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Review Article

Understanding Mesangial Damage and Repair: Insights from an Experimental Model of Immunoglobulin Light Chain-Associated Mesangiopathy

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Abstract

Mesangial injury represents a crucial event in the pathogenesis of light chain-associated glomerulopathies in patients with plasma cell dyscrasias. The glomerulopathic light chains interact with mesangial cells where purported receptors regulate the downstream cellular mechanisms that will be activated and result in glomerular alterations. The physicochemical and conformational characteristics of the abnormal light chains are primarily responsible for the downstream events affecting the mesangial milieu.

Different light chains are responsible for two diseases with diametrically opposite mesangial alterations: Light chain deposition disease which results in the expansion of the mesangium due to accumulation of matrix proteins not present in the normal mesangium and AL (light chain-associated) amyloidosis where the native mesangial matrix is replaced by fibrils (amyloid). In both cases there is enhancement of mesangial cell apoptosis and the altered mesangium has a marked decrease of mesangial cells, most undergoing apoptosis.

The repair of the damaged mesangium is difficult due to the absence of enough mesangial cells that can participate in the process and also the damage can be so extensive that the intrinsic processes that are available for repair (i.e. recruitment of stem cells from bone marrow and precursor stem cells in renal niches)

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cannot effectively carry out the recovery. Introducing exogenous mesenchymal stem cells represents a novel therapeutic avenue that has been experimentally tested with promising results.

In-vitro, *ex-vivo* and *in-vivo* animal models have been created to study these disorders providing excellent platforms to elucidate pathogenesis and provide insightful information that can be translated to the practice of renal pathology, as the *in-vitro* and *in-vivo* platforms corroborate each other. The article addresses how mesangial injury occurs in LCDD and AL-amyloidosis as examples of generic mesangiopathies and summarizes existing data regarding the experimental use of exogenous mesenchymal stem cells in mesangial healing.

Keywords: AL - Amyloidosis; Electron microscopy; Light chains; Light chain deposition disease; Mesangial injury; Mesangial repair; Mesangial homeostasis; Nodular glomerulosclerosis

Glomerulopathic Light Chains Produce Mesangial Injury

Approximately 85% of the monoclonal proteins in patients with plasma cell dyscrasia are associated with renal disease. Involvement of the tubular interstitial compartment is more common (70% of all cases) than that of the glomerular (about 30%) or vascular compartments. In some patients there is involvement of more than one compartment, but the manifestations usually are centered in one of the renal compartments [1-5].

Investigations performed in the last 20 years have provided detailed insights into how these abnormal light chains interact with the mesangium to produce glomerular injury. A number of experimental platforms have been used in our laboratory to delineate the sequence of events that results in glomerular alterations which are centered in the mesangium. In all these, light chains purified from the urine of patients with renal biopsy-proven conditions have been used. The glomerulopathies associated with monoclonal light chains are light chain deposition disease (LCDD) and AL (light chain-related) amyloidosis, each creating completely different alterations in the mesangium. The findings that have been elucidated have paved the way for the design of novel therapeutic interventions aimed at stopping, slowing down or avoiding renal damage in patients with plasma cell dyscrasias.

How Light Chains Interact With the Glomerulus

Normal light chains are small molecular weight proteins that are filtered through the peripheral capillary walls and are delivered to the proximal tubules where they are catabolized by endosomal activity in the apical portions of the tubules and their amino acids are then returned to the circulation. Normal and tubulopathic light chains do not interact with the mesangium thus mesangial homeostasis is maintained intact.

Glomerulopathic light chains exert their initial or most significant pathogenic effect at the level of the mesangium. Although interactions occur early in the development of these disorders with the peripheral capillary walls, there is no doubt that the two main diseases involved in glomerular alterations focus on the mesangium as the center for

pathogenetic development and progression, eventually leading to glomerulosclerosis/glomerular obsolescence. These light chains are attracted to the mesangium because of certain physicochemically and conformational characteristics that make them interact with the mesangial cells [6].

Studies have shown that there are saturable receptors on the mesangial cells that are utilized by the glomerulopathic light chains to effect their pathogenetic role (Figure 1) [7-8]. This receptor is not yet fully characterized and it appears to be of a high molecular weight (about 450-470 kd). It is not the cubilin-megalín receptor on proximal tubular cells used by light chains for their normal metabolism. When rat or human mesangial cells are incubated with glomerulopathic (but not tubulopathic) light chains, the number of caveolae on their surfaces increases dramatically [7]. The interactions of the glomerulopathic light chains occur with caveolae which are located on the mesangial cell surface (Figure 2). Caveolae play a role in signaling [8-11].

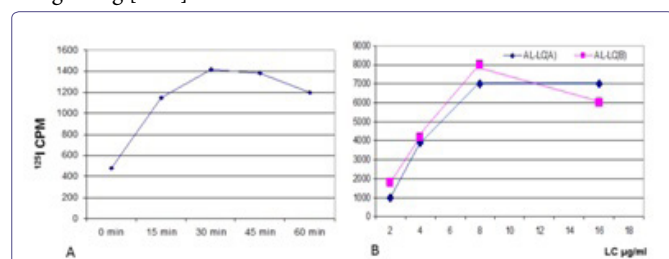


Figure 1: Time course and dose dependent rate of glomerulopathic light chain binding to human mesangial cells obtained from renal cortices of nephrectomy specimens for renal tumors (cortical tissue taken from areas away from the neoplasms).

Glomerulopathic light chains bind to mesangial cells and by 30 minutes the receptors are saturated, as shown in A. The binding for amyloidogenic and LCDD light chains follow a similar time course. Receptor saturation is dose dependent, as shown with two amyloidogenic light chains in B.

