Renal co-morbidity in patients with rheumatic diseases

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Abstract
Renal co-morbidity is common in patients with rheumatic disease based on regular assessment of serum and urine parameters of renal function. When patients present with both arthritis and renal abnormalities the following questions have to be addressed. Is kidney disease a complication of rheumatic disease or its management, or are they both manifestations of a single systemic autoimmune disease? Is rheumatic disease a complication of kidney disease and its management? How do rheumatic disease and kidney disease affect each other even when they are unrelated? The present review provides an overview of how to address these questions in daily practice.

Introduction
Rheumatic disease and kidney disease are both common in the general population. Rheumatologists are thus frequently exposed to patients with concomitant renal disease. In fact, 18% of rheumatology clinic patients were reported to have a glomerular filtration rate (GFR) of 60 ml/minute or less as compared with the 5% reported within the general population [1]. When patients present with both, arthritis and kidney disease, the following questions have to be addressed. Is kidney disease a complication of rheumatic disease or its management, or are they both manifestations of a single systemic autoimmune disease? Is rheumatic disease a complication of kidney disease and its management? How do rheumatic disease and kidney disease affect each other even when they are unrelated? The present review addresses these questions and may help attending specialists, either rheumatologists or nephrologists, to manage patients with concomitant rheumatic disease and kidney disease.

How to diagnose and monitor renal co-morbidity?
Renal co-morbidity may not be apparent in terms of signs and symptoms, so functional parameters must be routinely measured. Serum creatinine is the most commonly used parameter for renal excretory function and is therefore used to predict the GFR; for example, by applying the Modification of Diet in Renal Disease formula. Nowadays, the estimated GFR defines the stage of chronic kidney disease (CKD) and most laboratories routinely report the estimated GFR along with serum creatinine results [2]. The Modification of Diet in Renal Disease formula as well as other formulas used for GFR prediction, however, have not been validated to reliably predict GFR in patients with rheumatic diseases. In fact, the Modification of Diet in Renal Disease formula may underestimate renal function in patients with rheumatoid arthritis (RA) [3,4] or with lupus nephritis [5]. Concomitant elevations of blood urea nitrogen and uric acid imply renal excretory failure but only at an advanced stage of kidney damage.

Excretion is only one of the many renal functions (Table 1), and the aforementioned parameters can fail to indicate serious and potentially fatal kidney disease. For example, serum parameters may not indicate damage to the glomerular filtration barrier, which only becomes evident on urinalysis. Proteinuria and hematuria, as detected by urine dipstick analysis, is suggestive of glomerulonephritis (GN) – in systemic lupus erythematosus (SLE) or anti-neutrophil cytoplasmic antibody (ANCA) vasculitis, for example – and massive proteinuria is a hallmark of membranous lupus nephritis, amyloid-A amyloidosis or certain drug toxicities (Table 2). Dipstick urinalysis is routinely performed during the initial evaluation and monitoring visits by most rheumatologists [6], although their lack of accuracy to detect albuminuria advocates other methods [7]. A positive dipstick result at least requires further testing.

A number of tests can help to discriminate whether proteinuria originates either from overflow of serum proteins, glomerular damage or predominant tubulo-interstitial damage. A simple screening test is quantitative urinary protein, albumin, and creatinine taken from a spot urine sample. Quantitative urinary protein will be
measured by the salicylate method, which also detects immunoglobulin light chains that are missed by the dipstick analysis. If albuminuria accounts for most of the proteinuria, a glomerular disease can be anticipated. **Vice versa**, if albuminuria accounts only for a minor proportion, the proteinuria will originate from either protein overflow or tubular disease. Because urinary protein and albumin concentrations are confounded by fluid intake and water clearance, these parameters are divided by the urinary creatinine concentration to calculate the urinary protein/creatinine ratio or the urinary albumin/creatinine ratio. The spot urine albumin/creatinine ratio was already shown in 1983 to be a suitable surrogate parameter for the 24-hour urine protein excretion [8]. As a consequence, most institutions no longer collect 24-hour urine levels. Several studies document the accuracy of this approach; for example, for monitoring lupus nephritis [9,10]. Whether or not a kidney biopsy is required should be discussed with a nephrologist.

