

Expert consensus for the diagnosis and treatment of patient with hyperuricemia and high cardiovascular risk: 2021 update

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While fundamental concepts outlined in the 2018 consensus [1] will be revisited, emphasis will be placed on the more recent developments.

Definition and epidemiology: Constantly increasing burden and importance of hyperuricemia

Uric acid (UA) is the end product of purine metabolism, and its concentration in blood can increase in humans, great apes, and dalmatian dogs as a consequence of a genetic mutation that occurred millions of years ago and contribute to human evolution from less evolved species [2]. These elevated plasma levels of UA are the final result of almost three different mechanisms under genetic control and involve UA production, renal excretion, and gut absorption [3]. Under physiological conditions, UA synthesis and excretion are balanced in the body. Once this balance is disturbed, it leads

to hyperuricemia (HU). Typically, male UA levels greater than 7 mg/dL (420 μ mol/L) and female UA levels greater than 6 mg/dL (360 μ mol/L) are considered hyperuricemia.

The latest scientific data published after our first consensus [4, 5] indicates that mean serum uric acid (sUA) has increased constantly according to the prevalence of concomitant diseases in many populations, while the prevalence of HU increases with age and is higher in men than premenopausal women due to estrogen's positive influence on urate excretion by the kidneys [4]. The United States National Health and Nutrition Examination Survey (NHANES 2007–2016) estimated the HU prevalence to be 20.2% for men (22.8 million) and 20.0% for women (24.4 million). In simple terms, 1 in 5 men and 1 in 5 women suffer from HU. The prevalence of sUA levels > 6.0 mg/dL was 32.3% overall (75.8 million), 49.5% among men (55.8 million), and 16.4% among women (20.0 million).

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The overall mean sUA level was 5.39 mg/dL (95% confidence interval [CI] 5.34–5.45), with mean sUA levels of 6.04 mg/dL and 4.79 mg/dL among men and women, respectively. Moreover, the prevalence rates of HU remained stable between 2007 and 2016 (p for trend > 0.05) [5]. Of note, the prevalence of HU increased with age, with the highest being 27.8% (3.1 million) among individuals aged 80 years or older. Among patients aged 65 years or older, the prevalence of HU was 27.2% (12.6 million) [5]. Recently published data by the Irish health system indicate that from 2006 to 2014, the prevalence of HU increased from 19.7% to 25.0% in men and from 20.5% to 24.1% in women ($p < 0.001$). Moreover, age-specific prevalence increased in all groups from 2006 to 2014, and the magnitudes of the increases were similar for each age category [6].

The prevalence of HU increased significantly with worsening renal function, from 12.2% in patients with estimated glomerular filtration rate (eGFR) > 90 mL/min to 63.9% in patients with eGFR < 15 mL/min [6].

The adoption of a Western lifestyle by natives of other countries and cultures and a change in the socioeconomic background through immigration to Western countries and movement from rural to urban communities have influenced sUA levels [7, 8].

The pathophysiological effect of hyperuricemia on cardiovascular disease: What matters more? Overproduction or underexcretion?

First and foremost, an increased sUA level is the result of a purine/fructose-rich diet, genetic or environmental factors, metabolic disorders, as well as either: (1) its endogenous overproduction (due to, e.g., purine-rich diet, an error of purine metabolism affected by phosphoribosyl-pyrophosphate synthetase or purine salvage pathway influenced by hypoxanthine-xanthine phosphoribosyl transferase; cell breakdown or excessive purine turnover: lymphoproliferative diseases, myeloproliferative disease, polycythemia vera, Paget disease [9], psoriasis, tumor lysis, hemolysis, rhabdomyolysis, and exercise; [10] or — in most cases — by (2) insufficient excretion (caused by acute or chronic kidney disease [CKD], acidosis: lactic acidosis, ketoacidosis; hypovolemia, medication/toxin: diuretic, niacin, pyrazinamide, ethambutol, cyclosporin, beryllium, salicylates, lead, alcohol; sarcoidosis, hyperparathyroidism, hypothyroidism, Bartter syndrome, and Down syndrome [11–13].

It should be emphasized that due to the limitations of the current knowledge on what is more important for unfavorable cardiovascular effect of HU — overproduction of UA (increased activity of xanthine oxidase with reactive oxygen species formation) or limited excretion (effects of UA per se), the topic has become one of the hottest in the field of cardiovascular research and hyperuricemia. However, the mechanism of impaired oxidative mechanism which was widely described in a previous version of this document seems to be more consistent.

Genetics: Possibilities for individualized diagnoses and care strategies?

