A comparative study of the pathergy reaction among Turkish and British patients with Behçet's disease

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SUMMARY Forty-eight Turkish and 12 British patients along with 24 Turkish and 7 British healthy controls were assessed for the presence of the pathergy phenomenon. A photographic method with 'blind' reading was utilised. The pathergy phenomenon was present only among the Turkish patients.

The hyperreactivity of the skin to a needle prick (the pathergy test) is a curious phenomenon almost unique to Behçet's disease or syndrome (BS).¹ We have previously demonstrated its sensitivity and specificity,¹ the inter- and intraobserver variations,² and diagnostic usefulness^{3,4} in patients with BS from Turkey. In Japan, although formal figures are lacking, the pathergy phenomenon is reported to be present in the majority of the patients.⁵ However, other students of this disease doubted its existence.^{6,7} We believe the lack of positivity of this phenomenon among the British and American patients has been the main cause of this scepticism. We therefore compared the prevalence of the pathergy test in 2 groups of patients and in parallel controls from Istanbul, Turkey, and Leeds, England, in a blind study by means of photography.

Patients and methods

Forty-eight patients from Istanbul and 12 from Leeds were studied. Twenty-four apparently healthy Turkish and 7 British hospital staff served as controls. All Turkish and 11 British patients fulfilled O'Duffy's criteria⁸ for diagnosis. One British patient had aphthae and genital ulceration only. All Turkish and British patients were Caucasians except for one Pakistani in the British group. The pathergy test was done as previously described.¹ Needles of the same kind (21 G) and manufacturer were used in both countries. 1X1 colour slides of the puncture sites

were taken at 48 hours. To obviate differences in background due to light effects observed on some slides black paper was used to cover all parts of the slides except the needle site and its immediate vicinity, before the blind reading. Two observers, unaware of the hypothesis being tested, interpreted the slides on a 0–3+ scale (0 = only a needle mark, 1+ = papule only, 2+ = a 'small' pustule, 3+ = a 'large' pustule) in a blind fashion in a single session. For statistical analysis only a 2+ or 3+ result was considered typically positive in that pustule formation was present. Either observer interpreted each slide twice. The inter- and intraobserver errors (only positive–negative variability being considered) were calculated as previously described.²

Results

Inter- and intraobserver errors of the method were 22% and 15.5% respectively. Observer 1 read all controls as 0. This was also the case for observer 2 except in 2 instances, where she read a 1+ for a Turkish and a British control on her second reading. Turkish probands had significantly fewer 0 readings on all 4 occasions, whereas this was not observed among the British (Table 1). Among the British probands 2 had a 1+ score once and 1 had a 1+ score twice among the 4 occasions his/her reaction was interpreted. Nine British patients had a 0+ reading throughout. Among the Turkish probands there were again 9 patients whose readings were all 0+ (9/48 vs. 9/12, $\chi^2 = 11.91$, $p < 0.001$). The remaining 39 Turkish patients in 156 (39 × 4) observations had 0+ =

Table 1 *Pathergy readings*

	n*	0 score on all four readings	2 + or 3 + score in at least one reading
Turkish			
Probands	48	9	28
Controls	24	23	
		$\chi^2 = 5.73, p < 0.02$	
British			
Probands	12	9	0
Controls	7	6	0
		$\chi^2 = 0.92, p > 0.05$	

*Numbers refer to individuals tested.

28; 1+ = 72; 2+ = 41; and 3+ reading 15 times. If one considered a 2+ or 3+ reading at least once in 4 observations as a positive pathergy test, then 28/48 (58%) of the Turkish probands would have had a positive pathergy test, while all British patients and controls from either country would have had negative pathergy reactions.

Discussion

The photographic method of reading the pathergy reaction increases the variability² and decreases the sensitivity^{1,3} of the test. We believe this is mainly due to lessening of depth appreciation on a slide; thus the interpretation of papules becomes difficult. Nevertheless, it served its purpose of bringing out the differences between the 4 groups of subjects.

