

## STEM CELLS FOR TOOTH ENGINEERING

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### Abstract

Tooth development results from sequential and reciprocal interactions between the oral epithelium and the underlying neural crest-derived mesenchyme. The generation of dental structures and/or entire teeth in the laboratory depends upon the manipulation of stem cells and requires a synergy of all cellular and molecular events that finally lead to the formation of tooth-specific hard tissues, dentin and enamel. Although mesenchymal stem cells from different origins have been extensively studied in their capacity to form dentin *in vitro*, information is not yet available concerning the use of epithelial stem cells. The odontogenic potential resides in the oral epithelium and thus epithelial stem cells are necessary for both the initiation of tooth formation and enamel matrix production. This review focuses on the different sources of stem cells that have been used for making teeth *in vitro* and their relative efficiency. Embryonic, post-natal or even adult stem cells were assessed and proved to possess an enormous regenerative potential, but their application in dental practice is still problematic and limited due to various parameters that are not yet under control such as the high risk of rejection, cell behaviour, long tooth eruption period, appropriate crown morphology and suitable colour. Nevertheless, the development of biological approaches for dental reconstruction using stem cells is promising and remains one of the greatest challenges in the dental field for the years to come.

**Keywords:** Stem cells, odontoblast, dentin, ameloblasts, enamel, tooth, incisor, human.

### Introduction

Teeth are highly mineralized organs resulting from sequential and reciprocal interactions between the oral epithelium and the underlying cranial neural crest-derived mesenchyme (Duailibi *et al.*, 2006; Mitsiadis, 2001) (Fig. 1). Tissue recombination experiments point out that the oral epithelium contains the inductive capability for odontogenesis. This potential allows conditioning of the underlying mesenchyme, which in turn regulates the differentiation of epithelial cells. The importance of cranial neural crest-derived cells in odontogenesis has been shown in experiments where transplantation of mouse neural crest cells into chick embryos allowed growth of tooth germs (Mitsiadis *et al.*, 2003). Numerous growth factors have been shown to be involved in different stages of the embryonic tooth development (i.e. initiation, morphogenesis, cytodifferentiation). Members of the Transforming Growth Factor beta (TGF $\beta$ ) superfamily such as Bone Morphogenic Protein 2 (BMP-2) and BMP-4 are key signalling molecules in regulating epithelial-mesenchymal interactions during odontogenesis (Kratochwil *et al.*, 1996; Nadiri *et al.*, 2004; Vainio *et al.*, 1993). Molecules of the Fibroblast Growth Factor (FGF) family such as FGF-3, FGF-4, FGF-8 and FGF-10 are involved in cell proliferation and regulate expression of specific target genes in teeth (Bei and Maas, 1998; Kettunen *et al.*, 1998, 2000). Wnt proteins such as Wnt-3, Wnt-7b, Wnt-10a and Wnt-10b have essential roles as regulators of cell proliferation, migration and differentiation during tooth initiation and morphogenesis (Dassule and McMahon, 1998). Other diffusible factors such as sonic hedgehog (shh) also contribute to both initiation and subsequent dental morphogenesis (Khan *et al.*, 2007).

Two major cell types are involved in dental hard tissue formation: the mesenchyme-originated odontoblasts that are responsible for the production of dentin and the epithelium-derived ameloblasts that form the enamel (Fig. 1). Odontoblasts are columnar post-mitotic cells that form a layer in contact with the dentin. Odontoblastic processes are formed at their distal part, penetrate the dentin and participate in the secretion of dentin matrix and minerals. The matrix is composed of collagen (90%) and non-collagenous proteins such as Dentin Sialophosphoprotein (DSPP) and Dentin Matrix Protein 1 (DMP-1). The deposition of apatite minerals on this matrix gives rise to the mature calcified dentin.

Enamel is secreted by ameloblasts along the dentino-enamel junction. Enamel is mainly composed of hydrophobic proteins such as amelogenin, ameloblastin,

