Pulmonary Involvement of ANCA-associated Vasculitis in Adult Chinese Patients.

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Abstract

Objective: The aim of this study was to clarify the clinical characteristics and long-term outcomes of ANCA-associated vasculitis (AAV) patients with pulmonary involvement from a single Chinese cohort.

Methods: Newly diagnosed AAV patients with pulmonary involvement, as defined by CT, were recruited from January 2010 to June 2020. Clinical data and CT images were collected retrospectively. Baseline CTs were evaluated and re-classified into four categories: interstitial lung disease (ILD), airway involvement (AI), alveolar hemorrhage (AH), and pulmonary granuloma (PG).

Results: A total of 719 patients were newly diagnosed with AAV, 366 (50.9%) of whom combined with pulmonary involvement at baseline. Among the AAV cases with pulmonary involvement, 55.7% (204/366) had ILD, 16.7% (61/366) had AI alone, 14.8% (54/366) had PG, and 12.8% (47/366) had AH alone. During follow-up of a median duration of 42.0 months, 66/366 (18.0%) patients died, mainly died from infections. Survival, relapse, and infection were all significantly different based on the radiological features. Specifically, the ILD group tends to have a poor long-term prognosis, the PG group is prone to relapse, and the AI group is apt to infection. The AH group has a high risk of both early infection and relapse, thus a poor short-term prognosis.

Conclusion: AAV patients with diverse radiological features have different clinical characteristics and outcomes. Therefore, the intensity of immunosuppressive therapy must be carefully valued by considering the baseline CT findings among AAV patients with pulmonary involvement.

1. Background

Antineutrophil cytoplasmic antibody (ANCA)-associated vasculitis (AAV) is a group of disorders characterized by necrotizing inflammation of small- and medium-sized blood vessels and the presence of circulating ANCA. Clinical disease phenotypes include granulomatosis with polyangiitis (GPA), microscopic polyangiitis (MPA), and eosinophilic granulomatosis with polyangiitis (EGPA) [1]. Pauci-immune glomerulonephritis is a seldom type of circulating ANCA-negative AAV. Ethnicity difference was known as MPO-ANCA is predominantly among Asian patients with both GPA and MPA [2].

Pulmonary involvement is frequent and prominent among various manifestation of AAV. Radiologic findings, mainly computed tomography (CT), is key for the identification of pulmonary involvement. Previous studies showed that 52–80% of patients with AAV had pulmonary abnormalities on chest CT [3–5], mainly including interstitial lung disease (ILD), granuloma (nodules, masses, with or without cavitation), alveolar hemorrhage (AH). In most publications, ILD was defined as usual interstitial pneumonia (UIP) or non-specific interstitial pneumonia (NSIP) according to radiologic pattern[6, 7]. MPA is the main AAV subtype associated to ILD [4, 8–10]. Granuloma is more prevalent in GPA[4, 11]. The incidence of AH is 7–45% in GPA and 10–30% in MPA, but is rare in EGPA [12, 13]. Airway involvement (AI) have been reported by some small size of cohorts [11]. However, there is still limited information
focus attention on the association between different radiologic patterns and long-term outcomes among the AAV patients with pulmonary involvement.

On this ground, the aim of this study was to further characterized the relationship between the clinical features and long-term outcomes according to the pulmonary radiologic patterns in a retrospective cohort spanning 10 years with 366 adult Chinese AAV patients with pulmonary involvement. Furthermore, we counted the relapses and infectious events based on different radiologic patterns and identified their predictors.

2. Methods

2.1. Study population

Seven hundred and nineteen patients newly diagnosed as AAV at the Peking University First Hospital, from January 2010 to June 2020, were screened retrospectively. Inclusion criteria were defined as follows: (1) patients fulfilled the Chapel Hill Consensus Conference definition for AAV and the entry criteria required by Watts’ algorithm\[14, 15]. (2) patients with pulmonary involvement, as defined by chest CT. Exclusion criteria were defined as follows: (1) age younger than 18 years; (2) patients with secondary vasculitis or with comorbid autoimmune disease; (3) other causes known to induce CT abnormality; and (4) inadequate clinical data or without baseline CT.

This study was conducted in compliance with the Declaration of Helsinki principles and was approved by the Ethical Committee of Peking University First Hospital. Informed consent was not required for this work because it consists retrospective data, and all treatment decisions were made prior to our evaluation.

2.2. Evaluation of chest CT images

The baseline chest CT images were reviewed independently by two pulmonologists (P.Z and J.M) and one expert pulmonary radiologist (L.G) who were blinded to clinical and laboratory findings. CT scans were obtained with a slice thickness of less than or equal to 1.25 mm at 10- to 20-mm intervals from the apices to the lung bases. All images were obtained using regular-dose CT.

According to CT images, we divided all patients into four groups: patients with ILD (the ILD group), patients with pulmonary granuloma (the PG group), patients with alveolar hemorrhage alone (the AH group), patients with airway involvement alone (the AI group). Of note, airway involvement in this study was defined as involvement of lower respiratory airway including trachea and bronchus, and small airways. Patients with AH at onset and no CT abnormality remained after treatment were also classified into the AH group. Specific abnormal CT findings were classified according to Fleisessenger Society guidelines\[16], including ground glass opacity, reticulation, honeycombing, nodules, masses, cavities, bronchial stenosis, bronchiectasis, tree-in-bud, air trapping, emphysema, pleural thickening. An expert radiologist (L.G) determined the global pattern of abnormalities in ILD patients according to the 2013 idiopathic interstitial pneumonias classification\[17].
2.3. Data collection and definitions