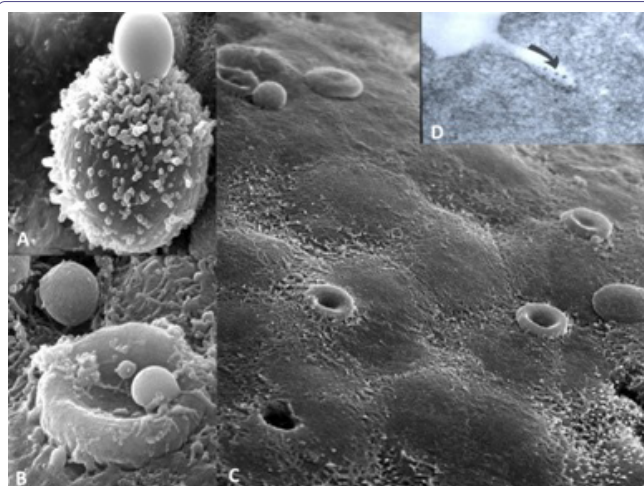


Figure 2: Light chain interacting with caveolae at the surface of human mesangial cells. Human mesangial cells incubated with LCDD - kappa light chain for 30 minutes Scanning electron microscopy A-C A - X50000, B - X45000, C - 15000 and D - Transmission electron microscopy, lead citrate and uranyl acetate Immunoelectron microscopy for kappa light chains, gold particles 20 nm X8500.

Note in A plasma cell releasing light chains. In B light chains are interacting with surface caveolae on mesangial cells, the latter well illustrated in C. On D, immunoelectron microscopy with kappa light chains labeled with gold particles interacting with surface membrane indentations (caveolae) on the surface of mesangial cell.

Both LCDD and amyloid forming light changes bind to receptors on mesangial cells and co-localize on the same sites along the membranes of mesangial cells, competing for the same receptor (Figure 3) [12]. However, while amyloidogenic light chains are avidly internalized into the mesangial cells (Figure 3B) using a clathrin-mediated mechanism (Figure 4), this is not the case with LCDD-light chains which activate cellular mechanisms when they interact with the surface receptors but are only minimally internalized [13,14]. Interaction with the surface receptors is responsible for their pathogenic effects downstream (Figures 3-5).

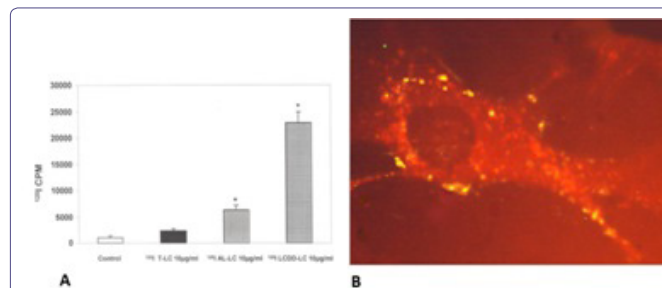


Figure 3: Binding of light chains to surface of human mesangial cells. Human mesangial cells incubated with amyloidogenic light chains (10µg/ml) for 30 minutes. A - Comparison of binding of different light chains B - X500- Co-localization on surface of mesangial cells of amyloidogenic and LCDD light chains. LCDD light chains marked with fluorescein and amyloidogenic light chains marked with Texas Red.

Note prominent binding of LCDD light chains to the surface of mesangial cells and the absence of binding in control or for tubulopathic light chains. Amyloidogenic light chains are not detected much at the surface of the mesangial cells as they are avidly internalized.

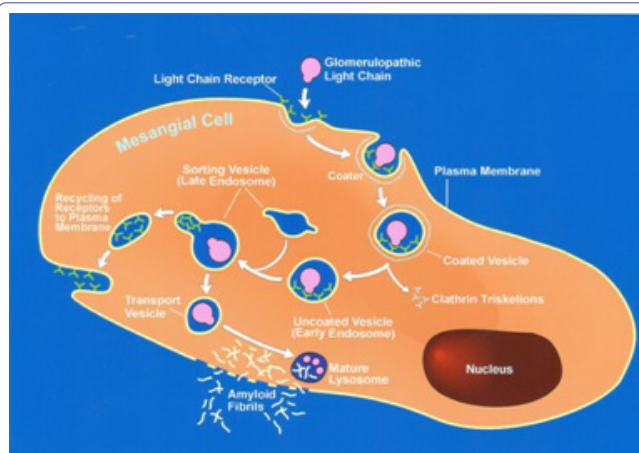


Figure 4: Schematic representation of clathrin-mediated endocytosis of amyloidogenic light chains.

The amyloidogenic light chains are endocytosed using a clathrin-mediated mechanism and they are delivered to the mature lysosomal compartment. The receptor is also internalized and returned to the surface.

