). Commercial scaffolds for regenerative cartilage are mainly made of natural materials, such as collagen or hyaluronic acid [40,44,156]. Due to the disadvantages of natural materials, research is underway to obtain scaffolds from synthetic and hybrid materials with the addition of biological components—membranes where the advantages of both synthetic and natural materials are taken into account [40,51,64]. This section presents scaffolds made of natural, synthetic, and hybrid materials, along with a short description of them (Table 3).
Table 3. Synthetic and hybrid scaffolds for cartilage regeneration.

| Scaffold Name          | Component         | Method            | Properties (Porosity (%), Pore Size (µm), Mechanical Properties) | Cell Source/Animal Model | Results                                                                                                                                 |
|------------------------|-------------------|-------------------|------------------------------------------------------------------|--------------------------|----------------------------------------------------------------------------------------------------------------------------------------|
| BioSeed®-C (Biotissue) | PGA/PLA, PDS      | Thermoplastic     | Good mechanical properties and adequate structure for cells    | Human articular chondrocytes | Assessed in clinical trials. In the results, the scaffolds featured significantly improved final postoperative values. This highlights their effectiveness in cartilage regeneration. |
| Spongy PU scaffold     | PU                | Freeze-drying     | 96.9% 126–186 µm Storage modulus: ~60.36 kPa                   | Chondrocytes, human MSCs | Biodegradable PU scaffold had better outcomes than PLA 3D membranes during culturing.                                                 |
| NSP-PCL scaffold       | PCL               | Freeze-drying     | The porosity of the scaffold was designed to promote cartilage ingrowth | Rabbit articular chondrocytes | The NSP-PCL scaffold indicated better results during in vitro and in vivo studies compared to the Chondro-Gide® scaffold.            |
| RO45 3DHIC             | PCL               | 3D printing       | RO45: 84.6% 135–285 µm Compressive modulus: 25.6 MPa 3DHIC: 83.8% 150–700 µm Compressive modulus: 3 MPa | Human adipose-derived MSCs | The RO45 scaffold was preferable for chondrogenic differentiation compared to 3DHIC, which indicated better cell proliferation, scaffold penetration, and more favorable mechanical properties in the final construct. |
| Polysulphonic scaffold | PES               | Non-solvent induced phase separation and porogen-leaching | 98.5% 60–300 µm | Rabbit model and human articular chondrocytes | A study with a rabbit model suggested that the scaffold is effective in repairing articular cartilage defects. In vitro study with human cells gave promise results. |
| PLLA-100 scaffolds     | PLLA              | Thermally induced phase separation | 93% 100 ± 20 µm | Human articular chondrocytes | The scaffold promoted the secretion of chondrogenic genes. It was better than the PLLA scaffold with larger pores (~200 µm).                |
| PLCL-2 scaffold        | PLCL              | Gel-pressing      | 80% 300–500 µm | Rabbit articular | The adequate structure of the scaffold showed that chondrocytes did not change.                                                       |
Young’s modulus: ~0.7 MPa  chondrocytes and mice model  their phenotypes during the in vitro study.  The in vivo study indicated that the scaffold would maintain mechanical integrity and guide cartilaginous tissue formation.

### Hybrid scaffold

| Scaffold | Component | Porosity | Young’s modulus | Biological effect |
|---|---|---|---|---|
| Chondrotissue® (Biotissue)[156,164] | PGA, HA | Freeze-drying | 99.1% 50–400 μm  Young’s modulus: ~9 kPa | Platelet-rich plasma and bone marrow concentrate  The one-step cartilage repair method is available for clinical use. Treatment results follow up to 5 years of good outcomes with the potential for future benefits. |
| IC scaffold [153] | PLGA, COL | Freeze-drying and cross-linking | ~150 μm  Young’s modulus: ~0.65 MPa | Bovine articular chondrocytes (BACs) and mice model  IC scaffold promoted cartilaginous gene expression, chondrocyte proliferation, and the regeneration of cartilage tissue with high mechanical properties. It seems to be promising for cartilage tissue applications. |
| Gel/PCEC-TGFβ1 hydrogel scaffold [165] | Gelatin, PCEC, TGFβ1 | Cross-linking, freeze-drying | ~85%  ~10 μm; ~450 μm  Young’s modulus: ~0.21 MPa | Rabbit articular chondrocytes  Scaffold with a controlled structure, good biocompatibility, elasticity, and mechanical properties, as well as potential in cartilage regeneration. |
| PLCL-COLI [166] | PLCL, COL | 3D printing | >85%  ~10 μm; ~450 μm  Young’s modulus: ~0.21 MPa | Bovine articular cartilage chondrocytes  A hybrid scaffold with high porosity, good mechanical strength, and interconnected pore network. It has potential as a scaffold for cartilage tissue engineering. |
| C2CHH scaffold [167] | PLA, COL, CH | Freeze-drying and melt-spun | >85%  Young’s modulus: 52.3 kPa | Rat mesenchymal stem cells (MSCs) and rat model  The in vitro study showed good properties of attachment, proliferation, and differentiation of the MSCs. Involved the implantation of a cell with MSCs and type II collagen mRNA expression. The in vivo study indicated the regeneration of tissue to hyaline cartilage. The scaffold could be promising for cartilage regeneration therapy. |
| ECM-PLGA scaffold [168,169] | PLGA, ECM | SCPL | 90% | Pig articular chondrocytes  Scaffold with high porosity and repetitive pore structure. In vitro and in vivo study |
CH/PLLA/PC scaffold [110] PLLA, CH, PC Freeze-drying and cross-linking 79–84% 49–170 μm Rabbit articular chondrocytes Outcomes from the in vivo study showed the suitability of the scaffold for cartilage tissue regeneration.

Chitosan-modified PLCL scaffold[171] PLCL, CH Porogen-leaching, lyophilization, and cross-linking ~85% 200–500 μm Young’s modulus: 0.04 MPa Pig articular chondrocytes Biodegradable scaffolds with high porosity, good mechanical strength, and interconnected pore structure. Supplied a good environment for chondrocyte adhesion, proliferation, differentiation, and ECM secretion. The results were good but still require further research.

CSMA/PECA/GO (S2) scaffold [172] CSMA, MPEG-PCL-AC (PECA), GO ~70% Mean 175.2 μm Compressive modulus: 0.48 MPa Rabbit articular chondrocytes Scaffold with an appropriate structure with biological components; provided an adequate environment for cells. The in vivo results were promising with great potential for the future.

CH—chitosan; COL—collagen, PU—polyurethane; PC—pectin based; PDS—poly-p-dioxanone; CS—chondroitin sulfate; CSMA—methacrylated chondroitin sulfate; HA—hyaluronic acid; PEG—poly(ethylene glycol); PCL—polycaprolactone; PLA—polylactic acid; PLLA—poly(l-lactide); PGA—poly(glycolic acid); PES—polyethersulfone; PLGA—poly(lactic-co-glycolic acid); PCEC—polycaprolactone-polyethylene glycol; ECM—extracellular matrix; PLCL—poly(l-lactide-co-ε-caprolactone); SCPL—solvent casting and particulate leaching method; AC—acryloyl chloride; GO—graphene oxide; PECA—poly(ethylene glycol) methyl ether-ε-caprolactone-acryloyl chloride.
4.1. Natural Scaffolds

Natural scaffolds are characterized by high bioactivity, biocompatibility, and biodegradability to non-toxic components. Due to their composition of natural materials, these scaffolds are similar to native tissue, which means that their presence creates an ideal environment for cells. Thus, the main advantage of natural polymers is their similarity to cartilage’s ECM components. Their presence appropriately stimulates chondrogenesis and the maintenance of the cellular phenotype of chondrocytes. These scaffolds affect the adhesion and proliferation of the cell and cell proliferation. Therefore, products used in cartilage regenerative medicine are mainly made of natural materials. Table 4 shows examples of commercial scaffolds, including their materials and basic characteristics. Scaffolds are mostly made of collagen, the main component of cartilage ECM. Unfortunately, these scaffolds often do not meet the necessary requirements, as they quickly lose their structure (sensitivity to an aquatic environment) and transform into a gel-like form. They are also not mechanically strong enough to support the cells and regenerated tissue. Thus, this kind of membrane does not have suitable properties to create hyaline cartilage. As a result of regeneration, non-valuable fibrous cartilage is obtained, which is susceptible to future damage [40,50,51,57,76,111,156,173].

Table 4. Natural scaffolds approved for medical use for cartilage tissue engineering.

| Product (Company)       | Materials                  | Characteristic                                                                                                                                 |
|-------------------------|----------------------------|------------------------------------------------------------------------------------------------------------------------------------------------|
| Hyalofast® (Anika) [110,154,174,175] | Benzyl ester of hyaluronic acid | A bioresorbable 3D scaffold used through a one-step procedure after microfracture. It can be used even for deep cartilage lesions. The scaffold’s non-woven structure allows it to be cut and adaptively matched into uneven lesions. |
| NeoCart® (Histogenics) [44,110,154] | Bovine type I collagen | Bioresorbable electrospun scaffold used in MACI, a two-step procedure. The patient’s chondrocytes are expanded into scaffolds. Then, they are incubated in the Tissue Engineering Processor (TEP), which simulates the variation of mechanical forces and reduces oxygen pressure, allowing the maintenance of the chondrocyte phenotype forming the appropriate proteins of the ECM. |
| ChondroGide (Geistlich) [110,154] | Type I/III collagen | The first described matrix for the ACI method. It is used in a one-step procedure. ChondroGide’s role is to support and promote the chondrogenic differentiation of MSCs released after the microfracture method. |
| ACI-Maix™ (MACI) [44,45] | Type I/III collagen | The procedure is a two-step process. Expanded autologous chondrocytes (2 or 3 passage) are cultured into the scaffold for 3 or 4 days before implantation into the patient. |
| Cartipatch® (Xizia Biotech) [44,156,173] | Agarose and alginate | The cylindrical scaffold of a single layer of hydrogel with expanded cartilage cells. The clinical procedure is the same as that for the two-step method. The alginate polymer provides elasticity to the matrix, which facilitates handling during the surgical procedure. |
| NOVOCART® 3D—AesculapOrthopaedics (BBraun) [44,64,111,156] | Type I collagen, chondroitin | A sponge scaffold with a bilayer structure and interconnected pores, used in a two-step procedure. This scaffold is desirable in young patients, <16 years |
Because natural materials usually have poor mechanical properties, they are often insufficient to regenerate a given tissue. An additional disadvantage is their limited processing methods resulting from the low resistance of the materials to changes in process parameters (e.g., pH, high-temperature, and pressure). For this reason, natural scaffolds do not have the desired parameters. Additionally, the regenerated cartilage is not hyaline cartilage but fibrocartilage with inferior properties.