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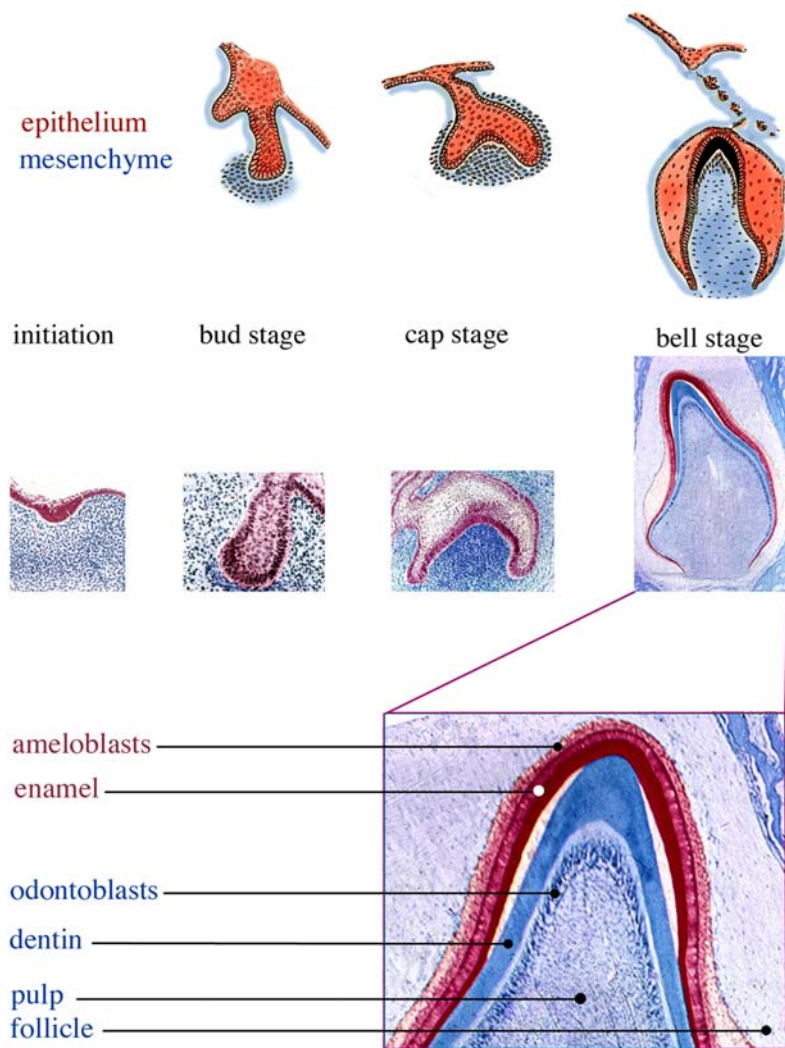
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**Figure 1.** Schematic illustration and histological sections showing the different stages of tooth development in humans. The sections were stained with haematoxylin. Orange colour indicates epithelial tissues and epithelial derivatives (ameloblasts and enamel) while blue colour shows mesenchymal tissues and mesenchyme derivatives (odontoblasts, dentin, dental pulp and follicle).

enamelin, amelotin, tuftelin and ODAM (odontogenic ameloblast associated proteins) (Sire *et al.*, 2007). Shortly after enamel deposition, the formation of the root starts as a consequence of cell proliferation in the inner and outer dental epithelia at the cervical loop area. Cells from the dental follicle give rise to cementoblasts (forming the cementum that covers the dentin of the root), fibroblasts (generating the periodontal ligament) and osteoblasts (elaborating the alveolar bone). Cementum, periodontal ligament and alveolar bone are the periodontal tissues that support teeth into the oral cavity.

Tooth loss or absence is a common and frequent situation that can result from numerous pathologies such as periodontal and carious diseases, fractures, injuries or even genetic alterations. In most cases this loss is not critical, but for aesthetical, psychological and medical reasons (e.g. genetic aberrations) replacement of the missing teeth is important. Recent efforts made in the field of biomaterials have led to the development of dental implants composed of biocompatible materials such as titanium that can be inserted in the maxillary and/or mandibular bone to replace the missing teeth. However, implants are still not completely satisfactory and their successful use greatly depends on their osteointegration. The quantity and quality of the bone, as well as its interaction with the surface of the implant are some crucial

parameters that can influence the achievement of the operation. Although innovative materials and techniques (e.g. surface treatment) have been used for the improvement of implant osteointegration (reviewed by Le Guehennec *et al.*, 2007), the metal/bone interface does not ensure complete integration of the implant, thus reducing its performance and long-term stability. Furthermore, dental implant technology is dependent on bone volume, as devices (i.e., peg) can be implanted only in patients possessing a sufficient amount of bone. Quite often there is a need for alveolar bone volume increase before any implant fixing. To overcome these difficulties, new ideas and approaches have emerged recently from the quickly developing fields of stem cell technology and tissue engineering.