Initial Effector Molecular Mechanisms that are Activated

Interactions between glomerulopathic light chains and mesangial cells in culture (parenthesis grown as monolayers on coverslips or on Matrigel) activate c-fos and NF-κβ which normally display a cytoplasmic localization in normal mesangial cells (Figure 5) and once stimulation occurs are translocated to the nuclei. C-fos has been found to participate in the activation of growth factors responsible for proliferative cellular activities (PDGF-β) and for cytoskeletal changes,

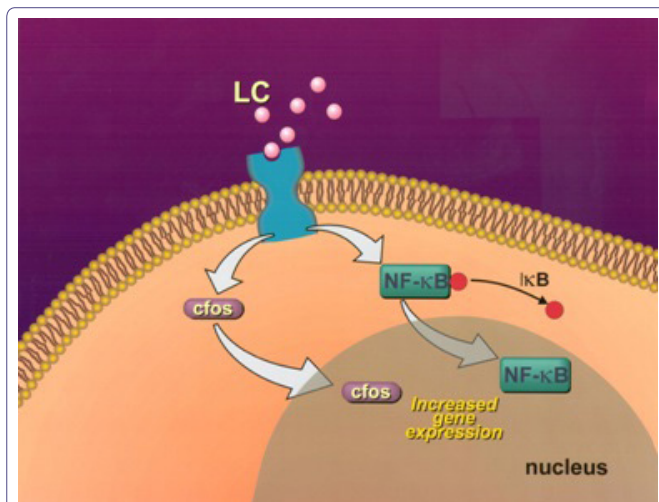


Figure 5: Schematic representation of the interactions of glomerulopathic light chains with the surface of mesangial cells.

Interactions with receptors at the surface of mesangial result in translocation of NF- κ B and c-fos to the nuclei where they activate increased expression of some existing genes as well as activation of genes that are normally silent in mesangial cells. This results in downstream effects which are different for LCDD and amyloidogenic light chains.

including phenotypic transformation, in mesangial cells. NF- κ B plays a crucial role in regulating Monocyte Chemoattractant Protein (MCP)-1 production by mesangial cells [7].

Growth Factors Participating in the Development of These Renal Diseases

Initially, in both situations, there is production of PDGF- β by mesangial cells and activation leading to mesangial cell proliferation once c-fos signaling has occurred (Figure 6,7). Afterwards, the two conditions engage different growth factors in their progression as a result of the signaling mechanisms involved being different. In the case of LCDD, transforming growth factor (TGF)- β is activated and becomes actively secreted by the mesangial cells. This growth factor is fundamental in the production of extracellular matrix and also exerts an inhibitory effect on mesangial cell proliferation (Figure 7) [15-18]. In contrast, amyloid formation occurs when mesangial cells are incubated with amyloidogenic light chains, and amyloid potentiates metalloproteinases which destroy the native mesangium and INHIBIT TGF- β , preventing matrix rebuilding. The end result is abundant matrix deposition in LCDD and amyloid fibrils in the affected mesangium in AL-amyloidosis. The presence of these two fundamental growth factors has been documented in glomeruli of patients with the two entities, highlighting the translational value and repercussion of these studies [19].

How Different Glomerulopathic Light Chain Affect Mesangial Cell Phenotype and Traffic in Mesangial Cells

When human or rat mesangial cells are incubated with glomerulopathic light chains extracted and purified from the urine of patient with renal biopsy-proven LCDD and AL-amyloidosis, there are dramatic changes that can be noted in each of the two conditions. These alterations in the mesangial "milieu" are crucial to understand the pathogenesis of these disorders and to translate the findings to clinical practice.

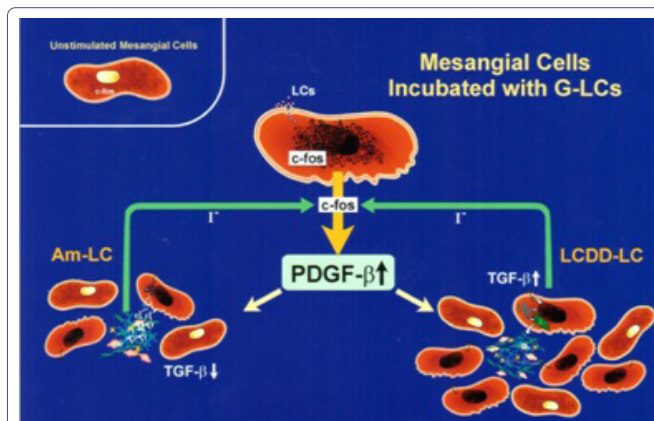


Figure 6: Interplay between PDGF- β and TGF- β in mesangial cells incubated with both glomerulopathic light chains.

In the case of Amyloidogenic Light Chains (Am-LC), there is initial production of PDGF- β and mesangial cell proliferation. Once amyloid is formed, it inhibits c-fos and ultimately cellular proliferation. In contrast, when mesangial cells are exposed to LCDD light chains after initial activation and secretion of PDGF- β there is production of TGF- β , leading to matrix formation and resulting in inhibition of PDGF- β .

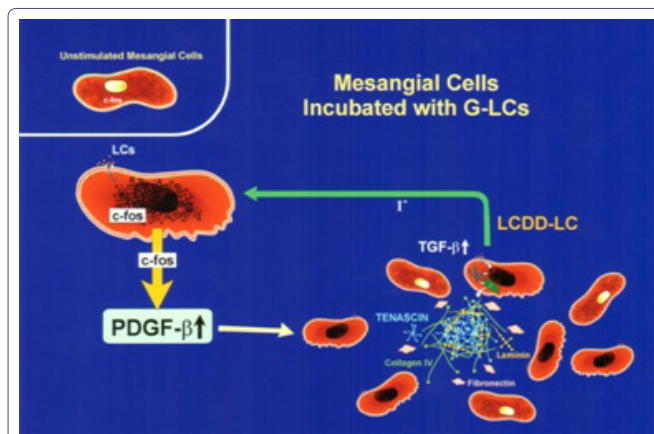


Figure 7: Interplay of growth factors in mesangial cells incubated with LCDD light chains.

The secretion and activation of TGF- β mediated by c-fos leads to increased tenascin-rich matrix production and the formation of mesangial nodules.