Data were collected from diagnosis and during follow-up until death or June 2020. All ANCA-negative vasculitis was pathologically diagnosed. Disease activity was assessed in accordance with the Birmingham Vasculitis Activity Score (BVAS)\cite{18}. AH was defined as bilateral alveolar infiltrates on radiological imaging without alternative explanation, plus at least one of the following: hemoptysis, unexplained decrease in hemoglobin (>2 g/dL), anemia (hemoglobin <10 g/dL), increased DLCO, or bronchoscopy evidence\cite{19}. Respiratory failure was defined by an arterial oxygen tension (\(P_aO_2\)) of <60 mmHg (8.0 kPa), with or without an arterial carbon dioxide tension (\(P_aCO_2\)) of >50 mmHg (6.0 kPa), when breathing atmospheric air at sea level under resting condition\cite{20}, or the need for any type of mechanical ventilation \cite{21}. Central nervous system (CNS) involvement was defined as new-onset neurologic deficits plus abnormal radiological and cerebrospinal fluid findings without other explainable causes\cite{22}. Infection was recorded if hospitalization was required. Relapse was defined by the recurrence of symptoms of active vasculitis in any organ system after remission is achieved with other causes excluded.

2.4. Statistical analyses

We used an analysis of variance (ANOVA) to assess differences among subject groups, and post hoc comparisons were made using the Bonferroni/Dunn test. Differences of qualitative results were compared using the chi-square test, and post hoc comparisons (Bonferroni correction) were performed to detect differences between the groups. Multivariable Cox regression analysis were performed to evaluate the predictors of infection, relapse and mortality. Variables associated with a given outcome at a p value<0.1 in Kaplan-Meier analysis (categorical variables) or univariable Cox regression (continuous variables) were entered into the final multivariable model. Results were expressed as hazard ratios with 95% confidence intervals. The difference was considered significant if the p-value < 0.05. Analysis was performed with SPSS software (v 27.0; SPSS, Chicago, IL, USA).

3. Results

3.1. Clinical characteristics

Three hundred and sixty-six patients newly diagnosed AAV with pulmonary involvement were eventually enrolled (Figure 1). The mean age at onset was 66[IQR 58-72] years old. Two hundred and ninety-nine (81.7%) patients were MPA, 60(16.4%) were GPA and 7(1.9%) were EGPA. Three hundred and fifty-eight (97.8%) patients were ANCA positive; 299(81.7%) were MPO-ANCA positive, 44(12.0%) were PR3-ANCA positive, and 11(3.0%) were double positive. We divided the AAV patients into four groups: the ILD group (n=204), the PG group (n=54), the AH group (n=47) and the AI group (n=61).

The baseline clinical characteristics and laboratory findings are summarized in Table 1. The mean age was significantly higher in the ILD group compared with the other three groups. The incidence of cardiovascular complication was also higher in the ILD group compared to the PG group. The MPO-ANCA
was predominant in all the four groups. In the comparison among the four groups, the positive rate of MPO-ANCA was higher in the ILD and AI groups than in the AH and PG groups, and PR3-ANCA was detected more in the PG and AH groups than in the other two groups, respectively.
Table 1
Clinical characteristics and laboratory findings of patients with different radiological patterns.