### Stem cells in regenerative medicine

A stem cell is defined as a cell that can continuously produce unaltered daughters and, furthermore, has the ability to generate cells with different and more restricted properties. Stem cells can divide either symmetrically (allowing the increase of stem cell number) or asymmetrically. Asymmetric divisions keep the number of stem cells unaltered and are responsible for the generation of cells with different properties. These cells can either multiply (progenitors or transit amplifying cells)

or be committed to terminal differentiation. Progenitors and transit amplifying cells have a limited lifespan and therefore can only reconstitute a tissue for a short period of time when transplanted. In contrast, stem cells are self-renewing and thus can generate any tissue for a lifetime. This is a key property for a successful therapy. The capacity to expand stem cells in culture is an indispensable step for regenerative medicine, and a considerable effort has been made to evaluate the consequences of the cultivation on stem cell behaviour.

Stem cells cannot be identified with certainty in any tissue: scientists rely on indirect properties such as the expression of a repertoire of surface proteins, slow cell cycle, clonogenicity, or an undifferentiated state. However, none of these criteria are specific. The evaluation of self-renewal is the ultimate way to show “stemness”, which relies on the isolation and transplantation of a putative stem cell (clonal analysis) followed by its serial transplantation and long-term reconstitution of a tissue.

During recent years, stem cells have been used extensively in many medical disciplines for the repair and/or regeneration of defective tissues and organs (e.g. bone, ligament, heart). New therapeutic approaches are largely inspired and based on our knowledge of embryonic development. The aim of regenerative medicine is to stepwise re-create *in vitro* all the mechanisms and processes that nature uses during initiation and morphogenesis of a given organ. In this context, stem cell research offers an amazing and seductive potential for body homeostasis, repair, regeneration and pathology. The possibility of manipulating stem cells *in situ* using specific signalling molecules or by expanding them *ex vivo* is an exciting outcome of basic research. Hence, regenerative medicine has become a fashionable field and the isolation and manipulation of embryonic and adult (or post-natal) stem cells for the creation of new functional organs that will replace the missing or defective organs constitutes an enormous challenge. Embryonic and adult stem cells have been under intense investigation that focuses on the *in vitro* development of new organs such as hair, skin and bone. Adult stem cells (ASC), which possess a restricted potential of differentiation, can easily be isolated from a patient and after *in vitro* amplification and/or differentiation could be re-injected to the same patient thus avoiding immune rejection, as is the case for allografts or xenografts. Since numerous problems remain, the ideal protocol for human pathologies is far away from being used. However, the knowledge in stem cell technology is increasing quickly in all medical disciplines and dictates the need for new strategic approaches in all fields, including reparative dentistry. Stem cell therapy constitutes a common challenge for dentists as well as biologists.

#### **Dental stem niches and other stem cell sources for the development of teeth *in vitro* or *ex vivo***

As tooth formation results from epithelial-mesenchymal interactions, two different populations of stem cells have to be considered: epithelial stem cells (EpSC), which will give rise to ameloblasts, and mesenchymal stem cells (MSC) that will form the odontoblasts, cementoblasts,

osteoblasts and fibroblasts of the periodontal ligament. Thus, tooth engineering using stem cells is based on their isolation, association and culture as recombinants *in vitro* or *ex vivo* conditions to assess firstly tooth morphogenesis and secondly cell differentiation into tooth specific cells that will form dentin, enamel, cementum and alveolar bone. Various approaches could be used according to the origin of stem cells. Many recent studies have focused on the localization of sites of adult tissues/organs where specific ASC populations reside. ASC are quiescent, slow-cycling, undifferentiated cells, which are surrounded by neighbouring cells and extracellular matrix. This microenvironment is specific for each stem cell compartment but is likely to be influenced by common factors such as vasculature or loading pressure. The specialized microenvironment, housing ASC and transient-amplifying cells (TAC), forms a “niche”. Understanding these microenvironments and their regulation is the key for the successful reproduction of such niches and for the *ex vivo* engineering of an organ with ensured functional homeostasis.

In teeth, two different stem cell niches have been suggested: the cervical loop of rodent incisor for EpSC (Harada *et al.*, 1999; Mitsiadis *et al.*, 2007) and a perivascular niche in adult dental pulp for MSC (Shi and Gronthos, 2003). In rodent incisors the proliferation of EpSC, which is located at the cervical loop area, is governed by signals from the surrounding mesenchyme. FGF signalling (mainly FGF-3 and FGF-10) is of particular importance since it is linked to the Notch pathway (Thesleff *et al.*, 2007). Molecules such as BMPs, Activin and Follistatin are also expressed inside the stem cell niche and are known to regulate its maintenance and functionality through a complex integrative network (Mitsiadis *et al.*, 2007; Thesleff *et al.*, 2007; Wang *et al.*, 2007).