Normal mesangial cells exhibit characteristics of smooth muscle cells with abundant intracytoplasmic filaments maintained together by dense bodies or spindle densities (so-called myofilaments) and attachments plaques at their surface (Figure 8A,B). Normal mesangial cells express smoothelin and in some instances smooth muscle actin, the latter most noticeably when an activated phenotype develops. LCDD-light chains induce transformation of mesangial cells from their smooth muscle to a myofibroblastic phenotype (some consider this phenotype as an embryonic phenotype). This occurs by loss of some of the contractile elements and acquisition of cisternae of rough endoplasmic reticulum endowing the mesangial cells with the ability to increase their production of extracellular matrix proteins [16,20]. These activated mesangial cells typically express smooth muscle actin and maintain expression of smoothelin (Figure 8C).

In contrast, the mesangial cells transform to a macrophage phenotype when incubated with amyloidogenic light chains (Figure 9). This change is morphologically characterized by the loss

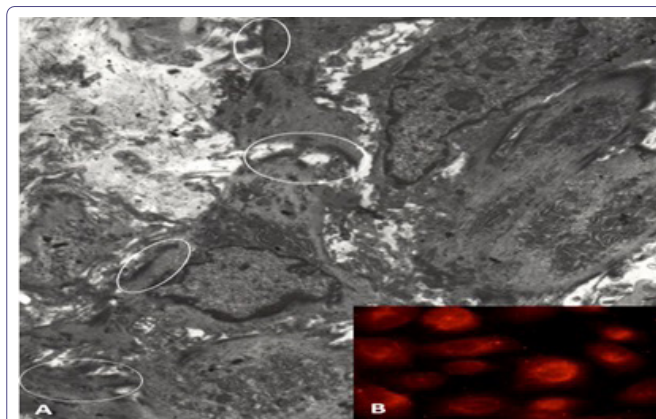


Figure 8: Normal human mesangial cells in culture- monolayer, obtained from uninvolved renal cortices from nephrectomies performed for renal tumors. A - X8000; B - X12000. Transmission electron microscopy, lead citrate and uranyl acetate. B - X750 Immunofluorescence labeling for smoothelin in normal mesangial cells.

Normal mesangial cells exhibit abundant intracytoplasmic Myofilaments maintained together by spindle densities or dense bodies. They also have Attachment Plaques (circles) at their surface. Normal mesangial cells express smoothelin reflecting the presence of intracytoplasmic myofilaments.

of contractile elements in their cytoplasm and the appearance of numerous lysosomes resulting in marked expansion of the lysosomal compartment, endowing these transformed mesangial cells with the machinery needed for active endocytosis and protein catabolism to take place. Rab proteins play a key role in transporting the light chains to the mature lysosomal compartment (Figure 10A) where they are processed and transformed into fibrils [12]. The transformed mesangial cells initially displayed Early Endosomal Antigen (EEA)-1 and later Lysosomal - Associated Membrane Protein (LAMP) and cathepsin D (characteristic of mature lysosomes) in their cytoplasm (Figure 10B). These mesangial cells acquire phagocytic/macrophage markers such as CD68 and Lysotracker (Figure 11), as a result of the acquisition of numerous lysosomes. During the process of mesangial cell transformation, hybrid cells with macrophage/smooth muscle characteristics can be identified attesting to the in-situ cellular transformation process and mitigating against the idea that these macrophage like cells are transported from the bone marrow, as

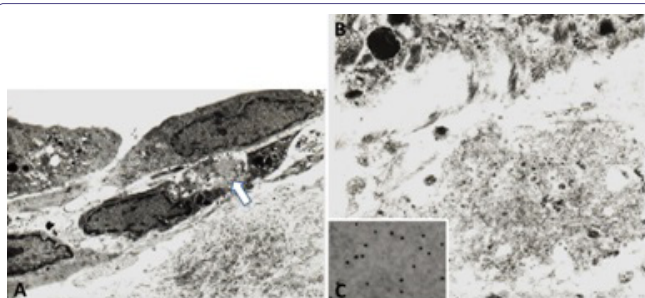


Figure 9: Human mesangial cells growing on Matrigel incubated with amyloidogenic light chain (10µg/ml) for 4 days. Transformed mesangial cells with macrophage phenotype engaged in formation of amyloid fibrils. A - X7500; B - X13500. Transmission electron microscopy, lead citrate and uranyl acetate. C - X8500- Immunogold labeling for lambda light chains, gold particles 20 nm.

Note the abundance of lysosomes in the cytoplasm of transformed mesangial cells participating in amyloid formation (arrow). The mesangial cell on top; however, remains non-transformed and exhibits typical characteristics of smooth muscle cells. In B, close up of area where amyloid is formed. Fibrils are labeled for lambda light chains (gold particles) demonstrating the derivation of the fibrils from lambda light chains.

was suggested could happen *in-vivo* (Figure 12). This plasticity of mesangial cells is crucial to drive pathologic events that occur in association with monoclonal light chain-associated mesangiopathies [20].

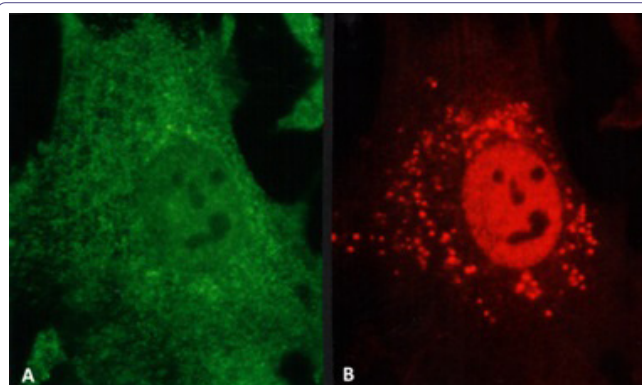


Figure 10: A - X500- Mesangial cells growing on monolayers incubated with amyloidogenic light chain (10µg/ml) for 1.5 hours. Immunofluorescence labeling of mesangial cells for Rab 7 with fluorescein; B - X500- Immunofluorescence labeling of mesangial cells for cathepsin D with Texas red.

