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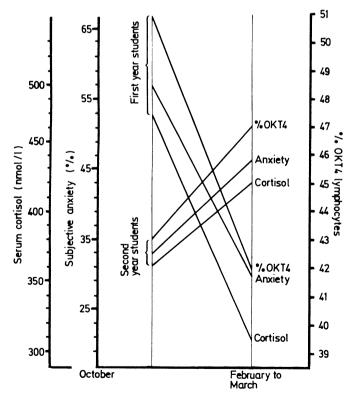
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# Stress, cortisol concentrations, and lymphocyte subpopulations

The aetiology of rheumatoid arthritis is almost certainly multifactorial, so that established genetic factors and suspected infective agents do not preclude the possibility of psychological factors also influencing the onset of disease.1 This influence may be mediated by stress modifying the responses of the immune system.<sup>2</sup> Before starting further studies of people with disease we therefore decided to investigate the responses of T lymphocyte populations in normal students under stress.

#### Subjects, methods, and results

We invited medical students on the second day of their clinical course in a medical school new to them to participate in the study. All 35 with surnames beginning A to G were approached: 33 accepted, attending during their first week (October 1983). Twenty eight second year students served as



Subjective anxiety scores, serum cortisol concentrations, and proportions of OKT4 lymphocytes in first and second year students.

Conversion: SI to traditional units—Cortisol: 1 nmol/1 ≈ 0.036 μg/100 ml.

controls. Thirty one of the first year students and 22 of the second year students were tested again about four months later (in February and March 1984). Each student assessed his or her subjective anxiety on that day on an uncalibrated 10 cm visual analogue scale, the extremes of which were marked "The day of my life I felt least/most anxious or tense." The score was recorded as zero (least) to 100 (most). Personal and family histories of atopic disorder were asked for and no difference was found between first and second year students.

All students were seen between 11 am and 1 pm to avoid the effects of diurnal variation. Total leucocyte and lymphocyte counts were determined using a Coulter S Plus counter. Parallel heparinised blood samples were separated on discontinuous Ficoll-metrizoate (Triosil) gradients, incubated with fluoresceinated monoclonal anti-T cell antibodies (Ortho-Mune OKT series) without the addition of any fetal calf serum, and analysed on the cytofluorograph (FACS 420; Becton Dickinson).3 Scatter gates set on the 90° light scatter profile were used to minimise contamination of cell preparations with non-lymphocytic cells. The heparinised plasma was frozen and used to measure serum cortisol concentrations.

At the first testing nine first year students and one second year student did not have lymphocyte subpopulation estimations done, but their anxiety and cortisol measurements were included in the calculation of those means

The Mann-Whitney test, two tailed, was used to compare the first with the second year students at the first and second visits, while Wilcoxon's matched pairs test, two tailed, was used to compare the first with the second visit of both groups of students.

The figure illustrates the differences in means among the four sets of results (though the means were not used in the statistics). The new students showed significantly higher subjective anxiety scores and cortisol concentrations than the second year students and than themselves at the second visit (all p values < 0.005). This finding was associated with a significantly raised proportion of OKT4 cells when compared with the second year students at the same time (p=0.0001) and with themselves a few months later (p<0.005). The second year students showed increases in anxiety, cortisol values, and proportions of OKT4 cells at their second visit, probably because they were approaching an examination, but the increases did not reach significance. There were no significant changes in white cell counts, lymphocyte counts, or proportions of OKT8 cells.

### Comment

Students during their first week experienced a high level of psychological and physiological arousal, shown by raised subjective anxiety scores and serum cortisol concentrations. These features were associated with a significant increase in the proportion of OKT4 cells (helper T lymphocytes).

Not all psychological states produce similar responses: for example, Kronfol and House, commenting on our preliminary results,4 reported that they had found no difference in the percentage of lymphocyte subsets when comparing depressed people with controls.5

Probably our finding is only part of a far more complicated immune response to stress. In many diseases (well represented by rheumatoid arthritis) the influence of stress and other emotional factors on the immune system may prove to be of fundamental importance in both susceptibility to disease and subsequent fluctuations during the course of disease.

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