| Characteristic                              | Alveolar hemorrhage (n=47) | Interstitial lung disease (n=204) | Pulmonary granuloma (n=54) | Airway involvement (n=61) |
|--------------------------------------------|----------------------------|----------------------------------|---------------------------|---------------------------|
| Age, median (IQR) (years)                  | 56(47-64)                  | 68(62-75) ^a,b,c                 | 59(53-69)                 | 61(58-68)                 |
| Sex (male), no. (%)                        | 21(44.7)                   | 110(53.9)                        | 28(51.9)                  | 24(39.3)                  |
| Ever smoker, no. (%)                       | 13(27.8)                   | 85(41.7)                         | 19(35.2)                  | 17(27.9)                  |
| MPA, no. (%)                               | 36(76.6)                   | 187(91.7) ^a                     | 25(46.3) ^a,d,e           | 51(83.6)                  |
| GPA, no. (%)                               | 11(23.4)                   | 14(6.9) ^b                       | 26(48.1) ^a,d,e           | 9(14.8)                   |
| EGPA, no. (%)                              | 0                          | 3(1.5)                           | 3(5.6)                    | 1(1.6)                    |
| Time of onset to diagnosis, median (IQR) (months) | 1(0-2)                  | 2(0-8)                           | 1(0-4)                    | 1(0-8)                    |
| Follow-up, median (IQR) (months)           | 39(17-64)                  | 40(17-75)                        | 52(26-72)                 | 42(19-84)                 |
| **Comorbidities, no. (%)**                 |                            |                                  |                           |                           |
| Cardiovascular event ^1,                   | 18(38.3)                   | 112(54.9) ^a                     | 18(33.3)                  | 24(39.3)                  |
| Malignancy                                 | 1(2.1)                     | 4(2.0)                           | 0                         | 3(4.9)                    |
| **Organ involvement at diagnosis**         |                            |                                  |                           |                           |
| Fever, no. (%)                             | 23(48.9)                   | 74(36.3)                         | 22(40.7)                  | 28(45.9)                  |
| Weight loss (≥2 kg), no. (%)               | 18(38.3)                   | 95(46.6)                         | 24(44.4)                  | 32(52.5)                  |
| Cutaneous, no. (%)                         | 5(10.6)                    | 9(4.4)                           | 5(9.3)                    | 2(3.3)                    |
| Ear, nose and throat, no. (%)              | 14(29.8)                   | 57(27.9) ^a                      | 26(48.1)                  | 24(39.3)                  |
| Eye, no. (%)                               | 6(12.8)                    | 10(4.9) ^a                       | 10(18.5)                  | 7(11.5)                   |
| Cardiovascular, no. (%)                    | 2(4.3)                     | 4(1.2)                           | 1(1.9)                    | 2(3.3)                    |
| Gastrointestinal, no. (%)                  | 2(4.3)                     | 3(1.5)                           | 3(5.6)                    | 2(3.3)                    |
| Central nervous system, no. (%)            | 3(6.4)                     | 5(2.5)                           | 4(7.4)                    | 3(4.9)                    |
| Peripheral nervous system, no. (%)         | 3(6.4)                     | 34(16.7)                         | 10(18.5)                  | 13(21.3)                  |
| Characteristic                                      | Alveolar hemorrhage (n=47) | Interstitial lung disease (n=204) | Pulmonary granuloma (n=54) | Airway involvement (n=61) |
|---------------------------------------------------|-----------------------------|-----------------------------------|---------------------------|---------------------------|
| Renal 2, no. (%)                                  | 46(97.9)                    | 165(80.9)                         | 44(81.5)                  | 52(85.2)                  |
| Renal insufficiency 3, no. (%)                    | 20(42.6)                    | 50(24.5)                          | 18(33.3)                  | 21(34.4)                  |
| Laboratory findings at diagnosis                  |                             |                                   |                           |                           |
| MPO, no. (%)                                       | 33(70.2)                    | 179(87.7)                         | 34(63.0)                  | 53(86.9)                  |
| PR3, no. (%)                                       | 11(23.4)                    | 12(5.9)                           | 17(31.5)                  | 4(6.6)                    |
| Double positive, no. (%)                          | 0                           | 9(4.4)                            | 0                         | 2(3.3)                    |
| ANCA negative, no. (%)                            | 2(4.3)                      | 3(1.5)                            | 2(3.7)                    | 1(1.6)                    |
| Serum creatinine, mean±SD, µmol/L                 | 424.1±312.9                 | 323.6±265.3                       | 315.1±249.4               | 398.0±355.7               |
| C-reactive protein, mean±SD, mg/L                 | 54.4±53.9                   | 45.9±49.7                         | 55.3±52.0                 | 45.1±50.0                 |
| BVAS at onset, median (IQR)                       | 20(17-24)                   | 17(14-21)                         | 20(15-25)                 | 20(14-24)                 |
| Five-Factor Score, no. (%)                        |                             |                                   |                           |                           |
| 0                                                 | 5(10.6)                     | 60(29.4)                          | 15(27.8)                  | 19(31.1)                  |
| 1                                                 | 9(19.1)                     | 53(26.0)                          | 9(16.7)                   | 17(27.9)                  |
| ≥2                                                | 33(70.2)                    | 91(44.6)                          | 28(51.9)                  | 25(41.0)                  |
| Bronchoalveolar lavage results at diagnosis, no. (%) |                             |                                   |                           |                           |
| Lymphocytes count, mean ± SD                      | 8.7±8.0                     | 9.4±11.1                          | 10.8±14.8                 | 5.8±7.7                   |
| Lymphocytes >20%, no. (%)                         | 2/18(11.1)                  | 7/59(11.9)                        | 2/20(10.0)                | 1/15(6.7)                 |
| Neutrophils count, mean ± SD                      | 43.9±32.6                   | 44.3±30.7                         | 32.8±33.1                 | 62.1±31.0                 |
| Neutrophils >5%, no. (%)                          | 16/18(88.9)                 | 55/59(93.2)                       | 17/20(85.0)               | 14/15(93.3)               |
| Hemosiderin-laden macrophages count, mean ± SD     | 60.2±26.8                   | 12.7±20.4                         | 37.8±31.7                 | 5.1±17.7                  |
| Hemosiderin-laden macrophages ≥20%, no. (%)       | 16/18(88.9)                 | 15/59(25.4)                       | 12/20(60)                 | 1/15(6.7)                 |
| Characteristic                                      | Alveolar hemorrhage (n=47) | Interstitial lung disease (n=204) | Pulmonary granuloma (n=54) | Airway involvement (n=61) |
|-----------------------------------------------------|-----------------------------|----------------------------------|---------------------------|---------------------------|
| Pulmonary function test results at diagnosis, no. (%)|                             |                                  |                           |                           |
| Obstruction, no. (%)                                | 1/9 (11.1)                  | 7/66 (10.6) \(^a\)              | 9/19 (47.4)               | 4/12 (33.3)               |
| Restriction, no. (%)                                | 4/9 (44.4)                  | 43/66 (65.2)                     | 7/19 (36.8)               | 8/12 (66.7)               |
| FEV1/FVC, mean ±SD                                  | 74.9±9.3                    | 80.0±8.6 \(^a,c\)               | 70.1±12.3                 | 66.1±14.0                 |
| FEV1% pred, mean ±SD                                | 88.1±27.7                   | 83.4±20.4                       | 79.4±21.8                 | 70.1±24.2                 |
| FVC% pred, mean ±SD                                 | 98.5±27.3                   | 79.6±23.8 \(^b\)               | 86.9±23.1                 | 80.8±14.7                 |
| TLC% pred, mean ±SD                                 | 94.7±25.9                   | 73.7±17.6 \(^b\)               | 85.4±19.5                 | 79.5±15.5                 |
| FEF 25-75% pred, mean ±SD                           | 88.1±4.0                    | 62.8±25.5                       | 38.4±30.3                 | 23.78±11.1 \(^c,f\)      |
| FEF50% pred, mean ±SD                                | 66.3±33.6                   | 75.3±34.5 \(^a\)               | 50.0±35.8                 | 38.5±21.8 \(^c\)         |
| FEF25% pred, mean ±SD                                | 54.5±22.8                   | 69.8±44.5                       | 45.4±30.9                 | 42.1±18.7                 |
| DLCO % pred, mean ±SD                                | 65.1±28.8                   | 51.6±17.1                       | 66.4±21.9                 | 57.3±16.7                 |
| Treatment, no. (%)                                   |                             |                                  |                           |                           |
| Induction                                           |                             |                                  |                           |                           |
| Glucocorticoids pulses                              | 32 (68.1)                   | 78 (38.2) \(^b\)               | 23 (42.6)                 | 33 (54.1)                 |
| Immunosuppressant                                   | 41 (87.2)                   | 147 (72.1)                      | 37 (68.5)                 | 47 (77.0)                 |
| Cyclophosphamide                                    | 38 (80.9)                   | 142 (69.6)                      | 35 (64.8)                 | 45 (73.8)                 |
| Rituximab                                           | 0                           | 1 (0.5)                         | 1 (1.9)                   | 0                         |
| Plasma exchanges                                    | 22 (46.8)                   | 48 (23.5) \(^b\)               | 17 (31.5)                 | 18 (29.5)                 |
| Dialysis                                            | 21 (44.7)                   | 50 (24.5) \(^b\)               | 17 (31.5)                 | 21 (34.4)                 |
| Maintenance                                         |                             |                                  |                           |                           |
| Immunosuppressant                                   | 21/21 (100.0)               | 99/105 (94.3)                   | 36/37 (97.3)              | 28/32 (87.5)              |
| Cyclophosphamide                                    | 21/21 (100.0)               | 95/105 (90.5)                   | 34/37 (91.9)              | 27/32 (84.3)              |
| Rituximab                                           | 4/21 (19.0)                 | 3/105 (2.9) \(^b\)             | 5/37 (13.5)               | 2/32 (6.3)                |
| Azathioprine                                        | 3/21 (14.3)                 | 28/105 (26.7)                   | 9/37 (24.3)               | 5/32 (15.6)               |
| Characteristic                   | Alveolar hemorrhage (n=47) | Interstitial lung disease (n=204) | Pulmonary granuloma (n=54) | Airway involvement (n=61) |
|---------------------------------|-----------------------------|----------------------------------|---------------------------|---------------------------|
| Mycophenolate mofetil          | 0                           | 8/105 (7.6)                      | 2/37 (5.4)                | 2/32 (6.3)                |

AAV: ANCA-associated vasculitis; AI: airway involvement; ANCA: antineutrophil cytoplasm antibodies; BVAS: Birmingham Vasculitis Activity Scores; DLCO: diffusing capacity for carbon monoxide; EGPA: eosinophilic granulomatosis with polyangiitis; FEF<sub>25–75</sub> = forced expiratory flow at 25–75% of FVC; FEF<sub>50</sub> = forced expiratory flow at 50% of FVC; FEF<sub>25</sub> = forced expiratory flow at 50% of FVC; FEV1: forced expiratory volume; FVC: forced vital capacity; GPA: granulomatosis with polyangiitis; IQR: interquartile range; MPA: microscopic polyangiitis; no: number; MPO: myeloperoxidase; PR3: proteinase-3; SD: standard deviation; TLC: total lung capacity; % pred: % predicted.

1 Cardiovascular event was defined as the presence of one or more of the following conditions at the time of AAV diagnosis: ischemic heart disease, congestive heart failure, hypertension, stroke, thromboembolism.

2 Renal involvement was defined as the presence of hematuria (≥10 red blood cells per high power field), proteinuria (≥500 mg/24 h), rise in serum creatinine >30%, biopsy-proven glomerulonephritis, or need to initiate renal replacement therapy all attributable to active vasculitis.

3 Renal insufficiency was defined as serum creatinine level ≥500µmol/L or undergoing dialysis.

p< 0.05: a ILD vs. PG, b ILD vs. AH, c ILD vs. AI, d PG vs. AH, e PG vs. AI, f AH vs. AI

Patients in the PG group were more commonly associated with systemic AAV manifestations, such as ENT and eye symptoms compared to the ILD group. No significant difference regarding the prevalence of renal involvement and renal insufficiency between four groups. The ILD group had a significantly lower BVAS compared to the PG and AH groups, and the AH group had a higher Five Factor Score than the ILD and AI groups at baseline.

Bronchoalveolar lavage was performed in 112(30.6%) patients. The neutrophils count was significantly higher in the AI group compared to the PG group. The hemosiderin cells count was significantly higher in the AH group than other three groups, and also higher in the PG group compared to the ILD and AI groups.

A total of 101 patients had pulmonary functional tests at baseline. The proportion of obstructive ventilation dysfunction was higher in the PG group compared to the ILD group. There were no significant differences in the proportion of restrictive ventilation dysfunction. However, the mean value of FVC% pred and TLC% pred were both lower in the ILD group than the PG group. Intriguingly, surrogate markers of small airway disease, such as FEF<sub>25–75</sub> and FEF<sub>50</sub>, were significantly decreased in the AI group. FEF<sub>50</sub> was also decreased in the PG group compared to the ILD group.

There were no significantly differences in the uses of immunosuppressants for induction therapy. However, the percentage of use of glucocorticoids pulse, plasma exchange and dialysis were higher in the
AH group than the ILD group. The percentage of use of rituximab was also higher in the AH group than the ILD group during maintenance therapy.

Ninety-nine (27.0%) patients occurred respiratory failure during follow-up, 63.6% (63/99) of whom from the ILD group, 8.1% (8/99) of whom from the PG group, 16.2% (16/99) from the AH group and 12.1% (12/99) from the AI group.