4.2. Hydrogel Scaffolds

Scaffolds of a hydrogel form are of great interest in cartilage regenerative engineering. These scaffolds are formed as a result of the cross-linking of natural and synthetic (or both) polymers and are characterized by their ability to absorb water or biological fluids. All these features make such scaffolds very similar to natural cartilage ECM [155,178,179].

Unfortunately, like natural scaffolds, hydrogel scaffolds have one major disadvantage. Due to their solubility in aquatic conditions, these scaffolds have low mechanical strength, which makes them difficult to handle [105,178]. Intensive studies are currently underway on the development of hydrogels from synthetic and hybrid materials [49,53,176,177,180]. Yang et al. obtained a synthetic hydrogel scaffold with the strength and modulus of native cartilage. This scaffold was composed of a bacterial cellulose (BC) nanofiber network with a PVA–poly(2-acrylamido-2-methyl-1-propanesulfonic acid sodium salt) (PAMPS) double-network hydrogel. BC was chosen as the nanofiber network due to its high tensile strength, biocompatibility, and lack of the enzymes necessary to degrade cellulose in the human body. Moreover, BC mimics collagen. The second layer of the PVA hydrogel provides elasticity, viscoelastic energy dissipation, and tensile resistance by allowing the BC fibers to share the load in the composite framework. This is an example of a scaffold with promising performance for further research in cartilage tissue engineering [180]. An example of a hybrid hydrogel scaffold is gelatin/polycaprolactone–polyethylene glycol (Gel/PCEC-TGFβ1) (Table 3). This scaffold uses both natural and synthetic polymers and growth factors. It was prepared and evaluated for human mesenchymal stem cells derived from adipose tissue (h-AD-MSCs). During the study, the tests indicated the expression of cartilage-specific genes, such as collagen type II and aggrecan, showing promising results and potential for further research on cartilage regeneration [165].

So-called injectable hydrogels have gained interest in medicine for local deformation in cartilage. In this process, a mixture of the patient’s expanded cells with the hydrogel is injected into the cartilage-damaged area. In the body, cells gradually multiply and the hydrogel is degraded. The advantage of this method is its low invasiveness and the possibility of its precise adjustment to the defect [49,53,179]. There are numerous ongoing/recruiting clinical trials using sealant gel-based MSC constructs for cartilage regeneration [49,176]. Some of them (CARTISTEM®, CaReS®, and Cartipatch®) were approved for clinical usage (Table 4) [49,176,177].
4.3. Synthetic Scaffolds

Research on using synthetic scaffolds for the regeneration of cartilage has been described in numerous studies. These scaffolds are characterized by their biocompatibility, biodegradability, and good mechanical properties and can be obtained by various methods due to their better resistance to physicochemical properties compared to natural membranes. Currently, few synthetic scaffolds are being tested in clinical trials for their use in cartilage regeneration [51].

An example of a commercial scaffold made from synthetic materials is the BioSeed®-C (Biotissue) membrane. This membrane is composed of PGA/PLA and PDS materials and is characterized by bioresorbability, elasticity, and the ability to be cut without fraying. This membrane’s adequate 3D structure and the stability of the environment during culturing stimulate the patient’s cells to differentiate [156]. The clinical outcomes at 4 years after implantation showed promising curative results for cartilage defects of the knee [157].

The most commonly used synthetic materials for the production of scaffolds for cartilage repair are PLA [66,181,182], PCL [91,159,160,183,184], and copolymers such as PLGA [185–188] and PLCA [154,163,189]. Christensen et al. used a nanostructured porous polycaprolactone (NSP-PCL) scaffold [159] and compared its in vivo and in vitro outcomes in a rabbit model with a commercial Chondro-Gide® scaffold. The observation time was 13 weeks, and the results were better for the synthetic scaffolds than the commercial ones. This scaffold had higher chondrogenic markers during the in vitro study and better in vivo histological scores. Thus, NSP-PCL seems to be an adequate scaffold for cartilage repair [159]. Research was also conducted with other synthetic materials, such as a spongy PU scaffold [160]. Scaffolds made from PU material had good hydrophilicity and porosity with interconnected pores and adequate mechanical strength. In a previous study, a PU scaffold was compared with a conventional PLA scaffold. The suitability of the scaffold for cartilage regeneration was evaluated during culturing with chondrocytes and human mesenchymal stem cells (MSCs). The chondrocytes grew better and secreted more glycosaminoglycan in the PU scaffolds than in the PLA scaffolds. Moreover, the human MSCs showed greater chondrogenesis in the PU scaffolds than in the PLA membranes. Degradable PU scaffolds thus have potential in cartilage tissue engineering applications [158]. Another example is PES materials. Polysulphonic membranes are an example of scaffolds that offer promising results for the regeneration of cartilage, as these membranes have an interconnected pore network, good elasticity, and excellent mechanical properties [97,98,161,162]. A study with a rabbit model showed that this membrane was better than a commercial Chondro-Gide® scaffold [162]. Unfortunately, the main disadvantage of synthetic polymers is their degradation, which leads to the release of acids. This can cause inflammation in the body. Additionally, in some cases, the degradation is too fast, causing the membranes to break or even crumble; moreover, the membranes do not have adequate biological properties. Therefore, research is being done to obtain scaffolds with a combination of synthetic and natural materials [59,86,103,111,115,116].

4.4. Hybrid Scaffolds

Considering the advantages of synthetic and natural materials, scaffolds with good mechanical and biological properties can be obtained. Currently, hybrid scaffolds are one direction of research in pursuit of a suitable implant for articular cartilage regeneration. The membranes of such scaffolds can be improved by inserting other biologically active additives, such as growth factors, and through a selection of appropriate kinds of cells [25,40,49–51,64,115,190]. This section presents and discusses scaffolds made of hybrid materials. Some of these scaffolds are outlined in Tables 3 and 4.

Currently, few hybrid scaffolds have undergone clinical trials. One of them is Chondrotissue® (Biotissue) [158,166]. This resorbable membrane is composed of PGA and HA and is used in clinical contexts through a one-step treatment method. This membrane’s elasticity is due to the addition of autologous platelet-rich plasma (PRP) or serum enriched with platelets. This method relieves pain, improves mobility, and supports cartilage regeneration. Five years of clinical trials confirmed the
good outcomes of this one-step procedure with Chondrotissue®, which provides stable results with future potential in hyaline cartilage regeneration.

Rofiqoh et al. developed an IC hybrid scaffold composed of PLGA and collagen. This scaffold features a high porosity membrane with an interconnected pore network and good mechanical properties. The studies were carried out with the use of bovine articular chondrocytes and invivo implantation into mice. The results showed the regeneration of cartilage-like tissue with high potential for further work [153]. Another example of a hybrid scaffold is PLCL-COLI. In a previous study, a PLCL membrane was printed, treated with alkali, and coated with collagen type I (COLI). The obtained scaffold had high porosity with a controlled structure. This scaffold provided good biocompatibility and elastic and mechanical properties. The compressive modulus of the membrane was, moreover, 0.21 MPa (similar to human cartilage). This scaffold provides good outcomes and has promise as an implant in cartilage repair [166]. Next, a C2C1H scaffold was obtained and characterized by Haaparanta et al. This scaffold was composed of collagen, chitosan, and PLA. A synthetic polymer used as a 3D mesh gave the scaffold good mechanical strength, and the natural components mimicked an appropriate environment for chondrocytes. The researchers studied eight scaffolds, determining C2C1H to be the best. This scaffold had a highly porous structure with interconnected pores and good mechanical strength with appropriate stiffness. A culture with isolated bovine chondrocytes showed promising results, with promise for further work towards cartilage regeneration [167]. Another example is the ECM-coated polyactic-co-glycolic acid (ECM-PLGA) scaffold designed by Nogami et al. In this scaffold, a synthetic polymer established the appropriate mechanical properties, while the use of ECM provided an appropriate environment for the cells. This scaffold’s structure achieved the relevant properties for cartilage regeneration. The in vitro study showed attachment, growth, and differentiation of the MSCs. In the invivo study, cell-free scaffolds were implanted into the osteochondral defects of rat knees. Research workers demonstrated that the scaffolds promoted the regeneration of hyaline-like cartilage, which was better than the cartilage in the empty control group. An ECM-PLGA implant may be a good component for use in a one-step method for cartilage regeneration, but more research is required [168,169].

In the literature, there are many examples of hybrid scaffolds used for cartilage tissue engineering, where a biodegradable synthetic polymer provides the housing framework. Mainly, these scaffolds use porous membranes to provide the necessary mechanical properties to support tissue growth, while the additives include natural composites (bioactive fillers). These fillers produce bioactive signals that supply the required information for chondrogenesis and maintain the proper phenotypes of the chondrocytes [25,49,51,110,115,190–192]. These are mostly components that naturally occur in the cartilage, such as HA [156,164,193,194], CS [172], COL [64,151,153,167,170,195,196], and ECM [168,197].

5. Conclusions

Currently, the most promising method for cartilage regeneration is the transplantation of implants with or without cells in the area with the damage. Therefore, research is underway to obtain an appropriate scaffold. There are many commercial scaffolds used in orthopedics, which, unfortunately, do not completely fulfill their proper roles in the regeneration of hyaline cartilage. New solutions are constantly being sought, including new scaffolds, growth factors, and sources of cells, as well as methods for delivering the implants to damaged areas. Currently, according to the literature, hybrid scaffolds provided the most promising results in research on articular cartilage regeneration. A combination of synthetic materials to ensure adequate mechanical strength and natural components to ensure proper chondrogenesis and preserve the phenotype has the greatest probability of obtaining hyaline cartilage in a damaged area. In addition, the literature provides information on the search for an appropriate method/improvement of current methods for scaffolding production. Boosters that can be added to the scaffolds or the medium (e.g., growth factors) are also being sought. The selection of appropriate types of cells is also under investigation, mainly focusing on MSCs and human autologous chondrocytes. Proper selection of all the above-
mentioned factors could ensure that the appropriate articular cartilage regeneration is obtained. The problems to be solved are significant due to the number of people with cartilage problems, such as osteoarthritis. There are several scaffolds designed for orthopedics, but no one solution can guarantee the reconstruction of hyaline cartilage, as most interventions yield fibrocartilage, which is susceptible to further damage. Thus, patients eventually return to the starting point. Consequently, it is important to obtain an appropriate scaffold and method for the regeneration of hyaline cartilage.