| Renal manifestation | Drug |
|---------------------|------|
| Acute renal failure | NSAIDs, cyclosporine overdose, biphosphonates |
| Chronic renal failure | NSAIDs, cyclosporine overdose |
| Proteinuria/nephrotic syndrome | NSAIDs, gold |
| Sodium retention/edema/hypertension | NSAIDs |
| Arterial hypertension | Leflunomide, cyclosporine |

NSAID, nonsteroidal anti-inflammatory drug.

measured by the salicylate method, which also detects immunoglobulin light chains that are missed by the dipstick analysis. If albuminuria accounts for most of the proteinuria, a glomerular disease can be anticipated. **Vice versa**, if albuminuria accounts only for a minor proportion, the proteinuria will originate from either protein overflow or tubular disease. Because urinary protein and albumin concentrations are confounded by fluid intake and water clearance, these parameters are divided by the urinary creatinine concentration to calculate the urinary protein/creatinine ratio or the urinary albumin/creatinine ratio. The spot urine albumin/creatinine ratio was already shown in 1983 to be a suitable surrogate parameter for the 24-hour urine protein excretion [8]. As a consequence, most institutions no longer collect 24-hour urine levels. Several studies document the accuracy of this approach; for example, for monitoring lupus nephritis [9,10]. Whether or not a kidney biopsy is required should be discussed with a nephrologist.

**Renal and rheumatic manifestations of systemic autoimmune disease**

Rheumatic diseases are systemic autoimmune diseases that preferentially affect joints and soft tissues, but the inflammatory process regularly involves solid organs, including the kidney (Table 3). In addition, toxicity of disease-modifying antirheumatic drugs (DMARDs) frequently affects the kidney. The attending specialist is left with the dilemma of these differential diagnoses that require either intensifying or reducing drug therapy (Table 2).

**Rheumatoid arthritis**

Owing to their high prevalence, RA and renal disease often coincide. The renal toxicity of antirheumatic drugs (for example, NSAID or cyclosporine toxicity), secondary renal disease induced by the chronic inflammatory process (especially renal amyloidosis) and, potentially, renal manifestations of the primary disease process, however, are important differential diagnoses.

In a prospective study of renal disease in 235 patients with early RA, 7% developed persistent proteinuria and 6% presented elevated serum creatinine with or without
proteinuria during a 42-month observation period [11]. These abnormalities were mostly drug-related and reversible after discontinuation of the treatment. D-Penicillamine and gold (but not methotrexate) were most often associated with proteinuria. Drug-induced proteinuria was associated with age over 50 years and elevation of C-reactive protein levels or the erythrocyte sedimentation rate. Seventeen percent of patients displayed microhematuria at study entry, with intermittent or persistent hematuria occurring in 42% of patients during the study period. Hematuria was again more common in patients older than 50 years and was associated with RA activity rather than treatment [11].

In the more recent Methotrexate and Renal Insufficiency (MATRIX) study, elevated serum creatinine values were found in 19% of prevalent RA patients. Among these, 20% were in stage 2 and 15% were in stage 3 of CKD. Proteinuria, hematuria and leucocyturia were observed in 16%, 17% and 20% of the patients, respectively [12]. The causes of renal pathology were not determined in this study. In 1995, however, Helin and colleagues reported a retrospective analysis of renal biopsies from 110 RA patients in whom the clinical renal disease was probably due to antirheumatic therapy or RA itself [13]. The most common histopathologic finding was mesangial GN in 36%, followed by amyloidosis (30%) and membranous GN (17%). Focal proliferative GN (4%), minimal-change nephropathy (3%), and acute interstitial nephritis (1%) were less common. Amyloidosis was the most frequent finding in patients with nephrotic syndrome. In patients with isolated proteinuria, amyloidosis, membranous GN, and mesangial GN were almost equally common [13]. Similar results were reported in 1998 from Japan [14]. Novel DMARDs including biological drugs are more effective in suppressing systemic inflammation, however, and the incidence of renal amyloidosis has therefore declined [15]. Whether mesangial GN, which was frequently characterized by hematuria and glomerular deposition of IgM-containing immune complexes, is directly related to RA is still a matter for debate. In contrast, rheumatoid vasculitis, a rare but severe complication of chronic, erosive RA, may directly affect the kidneys. This vasculitis causes a necrotizing immune complex GN that requires intense immunosuppression [16,17].