There is growing evidence that genetics and environmental factors play a key role in HU's development [14]. Considering HU's pathophysiological aspects, it can be divided into overproduction (liver) and underexcretion types (gut, kidney). Indeed, genome-wide association studies identified the genetic basis of HU as dominated by loci containing urate transporters and interacting proteins involved in the excretion of urate; among them SLC2A9 (GLUT9), ABCG2, SLC22A11, SLC17A1–SLC17A4, and PDZK1; and proteins associated with metabolic pathways (e.g., GCKR, A1CF, IGF1R) [15]. Among these, GLUT9 and ABCG2 were recognized as the most significant [16]. Indeed, SLC2A9 (GLUT9) has a key role in urate transport and reabsorption. GLUT9-encoded protein is useful in urate excretion into urine and the reabsorption of urate into the blood. Differences among variants of GLUT9 also influence the excretion of UA in urine and its reabsorption to blood [17]. Systematic analysis of GLUT9's variants confirms its key role in the treatment of HU. Moreover, the ABCG2 gene (BCRP) is engaged in intestinal excretion and UA transport in proximal tubule epithelial cells [18]. Mutations in ABCG2 impede proper sUA regulation and lead to HU. Of note, in hemochromatosis patients, iron/heme overload enhances the activity of xanthine oxidase and — through p53 — causes the reduction of ABCG2 expression. This leads to a reduction in UA intestinal excretion and subsequent accumulation in tissue and serum, causing hereditary hemochromatosis-associated arthritis [19].

Among many others, one should acknowledge organic anion transporters 10 (OAT10, SLC22A13), acting as a key part of urate transport from urine to the blood; lactate dehydrogenase D (LDHD), decreasing excretion of UA [20];

hypoxanthine-guanine phosphoribosyltransferase (HGPRT) whose deficiency caused by an HPRT1 mutation leads to elevated UA levels in the blood, which are associated with Kelley-Seegmiller syndrome, Lesch-Nyhan syndrome, and HU [21]; mitochondrial seryl-tRNA synthetase precursor, a member of the class II tRNA synthetase family, which is involved in the ligation of serine to tRNA (Ser) and is involved in selenocysteinyl-tRNA (sec) biosynthesis in mitochondria [22]; xanthine dehydrogenase (XDH), influencing the oxidation of hypoxanthine to xanthine and the oxidation of xanthine to UA [23], therefore reducing the levels of xanthine oxidoreductase. Drabkin et al. [20] confirmed that a mutation could cause HU in LDHD within the putative catalytic site of the encoded d-lactate dehydrogenase, which results in increased blood levels of d-lactate — typically present in blood in miniscule amounts. As a consequence, excessive renal secretion of d-lactate in exchange for UA reabsorption culminates in HU. In line with the human phenotype, injection of d-lactate into naive mice resulted in HU [20].

These advances lead to a clear-cut approach to individualized patient care; genetic data can be informative about the prognosis in patients suffering from HU and help clinicians select dosage of urate-lowering therapy (ULT) and offer the correct advice on lifestyle changes.

Recent studies linking hyperuricemia to cardiovascular disease

Hyperuricemia and ischemic heart disease

We look forward to the results of the ALL-HEART study, which is a multicenter, controlled, prospective, randomized trial, examining the effects of allopurinol (up to 600 mg daily) vs. no treatment on cardiovascular outcome (non-fatal myocardial infarction, non-fatal stroke or cardiovascular death) in patients with coronary artery disease. The secondary goals are to determine the cost-effectiveness of adding allopurinol to usual therapy, determine whether allopurinol improves the quality of life, and to determine the safety and tolerability of giving allopurinol to patients with ischemic heart disease (without a history of gout). The main inclusion criteria were patients of 60 years of age and over and ischemic heart disease. The main exclusion criteria were history of gout, eGFR < 30 mL/min, and moderate-to-severe heart failure and significant hepatic disease [24].

Hyperuricemia and hypertension

An ample body of evidence widely acknowledges that the association between an increase in relative risk of hypertension and high levels of sUA remains independent of traditional risk factors [25–33]. A substantial meta-analysis of 18 studies confirmed an increase of 13% in the incidence of new-onset hypertension for every 1% increase in sUA levels [25]. The PAMELA (Pressioni Arteriose Monitorate e Loro Associazioni) study confirmed that a rise in sUA by 1 mg/dL was associated with a significant increase in the risk of developing the new-onset home and ambulatory hypertension (odds ratio [OR] 1.34, 95% CI 1.06–1.7, $p = 0.015$; OR 1.29, 95% CI 1.05–1.57, $p = 0.014$; respectively) [26]. Finally, the Saku study confirmed that HU predicted the risk of developing hypertension was independent of alcohol drinking status [34].

Hyperuricemia and stroke

Serum uric acid plays a key and influential role in the physiopathology of stroke [35]. Kim et al. [36] reported that HU was associated with a significantly higher risk of both stroke incidence (relative risk [RR] 1.41) and mortality (RR 1.36) in their meta-analyses of unadjusted study estimates. Accordingly, Zhong et al. [37] in their meta-analysis confirmed similar results; elevated sUA levels were significantly associated with an increased risk of stroke in both men (RR 1.10 per 1 mg/dL increase in sUA) and women (RR 1.11 [1.09–1.13]). In the newest study to date; CIRCS investigators presented that elevated sUA level is an independent predictor of total stroke in women but not in men. The positive association present in women was mostly attributable to ischemic stroke and was more pronounced among nonusers of antihypertensive medication [38].

