

“CLINICAL DIAGNOSIS OF HYPERSENSITIVITY PNEUMONITIS”

The probability of hypersensitivity pneumonitis ranged from 0% in patients with none of these features to 98% in patients with all six

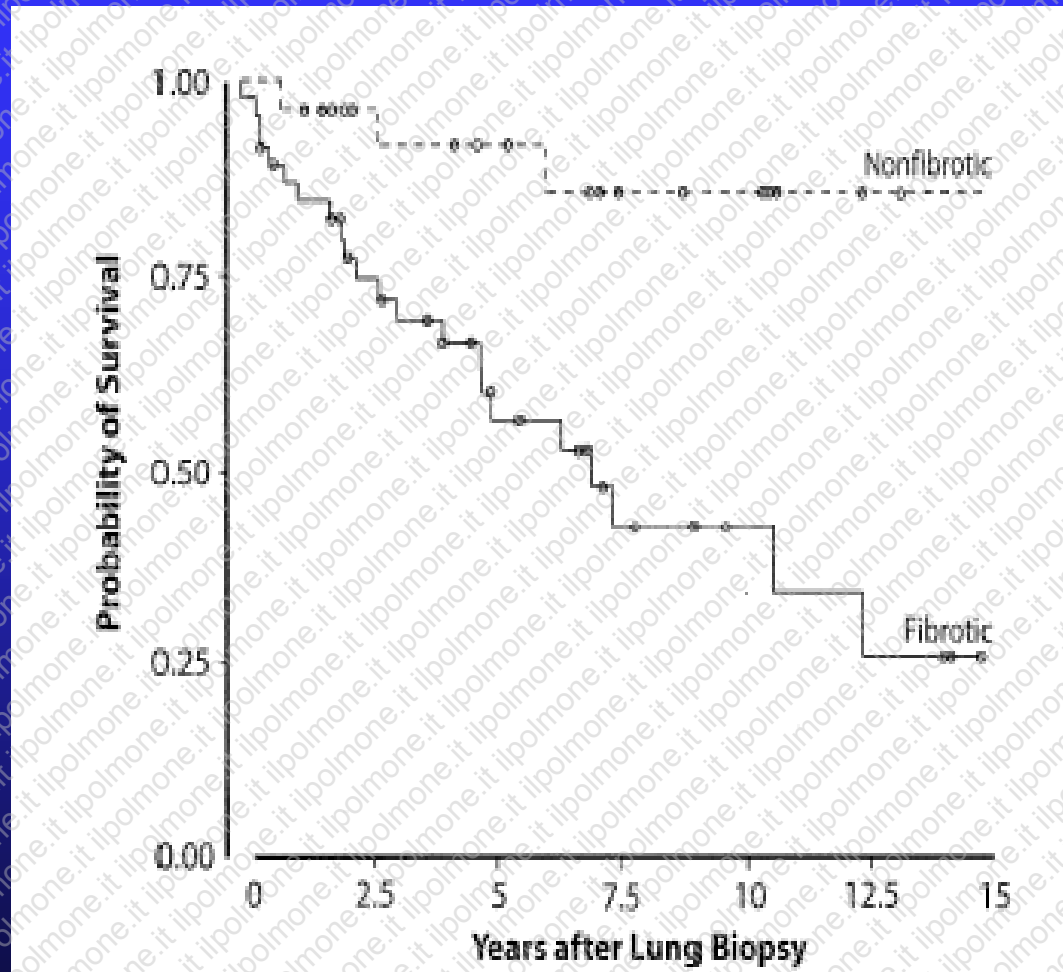
Lacasse Y et al. *Am J Respir Crit Care Med* 2003; 168:952

DIFFERENTIAL DIAGNOSIS

- Viral or bacterial pneumonia
- UIP and NSIP (fibrotic pattern)
- Collagen vascular disease with pulmonary involvement
- Organic dust toxic syndrome (or pulmonary mycotoxicosis)
- Eosinophilic pneumonia
- Allergic bronchopulmonary aspergillosis

The effect of pulmonary fibrosis on survival in patients with hypersensitivity pneumonitis

Vourlekis JS et al. AJM 2004;116:662



Conclusion: Fibrosis was the best predictor of diminished survival of the variables tested, including symptom duration, antigen type, smoking status, and pulmonary function

Prognosis

Authors	N°	UIP Like	Fibrotic NSIP like	Cellular NSIP like	Granu lomas	Giant Cells
Vourlekis J et al AJM 2002;112:490	6	1 (17%)	3 (50%)	2 (33%)	0	1
Ohtani Y et al Thorax 2005;60:665	24	11 (46%)	8 (33%)	5 (21%)	5	19
Churg A et al AJSP 2006;30:201	13	9 (69%)	4 (31%)	-	7	11
Total	43	21 49%	15 35%	7 16%	12 28%	31 72%

Prognosis

43 Patients

21 UIP-like

15 fib. NSIP-like

7 cell. NSIP-like

Follow up in 34 Patients

12 (35%) Died

7 (20%) Stable

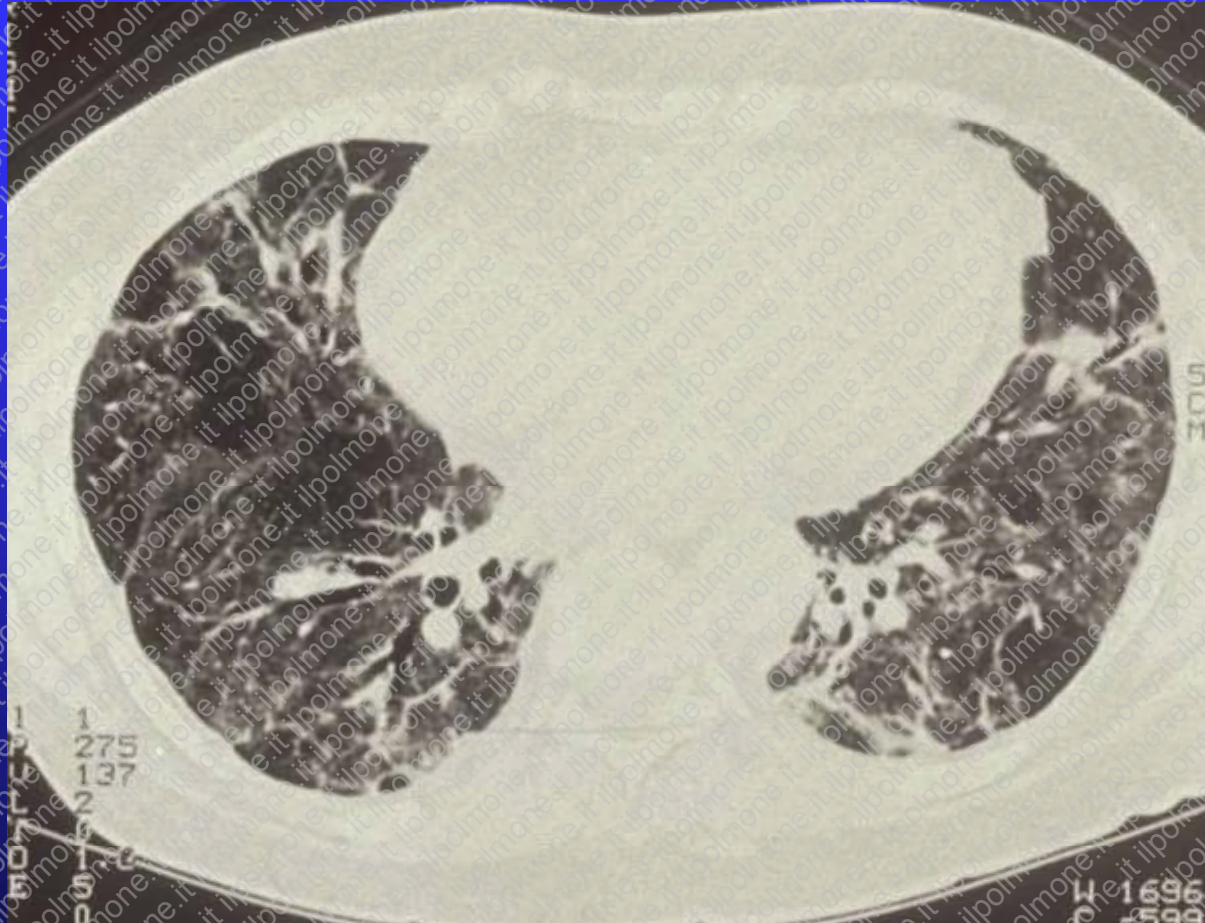
15 (45%) Improved

3.3 years

Vourlekis J et al. AJM 2002;112:490
Ohtani Y et al. Thorax 2005;60:665
Churg A et al. AJSP 2006;30:201

