Results All 13 patients had underlying diseases, 12 had been treating with immunosuppressive drugs, except one case who had underlying OA. SLE was the most common underlying disease found in our study. Acute and chronic onset monoarthritis occurred in a similar numbers. Chronic long standing joint infection mostly occurred in the patients with partial response to the first line antibiotic treatment. Knee and hip joint were the common sites of infection. Arthroscopic drainage or arthrotomy was performed in all but one patient who was treated by repeated arthrocentesis of the knee joint. All patients received at least 2-week parenteral antibiotic. Three patients were excluded from antibiotic treatment evaluation because of lost follow up after being discharged from the hospital.

Clinical response at the early phase: In 16 events of *Salmonella* septic arthritis, 6 were treated with cotrimoxazole, 7 with quinolone compound and 3 with beta-lactam antibiotics. Dramatic clinical response was found only in the patients who were treated with cotrimoxazole or quinolone compound antibiotics. 2 out of 3 patients who were treated with beta lactam antibiotics partially responded to antibiotics given. After 8 weeks, these 2 patients were retreated with another course of cortimoxazole or quinolone compound.

Long term Clinical outcome: In all patients who were treated with 3 month course of quinolone compound, there was no relapse after 1–5 years follow up. Relapses were more common in cotrimoxazole treatment group (4 out of 6) and usually occurred within 4 months after antibiotic was stopped. Only one case was completely responded to beta-lactam antibiotic treatment without relapse. AVN was the most common complication found in *Salmonella* septic hip, and chronic osteomyelitis was found in a case with septic knee.

Conclusion We suggest to treat *Salmonella* septic arthritis in immunocompromised host with adequate drainage and 3 months of quinolone compound antibiotics. Alternative regimen is parenteral cotrimoxazole in an early phase followed by oral quinolone compound for totally 3 months. Long term treatment with cotrimoxazole frequently resulted in relapse.

SAT0221 DIAGNOSTIC VALUE OF SERUM AND SYNOVIAL PROCALCITONIN (PCT) IN BACTERIAL, CRYSTAL-ASSOCIATED AND RHEUMATOID ARTHRITIS

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Background Procalcitonin (PCT), the calcitonin peptide precursor, is a new specific marker of severe bacterial and fungal infections. PCT is not or slightly elevated in others inflammatory diseases. PCT measurements could help to differentiate bacterial arthritis to other forms of acute arthritis.

Objectives To determine the interest of serum and synovial PCT for the diagnosis of acute bacterial arthritis.

Methods

- A one year prospective study (RR study group): sera and synovial fluid from 32 adults with acute arthritis were immediately frozen (-20°C) before treatment. for all patients, the diagnosis was confirmed by specific synovial analysis.
- PCT (N < 0.5 mg) was measured by a specific and ultrasensitive immunoluminometric yesy (BRAHMS diagnostica, Berlin, Ger.)

Results

Abstract SAT0221 Table 1

	serum PCT ng/ ml	synovial PCT ng/ml	CRP mg/l	ESR mm/h
Bacterial arthritis (n = 10)	0.59 (<0.08–7.91)	0.22 (0–1.31)	37 (61–493)	80 (30–120)
Crystal arthritis (n = 11)	<0.08**(<0.08–0.69)	0.12 (NS) (0.08–0.88)	91*(4–355)	42 (NS) (3–98)
Rheumatoid arthritis (n = 11)	<0.08**(<0.08–0.61)	<0.08–0.6)	50.4** (3.2–90)	50 (NS) (9–130)

Results are expressed in median and range. Values are compared between septic group and crystal or rheumatoid group by Mann-Whitney test: **p < 0.01; *p < 0.05.

Conclusion

- These results show that plasma PCT levels are higher in patients with septic arthritis, but PCT sensitivity remains insufficient. However, in case of bacterial arthritis, PCT in association with CRP, could be of great interest for clinicians, as its increase is in favour of a bacterial origin.
- PCT is not present in all biological fluids. If synovial PCT does not seem to be helpful in clinical diagnosis, this study is, to our knowledge, the first to demonstrate PCT existence in synovial fluid.

SAT0222 TOTAL BODY AND REGIONAL BONE MINERAL AND SOFT-TISSUE COMPOSITION IN HIV DISEASE

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Background Osteopenia and adipose tissue maldistribution are two complications of highly active anti-retroviral therapy (HAART: 1 protease inhibitor + 2 nucleoside reverse transcriptase inhibitors). Dual energy x-rays absorptiometry (DEXA) is a reliable tool to assess total body and regional bone mineral and soft-tissue composition.

Objectives (1) To compare the results of DEXA in HIV-infected patients treated or not with HAART, with or without clinical lipodystrophy and (2) to evaluate DEXA as a tool for early diagnosis of body fat redistribution.

Methods A cross-sectional analysis was performed on 220 men. All subjects gave written informed consent. Controls (C): 106 healthy uninfected volunteers without medical or surgical history, nor immobilisation and therapies which may modify bone mineral density, mean age 38 years. One hundred and fourteen HIV-infected patients at steady-state, without opportunist disease, nor prophylactic drugs? use were divided in 3 groups: (i) 20 naive HIV-patients (N: mean age 38 years, disease duration 43 mths, viral load 3.7 log, CD4⁺ 417/mm³), (ii) 32 HIV-treated patients with no clinical lipodystrophy (n-CLD: mean age 35, disease duration 72 mths, HAART duration 15 mths, viral load 2.3 log, CD4⁺ 524/mm³), and (iii) 62 HIV-treated patients with clinical lipodystrophy (CLD: mean age 42, disease duration 94 mths, HAART duration 26 mths, viral load 1.8 log, CD4⁺ 483/mm³). Total body and regional soft-tissue composition were measured with DEXA (DPX-L, Lunar Corp. Acquisition and analysis software 4.6). The following parameters were studied: