

Σοβαρό άσθμα και Αλλεργική Βρογχοπνευμονική Μυκητίαση

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Πνευμονολόγος, Επιμελητής Β' ΕΣΥ 6^η Πνευμονολογική Κλινική ΓΝΝΘΑ «Η ΣΩΤΗΡΙΑ»

09:00-10:30 Στρογγύλη Τράπεzα

Το σοβαρό Βρογχικό Άσθμα ως συννοσηρότητα Προεδρείο: Ε. Ζέρβας - Π. Μπακάκος

- Σοβαρό άσθμα και αππεργική βρογχοπνευμονική μυκητίαση Κ. Σάμιταs
- Ηωσινοφιλική κοκκιωμάτωση με πολυαγγειτιδα (EGPA) και σοβαρό άσθμα
 Ε. Φούκα
- Άσθμα ως συννοσηρότητα στη ΧΑΠ Γ. Χειθάς
- Άσθμα και ρινικοί ποιθύποδες Π. Μαραγκουδάκης

Conflict of interest (Δήλωση σύγκρουσης συμφερόντων)

- Honorarium (τιμητικές αμοιβές) ως προσκεκλημένος ομιλητής από τις φαρμακευτικές εταιρείες Novartis, Elpen, Bristol, Medi-Globe, AstraZeneca, Boehringer Ingelheim, Chiesi
- Ερευνητικές επιχορηγήσεις (unrestricted research grants) από τις φαρμακευτικές εταιρείες GSK, Novartis
- Κάλυψη εξόδων συμμετοχής σε ελληνικά και διεθνή συνέδρια από τις φαρμακευτικές εταιρείες Menarini, Novartis, Elpen, GSK, Demo, Pharmathen, AstraZeneca

Σοβαρό άσθμα και Αλλεργική Βρογχοπνευμονική Μυκητίαση

- **✓** ABPA: Introduction
- ✓ Aspergillus-induced Asthma (AIA) & Severe Asthma with Fungal Sensitization (SAFS)
- **✓ ABPA: Pathogenesis**
- **✓ ABPA: Clinical features**
- **✓ ABPA: Diagnosis**
- **✓** ABPA: Treatment
- **✓ ABPA: Conclusions**

BRONCHO-PULMONARY ASPERGILLOSIS* A REVIEW AND A REPORT OF EIGHT NEW CASES

K. F. W. HINSON, A. J. MOON, AND N. S. PLUMMER From the London Chest Hospital

(RECEIVED FOR PUBLICATION APRIL 4, 1952)

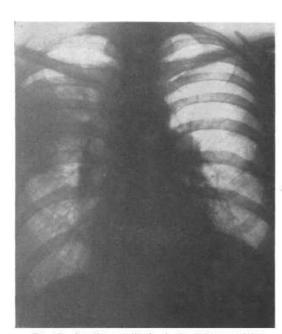


Fig. 13.—Case 6: consolidation in the right upper lobe (20.10.47).

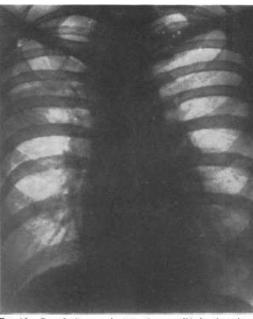


Fig. 14.—Case 6: three weeks later the consolidation has almost resolved (12.11.47).

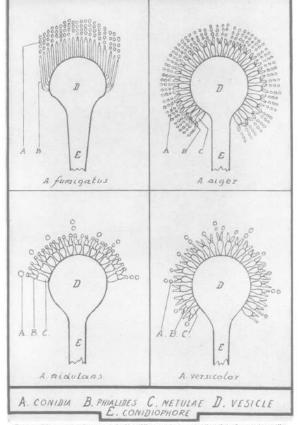


Fig. 1.—Diagram showing morphological differences between strains of the fungus Aspergillus.



Fig. 2 .- A sputum plug from Case 7 separated for display-

We are indebted to Dr. E. H. Hudson and Dr. J. Smart for allowing us to study Cases 5 and 7, and to Dr. E. N. Davey and Dr. Bertram H. Jones and Dr. F. J. D. Knights for providing clinical details in Cases 4 and 8 relating to pathological specimens studied by us.

We are grateful for help in various ways from Dr. L. J. Grant, Dr. I. M. Hughes, Dr. R. Hastings James, Dr. G. Phipps, and Mr. Vernon C. Thompson, and to Miss M. Davis for her patient secretarial

Messrs. May and Baker kindly supplied the drug M & B 938, and we would like to thank Dr. W. R. Thrower for his help in the therapeutic trials.

Remounces

Abbott, J. D., Festando, H. V. J., Guilleg, K., and Maude, B. W. (1972). 20th and J., 1, 223.

Artis, C. J., and Black, F. (1990). The str. path Sec. Lines., 47, 8. Secretal, J. Hagher (1982). Duran nay. Sec. Lines., 12, 277.

Secretal, J. Hagher (1982). Duran nay. Sec. Lines., 18, 277.

Secretal, J. (1982). A sec. lines of the street o

Holden, G. W. (1975). Trans. Assoc. clim. clim. Ass., 21, 97. Jacobson, H. P. (1972). Pargous Observa, p. 270. London

Rigge, G. F. (1816). Drive. Arck. Physiol. (Mirchel), 1, 254. Cited by Réson (1897). Kampseler, R. H., and Black, H. A. (1934). Journ Ser. Tubers, 36.

Harfman, E. (1922). Lebehark der apsziellen pathologischen Am-rosen, Eth. ed., vol. 1. Berlin. Cloud by Manthen (1930). Kabenger, K. (1920). Deach word. Weeber, 44, 1130. Karbensmater, F. (1933). Die in aud au dem Kileper den Februaliu. Kinkenmenter, F. (1873). The is and as down Mayor des relevands a Members with controlled Pastavier. Lecture 1, 1988. As a control of the Indiana. No. 6, 1978. J. Aser Son. 14, 428. Lathaux, M. 6, 1978. J. Aser seed As. 8, 27, 1034. Leavann, W. Stoten van, Bien, Z., Kermer, W. and Varekamp, H. (1925). Z. Leavandowski, M. Lee, M. 1988. Leavann, M. Stoten van, Bien, Z., Kermer, W. and Varekamp, H. (1925). Exhauster Controlled Controlle

(1962). "See qualquet master d'errest dans la déstruction (1962). Ministy, 8. (1961). "See qualquet est l'homene." Tobre Fac. Ned. 1960 (1962). "A service de la propulet de l'homene." Tobre Fac. Ned. 1960 (1962). "A service de la propulet de la p

Handers State (1851). M. Anat. Fired. Parel Anim. Cited by Reines Robin. C. (1851). Missone meta-risk det nighteur perceiter. Paris. Cited by Reines (1871). Robrigaer Villagus, R., and Schren. A. T. (1941). Sim mid., R. Ann. C. F. (1951). Paris. Robrigaer Villagus, R., and Schren. A. T. (1941). Sim mid., R. Ann. C. F. (1951). Paris. Rev., 43, 409.
Ron. C. F. (1951). Paris. Rev., 43, 409.
Ron. C. F. (1951). Paris. Rev., 44, 409.
Robrigaer Villagus, R. (1952). Section of the Circle, p. 155. Paris. Schrenzel, R. (1952). Section of Control, Schreder, J. (1952). Ann. J. Revegend, 37, 337.
Septer, R. R. and Meta-rebete, F. V. (1952). Ann. J. Revegend, 37, 337.
Schreder, L. V. (1988). Ann. Rev. Therm., 12, 362.
Shaper, F. T. (1952). "De vegetabilities arguments antivially state—lang Review, p. 14. (1964) of Honor Histories down Dan. Dan. Control, p. 14. (1964) of Honor Histories down Dan. Dan. Control, p. 14. (1964) of Honor Histories down Dan. Dan. Control, p. 14. (1964). A Handes of the Atjumphi. Technol. Control, p. 14. (1964). A Handes of the Atjumphi. Technol. Control, p. 14. (1964). A Handes and Car., 16, 314.
Tetrag, J. W., and Kericky, P. (1951). In A Tambook of News Dangard, ed. Stanks, S. C. (1964). Not Control, ed. Stanks, S. C. (1964). Not Control, ed. Stanks, S. C. (1964). Indian and Car., 16, 312.
Vineausthas, R. (1964). Indian and Car., 16, 312.
Weingerton, R. J. (1963). Lancer, E. (190.).

Henbringson, J. H. (1943). Amer. Rev. Tabers., 43, 107. Hensinger, C. F. (1826). Smither ron der königlichen zummeinden Artialt zu Witchung 1824-25. Werzburg, Chied by Rimon (1897). M. B. (1981). Lorent. 1 (10)
 Morgarien, A. and Leine, S. (1981). Therefore, Lond. 22, 98.
 Wassen, S. W. (1898). Treat park Soc. Lond. 41, 34.
 Wassen, S. W. (1898). Treat park Soc. Lond. 41, 34.
 Young, F. H. (1983). Lonent, 1, 198.
 Young, R. A., and Bearment. G. E. (1994). A Terriposk of the Parather of Medium, ed. Phys. F. W. (8), 44, p. 1278. London.

List of Aspergillus species

From Wikipedia, the free encyclopedia

List of fungus species in the genus Aspergillus.[1]

Species [edit]

This list is incomplete; you can help by expanding it.

The genus Aspergillus includes several hundred fungus species, including:

Contents

A.B.C.D.E.F.G.H.I.J.K.L.M.N.O.P.Q.R.S.T.U.V.W.X.Y.Z

A [edit]

Aspergillus acidus ^[2]	Aspergillus alliaceus		
Aspergillus aculeatinus[2]	Aspergillus amazonicus 2		
Aspergillus aculeatus	Aspergillus ambiguus ^[2]		
Aspergillus aeneus[2]	Aspergillus amoenus ^[2]		
Aspergillus affinis ^[2]	Aspergillus amstelodami ^[2]		
Aspergillus alabamensis[2]	Aspergillus amyloliquefaciens [2]		
Aspergillus albertensis	Aspergillus amylovorus[2]		
Market Million Control (Control (Co.)	Aspergillus anomalus 2		

Top · A · B · C · D · E ·	F - G - H - I - J - K - L - M - N -	. O . P . Q . R . S . T . U	J · V · W · X · Y · Z
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Academillus harthallative[4]

B [edit]

Aspergillus baarnensis ^[4]	Aspergillus pertriolledus		
Aspergillus baeticus ^[2]	Aspergillus biplanus ^[2]		
Aspergillus bahamensis[2]	Aspergillus bisporus[2]		
risperginal barratrerios	Aspergillus bombycis 2		

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C [edit]

Aspergillus caatingaensis ^[4]	Aspergillus caninus
Aspergillus caelatus ^[2]	Aspergillus capensis[4]
Aspergillus caesiellus	Aspergillus capensis[2]
Aspergillus caespitosus	Aspergillus capsici ^[4]
Aspergillus calidoustus[2]	Aspergillus carbonarius[2]
Asperallius californicus[4]	Aspergillus carneus

Aspergillus anthodesmis[2]	Aspergillus asperescens[2]
Aspergillus angustatus ^[3]	Aspergillus assulatus[4]
Aspergillus apicalis[2]	Aspergillus astellatus[4]
Aspergillus appendiculatus[2]	Aspergillus aurantiobrunneus[2]
Aspergillus arachidicola[2]	Aspergillus aurantiopurpureus[3]
Aspergillus arenarius[2]	Aspergillus aureolatus ^[2]
Aspergillus arvil(2)	Aspergillus aureoterreus[2]
Aspergillus askiburgiensis ^[3]	Aspergillus aureus ^[2]

Aspergillus botswanensis	Aspergillus brevipes[2]
Aspergillus brasiliensis ^[2]	Aspergillus brevistipitatus[2]
Aspergillus brevistipitatus[4]	Aspergillus bridgeri ^[2]
Aspergillus brevijanus	Aspergillus brunneo-uniseriatus

Aspergillus cavernicola[2]	Aspergillus clavatus
Aspergillus cervinus[2]	Aspergillus conicus[2]
Aspergillus chevalier ⁽²⁾	Aspergillus conjunctus[2]
Aspergillus chinensis ^[4]	Aspergillus conversis[2]
Aspergillus chungil ^[2]	Aspergillus coreanus ^[2]
Aspergillus cibarius[2]	Aspergillus coremiiformis[2]

Aspergillus auricomus^[2]
Aspergillus australensis^[4]
Aspergillus austroafricanus^[2]
Aspergillus avenaceus^[2]
Aspergillus awamor^[2]

Aspergillus brunneus
Aspergillus brunneoviolaceus^[2]

Aspergillus cretensis^[2]
Aspergillus cristatus^[2]
Aspergillus crustosus^[2]
Aspergillus crystallinus^[2]
Aspergillus cumulatus^{**}
Aspergillus cvjetkovicii^[2]

Aspergillus fumigatus

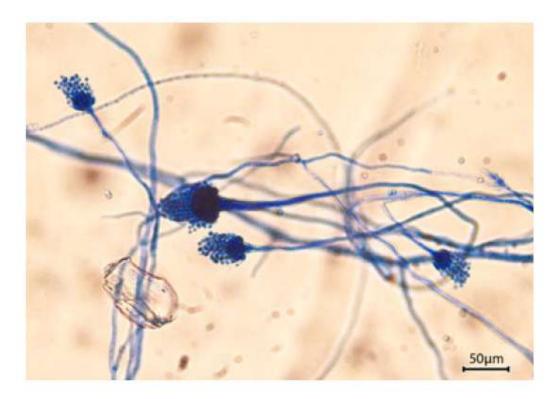


Fig. 1. Photomicrograph of Aspergillus fumigatus under lactophenol cotton blue mount $(100\times)$.

