Poster Round

海報目錄

時 間:113年12月14日(星期六)15:15-15:40

地 點:新竹喜來登大飯店 3F 海報區

主持人:許鐘元醫師

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Difference between radiographic and non-radiographic axial Spondyloarthritis in Chinese patients: from Taiwan AS Study Group Cohort

華人僵直性脊椎炎及無放射線學異常之中軸型脊椎關節炎臨床表現比較:來自台灣僵直性脊椎炎研究小組世代

<u>Yu-Sheng Chang</u>¹, Hung-Ta Wu², Hsien-Tzung Liao³, Kuan-Chia Lin⁴, Chun-Hsiung Chen⁵, Horng-An Chen⁶, Chun-Chun Wang⁷, Lin Sheng-Hong^{1,2}, Su Chin-Fang^{1,2}, Kao Jui-Hung^{1,2}, Chung-Tei Chou³

張又升, 吳宏達, 廖顯宗, 林寬佳, 陳俊雄,陳宏安, 王淳峻, 林聖閎, 蘇勤方, 高瑞鴻, 周昌德

- ¹ Division of Allergy, Immunology, and Rheumatology, Department of Internal Medicine, Shuang Ho Hospital, Taipei Medical University
- ² Department of Radiology, Taipei Veterans General Hospital, Taipei, Taiwan, ROC.
- ³ Division of Allergy, Immunology and Rheumatology, Department of Medicine, Taipei Veterans General Hospital,
- ⁴ Community Medicine Research Center, Institute of Hospital and Health Care Administration, National Yang Ming Chiao Tung University, and Cheng Hsin General Hospital, Taipei, Taiwan, ROC.
- ⁵ Division of Allergy, Immunology and Rheumatology, Taipei Tzu Chi Hospital, Buddhist Tzu Chi Medical Foundation, and School of Medicine, Tzu Chi University
- ⁶ School of Medicine, College of Medicine, National Sun Yat-sen University, Kaohsiung, and Chi Mei Medical Center, Taiwan.
- ⁷ Taoyuan General hospital, ministry of health and welfare
- 1衛生福利部雙和醫院(委託臺北醫學大學興建經營)過敏免疫風濕科
- 2臺北榮民總醫院放射線部
- 3臺北榮民總醫院過敏免疫風濕科
- 4振興醫院、陽明交通大學
- 5 台北慈濟醫院、慈濟大學
- 6奇美醫院、中山大學
- 7 衛生福利部桃園醫院過敏免疫風濕科

Background: Axial Spondyloarthritis (axSpA) encompasses both radiographic axSpA (r-axSpA), previously known as ankylosing spondylitis, and non-radiographic axSpA (nr-axSpA). Prior researches have highlighted significant differences between these subgroups, yet data specific to the Chinese population remains limited. This study aims to compare the demographic and clinical characteristics, disease onset, and clinical features of Taiwanese patients with r-axSpA and nr-axSpA.

Methods: In this cross-sectional study, patients with axSpA were recruited from the outpatient clinics of Central Clinic and Hospital, Shuang-Ho Hospital, and Taipei Veterans General Hospital from 2021 to 2024. Demographic information, disease characteristics, and axSpA activity and functionality were recorded and compared.

Results: A total of 492 r-axSpA and 119 nr-axSpA patients were included. The nr-axSpA group had a significantly lower male rate (52.9% vs. 76.8%, p<0.001), older age at onset (31.2 years vs. 26.59 years, p<0.001), and shorter diagnostic delay (2.35 years vs. 3.56 years, p<0.001). Initial symptoms were similar between groups, except nr-axSpA patients reported lower rate of buttock pain but higher rate of fatigue.(Table2) Disease activity indices (BASDAI) were comparable; however, the nr-axSpA group had significantly lower ESR(15.66mm/hr vs. 23.51mm/hr, p<0.001), CRP(11.19 mg/dL vs. 4.17mg/dL, p<0.001), and ASDAS, and better functionality (BASFI 1.92 vs. 2.71, p=0.0014).

Conclusions: In this Taiwanese axSpA cohort, nr-axSpA patients were less male-dominant, presented at an older age, experienced more buttock pain and fatigue, had lower ESR/CRP and ASDAS, and demonstrated better functional outcomes compared to r-axSpA patients.

