COPENHAGEN — New European SLE Guidelines for Clinical Practice Announced

The 10 recommendations include:

► **Patient assessment.** In addition to routine clinical practice, “the assessment of SLE patients must include an evaluation of disease activity with a validated index at each visit and an annual evaluation of organ damage,” said Dr. Mosca.

Additionally, general quality of life, as ascertained by patient history and by a 0-10 visual analog scale, comorbidities, and drug toxicity should be assessed at each visit.

► **Cardiovascular risk factors.** At the baseline visit, and at least once annually during follow-up, “ask patients about smoking, vascular events, physical activity, their use of oral contraceptives and other hormonal therapies, and family history,” Dr. Mosca said.

Lipid profile and serum glucose measurement should be done at baseline and annually thereafter, as should examination of blood pressure and body mass index or waist circumference. “Depending on the findings, a patient may require more regular follow-up for specific conditions,” she said.

► **Other comorbidities.** Individuals with lupus are at increased risk for certain comorbidities, in particular osteoporosis, said Dr. Mosca. “Corticosteroid medications can trigger bone loss; disease-associated pain and fatigue can lead to inactivity, further increasing the osteoporosis risk; and bone loss may occur as a direct result of the disease.”

The guidelines recommend assessing all SLE patients for adequate calcium and vitamin D intake, regular exercise, and smoking status.

SLE patients should be screened and followed for osteoporosis according to the guidelines for postmenopausal women, for patients on steroids, or for patients on any other drug that may interfere with bone mineral density, she said at the meeting.

Studies have also shown that SLE patients are at an increased risk for certain cancers, “yet lupus patients tend to undergo screening less often than do individuals in the general population,” possibly because lupus-related concerns may take precedence, said Dr. Mosca. The guidelines recommend cancer screening according to the guidelines for the general population.

“It’s up to the clinicians who care for these patients to encourage appropriate screening,” she said.

► **Infection risk.** Lupus patients should be screened for HIV based on individual risk factors, and they should be screened for the hepatitis C and the hepatitis B viruses and for tuberculous according to local guidelines before beginning immunosuppressive therapy, according to the recommendations.

During immunosuppressive therapy, selected SLE patients should be tested for cytomegalovirus infection, because it increases the degree of immunosuppression of cell-mediated immunity, Dr. Mosca said.

Because of the increased risk of infection in SLE, patients should receive only inactivated pneumococcal and influenza vaccines, according to CDC guidelines for immunosuppressed patients, “preferably during periods of inactive disease,” she said.

► **Frequency of assessment.** The recommendations suggest patient assessments every 6-12 months for individuals with inactive disease, no organ

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PsA, Others Join Diabetes as CVD Risk Factors

BY MITCHEL L. ZOLER

COPENHAGEN — Psoriatic arthritis, rheumatoid arthritis, and ankylosing spondylitis are as strong as diabetes as risk factors for cardiovascular disease, prompting a European League Against Rheumatism task force to issue the group’s first consensus recommendations for managing cardiovascular risk in these patients. “In our view, rheumatoid arthritis [RA], ankylosing spondylitis [AS], and psoriatic arthritis [PsA] should be seen as new, independent cardiovascular risk factors,” Dr. Michael T. Nurmohamed said at the annual European Congress of Rheumatology. “The risk is comparable to type 2 diabetes,” added Dr. Nurmohamed, of Free University Medical Center in Amsterdam.

Cardiovascular risk management is currently necessary in patients with RA, AS, or PsA, and should involve assessing and treating conventional cardiovascular disease (CVD) risk factors as well as suppressing the underlying inflammatory process by treatment with disease-modifying antirheumatic drugs (DMARDs). “Most important is to decrease the inflammatory burden as much as possible,” through the use of biologic and/or synthetic DMARDs, he said in an interview. “The extent to which antirheumatic treatment decreases the risk is not known.”

Just as cardiovascular disease is now the most feared outcome of diabetes, it may be time to expand the definition of clinical impact of RA, AS, and PsA to include the extra CVD burden they trigger, Dr. Nurmohamed said.

Designation of RA, AS, and PsA as CVD risk factors for a task force of the European League Against Rheumatism (EULAR) is the first time a major medical group has singled out these conditions in this way. The extra risk from these disorders is substantial.

When a clinician uses the European SCORE (Systemic Coronary Risk Evaluation) formula to calculate an RA patient’s 10-year risk for cardiovascular disease death, this number is increased by 50% to get the patient’s actual risk when at least two of three criteria are present: disease duration more than 10 years, positivity for rheumatoid factor or anti-cyclic citrullinated peptide antibody, or extra-articular manifestations.