In the dental pulp, MSCs are thought to reside in a perivascular niche (Shi and Gronthos, 2003), but little is known on the exact location and molecular regulation of this niche. The Eph receptor tyrosine kinase family of guidance molecules appears to be involved in the maintenance of the human dental pulp perivascular niche (Stokowski *et al.*, 2007). EphB and its ligand Ephrin-B were shown to inhibit MSC migration and attachment via the MAPK pathway through unidirectional and bidirectional signalling respectively (Stokowski *et al.*, 2007).

In addition to the dental pulp MSC (Gronthos *et al.*, 2000; Miura *et al.*, 2003), other MSC populations have been isolated from human dental tissues such as the periodontal ligament (Seo *et al.*, 2004, 2005) and the dental follicle (Morsczech *et al.*, 2005), but nothing is known about the existence of a niche in these tissues.

#### **Mesenchymal stem cells**

MSC can be isolated from different sources. First described in bone marrow (Friedenstein *et al.*, 1970), MSC have been extensively characterized *in vitro* by the expression of markers such as STRO-1, CD146 or CD44 (Pittenger *et al.*, 1999). STRO-1 is a cell surface antigen used to identify

osteogenic precursors in bone marrow, CD146 a pericyte marker, and CD44 a mesenchymal stem cell marker. MSC possess a high self-renewal capacity and the potential to differentiate into mesodermal lineages thus forming cartilage, bone, adipose tissue, skeletal muscle and the stroma of connective tissues (Prockop, 1997). The potential of dental MSC for tooth regeneration and repair has been extensively studied in the last years. Below, we discuss mesenchymal progenitors that have been assessed for tooth engineering purposes, such as progenitors derived from teeth and bone marrow.

**Stem cells from human exfoliated deciduous teeth (SHED).** The isolation of post-natal stem cells from an easily accessible source is indispensable for tissue engineering and clinical applications. Recent findings demonstrated the isolation of mesenchymal progenitors from the pulp of human deciduous incisors (Miura *et al.*, 2003). These cells were named SHED (Stem cells from Human Exfoliated Deciduous teeth) and exhibited a high plasticity since they could differentiate into neurons, adipocytes, osteoblasts and odontoblasts (Miura *et al.*, 2003). *In vivo* SHED cells can induce bone or dentin formation but, in contrast to dental pulp, DPSC failed to produce a dentin-pulp complex.

**Adult dental pulp stem cells (DPSC).** After a dental injury, dental pulp is involved in a process called reparative dentinogenesis, where cells elaborate and deposit a new dentin matrix for the repair of the injured site (Mitsiadis and Rahiotis, 2004). It has been shown that adult dental pulp contains precursors capable of forming odontoblasts under appropriate signals (About *et al.*, 2000; About and Mitsiadis, 2001; Alliot-Licht *et al.*, 2005; Gronthos *et al.*, 2000, 2002; Miura *et al.*, 2003; Teles *et al.*, 2005). Among these signals are the calcium hydroxide or calcium phosphate materials, which constitute pulp-capping materials used by dentists for common dental treatments. Dental pulp progenitors have not been clearly identified but some data suggest that pericytes, which are able to differentiate into osteoblasts, could also differentiate into odontoblasts (Alliot-Licht *et al.*, 2005; Lovschall *et al.*, 2007; Shi and Gronthos, 2003). Tooth repair is a lifetime process thus suggesting that MSC might exist in adult dental pulp. The *in vivo* therapeutic targeting of these adult stem cells remains to be explored.

**Stem cells from the apical part of the papilla (SCAP).** Stem cells from the apical part of the human dental papilla (SCAP) have been isolated and their potential to differentiate into odontoblasts was compared to that of the periodontal ligament stem cells (PDLSC) (Sonoyama *et al.*, 2006). SCAP exhibit a higher proliferative rate and appears more effective than PDLSC for tooth formation. Importantly, SCAP are easily accessible since they can be isolated from human third molars.

**Stem cells from the dental follicle (DFSC).** DFSC have been isolated from follicle of human third molars and express the stem cell markers Notch1, STRO-1 and nestin (Morsczeck *et al.*, 2005). These cells can be maintained in culture for at least 15 passages. STRO-1 positive DFSC can differentiate into cementoblasts *in vitro* (Kemoun *et al.*, 2007) and are able to form cementum *in*

*vivo* (Handa *et al.*, 2002). Immortalized dental follicle cells are able to re-create a new periodontal ligament (PDL) after *in vivo* implantation (Yokoi *et al.*, 2007).