Rab proteins participate in the trafficking of the amyloidogenic light chains to the mature lysosomes and they can be detected in the cytoplasm of mesangial cells at different stages of the process of amyloidogenic. Once the mesangial cells acquire lysosomes, they also acquire the expected enzymes which include cathepsin D.

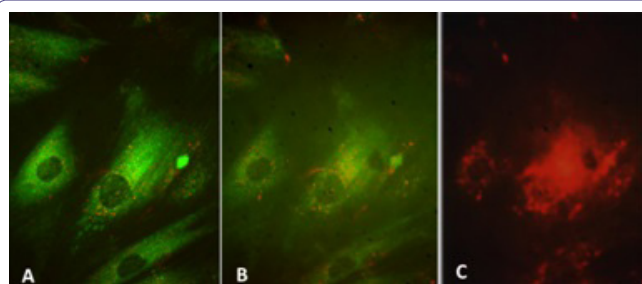


Figure 11: Amyloidogenic light chain (10µg/ml) incubated with human mesangial cells for 3 hours. Trafficking of amyloidogenic light chains to the mature lysosomal compartment in mesangial cells. Chase of amyloidogenic light chains labeled with Texas red (red) into the mature lysosomal compartment, previously marked with fluorescein (green) X350.

Internalized light chains appear as red dots that traffic through the mesangial cell eventually being delivered to the mature lysosomal compartment (co-localization represented by yellow color).

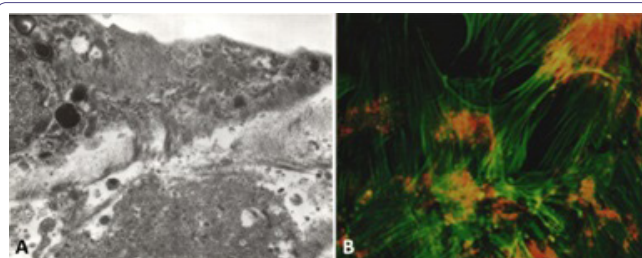


Figure 12: Human mesangial cells growing on Matrigel incubated with amyloidogenic light chain (10µg/ml) for 3 hours. Hybrid mesangial cells (with smooth muscle and macrophage phenotypes). A - X17500 Lead citrate and uranyl acetate and B - X500 Double labeling with muscle specific actin (fluorescein) and CD68 (Texas red).

In A, the mesangial cell illustrated at the top shows intracytoplasmic myofilaments (top right) and abundant lysosomes with loss of filaments (bottom left.) In B, co-expression of muscle specific actin and CD68 with co-localization clearly depict hybrid cells in the process of amyloidogenesis.

The two different types of glomerulopathic light chains though they use and compete for the same receptor on mesangial cells, exhibit entirely different intracellular trafficking patterns [12]. In the case of LCDD light chains, the minimal internalization that occurs leads to endosomal catabolism, with the receptor being returned to the cell membranes (Figure 13). The amyloidogenic light chains are taken in by the mesangial cells and the endosomal system is unable to catabolize them, directing the internalized light chains to the mature lysosomal system where catabolism leads to the formation of fibrils (Figures 11,13). The lysosomes must maintain an acidic environment to process the internalized light chains and convert them into fibrils. If the pH of the lysosomes is increased, their ability to form fibrils is impaired [21]. These fibrils are then extruded into the extracellular compartment when the lysosomes fuse with the cellular membranes and actively expel their contents [22].

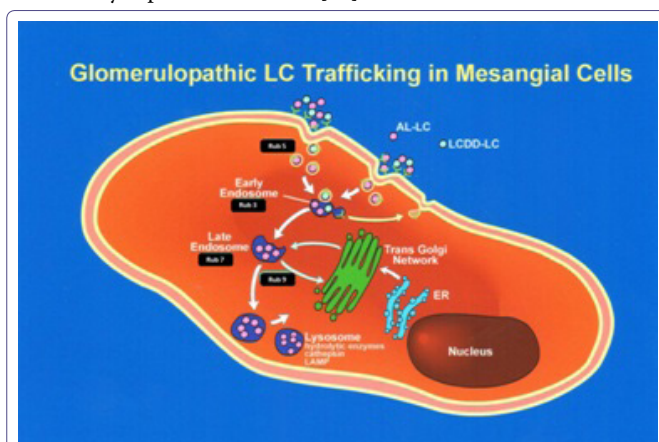


Figure 13: Schematic representation showing trafficking of LCDD and amyloidogenic light chains in mesangial cells.

While LCDD light chains are only minimally internalized into the mesangial cells and when such catabolized by endosomes, amyloidogenic light chains are delivered to the mature lysosomal compartment where their catabolism results in the formation of amyloid fibrils.

What are the Morphologic Parameters that Allow to Make a Diagnosis of these Diseases in the Clinical Arena?

It is important to understand how these diseases manifest their pathologic effects in the glomerular compartment to relate findings to those observed in experimental platforms. Renal biopsies from patients with LCDD and AL-amyloidosis reveal completely divergent mesangial abnormalities. Both diseases evolve from minimal manifestations to advanced disease and the sequence of events that must take place have now been elucidated in the laboratory where these diseases have been recreated for the purposes of studying the molecular mechanisms at play.