Association of uveitis with disease manifestations in Chinese patients with spondyloarthritis: from Taiwan AS Study Group Cohort

Chun-Hsiung Chen¹, <u>Yu-Sheng Chang</u>², Hsien-Tzung Liao³, Horng-An Chen⁴, Chin-Fang Su², Chun-Chi Lu⁵, Hung-Ta Hondar Wu³, Kuan-Chia Lin⁶ and Chung-Tei Chou³

<u>陳俊雄</u> ¹ ,張又升 ² ,廖顯宗 ³ ,陳宏安 ⁴ ,蘇勤方 ² ,盧俊吉 ⁵ ,林寬佳 ⁶ ,王淳俊 ⁷ 、吳宏達 ² ,周昌 德 ³

¹Division of Allergy, Immunology and Rheumatology, Taipei Tzu Chi Hospital, Buddhist Tzu Chi Medical Foundation, Taiwan, and School of Medicine, Tzu Chi University, Hualien, Taiwan. ²Department of Internal Medicine, Shuang Ho Hospital, Taipei Medical University. ³National Yang-Ming University, and Taipei Veterans General Hospital, Taiwan. ⁴School of Medicine, College of Medicine, National Sun Yat-sen University, Kaohsiung, and Chi Mei Medical Center, Taiwan. ⁵Tri-Service General Hospital, National Defense Medical Center, Taipei, Taiwan, ⁶National Yang-Ming University, and Cheng Hsin General hospital, ⁷Tao-Yuan General Hospital.

¹ <u>台北慈濟醫院、慈濟大學</u>,² 雙和醫院、台北醫學大學,³ 台北榮民總醫院、陽明交通大學,⁴ 奇美醫院、中山大學,⁵ 三軍總醫院,⁶ 振興醫院、陽明交通大學,⁷ 部立桃園院

Abstract

Background: To investigate whether uveitis has association with disease manifestations, disease activity and co-morbidities in patients with Spondyloarthritis (SpA).

Methods: 615 SPA patients were enrolled in the cross-sectional study. 492 patients with Ankylosing spondylitis, and 123 patients with non-radiographic SpA. Patients visited the Outpatient Department of Veterans General Hospital-Taipei, Shuang Ho Hospital, and Central clinics. Patients completed the questionnaires, containing disease activity (BASDAI), functional ability (BASFI) and patient's global assessment (BASG). Uveitis was accepted only if diagnosed by an ophthalmologist.

Results: 172 patients had uveitis, and 443 patients had non-uveitis. Prevalence of uveitis in this SpA cohort was 28.0% (172/615). Patients with uveitis had higher percentage of age ≥ 45yr than non-uveitis [103(59.88%) vs. 171(38.60%), p<0.001]. Patients with uveitis had older age of definite diagnosis [32.60(13.17) vs. 30.10(13.09) y/o, p=0.0338] and longer disease duration [18.72(13.00) vs. 14.64(11.87) yrs, p=0.0002] than non-uveitis. Patients with uveitis had higher percentage of anterior chest wall pain, neck soreness, neck limitation, enthesitis, peripheral arthritis, dactylitis, inflammatory bowel disease, psoriasis, sleeping, mood disorders than non-uveitis (all p<0.05). Patients with uveitis had higher risk of DM, AMI, and renal disease than with non-uveitis (all p<0.05). Patients with uveitis had higher BASDAI-1 (fatigue), BASDAI-3 (peripheral arthritis), BASDAI-4 (enthesitis), BASFI-3 (reaching a high shelf without help), BASFI-8 (looking over without turning body) than non-uveitis (all p<0.05).

Conclusion: Comparing to non-uveitis, the presence of uveitis in SpA patients was associated with older age, longer disease duration, fatigue, peripheral arthritis, enthesitis, dactylitis, neck limitation, extra-articular manifestations, sleeping/mood disorders and specific co-morbidities.

Septic Arthritis Caused by *Cutibacterium acnes* Masquerading as Ankylosing Spondylitis Flare-up with Peripheral Articular Involvement: A Case Report

吳柏佑 Po-Yu Wu

1. 長庚大學醫學院醫學系 School of Medicine, College of Medicine, Chang Gung University, Taoyuan, Taiwan.

黄冠霖 Guan-Lin Huang

2. 長庚大學醫學院中醫學系 School of Chinese Medicine, College of Medicine, Chang Gung University, Taoyuan, Taiwan.