- Spore forming fungi
- Thermophilic, survival at temperatures up to 70°C
- Soil, compost, garbage collection, water damaged structures, damp basements, barns, sewage treatment facilities
- The spores are dispersed by wind in the atmosphere
- Inhalation is unavoidable
- Size of spores: 3-5μm (reach lower airways)

Aspergillus associated respiratory disorders

Table 1. Aspergillus-associated respiratory disorders^{1,2}

- Upper respiratory tract
- 1. Allergic aspergillosis
 - Allergic Aspergillus sinusitis (AAS)
- 2. Saprophytic colonisation
 - Sinus fungal balls
- 3. Invasive disease
 - Acute fulminant invasive sinusitis
 - Chronic invasive sinusitis
 - Granulomatous invasive sinusitis
- II. Lower respiratory tract
- 1. Allergic aspergillosis
 - (IgE mediated) Aspergillus induced asthma (AIA)
 - Allergic bronchopulmonary aspergillosis (ABPA)
 - Hypersensitvity pneumonitis
- 2. Saprophytic colonisation
 - Aspergilloma simple complex (chronic cavitary pulmonary aspergillosis)
- 3. Invasive disease
 - Invasive pulmonary aspergillosis
 acute
 subacute (chronic necrotising pulmonary aspergillosis)

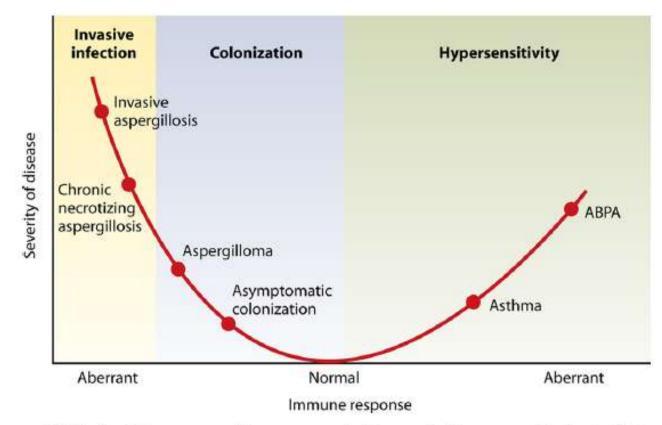


FIG. 1. Diagrammatic representation of diseases attributed to Aspergillus species as a function of the host's immune response. ABPA, allergic bronchopulmonary aspergillosis.

Aspergillus Induced Asthma-AIA

- √ Asthma + hypersensitivity to Aspergillus (Aspergillus Sensitization-AS)
- ✓ SPTs or elevated s.IgE levels
- ✓ 16% to 38% in different geographical regions -pooled prevalence of 25%
- ✓ Exclusion of ABPA

Aspergillus Induced Asthma-AIA

Variables	Group A $(n = 26)$	Group B $(n = 26)$	Group C $(n = 49)$	Group D $(n = 22)$	Group E $(n = 8)$	p Value
TLC, cells/μL Range	5,900-9,900	4,400–12,600	3,500-13,100	6,500-13,000	5,800-19,000	E vs B, < 0.005; E vs C, < 0.05 E vs D, < 0.0168
Mean ± SD AEC, cells/μL	$7,300 \pm 1,400$	$7,870 \pm 1,948$	$8,154 \pm 1,951$	$8,500 \pm 1,400$	$1,000 \pm 4,100$	E vs B, E vs C, E vs D, < 0.05;
Range Mean ± SD	80-300 250 ± 61	190-2,210 1,500 ± 690	110-3,720 $2,599 \pm 895$	240–1,900 11,300 ± 450	800-2,300 $1,400 \pm 595$	D vs B, D vs C, < 0.0001
PFT Disease, No. (%)			300		E vs BCD, < 0.05
Mild	,	18 (69.2%)	28 (57.1%)	11 (50%)	3 (38%)	
Moderate		6 (23%)	12 (24.4%)	8 (36.3%)	1 (12.5%)	
Severe FEV ₁ , L		2 (7.69%)	9 (18.3%)	3 (13.7%)	4 (50%)	
Range	2.8-4.4	1.12-4.0	0.56-4.3	1.4-4.6	0.89-4.6	
Mean ± SD	3.6 ± 0.54	2.46 ± 0.73	2.4 ± 0.8	2.6 ± 0.97	2.1 ± 0.87	
FVC, L	22.40	214 446	1.40.40	22.50	24.4	
Range	3.3-4.9	2.14-4.46	1.49-4.9	2.2–5.0	2.4-4.4	
Mean ± SD PEFR, L/s	4.2 ± 0.57	3.23 ± 0.7	3.2 ± 0.8	3.5 ± 0.89	3.2 ± 0.72	
Range	7.6-12	2.61-10.4	1.58-12.12	3.5-10	1.4-8.4	
Mean ± SD	11 ± 1.6	6.19 ± 2.1	5.8 ± 2.4	6.2 ± 1.8	5.3 ± 2.6	
Serum total IgE, IU/mL						E vs B, E vs C, E vs D, < 0.05; D vs B, D vs C, < 0.05
Range	17.27-155.50	17.27-2,195	17.27-2,057		1,676.62-2,489.11	11 - 18 11 18
Mean ± SD	72.55 ± 51.38	$1,063.56 \pm 585.63$	$1,052 \pm 580.30$	$1,532 \pm 432.56$	$1,987.78 \pm 319.42$	

^{*}PFT, pulmonary function test; PEFR = peak expiratory flow rate.

105 asthmatic patients

- B: asthmatic subjects, SPTs-
- C: asthmatic subjects , SPTs +, Aspergillus Ag –
- D: SPTs +, Aspergillus Ag +
- E: ABPA (Aspergillus Ag +)

105 patients with bronchial asthma:

28.5% (30) sensitized to Aspergillus antigens = more severe form of asthma

- ➤ Higher mean duration of illness (p,0.001),
- ➤ Higher mean eosinophil count (p,0.0001),
- ➤ Higher mean total IgE (p,0.05)
- More usage of oral corticosteroids per year (p,0.004).
- > increased incidence of bronchiectasis

Sensitization to moulds and asthma severity

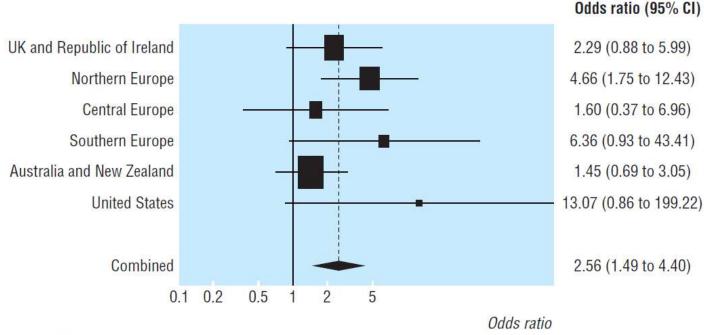


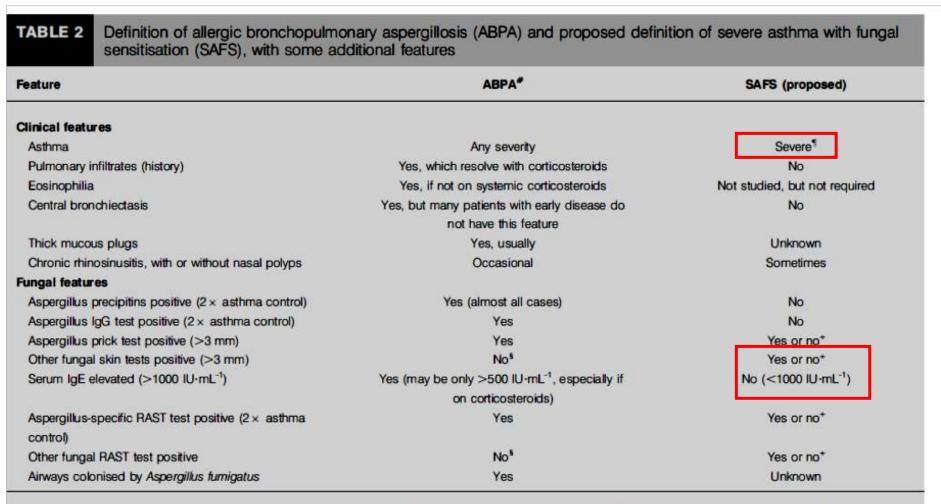
Fig 2 Multivariable adjusted odds ratios (95% confidence interval) for association of severe versus mild asthma with sensitisation to moulds (either *Alternaria alternata* or *Cladosporium herbarum*, or both) by region (adjusted within region for age, sex, smoking habits, passive smoking, and parental history of asthma) with combined odds ratio from model with region included as random effect

Sensitisation to airborne moulds and severity of asthma: cross sectional study from European Community respiratory health survey

Mahmoud Zureik, Catherine Neukirch, Bénédicte Leynaert, Renata Liard, Jean Bousquet, Françoise Neukirch, on behalf of the European Community Respiratory Health Survey

- ✓ The frequency of sensitisation to moulds (Alternaria alternata or Cladosporium herbarum or both) increased significantly with increasing asthma severity
- ✓ odds ratio 2.56 for severe vs mild asthma

Severe asthma with fungal sensitization (SAFS) – Diagnostic Criteria



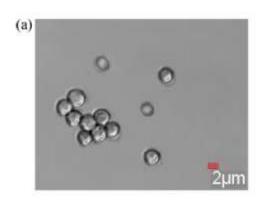
Ig: immunoglobulin; RAST: radioallergosorbent test. *: as defined by Ricketti et al. [126] and PATTERSON et al. [127]; *: typically British Thoracic Society level 4 or equivalent; *: at least one fungal skin or RAST test positive (better and more specific tests may emerge in the future); *: there are rare instances of bronchopulmonary mycosis due to other fungi, with typical clinical features.