Patients with LCDD and renal involvement can present with a variety of glomerular manifestations depending on the stage of the disease process. This is easy to understand now that we have a clear appreciation of the events that must take place to arrive to the advanced morphologic expression known as nodular glomerulosclerosis (Figure 14), a process that mimics morphologically what is seen in advanced diabetic nephropathy, suggesting similar molecular mechanisms involved in their pathogenesis [18]. LCDD is preferentially associated with kappa light chains, mostly kappa I and IV, but there are cases also associated with lambda light chains [23-25]. These light chains manifest morphologically as punctate

electron dense material that is sometimes (if not electron dense enough) difficult to identify, as it tends to blend with the increased mesangial matrix (Figure 14). While in the initial phases of LCDD there are proliferative mesangial changes mediated by PDGF- β , as the disease progresses and matures, mesangial matrix deposition dominates the pathologic picture, driven by TGF- β , which in turn exerts a negative feedback on mesangial cellular proliferative activity resulting in the formation of mesangial nodules and progressive loss of mesangial cells as a direct effect of the glomerulopathic light chains [14,17]. Depending on which growth factor predominates in the mesangial milieu at the time of the renal biopsy, the morphologic findings will vary accordingly with either more or less mesangial cell proliferation and/or matrix deposition [19].

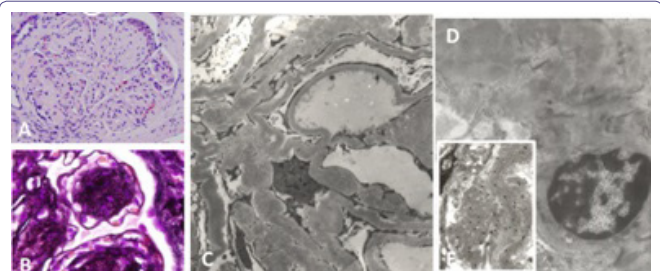


Figure 14: Renal manifestations of LCDD in renal biopsy material. A - X500 - Hematoxylin and eosin stain; B - X750; silver methenamine stain, C - X5800, D - X15500, E - X8000 transmission electron microscopy, lead citrate and uranyl acetate; E-Immunogold labeling for kappa light chains, gold particles 20 nm.

The typical appearance of LCDD in glomeruli is referred to as nodular glomerulosclerosis, because of the presence of mesangial nodules (appreciable in A and B) which contain abundant extracellular matrix (black staining in B). Ultrastructurally light chains appear as punctate, electron dense and tend to blend with the mesangial matrix (C and D). In E, gold particles mark the kappa light chain deposits.

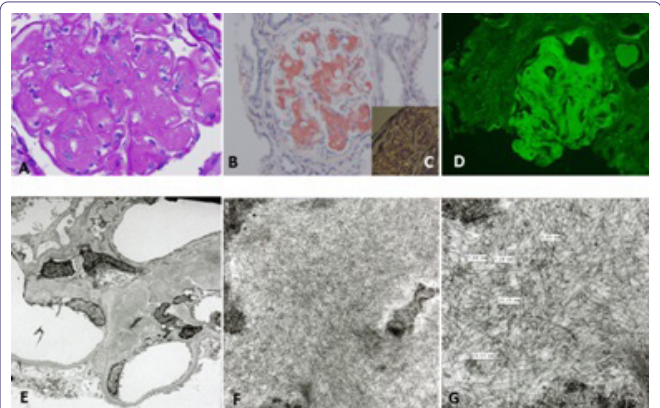


Figure 15: Renal manifestations of AL-amyloidosis in renal biopsy material. A - X500. Periodic Acid Schiff stain (PAS), B - X350 - Congo red stain; insert polarization C - X350 - Immunofluorescence of lambda light chains; D - X6500, E - X18500; F - X22500 - Transmission electron microscopy, lead citrate and uranyl acetate.

Mesangium is expanded with PAS+ amorphous material (A) which is Congo red positive (salmon pink color) and polarized with apple green birefringence (C). Also note the absence of mesangial cells. In D, the material is noted to stain for lambda light chains. Ultrastructurally, fibrillar material results in expansion of mesangial cells with replacement of normal mesangial matrix (E-G). The fibrils measure between 7-12 nm in diameter.

Patients with glomerular AL-amyloidosis progressively replace their normal mesangial matrix by non-branching fibrils with characteristic disposition (randomly disposed) and measuring

7-12 nm in diameter (Figure 15). This process; however, occurs slowly and in the early phases of this disease the replacement of normal mesangium may be very focal and segmental. The increased mesangial deposition of amyloid serves to further enhance apoptosis of mesangial cells which also occurs as a direct consequence of the amyloidogenic light chains [26-30]. Amyloidosis is characteristically associated with lambda light chains and glomerular amyloid is generally derived from $\lambda 6$ light chains [31,32]. However, kappa-associated amyloidosis also occurs, though much less commonly.

Extracellular Matrix in these Disorders

The experimental platforms clearly showed how the mesangial matrix is affected in these two conditions and the molecular mechanisms involved in the pathogenesis of these alterations. In LCDD, new matrix was produced by mesangial cells resulting in expansion of the mesangium. However, the matrix was noted to be abnormal. The normal collagen IV-rich mesangium was replaced by a tenascin-rich matrix. Genes normally quiescent in normal mesangial cells were activated leading to the production of abnormal extracellular matrix with fibrillary collagens, and most importantly, tenascin. Tenascin is a tenacious protein with an octopus-like appearance which binds strongly to integrins and matrix constituents [14]. It is a very difficult matrix protein to destroy and dispose of.

In the case of amyloidosis, it has been conclusively shown experimentally, that amyloid activate matrix metalloproteinases leading to destruction of the normal mesangial matrix which is then replaced by amyloid fibrils [30,33,34].

