Quantification and localization of M2 macrophages in human kidneys with acute tubular injury

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Abstract: This study addresses for the first time the question whether there is significant macrophage population in human kidney sections from patients with acute tubular injury (ATI). We examined therefore the interstitial macrophage population in human kidney tissue with biopsy-proven diagnosis of ATI, minimal change disease (MCD), and MCD with ATI. Kidney biopsies from patients with the above diagnoses were stained with antibodies directed against CD68 (general macrophage marker), CD163 (M2 marker), and HLA-DR (M1 marker) and their respective electron microscopy samples were evaluated for the presence of interstitial macrophages. Our study shows that patients with ATI have significantly increased numbers of interstitial CD68⁺ macrophages, with an increase in both HLA-DR⁺ M1 macrophages and CD163⁺ M2 macrophages as compared to patients with MCD alone. Approximately 75% of macrophages were M2 (CD163⁺) whereas only 25% were M1 (HLA-DR⁺). M2 macrophages, which are believed to be critical for wound healing, were found to localize close to the tubular basement membrane of injured proximal tubule cells. Ultra structural examination showed close adherence of macrophages to the basement membrane of injured tubular epithelial cells. We conclude that macrophages accumulate around injured tubules following ATI and exhibit predominantly an M2 phenotype. We further speculate that macrophage-mediated repair may involve physical contact between the M2 macrophage and the injured tubular epithelial cell.

Keywords: macrophages, acute kidney injury, CD163, HLA-DR, CD68, electron microscopy

Introduction

The healing response of mammalian tissues to acute injury is carried out by a complex interplay of diverse cell types including those of the innate immune system. Monocyte-derived tissue macrophages have recently been identified as important regulators of wound healing from early initiation of an inflammatory response to later facilitation of tissue repair and downregulation of inflammation.¹ These alternative functions of macrophages are attributed to plasticity of macrophage phenotypes, which have been described as a spectrum from classically activated macrophages (M1 phenotype) to alternatively activated macrophages (M2 phenotype).² Although the M1–M2 paradigm may be a simplification of the true in vivo picture, it represents a useful model for discerning distinct macrophage functions at different phases of wound healing and tissue repair.

Acute renal tubular injury is a clinicopathologic entity potentially caused by a host of insults to the kidney including ischemia and toxins.³ The resulting injury to renal
epithelial cells ranges from mild membrane and cytoplasmic alterations to frank necrosis. The ability of renal tubular epithelium to regenerate after such acute injury can be applied as a model to study the dynamics of wound healing and the roles of various cell types in tissue regeneration.\textsuperscript{4} Macrophages have been implicated in the process of initiating an inflammatory response as well as providing signals to damaged epithelial cells to proliferate and repair and prevent fibrosis.\textsuperscript{5,6} The observation that distinct macrophage subsets serve distinct functions in repair after tubular injury suggests that responses more heavily weighted by M2 phenotypes may result in more rapid and complete epithelial regeneration.\textsuperscript{7}

To date, the majority of this data has been obtained using animal models of kidney injury, with almost no quantitative studies of macrophage numbers in human patients with acute tubular injury (ATI). Thus, better understanding the role of macrophage responses to human ATI would support the potential use of macrophage phenotype analysis in biopsies to provide prognostic information, and may ultimately serve to support the idea of manipulating macrophage phenotypes to improve clinical outcomes after renal tubular injury.

To address the role of macrophage responses in the repair of injured human kidney, we have used immunohistochemistry analysis directed against CD68, HLA-DR, and CD-163 to quantify M1 and M2 macrophages respectively in biopsies from patients with ATI alone or minimal change disease (MCD) with or without ATI.

Material and methods

The protocol for case identification, scoring, and history review was approved by the Institutional Review Board at Yale New Haven Hospital. Pathology records were searched to obtain cases with biopsy diagnosis of ATI only or MCD, with or without ATI. Cases that showed any degree of overt interstitial nephritis or interstitial fibrosis were excluded from the study. Data were gathered on renal function and proteinuria levels at the time of biopsy and 1 month postbiopsy.