詹天明 Tien-Ming Chan

3. 林口長庚紀念醫院內科部風濕免疫科 Division of Rheumatology, Allergy and Immunology, Department of Internal Medicine, Chang Gung University and Chang Gung Memorial Hospital, Taoyuan, Taiwan.

Abstract

Background:

As of now, ankylosing spondylitis flare-up still lacks definitive diagnostic criteria. Considering that symptoms specific to AS and fluctuation of inflammatory markers may not always be present in a flare-up, it is sometimes difficult to differentiate between AS and septic arthritis. We report a patient who developed septic arthritis caused by *Cutibacterium acnes*, masquerading as an AS flare-up with peripheral arthritis.

Case Report:

A 23-year-old male, who was diagnosed with AS, presented with right shoulder pain without morning stiffness for 1 week at a clinic, along with fever. Right knee pain and bilateral Achilles tendon pain were noted in prior months. AS flare-up was suspected, though septic arthritis had to be considered. After the patient was admitted to a medical center, blood test revealed high CRP and ESR levels, and empirical ciprofloxacin, 400mg, q12h was given. Arthroscopic debridement and tissue culture were performed, from which grampositive *Cutibacterium acnes* grew. His symptoms and inflammatory markers improved partially after treating with daptomycin, 750mg, qd. Due to fluctuating inflammation, his DMARD regimen in outpatient follow-ups was adjusted. CRP and ESR levels have since returned to baseline, and BASDAI score have improved from 7.3 (upon admission) to 5.3 when discharged. His symptoms have been stable since then.

Conclusion:

When presented with fever, high CRP levels and pain without morning stiffness, septic arthritis should be considered if the symptoms were refractory to high DMARD dosage. Arthrocentesis, microbial cultures and thorough examination of medical history also can assist in differentiating, thus enabling proper treatment.

Image of upper cervical spine osteomyelitis and epidural abscess in a HLA-B 27 positive patient

HLA-B 27 陽性患者之高位頸椎骨髓炎與硬腦膜外膿瘍影像

Yu Chen¹, Chien-Sheng Wu¹

陳遊1、吳建陞1

新北市亞東紀念醫院

Affiliations

¹ Division of Rheumatology, Allergy and Immunology, Far Eastern Memorial Hospital, New Taipei City, Taiwan, R.O.C.

Background

Neck pain is a common symptom in spinal disorders, but unusual neck pain accompanied with neurological deficits can be considered as red flag of infection.

Methods

We reported a case of upper cervical osteomyelitis and epidural abscess in a HLA-B 27 positive female patient through chart review after informed consent.

Results

This 44-year-old woman as a hepatitis B carrier with liver cirrhosis had neck pain and headache for 2 months. Ankylosing spondylitis was suspected and a HLA-B27 test showed positive. However, her neck pain responded poorly to NSAID. Due to severe neck pain, she was brought to ER and was admitted on the next day to rheumatology ward. After admission, gait disturbance, dysarthria, and swallow difficulty were noted. On the second day of admission, involuntary movement was noted and emergent C spine MRI was performed which showed left side condyle and C1 osteomyelitis with epidural abscess formation. Fluoroscopy interventions including sac decompression, drainage of the epidural abscess arranged on the third day of admission. Finally, blood culture and surgical site culture revealed Staphylococcus aureus. Patient recovered well after long-term antibiotics treatment and regular followed up at outpatient department.

Conclusion

Upper cervical spine osteomyelitis and epidural abscess can be earlier awareness by concurrent neurologic deficits. Surgical intervention prevents the progression of neurologic impairment and decrease the risk of subsequent atlantoaxial subluxation.

Clinical Images: C-spine osteomyelitis and epidural abscess



left side condyle and C1 osteomyelitis with epidural abscess formation

Motion and Movement's Influence on Pain 身體的動作和移動對疼痛之影響

Ying-Yang Tsai¹, Shih-Ying Tsai², Lu-Chin Wei³, I-Hsien Lin⁴ 蔡瀛陽、蔡世英、魏呂瑾、林宜嫻

Chei-Shen Clinic, Taipei City 潔生診所(台北市)