In both circumstances, mesangial cells become apoptotic (Figure 16) as a direct consequence of the monotypical glomerulopathic light chains, resulting in significant decrease and eventually almost total disappearance of mesangial cells [35]. This is clearly evident in experimental platforms and in renal samples from patients with advanced LCDD and AL-amyloidosis (Figures 14,15).

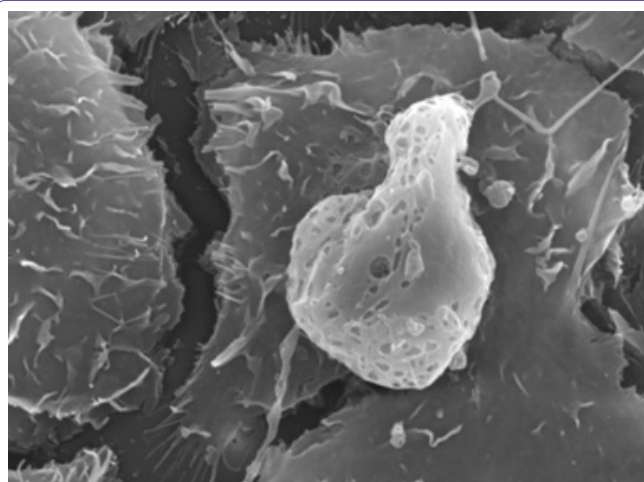


Figure 16: Apoptotic mesangial cell in mesangial cell culture incubated with amyloidogenic light chain (10 μ g/ml). Scanning electron microscopy X9800
Glomerulopathic light chains induce apoptosis of mesangial cells. An apoptotic mesangial cell is illustrated.

Reproduction of Pathologic Changes in the Mesangium in *In-Vitro*, *Ex-Vivo* and *In-Vivo* Experimental Platforms

The various platforms allowed for complete reproduction of the two disease processes [14,15,30,34-37]. Mesangial nodularity with increased extracellular matrix proteins, predominantly tenascin (Figure 17) and replacement of the normal mesangial matrix by amyloid fibrils was observed in the *in-vitro*, *ex-vivo* and *in-vivo* models using LCDD and amyloidogenic light chains, respectively. Each one of these platforms permitted a unique appreciation of the events taken place. Using the 6D live image system, the sequence of events taking place could be clearly depicted and the process of repair could be evaluated sequentially to completion. Each of the models was particularly useful for various purposes. For example, the cell culture model using coverslips and Matrigel demonstrated the basic principles at play and were particularly useful to delineate early events (Figure 18) [37]. The *ex-vivo* model provided a well-controlled system to create mesangial damage [34,36] and then repair with monitoring of key elements, including amounts of glomerulopathic light chains and then mesenchymal stem cells delivered to the kidney [38]. Finally, the *in-vivo* animal model where samples were studied with transmission and scanning microscopy was unique in depicting the process of amyloid secretion into the extracellular matrix [22]. In conjunction, the models provided a unified conglomerate of data reproducing the key cellular events in each case and substantiated the fact that what was observed *in-vitro* also occurred *in-vivo*. This fact further demonstrated the translational nature of these studies [39] and the applicability to design therapeutic strategies for patients with glomerulopathies associated with plasma cell dyscrasias.

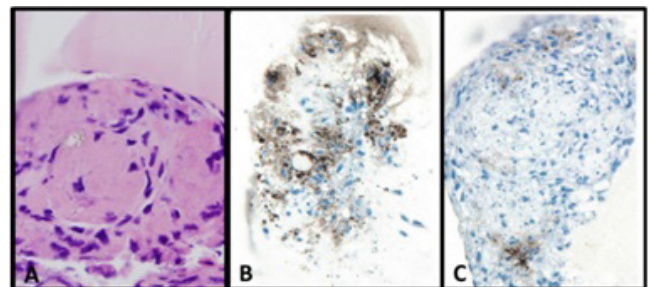


Figure 17: Human mesangial cells growing on Matrigel incubated with LCDD light chain (10 μ g/ml) for 4 days. Hematoxylin and eosin stain A - X500; Immunohistochemical stains for tenascin B - X350 and collagen IV C - X350.

Note formation of mesangial nodules similar to those seen in the renal biopsies from patients with LCDD and renal involvement (A). The mesangial nodules contain predominantly tenascin and the collagen IV initially present in the normal mesangium is almost totally gone and what little remains at the periphery of the nodules.

The Challenge of Repairing the Damaged Mesangium

If the glomeruli are not repaired, renal failure eventually occurs. As a result, the damaged mesangial areas in need of repair present several challenges: 1- Lack of mesangial cells to engage in the process; 2- Debris and abnormal material including fibrils and abnormal extracellular matrix proteins need to be disposed of, and 3-Lack of necessary growth factors, such as PDGF- β and TGF- β and other effector molecules to participate in and facilitate the healing process [38,40].

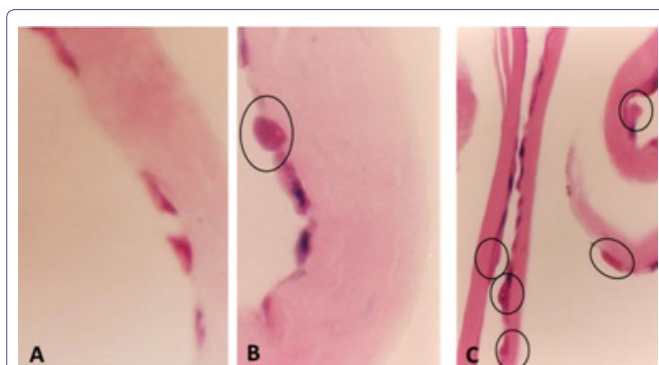


Figure 18: Mesangial cells growing on Matrigel incubated with tubulopathic (A) and amyloidogenic light chain (B and C) for 3 days (A), 1 day (B) and 2 days (C).

