Fibroblast Heterogeneity in Periodontium – a Review

Dr. A. Archana¹, Dr. Venkata Srikanth², Dr. Sasireka², Dr. Bobby Kurien²*, Dr. Ebenezer²

¹Department of Periodontics, Madha Dental College, Chennai, India
²Department of Periodontics, Adhiparasakthi Dental College, Chennai, India
Corresponding author: bobbds2000@gmail.com

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Abstract Fibroblast heterogeneity applies to both phenotypic and functional properties exhibited by a fibroblast population within or across tissues (Mc Culloch et al) There is significant evidence that fibroblasts are heterogeneous with respect to functional properties, and that certain subpopulations of these cells may be clonally selected and expanded in diseased tissues. There are wide variations of gene expression and strikingly different responses to extracellular signals among different fibroblast populations. This has prompted a large number of in vitro studies which suggest that fibroblasts are not homogeneous but instead comprise multiple subpopulations with extensive site-to-site and intra-site variations.

Keywords: periodontal ligament, fibroblasts, collagen, Alkaline phosphatase

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1. Introduction

1.1. Background

Fibroblasts are the predominant cells of the periodontal ligament (PL) and have important roles in the development, function, and regeneration of the tooth support apparatus. Biological processes initiated during the formation of the PL contribute to the long-lasting homeostastic properties exhibited by PL fibroblast populations Over the past decade there has been increasing support for the hypothesis that PDL is directly involved in periodontal repair processes. The formation of the PL is likely controlled by epithelial-mesenchymal and epithelial hard tissue interactions, but the actual mechanisms that contribute to the development of cellular lineages in the PL are unknown.

1.2. Fibroblast Heterogeneity in Systemic tissues

Morphologic differences were identified in a fetal lung fibroblast line, WI – 38, has been shown to be heterogeneous in cell shape, nuclear morphology, and distribution of organelles

1.2.1. Collagen Production

Human skin fibroblast clones have been shown to be different in collagen production as much as 3-fold. Studies by Mollenham and Bay reuther observed 3 distinct fibroblast cell types with respect to collagen production. Fibroblasts termed FI produced very little Type I and type III collagen. FII cells synthesized slightly more collagen and large quantities were made by FIII fibroblasts.

1.2.2. Expression of Diverse Intermediate Filament

(Gabbioani et al). The concept of the myofibroblasts were distinguished from other fibroblasts by their expression of a smooth muscle actin and/ or desmin

Figure 1. Macroscopic tooth structure

Figure 2. Ultramicroscopic periodontal ligament fibres
whereas typical fibroblasts express only vimentin, a high percentage of fibroblast in granulation issue acquire α smooth muscle actin becoming contractile and thus promoting wound closure. IFN-γ inhibits fibroblast expression of α smooth muscle actin and reduce retractive conditions and hypertrophic scars. The observation that not all fibroblasts in the wound microenvironment acquire a myofibroblast phenotype supports the concept of fibroblast heterogeneity in a single anatomic site.

1.2.3. Fluorescence Activated Cell Sorting (FACS)
strategy employs antibodies or ligands to cell surface molecules
FACS has been used to separate subpopulation on the basis of:
• Thy 1,
• Class II MHC antigen,
• collagen receptors,
• Clq receptors and
• estrogens receptors.

2. Evidence of Fibroblast Heterogeneity in Periodontal Tissues

Two type of periodontal fibroblast defined by tissue localization.
The Gingival Fibroblast (GF) construct the soft connective tissue which surround the alveolar bone. These fibroblasts produce and maintain the extra cellular components which provide for the integrity of the tissue.
The PDL fibroblasts (PDLF) produces and maintains the connective tissue attachment which firmly anchors the tooth to the alveolus.

2.1. Major Themes

One is that fibroblasts from different areas of the periodontium are different, namely PDL and gingival.
The second theme is that within each of these specialized tissues there are likely to be multiple subsets of fibroblasts.