Abstract

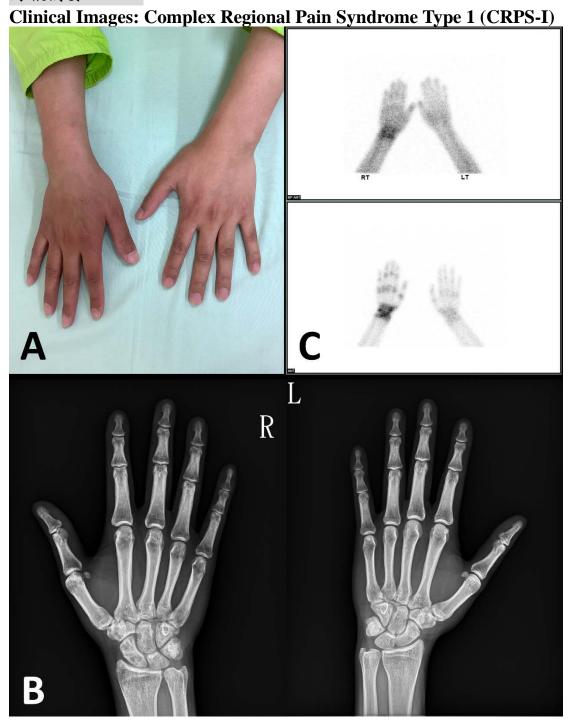
Backgound: 觀察病患的走路以瞭解人體生物力學之改變與肌肉骨骼疼痛之關係。

Method: 給予病患不同物理性刺激,並拍攝病患走路時的影像,再進行比對。

Result: 經長期追蹤,走路應分為著地期 ST(特指雙腳承重)和擺盪期 SW(特指單腳承重)。ST 的影響因素是重力(地心引力),作用在關節間隙,SW 的影響因素是向心力和離心力,即呼吸和軀幹肢體的互動,作用在骨頭之延伸和收復,帶動肌肉之向心與離心收縮。另一橫跨 ST 和 SW 的是交副感神經,使走路重心在後(交感),重心在前(副交感),牽動關節和骨頭。

Conclusion:大腦才是主要的影舞者,大腦的記憶才是治療疼痛的標的。

海報摘要 TCR18



A 41-year-old previously healthy man presents with a 10-month history of persistent pain in his right wrist following a traumatic injury. The patient reports increased pain in the right wrist after physical activity, without morning stiffness. On examination, the right wrist shows mild swelling compared to the left, feels warmer to the touch, and has darker skin coloration (A). Range of motion in the right wrist is limited due to pain. Laboratory results indicate normal inflammatory markers, including C-reactive protein and erythrocyte sedimentation rate. Tests for antinuclear antibody, rheumatoid factor, anti-CCP antibody, and HLA-B27 are all negative. The plain radiograph shows regional osteopenia in the right wrist, while the left side appears normal (B). The three-phase bone scan reveals increased uptake during early blood flow and blood pool phases, as well as in the delayed bone phase of the right wrist (C). These findings support a diagnosis of complex regional pain syndrome type 1 (CRPS-I).

Jheng-Wei Lin, MD 林政緯 National Taiwan University Hospital, Taipei, Taiwan 國立臺灣大學醫學院附設醫院

Dipeptidyl Peptidase-4 Inhibitors Versus Sulfonylureas for Psoriasis Prevention in Patients With Type 2 Diabetes Receiving Metformin

Hsiang-Gyen Lee¹, Wei-Sheng Chen², Yu-Chen Chen¹, Wei-Jui Lin¹, Ke-Yu Chang¹, Ke-Hung Lin¹, Ching-Kuei Chang¹, Yu-Chuan Shen³, Hui-Ching Hsu^{3,5}, Yu-Sheng Chang^{4,5}, Tzu-Min Lin^{1,5}, Chi-Ching Chang^{1,5}
¹Division of Allergy, Immunology and, Rheumatology Taipei Medical University
Hospital

²Division of Allergy, Immunology, and Rheumatology, Department of Internal Medicine, Taipei Veterans General Hospital, National Yang-Ming University, Taipei, Taiwan

³Division of Rheumatology, Immunology and Allergy, Taipei Medical University Wang Fang Hospital

二肽基肽酶-4 抑制劑與磺烯基尿素類降低使用 Metformin 糖尿病患者罹患乾癬之風險性

李向嚴 1 陳瑋昇 2 朱有晨 1 林韋睿 1 張克宇 1 林科宏 1 張晉魁 1 沈佑銓 3 許惠晴 $^{3.5}$ 張又升 $^{4.5}$ 林子閔 $^{1.5}$ 張 棋楨 $^{1.5}$

台北醫學大學附設醫院過敏免疫風濕科1

台北榮民總醫院過敏免疫風濕科2

台北醫學大學萬芳醫院過敏免疫風濕科3

台北醫學大學衛福部雙和醫院過敏免疫風濕科4

台北醫學大學醫學院醫學系內科學科過敏免疫風濕學科5

Abstract

OBJECTIVE: To compare the risk of incident psoriasis in patients with T2DM who initiate DPP-4is or TZDs with the risk in those who initiate sulfonylureas, the most common second-line glucose-lowering therapy, in addition to metformin monotherapy.