Long-term Risk of Emphysema in Patients with Farmer's Lung and Matched Control Farmers

Pekkanen R et al. Am J Respir Crit Care Med 1998;158:662



Conclusion: the largest difference between the study groups was observed in emphysema. An average of 14 yr after the diagnosis of FL, the HRCT examination revealed that 23% of the FL patients had emphysema compared with only 7% of the control farmers. Quite surprisingly, no significant difference was noted in the prevalence of fibrosis

TREATMENT

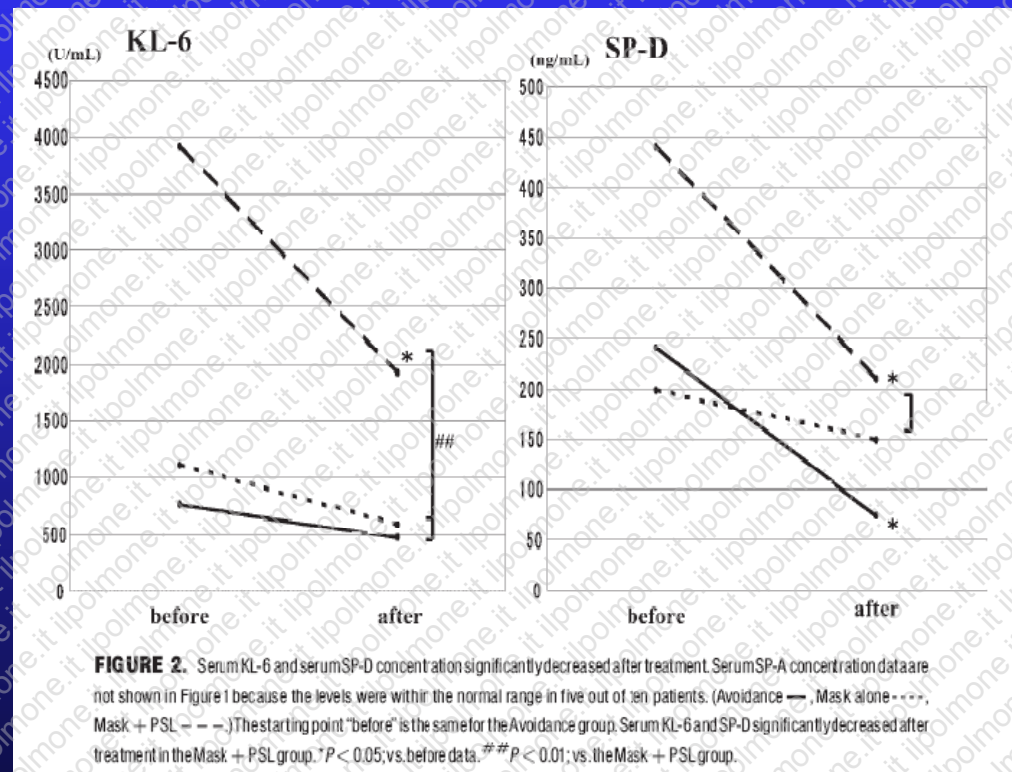
- Antigen exposure avoidance
- Corticosteroids
 - Prednisone 0.5-1 mg/kg/day followed by a gradual reduction until a maintenance dose of 10 to 15 mg/day ?
 - Inhaled steroids?
- Antifibrotic drugs for chronic cases?
- Variable prognosis

Therapeutic effects for hypersensitivity pneumonitis induced by Japanese mushroom (*Bunashimeji*)

Tsushima K et al. *Am J Ind Med.* 2006;49:826

CONCLUSIONS:

Complete cessation was the best treatment for hypersensitivity pneumonitis. The use of a mask was ineffective for patients with a high serum KL-6 and SP-D concentration and severe ground-glass opacity on chest HRCT. Initial treatment with steroids is recommended for these patients with high levels of total cell counts in BAL fluid





ALTERNATIVE TREATMENT ?

Carlsen KH et al. Allergic alveolitis in a 12-year-old boy: treatment with budesonide nebulizing solution. *Pediatr Pulmonol* 1992;12:257

Blanchet MR et al. Inhibitory effect of nicotine on experimental hypersensitivity pneumonitis in vivo and in vitro. *Am J Respir Crit Care Med* 2004;169:903.