2.2. Heterogenetic Features

Rose et al reported that PDL possessed glycogen pools within cytoplasm with many bands of contractile type of microfilaments. They found no contractile microfilaments in GF.
Mariotti & Cochran reported slight differences in proliferative rates with GF growing more rapidly than PDL fibroblasts. Of greater significance were the distribution of GAGs in the cellular fraction of PDL tissue indicative of fibroblasts heterogeneity.
Variations in cell morphology
- Mass-cultured GF are relatively small and fusiform in shape.
- GF-C1: cells are large and fusiform
- GFCS (GF clone 5); cells are large and elongated
- GF-C6 (GF clone 6); polygonal cells are constantly found in this clone.
- Mass cultured PF; cells are small and slender in shape.
- PF-C1 (PF clone 1); cells are large and relatively elongated in shape.

2.3. Cytokines

IL-8 mRNA was found to be highly expressed in GF compares with PDLF with a differential expression of 85.1 fold. Studies have shown that constitutively high levels of IL-8 are observed in culture human gingival fibroblasts (Kent et 1996) possibly due to
1) GF is more engaged to respond to inflammatory stimuli than PDLF
2) neutrophil mediated proinflammatory processes maybe regulated in part by GF in the cytokine network of immunoparticipant cells. (Takashiba 1992)
3) GF may play a role a role in IL-8 production in the formation of cytokine network (Takigawa et al 1994)

4. Differences in Growth Characteristics of Fibroblasts in Periodontal Regeneration

There are specific differences in the growth characteristics of human fibroblasts derived from periodontal ligament and gingiva in culture. (Angelo J. Mariotti 1990) Similar growth conditions likely result after periodontal surgery or in cases of pathological loss of connective tissue. Fibroblasts derived from gingiva became confluent by day 4 whereas fibroblasts from periodontal ligament were not confluent until day 6. An explanation for gingival cells reaching confluence earlier than periodontal ligament fibroblasts, despite similar cell generation times (slope of logarithmic growth curve), may be that gingival cells are larger than periodontal ligament cells.

4.1. Growth Characteristics

The synthesis of total protein and collagen over seven days was similar between fibroblasts derived from periodontal ligament and gingiva. Analysis of collagen and noncollagen protein synthesis revealed a greater trend in noncollagen protein synthesis in the GF cultures compared to PDLF cultures, which helped to observe Significant functional differences between gingival connective tissue and periodontal ligament connective tissue.

4.2. Apoptotic Rate

In periodontal tissue sections, a significantly reduced apoptotic rate was demonstrated in PDLF compared with GF. In vitro, IGF-1 substantially enhanced cell survival in PDLF compared with GF by the up-regulation of anti-apoptotic molecules and the down-regulation of pro-apoptotic molecules.

4.3. Healing after GTR

Healing after guided tissue regeneration (GTR) may be explained by differences in functional activities of gingival and periodontal ligament fibroblasts (C. Giannopoulou and G. Cimasoni) GF and PDLF appeared identical under the SEM. Most ECM components increased the proliferation rate of GF and the biosynthetic activity of PDLF. Epithelial cells increased the proliferation of both GF and PDLF but had no effect on their biosynthetic activity.

4.4. Flow Cytometry Analysis of GTR Associated Periodontal Cells

The culture human regenerative cells were found to have a fibroblast like morphology similar to cells derived from normal PDL and gingival. Flow cytometry analysis using a panel of antibodies: Collagen type I, fibronectin and tenascin were expressed by all three cells but not equally or uniformly suggesting the presence of subsets. (L. Kuru et al) . Thus a fibronectin-high subset of cells in the membrane associated and regenerated tissue cultures indicated by elevated fibronectin expression.

5. Phenotypic Differences in Alkaline Phosphatase Expression

The heterogeneity of periodontal fibroblastic cells in vivo and in vitro indicates that periodontal cells could express either osteoblastic phenotypes (Kawase T, Nojima N) or gingival fibroblast characteristics (Piche JE 1989). HPLF showed a gradual increase in spontaneous alkaline phosphatase activity between proliferative and confluence states, (Matsuda et al ) The ALP expressed by periodontal ligament fibroblastic cells is clearly lower than ROS 17/2.8 (Alliot-Licht B 1991) but is not null, as on gingival fibroblastic cells (Kawase T).