Dr. Nurmohamed based his recommendation on findings from an analysis done with his associates that found a greater than twofold increased risk for CVD in patients with RA, compared with people without RA. The higher level of conventional risk factors among the RA patients in the study explained roughly half of the doubled risk. The other half of the increased risk was directly attributable to RA, he said.

Similarly, a person’s Framingham risk score for having a cardiovascular event should also be boosted by about 50% if RA, AS, or PsA is present, he said. Major evidence for the increased CVD risk in patients with inflammatory arthritis is now well recognized. “It is awareness that something should be done,” he said in the interview. But the extent to which the new guidelines are already routinely followed in Europe by physicians who manage these patients is variable, he said. “Cardiovascular risk management in patients with rheumatoid diseases is uncommon.”

He acknowledged that the evidence supporting a CVD effect is stronger for RA than for AS and PsA, but added that adequate evidence exists to support including AS and PsA in the recommendations. Dr. Nurmohamed itemized the other eight task force recommendations:

- Adequate control of rheumatoid activity is necessary to lower a patient’s CVD risk.
- A CVD risk assessment following evidence-based EULAR guidelines is recommended annually for all RA patients, and should be considered for all patients with AS and PsA.
- CVD risk score models should be multiplied by 1.5 when an RA patient has at least two of the following three criteria: disease duration of more than 10 years, positivity for rheumatoid factor or anti-cyclic citrullinated peptide antibody, and extra-articular manifestations.
- The total cholesterol-HDL cholesterol ratio should be used in the formula for estimating CVD risk.
- Interventions with lipid-lowering drugs and with antihypertensive medications should follow national guidelines.
- Statins, ACE inhibitors, and angiotensin receptor blockers are the preferred treatment agents because of their pleiotropic effects, he said.
- With corticosteroids are prescribed, they should be at the lowest possible dose. Dr. Nurmohamed reported having no financial conflicts.

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damage, and no comorbidities. The treating clinician should emphasize prevention at the time of these assessments, Dr. Mosca said.

■ Laboratory assessment. According to the guidelines, baseline laboratory assessment should include testing for antinuclear antibodies (ANA), anti-phospholipid (aPL) antibodies, Complement 3 (C3) and Complement 4 (C4), as well as the following autoantibodies: anti-double stranded DNA (anti-dsDNA), anti-Ro, anti-La, and anti-ribonuclear protein (RNP).

Prior to pregnancy, previously negative patients should be re-evaluated for aPL, anti-Ro, and anti-La antibodies. Prior to surgery, transplant, or the initiation of estrogen-containing treatments, or in the presence of a new neurologic or vascular event, previously negative patients should be tested for aPL, according to Dr. Mosca.

A 12-month intervals in patients with inactive disease, “we recommend performing a complete blood cell count, erythrocyte sedimentation rate, C-reactive protein, serum albumin, serum creatinine, and urinalysis,” she said. “Monitoring should be tailored to specific treatment drugs, when necessary.”

■ Mucocutaneous involvement. “Mucocutaneous lesions should be characterized, according to existing classification systems, as to whether they may be lupus-specific, lupus non-specific, lupus mimickers, or drug related,” Dr. Mosca reported. “All lesions should be assessed for activity and damage using validated indexes.”

■ Kidney involvement. Monitoring recommendations in this domain depend on kidney status.

“Patients with persistently abnormal urinalysis or creatinine should have a urine protein/creatinine ratio or 24-hour proteinuria test,” she said. “A urinalysis should be done at least every 3 months.”

■ Eye assessment. “Eye damage in patients with lupus varies from minor problems to severe retinopathy. A small percentage of lupus patients develop scleritis, retinal vasculitis, cotton wool spots at the back of the eyeball, or retinal bleeding and swelling of the optic disc. According to the guidelines, patients on steroids or antimalarial drugs should undergo a baseline eye examination according to standard recommendations. Annual follow-up eye exams are recommended in selected patients taking steroids and those at high risk for eye problems.”

“In patients taking antimalarial drugs who are low risk for eye problems, no further testing is required until after 5 years from baseline, at which point yearly examinations are recommended,” Dr. Mosca said.

In addition to facilitating good clinical practice, monitoring SLE are expected to “improve the quality control of care for lupus patients and to standardize the collection and comparison of data in observational studies,” Dr. Mosca concluded.

The economic implications of the guidelines are expected to be published in the Annals of Rheumatic Disease later this year, were developed by an expert panel using a three-staged consensus approach comprising a discussion of relevant categories, a comprehensive literature review of specific assessment, and the integration of the evidence with expert opinion, said Dr. Mosca.

She reported having no financial conflicts of interest to disclose.

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