3.2. Prevalence of lung abnormalities on chest CT images

The CT findings and comparison are summarized in Supplementary Table 1. Common CT abnormalities included ground-glass opacity (53.3%), honeycombing (39.9%), bronchiectasis (18.6%), air trapping (16.7%), bronchial stenosis (14.2%), reticulation (12.8%) and tree-in-bud (12.0%).

Comparing between four groups, honeycombing and reticulation was more frequent in the ILD group than other three groups. The PG group was manifested more frequently as nodules, masses and cavities. It is noteworthy that airway lesions occurred among all groups. Airway-related abnormalities, including bronchiectasis, bronchial stenosis, tree-in-bud and air trapping were all significantly common in the AI group. Moreover, air trapping and bronchial stenosis were both more frequent in the PG group than ILD group. Air trapping was more common in the AH group compared to the ILD group.

Of the 204 patients in the ILD group, UIP was the majority (71.6%), NSIP was noted in 39(19.1%), OP was noted in 3(1.5%), and unclassified IP was noted in 16(7.8%).

3.3. Predictors of mortality

For the whole cohort, infection was the most prevalent cause of death (39/66,59.1%), active vasculitis only accounted for 19.7% (13/66) of the death. There were no statistically differences on the causes of death between four groups (Supplementary Table 2).

During follow-up of a median duration of 42.0 months, 66(18.0%) patients eventually died. The cumulative survival rate of overall patients at 1, 2, and 5 years was 90.4% (95%CI 88.9%- 91.9%), 87.5% (95%CI 85.7%-89.3%) and 84.3% (95%CI 82.2% -86.4%), respectively (Figure 2). Survival was significantly different based on the radiologic features. One- and 5-year overall survival rates, respectively, were: 89.2% and 84.8% for ILD, 96.7% and 83.3% for AI, 85.1% and 83.0% for AH, 92.6% and 90.7% for PG (Figure 3A, p=0.033).

In the multivariable Cox regression analysis, the independent predictors of all-cause mortality were UIP pattern, age≥ 65 years at diagnosis, respiratory failure, infection requiring hospitalization and AH (Table 2).
Table 2
Cox regression analysis of predictors for all-cause mortality in patients with ANCA-associated vasculitis with pulmonary involvement.

| Radiological pattern | Univariate | Multivariate |
|----------------------|------------|--------------|
|                      | Hazard ratio (CI 95%) | p value | Hazard ratio (CI 95%) | p value |
| Interstitial lung disease | 3.642(1.318-10.066) | 0.013 | 3.369(1.922-5.906) | <0.001 |
| UIP pattern           | 2.548(1.515-4.284) | <0.001 | 3.369(1.922-5.906) | <0.001 |
| Pulmonary granuloma   | 0.484(0.209-1.122) | 0.091 |           |         |
| Alveolar hemorrhage   | 3.028(1.757-5.218) | <0.001 | 2.658(1.491-4.738) | 0.001 |
| Airway involvement    | 1.806(0.893-3.650) | 0.100 |           |         |
| Age≥65 years at diagnosis | 2.231(1.237-4.024) | 0.008 | 2.118(1.151-3.896) | 0.016 |
| Sex (female)          | 1.156(0.686-1.947) | 0.587 |           |         |
| MPA                   | 1.588(0.752-3.354) | 0.226 |           |         |
| GPA                   | 0.619(0.280-1.365) | 0.234 |           |         |
| MPO-ANCA              | 2.349(0.938-5.883) | 0.068 |           |         |
| PR3-ANCA              | 0.756(0.302-1.894) | 0.551 |           |         |
| Cardiovascular comorbidities | 1.240(0.737-2.086) | 0.418 |           |         |

| Organ involvement at onset | Univariate | Multivariate |
|----------------------------|------------|--------------|
| Renal                      | 1.733(0.742-4.048) | 0.204 |           |         |
| Ear, nose and throat       | 0.521(0.285-0.954) | 0.035 |           |         |
| Central nervous system     | 2.840(1.217-6.629) | 0.016 |           |         |
| Peripheral nervous system  | 1.213(0.641-2.293) | 0.553 |           |         |
| Cardiovascular             | 4.646(1.131-19.085) | 0.033 |           |         |
| Gastrointestinal           | 2.830(1.024-7.823) | 0.045 |           |         |
| Eye                        | 0.596(0.215-1.647) | 0.318 |           |         |

ANCA: antineutrophil cytoplasm antibodies; BVAS: Birmingham Vasculitis Activity Scores; FFS: five factors score; GPA: granulomatosis with polyangiitis; IS: immunosuppressant; MPA: microscopic polyangiitis; MPO: myeloperoxidase; PR3: proteinase-3; UIP: usual interstitial pneumonia.
|                                | Univariate               | Multivariate             |
|--------------------------------|--------------------------|--------------------------|
| Respiratory failure            | 12.425(7.036-21.944)     | <0.001                   |
|                                |                          | 7.077 (3.731-13.423)     | <0.001                   |
| Initial serum creatine         | 1.001(1.000-1.002)       | 0.082                    |
| Renal insufficiency            | 1.518(0.892-2.583)       | 0.123                    |
| BVAS at onset                  | 1.014(0.989-1.041)       | 0.278                    |
| **FFS (ref. = 0)**             |                          |                          |
| FFS=1                          | 1.297(0.758-2.219)       | 0.342                    |
| FFS≥2                          | 1.112(0.647-1.913)       | 0.700                    |
| Infection requiring hospitalization | 6.533(2.977-14.337)     | <0.001                   |
|                                |                          | 2.529(1.042-6.137)       | 0.040                    |
| Relapse                        | 1.010(0.519-1.968)       | 0.976                    |
| IS for induction therapy       | 0.617(0.365-1.043)       | 0.071                    |
| IS for maintenance therapy     | 1.560(0.710-3.430)       | 0.269                    |

ANCA: antineutrophil cytoplasm antibodies; BVAS: Birmingham Vasculitis Activity Scores; FFS: five factors score; GPA: granulomatosis with polyangiitis; IS: immunosuppressant; MPA: microscopic polyangiitis; MPO: myeloperoxidase; PR3: proteinase-3; UIP: usual interstitial pneumonia.