Regardless of the etiology, concurrent renal disease is a predictor of mortality in RA patients. Compared with controls the hazard ratio for mortality was 1.78 in one population-based cohort study from Finland [18]. In this study, nephropathy presenting with combined hematuria and proteinuria, proteinuria alone, microalbuminuria, or histologically confirmed amyloidosis was associated with increased mortality in RA patients, whereas mortality was within expected limits in those with isolated hematuria or mesangial GN [18].

**Spondyloarthritis and psoriatic arthritis**

Thirty-five percent of patients with ankylosing spondylitis display abnormal serum creatinine measurements or urinalyses, often resulting from glomerular deposition of amyloid A or IgA [19]. Drug-induced nephrotoxicities – for example, interstitial nephritis or NSAID-associated membranous GN – are important differential diagnoses that deserve different therapeutic approaches. Amyloid-A nephropathy requires aggressive control of the underlying rheumatic disease, whereas NSAID-induced interstitial nephritis or membranous GN requires cessation of these drugs. In the latter case, glucocorticoids or TNF-blocking biologicals are alternative options for antiphlogistic therapy, and acetaminophen, metamizol/

### Table 3. Renal manifestations of systemic diseases commonly seen by rheumatologists

| Disease                              | Renal manifestation                                                                 |
|--------------------------------------|-------------------------------------------------------------------------------------|
| Rheumatoid arthritis                 | Mesangial GN, renal amyloidosis, membranous GN                                      |
| Spondylarthropathy/psoriatic arthritis| Renal amyloidosis, IgA nephropathy                                                 |
| Systemic lupus erythematosus         | Lupus nephritis: proliferative immune complex GN, membranous GN                     |
| Sjögren’s syndrome                   | Interstitial nephritis with renal tubular acidosis (type 1)                         |
| ANCA vasculitis                      | Crescentic (pauci-immune) GN, interstitial nephritis                                |
| Giant cell arteritis                 | Renal artery stenosis                                                               |
| Immune complex disease               | Immune complex GN                                                                  |
| Diffuse cutaneous systemic scleroderma| Scleroderma renal crisis, crescentic GN (with myeloperoxidase ANCA), interstitial nephritis, chronic (ischemic) kidney disease |
| Sarcoidosis                          | Nephrocalcinosis, nephrolithiasis, granulomatous interstitial nephritis              |
| Malignancy                           | Nephrotic syndrome (membranous GN, minimal change disease, focal glomerulosclerosis) |
| Diabetes                             | Chronic kidney disease: diabetic nephropathy                                        |
| Hypertension                         | Chronic kidney disease: hypertensive nephropathy                                    |

ANCA, anti-neutrophil cytoplasmic antibody; GN, glomerulonephritis.
dipyridamole or opioids should be used as alternative analgesic medications. Importantly, amyloid-A nephropathy and NSAID-induced membranous GN may both present through nephrotic syndrome, requiring a renal biopsy to make a diagnosis. Patients with psoriatic arthritis and other spondyloarthritides (for example, reactive arthritis) may also present with IgA nephropathy or amyloid-A nephropathy. In psoriatic arthritis patients, elevation of serum creatinine values and/or proteinuria correlates with age, disease activity and duration [20].

Systemic lupus erythematosus
Lupus nephritis determines long-term morbidity and mortality in SLE because renal failure is associated with a large number of secondary and tertiary complications, as discussed in detail below. Lupus nephritis affects up to 50% of patients with SLE, and screening SLE patients for renal manifestations of lupus is therefore mandatory at each visit. Screening intervals might be longer in patients with discoid lupus because they experience lupus nephritis in less than 5% of cases [21]. No reliable biomarker can predict the development of lupus nephritis in a non-nephritic patient to date, but the presence of U1snRNP antibodies, C1q antibodies, inherited complement deficiencies or high levels of anti-dsDNA antibodies are associated with a higher prevalence of nephritis in SLE patients.