Ultimately, two important conclusions can be highlighted. Thus far, no scaffolds have been obtained that achieve the optimal conditions for the regeneration of articular cartilage. The obtained results suggest that due to our poor ability to modify natural materials, hybrid scaffolds and composite ones combining the properties and advantages of several natural and synthetic materials are the most promising options.

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**References**

1. Gurtner, G.C.; Werner, S.; Barrandon, Y.; Longaker, M.T. Wound repair and regeneration. *Nature* **2008**, *453*, 314–321. doi:10.1038/nature07039.
2. Dzobo, K.; Thomford, N.E.; Senthobane, D.A.; Shipanga, H.; Rowe, A.; Dandara, C.; Pillay, M.; Motaung, K.S.C.M. Advances in regenerative medicine and tissue engineering: Innovation and transformation of medicine. *Stem Cells Int.* **2018**, *2018*, doi:10.1155/2018/2495848.
3. Duan, B. State-of-the-Art Review of 3D Bioprinting for Cardiovascular Tissue Engineering. *Ann. Biomed. Eng.* **2017**, *45*, 195–209. doi:10.1007/s10439-016-1607-5.
4. Scarritt, M.E.; Pashos, N.C.; Bunnell, B.A. A review of cellularization strategies for tissue engineering of whole organs. *Front. Bioeng. Biotechnol.* **2015**, *3*, 43. doi:10.3389/fbioe.2015.00043.
5. Yan, Q.; Dong, H.; Su, J.; Han, J.; Song, B.; Wei, Q.; Shi, Y. A Review of 3D Printing Technology for Medical Applications. *Engineering* **2018**, *4*, 729–742. doi:10.1016/j.eng.2018.07.021.
6. Giwa, S.; Lewis, J.K.; Alvarez, L.; Langer, R.; Roth, A.E.; Church, G.M.; Markmann, J.F.; Sachs, D.H.; Chandraker, A.; Wertheim, J.A.; et al. The promise of organ and tissue preservation to transform medicine. *Nat. Biotechnol.* **2017**, *35*, 530–542. doi:10.1038/nbt.3889.
7. Popoola, J.; Greene, H.; Kyegombe, M.; MacPhee, I.A. Patient involvement in selection of immunosuppressive regimen following transplantation. *Patient Prefer. Adherence* **2014**, *8*, 1705–1712. doi:10.2147/PPA.S38987.
8. Feinberg, A.W. Engineered tissue grafts: Opportunities and challenges in regenerative medicine. *Wiley Interdiscip. Rev. Syst. Biol. Med.* **2012**, *4*, 207–220. doi:10.1002/wsbm.164.
9. Park, K.M.; Shin, Y.M.; Kim, K.; Shin, H. Tissue Engineering and Regenerative Medicine 2017: A Year in Review. *Tissue Eng. Part B Rev.* **2018**, *24*, 327–344. doi:10.1089/ten.teb.2018.0027.
10. Murphy, S.V.; Atala, A. Organ engineering—Combining stem cells, biomaterials, and bioreactors to produce bioengineered organs for transplantation. *BioEssays* **2013**, *35*, 163–172. doi:10.1002/bies.201200062.
11. Mazza, G.; Al-Akkad, W.; Rombouts, K.; Pinzani, M. Liver tissue engineering: From implantable tissue to whole organ engineering. *Hepatol. Commun.* **2018**, *2*, 131–141. doi:10.1002/hepc.11316.
12. Tarassoli, S.P.; Jessop, Z.M.; Al-Sabah, A.; Gao, N.; Whitaker, S.; Doak, S.; Whitaker, I.S. Skin tissue engineering using 3D bioprinting: An evolving research field. *J. Plast. Reconstr. Aesthetic Surg.* **2018**, *71*, 615–623. doi:10.1016/j.bjps.2017.12.006.
13. Farina, M.; Alexander, J.F.; Thekkedath, U.; Ferrari, M.; Grattoni, A. Cell encapsulation: Overcoming barriers in cell transplantation in diabetes and beyond. *Adv. Drug Deliv. Rev.* **2019**, *139*, 92–115. doi:10.1016/j.addr.2018.04.018.
14. Liu, Y.; Luo, J.; Chen, X.; Liu, W.; Chen, T. *Cell Membrane Coating Technology: A Promising Strategy for Biomedical Applications*; Springer: Singapore, 2019; Volume 11, ISBN 0123456789.
15. Hollister, S.J. Porous scaffold design for tissue engineering. Nat. Mater. 2005, 4, 518–524, doi:10.1038/nmat1421.
16. O’Brien, F.J. Biomaterials & scaffolds for tissue engineering. Mater. Today 2011, 14, 88–95, doi:10.1016/S1369-7021(11)70058-X.
17. Bruzaukaitė, I; Bironaity, D; Bagdonas, E; Bernotienė, E. Scaffolds and cells for tissue regeneration: Different scaffold pore sizes—Different cell effects. Cytotechnology 2016, 68, 355–369, doi:10.1007/s10616-015-9895-4.
18. Dhandayuthapani, B; Yoshida, Y; Maekawa, T; Kumar, D.S. Polymeric scaffolds in tissue engineering application: A review. Int. J. Polym. Sci. 2011, 2011, doi:10.1155/2011/290602.
19. Loh, Q.L; Choong, C. Three-dimensional scaffolds for tissue engineering applications: Role of porosity and pore size. Tissue Eng. Part B Rev. 2013, 19, 485–502, doi:10.1089/tmb.2012.0437.
20. Jafari, M; Paknejad, Z; Rad, M.R; Motamedian, S.R; Eghbal, M.J; Nadjmi, N; Khojasteh, A. Polymeric scaffolds in tissue engineering: A literature review. J. Biomed. Mater. Res. Part B Appl. Biomater. 2017, 105, 431–459, doi:10.1002/jbm.b.33547.
21. Zhao, F.; Gu, H.; Mi, H.; Rao, C.; Fu, J.; Turng, L. Fabrication of scaffolds in tissue engineering: A review. Front. Mech. Eng. 2018, 13, 107–119, doi:10.1007/s11465-018-0496-8.
22. Walter, S.G.; Oscendorff, R; Schildberg, F.A. Articular cartilage regeneration and tissue engineering models: A systematic review. Arch. Orthop. Trauma Surg. 2019, 139, 305–316, doi:10.1007/s00402-018-3057-z.
23. Zhang, L.; Hu, J.; Athanasiou, K.A. The role of tissue engineering in articular cartilage repair and regeneration. Crit. Rev. Biomed. Eng. 2009, 37, 1–57, doi:10.1615/CritRevBiomedEng.v37.i1-2.10.
24. Kwon, H.; Brown, W.E.; Lee, C.A.; Wang, D.; Faschos, N.; Hu, J.C.; Athanasiou, K.A. Surgical and tissue engineering strategies for articular cartilage and meniscus repair. Nat. Rev. Rheumatol. 2019, 15, 550–570, doi:10.1038/s41584-019-0255-1.
25. Zhao, Z.; Fan, C.; Chen, F.; Sun, Y.; Xia, Y.; Ji, A.; Wang, D.A. Progress in Articular Cartilage Tissue Engineering: A Review on Therapeutic Cells and Macromolecular Scaffolds. Macromol. Biosci. 2020, 20, 1900278, doi:10.1002/mabi.201900278.
26. Sophia Fox, A.J.; Bedi, A.; Rodeo, S.A. The basic science of articular cartilage: Structure, composition, and function. Sports Health 2009, 1, 461–468, doi:10.1177/1941738109350438.
27. Bhosale, A.M.; Richardson, J.B. Articular cartilage: Structure, injuries and review of management. Br. Med Bull. 2008, 87, 77–95, doi:10.1093/bmb/ldn025.
28. Collins, K.H.; Herzog, W.; Macdonald, G.Z.; Reimer, R.A. Obesity, Metabolic Syndrome, and Musculoskeletal Disease: Common Inflammatory Pathways Suggest a Central Role for Loss of Muscle Integrity. Front. Physiol. 2018, 9, doi:10.3389/fphys.2018.00112.
29. Krishnan, Y.; Grodzinsky, A.J. Cartilage diseases. Matrix Biol. 2018, 71–72, 51–69, doi:10.1016/j.matbio.2018.05.005.
30. Oliveira, M.C.; Ph, D.; Vullings, J.; van de Loo, F.A.J.; Ph, D. Osteoporosis and osteoarthritis are two sides of the same coin paid for obesity. Nutrition 2020, 70, 110486, doi:10.1016/j.nut.2019.04.001.
31. Accadbled, F.; Vial, J.; Gauzy, J.S. De Osteochondritis dissecans of the knee. Orthop. Traumatol. Surg. Res. 2018, 104, S97–S105, doi:10.1016/j.otsr.2017.02.016.
32. Barendregt, A.M.; Mazzoli, V.; Berg, J.M. Van Den; Kuijpers, T.W.; Maas, M. T 1 q-mapping for assessing knee joint cartilage in children with juvenile idiopathic arthritis—Feasibility and repeatability. Pediatric Radiol. 2020, 50, 371–379.
33. Brody, L.T. Knee osteoarthritis: Clinical connections to articular cartilage structure and function. Phys. Ther. Sport 2015, 16, 301–316, doi:10.1016/j.ptsr.2014.12.001.
34. Silverwood, V.; Blagovev-Bucknall, M.; Jinks, C.; Jordan, J.L.; Protheroe, J.; Jordan, K.P. Current evidence on risk factors for knee osteoarthritis in older adults: A systematic review and meta-analysis. Osteoarthr. Cartil. 2015, 23, 507–515, doi:10.1016/j.joca.2014.11.019.
35. Johnson, C.I.; Argyle, D.J.; Clements, D.N. In vitro models for the study of osteoarthritis. Vet. J. 2016, 209, 40–49, doi:10.1016/j.tvjl.2015.07.011.
36. Appleton, C.T. Osteoarthritis year in review 2017: Biology. Osteoarthr. Cartil. 2018, 26, 296–303, doi:10.1016/j.joca.2017.10.008.
37. Armiento, A.R.; Alini, M; Stodard, M.J. Articular fibrocartilage—Why does hyaline cartilage fail to repair? Adv. Drug Deliv. Rev. 2019, 146, 289–305, doi:10.1016/j.addr.2018.12.015.
Membranes 2020, 10, 348

38. Bs, C.S.; Kweon, C.Y. Classifications in Brief: Outerbridge Classification of Chondral Lesions. Clin. Orthop. Relat. Res. 2018, 2101–2104, doi:10.1007/s11999-000000000000255.