The absence of the pathergy phenomenon among the British patients is another major point of difference between the Turkish and British patients with BS, in addition to the already established different HLA association.⁹ We had previously demonstrated that HLA B5 and pathergy positivity did not go together among the Turkish patients.³ Thus the absence of a positive pathergy test among the British patients cannot readily be explained by different HLA association.

It is to be noted that there are other regional differences in clinical and laboratory findings in BS. Japanese and Turks both share the high prevalence of HLA B51 but the HLA DR5 association is unique to the Japanese patients.^{9,10} Moreover involvement of the colon is common among Japanese but rare among Turkish and British patients.^{5,11,12} These findings justify the use of the term syndrome rather than disease

in describing the entity originally described by Behçet.¹³ Furthermore the absence of the pathergy reaction among British patients would suggest that it does not have an important role in the pathogenesis of this entity.

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References

- 1 Tüzün Y, Yazıcı H, Pazarlı H, Yalçın B, Yurdakul S, Müftüoğlu A. The usefulness of the nonspecific skin hyperreactivity (the pathergy test) in Behçet's disease in Turkey. *Acta Derm Venereol (Stockh)* 1979; **59**: 77-9.
- 2 Altaç M, Tüzün Y, Yurdakul S, Binyıldız P, Yazıcı H. The validity of the pathergy test (non-specific skin hyperreactivity) in Behçet's disease: a double blind study by independent observers. *Acta Derm Venereol (Stockh)* 1982; **62**: 158-9.
- 3 Yazıcı H, Tüzün Y, Pazarlı H, Yalçın B, Yurdakul S, Müftüoğlu A. The combined use of HLA-B5 and pathergy test as diagnostic markers of Behçet's disease in Turkey. *J Rheumatol* 1980; **7**: 206-10.
- 4 Özyazgan Y, Pazarlı H, Yazıcı H, Yurdakul S, Tüzün Y, Altaç M, Ülkü B and Müftüoğlu A. Usefulness of HLA determination and pathergy test in the diagnosis of uveitis in Turkey. *Br J Ophthalmol* 1981; **65**: 798-9.
- 5 Shimizu T. Clinicopathological studies on Behçet's disease. In: Dilşen N, Koniçe M, Övül C, eds. *Behçet's disease*. Amsterdam: Excerpta Medica, 1979: 467: 9-43.
- 6 O'Duffy JD. Summary of international symposium on Behçet's disease. *J Rheumatol* 1978; **5**: 229-33.
- 7 Lehner T, Barnes CG. Criteria for diagnosis and classification of Behçet's syndrome. In: Lehner T, Barnes CG, eds. *Behçet's syndrome* London: Academic Press, 1979: 1-7.
- 8 O'Duffy JD. Suggested criteria for diagnosis of Behçet's disease. *Book of abstracts, VI Pan American congress of rheumatic disease*. Toronto: 1974: 18.
- 9 Yazıcı H, Chamberlain M A, Schreuder I, D'Amato J, Müftüoğlu A. HLA antigens in Behçet's disease: a reappraisal by a comparative study of Turkish and British patients. *Ann Rheum Dis* 1980; **39**: 344-8.
- 10 Ohno S, Ohguchi M, Matsuda H, Wakisaka A, Aizawa M. Close association of HLA-Bw51, MT2 and Behçet's disease. International conference on Behçet's disease, Tokyo, Japan, October 1981.
- 11 Dilşen N, Koniçe M, Övül C. Clinical evaluation of 106 cases of Behçet's disease. In: Dilşen N, Koniçe M, Övül C, eds. *Behçet's disease*. Amsterdam: Excerpta Medica, 1979: 467: 124-9.
- 12 Sladen G E, Lehner T. Gastro-intestinal disorders in Behçet's syndrome and a comparison with recurrent oral ulcers. In: Lehner T, Barnes C G, eds. *Behçet's syndrome* London: Academic Press, 1979: 151-8.
- 13 Behçet H. Über residivierende aphtose, durchein Virus verursachte Geschwüre am Mund, am Auge und an den Genitalien. *Dermatol Monatsschr* 1937; **105**: 1152-7.