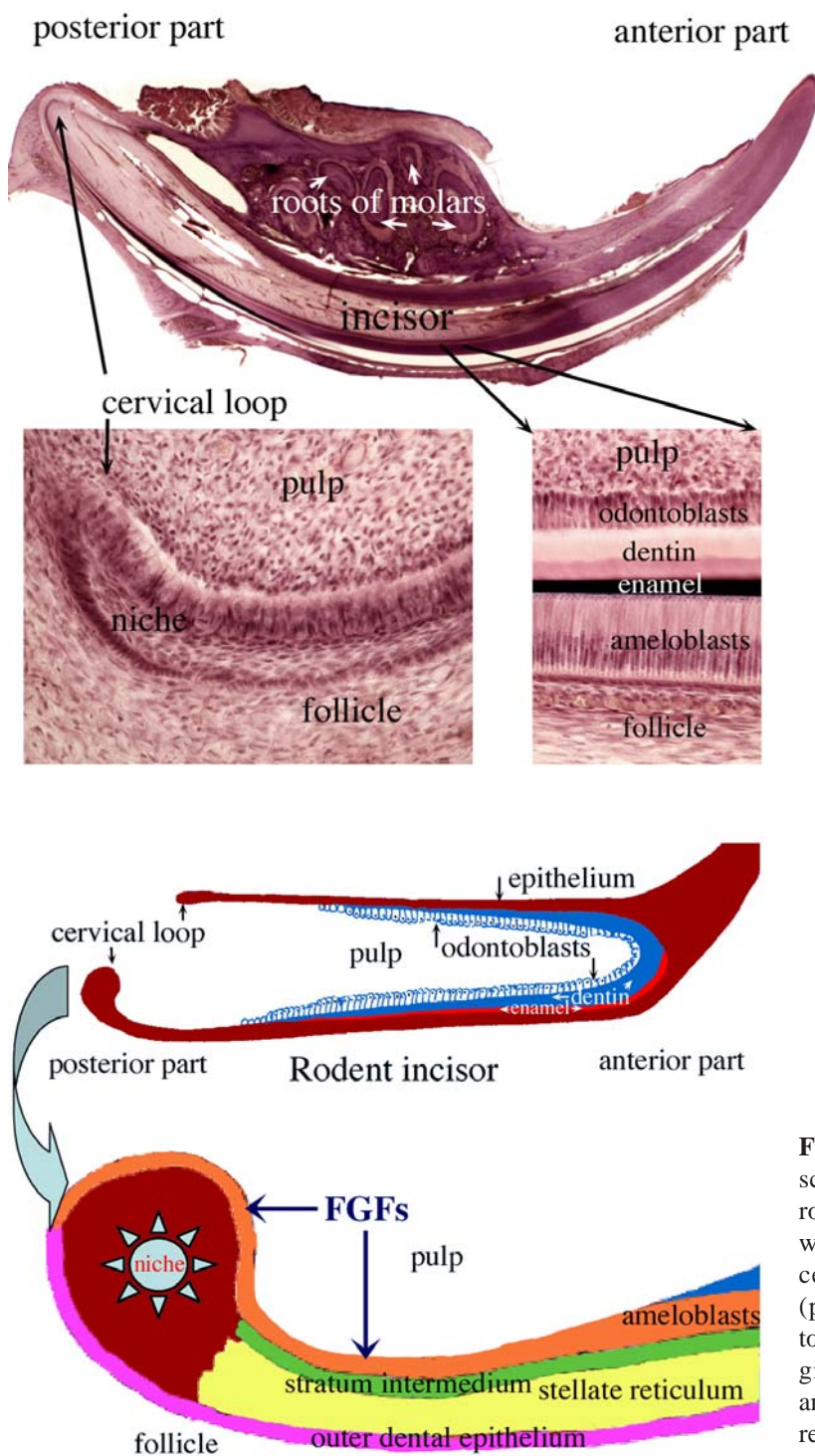
**Periodontal ligament stem cells (PDLSC).** The PDL is a specialized tissue located between the cementum and the alveolar bone and has as a role the maintenance and support of the teeth. Its continuous regeneration is thought to involve mesenchymal progenitors arising from the dental follicle. PDL contains STRO-1 positive cells that maintain certain plasticity since they can adopt adipogenic, osteogenic and chondrogenic phenotypes *in vitro* (Gay *et al.*, 2007). It is thus obvious that PDL itself contains progenitors, which can be activated to self-renew and regenerate other tissues such as cementum and alveolar bone (Seo *et al.*, 2004).