39. Posadzy, M.; Desimpel, J.; Vanhoenacker, F. Staging of Osteochondral Lesions of the Talus: MRI and Cone Beam CT. J. Belg. Soc. Radiol. 2017, doi:10.5334/jbr-btr.1377.

40. Medvedeva, E.V.; Grebenik, E.A.; Gornostaeva, S.N.; Telpuhov, V.I.; Lychagin, A.V.; Timashev, P.S.; Chagin, A.S. Repair of damaged articular cartilage: Current approaches and future directions. Int. J. Mol. Sci. 2018, 19, 2366, doi:10.3390/ijms19082366.

41. Mirza, U.; Shubeena, S.; Shah, M.S.; Zaffer, B. Microfracture: A technique for repair of chondral defects. J. Entomol. Zool. Stud. 2018, 6, 1092–1097.

42. Dunkin, B.S.; Lattermann, C. New and emerging techniques in cartilage repair: Matrix-induced autologous chondrocyte implantation. Oper. Tech. Sports Med. 2013, 21, 100–107, doi:10.1053/j.otsm.2013.03.003.

43. Brittberg, M. Symposium Scaffold based Autologous Chondrocyte Implantation: The Surgical Technique. Asian J. Arthrosc. 2019, 4, 23–26, doi:10.13107/aja.2456-1169.v04i01.006.

44. Huang, B.J.; Hu, J.C.; Athanasiou, K.A. Cell-based tissue engineering strategies used in the clinical repair of articular cartilage. Biomaterials 2016, 98, 1–22, doi:10.1016/j.biomaterials.2016.04.018.

45. Brittberg, M.; Recker, D.; Ilgenfritz, J.; Saris, D.B.F. Matrix-Applied Characterized Autologous Cultured Chondrocytes Versus Microfracture: Five-Year Follow-up of a Prospective Randomized Trial. Am. J. Sports Med. 2018, 46, 1343–1351, doi:10.1177/0363546518756976.

46. Faby, N.; Alini, M.; Stoddart, M.J. Mechanical stimulation of mesenchymal stem cells: Implications for cartilage tissue engineering. J. Orthop. Res. 2019, 36, 52–63, doi:10.1002/jor.23670.

47. Mastrolia, I.; Foppiani, E.M.; Murgia, A.; Candini, O.; Samarelli, A.V.; Grisendi, G.; Veronesi, E.; Horwitz, E.M.; Dominici, M. Challenges in Clinical Development of Mesenchymal Stromal/Stem Cells: Concise Review. Stem Cells Transl. Med. 2019, 8, 1135–1148, doi:10.1002/sc tm.19-0044.

48. Charlier, E.; Deroyer, C.; Ciregia, F.; Malaise, O.; Neuville, S.; Plener, Z.; Malaise, M.; de Seny, D. Chondrocyte dedifferentiation and osteoarthritis (OA). Biochem. Pharmacol. 2019, 165, 49–65, doi:10.1016/j.bcp.2019.02.036.

49. Lam, A.T.L.; Reuveny, S.; Oh, S.K.W. Human mesenchymal stem cell therapy for cartilage repair: Review on isolation, expansion, and constructs. Stem Cell Res. 2020, 44, 101738, doi:10.1016/j.stemcr.2020.101738.

50. Demoor, M.; Ollitrault, D.; Gomez-Leduc, T.; Bouyoucef, M.; Hervieu, M.; Fabre, H.; Lafont, J.; Denoix, J.M.; Audigié, F.; Mallevin-Gerin, F.; et al. Cartilage tissue engineering: Molecular control of chondrocyte differentiation for proper cartilage matrix reconstruction. Biochim. Et Biophys. Acta Gen. Subj. 2014, 1840, 2414–2440, doi:10.1016/j.bbagenen.2014.02.030.

51. Ahmadi, F.; Giti, R.; Mohammadi-Samani, S.; Mohammadi, F. Biodegradable Scaffolds for Cartilage Tissue Engineering. Calen Med. J. 2017, 6, 70–80, doi:10.22086/GMJ.V6I2.696.

52. Zhan, R.; Ma, J.; Han, J.; Zhang, W.; Ma, J. Mesenchymal stem cell related therapies for cartilage lesions and osteoarthritis. Am. J. Transl. Res. 2019, 11, 6275–6289.

53. Le, H.; Xu, W.; Zhuang, X.; Chang, F.; Wang, Y.; Ding, J. Mesenchymal stem cells for cartilage regeneration. J. Tissue Eng. 2020, 1, 1–22, doi:10.1177/2041731420943839.

54. Koh, Y.G.; Choi, Y.I.; Kwon, O.R. Second-Look Arthroscopic Evaluation of Cartilage Lesions after Mesenchymal Stem Cell Implantation in Osteoarthritic Knees. Am. J. Sports Med. 2014, 42, 1628–1637, doi:10.1177/0363546514529641.

55. Francis, S.L.; Duchi, S.; Onofrillo, C.; Bella, C. Di; Choong, P.F.M. Adipose-Derived Mesenchymal Stem Cells in the Use of Cartilage Tissue Engineering: The Need for a Rapid Isolation Procedure. Stem Cells Int. 2018, 2018, 13–16.

56. Augustyniak, E.; Trzciak, T. The role of growth factors in stem cell-directed chondrogenesis: A real hope for damaged cartilage regeneration. Int. Orthop. 2015, 39, 995–1003, doi:10.1007/s00264-014-2619-0.

57. Kalkan, R.; Nwekwo, C.W.; Adali, T. The Use of Scaffolds in Cartilage Regeneration. Eukaryot. Gene Expr. 2018, 28, 343–348.

58. Panadero, J.A.; Lanceros-Mendez, S.; Ribelles, J.L.G. Differentiation of mesenchymal stem cells for cartilage tissue engineering: Individual and synergistic effects of three-dimensional environment and mechanical loading. Acta Biomater. 2016, 33, 1–12, doi:10.1016/j.actbio.2016.01.037.

59. Eltom, A.; Zhong, G.; Muhammad, A. Scaffold Techniques and Designs in Tissue Engineering Functions and Purposes: A Review. Adv. Mater. Sci. Eng. 2019, doi:10.1155/2019/3429527.
60. Okubo, R.; Asawa, Y.; Watanabe, M.; Nagata, S.; Nio, M. Proliferation medium in three-dimensional culture of auricular chondrocytes promotes effective cartilage regeneration in vivo. Regen. Ther. 2019, 11, 306–315, doi:10.1016/j.reth.2019.10.002.

61. Takahashi, T.; Ogawara, T.; Asawa, Y.; Mori, Y.; Uchiuma, E.; Takato, T.; Hoshi, K. Three-Dimensional Microenvironments Retain Chondrocyte Phenotypes During Proliferation Culture. Tissue Eng. 2007, 13, 1583–1592, doi:10.1089/ten.2006.0322.

62. Schulze-Tanzil, G.; De Souza, P.; Villegas Castrejon, H.; John, T.; Merker, H.J.; Scheid, A.; Shakibaei, M. Redifferentiation of dedifferentiated human chondrocytes in high-density cultures. Cell Tissue Res. 2002, 308, 371–379, doi:10.1007/s00441-002-0562-7.

63. Koh, Y.G.; Lee, J.A.; Kim, Y.S.; Lee, H.Y.; Kim, H.J.; Kang, K.T. Optimal mechanical properties of a scaffold for cartilage regeneration using finite element analysis. J. Tissue Eng. 2019, 10, doi:10.1177/2041731419832133.

64. Irawan, V.; Sung, T.C.; Higuchi, A.; Ikoma, T. Collagen Scaffolds in Cartilage Tissue Engineering and Relevant Approaches for Future Development. Tissue Eng. Regen. Med. 2018, 15, 673–697, doi:10.1007/s13770-018-0135-9.

65. Lai, Y.S.; Chen, W.C.; Huang, C.H.; Cheng, C.K.; Chan, K.K.; Chang, T.K. The effect of graft strength on knee laxity and graft in-situ forces after posterior cruciate ligament reconstruction. PLoS ONE 2015, 10, doi:10.1371/journal.pone.0127293.

66. Conoscenti, G.; Schneider, T.; Stoelzel, K.; Carfi Pavia, F.; Brucato, V.; Goegele, C.; La Carrubba, V.; Schulze-Tanzil, G. PLLA scaffolds produced by thermally induced phase separation (TIPS) allow human chondrocyte growth and extracellular matrix formation dependent on pore size. Mater. Sci. Eng. C 2017, 80, 449–459, doi:10.1016/j.msec.2017.06.011.

67. Zhao, Y.; Tan, K.; Zhou, Y.; Ye, Z.; Tan, W.S. A combinatorial variation in surface chemistry and pore size of three-dimensional porous poly(e-caprolactone) scaffolds modulates the behaviors of mesenchymal stem cells. Mater. Sci. Eng. C 2016, 59, 193–202, doi:10.1016/j.msec.2015.10.017.

68. Nava, M.M.; Draghi, L.; Giordano, C.; Pietrabissa, R. The effect of scaffold pore size in cartilage tissue engineering. J. Appl. Biomater. Funct. Mater. 2016, 14, e223–e229, doi:10.5301/jabfm.5000302.

69. Zhang, Q.; Lu, H.; Kawazoe, N.; Chen, G. Pore size effect of collagen scaffolds on cartilage regeneration. Acta Biomater. 2014, 10, 2005–2013, doi:10.1016/j.actbio.2013.12.042.

70. Matsiko, A.; Gleeson, J.P.; O’Brien, F.J. Scaffold mean pore size influences mesenchymal stem cell chondrogenic differentiation and matrix deposition. Tissue Eng. Part A 2015, 21, 486–497, doi:10.1089/ten.tea.2013.0535.