RESEARCH DESIGN AND METHOD: This sequential, propensity-score-matched, new-user comparative effectiveness study employed a target trial emulation framework. The study included adults with T2DM receiving metformin monotherapy and utilized data from January 1, 2006, to December 31, 2015, from a general population database with data on Taiwanese citizens. The primary outcome was the incidence of psoriasis, which was determined through diagnoses recorded in urgent care, hospital, and outpatient department records. We employed Cox proportional hazards and Poisson regressions with 1:4 propensity score matching for the target trial emulations. Cox proportional hazards regression models were employed to evaluate the risk factors for psoriasis after adjustment for comorbidities and the use of other medications.

RESULTS: In the 49 810 propensity score-matched adults with T2DM (using DPP4is or sulfonylureas, of whom 27 630 [55.4%] were men, with a mean [standard deviation] age of 57.5 [11.8] years) who were identified on the database, the incidence rate of psoriasis in DPP4i users (1.88 cases per 1000 person-years) was lower than that in sulfonylurea users (4.67 cases per 1000 person-years), with a hazard ratio of 0.422 (95% CI, 0.273-0.716). However, no significant difference was observed for those using TZDs.

CONCLUSIONS: The psoriasis benefits associated with DPP4is can guide selection of glucose-lowering therapy in patients with T2DM who are at risk of psoriasis.

⁴Division of Rheumatology, Im 溪 munology and Allergy, Taipei Medical University Shuang Ho Hospital

⁵Division of Allergy, Immunology and Rheumatology, Department of Internal Medicine, School of Medicine, College of Medicine, Taipei Medical University, Taipei, Taiwan

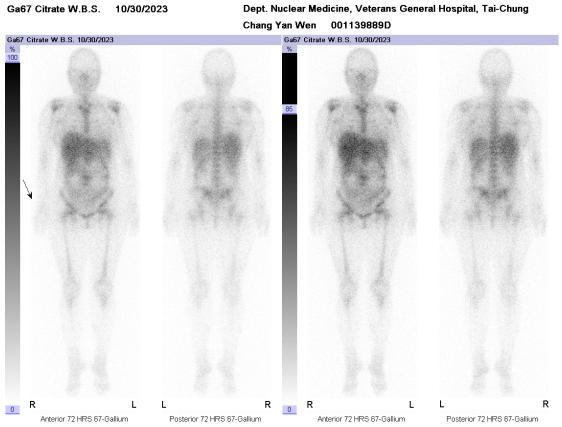
鎵-67 腫瘤掃描與正電子放射斷層造影在一位新診斷 γδT 细胞淋巴瘤乾癬患者中的應用

Gallium-67 and PET Imaging in a Psoriasis Patient Newly Diagnosed with Gamma-Delta T Cell Lymphoma

Yu-Hsien Li ¹, Tsu-Yi Hsieh ¹²³

- ¹ Division of Allergy, Immunology and Rheumatology, Taichung Veterans General Hospital, Taichung, Taiwan
- ² Department of Medical Education, Taichung Veterans General Hospital, Taichung, Taiwan
- ³ Department of Post-Baccalaureate Medicine, College of Medicine, National Chung Hsing University, Taichung, Taiwan

A 38-year-old woman with a 34-year history of psoriasis vulgaris, currently under treatment with brodalumab and exhibiting a good response in the dermatology clinic, presented to the rheumatology clinic with a fever, indurated nodules on her upper back, and erythematous macules on her face, limbs, and trunk. A skin biopsy of the left upper back revealed primary cutaneous gamma-delta T cell lymphoma. Gallium-67 (Ga-67) and positron emission tomography (PET) scans showed uptake in different regions of the body.



Picture 1.

A Ga-67 scan was performed to investigate the cause of an unknown fever. The report indicated inflammation in the bilateral shoulders, sternoclavicular junctions, hips, and sacroiliac joints.



Picture 2.A PET scan conducted one month after the Ga-67 scan suggested malignant lesions (grade 4 increased uptake) in the abdominal and pelvic walls, bilateral back, and left thigh.