- 1) severe (poorly controlled) asthma
- 2) a positive skin-prick test result for fungi or antifungal IgE>0.4 kU/L(not necessarily to Aspergillus species)
- 3) a total IgE <1000 kU/L, no bronchiectasis, no mucous plugging (exclusion of ABPA)

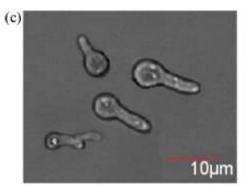
Allergic Bronchopulmonary Aspergillosis (ABPA)- Pathogenesis

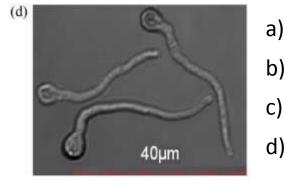
Allergic Bronchopulmonary Aspergillosis (ABPA)- Pathogenesis

- Predominantly affects patients with asthma and cystic fibrosis
- airway colonisation in susceptible hosts that elicits an allergic response.
 - Mainly type I (IgE-mediated hypersensitivity)
 - Tissue invasion does not occur

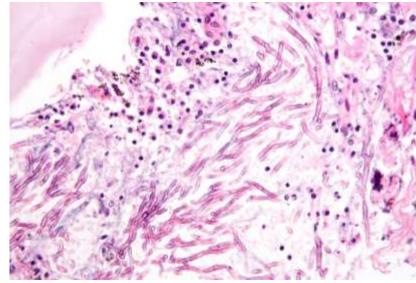








- a) Resting conidia
- b) Swelling of conidia
 - Germination
 - Hyphae formation



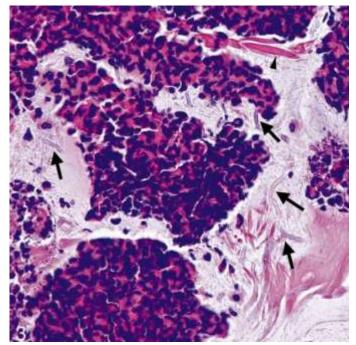
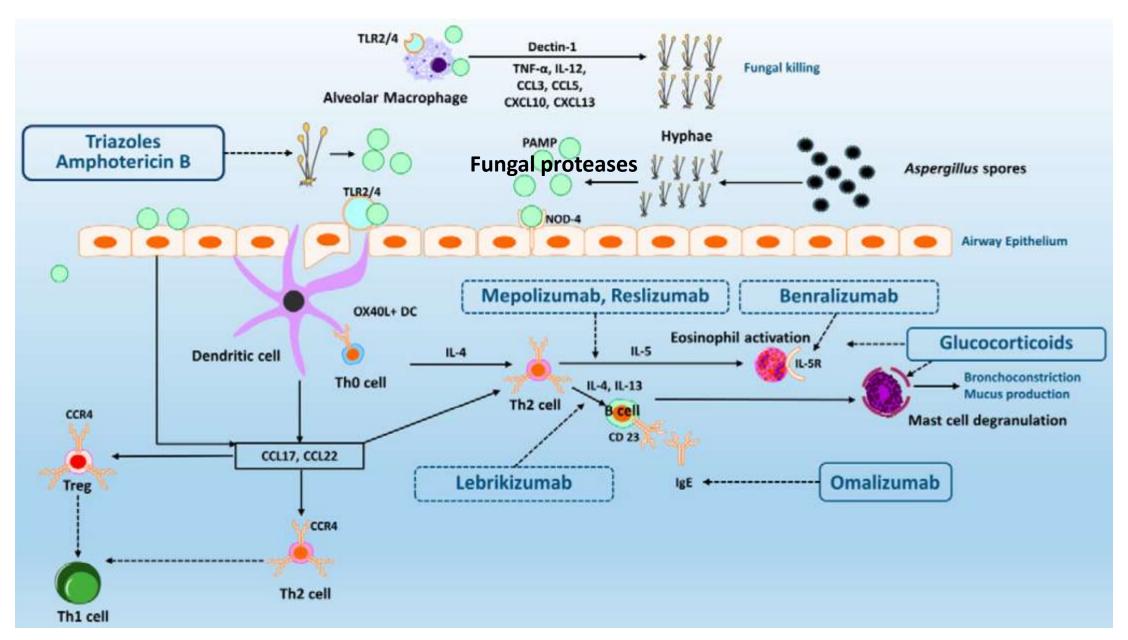


Figure 3. The differing morphological stages of A. fumigatus growth, as time proceeds, resting conidia (3a) begin to swell (3b) and germinate (3c), eventually forming hyphae (3d). [A. fumigatus conidia (1 × 10⁷ ml) were added to minimal essential medium (Sigma) supplemented with 5% fetal calf serum and incubated at 37°C. A 1 ml aliquot was withdrawn at the times indicated, diluted in ice cold PBS to halt any further development and representative images were captured using an Olympus BX51 Colorview soft imaging system).

Allergic Bronchopulmonary Aspergillosis (ABPA)- Immune response



Allergic Bronchopulmonary Aspergillosis (ABPA)- Pathogenesis

	Mutations/ polymorphisms	Population	Number of patients	Control population	Significance OR (95% confidence intervals)	Author/reference
	HLA (6p21_3)	1.0400.000.000				
	DR4	Caucasian	16 ABPA (asthma)	56 allengy; 39 controls	Allergy: 0.9 (0.3-2.9), P = 0.9; Control: 22.8	Ame et al. [49]
DF	DRS	Caucasian	16 ABPA (asthma)	56 allengy; 39 controls	(2.5-211.8), P = 0.002 Control: 5.3 (1.4-20.7), P = 0.02	Ame et al. [49]
		Caucasian	35 ABPA (asthma and CF)	50 Af sensitized asthma/CF; 90 controls	Asthma: 1.8 (0.7-4.9), P = 0.2; Control: 2.8 (1.1-6.8), P = 0.03	Chauhan et al. [50]
	DR7	Caucasian	16 ABPA (asthma)	56 allengy; 39 controls	Allergy: 1.7 (0.5-5.7), P = 0.4; Control: 35 (1.8-691.4), P = 0.004	Aron et al. [49]
	DR2	Caucasian	35 ABPA (both asthma and CF related)	50 Af sensitized asthma or CF; 98 controls	Asthma: 4.9 (1.8-13.Q, P= 0.001 Control: 17 (1.6-8.4), P= 0.001	Chashan et al. [50]
	DR2/DRS	Caucasian	35 ABPA (both asthma and CF related)	50 Af sensitized asthma or OF; 98 controls	Asthma: 5, 1 (1.9–13.3), P = 0.0005; Control: 5,4 (2.3–12.9), P < 0.0001	Chauhan et al. [50]
	DRB1*1501	Caucasian	35 ABPA (both asthma and CF related)	50 Af sensitized asthma or CF; 9ft controls	Asthma: 1.1 (0.9–10.1), P = 0.05 Control: 4.5 (2.1-9.7), P = 0.0001	Chauhan et al. [50]
	DRB1*1503	Caucasian	35 ABPA (both arthma and CF related)	50 Af sensitized arthma or CF; 98 controls	Asthma: 24.8 (1.4-452.7), P = 0.008 Control: 37.5 (4.4-316.8), P < 0.0001	Chauhan et al. [50]
_	DRB1*0701, DRB1*1501, DQB1*0602, DQB1*0201	Caucanian	38 ABPA (CF)	46 CF, 106 asthma, 176 controls	DRB 1*0701, DRB 1*1501, DQB 1*0602 associated with ABPA sasceptibility, while DQB 1*0201 associated with possible protection	Muro et al. [64]
	Mannose-binding lectin (1	Oq11.2-q21)			A PORCH CONTRACTOR AND ADDRESS OF THE PROPERTY	
	31011A in intron 1	Indian	11 ABPA (esthma)	49 allergic individuals; 84 controls	Allergy: 1.3 (0.5–3.1), P = 0.7 Control: 0.2 (2.8–23. 6, P < 0.0001	Kaur et al. (60)
	Exon 1 (R52C, G54D, G57E) Promoter B1/L -550,	Caucarian	38 allergic fungal disease (28 ABPA, 7 SAFS, 3 NOS)	Historical controls	No significant relationship, P > 0.05	Harrison et al. [61]
•	Y/X -221, P/Q + 4) Surfactant Protein A2 (10)	. 2.2.21				
,	01649C in exon 4	Indian	32 ABPA (asthma)	34 controls	2.6 (1.2-5.7), P = 0.01	Saxena (2003)[53]
	arment in table 4	Caucasian	7 ABPA (asthma)	46 controls	2.7 (0.3-21.9), P = 0.6	Vaid (2007)[5/l]
	f1492C in intron 3	Indian	32 ABPA (asthma)	34 controls	4.8 (1.1-21.6), P = 0.03	Saxena (2003)[53]
_	A 1660G in exon 4	Caucasian Indian	7 ABPA (asthma) 27 ABPA, 119 Af colonitors	46 controls	3.5 (0.7-16.7), P = 0.2 5.3 (1.7-16.9), 0.002	Vaid (2007)[58] Saxena et al. [53]
.9	foll-like receptor 9 (3p21.	33				
	f1237C in 5' promoter	Caucasian	22 ABPA (asthma)	14 SAFS, 80 controls	SAFS: 6.9 (0.8-58.2), P = 0.09; Control: 2.5 (1.01-6.1), P = 0.04	Cavalho et al. [59]

	200100000000000000000000000000000000000					
	Mutations/ polymorphisms	Population	Number of patients	Control population	Significance OR (95% confidence intervals)	Author/reference
	IL-4Rx (16p12.1-p11.2)		POST SERVICE SE	ACCURATION OF THE PARTY OF THE		VII
/10	-4G>A file75val) in promoter IL-10 (13q13)	Caucasian	40 ABPA (14 asthma, 26 CF)	56 non-ABPA (23 asthma, 33 CF)	3.3 (1.8-6.1), P = 0.008	Knutsen et al. [56]
	-1082 G>A in promoter	Caucasian	27 ABPA (CF)	BI OF	GG genotype: 1.67 (0.64-4.36); AG genotype: 0.43 (0.15-1.18)	Brouard et al. [54]
		Caucasian	9 ABPA	24 CCPA	0.38 (0.21-0.67), P = 0.0006	Sambatakou (2000 [57]
	TGF-\$ (19q13.1, 13.2)					
F-β	TBMC in exon 1	Caucasian	9 ABPA	24 CCPA	0.42 (0.24-0.75), P = 0.003	et al. [57]
-	CFTR mutations (7q31.2)					
	CHff1 gene (1q31-32)	Caucasian	79 ABPA in asthma	268 controls 94 asthmatics	Control: 10.4 (4.4-24.8) Asthma: 5.5 (1.6-18.8)	Müler et al., Aron et al., Marchand et al., Eaton et al., Agarwal et al. [48, 49, 51, 52, 63
	24 bp duplication in exon 10	NA	6 ABPA	5	All six children had 24 bp duplication	Vicencio et al. [63]

Af, Aspergillus fumigatus; CCPA, chronic cavitary pulmonary aspergillosis; CFTR, CF transmembrane conductance regulator; HLA, human leucocyte antigen; IL, interleukin; MBL, mannose-binding lectin; NOS, not otherwise specified; OR, odds ratio; SAFS, severe asthma with fungal sensitization; SNP, single nucleotide polymorphism; SP, surfactant protein; TGF, transforming growth factor; TLR, toll-like receptor; TNF, tumour necrosis factor.

Genetic susceptibility in ABPA complicating asthma and CF

(mutations/polymorphisms HLA/DR, MBL, SPA2, TLR-9, IL-4Ra, IL-10, TGF-β)

Allergic Bronchopulmonary Aspergillosis (ABPA)- Epidemiology

Table 1. Prevalence of Aspergillus sensitization (AS) and allergic bronchopulmonary aspergillosis (ABPA) complicating asthma in studies conducted in this millennium

Study	Country	Type of study	Skin test/antigen	Prevalence of AS, n/N (%; 95% CI)	Prevalence of ABPA, n/N (%; 95% CI)
Eaton et al. [25]	New Zealand	Prospective	SPT/commercial (Hollister-Stier, USA)	47/255 (18.4; 14.1–23.7)	12/243 (4.9; 2.8–8.5)
Kumar et al. [30]	India	Prospective	Intradermal/indigenous	47/200 (23.5; 18.1-29.9)	32/200 (16; 11.5-21.8)
Al-Mobeireek et al. [26]	Saudi Arabia	Prospective	SPT/commercial (SoluPrick, ALK labs)	12/53 (22.6; 13.3–35.8)	7/264 (2.7; 1.3–5.5)*
Maurya et al. [31]	India	Prospective	Intradermal/indigenous	30/105 (28.6; 20.8-37.9)	8/105 (7.6; 3.9-14.5)
Agarwal et al. [32]	India	Prospective	Intradermal/commercial (Hollister-Stier)	291/755 (38.5; 35.1–42.1)	155/755 (20.5; 17.8–23.6)
Prasad et al. [33]	India	Prospective	Intradermal/not available	74/244 (30.3; 24.9-36.4)	18/244 (7.4; 4.7-11.4)
Agarwal et al. [34]	India	Prospective	Intradermal/indigenous	87/242 (35.9; 30.2-42.2)	54/242 (22.3; 17.5-28)
Ghosh et al. [35]	India	Prospective	Intradermal/indigenous	54/215 (25.1; 19.8-31.3)	15/215 (6.9; 4.2-11.2)
Sarkar et al. [36]	India	Prospective	SPT/commercial (Creative Drug Industries, India)	40/126 (31.7; 24.2–40.4)	10/126 (7.9; 4.3–14.1)*
Ma et al. [27]	China	Prospective	_	11/200 (5.5; 3.1-9.7)	5/200 (2.5; 1.0-5.9)
Pooled prevalence	**************************************			25.1 (19.6-31.6)	8.4 (5.3-13.1)

^{*}Allergic bronchonulmonary mycosis

- AS complicating asthma (AIA): 5.5% to 38.5% with a pooled prevalence of 25%
- ABPA in asthma ranges between 2.5 and 22.3% with a pooled prevalence of 8.4%

Allergic Bronchopulmonary Aspergillosis (ABPA)- Clinical Features

- √ poorly controlled asthma
- ✓ golden-brown sputum (56%),
- √ peripheral eosinophilia

1/3 relatively asymptomatic despite extensive radiological lesions

113 patients with ABPA

mean age: 32 years,

mean age of onset of asthma:21 years.