**Bone marrow derived mesenchymal stem cells (BMSC).** BMSC have been tested for their ability to re-create periodontal tissue. These cells are able to form *in vivo* cementum, PDL and alveolar bone after implantation into defective periodontal tissues. Thus, bone marrow provides an alternative source of MSC for the treatment of periodontal diseases (Kawaguchi *et al.*, 2004). BMSC share numerous characteristics with DPSC and are both able to form bone-like or tooth-like structures. However, BMSC display a lower odontogenic potential than DPSC (Yu *et al.*, 2007), indicating that MSC from different embryonic origins are not equivalent. Indeed, DPSC derive from neural crest cells, whereas BMSC originate from the mesoderm. Furthermore, the comparison of the osteogenic and adipogenic potential of MSC from different origins shows that, even if cells carry common genetic markers, they are not equivalent and are already committed toward a specific differentiation pathway (Musina *et al.*, 2005, 2006). Commitment could arise from conditioning of stem cells by their specific microenvironment or stem cell niche. MSC can also be obtained from several other sources such as synovial membrane (De Bari *et al.*, 2001, 2003, 2004) and periosteum (De Bari *et al.*, 2006). As these cell populations display distinctive biological properties depending upon their tissue of origin (De Bari and Dell'accio, 2008), it remains to be explored which source might be used for an optimal tooth development for clinical application.

#### **In search of epithelium-originated dental stem cells**

Although significant progress has been made with MSC, there is no information available for dental EpSC in humans. The major problem is that dental epithelial cells such as ameloblasts and ameloblast precursors are eliminated soon after tooth eruption. Therefore, epithelial cells that could be stimulated *in vivo* to form enamel are not present in human adult teeth. Stem cell technology appears to be the only possibility to re-create an enamel surface.

**Epithelial stem cells from developing molars.** Several studies describe the use of EpSC isolated from newborn or juvenile animals, usually from third molar teeth. In these studies, epithelia were removed and cells dissociated enzymatically. Precursors were then amplified and associated with MSC (originated from the same tooth) *in*



**Figure 2.** Histological sections and schematic representation of a mandibular rodent incisor. The sections were stained with haematoxylin/eosin. Epithelial stem cells located in the cervical loop area (posterior part of the incisor) migrate towards the anterior part of the incisor and give rise to four epithelial cell layers: ameloblasts, stratum intermedium, stellate reticulum and outer dental epithelium.

*in vitro* in contact with biomaterials such as collagen sponges or synthetic polymers (Honda *et al.*, 2005, 2007b; Young *et al.*, 2002). These approaches are promising for tooth formation and/or regeneration. However, the clinical application is difficult, if not unrealistic, since it would require the donation of a tooth germ from children. The use of autologous stem cells is desirable but raises the question of a good and reliable source.

**Epithelial stem cells from the labial cervical loop of rodent incisor.** The rodent incisor is a unique model for studying dental EpSC since, in contrast to human incisors or other vertebrates, this tooth grows throughout life. An

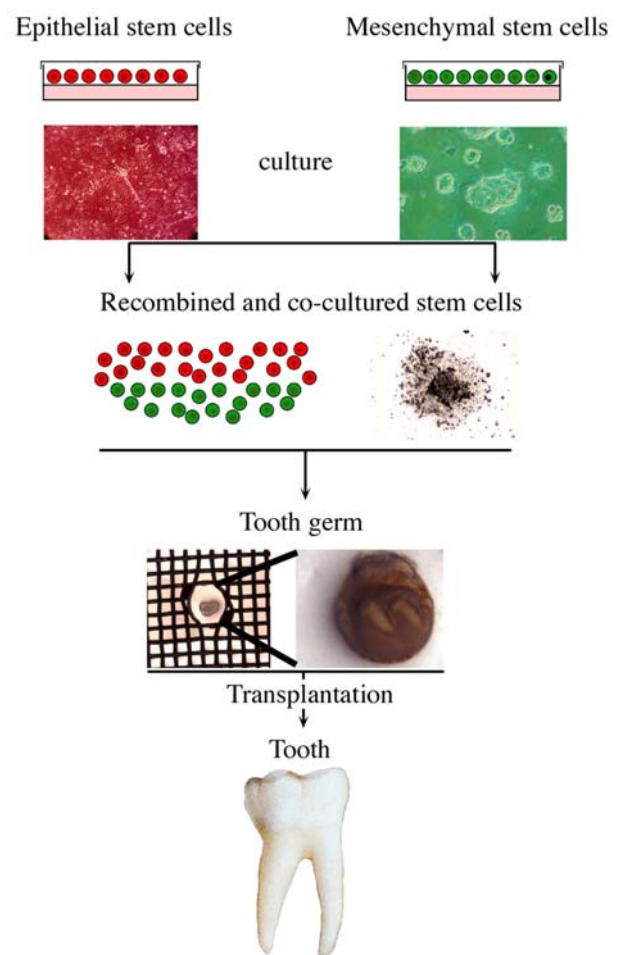
EpSC niche, which is located in the apical part of the rodent incisor epithelium (cervical loop area), is responsible for a continuous enamel matrix production (Harada *et al.*, 1999; Kawano *et al.*, 2004; Mitsiadis *et al.*, 1998, 2007; Smith and Warshawsky, 1975). In this highly proliferative area, undifferentiated epithelial cells migrate toward the anterior part of the incisor and give rise to ameloblasts (Fig. 2). Although these findings are important for understanding the mechanisms of stem cell homing, renewal and differentiation, this source of dental EpSC cannot be used for treatment in humans since it would require the introduction of rodent cells in the human mouth.