Lupus nephritis can range from mild immune complex disease to diffuse proliferative GN as classified by the current International Society of Nephrology (ISN)/Renal Pathology Society (RPS) classification [22]. Immune complex deposition within the mesangium mainly activates mesangial cells and causes mesangiproliferative GN with hematuria and various degrees of proteinuria. The severity of renal disease correlates with the extent of immune complex deposits inside the glomerular mesangium and capillary network (ISN/RPS classes 1 to 4). When lupus immune complexes deposit outside the glomerular basement membrane, they mainly activate podocytes and cause large-scale proteinuria; that is, nephrotic syndrome (ISN/RPS class 5), which can occur together with any of the other classes. ISN/RPS class 6 represents glomerulosclerosis, a postinflammatory state of fibrotic scar formation just like lupus nephritis without major inflammatory disease activity. Additional parameters indicate activity and chronicity of the lesions which may provide important clues for individual treatment strategies [22].

There are three main reasons that argue for performing a renal biopsy in SLE patients with abnormal urinalysis. First, several studies document that ISN/RPS class 3 and class 4 lupus nephritis can be present in patients with only minor urine abnormalities, such as microhematuria or a protein/creatinine ratio of 0.2 to 0.5 [23]. Second, only ISN/RPS classes 3 to 5 with relevant disease activity are at risk for developing renal failure and might benefit from immunosuppressive therapy. As such, a renal biopsy may not be needed to confirm the diagnosis of lupus nephritis, but to enforce mandatory therapy despite side effects. On the contrary, unnecessary treatments are avoided when biopsy results indicate renal involvement with low inflammatory activity. Rheumatologists (and nephrologists) have to ask the pathologist to provide all necessary information included in the ISN/RPS classification of lupus nephritis, especially the lupus activity index and chronicity index. This information will be provided by specialized renal pathologists. Third, SLE patients may suffer from kidney diseases other than lupus nephritis [23]; for example, interstitial nephritis or thrombotic microangiopathy as a manifestation of antiphospholipid antibody syndrome.

Other immune complex diseases
Arthritis and nephritis are also common in other types of immune complex diseases such as postinfectious immune complex disease, Henoch–Schönlein purpura, cryoglobulinemia or endocarditis. While the latter example has its own diagnostic and treatment algorithms, all other types of immune complex disease may benefit from a renal biopsy, because the standard histological work-up of the obtained renal tissue often provides diagnostic clues to the underlying systemic disease. Examples include endocapillary GN in postinfectious immune complex disease, IgA nephropathy in Henoch–Schönlein purpura, and membranoproliferative GN in cryoglobulinemia.

Anti-neutrophil cytoplasmic antibody-associated vasculitis
ANCA-associated vasculitis involves the glomerular capillaries leading to GN in up to 80% of Wegener's patients. Abnormal serum creatinine/cystatin C levels and urinalysis indicate renal involvement. Urine microscopy displays dysmorphic erythrocytes, acanthocytes, and erythrocyte casts as a sign of destructive GN. On renal biopsy, ANCA-associated vasculitis typically presents as a pauci-immune (devoid of immune complexes), necrotizing and crescentic GN – which is why renal ANCA-associated vasculitis often presents as a rapid progressive disease with advanced stage of renal failure. Immunosuppressive therapy with cyclophosphamide and glucocorticoids must be initiated the same day to preserve renal function [24,25]. In addition, patients with end-stage renal disease should be treated with plasmapheresis to improve outcome [26].