Ye Q et al. Thalidomide reduces IL-18, IL-8 and TNF-alpha release from alveolar macrophages in interstitial lung disease. *Eur Respir J* 2006;28:824

Long-term outcome of pulmonary function in farmer's lung: a 14 year follow-up with matched controls

Pekkanen R et al. Eur Respir J 1997;10:2046

Table 2. – Spirometry and pulmonary transfer factor in farmer's lung (FL) patients and control farmers

		FL patients (n=89)	Control farmers (n=84)	p-value*
VC	L	3.45±0.86	3.48±0.84	0.53
	% pred	102±14	100±14	
FEV ₁	L	2.46±0.64	2.55±0.67	0.60
	% pred	93±14	94±15	
FEV ₁ /VC	%	72±7	73±7	0.12
	% pred	91±8	94±9	
MEF ₅₀	L·s ⁻¹	2.71±1.16	3.06±1.25	0.08
	% pred	70±26	77±27	
TL _{co}	mmol·min ⁻¹ ·kPa ⁻¹	7.46±1.98	8.61±1.89	<0.001
	% pred†	98±19	110±16	

Table 3. – Spirometry and pulmonary transfer factor in farmer's lung (FL) patients with a single episode of the disease and in patients with one or more recurrences

		Single episode	Recurrent episode(s)	p-value*
Patients	n	53	36	
Sex	M/F	15/38	5/31	
Age	yrs	60±9	60±8	
Nonsmokers	n	43	30	
Smokers†	n	10	6	
VC	L	3.57±0.94	3.27±0.70	0.67
	% pred	102±13	102±16	
FEV ₁	L	2.54±0.73	2.35±0.48	0.75
	% pred	92±14	94±15	
MEF ₅₀	L·s ⁻¹	2.73±1.32	2.69±0.90	0.31
	% pred	68±28	72±24	
TL _{co} ‡	mmol·min ⁻¹ ·kPa ⁻¹	7.95±2.01	6.71±1.70	0.02
	% pred	102±18	90±18	

- Conclusion:** impairment of the pulmonary transfer factor is the most important long-term consequence of farmer's lung. After a single episode the difference was of the order of 10% and after two or more episodes it had increased to about 20%

Table 3 Clinical characteristics and histological pattern in patients with chronic bird fancier's lung (BFL)

	Group A: BOOP-like or cellular NSIP-like lesions (N=7)	Group B: fibrotic NSIP-like lesions (N=8)	Group C: UIP-like lesions (N=11)
Age (years)	56.9 (4.3)	58.4 (1.8)	64.7 (2.1)
Cases of recurrent acute episodes (%)	85.7 $\pm$	50.0 $\S$	0.0 $\pm$ $\S$
Exertional dyspnoea (%)	85.7	100.0	90.9
Duration of symptoms before surgical lung biopsy (months)	19.3 (7.5)	46.3 (14.2)	24.2 (5.3)
Exposure periods (years)	11.6 (2.8)	18.0 (2.8)	11.0 (2.5)
Finger clubbing (%)	0.0 $\pm$	50.0	81.8 $\pm$
Anti-PDE or BDE antibodies (%)	85.7 $\dagger$	62.5	18.2 $\dagger$
Antigen induced lymphocyte proliferation (%)	100.0	87.5	90.9
VC (% pred)	80.2 (9.1)	60.6 (5.3)	74.8 (7.2)
Tlco (% pred)	58.7 (5.8)	49.1 (6.0)	52.3 (5.7)
Micronodules on HRCT (%)	57.1 $\dagger$	25.0	0.0 $\dagger$
Traction bronchiectasis on HRCT (%)	28.6** $\pm$	100.0**	100.0 $\pm$
Honeycombing on HRCT (%)	0.0 $\pm$	50.0	90.9 $\pm$
BAI lymphocytes (%)	77.0 (2.6)** $\pm$	40.8 (8.6)** $\S$	19.1 (2.9) $\pm$ $\S$
Favourable response to treatment (%)*	7/7 (100.0%)* $\pm$	1/7 (14.3%)* $\pm$	1/9 (11.1%) $\pm$
No response to treatment (%)*	0/7 (0.0%) $\dagger$ $\dagger$	5/7 (71.4%) $\dagger$	6/9 (66.7%) $\dagger$
Alive/dead	7/0	4/4	5/6