5.1. Chemotactic Responses of PDLF and GF to Polypeptide Growth Factors

It has been hypothesized that competition from gingival fibroblasts may reduce the potential of periodontal regeneration PDL cells and GF exhibited dose-dependent migratory responses when challenged with PDGF, IGF-I, IGF-II, EGF, and TGF-beta. Only PDL cells migrated in a specific dose-dependent manner (F. Nishimura and V. P. Terranova).

5.2. TGF-β1 Expression

TGF-β1 is expressed in developing alveolar bone, PDL, and cementum at all stages of tooth development, notably in osteoblasts, PDL fibroblasts, and cementoblasts near the root apex TGFβ-R-I and -RII were weakly expressed in fibroblasts present in normal human gingival connective and PDL tissues and upregulated in regenerated tissue biopsies. Fibroblasts derived from PDL exhibited increased cellular proliferation, levels of alkaline phosphatase, collagen and protein synthesis as well as chemotaxis in response to TGF-β.

5.3. Integrin Expression

Expression of fibronectin receptor (α4β1) higher for GF and PDLF than DF, however all fibroblasts adhered equally well to fibronectin. Expression of α3β1 fibronectin receptor was equally great for PDLF, but lower for GF. PDL cell expression was lower than GF or DF for α4β8 both a fibronectin and laminin receptor.

6. Periodontal Wound Healing

Phenotypic differences between gingival and periodontal ligament fibroblasts have also been well documented in terms of the number of parameters of potential relevance to wound healing, morphology and growth potential (Hakkinen and Larjava, 1992) including the synthesis of matrix macromolecules (Hou and Yaeger, 1993) and response to TGF-β (Dennison et al, 1994; Mailhot et al, 1995).
6.1. ‘Fetal-like’ Fibroblasts

With specific reference to wound healing, Bucala et al (1994) have provided evidence that a specific subpopulation of circulating fibroblast-like cells enter the wound site by egress from the vasculature. Oral mucosa appears to be a privileged site in the adult in that it tends to display a ‘fetal like’ pattern of regenerative and scarless wound healing (Shafer et al, 1974; Walter and Grundy, 1992) The possible persistence of ‘fetal-like’ fibroblasts in the oral mucosa and their potential contribution to the characteristic mode of wound healing at this site have recently been discussed (Irwin et al, 1994).

6.2. Papillary Fibroblast

Irwin et al dissected the lamina propria of attached human gingiva into papillary tips and deeper reticular layers and demonstrated that fibroblast from each layer differ in several ways. Papillary fibroblasts appeared small and spindly proliferated faster in primary culture than those isolated from the reticular layers.

6.2.1. Response to Migration Stimulating Factor (MSF)

Papillary gingival fibroblasts were similar to fetal skin fibroblasts in their production of the migration stimulating factor (MSF) in contrast to the reticular fibroblasts which displayed a phenotype similar to adult skin fibroblasts that do not produce MSF. The fetal like gingival papillary fibroblasts may thus contribute to migration beneficial non-scarring wound healing in the oral mucosa.

6.3. Granulation Tissues Fibroblast

Granulation tissue is formed in connective tissue during wound healing, chronic inflammation, and certain pathological conditions. Phenotypical and functional heterogeneity in HGF and GTF fibroblasts is well documented with differences in morphology, size, and growth rate.

produce different amounts of extracellular matrix components such as proteoglycans, collagens, and matrix metalloproteinases (MMPs); respond differently to inflammatory cytokines and drugs; and express different amounts of integrin-type surface-receptors (fibronectin is regulated by a5b1 integrin)

It has been postulated that granulation tissue arising from gingival connective tissue does not contribute to the formation of hard tissue essential for periodontal regeneration (Larjava et al)

7. Fibroblast Heterogeneity in Pathogenesis Periodontal Disease

7.1. Clonal Selection Hypothesis

Fibroblast-derived proinflammatory mediators and cytokines such as PGE2, IL-1b, IL- 6, or IL-8 may be directly or indirectly implicated in periodontal tissue destruction by promoting fibrosis, granuloma formation or bone resorption. Irreversible changes have been observed in vitro in various phenotypic characteristics of fibroblasts which may be attributed to a positive selection process which results in the predominance of a certain subset (s) of fibroblasts with a unique proliferative and/or synthetic phenotype. (Korn JH 1992) also known as clonal selection hypothesis, was originally proposed by Ko et al This subpopulation (s) may be characterized by a proinflammatory cytokine secretion profile contributing to the pathogenetic mechanisms of the disease.