Hyperuricemia and metabolic syndrome

As mentioned above, several studies demonstrated that sUA level is associated with metabolic syndrome, high body mass index (BMI), waist circumference, high fasting blood glucose levels, and dyslipidemia [39]. Shirasawa et al. [40] analyzed data derived from 96,863 participants and confirmed that the adjusted OR for HU was considerably increased in obesity (with central obesity) compared with normal weight, regardless of sex (men: OR 2.12, 95% CI 2.03–2.21; women: OR 3.54, 95% CI 3.21–3.90) and was statistically increased in normal weight with central obesity

compared to normal weight (men: OR 1.44, 95% CI 1.36–1.52; women: OR 1.41, 95% CI 1.27–1.57). They concluded that middle-aged Japanese adults with normal weight but having central obesity should be screened using a combination of BMI and waist to height ratio and be educated about how to prevent HU [40].

Hyperuricemia and atrial fibrillation

Currently, evidence on serum urate with the risk of atrial fibrillation (AF) is mainly from cross sectional studies, based on prevalent AF cases, and were limited by only a 1-time measurement of serum urate. However, in a large prospective cohort study with 123,238 participants conducted from 2006 to 2012, both an increased cumulative average and elevations in serum urate over time were associated with increased risk of incident AF (adjusted hazard ratio [HR] 1.91, 95% CI 1.32–2.76, $p = 0.001$ for trend). The combination of high sUA and high-sensitivity C-reactive protein levels was associated with a significantly increased risk of incident AF (adjusted HR 2.63, 95% CI 1.63–4.23). Li et al. [41] provided evidence of an association between a relatively common treatable metabolic alteration (higher serum urate) and a common cardiac rhythm disorder (AF) with substantial morbidity and mortality. Moreover, Hong et al. [42] confirmed that the association between sUA and AF was significant ($p = 0.001$) after adjusting for potential confounding factors.

Hyperuricemia and liver diseases

The relationship between HU and liver disease has not been clearly described. Undoubtedly, the increased level of sUA is the result of a diet rich in purine and fructose, genetic and environmental factors, metabolic disorders, as well as endogenous overproduction or — in most cases — impaired excretion of UA [43–46]. Uric acid synthesis is mainly influenced by phosphoribosyl pyrophosphate synthetase and the purine pathway [46]. Some publications indicate the association of increased levels of sUA with non-alcoholic fatty liver disease, which is a part of metabolic syndrome, and chronic hepatitis. In experimental studies, the UA-stimulated expression of aldose reductase in both cultured hepatocytes (HepG2 cells) and hyperuricemic rat livers were associated with endogenous fructose production triglyceride accumulation through UA-induced oxidative stress and stimulation of the nuclear transcription factor activated T5 cells (NFAT5). Uric acid also potentiated the effects of elevated glucose levels to stimulate the accumu-

lation of triglycerides in the liver. Hyperuricemic rats exhibited elevated hepatic aldose reductase expression, endogenous fructose accumulation, and fat accumulation was significantly reduced by allopurinol co-administration. Thus, HU is correlated with the occurrence of hypertriglyceridemia and non-alcoholic fatty liver disease, which secondarily induces the development of non-alcoholic steatohepatitis [47, 48].

Jang et al. [49] investigated UA levels and their relationship to disease progression in 373 patients with biopsy-confirmed chronic hepatitis C (CHC) enrolled in interferon-based antiviral therapy. In this study HU was defined as UA levels > 7 mg/dL in men and > 6.0 mg/dL in women. Hyperuricemia was found in 15.8% of CHC patients, but UA levels did not differ between CHC patients and the control group of healthy subjects matched for sex and age (5.54 ± 1.20 mg/dL vs. 5.45 ± 1.45 mg/dL, $p = 0.3$). Logistic regression analysis showed that factors related to HU in men included BMI (OR/CI 1.12/1.05–1.30, $p = 0.006$) and advanced fibrosis (F3–4) (OR/CI 0.27/0.09–0.83, $p = 0.02$), while factors associated with HU in women included eGFR (OR/CI 0.97/0.95–0.99, $p = 0.02$) and diabetes (OR/CI 3.03/1.11–8.25, $p = 0.03$). There was a significant downward trend in sUA levels with the progression of fibrosis in men (6.21 ± 1.03 mg/dL, 5.82 ± 1.16 mg/dL, and 5.44 ± 1.28 mg/dL in stages F0–2, F3 and F4, respectively, trend $p = 0.01$), indicating that HU was inversely related to the severity of liver disease in men with CHC [49]. Petta et al. [50] do not share this opinion; in a study of 496 patients with biopsy-confirmed CHC treated with pegylated interferon and ribavirin, there was no independent association between UA levels and necroinflammatory activity, fibrosis, nor sustained virologic response. However, the association of HU with the severity of steatosis was confirmed (OR 3.176, 95% CI 1.828–5.517, $p < 0.001$), which could potentially be a therapeutic target in the treatment of CHC [50]. Moreover, Jang et al. [51], in a group of 213 patients, where HU was defined as UA levels > 7.0 mg/dL in men and > 6.0 mg/dL in women, showed that sUA levels decreased significantly after eradication of hepatitis C virus by the use of direct antiviral drugs. Improvement was only observed in patients with fibrosis-4 ratio (FIB-4) < 6.5 (37.1% vs. 25.7%, $p = 0.001$). The multivariate analysis showed that factors associated with significantly reduced sUA levels were FIB-4 < 6.5 (OR/95% CI 3.22/1.04–9.95, $p = 0.04$) and eGFR < 60 mL/min/1.73 m² (OR/95% CI 4.34/1.94–9.73, $p < 0.001$ [51]).