Dental EpSC can be isolated from post-natal teeth but exhibit complex problems that strongly limit their clinical application in humans. Other sources are thus required. Ideally these sources should be easily accessible, available from adult individuals and the derived cells must have potential for enamel matrix production. The use of non-dental EpSC will only be possible with the transfer of genes, creating an odontogenic potential to non-dental epithelia prior to any association with mesenchymal cells. This is certainly one of the most exciting goals of the next decade in tooth engineering.

#### Association of epithelial and mesenchymal stem cells

Since teeth are formed from two different tissues, building a tooth logically requires the association/cooperation of odontogenic mesenchymal and epithelial cells. The recombination of dissociated dental epithelial and mesenchymal tissues leads to tooth formation both *in vitro* and *in vivo* (Amar *et al.*, 1989; Yoshida *et al.*, 1998). Numerous attempts have been made in order to form teeth *in vivo* with very promising results. Single cell suspensions obtained from rat, pig or mice tooth germs have been seeded onto the surface of selected biomaterials (e.g. collagen-coated polyglycolic acid, calcium phosphate material, collagen sponges) and successfully re-implanted into the omentum of immunocompromised animals (Duailibi *et al.*, 2004; Honda *et al.*, 2006, 2007a, 2007b; Hu *et al.*, 2006; Robey, 2005; Young *et al.*, 2002). All these reports describe the presence of both dentin and enamel. This indicates that the recombined cells could re-organize themselves and form individual layers and, furthermore, that they can differentiate properly into odontoblasts and ameloblasts. In most of these studies the cells were directly seeded onto biomaterials without any additional *in vitro* procedure. In studies including *in vitro* steps before the *in vivo* transplantation, the results could be influenced by several critical parameters such as the presence or absence of serum, the type of serum, the composition of culture media, the cell density and the ratio between epithelial and mesenchymal cells. For these reasons, a definitive and universal protocol for tooth formation does not exist so far.

Making entire teeth with enamel and dentin structures *in vivo* is a reality and not a utopia. However, these bioengineered teeth have been produced in ectopic sites and are still missing some essential elements such as the complete root and periodontal tissues that allow correct anchoring into the alveolar bone. Recently, a new approach has been proposed for growing teeth in the mouse mandible (Nakao *et al.*, 2007). In this study, epithelial and mesenchymal cells were sequentially seeded into a collagen gel drop and then implanted into the tooth cavity of adult mice. With this technique the presence of all dental structures such as odontoblasts, ameloblasts, dental pulp, blood vessels, crown, periodontal ligament, root and alveolar bone could be observed (Nakao *et al.*, 2007). Thus, the implantation of these tooth germs in the mandible allowed their development, maturation and eruption (Fig. 3) indicating that stem cells could be used in the future for the replacement of missing teeth in humans.



**Figure 3.** Use of stem cells for tooth formation *in vitro* and *ex vivo*. A tooth germ can be created *in vitro* after co-culture of isolated epithelial and mesenchymal stem cells. This germ could be implanted into the alveolar bone and finally develop into a fully functional tooth.