Adult-onset Still's disease mimicking long-COVID: a case report 個案報告:一位年輕女性在新冠肺炎感染後發生成人型史迪爾氏症

I-Shu, Chen¹, Li-Heng Huang ², Yu-Heng Lin ², Song-Chou Hsieh¹ 陳亦抒,黃立恒,林昱亨,謝松洲

- 1. Division of Allergy, Immunology and Rheumatology, National Taiwan University Hospital 國立台灣大學醫學院附設醫院風濕免疫過敏科
- 2. Division of Allergy, Immunology and Rheumatology, National Taiwan University Hospital Yun-Lin branch

國立台灣大學醫學院附設醫院雲林分院風濕免疫過敏科

Background

The diagnosis of Adult-onset Still's disease (AOSD) remains challenging due to its protean manifestations and rarity. Emergence of COVID-19 pandemic further complicated the landscape as clinical presentations of AOSD shares several features with the increasingly recognized long-COVID syndrome [1, 2]. The symptomatic overlap may lead to delayed recognition of AOSD. Here we report such a challenging case which AOSD closely resembled long-COVID.

Case presentation

A 32-year-old woman presented to our rheumatology clinic for subacute intermittent fever and joint pain. Four months ago, she had an episode of COVID-19 mild disease. Her discomforts soon resolved after taking NRICM101(清冠一號). Three months ago, she developed daily fever, pain at bilateral hand joints and sore throat. Antibiotics were given for suspected tonsilitis in a local clinic, but response was limited. Subsequently, she was admitted to internal medicine service in hospital A. Laboratory tests revealed elevated C-reactive protein, erythrocyte sedimentation rate, anemia, elevated aspartate aminotransferase, alanine transaminase, alkaline phosphatase, gamma-glutamyl transferase, and positive anti-mitochondrial antibody. Infection workups including blood/sputum/urine and mycobacterium cultures were negative. Abdominal ultrasound showed splenomegaly. Whole-body computed tomography revealed small bilateral inguinal lymph nodes. Tumor markers were unremarkable. Under the impression of primary biliary cholangitis and post-COVID hyperinflammation, methylprednisolone 40mg daily were given with low oral maintenance dose after discharge.

Two months prior to current evaluation, her fever, joint pain and sore throat recurred. She also noticed some faint rashes at her trunk and limbs when her temperature rises. She then came to our hospital for second opinion. Blood sampling revealed persistently elevated liver enzymes, inflammation markers (hsCRP 6.36mg/dL, ESR 42mm/hr), LDH, anemia, leukocytosis, and a markedly elevated ferritin level (4424.18 ng/mL, normal < 204). Splenomegaly also remained on serial abdominal sonography. Admission was arranged. Close observation later confirmed presence of salmon-pink rash. No new evidence of infections or malignancies was identified. She was re-classified as AOSD according to Yamaguchi's criteria. Hydroxychloroquine, etoricoxib and methotrexate (10mg/week) were initiated. A cycle of methylprednisolone mini-pulse therapy was given (Fig. 1). Tocilizumab was considered but not given at last due to quick response to other treatments. She was symptom free now and oral prednisolone was tapered to less than 7.5mg daily.

Discussion and Conclusion

AOSD remained one of the most common rheumatic diagnoses in patients with fever of unknown origin (FUO) [3]. As an autoinflammatory disorder, it is not surprising that the prevalence and incidence were increased in patients with recent COVID-19 infection or vaccination [4]. Among the cardinal manifestations of AOSD, fever, pharyngitis, lymphadenopathy, abnormal liver enzymes and elevated acute phase reactants may also be seen in acute COVID-19 or post-COVID condition. We want to emphasize the importance to exam organomegaly, skin eruptions and degree of ferritin elevation, which helped us to distinguish AOSD in this patient. The inflammatory response in AOSD may trigger severe organ involvement or complications, such as myocarditis or macrophage activation syndrome. Thus, careful evaluation of disease severity is essential

following diagnosis of AOSD. On top of glucocorticoid and conventional DMARDs, Tocilizumab, antiinterleukin-1 therapies and Janus-kinase inhibitors emerged as new biological/targeted-synthetic DMARDs that help us to manage patient with AOSD [5].