- ✓ Cough (99%)
- ✓ Breathlessness (99%)
- Expectoration (98%)
- ✓ Wheezing (97%)
- ✓ Haemoptysis (41%)
- ✓ Nasal symptoms 45%
- ✓ Expectoration of sputum plugs 37%
- ✓ Nasal plugs by 6%

Plain chest radiology

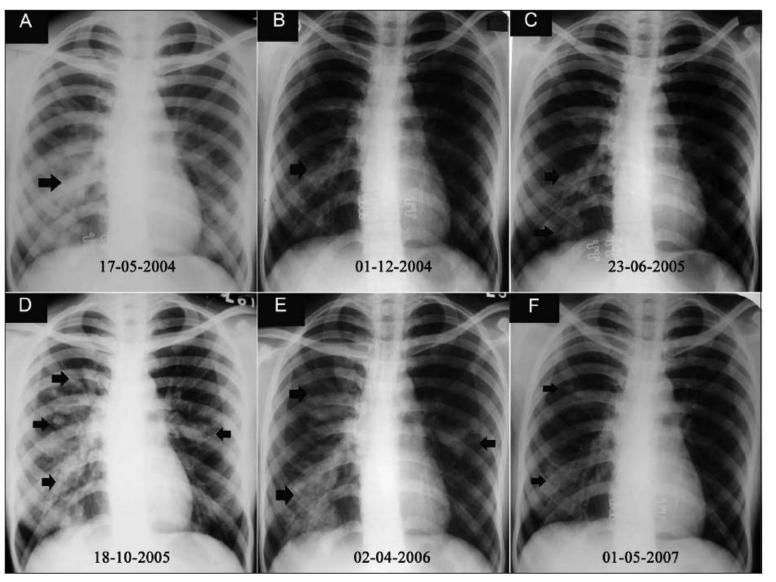
Transient changes

- Perihilar infiltrates simulating adenopathy
- Air-fluid levels from dilated central bronchi filled with fluid and debris
- Massive consolidation-unilateral or bilateral
- Radiologic infiltrates
- 'Toothpaste' shadows due to mucoid impaction in damaged bronchi
- 'Gloved finger' shadows from distally occluded bronchi filled with secretions
- 'Tramline' shadows representing oedema of the bronchial walls
- Collapse-lobar or segmental

Permanent changes

- Central bronchiectasis with normal peripheral bronchi
- Parallel-line shadows representing bronchial widening
- Ring-shadows 1-2 cm in diameter representing dilated bronchi en face
- Pulmonary fibrosis
- Late changes-cavitation, contracted upper lobes and localised emphysema

Consolidation (transient patchy-91%)



Y-shape and Finger-in-glove opacities (Mucoid impaction)

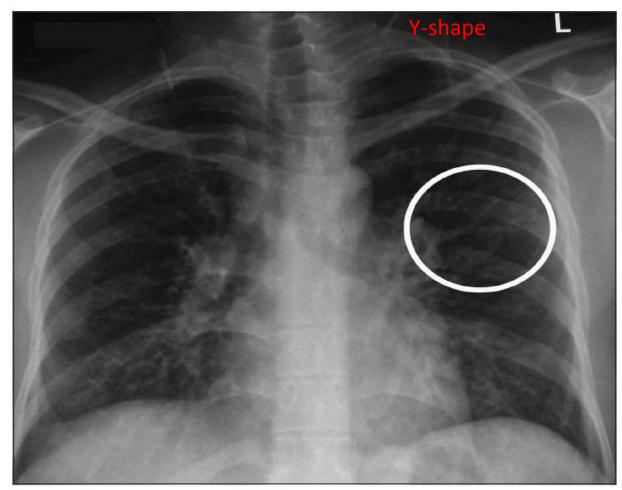


Figure 3: Chest radiograph shows a "Y-shaped" opacity (circle) that represent mucus-filled bronchi

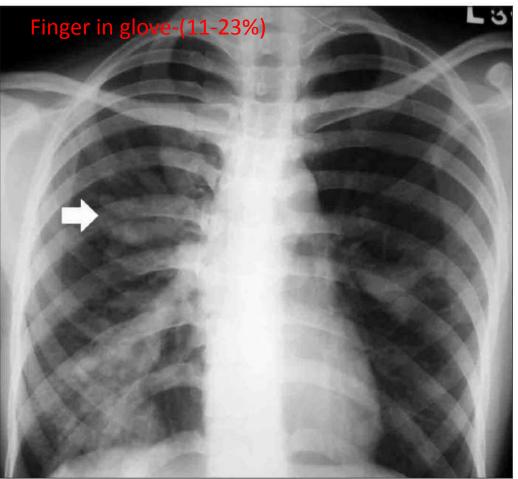
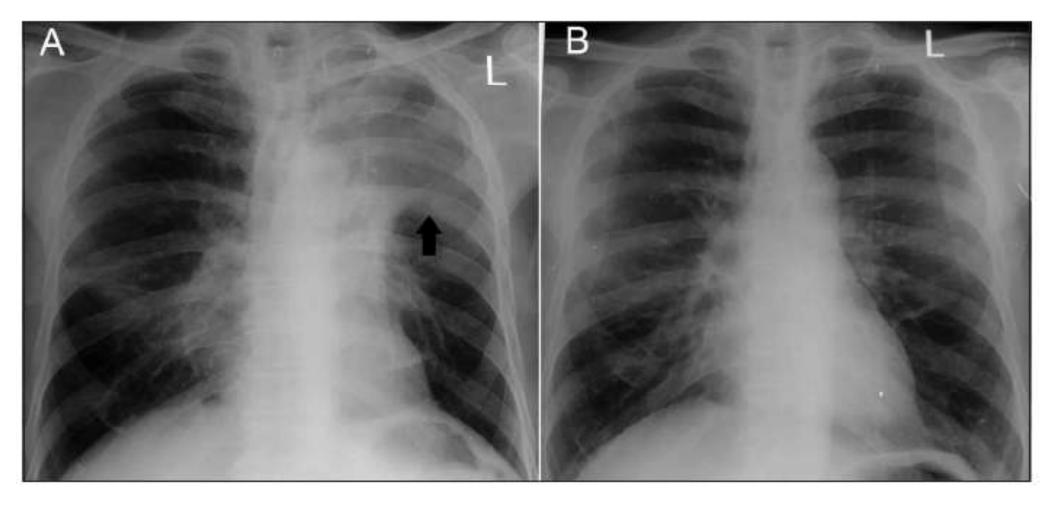


Figure 4: Chest radiograph shows mucoid impaction with the classic finger-in-glove pattern (arrow)

Atelectasis (14-39%)



Tram-line shadows (edema of the bronchial walls (45-92%)

Parallel lines (65-70%) - bronchial widening = permanent change

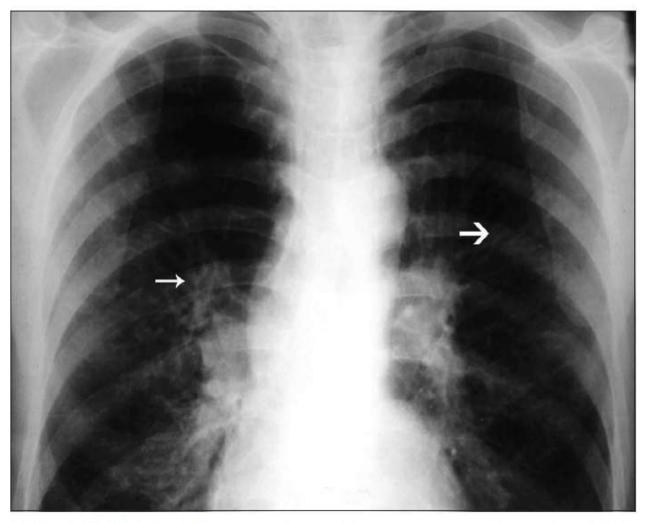


Figure 7: Chest radiograph shows the presence of tram-line (thick arrow) and parallel-line (thin arrow) shadows

Bronchiectasis

Figure 8: Chest radiograph shows central bronchiectasis (arrow) in the left mid-zone

Fibrosis

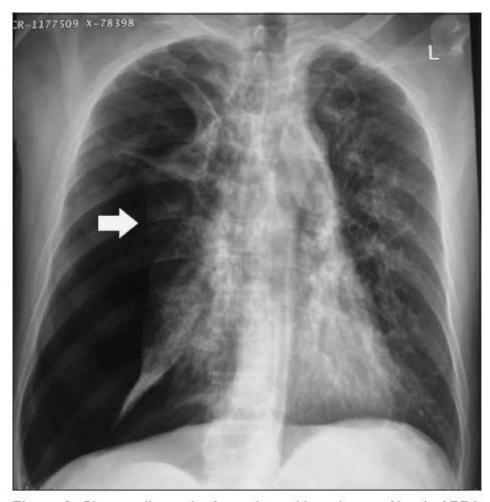


Figure 9: Chest radiograph of a patient with end-stage fibrotic ABPA who presented with a right-sided spontaneous pneumothorax (arrow)

R.Agarwal et al , Indian Journal of Radiology and Imaging / November 2011

Computed tomography findings

Bronchial abnormalities

- Bronchiectasis, usually central, as characterised by the 'signet ring' and 'string of pearls' appearances
- Dilated bronchi with or without air-fluid levels
- Totally occluded bronchi
- Bronchial wall thickening
- Parallel-line opacities extending to the periphery
- · High attenuation mucous plugs

Parenchymal changes

- Consolidation
- Non-homogeneous patchy opacities
- Parenchymal scarring of varying extent
- Segmental or lobar collapse
- Cavitation
- · Emphysematous bullae

Pleural involvement

- Pleural effusions
- Spontaneous pneumothorax
- Bronchopleural fistula
- Pleural fibrosis
- Pleural thickening

Central Bronchiectasis (CB-ABPA)

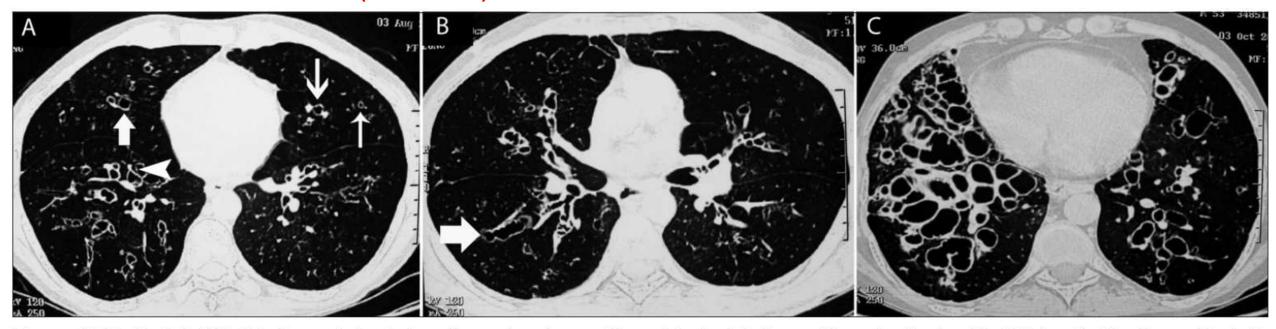
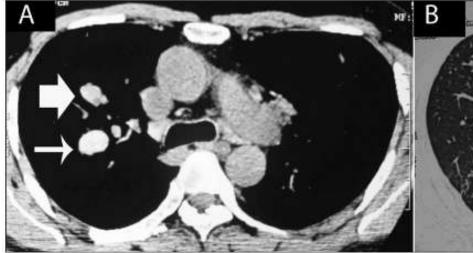
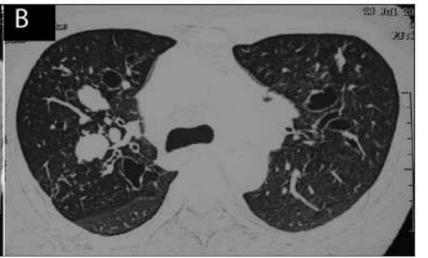


Figure 10 (A-C): Axial HRCTs (lung window) show the various types of bronchiectasis in three different patients with ABPA: cylindrical bronchiectatic cavities (thin arrow) of various sizes with the characteristic signet-ring appearance (thick arrow) (A), varicose bronchiectasis (arrows in B), and cystic bronchiectasis

- ✓ 26%–39% are associated with peripheral bronchiectasis
- ✓ usually upper lobes
- ✓ Cylindrical, varicose, cystic
- ✓ Central bronchiectasis (CB) is a sine qua non for the diagnosis of ABPA

Atelectasis and mucoid impaction









- filling of the airways by mucoid secretions
- generally hypodense
- may also have high CT attenuation values (HAM)-20%
- > pathognomonic of ABPA specificity of 100% sensitivity 19-32% should be considered as a radiological criteria separate from other findings.