Hyperuricemia and COPD

Identification of prognostic biomarkers for chronic obstructive pulmonary disease (COPD) may help improve therapy for patients at high risk. In univariate analysis, HU was associated with a higher risk of mortality in patients with COPD (HR 2.29, 95% CI 1.07–4.88, $p = 0.032$). Further analysis confirmed that HU was independently associated with a higher risk of mortality in patients with COPD (HR 2.68, 95% CI 1.18–6.09, $p = 0.019$) [52]. Moreover, it was recently published that sUA could activate the nucleotide-binding oligomerization domain-like receptor family pyrin domain-containing 3 (NLRP3) inflammasome, leading to interleukin (IL)-1 β secretion. In COPD patients, UA and uric ratio with creatinine (UCR) were positively associated with white blood cells, C-reactive protein, and IL-1 β . COPD smokers had lower UA and UCR values. Standard COPD therapy did not affect UA or UCR, while patients with cardiovascular diseases (CVD) had higher UA, but not UCR, levels. Multiparameter models of UA and UCR that included IL-1 β were able to correctly classify 86% and 90% of cases, respectively [53].

Current uric acid thresholds predicting cardiovascular events: Including improved cardiovascular risk estimation

As described in a previous document [1], sUA is considered an independent factor in the development of a wide variety of vascular disorders: hypertension [54], metabolic syndrome [55, 56], coronary artery disease [57], diabetes [58], cerebrovascular disease [59, 60], CKD [61], as well as other CVD [62, 63] and — conversely — these comorbidities increase the incidence of HU [64]. It is worth paying attention to other conditions, both those from the genre of metabolic syndrome (e.g., non-alcoholic fatty liver, non-alcoholic steatohepatitis) and those whose presence may affect the clinical course of disorders in the cardiovascular system (e.g., COPD [65] or incident asthma in men [66]).

What we know so far, is that some studies have found a robust association between sUA level and CVD not only in patients with clearly diagnosed HU but also in those with values considered normal to high > 5.2 – 5.5 mg/dL [67–69]. Moreover, this relationship concerned both subclinical and clinical manifestations of diseases [64], and remained highly significant even after renal function adjustment.

Of note, the Working Group on sUA and cardiovascular risk within the Italian Society of Hy-

per-tension designed the URRAH (Uric Acid Right for Heart Health) study, whose aim was to assess — in a group of 22,714 subjects — the level of UA above which the independent risk of CVD may increase in a significant manner in the general population [70]. Viridis et al. [71] confirmed that the threshold UA levels were **4.7 mg/dL** (95% CI 1.21–1.93 mg/dL) for increasing all-cause mortality, **5.6 mg/dL** (95% CI 4.99–6.21 mg/dL) for increasing cardiovascular mortality, and were significantly lower than levels referenced in clinical diagnostic criteria [71]. Considering the sex of the patients, the threshold for sUA for all-cause mortality was 5.4 mg/dL (95% CI 4.80–6.57) in men and 4.7 mg/dL (95% CI 4.40–5.10) in women. Most importantly for clinical practice is that the novel sUA thresholds allow for a significant net reclassification of — and improvement in — the Heart Score risk chart's present values for all-cause and cardiovascular mortality, 0.26 and 0.27, respectively.

In summation, this large-sample study led to a significant improvement in risk classification of the well-validated and guideline-recommended scale, the Heart Score. Further analysis for factors not included in the Heart Score such as hematocrit, diuretics, alcohol consumption, BMI, and eGFR did not significantly affect results [71]. The results reported by Viridis et al. [71] are largely superimposable over those obtained in several different populations of subjects enrolled in observational studies and show an increase in the relative risk of major cardiovascular events in the presence of sUA levels ranging between 4.5 and 5.5 mg/dL [72–75]. Previously reported investigations in smaller groups gave partially corroborative results specifically in the following contexts: in comparison to the Framingham Risk Score [72], in later life only [73], in patients undergoing percutaneous coronary interventions due to acute coronary syndrome [74], and in hypertensive patients; conflicting results were yielded in NHANES III [75]. The lower threshold level reported for CVD compared with gout can be explained by the causative role of the oxidative stress associated with the production of sUA by xanthine-oxidase that occurs with lower concentrations of serum urate and is largely independent of the inflammatory role of UA deposition.