Given that the high proportion of ILD in our cohort, and the decrease of FVC and DLCO were reported as independent risk factors for mortality in connective tissue disease-related-ILD patients[23], we conducted survival analysis of the subgroup of cases with pulmonary functional results (Supplementary Table 3). The only factor significantly associated with shorter survival was DLCO decreased (HR 0.970, p=0.019), and ROC curve analysis showed that the best cut-off value was 54.05% pred, with sensitivity and specificity of 66.7% and 63.4%, respectively (Supplementary Figure 1).

3.4. Predictors of relapse

During the follow up, 111(30.3%) patients experienced 186 relapses, all summarized in Table 3. The median durations from diagnosis to the first relapse was 12[IQR 5-26] months. Pulmonary (59.7%) was the most prevalent organ involvement at the time of relapse, followed by kidney (48.4%).
|                                | Alveolar hemorrhage (n=47) | Interstitial lung disease (n=204) | Pulmonary granuloma (n=54) | Airway involvement (n=61) |
|--------------------------------|-----------------------------|-----------------------------------|-----------------------------|---------------------------|
| Relapse cases, no (%)          | 15/47 (31.9)                | 55/204 (27.0)                     | 26/54 (48.1)                | 15/61 (24.6)              |
| Relapse events, no (%)         | 30                          | 89                                | 47                          | 20                        |
| First relapsing time, median (IQR) (months) | 13 (5-36)          | 12 (7-27)                         | 9 (5-23)                    | 9 (6-21)                  |
| **Organ involvement at relapse, no. (%)** |                             |                                   |                             |                           |
| Pulmonary                      | 21/30 (70.0)                | 48/89 (53.9)                      | 35/47 (74.5)                | 8/20 (40.0)               |
| Renal                          | 15/30 (50.0)                | 52/89 (58.4)                      | 12/47 (25.5)                | 11/20 (55.0)              |
| General ^1                     | 2/30 (6.7)                  | 12/89 (13.5)                      | 12/47 (25.5)                | 1/20 (5.0)                |
| Ear, nose, and throat          | 2/30 (6.7)                  | 2/89 (2.2)                        | 4/47 (8.5)                  | 6/20 (30.0) ^f            |
| Central nervous system         | 0                           | 2/89 (2.2)                        | 9/47 (19.1)                 | 0                         |
| Peripheral nervous system      | 2/30 (6.7)                  | 3/89 (3.4)                        | 3/47 (6.4)                  | 0                         |
| Gastrointestinal               | 1/30 (3.3)                  | 1/89 (1.1)                        | 3/47 (6.4)                  | 3/20 (15.0)               |
| Eye                            | 2/30 (6.7)                  | 5/89 (5.6)                        | 3/47 (6.4)                  | 1/20 (5.0)                |

^1 General included fever or weight loss (≥ 2 kg).

AH: Alveolar hemorrhage; AI: Airway involvement; ILD: interstitial lung disease; PG: pulmonary granuloma;

p < 0.05: ^a ILD vs. PG, ^d PG vs. AH, ^e PG vs. AI, ^f AH vs. AI

The percentage of relapse cases was significantly higher in the PG group than the AI group. Comparing organ involvements at relapse, pulmonary relapse was more common in the PG group than AI group. Renal relapse was more common in the ILD group compared to the PG group. ENT relapse was more common in the AI group compared to the ILD group. CNS relapse was the most common in the PG group.

Relapse free-survival rates between four groups were significantly different, patients with PG and AH were more likely to relapse (Figure 3B, p = 0.011). PR3-ANCA and CNS involvement were predictors of
relapse in the multivariate Cox regression analysis (Supplementary Table 4). Moreover, we analyzed predictors of relapse in each group. Female, PR3-ANCA and CNS involvement were the predictors of relapse in the PG group. ENT involvement was the predictor of relapse in the AI group. Multivariate analysis did not yield relevant clinical predictor of relapse in the I LD and AH groups (Supplementary Table 5-8).

Lastly, we investigated factors associated with pulmonary relapse, the leading organ involved at relapse. PG pattern and CNS involvement were risk factors of pulmonary relapse, but age $\geq 65$ years was seen to be a protective factor (Supplementary Table 9).