HAM (High Attenuation Mucus): denser than the paraspinal skeletal muscle

Centrilobular nodules / tree-in-bud / mosaic pattern

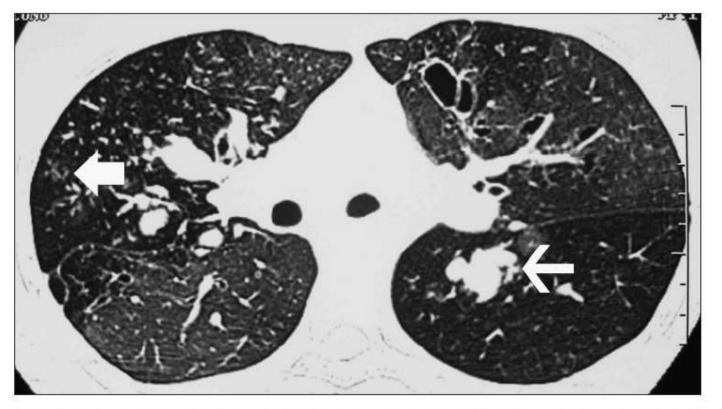


Figure 16: Axial HRCT (lung window) shows a mosaic pattern. There is central bronchiectasis with mucoid impaction in many of the bronchiectatic cavities (thin arrow). Also seen are centrilobular nodules in a tree-in-bud pattern (bold arrow)

Spontaneous pneumothorax, Fibrocavitary disease

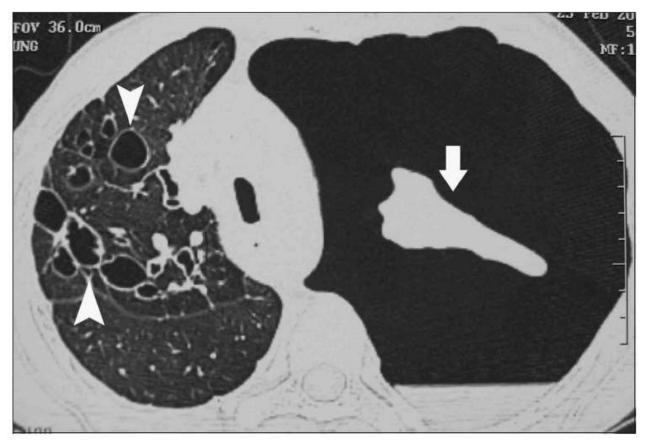


Figure 17: Axial HRCT (lung window) in a patient of ABPA who presented with a left-sided spontaneous pneumothorax (arrow). Extensive central and peripheral bronchiectasis is seen involving the right lung (arrowheads)

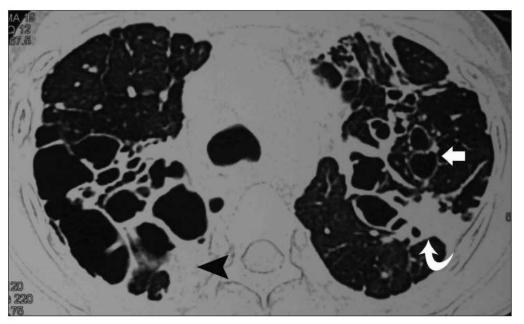
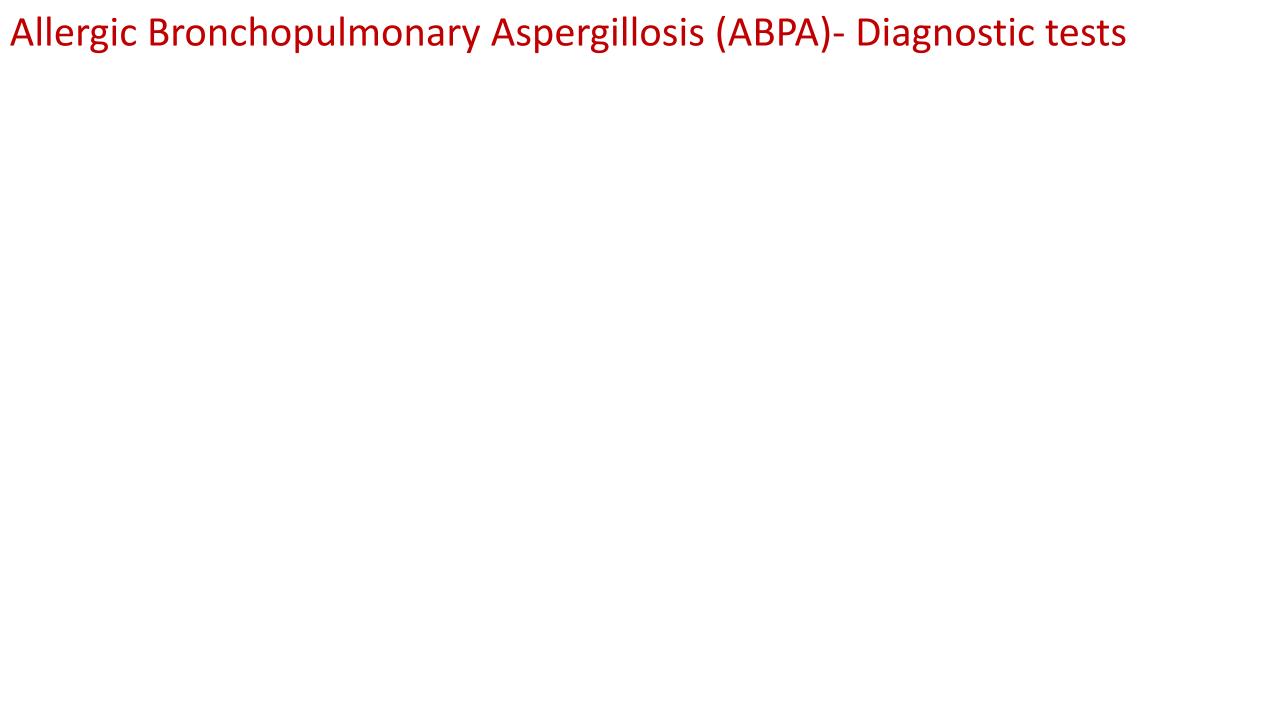


Figure 18: Axial HRCT (lung window) shows extensive bronchiectatic cavities (arrows), with pleural (arrowhead) and pulmonary fibrosis (curved arrow)



- > Aspergillus Skin test: is a surrogate marker for ABPA was regarded as hallmark of ABPA
 - Sensitivity 88- 94%
 - Should be replaced by Aspergillus s.IgE
- > A.Fumigatus specific IgE: are considered to be a hallmark of ABPA
 - level > 0.35 kUA/L sensitivity 100% (must be used as screening test) specificity 66.2%
 - Unreliable for follow up of ABPA
- > Total Serum IgE: diagnosis and follow up of ABPA
 - A normal serum IgE generally excludes active ABPA
 - The cut-off value remains speculative and needs validation
 - level > 1000IU/mL (2400ng/ml) (classic ABPA): sensitivity 39% specificity 100%
 - The lowest value after treatment (clinical and radiological improvement) is a 'new' baseline for an individual.
 - An increasing level (>50% of the 'new' baseline) of total IgE along with clinical and radiological worsening =exacerbation of ABPA

> Serum precipitins or specific IgG to A.Fumigatus:

- 10% of asthmatics with or without SAFS
- A.fumigatus-specific IgG (>27 mgA/L) is far more sensitive (89%) than Aspergillus precipitins (27%)

> Peripheral eosinophilia:

- > 500 cells/μL criterion for diagnosis of ABPA
- only 40% of patients with ABPA > 1000 cells/IL at diagnosis
- a low eosinophil count does not exclude ABPA

> Sputum cultures for A. Fumigatus:

- supportive but not diagnostic of ABPA
- 39 to 60% depending on the number of specimens examined
- vast majority of culture-negative ABPA patients have detectable A. fumigatus DNA in their sputum
- Susceptibility to antifungal agents

> Pulmonary function tests:

- helpful in categorizing the severity of asthma and the underlying lung disease.
- can be normal in ABPA
- normal spirometry should not exclude ABPA

> Recombinant Aspergillus antigens:

- Asp f1,3 in AS and ABPA, Asp f3,4,6 in ABPA
- 36-68% specificity

> Galactomannan detection:

- Polysaccharide component of aspergillus cell wall
- Sensitivity 25,7%
- Specificity 82%

			Greenbe			
		1986	1991-2	002	2013/2016	1999
1952 First Case Series (79)	1977 Diagnostic Criteria (43)	Rosenberg- Patterson criteria ^{46,47}	1991 Diagnostic Criteria: Revised (44)	'Truly minimal' criteria ⁷	ISHAM Working Group ²⁹	ABPA in CF ⁵⁵
Clinical features described	Asthma Total IgE elevated Immediate skin test positive Serum eosinophilia Precipitins Parenchymal infiltrates Central bronchiectasis	 Major criteria Asthma Presence of transient pulmonary infiltrates (fleeting shadows) Immediate cutaneous reactivity to Af Elevated total serum IgE Precipitating antibodies against Af Peripheral blood eosinophilia Elevated serum IgE and IgG to Af Central/proximal bronchiectasis with normal tapering of distal bronchi Minor criteria Expectoration of golden brownish sputum plugs Positive sputum culture for Aspergillus species Late (Arthus-type) skin reactivity to Af 	ABPA-CB Asthma Immediate skin test positive Total IgE elevated Specific IgE and IgG elevated Central bronchiectasis ABPA-S Asthma Immediate skin test positive Total IgE elevated Specific IgE & IgG elevated Additional findings Mucus plugs Sputum + aspergillus Precipitins Parenchymal infiltrates Delayed skin test positive	1. Asthma 2. Immediate cutaneous reactivity to Af 3. Total serum IgE > 1,000 ng/mL (417 kU/L) 4. CB in the absence of distal bronchiectasis	Predisposing conditions 1. Bronchial asthma 2. Cystic fibrosis Obligatory criteria (both should be present) 1. Type I Aspergillus skin test positive (immediate cutaneous hypersensitivity to Aspergillus antigen) or elevated IgE levels against Af 2. Elevated total IgE levels (>1,000 IU/mL)* Other criteria (at least two of three) 1. Presence of precipitating or IgG antibodies against Af in serum 2. Radiographic pulmonary opacities consistent with ABPA 3. Total eosinophil count >500 cells/µL in steroid naïve patients (may be historical) (*If the patient meets all other criteria, an IgE value <1,000 IU/mL may be acceptable)	Presence of two of the following three: (i) Immediate skin reactivity to Af antigens, (ii) Precipitating antibodies to Af antigens, (iii) Total serum IgE >1,000 IU/mL; and at least two of the following six: (i) Bronchoconstriction, (ii) Peripheral blood eosinophilia >1,000/µL, (iii) History of pulmonary infiltrates, (iv) Elevated specific IgE-Af/IgG-Af, (v) Af in sputum by smear or culture, (vi) Response to steroids

			Greenbe	erger		•
		1986	1991-2002		2013/2016	1999
1952 First Case Series (79)	1977 Diagnostic Criteria (43)	Rosenberg- Patterson criteria ^{46,47}	1991 Diagnostic Criteria: Revised (44)	'Truly minimal' criteria ⁷	ISHAM Working Group ²⁹	ABPA in CF ⁵⁵
Clinical features described	Asthma Total IgE elevated Immediate skin test positive Serum eosinophilia Precipitins Parenchymal infiltrates Central bronchiectasis	Major criteria 1. Asthma 2. Presence of transient pulmonary infiltrates (fleeting shadows) 3. Immediate cutaneous reactivity to Af 4. Elevated total serum IgE 5. Precipitating antibodies against Af 6. Peripheral blood eosinophilia 7. Elevated serum IgE and IgG to Af 8. Central/proximal bronchiectasis with normal tapering of distal bronchi Minor criteria 1. Expectoration of golden brownish sputum plugs 2. Positive sputum culture fo Aspergillus species 3. Late (Arthus-type) skin reactivity to Af	ABPA-CB Asthma Immediate skin test positive Total IgE elevated Specific IgE and IgG elevated Central bronchiectasis ABPA-S Asthma Immediate skin test positive Total IgE elevated Specific IgE & IgG elevated Additional findings Mucus plugs Sputum + aspergillus Precipitins Parenchymal infiltrates Delayed skin test positive	1. Asthma 2. Immediate cutaneous reactivity to Af 3. Total serum IgE >1,000 ng/mL (417 kU/L) 4. CB in the absence of distal bronchiectasis	Predisposing conditions 1. Bronchial asthma 2. Cystic fibrosis Obligatory criteria (both should be present) 1. Type Aspergillus skin test positive (immediate cutaneous hypersensitivity to Aspergillus antigen) or elevated IgE levels against Af 2. Elevated total IgE levels (>1,000 IU/mL)* Other criteria (at least two of three) 1. Presence of precipitating or IgG antibodies against Af in serum 2. Radiographic pulmonary opacities consistent with ABPA 3. Total eosinophil count >500 cells/µL in steroid naïve patients (may be historical) (*If the patient meets all other criteria, an IgE value <1,000 IU/mL may be acceptable)	Presence of two of the following three: (i) Immediate skin reactivity to Af antigens, (ii) Precipitating antibodies to Af antigens, (iii) Total serum IgE >1,000 IU/mL; and at least two of the following six: (i) Bronchoconstriction, (ii) Peripheral blood eosinophilia >1,000/µL, (iii) History of pulmonary infiltrates, (iv) Elevated specific IgE-Af/IgG-Af, (v) Af in sputum by smear or culture, (vi) Response to steroids

ISHAM working group criteria - 2013

Table 4. Newly proposed diagnostic criteria for allergic bronchopulmonary aspergillosis

Predisposing conditions

Bronchial asthma, cystic fibrosis

Obligatory criteria (both should be present)

Type I Aspergillus skin test positive (immediate cutaneous hypersensitivity to Aspergillus antigen) or elevated IgE levels

against Aspergillus fumigatus

Elevated total IgE levels (> 1000 IU/mL)*

Other criteria (at least two of three)

Presence of precipitating or IgG antibodies against A. fumigatus in serum

Radiographic pulmonary opacities consistent with ABPA†

Total eosinophil count > 500 cells/ μ L in steroid naïve patients (may be historical)

*If the patient meets all other criteria, an IgE value < 1000 IU/mL may be acceptable.

[†]The chest radiographic features consistent with ABPA may be transient (i.e. consolidation, nodules, tram-track opacities, toothpaste/finger-in-glove opacities, fleeting opacities) or permanent (i.e. parallel line and ring shadows, bronchiectasis and pleuropulmonary fibrosis).

doi: 10.1111/cea.12141

Clinical & Experimental Allergy, 43, 850-873

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OPINIONS IN ALLERGY

Allergic bronchopulmonary aspergillosis: review of literature and proposal of new diagnostic and classification criteria

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Proposed diagnostic criteria-ISHAM working group 2016

A. Predisposing conditions

Bronchial asthma, cystic fibrosis, COPD, post-tuberculous fibrocavitary disease

B. Essential criteria (both must be met)

- i. Serum Aspergillus fumigatus-specific IgE levels >0.35 KUA/L ‡
- ii. Elevated serum total IgE levels >1000 IU/mL*

Additional criteria (at least two of three)

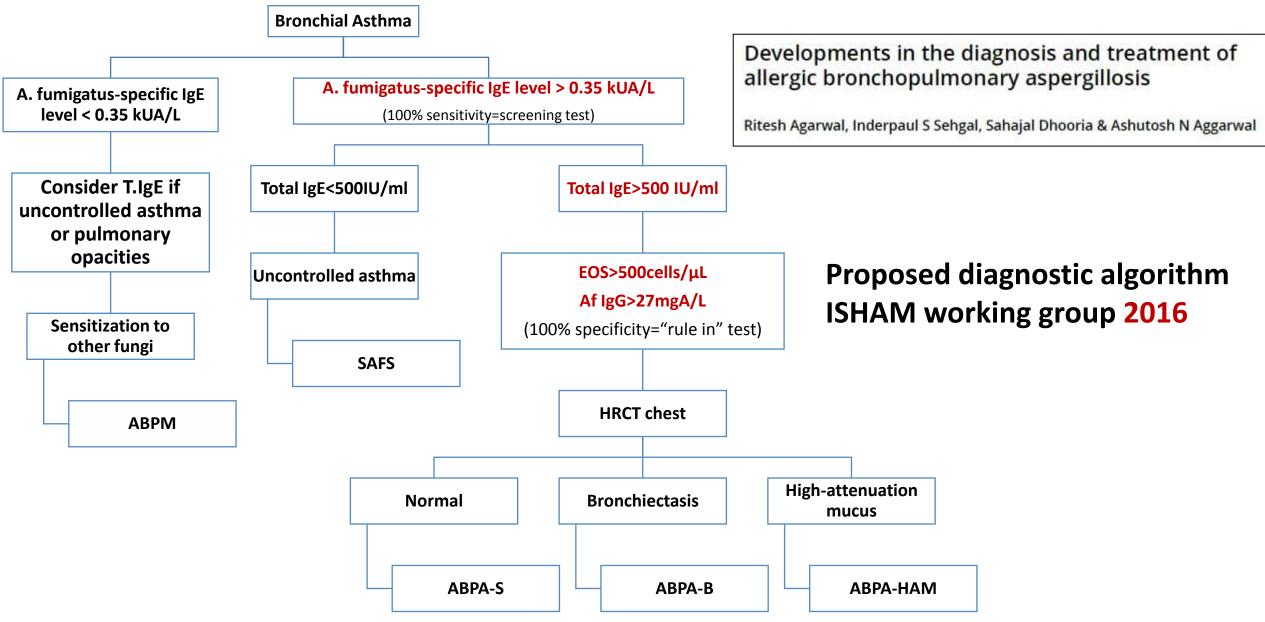
- i. Serum Aspergillus fumigatus-specific IgG levels >27 mgA/L
- ii. Thoracic imaging findings consistent with ABPA†
- iii. Peripheral blood eosinophil count >500 cells/ μ L (may be historical)

^{*}An IgE value <1000 IU/mL may be acceptable, if all other criteria are met (especially if the serum Aspergillus fumigatus-specific IgG levels >27 mgA/L)

[†]Features on HRCT chest and/or chest radiograph consistent with ABPA include transient abnormalities (i.e. nodules, consolidation, mucoid impaction, hyperattenuating mucus, fleeting opacities, toothpaste/gloved finger opacities, tramtrack opacities) or permanent (i.e. parallel lines, ring shadows, bronchiectasis and pleuropulmonary fibrosis).

‡A positive type I Aspergillus skin test may be considered as a criterion in the place of serum Aspergillus fumigatus-specific IqE levels only if the latter test is not available

Allergic Bronchopulmonary Aspergillosis (ABPA)- Diagnostic Algorithm



Allergic Bronchopulmonary Aspergillosis (ABPA)

Proposed Clinical Staging/Course - ISHAM working group 2016

Clinical stagi	Clinical staging of allergic bronchopulmonary aspergillosis (ABPA) in patients with asthma					
Stage	Definition	Features				
0	Asymptomatic	 No previous diagnosis of ABPA Controlled asthma (according to GINA/EPR-3 guidelines) Fulfilling the diagnostic criteria of ABPA 				
1	Acute	 No previous diagnosis of ABPA Uncontrolled asthma/symptoms consistent with ABPA Meeting the diagnostic criteria of ABPA 				
1a	With mucoid impaction	Mucoid impaction observed on chest imaging or bronchoscopy				
1b	Without mucoid impaction	Absence of mucoid impaction on chest imaging or bronchoscopy				
2	Response	 Clinical and/or radiological improvement AND Decline in IgE by ≥25% of baseline at 8 weeks 				
3	Exacerbation	 Clinical and/or radiological worsening AND Increase in IgE by ≥50% from the baseline established during response/remission 				
4	Remission	 Sustained clinico-radiological improvement AND IgE levels persisting at or below baseline (or increase by <50%) for≥6 months off treatment 				
5a	Treatment-dependent ABPA	 ≥2 exacerbations within 6 months of stopping therapy OR Worsening of clinical and/or radiological condition, along with immunological worsening (rise in IgE levels) on tapering oral steroids/azoles 				
5b	Glucocorticoid -dependent asthma	Systemic glucocorticoids required for control of asthma while the ABPA activity is controlled (as indicated by IgE levels and thoracic imaging				
6	Advanced ABPA	 Extensive bronchiectasis due to ABPA on chest imaging AND Complications (cor pulmonale and/or chronic type II respiratory failure) 				

Allergic Bronchopulmonary Aspergillosis (ABPA)

Treatment of ABPA ISHAM

Treatment goals

- 1) control of symptoms of asthma or cystic fibrosis (CF)
- 2) prevent or treat pulmonary exacerbations of ABPA
- 3) reduce or remit pulmonary inflammation; and
- 4) mitigate progression to end-stage fibrotic or cavitary disease

Treatment goals

Early and aggressive treatment

- 1) control of symptoms of asthma or cystic fibrosis (CF)
- 2) prevent or treat pulmonary exacerbations of ABPA
- 3) reduce or remit pulmonary inflammation; and
- 4) mitigate progression to end-stage fibrotic or cavitary disease

1. Glucocorticoids

A) Oral corticosteroids

Treatment of choice for ABPA

Regimen 1 (low dose)

Prednisolone 0.5 mg/kg/day for one to two weeks, then on alternate days for six to eight weeks
Then taper by 5–10 mg every 2 weeks and discontinue

Regimen 2 (medium dose)

Prednisolone, 0.75 mg/kg for 6 weeks, 0.5 mg/kg for 6 weeks, then tapered by 5 mg every 6 weeks to continue for a total duration of at least 6–12 months

- 13% may not respond and may require escalation of steroid dose or other therapies
- 50% of patients relapse when they are tapered
- 20–45% glucocorticoid dependent (stage 5b)
- After discontinuation of prednisolone –monitoring every 6-8 weeks to ensure remission is maintained

1. Glucocorticoids

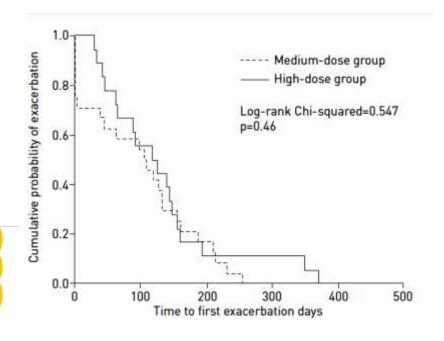
A) Oral corticosteroids

A randomised trial of glucocorticoids in acute-stage allergic bronchopulmonary aspergillosis complicating asthma

Ritesh Agarwal¹, Ashutosh N. Aggarwal¹, Sahajal Dhooria¹, Inderpaul Singh Sehgal¹, Mandeep Garg², Biman Saikia³, Digambar Behera¹ and Arunaloke Chakrabarti⁴

92 subjects (high-dose n=44, medium-dose n=48) were included in the study. The numbers of subjects with exacerbation after 1 year (high-dose 40.9% versus medium-dose 50%, p=0.59) and glucocorticoid-dependent ABPA after 2 years (high-dose 11.4% versus medium-dose 14.6%, p=0.88) were similar in the two groups. Although composite response rates were significantly higher in the high-dose group, improvement in lung function and time to first exacerbation were similar in the two groups. Cumulative glucocorticoid dose and side-effects were significantly higher in the high-dose group.

Medium-dose oral glucocorticoids are as effective and safer than high-dose in treatment of ABPA.



Low-dose oral glucocorticoids are as effective and safer than medium-dose in treatment of ABPA.

1. Glucocorticoids

B) Inhaled clucocorticoids

- High doses of ICS alone have a little role in the management of ABPA
- They can be used for asthma control

C) Intravenous pulse doses of glucocorticoids

- 15mg/kg methylprednisolone- max=1gr intravenously for 3 consecutive days
- Pediatric patients (steroid-sparing modality)
- Refractory asthma exacerbations

2. Antifungal agents • steroid-sparing agents

A) Itraconazole

- The most widely used
- Poor bioavailability, interactions with several drugs (+glucocorticoids)
- 200 mg twice a day, with therapeutic drug monitoring for at least 16 weeks.
- Response often takes longer than 16 weeks
- Tapered after 4-6 months (over 4 to 6 months)

Allergic Bronchopulmonary Aspergillosis (ABPA)

Treatment of ABPA ISHAM

2. Antifungal agents

A Randomized Trial of Itraconazole vs Prednisolone in Acute-Stage Allergic Bronchopulmonary Aspergillosis Complicating Asthma **≋**CHEST

Ritesh Agarwai, MD, DM; Sahajai Dhooria, MD, DM; Inderpaul Singh Sehgai, MD, DM; Ashutosh N. Aggarwai, MD, DM; Mandeep Garg, MD; Birnan Saikia, MD; Digambar Behera, MD; and Arunaloke Chakrabarti, MD

CONCLUSIONS: Prednisolone was more effective in inducing response than itraconazole in acute-stage ABPA. However, itraconazole was also effective in a considerable number and, with fewer side effects compared with prednisolone, remains an attractive alternative in the initial treatment of ABPA.