A large body of evidence confirms the contribution of HU to the worsening of cardiovascular, diabetic, lipid, and renal diseases [76–78]; thereby supporting the conclusion that increased sUA levels correlate with elevated cardiovascular risk [79, 80] and further stressing the importance of a redefinition of threshold levels referred to during

identification of patients at risk of CVD in the presence of hyperuricemia.

URRAH researchers identified the prognostic cut-off values of sUA in predicting fatal and morbid heart failure; sUA more than 5.34 mg/dL (CI 4.37–5.6, sensitivity 52.32, specificity 63.96, $p < 0.0001$) was the univariate prognostic cut-off value for all heart failure, whereas sUA more than 4.89 mg/dL (CI 4.78–5.78, sensitivity 68.29, specificity 49.11, $p < 0.0001$) was the prognostic cut-off value for fatal heart failure [81]. Moreover, Huang et al. [82] analyzed ten studies involving 12,854 acute heart failure patients and confirmed that acute heart failure patients with the highest sUA levels had an increased risk of all-cause mortality (risk ratio [RR] 1.43, 95% CI 1.31–1.56) and a combined endpoint of death or readmission (RR 1.68, 95% CI 1.33–2.13), after adjusting for potential variables. Therefore, elevation in sUA levels significantly increased risks of all-cause mortality and combined endpoint of death or readmission by 11% and 12%, respectively, for every 1 mg/mL, they were elevated [82]. Also, URRAH researchers adjusted for such confounders as age, arterial hypertension, diabetes, CKD, smoking habits, ethanol intake, BMI, hematocrit, low-density lipoprotein cholesterol, and use of diuretics in multivariate Cox regression analyses and identified an independent association between sUA and fatal myocardial infarction across the whole database (HR 1.381, 95% CI 1.096–1.758, $p = 0.006$) and in women, specifically (HR 1.514, 95% CI 1.105–2.075, $p < 0.01$), but not in men [83].

Hyperuricemia and cardiovascular events: A high serum uric acid level and its influence on cardiovascular outcome

As we mentioned earlier, several studies have confirmed the relationship between sUA and CVD mortality [26, 84]. However, Rahimi-Sakak et al. [85] performed a meta-analysis of 44 prospective cohort studies with dose-response analysis published between 2000 and 2018 to determine the relationship between sUA and CVD mortality. Pooled results confirmed a significant positive association between sUA levels and risk of CVD mortality (HR 1.45, 95% CI 1.33–1.58, $I^2 = 79\%$). Sub-group analysis yielded that this association was stronger in women compared to men. Also, there was a significant non-linear association between sUA levels and CVD mortality risk ($r = 0.0709$, $p = 0.001$) [85]. In a cross-sectional study, Lee et al. [86] investigated the relationship of sUA with CVD risk in the Korean adult general

population (8781 participants from first and second years of the Seventh Korea National Health and Nutrition Examination Survey 2016–2017). There was a significant association of sUA with 10-year CVD risk scores after adjusting for physical activity, BMI, serum creatinine, and alcohol consumption in both sexes ($p < 0.001$); at sUA levels of 6.9 mg/dL, the CVD risk was lowest [86].

Hyperuricemia treatment and cardiovascular outcomes: Allopurinol remains first-line urate-lowering therapy

In a systematic review of 24 guidance documents, 19 of them provided target levels for long-term sUA control, most of which recommended 6.0 mg/dL (or 360 $\mu\text{mol/L}$), except the South African guidelines, which recommended 5.0 mg/dL (300 $\mu\text{mol/L}$) [87]. Also, the Polish Society of Hypertension Guidelines 2019 recommend 5.0 mg/dL level for long-term sUA control [88]. Still, the definition of HU varies greatly across clinical trials, making epidemiological reports somewhat inconsistent and difficult to compare.

Xanthine-oxidase inhibitors (XOI), especially allopurinol, are still recommended by almost all guidelines as a **first-line ULT**. The results on febuxostat have recently been implemented after publication of the FAST trial (long-term cardiovascular safety of febuxostat compared with allopurinol in patients with gout: a multicenter, prospective, randomized, open-label, non-inferiority trial) [89, 90] that reported divergent results from those described by the CARES (Cardiovascular Safety of Febuxostat and Allopurinol in Patients with Gout and Cardiovascular Morbidities) study [91].

Febuxostat: Further clinical trials needed?