71. Chwojnowski, A.; Kruk, A.; Wojciechowski, C.; Łukowska, E.; Dulnik, J.; Sajkiewicz, P. The dependence of the membrane structure on the non-woven forming the macropores in the 3D scaffolds preparation. Desalin. Water Treat. 2017, 64, 324–331, doi:10.5004/dwt.2017.11394.

72. Kruk, A.; Gadomska-Gajadzur, A.; Ruśkowski, P.; Chwojnowski, A.; Dulnik, J.; Synoradzki, L. Preparation of biodegradable semi-permeable membranes as 3D scaffolds for cell cultures. Desalin. Water Treat. 2017, 64, 317–323, doi:10.5004/dwt.2017.11415.

73. Przytulskas, M.; Kulikowski, J.L.; Wasyleczko, M.; Chwojnowski, A.; Piętka, D. The evaluation of 3D morphological structure of porous membranes based on a computer-aided analysis of their 2D images. Desalitin. Water Treat. 2018, 128, doi:10.5004/dwt.2018.22569.

74. Sikorska, W.; Wojciechowski, C.; Przytulskas, M.; Rokicki, G.; Wasyleczko, M.; Kulikowski, J.L.; Chwojnowski, A. Polysulfone–polyurethane (PSf–PUR) blend partly degradable hollow fiber membranes: Preparation, characterization, and computer image analysis. Desalin. Water Treat. 2018, 128, doi:10.5004/dwt.2018.23101.

75. Malik, T.; Razzaq, H.; Razzaque, S.; Nawaz, H.; Siddiqua, A.; Siddiqua, M.; Qaisar, S. Design and synthesis of polymeric membranes using water-soluble pore formers: An overview. Polym. Bull. 2019, 76, 4879–4901, doi:10.1007/s00289-018-2616-3.

76. Armiento, A.R.; Stoddart, M.J.; Alimi, N.; Eglin, D. Biomaterials for articular cartilage tissue engineering: Learning from biology. Acta Biomater. 2018, 65, 1–20.

77. Rares, H.; Benea, C.; Earar, K.; Lattanzi, W.; Quercia, V.; Berce, C.; Mohan, A. Collagen Scaffold and Liposuspension Fluid—Derived Stem Cells for the Treatment of Cartilage Defects in a Rabbit Model. Rev. Chim. 2015, 69, 515–520.
78. Gobbi, A.; Whyte, G.P. Long-term Clinical Outcomes of One-Stage Cartilage Repair in the Knee With Hyaluronic Acid—Based Scaffold Embedded With Mesenchymal Stem Cells Sourced From Bone Marrow Aspirate Concentrate. *Am. J. Sports Med.* 2019, 47, 1621–1628, doi:10.1177/0363546519845362.

79. Lin, H.; Beck, A.M.; Fritch, M.R.; Tuan, R.S.; Deng, Y.; Kilroy, E.J.; Tang, Y.; Alexander, P.G. Optimization of photocrosslinked gelatin/hyaluronic acid hybrid scaffold for the repair of cartilage defect. *J. Tissue Eng. Regen. Med.* 2019, 13, 1418–1429, doi:10.1002term.2883.

80. Wen, S.; Hung, K.; Hsieh, K.; Chen, C.; Tsai, C.; Hsu, S. In vitro and in vivo evaluation of chitosan—Gelatin scaffolds for cartilage tissue engineering. *Mater. Sci. Eng. C* 2013, 33, 2855–2863, doi:10.1016/j.msec.2013.03.003.

81. Mittal, H.; Sinha, S.; Singh, B.; Kaur, J.; Sharma, J.; Alhassan, S.M. Recent progress in the structural modification of chitosan for applications in diversified biomedical fields. *Eur. Polym. J.* 2018, 109, 402–434, doi:10.1016/j.eurpolymj.2018.10.013.

82. Chen, S.; Chen, W.; Chen, Y.; Mo, X.; Fan, C. Chondroitin sulfate modified 3D porous electrospun nano fiber scaffolds promote cartilage regeneration. *Mater. Sci. Eng. C* 2021, 118, 1–12, doi:10.1016/j.msec.2020.111312.

83. Zhou, F.; Zhang, X.; Cai, D.; Li, J.; Mu, Q.; Zhang, W.; Zhu, S. Silk fibroin-chondroitin sulfate scaffold with immuno-inhibition property for articular cartilage repair. *Acta Biomater.* 2017, 63, 64–75, doi:10.1016/j.actbio.2017.09.005.

84. Filardo, G.; Drobnic, M.; Perdisa, F.; Kon, E.; Hribernik, M.; Maracci, M. Fibrin glue improves osteochondral scaffold fi xation: Study on the human cadaveric knee exposed to continuous passive motion. *Osteoarthr. Cartil.* 2014, 22, 557–565, doi:10.1016/j.joca.2014.01.004.

85. Kim, J.; Hwang, T.; Lim, D.; Yeon, S.; Lee, Y.; Koh, Y.J. Biochemical and Biophysical Research Communications Chondrogenic differentiation of human hASCs by stiffness control in 3D fi brin hydrogel. *Biochem. Biophys. Res. Commun.* 2020, 522, 213–219, doi:10.1016/j.bbrc.2019.11.049.

86. Chung, C.; Burdick, J.A. Engineering cartilage tissue. *Adv. Drug Deliv. Rev.* 2008, 60, 243–262, doi:10.1016/j.addr.2007.08.027.

87. Asghari, F.; Samiei, M.; Adibkia, K.; Akbarzadeh, A.; Davaran, S. Biodegradable and biocompatible polymers for tissue engineering application: A review. *Artif. Cells Nanomed. Biotechnol.* 2017, 45, 185–192, doi:10.3109/21691401.2016.1146731.

88. Fan, C.; Wang, D. A biodegradable PEG-based micro-cavitary hydrogel as scaffold for cartilage tissue engineering. *Eur. Polym. J.* 2015, 72, 651–660, doi:10.1016/j.eurpolymj.2015.02.038.

89. Janmohammadi, M.; Noroubkash, M.S. International Journal of Polymeric Materials and Electrospun polycaprolactone scaffolds for tissue engineering: A review. *Int. J. Polym. Mater. Polym. Biomater.* 2018, 1–13, doi:10.1080/00994037.2018.1466139.

90. Dao, T.T.; Vu, N.B.; Pham, L.H.; Bui, T.; Le, P.T.; Van Pham, P. In Vitro Production of Cartilage Tissue from Rabbit Bone Marrow-Derived Mesenchymal Stem Cells and Polycaprolactone Scaffold. *Adv. Exp. Med. Biol.* 2017, 1804, 45–60, doi:10.1007/5584.

91. Singhvi, M.S. Polyactic acid: Synthesis and biomedical applications. *J. Appl. Microbiol.* 2012, 127, 1612–1626, doi:10.1111/jam.14290.

92. Silva, D.; Kaduri, M.; Polely, M.; Adir, O.; Krinsky, N.; Shainsky-roitman, J.; Schroeder, A. Mini Review Biocompatibility, biodegradation and excretion of polyactic acid (PLA) in medical implants and theranostic systems. *Chem. Eng. J.* 2018, doi:10.1016/j.cej.2018.01.010.

93. Wen, Y.; Dai, N.; Hsu, S. Acta Biomaterialia Biodegradable water-based polyurethane scaffolds with a sequential release function for cell-free cartilage tissue engineering. *Acta Biomater.* 2019, 88, 301–313, doi:10.1016/j.actbio.2019.02.044.

94. Hung, K.; Tseng, C.; Hsu, S. Synthesis and 3D Printing of Biodegradable Polyurethane Elastomer by a Water-Based Process for Cartilage Tissue Engineering Applications. *Adv. Healthc. Mater.* 2014, 1578–1587, doi:10.1002/adhm.201400018.

95. Budak, K.; Sogut, O.; Sezer, U.A. A review on synthesis and biomedical applications of polyglycolic acid. *J. Polym. Res.* 2020, 27, 1–19.

96. Mahboudi, H.; Soleimani, M.; Enderami, S.E.; Kehtari, M.; Hanaee-Ahvac, H.; Ghanbarian, H.; Bandehpour, M.; Nojehdehi, S.; Mirzaei, S.; Kazemi, B. The effect of nanofibre-based polyethersulfone (PES) scaffold on the chondrogenesis of human induced pluripotent stem cells. *Artif. Cells Nanomed. Biotechnol.* 2018, 46, 1948–1956, doi:10.1080/21691401.2017.1396998.
97. Dudziński, K.; Chwojnowski, A.; Gutowska, M.; Płorczak, M.; Czubak, J.; Łukowska, E.; Wojciechowski, C. Three dimensional polyethersulphone scaffold for chondrocytes cultivation—The future supportive material for articular cartilage regeneration. *Biochern. Biomed. Eng.* 2010, 30, 65–76.

98. Płorczak, M.; Czubak, J.; Hoser, G.; Chwojnowski, A.; Kawiak, J.; Dudziński, K.; Czumińska, K. Repair of articular cartilage full thickness defects with cultured chondrocytes placed on polysulphonic membrane—Experimental studies in rabbits. *Biochern. Biomed. Eng.* 2008, 28, 87–93.

99. Irfan, M.; Idris, A. Overview of PES biocompatible/hemodialysis membranes: PES-blood interactions and modification techniques. *Mater. Sci. Eng. C* 2015, 56, 574–592, doi:10.1016/j.msec.2015.06.035.

100. Filimon, A.; Olaru, N.; Doroftei, F.; Logigan, C.; Dunca, S. Design of Biologically Active Surfaces Based on Functionalized Polysulphones by Electrospinning. *Proceedings* 2019, 41, 35, doi:10.3390/ecsoc-23-06495.

101. Filimon, A.; Avram, E.; Dunca, S. Surface and Interface Properties of Functionalized Polysulphones: Cell-Material Interaction and Antimicrobial Activity. *Polym. Eng. Sci.* 2015, 55, 2184–2894, doi:10.1002/pen.26772.

102. Ye, H.; Zhang, K.; Kai, D.; Li, Z.; Loh, X.J. Polyester elastomers for soft tissue engineering. *Chem. Soc. Rev.* 2018, 47, 4545–4580, doi:10.1039/c8cs00161h.

103. Puppi, D.; Chiellini, F.; Piras, A.M.; Chiellini, E. Progress in Polymer Science Polymeric materials for bone and cartilage repair. *Prog. Polym. Sci.* 2010, 35, 403–440, doi:10.1016/j.progpolymsci.2010.01.006.