TABLE 2 Outcomes of Study Subjects Treated With Prednisolone or Itraconazole (N = 131)

Outcome	Prednisolone Group (n = 63)	Itraconazole Group (n = 68)	Estimated Difference (95% CI)	P Value
Primary outcomes				
Subjects with response following 6 wk of treatment ^a	63 (100%)	60 (88.2%)	-11.8 (-21.5 to -3.7)	(.007)
Subjects with response following 3 mo of treatment	63 (100%)	60 (100%)	0 (-0.06 to 0.06)	

- ✓ Prednisolone more effective in inducing response
- ✓ No difference in Serum IgE / exacerbations

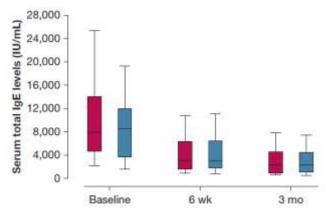
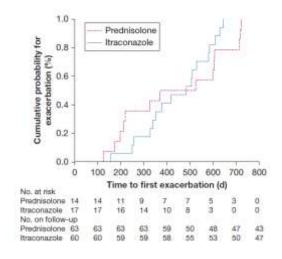


Figure 2 – Box and whisker plots showing the IgE levels at baseline, 6 weeks, and 3 months in the two groups (prednisolone: red plots; itraconazole: blue plots). Box plots represent the 25th and 75th per-



Adverse Reaction	Prednisolone Group (n = 63)	Ibraconazole Group (n = 60) ^a	Estimated Difference (95% CI)	P Value
Discontinuation of study drug	0	0	100	100
Cushingoid habitus	52 (82.5%)	0	82.5 (69.9 to 89.9)	.0001
Hypertension	0	0	444	1
Hyperglycemia	2 (3.2%)	0	3.2 (-3.3 to 10.9)	.50
Hypertrichosis	12 (19.1%)	0	19.1 (9.2 to 30.4)	.002
Acne	11 (17.5%)	0	17.5 (7.9 to 28.6)	.002
Striae	8 (12.7%)	0	12.7 (4.1 to 23.1)	.003
Weight gain (> 10% of baseline) at 6 wk	37 (58.7%)	2 (3.3%)	55.4 (40.7 to 66.9)	.0001
Mood changes	3 (4.8%)	0	4.8 (-2.0 to 13.1)	.24
Fatigue	3 (4.8%)	8 (13.3%)	-8.6 (-19.9 to 1.9)	.26
Liver function test abnormalities	0	9 (15%)	-15 (-26.1 to -6.0)	.001
Nausea	0	2 (3.3%)	-3.3 (-11.4 to 2.9)	.24

TABLE 3 1 Advance Boartions Noted in Chick Subjects Treated With Producedons or Iterapayole (n.

2. Antifungal agents • steroid-sparing agents

A) Itraconazole

- The most widely used
- Poor bioavailability, interactions with several drugs (+glucocorticoids)
- 200 mg twice a day, with therapeutic drug monitoring for at least 16 weeks.
- Response often takes longer than 16 weeks
- Tapered after 4-6 months (over 4 to 6 months)

2. Antifungal agents • steroid-sparing agents

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B) Nebulized amphotericin B

- No systemic absorption adverse events
- Limited efficacy
- Use when alternative options have been exhausted
- May prevent ABPA exacerbations

Nebulized amphotericin B

Amphotericin B deoxycholate

Daily: 5-40 mg twice daily

Intermittent: 20 mg (10 mg twice daily) thrice weekly

Liposomal amphotericin B

Intermittent: 25 mg twice weekly Amphotericin B lipid complex

Intermittent: 50 mg twice weekly

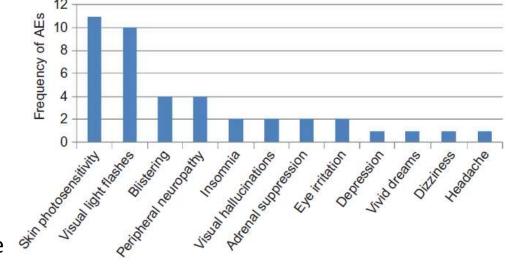
Allergic Bronchopulmonary Aspergillosis (ABPA)

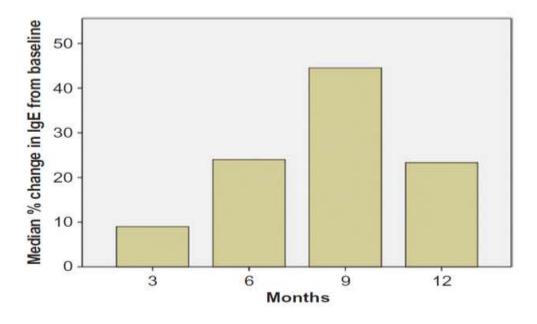
Treatment of ABPA ISHAM

2. Antifungal agents

C) Newer Azoles (Voriconazole, Posaconazole)

- Few studies
- Clinical improvement in 70-75%
- Better bioavailibity, less adverse reactions
- Reduction in OCS, improvement in asthma control, decline in IgE
- Symptomatic patients despite treatment, adverse effects to itraconazole





		3 month	rs (%)	6 month	is (%)	12 mc	onths (%)
Clinical or health-care utilization feature		Vori (n = 25)	Posa (n = 9)	Vori (n = 19)	Posa (n = 9)	Vori (n = 17)	Posa (n = 9
Symptoms	Reduction in cough frequency (%)	17/24 (70)	7/9 (78)	15/19 (78)	6/9 (67)	7/17 (41)	8/9 (89)
	Reduction in breathlessness (%)	10/24 (41)	5/9 (56)	12/19 (63)	4/9 (44)	7/17 (41)	4/9 (44)
	Increased energy (%)	8/24 (33)	4/9 (44)	8/19 (42)	4/9 (44)	7/17 (41)	5/9 (56)
	Reduced chest infections (%)	17/24 (70)	7/9 (78)	9/19 (47)	7/9 (78)	9/17 (53)	7/9 (78)
Medication use	Reduction in oral antibiotics use (%)	16/24 (67)	7/9 (78)	11/19 (58)	7/9 (78)	11/17 (64)	6/9 (78)
	Reduction in OCS use (%)	4/18 (22)	2/9 (29)	5/18 (28)	2/7 (29)	5/17 (29)	2/7 (29)
	Discontinuation of OCS (%)	8/18 (33)	4/7 (57)	12/18 (67)	4/7 (57)	15/17 (88)	3/7 (43)
	Reduction in SABA use (%)	12/25 (48)	6/9 (67)	8/19 (42)	5/9 (56)	10/17 (58)	7/9 (78)
Health-care service use	Reduction in hospital admissions (%)	9/10 (90)	1/2 (50)	9/10 (90)	1/2 (50)	9/10 (90)	2/2 (100)
	Reduction in GP/emergency visits (%)	13/25 (52)	6/9 (67)	11/19 (58)	8/9 (89)	12/17 (71)	6/9 (67)
Quality of life	Reduction in patients' overall symptoms (%)	18/25 (72)	7/9 (78)	13/19 (68)	7/9 (78)	10/17 (58)	7/9 (78)
78 (145 (147 KB)	Increased exercise tolerance (%)	7/25 (28)	4/9 (44)	6/19 (31)	5/9 (56)	5/17 (29)	4/9 (44)
	Increased QOL (%)	18/25 (72)	7/9 (78)	13/19 (68)	7/9 (78)	10/17 (58)	7/9 (78)

Allergic Bronchopulmonary Aspergillosis (ABPA)

Treatment of ABPA ISHAM

3. Other agents

A. Omalizumab (375mg sc /2w)

Table 3
Baseline characteristics of 102 individuals.

Baseline characteristics	No of data gained	
Age (years)	N = 102 (100)	
Mean (SD)		41 (19)
Median (range)		41 (7, 76)
Gender n (%)		A. Commence
Male		48 (47.1)
Female		54 (52.9)
Race n (%)		
Caucasian		98 (96.07)
Melanoderm		3 (2.94)
Xanthoderm		1 (0.98)
Clinical history n (%)		
with TB		2 (1.96)
with Asthma		17 (16.67)
with CF		40 (39.21)
ABPA duration prior to Anti-IgE(yrs)	N = 59 (57.8)	
Mean (SD)		5.4 (4.26)
Anti-fungal treatment		47 (46.08)
Treatment failure with systemic		101 (99.03)
steroids or itraconazole prior to treatment n (%)		
Total eosinophil count	N - 22 (21.57)	
Mean (SD)	15 15	676.36 (190.16)
Median (range)		676 (317, 1100)
Total IgE (IU/ml)	N = 97 (95.1)	The state of the s
Mean (SD)	277	1901 (1971.67)
Median (range)		1901 (131, 10,2)
Specific IgE for A.f (IU/ml)	N = 48 (47.05)	estation in the last services and
Mean (SD)		31.72 (24.16)
FEV1 of predicted (%)	N = 93 (91.17)	
Mean (SD)		59.63 (19.34)
Median (range)		60 (21, 115)
FVC of predicted (%)	N = 24(23.5)	
Mean (SD)		83.4 (21.6)
Median (range)		83 (45, 95)
FEV1/FVC	N - 31 (30.39)	
Mean (SD)		56.93 (14.16)
Median (range)		57 (41, 85)
Exacerbations prior	N = 98 (96.07)	
Mean (SD)		2.74 (2.31)
Median (range)		3 (0, 10)

102 ABPA patients

1091 IU/ml mean IgE

99% treatment failure to steroids/itraconazole

83% intravenous/16,67% sc Omalizumab

Dose: 225 mg to 750 mg, most commonly used dose was 375 mg every

two weeks

Table 4
Effect of omalizumab on ABPA patients.

	prior			poster			P value
	Mean	SD	N	Mean	SD	N	e
Total IgE (IU/ml)	1901	1971.67		804.5	514.7		< 0.001
Exacerbation rate (per year per patient)	2.7404	2.3117		0.38	0.698		< 0.001
FEV1 of predicted (%)	59.63	19.34		72.21	19		< 0.001
FVC%	83.4	21.6		94.83	22.11		0.0767
FENO(ppm)	31.4	24.36		17.66	11.24		0.0713
ACT score	11.367	6.2		18.53	9.5		0.0099
Prednisone dosage (mg/d)	16.39	13.47		1.63	2.25		< 0.001
No.of PSL use			96			67	< 0.001

86% decrease in exacerbations

30% discontinuation of steroids

70% reduction of steroids to <90% initial dose

3. Other agents

B. Mepolizumab

PubMed

Allergic bronchopulmonary aspergillosis Mepolizumab

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C. Therapeutic Bronchoscopy:

When atelectasis persists after 4 weeks of OCS treatment

D. Environmental Control:

Avoidance of gardening, farm related activities, renovations, compost, use of a mask

Allergic Bronchopulmonary Mucosis (ABPM)

- ABPA-like syndrome caused by fungi other than fumigatus (Candida albicans most often)
- Less than 150 cases reported globally
- Diagnostic criteria similar to ABPA (sensitization to the specific fungi)
- Clinical and lab findings similar to ABPA

Treatment similar to ABPA (antifungals according to their efficacy against a particular etiologic agent)

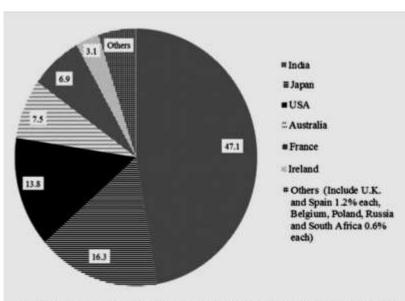
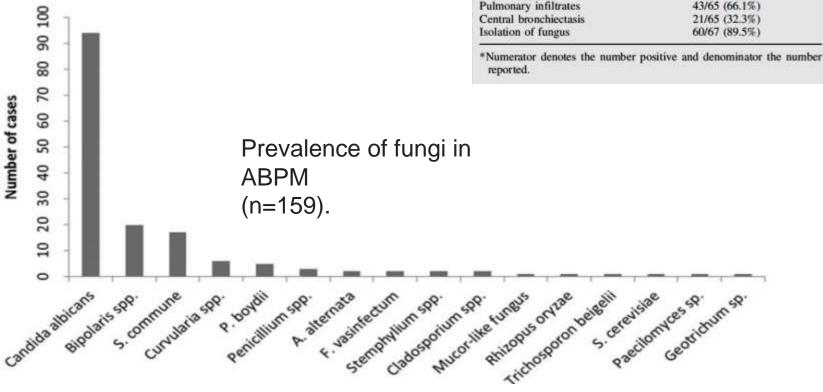


Figure 1. Geographic distribution (%) of 159 reported cases of allergic bronchonulmonary mycosis



A. Chowdhary et al., Crit Rev Microbiol, Early 2013

Table 6. Synopsis of clinical and laboratory diagnostic profiles of

allergic bronchopulmonary mycosis cases reported in English (n = 143).