Febuxostat, a nonpurine XOI, is recommended in patients refractory or intolerant to allopurinol and requires no dose limitation in CKD stages 1–3. Febuxostat induces potent inhibition of XO and greater hypouricemic activity than the commonly used allopurinol doses [92]. However, preliminary results from a safety trial with febuxostat versus allopurinol, mainly based on a large-scale, randomized study design has suggested a modestly higher rate of cardiovascular events with febuxostat [93]. Based on the preceding study's findings, treatment with febuxostat in patients at high cardiovascular risk has not been recommended. Another safety oriented randomized controlled trial reporting results of a 32-month follow-up ($n = 6190$), demonstrated the compa-

rable effects of febuxostat and allopurinol on the primary cardiovascular end-point with a higher rate in the secondary objectives of all-cause and cardiovascular mortality in the febuxostat group than in the allopurinol group (HR for death from any cause 1.22, 95% CI 1.01–1.47; HR for cardiovascular death 1.34, 95% CI 1.03–1.73) [91]. In the meantime, a meta-analysis of 35 studies did not show a significant difference between febuxostat and allopurinol in cardiovascular events (RR 1.69, 95% CI 0.54–5.34, $p = 0.37$) [94]. Moreover, in the FREED study of over 1000 elderly patients with HU, 25% relative risk reduction in composite with death to any cause, cerebrovascular disease, non-fatal coronary artery disease, heart failure requiring hospitalization, an arteriosclerotic disease requiring treatment, renal impairment, and AF was observed in the febuxostat group compared to the non-febuxostat group. There was no difference in cardiovascular clinical outcomes examined separately with febuxostat versus control treatment.

In contrast, the European Medicines Agency (EMA)-required Febuxostat versus Allopurinol Streamlined Trial (FAST), published in the *Lancet*, does not support the finding of an increased cardiovascular risk of febuxostat, despite using higher (EMA-approved) dosages compared to the CARES trial [90]. In 6128 patients (with previous CVD), for the incidence of the primary endpoint (a composite of hospitalization for non-fatal myocardial infarction or biomarker-positive acute coronary syndrome, non-fatal stroke, or cardiovascular death), on-treatment, febuxostat (172 patients [1.72 events per 100 patient-years]) was non-inferior to allopurinol (241 patients [2.05 events per 100 patient-years]; adjusted HR 0.85 [95% CI 0.70–1.03], $p < 0.0001$) [90]. In editorial comments, Bardin and Richette [89] underlined that patients in the CARES study had more severe gout than those in the FAST study and that all patients in CARES had a history of CVD in contrast to 2046 (33.4%) of 6128 in FAST. No excess of deaths was observed in this subgroup of patients in FAST. Still, their group size might be insufficient to fully assess febuxostat risk in patients with severe CVD [89]. Thus, further clinical trials are needed to clarify this issue and provide clear evidence for the withdrawal of the Food and Drug Administration alert in this issue.

Uric acid and COVID-19

There are not many studies on the relationship between sUA and coronavirus disease 2019 (COVID-19). However, of note that favipiravir, a purine nucleic acid analog and antiviral agent

studied in Japan to treat COVID-19, has frequent side effects [95]. Favipiravir's action causes it as a moderate inhibitor of organic anion transporters 1 and 3 (OAT1 and OAT3), which are involved in UA excretion in the kidney, as well as by its influence on UA reuptake via urate transporter 1 (URAT1) in the proximal renal tubules. Elevated UA levels were found to return to normal after discontinuation of favipiravir, and favipiravir is not recommended for long periods to treat viral infection [96, 97]. Moreover, in a Japanese, prospective, randomized, open-label trial of early versus late favipiravir treatment in hospitalized patients with COVID-19, 84.1% developed transient HU [98].

Management strategies: Updated five-step recommendations for the treatment of patients with increased serum uric acid levels (Fig. 1)

STEP 1: Assess serum uric acid level

The measurement of sUA concentration is recommended as a part of screening in cardiac/hypertensive patients by experts of both the European Society of Cardiology and the European Society of Hypertension [99].

Our recommendation remains unchanged: the optimal target of sUA levels should be 6 mg/dL (360 $\mu\text{mol/L}$). Serum uric acid levels should be monitored regularly and maintained at < 6 mg/dL. Still, despite the lack of randomized controlled trials, one should consider an sUA target of < 5 mg/dL in patients with high cardiovascular risk comprised of at least two of the following: hypertension, diabetes, dyslipidemia, organ target organ damage or previous cardiovascular events.

STEP 2: Check comorbidities and active treatments and stop the administration of drugs that influence serum uric acid

Appropriate strategies should be defined and implemented in patients with HU, regarding more aggressive control of concomitant risk factors and the use of drugs indirectly affecting UA levels. Efficient management of concomitant diseases that influence sUA levels, such as hypertension, type 2 diabetes, metabolic syndrome, CKD, and CVD [56, 61–64] should be the procedure of choice. In clinical scenarios, feasible changes should be considered if the potential benefits exceed the potential harms, particularly:

- Diuretics, among them hydrochlorothiazide
 - switching from hydrochlorothiazide to alternative antihypertensive agents, if possible;

