Results

The study encompassed 20 patients presenting by severe asthma, with mean age of 9.9±2.4; range 7-14 years and 10 controls with mean age of 9.4±2.8; range 6.5-14 years. There were 12 male patients and 8 females; and 7 of controls were males and the other 3 were males, (Table 1). There was a non-significant difference between patients and controls as regards age, (Fig. 1) or sex, (Fig. 2) presentation.

Table (1): Demographic data of study participants

	Control	Patients
Number	10	20
	9.4±2.8	9.9±2.4
Age (years)	7:3	6:4
Sex; M:F		

Fig. (1): Mean age of patients and controls included in the study

Control
Patients

7
6
5
4
3
2
1

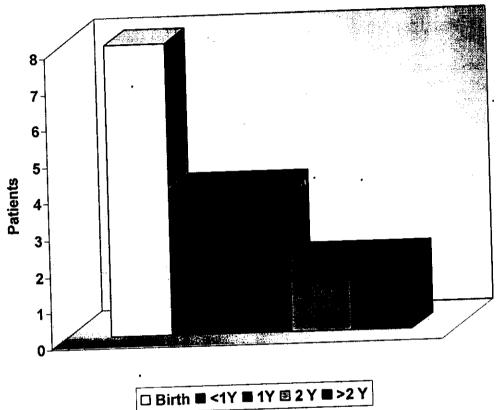
Age of onset of disease

In 8 patients (40%) disease was started to present since birth, in 4 patients (20%) disease started to symptomatize at age younger than one year, in other 4 patients at age of one year and in 2 patients at age of 2 years; while in the other 2 patients disease was presented at age of 4 years, (Table 2, Fig. 3).

Table (2): Patients' distribution according to age at onset of symptoms

Number	%
8	40
4	20
4	20
2	10
2	10
	Number

Fig. (3): Patients distribution according to age of onset of symptoms



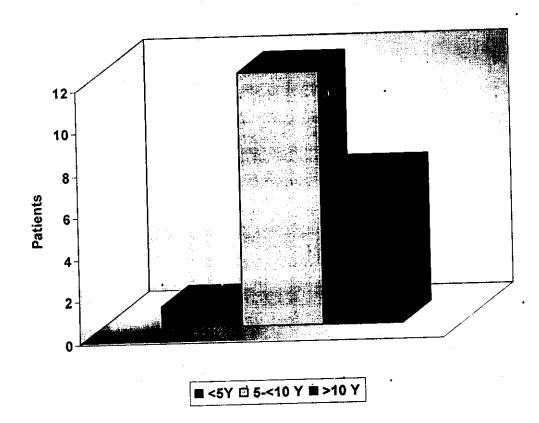
Duration of disease

Only one patient (5%) had disease duration of 3 years; while 12 patients (60%) had mean disease duration of 7.7±0.8; range: 6.3-9 years and the other 7 patients (35%) had diseased since a mean duration of 12.3±1.5; range: 10-14 years, (Table 3, Fig. 4).

Table (3): Patients' distribution according to duration of disease

Number (%)	Duration (years)	
1 (5%)	3	
12 (60%)	7.7±0.8	
7 (35%)	12.3±1.5	
	1 (5%) 12 (60%)	

Fig. (4): Patients distribution according to duration of disease



Routine laboratory Investigations

Hemoglobin concentration:

There was a significant (P<0.05) decrease of hemoglobin concentration in patients (11.2±1.2; range: 9.2-13.8 gm) compared to control levels, (13±1; range: 11.5-14 gm), (Table 4, Fig. 5).

Erythrocyte sedimentation rate (ESR)

ESR of patients (12±3.7; range: 7-19mm) showed significant (P<0.05) increase compared to that reported in controls (8±2.06; range: 6-12mm), (Table 3, Fig. 4).

o Total leucocytic count (TLC)

There was significant (P<0.05) increase of TLC in patients (7358±2100; range 3700-12,000 cell/cc) compared to count detected in controls, (5980±986; range 4500-7500 cell/cc), (Table 3, Fig. 4).