Note that the mesangial cells incubated with tubulopathic light chains are normally located on top of the matrix. The mesangial cells incubated with amyloidogenic light chains are forming amyloid (amorphous eosinophilic material-circled) and the amounts of amyloid increase significantly as more time passes (B,C circles).

Mechanisms of Mesangial Repair

Once glomerular damage occurs, there are a number of intrinsic mechanisms that are activated which will try to heal the damage. These mechanisms include migration of bone marrow mesenchymal stem cells to the kidneys and/or activation of progenitor stem cells that are present in renal niches [41-43].

These intrinsic mechanisms are limited and may not be entirely effective when the damage surpasses a certain degree of severity. The end result then, if no other therapies are employed, is for the glomeruli to become sclerotic or obsolescent with accompanying loss of their function. Ultimately the patient develops irreversible renal failure.

Exogenous Mesenchymal Stem Cells in the Repair of the Damaged Mesangium

Obviously it is important to begin treatment early, before irreversible glomerular changes take place. Injection of exogenous mesenchymal stem cells will provide not only cells that can be engaged in the process of repair but also important paracrine effects that will make the repair process much more efficient [38, 44-46].

One challenge is to make sure that the exogenous mesenchymal cells are delivered in adequate amounts to the sites of injury. Interestingly, they do so quite effectively at least when perfused through the renal artery but it may be more challenging to accomplish the desired delivery to certain sites, if they are diluted in the circulation. Pre-administration of curcumin enhances the migration of mesenchymal stem cells to where they are needed for repair [38].

Exogenous stem cells migrate to sites of injury (Figure 19) and rapidly engage in cleaning the damaged mesangium by acquiring a macrophage phenotype and phagocytosing the extraneous material. Once cleaning has taken place, the mesenchymal stem cells which are typically primitive and poorly differentiated with minimal amounts of cytoplasm with few organelles, differentiate into mature mesangial cells acquiring more cytoplasm in the form of cell process, and contractile elements such as myofilaments and attachment plaques [39,45,46]. This sequence of events was clearly depicted using the 6D liver cell imaging system (Figure 20) and confirmed in the *ex-vivo* platform. They eventually are able to perform contractile functions [46] and are morphologically akin to normal mesangial cells. Intrinsic

extracellular matrix properties regulate stem cell differentiation [47]. Then, transformed stem cells proceed to generate new mesangial matrix.

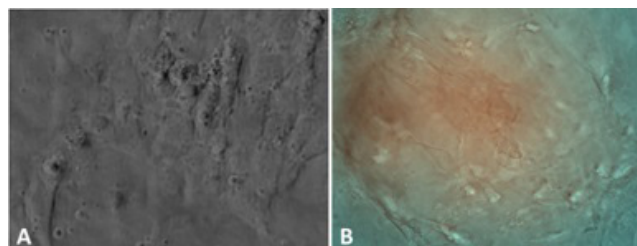


Figure 19: Mesangial cells incubated with LCDD light chains for 3 days and addition of mesenchymal stem cells afterward. A- phase contrast microscopy. B - Mesenchymal stem cells marked red with PKH-red

In A, the altered mesangial matrix with deposition of matrix proteins is shown. Mesenchymal stem cells marked red identify and migrate to the sites of damage to participate in the repair process (red staining in damaged mesangial area).

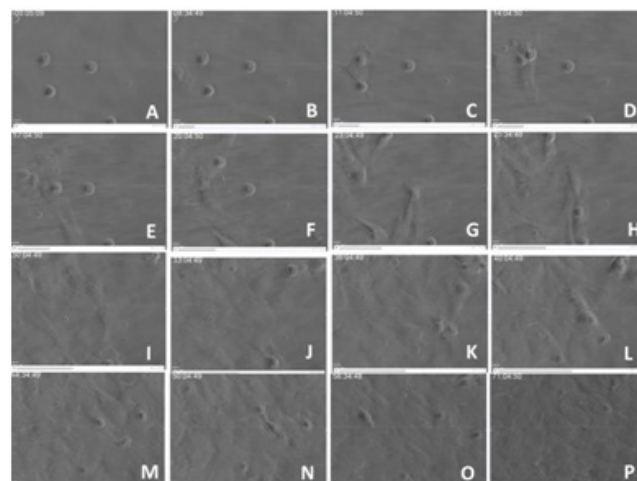


Figure 20: Sequence of events illustrating repair of damaged mesangium by amyloidogenic light chain using exogenous mesenchymal stem cells - 6D live cell platform

Mesenchymal stem cells clean the damaged mesangium by phagocytosing apoptotic mesangial cells, amyloid fibrils and debris and proceed to differentiate into mature mesangial cells lying down new matrix.

Conclusion

This manuscript summarizes how mesangial damage occurs in mesangiopathies and how it can be repaired, at least experimentally. The information that has been obtained should be of great value in the design of novel therapeutic avenues to address renal damage in plasma cell dyscrasias. The use of mesenchymal stem cells to repair the damaged mesangium appears to be a promising therapy to avoid irreversible renal damage in many glomerulopathies, not only those associated with plasma cell dyscrasias but also in other glomerular disorders with similar molecular pathogenesis, as the mechanisms involved are generic. Understanding mesangial damage represents a definitive advance in terms of future development of new therapeutic avenues to ameliorate or stop irreversible glomerulosclerosis.

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