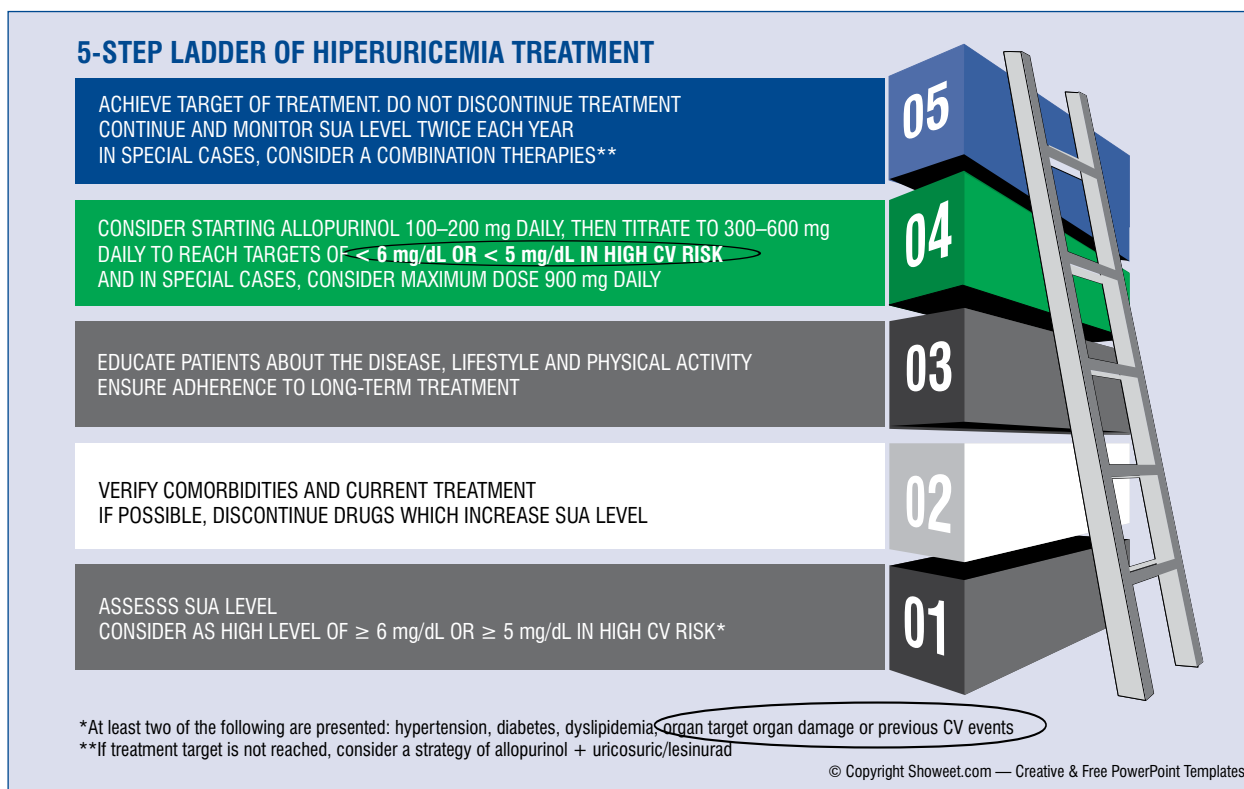


Figure 1. Management strategy for patients suffering from hiperuricemia; CV — cardiovascular; SUA — serum uric acid.

- Angiotensin II receptor blockers (ARBs) — although losartan is the only antihypertensive drug that lowers sUA level [100], switching from other ARBs to losartan is NOT recommended;
- Low-dose acetylsalicylic acid (ASA) — cessation of low-dose ASA in primary cardiovascular prevention patients or switching to alternatives, if possible; cessation of low-dose ASA in secondary cardiovascular prevention patients is NOT recommended;
- Cholesterol-lowering agents — switching from cholesterol-lowering drugs to fenofibrate is NOT recommended.

Building multidisciplinary teams for optimal diagnostics and management strategies, with an appropriate estimation of HU’s significance, is necessary. It is essential to improve adherence to clinical practice guidelines, raise awareness of HU and associated comorbidities, and prompt their more diligent and specific monitoring.

STEP 3: Recommended life-style changes

Among the most significant lifestyle changes are:

- Limiting intake of purines, including red meat and seafood;

- Limiting intake of high-fructose corn syrup. Based on the Third National Health and Nutrition Examination Survey, sUA levels increase with increasing sugar-sweetened soft drink intake. After adjusting for covariates, sUA levels associated with sugar-sweetened soft drink consumption categories (< 0.5, 0.5–0.9, 1–3.9, and ≥ 4 servings/day) were greater than those associated with no intake (by 0.08, 0.15, 0.33, and 0.42 mg/dL, respectively) (95% CI 0.11–0.73, p < 0.001 for trend) [101];
- Limiting alcohol. Limiting or abstaining from alcohol leads to decreased UA levels by 1.6 mg/dL compared to the control group [102];
- Weight loss and regular physical activity for patients who are overweight or obese [103, 104];
- Adding coffee, dairy products, cherries [100, 105], and ascorbic acid [106].

STEP 4: Give xanthine oxidase inhibitors as first-line therapy, titrated to achieve serum uric acid target

As mentioned earlier, allopurinol, XOI, is recommended as a first-line ULT in agreement with most of the guidelines. According to the summary of product characteristics of allopurinol, the rec-

ommended initial allopurinol dosage is 100 to 200 mg daily in mild conditions, 300 to 600 mg daily in moderate conditions, 700 to 900 mg daily in severe conditions. **The dosage should be titrated up to achieve the desired sUA target level** [107].