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Table (2): Results of routine laboratory investigations

Group	Control	Patients '
Parameter	(n=10)	(n=20)
Hemoglobin concentration (gm)	13±1	11.2±1.2*
Helilogiobili oorlooridadaa (3)	(11.5-14)	(9.2-13.8)
ECD (mm)	8±2.06	12±3.7*
ESR (mm)	(6-12)	(7-19)
TLC (cell/cc)	5980±986	7358±2100*
120 (Genroo)	(4500-7500)	(3700-12000)

^{*:} significant versus control

Fig. (5): Mean hemoglobin concentration of patients compared to control levels

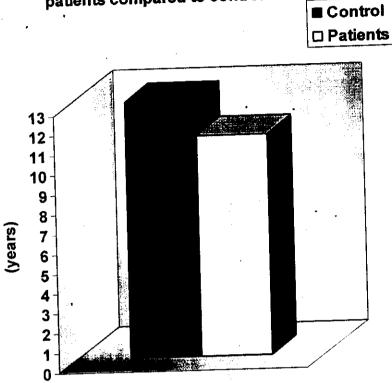


Fig. (6): Mean ESR in patients compared to control levels



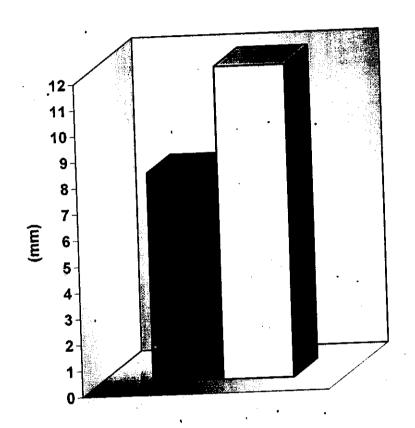
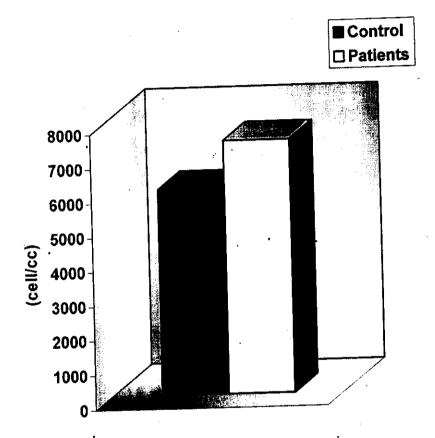


Fig. (7): Mean TLC in patients compared to control levels



AEC Eisinophy

Detection of Antigen presenting cells

Number of patients according to frequency of detection of APG

AEC cell/ce: In control group; no APC could be detected in 4 controls (40%), one had frequency of APC of 50 cell/cc, and the other 5 controls had >50-<100 cell/cc. On contrary, in patients, 15 patients had APC in frequency of >100-500 cells/cc; while another 5 patients had >500 APC cells/cc. There was a significant increase of frequency of detection of APC in patients compared to controls, (X2=9.15, P<0.001), (Table 5, Fig. 8).

o Erythrocyte sedimentation rate (ESR) ESR of patients (12±3.7; range: 7-19mm) showed significant (P<0.05) increase compared to that reported in controls (8±2.06; range: 6-12mm), /Table 3, Fig. 4).

Table (2): Results of routine laboratory investigations

		Control	Patients
		(n=10)	(n=20)
Number of patients	10	4	0
according to the	0-50	1	0
!	>50-100	5	0
frequency of detection of AFC	>100-500	0	15
cell/cc	>500	0	5
Mean±SD (range) (o	ell/cc)	53±42 (0-100)	367±203* (120-820)