104. Chen, F.M.; Liu, X. Advancing biomaterials of human origin for tissue engineering. *Prog. Polym. Sci.* 2016, 53, 86–168, doi:10.1016/j.progpolymsci.2015.02.004.

105. Janoušková, O. Synthetic Polymer Scaffolds for Soft Tissue Engineering, *Physiol. Res.* 2018, 67, S335–S538.

106. He, Y.; Wang, W.R.; Ding, J.D. Effects of L-lactic acid and D,L-lactic acid on viability and osteogenic differentiation of mesenchymal stem cells. *Chin. Sci. Bull.* 2013, 58, 2404–2411, doi:10.1007/s11434-013-5798-y.

107. Xu, Y.; Kim, C.; Saylor, D.M.; Koo, D. Polymer degradation and drug delivery in PLGA-based drug—Polymer applications: A review of experiments and theories. *J. Biomed. Mater. Res. Part B Appl. Biomater.* 2016, 105, 1692–1716, doi:10.1002/jbm.b.33648.

108. Jiang, L.; Xu, L.; Ma, B.; Ding, H.; Tang, C. Effect of component and surface structure on poly (L-lactide-co-e-caprolactone) (PLCA)-based composite membrane. *Compos. Part B* 2019, 174, 107031, doi:10.1016/j.compositesb.2019.107031.

109. Prasanna, S.; Narayan, B.; Rastogi, A.; Srivastava, P. Design and evaluation of chitosan/poly (L-lactide)/pectin based composite scaffolds for cartilage tissue regeneration. *Int. J. Biol. Macromol.* 2018, 112, 909–920, doi:10.1016/j.ijbiomac.2018.02.049.

110. Nofar, M.; Saciligil, D.; Carreau, P.J.; Kamal, M.R.; Heuzey, M.C. Poly (lactic acid) blends: Processing, properties and applications. *Int. J. Biol. Macromol.* 2019, 125, 307–360, doi:10.1016/j.ijbiomac.2018.12.002.

111. Jeukens, R.M.; Roth, A.K.; Peters, R.J.R.W.; van Donkelaar, C.C.; Thies, J.C.; van Rhiijn, L.W.; Emans, P.J. Polymers in cartilage defect repair of the knee: Current status and future prospects. *Polymers* 2016, 8, 219.

112. Daranarong, D.; Techakool, P.; Intatue, W.; Daengngern, R.; Thomson, K.A.; Molloy, R.; Kungwan, N.; Foster, L.J.R.; Boonyawan, D.; Punyodom, W. Effect of surface modification of poly(l-lactide-co-e-caprolactone) membranes by low-pressure plasma on support cell biocompatibility. *Surf. Coat. Technol.* 2016, 306, 328–335, doi:10.1016/j.surfcoat.2016.07.058.

113. Guo, C.; Cai, N.; Dong, Y. Duplex surface modification of poly(olactic acid) scaffold. *Mater. Lett.* 2013, 94, 11–14, doi:10.1016/j.matlet.2012.11.092.

114. Tsurumi, T.; Fuse, M. Enhancement of apatite precipitation on an alkaline hydrolyzed poly (lactic acid-co-Caprolactone) film in simulated body fluid. *J. Hard Tissue Biol.* 2014, 23, 15–19, doi:10.2485/jhtb.23.15.

115. Setayeshmehr, M.; Esfandiari, E.; Rafieinia, M.; Hashemiben, B.; Taheri-kafrani, A.; Samadikuchaksaraei, A. Hybrid and Composite Scaffolds Based on Extracellular. *Tissue Eng. Part B Rev.* 2019, 25, 202–224, doi:10.1089/ten.teb.2018.0245.

116. Zhang, X.; Wu, Y.; Pan, Z.; Sun, H.; Wang, J.; Yu, D.; Zhu, S.; Dai, J.; Chen, Y.; Tian, N.; et al. The effects of lactate and acid on articular chondrocytes function: Implications for polymeric cartilage scaffold design. *Acta Biomater.* 2016, 42, 329–340, doi:10.1016/j.actbio.2016.06.029.

117. Lu, T.; Li, Y.; Chen, T. Techniques of fabrication and construction three-dimensional scaffold. *Int. J. Nanomed.* 2013, 8, 337–350, doi:10.2147/IJNN.S38635.

118. Dutta, R.C.; Dey, M.; Dutta, A.K.; Basu, B. Competent processing techniques for scaffolds in tissue engineering. *Biotechnol. Adv.* 2017, 35, 240–250, doi:10.1016/j.biotechadv.2017.01.001.
119. Mannella, G.A.; Conoscenti, G.; Pavia, F.C.; Carrubba, V. La; Brucato, V. Preparation of polymeric foams with a pore size gradient via Thermally Induced Phase Separation (TIPS). Mater. Lett. 2015, 160, 31–33, doi:10.1016/j.matlet.2015.07.055.

120. Buzarovska, A.; Gualandi, C.; Parrilli, A.; Scandola, M. Effect of TiO2 nanoparticle loading on Poly(L-lactic acid) porous scaffolds fabricated by TIPS. Compos. Part B 2015, doi:10.1016/j.compositesb.2015.07.016.

121. Sultana, N.; Hassana, M.I.; Ridzuana, N.; Ibrahima, Z.; Sooon, C.F. Fabrication of Gelatin Scaffolds using Thermally Induced Phase Separation Technique. Int. J. Eng. 2018, 31, 1302–1307, doi:10.5829/ije.2018.31.08b.19.

122. Structure, M.M.; Mechanical, H.; Kim, J.; Shin, K.; Koh, Y.; Hah, M.J.; Moon, J.; Kim, H. Production of Poly(ε-Caprolactone)/Hydroxyapatite Composite Scaffolds with a Tailored Macro/Micro-Porous Structure, High Mechanical Properties, and Excellent Bioactivity. Materials 2017, 10, 1123, doi:10.3390/ma1011123.

123. Georgiadou, S.; Katsogiannis, K.A.G.; Vladisavljevic, G.T. Porous electrosprun polycaprolactone (PCL) fibres by phase separation. Eur. Polym. J. 2015, 69, 284–295, doi:10.1016/j.eurpolymj.2015.01.028.

124. Huang, C.; Thomas, N.L. Fabricating Porous Poly(lactic acid) Fibres via Electrosprinning. Eur. Polym. J. 2017, doi:10.1016/j.eurpolymj.2017.12.025.

125. Prasad, A.; Sankar, M.R.; Katiyar, V. ScienceDirect State of Art on Solvent Casting Particulate Leaching Method for Orthopedic Scaffolds Fabrication. Mater. Today Proc. 2017, 4, 898–907.

126. Plisko, T.V.; Penkova, A.V.; Burts, K.S.; Bilyukchevich, A.V.; Dmitrenko, M.E.; Melnikova, G.B.; Atta, R.R.; Mazur, A.S.; Zolotarev, A.A.; Missyul, A.B. Effect of Pluronic F127 on porous and dense membrane structure formation via non-solvent induced and evaporation induced phase separation. J. Membr. Sci. 2019, 580, 336–349, doi:10.1016/j.memsci.2019.03.028.

127. Gadomska-Gajadhur, A.; Kruk, A.; Ruśkowski, P.; Sajkiewicz, P.; Dulnik, J.; Chwojnowski, A. Original method of imprinting pores in scaffolds for tissue engineering. Polym. Adv. Technol. 2020, 1–13, doi:10.1002/pat.5091.

128. Taylor, P.; Yang, Q.; Chen, L.; Shen, X.; Tan, Z. Preparation of Polycaprolactone Tissue Engineering Scaffolds by Improved Solvent Casting/Particulate Leaching Method Preparation of Polycaprolactone Tissue Engineering Scaffolds by Improved Solvent Casting/Particulate Leaching. J. Macromol. Sci. Part B Phys. 2006, 45, 1171–1181, doi:10.1080/0022340601976873.

129. Sharifi, F.; Irani, S.; Azadegan, G.; Pezeshki-modarees, M. Bioactive Carbohydrates and Dietary Fibre Co-electrosprun gelatin-chondroitin sulfate/polycaprolactone nanofibrous scaffolds for cartilage tissue engineering. Bioact. Carbohydr. Diet. Fibre 2020, 22, 100215, doi:10.1016/j.bcdf.2020.100215.

130. Zhou, Y.; Chyu, J.; Zumwalt, M. Recent Progress of Fabrication of Cell Scaffold by Electrosprinning Technique for Articular Cartilage Tissue Engineering. Int. J. Biomater. 2018, 2018, doi:10.1155/2018/1953636.

131. Girão, A.F.; Semitela, A.; Ramalho, G.; Completo, A.; Marques, P.A.A.P. Mimicking nature- Fabrication of 3D anisotropic electrosprun polycaprolactone scaffolds for cartilage tissue engineering applications. Compos. Part B 2018, doi:10.1016/j.compositesb.2018.08.001.

132. Bdikin, I.; Marques, P.A.A.P. Electrospinning of bioactive polycaprolactone-gelatin nanofibres with increased pore size for cartilage tissue engineering applications. J. Biomater. Appl. 2020, 35, 459–470, doi:10.1177/0885328220940194.

133. Wang, Z.; Wan, L.; Liu, Z.; Huang, X.; Xu, Z. Enzymatic Enzyme immobilization on electrosprun polymer nanofibers: An overview. J. Mol. Catal. B 2009, 56, 189–195, doi:10.1016/j.molcatb.2008.05.005.

134. Yang, Z.; Si, J.; Cui, Z.; Ye, J.; Wang, X.; Wang, Q.; Peng, K.; Chen, W.; Chen, S. Biomimetic composite scaffolds based on surface modification of polydopamine on electrosprun poly (lactic acid)/cellulose nanofibres. Carbohydr. Polym. 2017, 174, 750–759, doi:10.1016/j.carbpol.2017.07.010.

135. Canton, T.T.; Kunert, L.R.; Suellen, B.; Ana, I.; Serafini, P. Nonwoven membranes for tissue engineering: An overview of cartilage, epithelium, and bone regeneration. J. Biomater. Sci. Polym. Ed. 2019, 30, 1026–1049, doi:10.1080/09205063.2019.1620592.