Panarteritis nodosa and giant cell arteritis
Vasculitic involvement of intermediate-sized and large vessels in panarteritis nodosa and giant cell arteritis, respectively, may occasionally lead to renal functional impairment [27,28]. Inflammation and formation of
microaneurysms in arteries of the renal parenchyma in panarteritis nodosa or vasculitic stenosis of renal arteries and their branches in giant cell arteritis will lead to chronic renal hypoperfusion and ischemia. Without involvement of the small glomerular capillaries these present clinically as progressive renal impairment and albuminuria, usually without gross hematuria and leukocyturia on urinalysis. Imaging studies rather than renal biopsy are required to confirm the diagnosis.

Scleroderma

Scleroderma can be complicated by a number of different forms of kidney disease [29]. The classical manifestation is scleroderma renal crisis, which represents a form of malignant hypertension with acute renal failure and potentially other organ manifestations due to excessive hypertension. Hypertensive crisis develops because the microvascular abnormalities in scleroderma cause renal hypoperfusion, triggering excessive release of renin that activates angiotensin. Immediate angiotensin blockade is therefore necessary. Renal crisis is an intriguing complication of diffuse cutaneous systemic sclerosis but affects only 2% of patients. Many more patients have ischemic nephropathy leading to slowly progressing CKD that is accompanied by hypertension and albuminuria. Blood pressure measurement and dipstick analysis are therefore mandatory at all routine check-ups of scleroderma patients. Normotensive acute renal failure should be evaluated for interstitial nephritis or ANCA vasculitis, a separate entity in scleroderma with poor outcome [30]. A renal biopsy is usually required to obtain the diagnosis in scleroderma patients with renal failure.

Sarcoidosis

Ectopic production of calcitriol, mainly by activated macrophages, causes hyperabsorption of dietary calcium in up to 50% of patients with sarcoidosis [31]. This excess calcium load leads to hypercalcuria, and in 2 to 20% of cases leads to hypercalcinemia. The increased urinary excretion of calcium may cause nephrolithiasis, nephrocalcinosis, and polyuria. Nephrolithiasis has been reported in up to 14% of patients with sarcoidosis and may be the initial presenting feature of as well as the cause for renal failure in sarcoidosis patients [31,32]. Granulomatous interstitial nephritis is another typical renal manifestation of active diffuse sarcoidosis [33]. In this renal disease, urinalysis can be either normal or show sterile leukocyturia or mild tubular proteinuria that responds to glucocorticosteroids [33]. Sarcoidosis-related glomerular diseases are rare and their linking mechanisms are unknown [34]. A nephritic sediment with erythrocyte casts or heavy proteinuria should differentiate these from interstitial nephritis, although the diagnosis will be obtained by renal biopsy only.

Renal and rheumatic manifestations of other systemic diseases

Paraneoplastic syndromes

A variety of renal and musculoskeletal disorders can be associated with malignancy. Both types of disorder may occur concomitantly in individual patients, and differential diagnosis may be difficult, especially with regard to systemic autoimmune disease. Membranous GN, minimal change nephropathy and focal glomerulosclerosis are the most common renal manifestations of malignant tumors [35]. The predominant clinical finding in these conditions will be proteinuria with or without elevated serum creatinine. More rarely, membranoproliferative GN, rapid progressive GN, thrombotic microangiopathy, and interstitial renal disease have been described in isolated patients with solid tumors or hematologic malignancies. These paraneoplastic conditions may additionally be accompanied by tumor-associated arthralgias, oligoarthritis or polyarthralgia, polymyalgia and dermatomyositis. More rarely, lupus-like syndromes, vasculitis and eosinophilic fasciitis may develop due to an underlying malignancy [36].

Establishing the correct diagnosis of a paraneoplastic syndrome rather than a systemic autoimmune disease with renal and rheumatic manifestation may be difficult, but is obviously of crucial importance for the patient’s treatment and prognosis. These paraneoplastic syndromes must be differentiated from disease that directly results from malignancy. For example, multiple myeloma may cause renal damage due to amyloidosis, glomerular light/heavy chain deposition disease or cast nephropathy, as well as musculoskeletal pain due to local tumor growth. In addition, tumor lysis syndrome can cause uric acid nephropathy as well as crystal arthropathy. Finally, end-stage renal disease and proteinuria cause an elevated erythrocyte sedimentation rate, which should not be suspected as active rheumatic disease or occult malignancy [37,38].