3.5. Predictors of infection

Since infection was the leading cause of death in AAV patients with pulmonary involvement and also a significant factor independently associated with shorter survival, we further collected the infection-related data. There were total of 347 infections in 237 patients during the follow-up. The median durations from diagnosis to the first infection was 0[IQR 0-3] month. As shown in Table 4, pulmonary infection was the leading infection in each group. Genitourinary infection took second place.
Table 4
Details of infections during follow-up.

|                         | Alveolar hemorrhage (n=47) | Interstitial lung disease (n=204) | Pulmonary granuloma (n=54) | Airway involvement (n=61) |
|-------------------------|-----------------------------|----------------------------------|---------------------------|---------------------------|
| **Infectious patients, no. (%)** | 34(72.3)                    | 105(51.5)                        | 43(79.7)                  | 55(90.2) c                |
| **First in-patient infection time, median (IQR)(months)** | 0(0-2)                      | 0(0-4)                           | 0(0-3)                    | 0(0-2)                    |
| **Infectious episodes, no. (%)** | 53                          | 147                              | 60                        | 87                        |
| **Infection pathogens, no. (%)** |                             |                                  |                           |                           |
| Bacteria                | 48/53(90.6)                 | 138/147(93.8)                    | 54/60(90.0)               | 82/87(94.3)               |
| Fungus                  | 20/53(37.7)                 | 38/147(25.9)                     | 15/60(25.0)               | 20/87(23.0)               |
| Pneumocystis jirovecii  | 0                           | 8/147(5.4)                       | 3/60(5.0)                 | 3/87(3.4)                 |
| Aspergillus             | 3/53(5.7)                   | 6/147(4.1)                       | 6/60(6.7)                 | 3/87(3.4)                 |
| Candida                 | 12/53(22.6)                 | 24/147(16.3)                     | 8/60(13.3)                | 16/87(18.4)               |
| Viral                   | 6/53(11.3)                  | 17/147(11.6)                     | 6/60(10.0)                | 4/87(4.6)                 |
| CMV                     | 2/53(3.8)                   | 6/147(4.1)                       | 6/60(10.0)                | 1/87(1.1)                 |
| EBV                     | 1/53 (1.9)                  | 4/147(2.7)                       | 4/60(6.7)                 | 0                         |
| Influenza virus         | 0                           | 2/147(1.4)                       | 0                         | 1/87(1.1)                 |
| **Locations of infection, no. (%)** |                             |                                  |                           |                           |
| Pulmonary               | 50/53(94.3)                 | 135/147(91.8)                    | 57/60(95.0)               | 85/87(97.7)               |
| Genitourinary           | 1/53 (1.9)                  | 17/147(11.7)                     | 2/60(3.3)                 | 7/87(8.0)                 |
| Gastrointestinal        | 4/53(7.5)                   | 5/147(3.4)                       | 2/60(3.3)                 | 2/87(2.3)                 |
| Catheter-associated     | 1/53 (1.9)                  | 4/147(2.7)                       | 1/60(1.7)                 | 2/87(2.3)                 |
| Sepsis                  | 1/53 (1.9)                  | 8/147(5.4)                       | 2/60(3.3)                 | 0                         |

AH: Alveolar hemorrhage; AI: Airway involvement; CMV: cytomegalovirus; EBV: Epstein-Barr virus; ILD: interstitial lung disease; PG: pulmonary granuloma;

p< 0.05: c ILD vs. AI

AAV: ANCA-associated vasculitis; AH: alveolar hemorrhage; AI: airway involvement; CT: computed tomography; CTD: connective tissue disease; ILD: interstitial lung disease; PG: pulmonary granuloma.
Regarding the pathogens of infection, bacterium was the most common pathogen (92.8%), followed by fungus and virus (26.8%, and 9.2%, respectively). Candida was the most frequent fungal infection (17.3%), followed by aspergillus (5.2%). Pneumocystis jirovecii was found in 14 cases (4.0%), 8 from ILD group and 3 from PG and AI group respectively, leading to 9 deaths in total (6 in the ILD group, 1 in the PG group and 2 in the AI group). Cytomegalovirus (4.3%) ranked first among viral pathogens, followed by Epstein-Barr virus and influenza virus (2.6%, and 0.9%, respectively).

The percentage of infectious patients in the AI group was higher than that in the ILD group. However, there were no significant differences regarding to the infection pathogens and locations between four groups.

The infection-free survival curve was especially steep within the first 12 months, especially the first 3 months. Three- and twelve-month infection-free survival were 55.7% and 48.4%. There was a significant difference on the infection-free survival between four groups (Figure 3C, p<0.001). Multiple Cox regression analysis showed that the following factors were independent predictors of secondary infections: respiratory failure, AI, AH, and glucocorticoids pulse for induction therapy (Supplementary Table 10). Immunosuppressants for both induction and maintenance therapy did not significantly associated with infection.

4. Discussion

To the best of our knowledge, this is currently the largest Chinese cohort focused on AAV-related pulmonary involvement with a long follow-up period and is therefore representative. We only enrolled the AAV patients with pulmonary involvement in this cohort to comprehensively clarify the clinical characteristics of these patients and if different radiological subgroups had differences in clinical features, survival, the risk of both relapse and secondary infection.

Our study demonstrates the clinical characteristics of patients with AAV-related lung involvement. Patient in the ILD group are the oldest and have more comorbid cardiovascular events (CVE). MPO-ANCA dominates in all the four groups. BVAS is lower in the ILD group and FFS is higher in the AH group. Considering the bronchoalveolar lavage results, neutrophils count is higher in the AI group and hemosiderin-laden macrophages count is higher in the AH group. Pulmonary function results suggest small airway obstruction was not rare among AAV patients, especially in the PG and AI subgroups. As for treatment, most patients used the standard regimen of glucocorticoid combined with immunosuppressants for induction, while the proportion of glucocorticoids pulses, plasma exchange, dialysis for induction and rituximab for maintenance are higher in the AH group.