^{*:} significant versus control

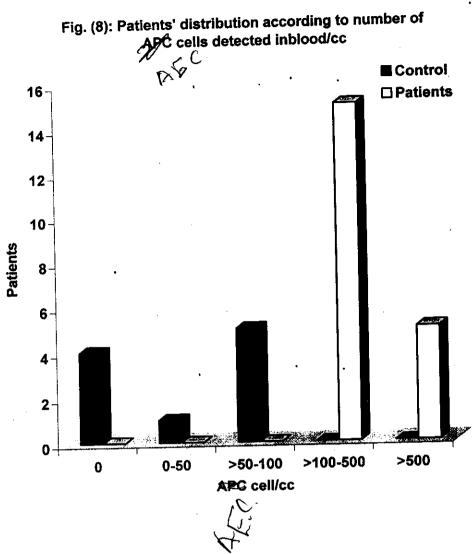
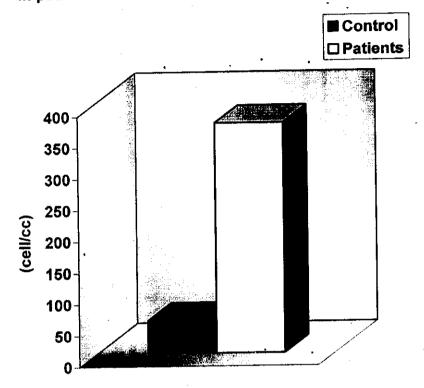


Fig. (9): Mean number of ACC (cell/cc)detected in patients and controls included in the study



Results of Flow cytometry

Results obtained by flow cytometry were presented in table (5)

Percentage of CD4:

- The percentage of CD4 in pre-treatment samples (45.2±3; range: 39.8-50.2%) showed a non-significant (P>0.05) decrease compared to control levels, (46.8±3.1; range: 42-51.1%).
- Furthermore, post-treatment samples showed a non-significant (P<0.05) decrease of percentage of CD4 (44.6±3.4; range: 39.7-51%) compared both to control and pre-treatment levels, (Fig. 10).

Percentage of T-helper cells

1. T-helper (Th₀) cell

- a. There was a significant (P<0.05) decrease of percentage of Th₀ in patients both pre-treatment sample (4.35±1.65; range: 1.9-6.6%) and post-treatment samples (5.02±1.26; range: 3-7.1%) compared to control levels, (6±1.1; range: 4.1-8%).
- b. There was a significant (P<0.05) decrease of the percentage of Th₀ in pre-treatment samples compared to post-treatment samples, (Fig. 11).

2. T-helper (Th₁) cell

- a. There was a significant (P<0.05) decrease of percentage of Th₁ in patients both pre-treatment sample (12.53±2.9; range: 8.2-18.5%) and post-treatment samples (12.9±2.4; range: 9.3-17.7%) compared to control levels, (21.2±2.2; range: 17.9-25.2%).
- b. There was a non-significant (P>0.05) difference between percentage of Th₁ between pre-treatment and post-treatment samples, (Fig. 12).

3. T-helper (Th₂) cell

- a. There was a significant (P<0.05) increase of percentage of Th₂ in pre-treatment sample (20.4±2; range: 16.9-23.6%) compared to both control, (17.4±1.3; range: 15.9-20.2%) and post-treatment levels (17.9±2.17; range: 14.2-21.3%).
- b. There was a non-significant (P>0.05) difference between percentage of Th₂ cell in post-treatment samples compared to control samples, (Fig. 13).

The T1/T2 ratio

- There was a significant (P<0.05) decrease of Th₁/Th₂ in patients samples; pre-treatment (0.62±0.14; range: 0.39-0.81%) and post-treatment (0.73±0.13; range: 0.47-0.95%) compared to control levels, (1.21±0.05; range: 1.1-1.3%).
- Moreover, there was a significant (P<0.05) increase of the Th₁/Th₂ ratio in post-treatment samples compared to pretreatment samples, (Fig. 14).