Diabetes mellitus

Diabetic nephropathy is the most common cause of end-stage renal disease in the western world. Glomerular sclerosis due to chronic hyperglycemia leads to proteinuria up to the nephrotic range, and is paralleled by a progressive rise in serum creatinine values. Diabetes may also cause musculoskeletal disorders, including diabetic cheiroarthropathy of the hands, and neuropathic Charcot arthropathy affecting the feet [39]. Other musculoskeletal complications seen in diabetic patients are Dupuytren's contracture and carpal tunnel syndrome, although in end-stage renal disease the latter may also be caused by dialysis-related amyloidosis due to local deposition of β2-microglobulin (see below). Adhesive capsulitis of the shoulder and diffuse idiopathic skeletal hyperostosis are also associated with diabetes [39].
Musculoskeletal manifestations of renal disease

Mineral and bone disorders are a central complication of CKD (referred to as CKD-MBD), because renal failure impairs renal phosphorus clearance and the activation of 25-OH vitamin D. The resulting hyperphosphatemia and hypocalcemia both induce secondary hyperparathyroidism, which is associated with two major consequences: bone mineral loss and extraskeletal calcifications. Parathyroid hormone-induced bone mineral loss (that is, renal osteodystrophy) is often associated with osteoporosis due to immobilization during hemodialysis and due to CKD-associated systemic inflammation. Both increase the risk of fracture-related morbidity [40].

The most common complication of CKD-MBD is accelerated atherosclerosis leading to cardiovascular and cerebrovascular events. Extraskeletal calcifications in CKD-MBD also include chondrocalcinosis and pseudogout due to the precipitation of calcium pyrophosphate dihydrate crystals. The international Kidney Disease Improving Global Outcomes initiative guidelines are available for evidence-based CKD-MBD management and inform about the use of vitamin D analogs, phosphate binders, and calcium supplements [40]. The impaired renal uric acid clearance leads to hyperuricemia. Concomitant use of diuretics and/or cyclosporin is common and can further impair uric acid clearance and increase the risk of gout attacks. Advanced CKD represents a state of systemic immunosuppression that can alter the clinical presentation of autoinflammatory diseases like gout. Acute gout attacks can be mitigated in CKD, and symptomatic patients should be examined for signs of chronic gout [41]. Treating crystal arthropathy is problematic in CKD as drugs commonly used to treat acute gout attacks, such as NSAIDs, should be avoided in CKD patients (Table 4). Colchicine should also be avoided because the narrow therapeutic range makes appropriate dose adjustments difficult [41]. We use 20 mg prednisolone for 3 to 5 days to control acute gout attacks and advise patients to start steroids immediately upon first symptoms of an attack. The central role of IL-1β in the pathophysiology of crystal arthritis suggests that IL-1 inhibitors are efficient in CKD patients [42,43]. Interval control of hyperuricemia and the treatment of chronic gout require dose adjustments of allopurinol or benzbromarone (when GFR >50 ml/minute/1.73 m²). Some reports propose febuxostat, uricase or anakinra as alternative treatments for gout, which may offer advantages for CKD patients [41,43,44].

Carpal tunnel syndrome, bone cysts, and destructive spondyloarthrophy or joint arthropathy used to be classical complications for end-stage kidney disease patients affected by dialysis-related amyloidosis, but their prevalence declined upon broad use of high-flux dialysers [45]. Impaired clearance of small proteins like β₂-microglobulin leads to β-sheet amyloid fibrils. Symptomatic amyloidosis rarely developed before 5 years of hemodialysis but the use of high-flux dialysis filters has recently delayed the onset of dialysis-related amyloidosis. β₂-Microglobulin amyloid deposition in bone, joints, and tendons causes pain, stiffness, and arthritis. Bone cysts can cause fractures and tendon deposits cause tears in ligaments and tendons [45].