136. Abdelaal, O.A.M.; Darwish, S.M.H. Review of Rapid Prototyping Techniques for Tissue Engineering Scaffolds Fabrication. In Characterization and Development of Biosystems and Biomaterials; Springer: Berlin/Heidelberg, Germany, 2013; Volume 29, pp. 33–54, doi:10.1007/978-3-642-31470-4.
137. Li, K.; Wang, D.; Zhao, K.; Song, K.; Liang, J. Electrohydrodynamic jet 3D printing of PCL/PVP composite scaffold for cell culture. *Talanta* **2020**, *120750*, doi:10.1016/j.talanta.2020.120750.

138. Daly, A.C.; Freeman, F.E.; Gonzalez-fernandez, T.; Critchley, S.E.; Nulty, J.; Kelly, D.J. 3D Bioprinting for Cartilage and Osteochondral Tissue Engineering. *Adv. Healthc. Mater.* **2017**, *6*, 1700298, doi:10.1002/adhm.201700298.

139. Seung, J.; Sang, H.; Jung, H.; Lee, H.; Hong, H.; Jin, Y.; Ji, Y.; Joo, O.; Hee, S.; Hum, C. 3D-printable photocurable bioink for cartilage regeneration of tonsil-derived mesenchymal stem cells. *Addit. Manuf.* **2020**, *33*, 101136, doi:10.1016/j.addma.2020.101136.

140. Marycz, K.; Smieszek, A.; Targonska, S.; Walsh, A.; Szustakiewicz, K.; Wiglus, R.J. Three dimensional (3D) printed PLA with nano-hydroxyapatite doped with europium(III) ions (nHAp-PLLA@Eu3+) composite for osteochondral defect regeneration and theranostics. *Mater. Sci. Eng. C* **2020**, *110634*, doi:10.1016/j.msec.2020.110634.

141. Wu, J.; Yang, R.; Zheng, J.; Pan, L.; Liu, X. Fabrication and improvement of PCL/alginate/PAAm scaffold via selective laser sintering for tissue engineering. *Micro Nano Lett.* **2019**, *14*, 852–855, doi:10.1049/mnl.2018.5806.

142. Aisenbrey, E.A.; Tomashke, A.; Kleinjan, E.; Muralidharan, A.; Pascual-garrido, C.; Mcleod, R.R.; Ferguson, V.L.; Bryant, S.J. A Stereolithography-Based 3D Printed Hybrid Scaffold for In Situ Cartilage Defect Repair. *Macromol. Biosci.* **2017**, *18*, 1–8, doi:10.1002/mabi.201700267.

143. Gauvin, R.; Chen, Y.; Woo, J.; Somani, P.; Zorlutuna, P.; Nichol, J.W.; Bae, H.; Chen, S.; Khademhosseini, A. Biomaterials Microfabrication of complex porous tissue engineering scaffolds using 3D projection stereolithography. *Biomaterials* **2012**, *33*, 3824–3834, doi:10.1016/j.biomaterials.2012.01.048.

144. Charterain, N.A.; Williams, C.B.; Whittington, A.R. A review on fabricating tissue scaffolds using vat photopolymerization. *Acta Biomater.* **2018**, *74*, 90–111, doi:10.1016/j.actbio.2018.05.010.

145. Longley, R.; Ferreira, A.M.; Gentile, P. Recent Approaches to the Manufacturing of Biomimetic Multi-Phasic Scaffolds for Osteochondral Regeneration. *Int. J. Mol. Sci.* **2018**, *19*, 1–17, doi:10.3390/ijms19061755.

146. Mohanty, A.K. Improving the Impact Strength and Heat Resistance of 3D Printed Models: Structure, Property, and Processing Correlations during Fused Deposition Modeling (FDM) of Poly(Lactic Acid). *ACS Omega* **2018**, doi:10.1021/acsomega.8b00129.

147. Murphy, S.V.; Atala, A. 3D bioprinting of tissues and organs. *Nat. Biotechnol.* **2014**, *32*, 773–785, doi:10.1038/nbt.2958.

148. Cheng, A.; Schwartz, Z.; Kahn, A.; Li, X.; Shao, Z.; Sun, M.; Ao, Y.; Boyan, B.D.; Chen, H.; Antonio, S.; et al. Advances in Porous Scaffold Design for Bone and Cartilage Tissue Engineering and Regeneration. *Tissue Eng. Part B Rev.* **2018**, *25*, 14–29, doi:10.1089/ten.TEB.2018.0119.

149. Chen, W.; Xu, Y.; Liu, Y.; Wang, Z.; Li, Y.; Jiang, G.; Mo, X.; Zhou, G. Three-dimensional printed electrospun fi ber-based scaffold for cartilage regeneration. *Mater. Des.* **2019**, *179*, 107886, doi:10.1016/j.matdes.2019.107886.

150. Garrigues, N.W.; Little, D.; Sanchez-adams, J.; Ruch, D.S.; Guilak, F. Electrospun cartilage-derived matrix scaffolds for cartilage tissue engineering. *J. Biomed. Mater. Res. Part A* **2014**, *59784*, 28–30, doi:10.1002/jbma.a.35068.

151. Xu, T.; Binder, K.W.; Albanna, M.Z.; Dice, D.; Zhao, W.; Yoo, J.J.; Atala, A. Hybrid printing of mechanically and biologically improved constructs for cartilage tissue engineering applications. *Biofabrication* **2013**, *5*, 1–11, doi:10.1088/1758-5082/5/1/015001.

152. Nielsen, L.E. Polymer Reviews Cross-Linking—Effect on Physical Properties of Polymers. *J. Macromol. Sci. Part C* **2008**, *3*, 69–103.

153. Rofiqoh, N.; Putri, E.; Wang, X.; Chen, Y.; Li, X.; Kawazoe, N.; Chen, G. Preparation of PLGA-collagen hybrid scaffolds with controlled pore structures for cartilage tissue engineering. *Prog. Nat. Sci. Mater. Int.* **2020**, doi:10.1016/j.pnsc.2020.07.003.

154. Laurent, P. Suitability of a PLCL fibrous scaffold for soft tissue engineering applications: A combined biological and mechanical characterisation. *J. Biomat. Appl.* **2018**, *32*, 1–13, doi:10.1177/0959053017757064.

155. Ahmed, E.M. Hydrogel: Preparation, characterization, and applications: A review. *J. Adv. Res.* **2015**, *6*, 105–121, doi:10.1016/j.jare.2013.07.006.
156. Bistolfi, A.; Ferracini, R.; Galletta, C.; Tosto, F.; Sgarminato, V.; Digo, E.; Vernè, E.; Massé, A. Regeneration of articular cartilage: Scaffold used in orthopedic surgery. A short handbook of available products for regenerative joints surgery. *Clin. Sci. Res. Rep.* **2017**, *1*, 1–7, doi:10.15761/CSRR.1000101.

157. Müller, S. Repair of Focal Cartilage Defects With Scaffold-Assisted Autologous Chondrocyte Grafts. *Am. J. Sports Med.* **2011**, *39*, 1697–1705, doi:10.1177/0363546511403279.

158. Tsai, M.; Hung, K.; Hung, S.; Hsu, S. Evaluation of biodegradable elastic scaffolds made of anionic polyurethane for cartilage tissue engineering. *Colloids Surf. B Biointerfaces* **2015**, *125*, 34–44, doi:10.1016/j.colsurfb.2014.11.003.

159. Borsøe, B.; Casper, C.; Foldager, B.; Møller, O.; Lind, M. A novel nano-structured porous polycaprolactone scaffold improves hyaline cartilage repair in a rabbit model compared to a collagen type I/III scaffold: In vitro and in vivo studies. *Knee Surg. Sports Traumatol. Arthrosc.* **2012**, *20*, 1192–1204, doi:10.1007/s00167-011-1692-9.

160. Theodoridis, K.; Aggelidou, E.; Vavilis, T.; Manthou, M.E.; Demiri, E.C.; Boukla, A.; Salpistis, C. Hyaline cartilage next generation implants from adipose—Tissue—Derived mesenchymal stem cells: Comparative study on 3D—Printed polycaprolactone scaffold patterns. *J. Tissue Eng. Regen. Med.* **2019**, *13*, 342–355, doi:10.1002/term.2798.

161. Płonczak, M.; Czubak, J. Culture of Human Autologous Chondrocytes on Polysulphonic Membrane—Preliminary Studies. *Biocybern. Biomed. Eng.* **2012**, *32*, 63–67, doi:10.1016/S0208-5216(12)70042-6.

162. Płonczak, M. The Value of Autogenous Cartilage Cell Transplants in the Experimental Treatment of Articular Cartilage Defects in Rabbits. Doctoral Dissertations, Medical Centre of Postgraduate Education in Warsaw, Warszawa, 28 May 2008.

163. Taylor, P.; Jung, Y.; Kim, S.H.; You, H.J. Application of an elastic biodegradable poly (L-lactide-co-ε-caprolactone) scaffold for cartilage tissue regeneration. *J. Biomater. Sci. Polym. Ed.* **2012**, *23*, 1073–1085, doi:10.1163/15685620124499336.

164. Siclari, A.; Mascaro, G.; Kaps, C.; Boux, E. A 5-Year Follow-Up After Cartilage Repair in the Knee Using a Platelet—Rich Plasma-Immersed Polymer-Based Implant. *Open Orthop. J.* **2014**, *8*, 346–354.

165. Asadi, N.; Alizadeh, E.; Rahmani, A.; Bakhsheyesh, D.; Mostafavi, E.; Akbarzadeh, A.; Davaran, S. Fabrication and in Vitro Evaluation of Nanocomposite Hydrogel Sca ffolds Based on Gelatin/PCL—PEG—PCL for Cartilage Tissue Engineering. *ACS Omega* **2019**, *4*, 449–457, doi:10.1021/acs.omega.8b02593.

166. He, Y.; Liu, W.; Guan, L.; Chen, J.; Duan, L.; Jia, Z.; Huang, J.; Li, W.; Liu, J.; Xiong, J.; et al. A 3D-Printed PLCL Scaffold Coated with Collagen Type I and Its Biocompatibility. *Biomater. Res. Int.* **2018**, *2018*, 1–10, doi:10.1155/2018/5147156.

167. Haaparanta, A.; Ja, E.; Fatih, I.; Ville, C.; Kiviranta, I.; Kelloma, M. Preparation and characterization of collagen/PLA, chitosan/PLA, and collagen/chitosan/PLA hybrid scaffolds for cartilage tissue engineering. *J. Mater. Sci. Mater. Med.* **2014**, *25*, 1129–1136, doi:10.1007/s10856-013-1529-5.