Immunohistochemistry

Five micron thick sections were cut on charged glass slides. They were then dried, deparaffinized in xylene, rehydrated in graded ethanol to distilled water. The slides were then placed in citrate buffer between 95\(^\circ\)C and 101\(^\circ\)C for 30 minutes (min). Once the slides had cooled they were placed in Tris-buffered saline (TBS). The slides were rinsed in TBS after each subsequent step. Endogenous peroxidase was quenched with hydrogen peroxide. Slides were then incubated in mouse anti-human CD68 (ThermO Cat# MS397) diluted to 1:600 for 30 min. The antibodies were then detected with Envision + (Mouse) (Dako K4007; Dako Denmark A/S, Glostrup, Denmark). This reaction is visualized with the included DAB in the kit. The slides are then placed in Tris-ethylenediaminetetraacetic acid (EDTA) at 90\(^\circ\)C to 101\(^\circ\)C for 30 min, then cooled and rinsed in TBS. The second antibody, either HLA-DR (AbCam Cat # ab20181) diluted to 1:500, or CD163 (Cell Marque cat # 163M-16) diluted 1:40, was applied for 30 min. This antibody is then detected with alkaline phosphatase labeled Envision G2 (Dako Cat# K5355). The antibody was detected with Envision secondary antibody-peroxidase reagent (Dako) then visualized with diaminobenzidine and counterstained with hematoxylin. The slides were then dehydrated, cleared, and mounted with resinous mounting medium.

Cases were scored at 200\(\times\) magnification (one high power field). The numbers of interstitial cells with positive cytoplasmic staining either for CD68, CD163 or HLA-DR, were recorded for ten adjacent high power fields (HPF) and then the average number of positive cells per HPF was calculated.

Electron microscopy

Kidney biopsy tissues were fixed in 2% formaldehyde, 2.5% glutaraldehyde in 0.1 M sodium cacodylate buffer, pH 7.4. They were then washed in 0.1 sodium phosphate pH 7.4 and exposed to 1% osmium tetroxide in the same buffer. The samples were then dehydrated through a series of graded ethanols to 100\%, and then passed through propylene oxide. Tissue was then infiltrated in 1:1 mixture of propylene oxide, and LX-112 resin, (Ladd Research Industries, Williston, VT, USA). The tissue was finally infiltrated with pure LX-112 resin, transferred to labeled capsules and polymerized overnight at 70\(^\circ\)F. Semi-thin, 0.2 \(\mu\)m sections were cut on a Reichert Ultracut E Ultramicrotome, and stained with toluidine blue for evaluation on the light microscope. Suitable areas were then thin sectioned on the same ultramicrotome and sections were collected on copper grids. Sections were then stained with 3% uranyl acetate and lead citrate and viewed utilizing a Carl Zeiss Libra 120 Electron Microscope (Carl Zeiss Meditec AG, Jena, Germany) at 80 KV. Images were taken at an initial magnification of 5,000\(\times\) with a charge-coupled device (CCD) camera and software (American Medical Technology, Boston, MA, USA).

Statistical analysis

Results are presented as mean \(\pm\) standard error of the mean of kidney biopsies from 8 patients. Differences were considered
CD163+ M2 macrophages were seen throughout the interstitium and often directly adjacent to injured tubular epithelial cells (Figure 2A and B). HLA-DR positive M1 macrophages were more sparse and often randomly distributed in the interstitium without specific adherence to tubular basement membranes (Figure 2B and C). Ultra structural examination of the peritubular interstitium and peritubular capillaries in patients with ATI showed macrophages migrating from the capillary lumen to the adjacent interstitium (Figure 3A–C) and adhering to the basement membrane of injured tubular epithelial cells (Figure 3D). Thus, the physical adherence of macrophages to the basement membrane of injured tubular epithelial cells may be an important step in the M2-mediated repair process of ATI.