Implications of unrelated chronic kidney disease in the rheumatic patient

Patients with impaired renal function have a substantial increase in cardiovascular risk that can be explained only in part by an increase in traditional risk factors such as hypertension and diabetes. It is now well recognized that CKD alone is an independent risk factor for cardiovascular complications [46,47]. International guidelines thus stress the need for preventing and treating cardiovascular risk factors in patients with CKD, as has been proposed for RA or SLE patients [48-51]. It is very likely that patients with rheumatic diseases and concomitant CKD are threatened by additive proatherogenic effects, and such individuals should hence be regarded as high-risk patients for cardiovascular complications that deserve rigorous control of their rheumatic disease activity. Several DMARDs cannot be used in CKD patients, however, due to their inherent nephrotoxicity. For example, NSAIDs (including cyclooxygenase-2 inhibitors) can cause acute deterioration of renal function, which is more common particularly in older patients, in CKD patients or in heart failure patients, and in states of volume depletion, because renal blood flow is dependent on renal prostaglandin synthesis [52,53]. Glucocorticoids, acetaminophen, metamizol/dipyrone or opioids can replace NSAIDs in CKD patients. Cyclosporine A is another potentially nephrotoxic DMARD. Cyclosporine causes vasoconstriction of the afferent and efferent glomerular arterioles, which leads to reductions in renal blood flow and the GFR. Chronic administration of cyclosporine can cause obliterative arteriolopathy, ischemic scarring, tubular atrophy and progressive loss of renal function. Risk factors include high dose, increased age and pre-existing CKD [54]. Cyclosporine is hence contraindicated in RA patients with renal dysfunction according to international guidelines [55,56].

Renal insufficiency also impairs the excretion of some DMARDs, which increases (nomenal) toxicity. For example, methotrexate is not nephrotoxic per se but it is excreted via the kidneys, and therefore accumulates and is increasingly toxic along the stages of CKD [57,58]. Some authors advocate dose reductions in CKD patients, but this can be dangerous because declines in the GFR can always happen with incident fluid losses – for example, through vomiting, diarrhea, use of diuretics or...
NSAIDs, sweating or fever – which further impair methotrexate clearance without a physician even being consulted. In view of the potential lethal side effects and usually available alternative treatment regimens (for example, leflunomide or biologicals), methotrexate should generally be avoided in CKD patients. Antimalarials, sulfasalazine, and azathioprine are only partially excreted by the kidneys, and therefore dose reductions were proposed at GFR <50 ml/minute (for details see [59,60]).

Some DMARDs have specific side effects that add to the cardiovascular risk of patients with rheumatic disease. For example, cyclosporine or leflunomide often increases blood pressure, which adds blood pressure monitoring and blood pressure control (<130/80 mmHg) to the responsibilities of the rheumatologist [49]. A typical side effect of tacrolimus and tocilizumab is hyperlipoproteinemia, which must be controlled by appropriate lipid-lowering therapy to levels below 70 to 100 mg/dl in rheumatic patients with CKD, most commonly with statins [48].

Anemia in patients with active rheumatic disease and renal insufficiency can be inflammation related and/or CKD related. Differential diagnosis may be difficult in individual patients, but persistent anemia despite control of inflammatory disease activity suggests a renal cause. These patients should receive iron supplementation and erythropoiesis-stimulating agents according to current guidelines [61].

It is also of note that the guidelines for the prevention of glucocorticoid-induced osteoporosis – that is, supplementation of calcium and vitamin D or bisphosphonate therapy – are not validated and therefore do not apply to rheumatic patients with CKD. Bone metabolism in CKD patients is totally different from that in nonrenal patients with its own needs for prophylaxis and therapeutic interventions. In addition, the use of bisphosphonates is mostly contraindicated in the CKD patient as it can increase the risk of adynamic bone disease and the risk of bisphosphonate-associated nephrotoxicity [62].