168. Nogami, M.; Kimura, T.; Seki, S.; Matsui, Y.; Yoshida, T.; Koike-Soko, C.; Okabe, M.; Motomura, H.; Gejo, R.; Nikaido, T. A human amnion derived extracellular matrix coated cell free scaffold for cartilage repair: In vitro and in vivo studies. *Tissue Eng. Part A* **2016**, *22*, 680–688.

169. Yamanaka, K.; Yamamoto, K.; Sakai, Y.; Suda, Y.; Shigemitsu, Y. Seeding of mesenchymal stem cells into inner part of interconnected porous biodegradable scaffold by a new method with a filter paper. *Dent. Mater.* **2015**, *31*, 74–85, doi:10.1016/dtm.2013.330.

170. Chen, C.; Shyu, V.B.; Chen, J.; Lee, M. Selective laser sintered poly-ε-caprolactone scaffold hybridized with collagen hydrogel for cartilage tissue engineering. *Biofabrication* **2014**, *01*, 015004, doi:10.1088/1758-5082/6/1/015004.

171. Taylor, P.; Li, C.; Wang, L.; Yang, Z.; Kim, G. A Viscoelastic Chitosan-Modified Three-Dimensional Porous Poly (L-Scaffold for Cartilage Tissue Engineering. *Biomater. Sci.* **2012**, *405–424*, doi:10.1038/n02050610X51970.

172. Liao, J.; Qu, Y.; Chu, B.; Zhang, X.; Qian, Z. Biodegradable CSMA/PECA/Graphene Porous Hybrid Scaffold for Cartilage Tissue Engineering. *Sci. Rep.* **2015**, *5*, 9879, doi:10.1038/srep09879.

173. Urbanek, O.; Kolbuk, D.; Wrobel, M. International Journal of Polymeric Materials and Articular cartilage: New directions and barriers of scaffolds development—Review. *Int. J. Polym. Mater. Polym. Biomater.* **2018**, *1–15*, doi:10.1080/00914037.2018.1452224.

174. Tan, S.I.; Jun, S.; Tho, W.; Tho, K.S. Biological resurfacing of grade IV articular cartilage ulcers in knee joint with Hyalofast. *J. Orthop. Surg.* **2020**, *28*, 1–7, doi:10.1177/2309499020905158.
175. Sofu, H.; Camurcu, Y.; Uçpınar, H.; Ozcan, S.; Yurten, H.; Sahin, V. Clinical and radiographic outcomes of chitosan-glycerol phosphate / blood implant are similar with hyaluronic acid-based cell-free scaffold in the treatment of focal osteochondral lesions of the knee joint. Knee Surg. Sports Traumatol. Arthrosc. 2019, 27, 773–781, doi:10.1007/s00167-018-5079-z.

176. Park, Y.B.; Ha, C.W.; Lee, C.H.; Yoon, Y.C.; Park, Y.G. Cartilage Regeneration in Osteoarthritic Patients by a Composite of Allogeneic Umbilical Cord Blood-Derived Mesenchymal Stem Cells and Hyaluronate Hydrogel: Results from Trial I and Results from a a Clinical Clinical Trial for Safety and Concept. Stem Cells Transl. Med. 2017, 6, 613–621.

177. Negoro, T.; Takagaki, Y.; Okura, H.; Matsuyama, A. Trends in clinical trials for articular cartilage repair by cell therapy. NPJ Regen. Med. 2018, doi:10.1038/s41536-018-0055-2.

178. Hoffman, A.S. Hydrogels for biomedical applications. Adv. Drug Deliv. Rev. 2012, 64, 18–23, doi:10.1016/j.addr.2012.09.010.

179. Ren, K.; He, C.; Xiao, C.; Li, G.; Chen, X. Injectable glycopolypeptide hydrogels as biomimetic scaffolds for cartilage tissue engineering. Biomaterials 2015, 51, 238–249, doi:10.1016/j.biomaterials.2015.02.026.

180. Yang, F.; Zhao, J.; Koshut, W.J.; Watt, J.; Riboh, J.C.; Gall, K.; Wiley, B.J. A Synthetic Hydrogel Composite with the Mechanical Behavior and Durability of Cartilage. Adv. Funct. Mater. 2020, 30, 1–8, doi:10.1002/adfm.202003451.

181. Rosenzweig, D.H.; Carelli, E.; Steffen, T.; Jarzem, P.; Haglund, L. 3D-printed ABS and PLA scaffolds for cartilage and nucleus pulposus tissue regeneration. Int. J. Mol. Sci. 2015, 16, 15118–15135, doi:10.3390/ijms160715118.

182. Cristian, M.; Conde, M.; Demarco, F.F.; Alcazar, J.C.; Nör, J.E.; Beatriz, S. Influence of Poly-L-Lactic Acid Scaffold’s Pore Size on the Proliferation and Differentiation of Dental Pulp Stem Cells. Braz. Dent. J. 2015, 26, 93–98.

183. Oh, S.H.; Kim, T.H.; Im, G. II; Lee, J.H. Investigation of pore size effect on chondrogenic differentiation of adipose stem cells using a pore size gradient scaffold. Biomacromolecules 2010, 11, 1948–1955, doi:10.1021/bm100199m.

184. Moura, C.S.; Silva, J.C.; Fernandes, P.R.; Lobato, C.; Manuel, J.; Cabral, S.; Linhardt, R.; Bártilo, P.J.; Ferreira, F.C. Chondrogenic differentiation of mesenchymal stem/stromal cells on 3D porous poly (ε-caprolactone) scaffolds: Effects of material alkaline treatment and chondroitin sulfate supplementation. J. Biosci. Bioeng. 2020, 129, 756–764, doi:10.1016/j.jbiosc.2020.01.004.

185. Sonomoto, K.; Yamaoka, K.; Kaneko, H.; Yamagata, K. Spontaneous Differentiation of Human Mesenchymal Stem Cells on Poly-Lactic-Co-Glycolic Acid Nano-Fiber Scaffold. PLoS ONE 2016, 11, e0153231, doi:10.1371/journal.pone.0153231.

186. He, A.; Liu, L.; Luo, X.; Liu, Y.; Liu, Y.; Liu, F.; Wang, X. Repair of osteochondral defects with in vitro engineered cartilage based on autologous bone marrow stromal cells in a swine model. Sci. Rep. 2017, 12, doi:10.1038/srep40489.

187. Zhang, Y.; Yang, F.; Liu, K.; Shen, H.; Zhu, Y.; Zhang, W.; Liu, W.; Wang, S.; Cao, Y.; Zhou, G. The impact of PLGA scaffold orientation on in vitro cartilage differentiation. Biomaterials 2012, 33, 2926–2935, doi:10.1016/j.biomaterials.2012.01.006.

188. Duan, P.; Pan, Z.; Cao, L.; He, Y.; Wang, H.; Qu, Z.; Dong, J. The effects of pore size in bilayered poly (lactide- ε-glycolide) scaffolds on restoring osteochondral defects in rabbits. J. Biomed. Mater. Res. Part A: Off. J. Soc. Biomater. Jpn. Soc. Biomater. Aust. Soc. Biomater. Korean Soc. Biomater. 2013, 102, 180–192.

189. Jonnalagadda, J.B.; Rivero, L.V.; Dertien, J.S. In vitro chondrocyte behavior on porous biodegradable poly (ε-caprolactone)/polyglycolic acid scaffolds for articular chondrocyte adhesion and proliferation. J. Biomater. Sci. 2015, 37–41, doi:10.1080/09205063.2015.1015864.

190. Safinsha, S.; Ali, M.M. Composite scaffolds in tissue engineering. Mater. Today Proc. 2020, 24, 2318–2329, doi:10.1016/j.matpr.2020.03.761.

191. Li, L.; Li, J.; Guo, J.; Zhang, H.; Zhang, X.; Yin, C. 3D Molecularly Functionalized Cell-Free Biomimetic Scaffolds for Osteochondral Regeneration. Adv. Funct. Mater. 2019, 29, 1807356, doi:10.1002/adfm.201807356.

192. Setayeshmehr, M.; Esfandiari, E.; Hashemibenii, B.; Hossein, A. Chondrogenesis of human adipose-derived mesenchymal stromal cells on the [devitalized costal cartilage matrix/poly (vinyl alcohol)/fi brin] hybrid scaffolds. Eur. Polym. J. 2019, 118, 528–541, doi:10.1016/j.eurpolymj.2019.04.044.
193. Tavakoli, E.; Mehdikhani-nahrkhalaji, M.; Hashemi-beni, B.; Zargar-kharazi, A. Preparation, Characterization and Mechanical Assessment of Poly (Lactide-Co-Glycolide)/Hyaluronic Acid/Fibrin/Bioactive Glass Nano-Composite Scaffolds for Cartilage Tissue Engineering Applications. *Procedia Mater. Sci.* 2015, 11, 124–130, doi:10.1016/j.mspro.2015.11.126.

194. Mintz, B.R.; Cooper, A.C., Jr. Hybrid hyaluronic acid hydrogel/poly (e-caprolactone) scaffold provides mechanically favorable platform for cartilage tissue engineering studies. *J. Biomed. Mater. Res. Part A* 2013, 102, 2918–2926, doi:10.1002/jbm.a.34957.

195. He, X.; Fu, W.; Feng, B.; Wang, H.; Liu, Z.; Yin, M.; Wang, W. Electrospun collagen—poly (l-lactic acid-co-e-caprolactone) membranes for cartilage tissue engineering. *Regen. Med.* 2013, 8, 425–436.

196. Paatela, T.; Meller, A.; Muhonen, V.; Salonius, E.; Haaparanta, A.; Elina, J. Articular Cartilage Repair With Recombinant Human Type II Collagen/Polylactide Scaffold in a Preliminary Porcine Study. *J. Orthop. Res.* 2016, 34, 745–753, doi:10.1002/jor.23099.

197. Levorson, E.J.; Hu, O.; Mountzaris, P.M.; Kasper, F.K.; Mikos, A.G. Cell Derived Polymer / Extracellular Matrix Composite Scaffolds for Cartilage Regeneration, Part 2: Construct Devitalization and Determination of Chondroinductive Capacity. *Tissue Eng. Part C Methods* 2014, 20, 358–372.

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