Importantly — due to its renal excretion — in patients with CKD, impaired renal function may lead to retention of the drug and/or its metabolites (oxypurinol) with subsequent prolongation of plasma half-lives. For this reason, in severe CKD, it may be appropriate to use less than 100 mg per day or to use single doses of 100 mg at longer intervals than 1 day. In special situations and the presence of appropriate equipment, the dose should be adjusted to maintain plasma oxypurinol levels below 100 $\mu\text{mol/L}$ (15.2 mg/L). If allopurinol is used in dialysis patients, it should be administered at a 300–400 mg dose immediately after dialysis, but without additional doses on other days [107].

STEP 5: Achieve targeted serum uric acid levels, do not stop treatment, continue monitoring serum uric acid levels twice per year; in special cases, consider combined therapy

Only 2 in 5 patients with HU reached the target of sUA with this therapy [108]. Suppose that the sUA target cannot be achieved. In this case, the dose should be escalated with supervision up to 900 mg of allopurinol, or the patient should be switched to benzbromarone or combined therapy of benzbromarone and allopurinol (**STEP 5**), except in patients with eGFR of $< 30 \text{ mL/min}$ [109]. However, escalations should be performed carefully to achieve optimal treatment goals, mainly due to allopurinol hypersensitivity syndrome and severe cutaneous allergic reactions, usually after 8 weeks of therapy [110–112]. Factors that are known to contribute to the development of this syndrome include initial doses that are too high, CKD, accompanying use of diuretics, and the presence of HLA-B*5801 [113, 114].

Although high-dose allopurinol ($\geq 300 \text{ mg/dL}$) is associated with a reduced risk of all-cause mortality [7, 115], consideration of an optimal dose seems to be a major factor design of future research.

Lesinurad is an oral selective inhibitor of URAT1 and OAT4 renal transporters, which increase renal UA excretion and lower sUA levels by inhibiting UA reabsorption. A dose of 200 mg daily is recommended in combination with XOIs in patients who do not achieve treatment targets. Adding lesinurad can increase the efficiency of XOIs

(compared to monotherapy) and help avoid maximal XOI dosages [116]. In a CLEAR study, lesinurad of 200 mg or 400 mg together with allopurinol significantly increased the proportion of patients who achieved the sUA target levels compared to allopurinol (54.2%, 59.2%, and 27.9%, respectively, $p < 0.0001$) [117]. The approval of lesinurad was based on data from three pivotal phase III studies (CLEAR 1, CLEAR 2, and CRYSTAL), which assessed lesinurad 200 mg and 400 mg doses. The target sUA level was achieved by significantly more patients on lesinurad 200 mg plus allopurinol group (CLEAR 1 and CLEAR 2 trials) or lesinurad 200 mg plus febuxostat group (CRYSTAL study) compared to patients who received either XOI alone. The safety profile of lesinurad 200 mg plus an XOI was comparable to allopurinol or febuxostat alone. To summarize, lesinurad, in combination with allopurinol, is a novel option for the treatment of HU in adults with gout who have not achieved their target sUA levels with allopurinol alone (STEP 5) [118, 119]. Once the sUA target is achieved continuously, ULT's dose should be maintained indefinitely with ongoing monitoring of sUA levels twice a year (**STEP 5**) [117, 120].

Many unresolved questions still remain: Areas in need of further study

First and foremost, UA's treatment target may still need to be reconsidered, especially since data from the URRAH study identified new cardiovascular thresholds and improved algorithms for assessing total cardiovascular risk. Still, there is a clear need for further evidence to support the treatment of asymptomatic HU, although a large body of evidence does show the beneficial effect of ULT on cardiovascular results.

Most relevant recommendations: The take home message for the clinical practitioners

In conclusion, we would like to summarize our opinions that should be helpful to clinicians treating patients suffering from hyperuricemia and at high cardiovascular risk:

- 1 in 5 patients suffers from HU. The prevalence of HU is continuously increasing;
- All patients with HU should be effectively informed about environmental and pharmacological factors influencing HU, comorbidities, and cardiovascular risk factors; be advised about the immediately required lifestyle and

diet modifications and weight loss, if necessary; and strict adherence to recommended treatments;

- Both patients and physicians of all specialties (especially primary care physicians, cardiologists, and pulmonologists) should strive to obtain and maintain lifelong sUA levels lower than 6 mg/dL; for patients at high cardiovascular risk, the target level should be 5 mg/dL;
- As mentioned earlier, allopurinol — a XO1 — is recommended as a first-line ULT. According to the summary of product characteristics of allopurinol, the recommended initial allopurinol dosage is 100 mg to 200 mg daily in mild conditions, 300 mg to 600 mg daily in moderate conditions, and 700 mg to 900 mg daily in severe conditions;
- The dosage of XOIs should be titrated up to achieve the desired sUA target level and monitored twice a year thereafter to establish the correct level of sUA;
- If the sUA target levels are not reached, combined therapy of allopurinol + lesinurad might be considered.

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