Finally, diagnostic procedures such as computed tomography or magnetic resonance imaging involving iodine-containing or gadolinium-containing contrast agents, respectively, must be used with great caution in CKD patients of stage 4 and stage 5. Radiocontrast agents often cause radiocontrast nephropathy that is associated with an increase of overall mortality, especially in CKD patients. If not avoidable, preventive measures such as discontinuation of diuretics and prehydration are mandatory to minimize the risk of radiocontrast nephropathy [63]. The potential benefits of prophylactic N-acetylcysteine for the prevention of radiocontrast nephropathy remain under debate. Nephrogenic systemic fibrosis is a rare but devastating complication in patients with end-stage renal disease that may develop after administration of linear gadolinium chelate-containing contrast agents [64]. An active inflammatory process at the time of gadolinium exposure is an important risk factor for the subsequent development of this condition. Magnetic resonance imaging using linear gadolinium chelates of patients with active rheumatic disease and end-stage renal disease must therefore be avoided, and cyclic gadolinium chelates may be preferred in stage 4 CKD patients.

**Conclusion**

Patients with rheumatic diseases must be routinely monitored by blood and urinary parameters for

### Table 4. Treatment of crystal arthropathies in chronic kidney disease and dialysis patients

| Drug                          | CKD stage 3 and stage 4 patients (GFR 15 to 59 ml/minute/1.73 m²) | Dialysis patients (GFR <15 ml/minute/1.73 m²) |
|-------------------------------|------------------------------------------------------------------|---------------------------------------------|
| **Pain control in acute gout attack** |                                                                  |                                             |
| NSAIDs                        | Avoid when possible or use with great caution                    | May be used with caution                    |
| Colchicine                    | Avoid when possible or use with great caution (for example, 0.5 mg three times per week) | Avoid whenever possible                     |
| Glucocorticoids              | Short-term use possible (for example, 20 mg/day)                 | Short-term use possible (for example, 20 mg/day) |
| Anakinra                     | Effective (100 mg/day for 3 days) but yet limited data published  | Not evaluated                               |
| **Maintenance therapy**       |                                                                  |                                             |
| Allopurinol                  | Reduce dose to 100 to 150 mg or avoid                            | Avoid                                       |
| Colchicine                   | Avoid when possible or use with great caution (for example, 0.5 mg three times per week) | Avoid whenever possible                     |
| Anakinra                     | Not evaluated                                                    | Effective in case reports (100 mg three times per week) |
| Febuxostat                   | Safe at GFR >30 ml/minute/1.73 m²                                 | Not evaluated                               |
| Benzbromarone                | Probably ineffective at GFR <50 ml/minute                        | Ineffective                                |

CKD, chronic kidney disease; GFR, glomerular filtration rate; NSAID, nonsteroidal anti-inflammatory drug.
concomitant CKD, which can be either a manifestation of the same systemic disease, a side effect of DMARD therapy, or an unrelated renal disease (Figure 1). All three entities affect the diagnostic and therapeutic strategies of the rheumatologist. It should therefore be left to the expertise of the nephrologist to determine whether renal biopsy is required for the diagnostic work-up. Patients should be subdivided into those needing more intense immunosuppressive treatment because of their kidney disease and those needing dose reduction or modification of their medication because of their kidney disease. Ideally, patients with rheumatic diseases and renal co-morbidity should be managed through close collaboration between a rheumatologist and a nephrologist.

Figure 1. Algorithm for screening and monitoring of renal co-morbidity in rheumatic disease patients. See text for details. 1) Regular screening may be performed on a 6-monthly to 12-monthly basis. CKD, chronic kidney disease; eGFR, estimate glomerular filtration rate; MTX, methotrexate; NSAID, nonsteroidal anti-inflammatory drug; UTI, urinary tract infection.

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Abbreviations
ANCA, anti-neutrophil cytoplasmic antibody; CKD, chronic kidney disease; DMARD, disease-modifying antirheumatic drug; GFR, glomerular filtration rate; GN, glomerulonephritis; IL, interleukin; iPTH, intact parathyroid hormone; ISN, International Society of Nephrology; MBD, mineral and bone disorders; NSAID, nonsteroidal anti-inflammatory drug; RA, rheumatoid arthritis; RPS, Renal Pathology Society; SLE, systemic lupus erythematosus; TNF, tumor necrosis factor.

Competing interests
The authors declare that they have no competing interests.

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