5000 **Ferritin** 4424.18 4000 3314.43 3000 2584.16 2000 1153.77 1000 159.10 Discharge Admission 2023/1/31 2023/2/5 2023/1/16 2023/1/21 2023/1/26 1# QD HCQ 200mg 1# BID Etoricoxib 60mg/tab 1# QD MTX 2.5mg/tab 4# QW 10mg 80mg **MTP** Q12H Q8H PSL 5mg/tab 2# BID

Figure 1. The treatment course and serial ferritin levels of the patient

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- 4. Gottschalk, M.N., Heiland, M., Nahles, S. et al. Orphanet J Rare Dis 2023; 18: 50.
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Clinical Images: Persistent pruritic eruptions associated with adult-onset Still's disease



The 44-year-old woman without systemic diseases experienced fever and progressive pruritic skin rash over face and neck (Figure A), four extremities (right upper arm, Figure B) and abdomen (Figure C) for three weeks. She had a similar episode eight months ago and improved after antihistamine and oral steroid use at dermatology outpatient clinic. She also reported sore throat and intermittent hand joint swelling (Left hand at ward in Figure D, Right hand at home in Figure E). Laboratory studies revealed abnormal serum white blood count 11960 /uL, ferritin 10909 ng/mL, CRP 10.48 mg/dL, ESR 65 mm/hr and IgE 57 KU/L. Autoimmune markers including ANA, anti-dsDNA, ANCA, SSA/SSB, RF, ACPA and autoimmune myositis panel were all negative. Bone marrow biopsy reported no neoplasm. Excisional biopsy of right neck lymph node revealed reactive hyperplasia. She received skin biopsy over right forearm and pathologic report showed superficial perivascular infiltrates and dyskeratosis in the upper epidermis, which are compatible with those of persistent pruritic eruption of adult-onset Still disease in appropriate clinical settings. She received oral prednisolone twice daily and Azathioprine once daily after pathology report. Her skin rash, joint swelling and fever subsided, and she is currently well control with Plaquenil twice daily. These images strengthen the importance of skin biopsy in common skin lesion in unique clinical manifestations.

Ping-Hsuan Kuo, MD, Yi-Syuan Sun, MD, Chien-Chih Lai, MD, Ph.D., Yen-Po Tsao, MD, Ph.D., Wei-Sheng Chen, MD, Hsien-Tzung Liao, MD, Ph.D., Ming-Han Chen, MD, Ph.D. Hung-Cheng Tsai, MD Taipei Veterans General Hospital, Taipei, Taiwan.

郭蘋萱, 孫易暄, 賴建志, 曹彥博, 陳瑋昇, 廖顯宗, 陳明翰, 蔡弘正臺北榮民總醫院

Impact of Hydroxychloroquine on Dose-Dependent Reducing Diabetes Mellitus Risk Among Primary Sjögren Syndrome: A Population-Based Study

Wei-Jui Lin¹, Wei-Sheng Chen², Ke- Yu Chang¹, Ke-Hung Lin¹, Hsiang-Gyen Lee¹, Ching-Kuei Chang¹, Yu-Chuan Shen³, Hui-Ching Hsu^{3,5}, Yu-Sheng Chang^{4,5}, Tzu-Min Lin^{1,5}, Chi-Ching Chang^{1,5}

¹Division of Allergy, Immunology and, Rheumatology Taipei Medical University Hospital

- ²Division of Allergy, Immunology, and Rheumatology, Department of Internal Medicine, Taipei Veterans General Hospital, National Yang-Ming University, Taipei, Taiwan
- ³Division of Rheumatology, Immunology and Allergy, Taipei Medical University Wang Fang Hospital
- ⁴Division of Rheumatology, Immunology and Allergy, Taipei Medical University Shuang Ho Hospital
- ⁵Division of Allergy, Immunology and Rheumatology, Department of Internal Medicine, School of Medicine, College of Medicine, Taipei Medical University, Taipei, Taiwan

烴氯奎寧劑量依賴性降低原發性乾燥症患者罹患糖尿病之風險性

林韋睿¹陳瑋昇²張克宇¹林科宏¹李向嚴¹張晉魁¹沈佑銓³許惠晴^{3,5}張又升^{4,5}林子閔^{1,5}張棋楨^{1,5} 台北醫學大學附設醫院過敏免疫風濕科¹

台北榮民總醫院過敏免疫風濕科2

台北醫學大學萬芳醫院過敏免疫風濕科3

台北醫學大學衛福部雙和醫院過敏免疫風濕科4

台北醫學大學醫學院醫學系內科學科過敏免疫風濕學科5

Abstract

Background: HCQ is an antimalarial drug that is widely prescribed for the treatment of Sjögren syndrome. Glucocorticoids may alter glucose metabolism and contribute to diabetes mellitus. Treatment for primary Sjögren syndrome (SS) sometimes requires glucocorticoids that may worsen glucose homoeostasis. HCQ can reduce diabetes risk in SLE and RA. This study aimed to investigate the association of HCQ use and diabetes mellitus risk in primary SS patients.