Table (5): Results of Flow cytometry

Group	' Control	Patients (n=20)	
Parameter	(n=10)	Before treatment	After treatment
CD4 (%)	46.8±3.1	45.2±3	44.6±3.4
	(42-51.1)	(39.8-50.2)	(39.7-51)
Th _o (%)	6±1.1	4.35±1.65*	5.02±1.26*†
	(4.1-8)	(1.9-6.6)	(3-7.1)
Th ₁ (%)	21.2±2.2	12.53±2.9*	12.9±2.4*
·	(17.9-25.2) 🗸	(8.2-18.5)	(9.3-17.7)
Th ₂ (%)	17.4±1.3	20.4±2*	17.9±2.17†
	(15.9-20.2)	(16.9-23.6)	(14.2-21.3)
Th₁/Th₂	, 1.21±0.05	0.62±0.14*	0.73±0.13*† ₂
	(1.1-1.3)	(0.39-0.81)	(0.47-0.95)

^{*:} significant versus control
†: significant versus before treatment

Fig. (10): Mean percentages of CD4 cells as detected by flow Cytometry in patients (pre- & post-treatment) compared to control levels

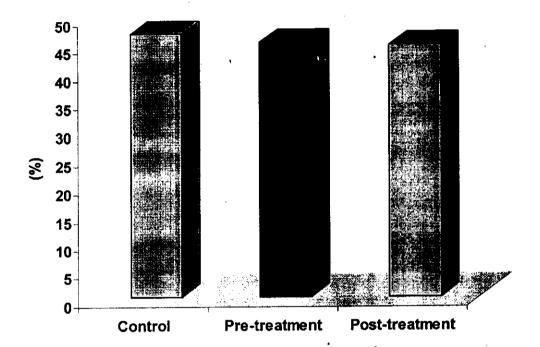


Fig. (11): Mean percentages of Th0 cells as detected by Flow Cytometry in patients (pre- & post-treatment) compared to control levels

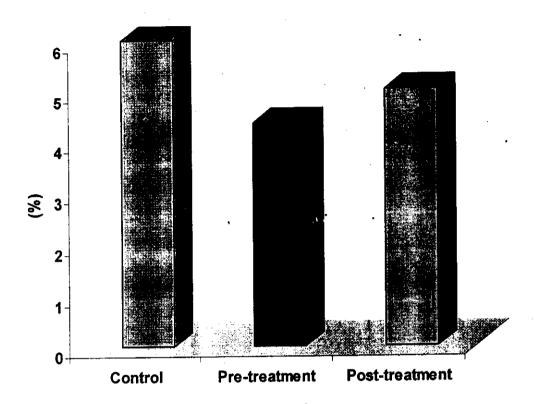


Fig. (12): Mean percentages of Th1 cells as detected by flowcytometry in patients (pre- & post-treatment) compared to control levels

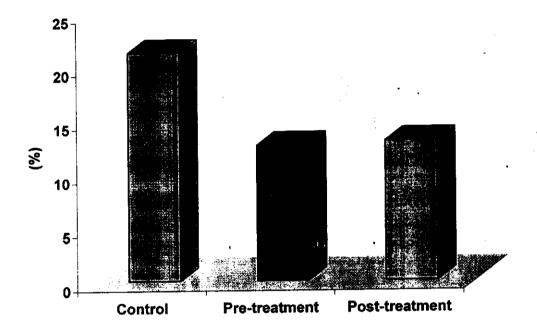


Fig. (13): Mean percentages of Th2 cells as detected by flowcytometry in patients (pre- & post-treatment) compared to control levels

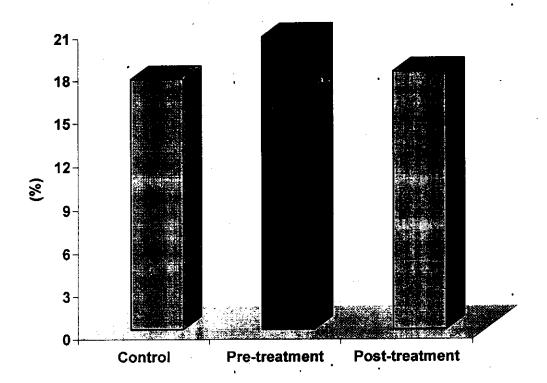


Fig. (14): Mean Th1/Th2 ratio as detected by flowcytometry in patients (pre- & post-treatment) compared to control levels

