

Abstract 428 Figure 2

sequelae. Compared with controls of 130 SLE patients without HZ, cases were significantly more likely to have received high-dose prednisone 65/65 (OR 16.41,  $p=0.0066$ ) with mean prednisone  $18.5 \pm 12$  mg/day and cyclophosphamide (Cyc) 19/65 (OR 7.05,  $p<0.0001$ ). IV Cyc with mycophenolate mofetil (MMF) conferred greatest risk for HZ infection. There was no association of disease activity with HZ risk, whereas hydroxychloroquine was a negative risk factor for HZ infection (OR 0.26,  $p=0.0005$ ).

**Conclusions** Immunosuppressive agents and corticosteroids are risk factors associated with development of HZ in SLE. On the other hand, hydroxychloroquine appeared to have a protective role against HZ.

#### 431 A MULTICENTER STUDY OF CLINICAL FEATURES AND REMISSION IN PATIENTS WITH SYSTEMIC LUPUS ERYTHEMATOSUS IN CHINA

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**Background and aims** To study the clinical characteristics and remission rate in systemic lupus erythematosus (SLE), and to investigate potential factors affecting remission. These data may provide evidence for rational medication of SLE.

**Methods** Clinical remission was defined as follows: SLEDAI  $\leq 4$ , with no SLEDAI activity in major organ systems (renal, central nervous system, cardiopulmonary, vasculitis, hemolytic anaemia, fever) and no gastrointestinal activity; Current prednisone (or equivalent) dose  $\leq 7.5$  mg daily; Well-tolerated standard maintenance doses of immunosuppressive drugs, hydroxychloroquine, and/or approved biologic agents. A cross-sectional survey was undertaken in 11 hospitals of China from October 2013 to April 2014. Clinical data of 485 consecutive SLE patients were collected.

**Results** 1. A total of 82 patients (17.5%) achieved clinical remission. Patients who received hydroxychloroquine or immunosuppressant therapy for more than 6 months yielded a higher remission rate of 24.0% (69/288). 2. The factors, including gender, age, marriage, education background, work environment, income and history of autoimmune diseases, had no significant correlations with the remission of SLE. 3. There are 51.3% of the patients with SLEDAI  $\leq 4$ . And 59.8% patients complained of symptoms, the most often symptoms were alopecia, Raynaud's phenomenon and arthritis. Anti-nuclear Antibody, anti-dsDNA, and hypocomplementaemia

were common seen in SLE patients. 4. Most SLE patients received small doses of glucocorticoid. Hydroxychloroquine is a common choice in the SLE therapeutics in China.

**Conclusions** The clinical remission among SLE patients is infrequent. Half of the patients are in a stable state. In order to target remission, prevent damage and improve quality of life, treating-to-target-in-SLE should be recommended.

#### 432 PERICARDIAL EFFUSION AND CARDIAC TAMPONADE IN SYSTEMIC LUPUS ERYTHEMATOSUS

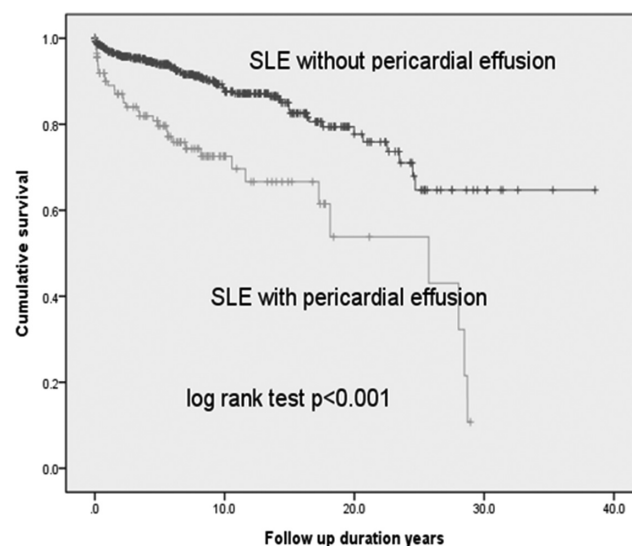
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**Background and aims** To investigate the factors associated with systemic lupus erythematosus (SLE)-related pericardial effusion/ cardiac tamponade and its long-term outcome in Chinese patients.

**Methods** Medical records of 690 SLE patients who admitted in Chang Gung Medical Centre from 2005 to 2012 were reviewed.

**Results** The mean ages at onset and at admission were  $36.3 \pm 16.4$  years and  $40.8 \pm 16.0$  years, respectively. Of the 690 patients, 113 (16.4%) had SLE-related pericardial effusion. Cardiac tamponade developed in 9.7% (11 of 113) patients with pericardial effusion or in 1.5% (11 of 690) of SLE patients. Moreover, 4 of the 11 patients represented with cardiac tamponade as initial presentation of SLE. Cox regression analysis indicated that age at admission  $>50$  years (HR 3.38, 95% CI 2.06–5.55,  $p<0.001$ ), pericarditis (HR 1.70, 95% CI 1.00–2.90,  $p=0.049$ ), pleuritis (HR 2.30, 95% CI 1.43–3.72,  $p=0.001$ ), leukopenia (HR 1.90, 95% CI 1.11–3.24,  $p=0.019$ ), thrombocytopenia (HR 3.28, 95% CI 1.82–5.89,  $p<0.001$ ), and seizure (HR 1.84, 95% CI 1.12–3.00,  $p=0.016$ ) were associated with mortality in SLE. The mortality rate was higher in the pericardial effusion group (30.1%; 34/113) than in the non-pericardial effusion group (11.3%;



Abstract 432 Figure 1