Despite the outstanding advances in tooth bioengineering, such a technology can not be applied to human restorative dentistry for one simple reason: the epithelial and mesenchymal cells used for tooth reconstruction are of dental origin and have been given by a donor. The challenge that remains is to find out new and easily accessible sources of both epithelial and mesenchymal stem cells that can be reprogrammed for an odontogenic potential and then associated to form a fully functional tooth. One alternative could be the use of genetically modified cells expressing specific genes (e.g. transgenes, siRNA) or with a specifically deleted gene (e.g. knock-in, knock-out). Ideally, this approach should provide a non-limited source of cells and introduce new genetic information to reprogram a non-dental cell to acquire odontogenic properties. For example, p53-deficient mice were used to establish dental epithelial clonal cell lines subsequently associated with mesenchymal cells to bioengineer teeth *in vivo* (Komine *et al.*, 2007). These cell lines demonstrated heterogeneous outcomes in terms of regeneration depending on their differentiation state. Although this technique provides us with an unlimited source of epithelial cells and shows the potential of

genetically modified cells that can be used for tooth engineering (Komine *et al.*, 2007), many questions have to be resolved. Which gene should be used to trigger an odontogenic program? Is only one gene enough to reprogram a cell toward a tooth specific cell?

### Conclusions

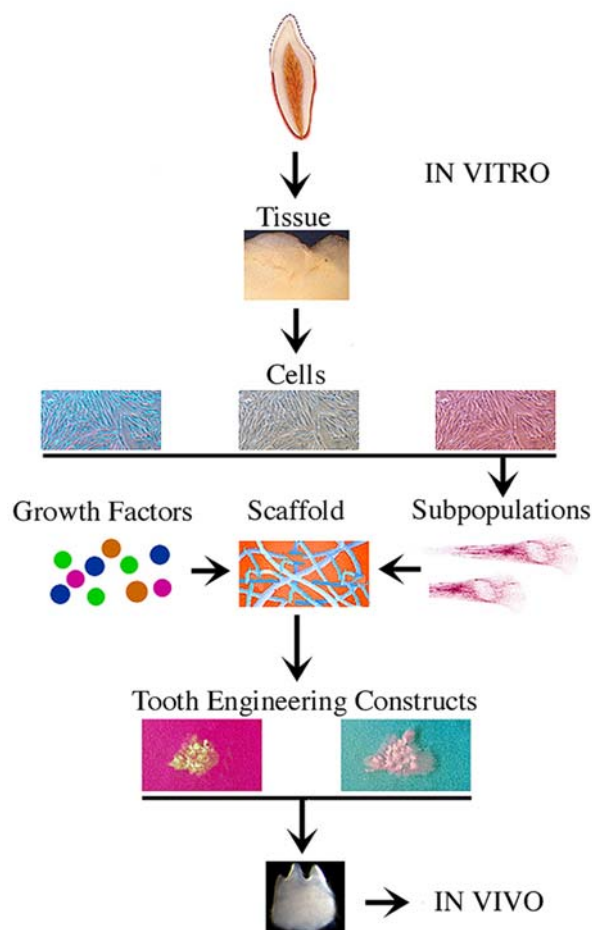
Taken together these recent findings clearly indicate that the control of morphogenesis and cytodifferentiation is a challenge that necessitates a thorough understanding of the cellular and molecular events involved in development, repair and regeneration of teeth. The identification of several types of epithelial and mesenchymal stem cells in the tooth and the knowledge of molecules involved in stem-cell fate is a significant achievement. *In vitro* and *in vivo* experiments using these cells have provided promising results illustrated by the generation of a complete tooth with all dental structures including cells and extracellular matrix deposition (Nakao *et al.*, 2007). However, many problems remain to be addressed before considering the clinical use of these technologies. The use of animal cells for human diseases is restricted by immune rejection risks. Additionally, isolating autologous stem cells requires a source of easily accessible cells without the need for a surgery. It may be possible to replace dental mesenchymal stem cells with stem cells of another origin. At present, it does not appear that this is the case for epithelial stem cells. A reliable source of EpSC for that purpose remains to be determined. Alternative solutions such as the use of artificial crowns are considered. The engineering of tri-dimensional matrices (either polylactic acid polymers or collagen sponges) which a composition more or less similar to that of the organs to reconstruct, and the addition of growth factors such as FGF, BMP or PDGF might facilitate the transplantation and the differentiation of stem cells (Fig. 4). However, the engineering of tooth substitutes is hard to scale up, costly, time-consuming and incompatible with the treatment of extensive tooth loss. Scientific knowledge is not enough and the main challenge in stem-cell therapy is to find a compromise between the benefits to the patients, regulatory agencies, increased stem cell requirements, costs, coverage by health insurance and the role of pharmaceutical companies.

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**Figure 4.** Construction of a bioengineered tooth. The association of tooth-derived stem cells with defined scaffolds in the presence of growth factors allows the creation of tooth specific constructs such as crown and root of missing parts of an injured tooth. These biological constructs could be used in dental clinics as substitutes for metal implants, crowns and restorative dental materials.

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