Results

1400 (80-37, 530, n=63)

 41.70 ± 18.97 (6-84, n = 71)

46/143* (32.1%)

51/143 (35.6%)

100/116 (86.2%)

62/67 (92.5%)

39/43 (90.6%)

35/39 (89.7%)

52/55 (94.5%)

1.33:1

Characteristics investigated

History of asthma

Raised total IgE

Type I skin test

Eosinophilia

Precipitins

Mean age ± SD (years; range) Sex distribution (male:female)

History of allergic disorders

Median total IgE (IU/ml; range)

Specific (IgE/IgG) antibodies

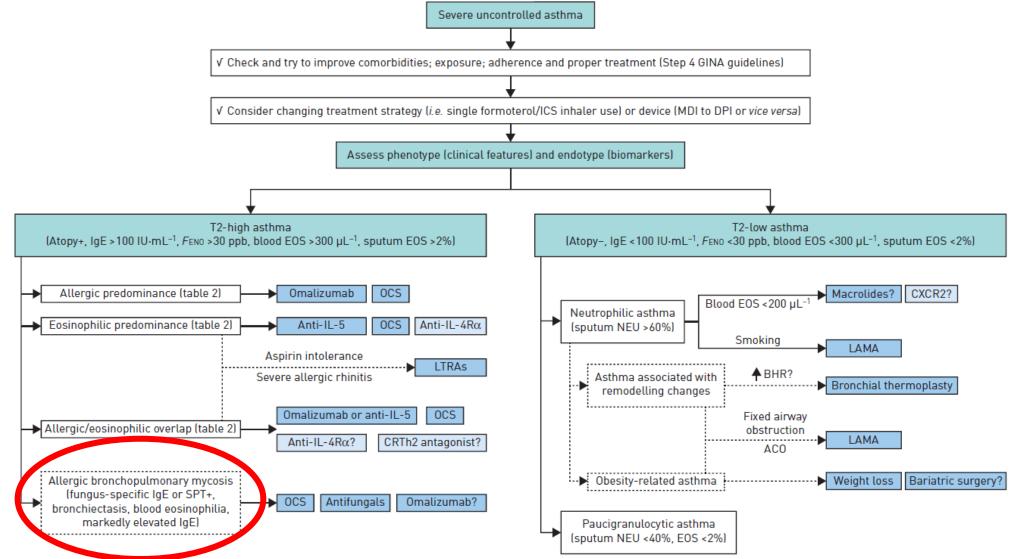
Allergic Bronchopulmonary Aspergillosis (ABPA)- Conclusions

- ❖ ABPA is a controllable, albeit chronic illness
- ❖ All asthmatic patients (regardless severity) should be routinely investigated for ABPA with A.Fumigatus specific IgE
- It is important to treat the disease aggressively during the early stages
- Glucocorticoids should be used as the first-line of therapy in ABPA, and itraconazole reserved in those with exacerbations and glucocorticoid-dependent disease.
- Newer therapies may be tried in those with recurrent exacerbations, glucocorticoid-dependent ABPA or in patients who develop treatment-related adverse reactions.

An algorithmic approach for the treatment of severe uncontrolled asthma

Eleftherios Zervas¹, Konstantinos Samitas¹, Andriana I. Papaioannou², Petros Bakakos³, Stelios Loukides² and Mina Gaga¹







Σοβαρό άσθμα και Αλλεργική Βρογχοπνευμονική Μυκητίαση

Κωνσταντίνος Σάμιτας MD PhD

Πνευμονολόγος, Επιμελητής Β΄ ΕΣΥ 6^η Πνευμονολογική Κλινική ΓΝΝΘΑ «Η ΣΩΤΗΡΙΑ»

09:00-10:30 Στρογγύλη Τράπεzα

Το σοβαρό Βρογχικό Άσθμα ως συννοσηρότητα Προεδρείο: Ε. Ζέρβας - Π. Μπακάκος

- Σοβαρό άσθμα και αππεργική βρογχοπνευμονική μυκητίαση Κ. Σάμπαs
- Ηωσινοφιλική κοκκιωμάτωση με πολυαγγειτιδα (EGPA) και σοβαρό άσθμα
 Ε. Φούκα
- Ασθμα ως συννοσηρότητα στη ΧΑΠ Γ. Χειβάς
- Άσθμα και ρινικοί πολύποδες Π. Μαραγκουδάκης



Allergic Bronchopulmonary Aspergillosis (ABPA)- CF

- Prevalence 2-15% in CF
- Prompt recognition is essential due to profound deterioration of lung function
- Wheezing, fleeting opacities, bronchiectasis, mucus plugging
- > Similar treatment

Practice points

Diagnosis in CF

- Acute or subacute deterioration in respiratory symptoms or lung function.
- Total serum IgE greater than 400 IU/ml.
- Skin-prick test positive to Af, together with either
 - o Positive Aspergillus precipitins
 - o Radiographic features consistent with ABPA.

Allergic Bronchopulmonary Aspergillosis (ABPA)- Proposed Scoring System- ISHAM 2016

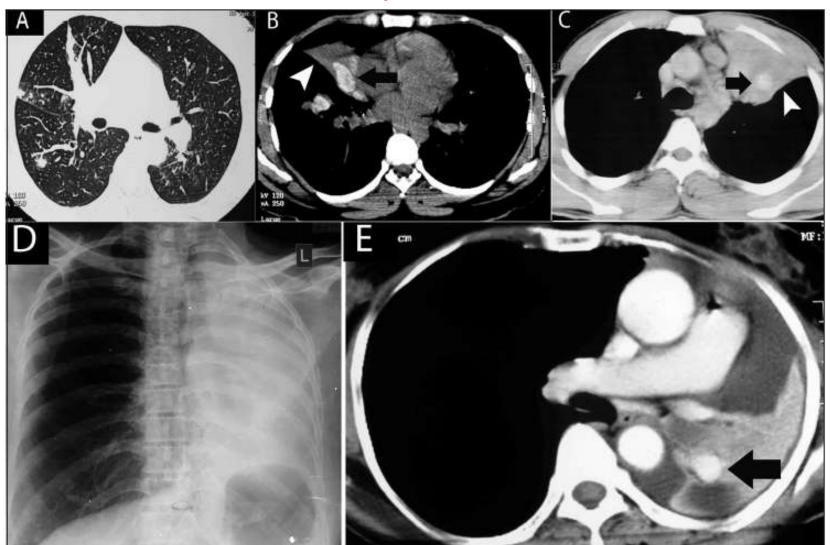
Immunological score	Value/findings	Score
A. fumigatus-specific IgE	<0.35 kUA/L	-7
	0.35-1.9 kUA/L	+1
	>1.9 kUA/L	+3
Total IgE	<417 IU/mL	-3/_
	417-1000 IU/mL	+1
	1000-2300 IU/mL	+2
	>2300 IU/mL	¥3
Peripheral blood eosinophil count	<500 cells/μL	(0)
Control of the Contro	500-1000 cells/μL) 1 3
	>1000 cells/μL	+4
A. fumigatus-specific IgG	<27 mg _A /L	0
1 500 1 10004	$>27 \text{ mg}_A/L$	+4
Radiological score		
HRCT chest*	Normal	0
	≥2 features of fibrosis	+2
	Bronchiectasis involving <3 lobes	+3
	Bronchiectasis involving ≥3 lobes	+4
	Extensive mucoid impaction	+4
	Hyperattenuating mucus	+5
Scoring		
Total score 8 with radiologic score 0	ABPA at risk	
Total score ≥9 with radiologic score of 0	ABPA-S (serological ABPA)	
Total score ≥9 with radiologic score of 2	ABPA-CPF (ABPA with chronic	
982	pleuropulmonary fibrosis)	
Total score ≥9 with radiologic score of 3 or 4	ABPA-B (ABPA with bronchiectasis)	
Total score ≥9 with radiologic score of 5	ABPA-HAM (ABPA with high attenuation	l)
	mucus)	

Allergic Bronchopulmonary Aspergillosis (ABPA)- Clinical Features

Clinical features	Behera et al/1994	Chakrabati et al/2002	Agarwal et al/2007
Patients, No.	35	89	155
Male/female gender, No	14/21	53/35	79/76
Mean age, yr	34.3	36.4	33.4
Mean duration of asthma, yr	11.1	12.1	8.9
History of asthma	94%	90%	100%
Absolute EOS count>500/μL	12/28 (43%)	100%	76.1%
Fleeting shadows	77%	74%	40%
Skin test against Asp (type I)	51%	85%	100%
Elevated IgE levels			100%
Serum precipitins against Aspergillus	77%	71.9%	85.6%
Central bronchiectasis	71%	69%	76.1%

Allergic Bronchopulmonary Aspergillosis (ABPA)- Radiological investigation - CT scan findings

Atelectasis and mucoid impaction



- A. Subsegmental atelectasis.
- B. Hyperattenuated mucus (arrow) with segmental collapse (arrowhead).
- C. High attenuation mucus (arrow) within a collapsed left upper lobe and lingual (arrowhead).
- D. E. left lung collapse which is due to hyperdense mucus (arrow) within the collapsed lung

Allergic Bronchopulmonary Aspergillosis (ABPA) Proposed Radiological staging ISHAM

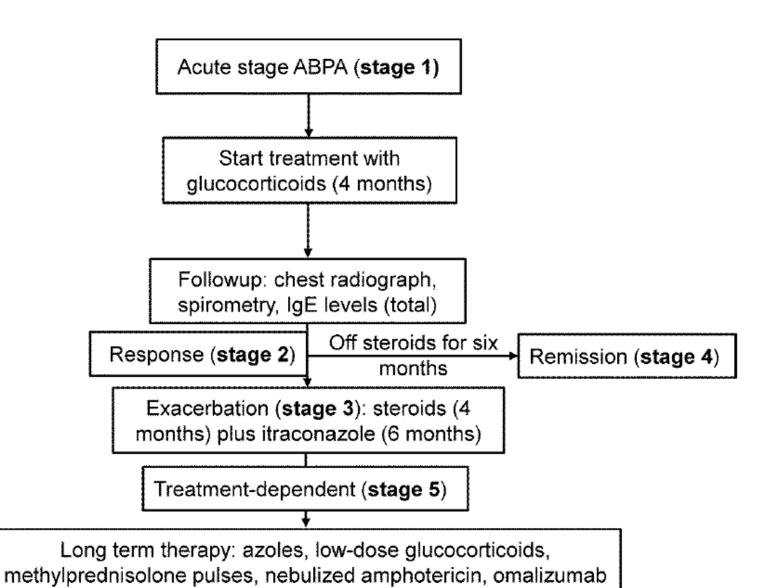
Table 6. Newly proposed radiological classification of ABPA based on computed tomographic (CT) chest
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Classification	Features
ABPA-S (Serological ABPA)	All the diagnostic features of ABPA (Table 4) but no abnormality resulting from ABPA on HRCT chest*
ABPA-B (ABPA with bronchiectasis)	All the diagnostic features of ABPA including bronchiectasis on HRCT chest
ABPA-HAM (ABPA with high- attenuation mucus)	All the diagnostic features of ABPA including presence of high-attenuation mucus
ABPA-CPF (ABPA with chronic pleuropulmonary fibrosis)	ABPA with at least two to three other radiological features such as pulmonary fibrosis, parenchymal scarring, fibro-cavitary lesions, aspergilloma and pleural thickening without presence of mucoid impaction or high-attenuation mucus

^{*}Findings resulting from co-existent disease, bullae from asthma, tracheomalacia, etc. should not be considered while labelling the patients as ABPA-S.

HRCT, high-resolution CT; ABPA, allergic bronchopulmonary aspergillosis.

Allergic Bronchopulmonary Aspergillosis (ABPA) Practical approach to treatment - ISHAM



- ✓ Well controlled asthma, no radiographic abnormalities (stage 0)-- follow up- no treatment
- ✓ Monitor clinical symptoms, chest radiograph and total IgE levels, every eight weeks
- ✓ Response (stage 2) = clinical and/or radiological improvement with at least 25% decline in IgE levels
- Monitor IgE frequently to establish the 'new' baseline level for an individual patient
- ✓ Exacerbation (Stage 3)= ◆ Clinical and/or radiological worsening along with 50% increase in IgE