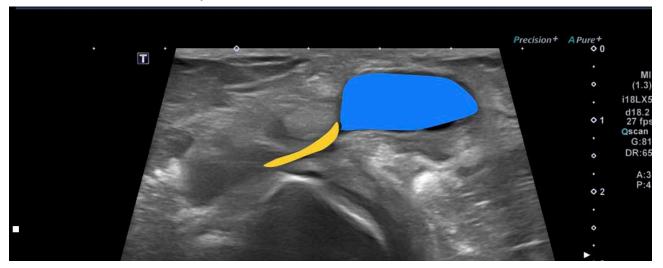
Methods:

This population-based cohort study utilized data from the Taiwan National Health Insurance Research Database, focusing on primary SS patients between 2006 and 2015. We employed multivariate and stratified analyses, using the Kaplan–Meier method and Cox proportional hazards regression, to evaluate the diabetes risk associated with HCQ and corticosteroids individually, as well as the dose-dependent diabetes risk when concurrent HCQ and corticosteroids use.

Results: Five hundred and thirty six newly diagnosed diabetes mellitus patients were identified among primary SS patients (6232 HCQ users and 3116 HCQ non-users), with an average follow-up period of 4.36 years. We observed that compared with those obtained for HCQ nonusers, the aHRs obtained for HCQ users in the $151 \le \text{cDDD} < 350$ and $\text{cDDD} \ge 351$ subgroups were 0.600 (95% CI, 0.454-0.794), and 0.326 (95% CI 0.246-0.433), respectively. Higher corticosteroid (≥ 151 cDDD) was associated with increased risk of developing diabetes mellitus [aHR 1.833(95% CI 1.410-2.383), which was reduced by concomitant HCQ use at a dose > 350 cDDD [aHR 0.632 (95% CI (0.421-0.948), P < 0.01].

Conclusions: This propensity score matched analysis of Taiwanese patients with primary SS found that patients with primary pSS who were administered high doses of HCQ exhibited a significantly reduced risk of developing diabetes mellitus. Conversely, high doses of corticosteroids were associated with a substantial increase in diabetes risk. Importantly, the concurrent use of high doses of HCQ effectively mitigated the diabetes risk associated with high corticosteroid dosages.

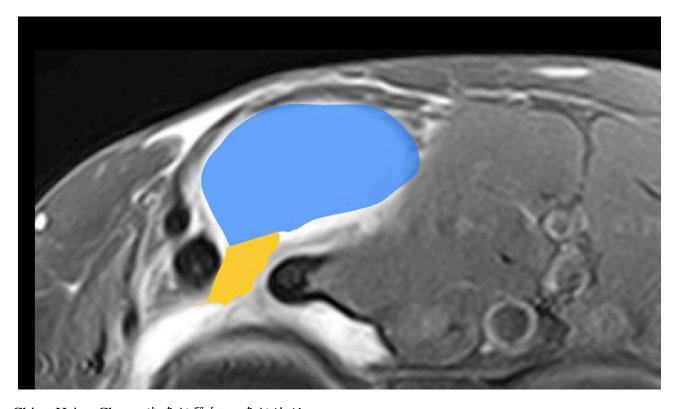
Clinical Images: How to simulate surgical treatment with the ultrasound to define the stalk location of baker cyst



In this context, the blue area represents the main body of the Baker's cyst, while the yellow area indicates the position of the stalk.

Ultrasound plays a crucial role in the localization and guidance of the removal process. The specific steps include:

- 1.Initial Examination: Use ultrasound equipment for an initial examination to determine the location and size of the stalk.
- 2.Localization: Precisely mark the location of the stalk through ultrasound images, aiding the doctor in accurately finding the target during the procedure.
- 3.Surgical Simulation: Before the stalk needling, simulate the removal process using ultrasound images to determine the optimal injection point.



Chien-Hsing Chen, 陳建行醫師。建行診所