7.2. Phagocytosis

Under physiologic conditions, fibroblasts degrade the collagen matrix by which they are surrounded primarily by a phagocytic pathway. EM study in rats show that the PDL fibroblasts contain more phagocytosed collagen than gingival fibroblasts (Svoboda ELA 1981) suggesting site specific differences in phagocytic capacity. Cultured PDL fibroblasts appear to synthesize more collagen and fibronectin than gingival fibroblasts. Interestingly fibronectin which coats collagen fibrils in vivo (Pitaru et al 1987) has been proposed initiate phagocytosis by acting as a recognition site for fibroblasts (Mc culloch et al 1990). Since fibroblasts phagocytosed fibronectin beads more rapidly than collagen Type I beads, a higher level of fibronectin in the PDL might result in increased phagocytic activity.

7.3. Nitric Oxide Release

Van der Paw et al 2000 studied that PDL fibroblasts respond to very weak forces with the release of Nitric oxide whereas gingival fibroblasts do not. This second messenger molecule has been proposed to play a role in tissue remodelling and maintenance of PDL space. Though the precise action is not known, it might play a role in collagen phagocytosis.

7.4. Collagen Remodeling

7.4.1. Collagen Synthesis

GF or PF showed positive intracellular staining for both Collagen type I and Fibronectin, and Collagen III and Fibronectin. However, there were variations in fluorescence intensity for CI and Fn, ranging from relatively weak to strongly positive. The greater expression of collagen type I and fibronectin in mass cultures and most clones of PDL fibroblasts, in contrast to gingival fibroblasts under the same culture conditions, suggests that PDL fibroblasts are more active biosynthetically than gingival fibroblasts. This observation is consistent with recent observations that synthetic activity (total protein and collagenase digestible protein) of mass-cultured PDL fibroblast is one and a half to two times greater than that of the similarly passaged gingival fibroblasts (Hokama MM, 1987;) This increased synthetic activity of PDL fibroblasts conforms to the reported rapid turnover of collagen in the PDL tissues (Sodek J).

7.4.2. Collagenase Activity

The functional activity of collagenase synthesized by PDL fibroblasts has been found to be less than that of gingival fibroblasts (Sodek J 1975).

7.5. Effects of Mechanical Force

Primary fibroblasts derived from human periodontium are more resistant to mechanical load than is porcine
formation of IP3 and redistribution of cPKCs [Ca2+]i signals that was coupled to a differential growth, shape, attachment, and movement. Present studies specific fibroblast subsets. C1q influences fibroblast granulation tissues by differentially regulating activities of mRNA s encoding for cellular integrin subunits were different (Bolcato-Bellemin et al 2000).

7.5.1. Fibroblast Heterogeneity of Signal Transduction Mechanisms to Complement-C1q

C1q may participate in the compositional change of oral granulation tissues by differentially regulating activities of specific fibroblast subsets. C1q influences fibroblast growth, shape, attachment, and movement. Present studies show that the cC1qR and gC1qR fibroblast subsets responded to purified C1q with a differential generation of [Ca2+]i signals that was coupled to a differential formation of IP3 and redistribution of cPKCs.

8. Conclusion

The study of fibroblast heterogeneity will make great contributions to the understanding of pathologic processes and wound healing. When the role of various fibroblast subsets is understood manipulation of the response of these cells to injury and environmental challenge may be possible, leading to new and specific therapies for periodontal disease and other connective tissue diseases. Such therapies may include stimulating growth and differentiation receptors on fibroblasts or seeding damaged tissue with fibroblast progenitor cells. Finally, exploring how the fibroblast interacts with host defenses will undoubtedly reveal new information about the pathogenesis of periodontal disease.

References