**Discussion**

Our study shows that patients with either ATI alone or MCD with ATI have a significant increase in interstitial macrophage density compared to patients with MCD alone. Based on CD68, HLA-DR and CD163 expression, these are predominately comprised of M2 macrophages, with lesser but significant numbers of M1 macrophages. Although we do not have definitive information on the time interval between the renal insult and the time of biopsy, our chart review demonstrates 5.3±12 days between the clinical diagnosis of AKI (as evidenced by an increased serum creatinine level) and the time of biopsy in these patients. We therefore believe that the biopsy was performed well after the initial insult and most likely during the early repair phase of AKI. These time intervals are similar to animal models of renal ischemia/reperfusion injury, where the M2 macrophage is the predominant macrophage present in the kidney interstitium in the reparative phase of ATI. We therefore hypothesize that the M2 macrophage is the predominant macrophage type involved in tubular cell injury repair in humans.

M2 macrophages have been associated with resolving inflammation and inducing repair. They can express anti-inflammatory cytokine profiles for more than 4 weeks in vitro. The mechanisms underlying the protective effects of M2 macrophages include suppression and downregulation in the cytokine profiles of M1 macrophages, leading to a decrease in the production of pro-inflammatory cytokines. Additionally, M2 macrophages promote wound healing by releasing growth factors and extracellular matrix proteins.

**Table 1** Average values of creatinine (mg/dL) and proteinuria (spot pr:cr ratio) in patients at time of biopsy and 1 month later

<table>
<thead>
<tr>
<th>Diagnosis</th>
<th>Creatinine at bx</th>
<th>Proteinuria at bx</th>
<th>Creatinine one month post bx</th>
<th>Proteinuria one month post bx</th>
</tr>
</thead>
<tbody>
<tr>
<td>ATI</td>
<td>4.9±1.5</td>
<td>2.27±1.0</td>
<td>1.8±0.3</td>
<td>ND</td>
</tr>
<tr>
<td>MCD</td>
<td>0.7±0.1</td>
<td>3.5±2.7</td>
<td>0.7±0.1</td>
<td>0.5±0.3</td>
</tr>
<tr>
<td>MCD + ATI</td>
<td>2.2±0.5</td>
<td>1±6.0</td>
<td>1.5±0.3</td>
<td>1.4±1.2</td>
</tr>
</tbody>
</table>

**Notes:** Values shown are mean ± SEM; creatinine values are serum creatinine.

**Abbreviations:** ATI, acute tubular injury; bx, biopsy; cr, creatinine; MCD, minimal change disease; ND, not detected; pr, proteinuria; SEM, standard error of the mean.
of pro-inflammatory cytokines and suppression of M1 macrophages.\(^2\) This may explain the relatively low number of M1 macrophages in our patients’ biopsies. Tubular epithelial cells produce cytokines such as CCL2 that induce monocyte mobilization to the kidney following injury.\(^12\)–\(^14\) In experimental models of acute kidney disease, certain cytokines exert M2 macrophage-mediated reparative effects of tubular structure and function.\(^15\) Similar mechanisms may play a role in human kidneys following ATI and may lead to extravascular migration of monocytes into the interstitium and subsequent phenotype switch to a M2 predominant macrophage population, which induces epithelial cell repair through stimulation of cytokine excretion.

Previous studies in ischemia/reperfusion models of ATI have documented migration of macrophages to the area of tubular injury.\(^16\) CD163 positive M2 macrophages specifically localize close to the tubular basement membrane of injured proximal tubule cells as shown in our immunohistochemistry and ultrastructural studies. We conclude from our results that physical adherence of the M2 macrophage to the injured tubular epithelial cell may play an important role in macrophage-mediated repair mechanisms.

The role of macrophages in tubular injury repair is increasingly recognized.\(^17\)–\(^18\) In situ expansion of specific subtypes of intrinsic macrophages has been recognized as an important mechanism in tubular injury repair.\(^8\) Recent advances in molecular immunology and insight into cytokine expression that enhance cell injury repair, have
opened possible therapeutic avenues which may lead to novel drug development. Further studies in human kidney tissues are needed to assess the role of modified cytokines’ expression patterns that may enhance macrophage-mediated tubular injury repair.

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Disclosure
The authors have no conflicts of interest to disclose.

References