Firstly, our cases are older compared to the previous European studies and roughly the same age as other Asian cohorts, but these studies only focused on AAV patients with renal involvement[24–27]. Comparing to the studies focusing on AAV-related lung involvement, similar characteristics are also observed, probably due to a higher proportion of ILD, especially UIP pattern, in Asian AAV patients [5, 28, 29]. Secondly, the ILD group had a lower BVAS, which was in contrast to a previous Japanese study[29]. This
indicated the vasculitis activities in AAV-related ILD patients are relatively low at diagnosis, careful dosing of immunosuppressant is called for induction treatment by weighing the infectious side effect. Finally, this is the first study to document the bronchoalveolar lavage and pulmonary function results of patients with AAV-related pulmonary involvement in detail, confirming that these results differ based on the radiological images, and that DLCO might partially predict patient outcomes. In summary, we observe that patients with different types of lung involvement have their unique clinical characteristics.

As for radiological features, our study highlights the importance of AI in AAV patients, and further determines the global pattern of abnormalities in ILD patients. In our cohort, 16.7% of patients (AI group) presented with lower respiratory tracts involvement alone, which is second only to the ILD. However, a study by Yamagata, et al. showed that airway lesions was as common as ILD, accounting for 66%[30]. This discrepancy may be explained by the fact that our study excluded ILD-related traction bronchiectasis as well as airway lesions due to other comorbidities. A few studies have focused on the association between AI and AAV, suggesting that AI is a comorbidity rather than one of the pulmonary manifestation of AAV[31, 32]. Conversely, some patients with AI substantially improved after the initial immunosuppressive treatment in our study, therefore we tend to consider AI as a part of systemic manifestation. Specifically, AAV-related AI is more frequent characterized by small airway disease.

Pulmonary functional results further confirm these radiologic findings, since FEF_{25-75} and FEF_{50}, widely regarded as markers of small airways obstruction, are significantly decreased in the AI and PG groups. Recent data suggest that airway could be a site of initiation of immune response in AAV, which might correlate with neutrophils activation[32, 33]. Correspondingly, our study reports that neutrophils count is relatively higher in bronchoalveolar lavage fluids of patients from AI group.

Another innovation point lies in the careful classification of all ILD patients. Our data show that UIP account for 39.9% of the total patients with AAV-related lung involvement. As for the ILD subgroup, UIP account for 71.6%, which is significantly higher than the previous studies (38%-57%) [29, 34, 35]. NSIP is found in only 19.1% of ILD patients, which is much lower than a European study[6]. Meanwhile, 3 cases presented as OP, 2 of them with MPO-ANCA positive and 1 with PR3-ANCA positive, which is a rare and unique imaging manifestation of AAV, with only a few previous cases reported[36–38]. Hence, our study suggests that the imaging findings of AAV can be diverse and OP can be a rare manifestation of AAV related ILD.

Although pulmonary involvement is known as a predictor of poor prognosis of AAV, the patients in our pulmonary-involvement cohort seemed have better prognosis than previous reports[39, 40]. The median duration of follow-up of our study was 42.0 months and 66 (18.0 %) deaths were recorded. The 1-year and 5-year survival rate are 90.4% and 84.3% respectively. Previous study mostly by nephrologists and rheumatologists reported that the cumulative survival was 77.4% - 88% and 61.9%-78% at 1 and 5 years [39, 40]. As same as Flossmann's report [43], infection and active vasculitis was the main cause of the 1-year death in our cohort. Furthermore, we also found the independent predictors of all-cause mortality during long-term follow-up in AAV patients with pulmonary involvement were UIP pattern, age≥65 years at diagnosis, respiratory failure, infection needing hospitalization and AH, but not including initial renal
function, higher BVAS, lower hemoglobin, and higher white cell count in other publications[39, 40]. This may be partly explained by the specificity of our cohort. Firstly, the patients without lung involvement were excluded, that might remove some patients with other important organ injuries and high BVAS, such as those with more severe renal failure. Secondly, we enrolled the patients in the last decade with more standardized treatment and biological agents which gave them more opportunities.

Survival is significantly different based on the radiologic features. ILD, especially UIP pattern, and AH groups tend to have poor prognosis. AH is associated with early prognosis whereas ILD is related to long-term outcome, which is in line with a previous Japanese study [28]. Our cohort is a representative Chinese cohort with updated treatment and longer follow-up duration than that in previous study. In addition to the ILD patients who died from respiratory failure, some also died from CVE. A recent meta-analysis reported that excess mortality due to CVE ranges from about 2 to 4-fold in AAV and is higher in the elderly[41]. Patients with Al have better early prognosis but a decreasing trend in long-term outcome, probably due to an increasing risk of infection during follow-up. Therefore, carefully identifying the baseline chest imaging characteristics of AAV patients can predict their prognosis.

Relapse is another major concern. Pulmonary involvement is regarded as a risk factor of relapse[1]. However, there is no study pay attention to the specific relapse events among these patients. Beyond that, the value of measuring ANCA to predict a relapse is doubtable. Kemna et al. reported that an ANCA rise was related to a relapse only in patients with renal involvement[42]. Our study reveals that AAV patients with lung involvement have significantly higher risk of relapse than previously reported in patients with renal involvement[27]. Comparing between subgroups, relapse is more frequent in the PG and AH groups and less common in the ILD and Al groups. Regarding to the relapse organs, patients in the PG and AH groups are more likely to have pulmonary relapse, whereas patients in the ILD and Al groups are more prone to renal relapse. We further identify that PR3-ANCA positive and CNS involvement are as risk factors of relapse in AAV-related lung involvement patients. Moreover, predictors of relapse are distinct in each imaging subgroups. Of note, PG pattern is an independent risk factor for pulmonary relapse. This is the first study to elaborate on the relapse of AAV-related lung involvement patients and to demonstrate that there is a correlation between relapse and imaging patterns.

Infection was the leading cause of death in AAV patients[43, 44], and underlying pulmonary involvement was regarded as an independent predictor of secondary infection[40]. In the present study, more than half of the patients died from infection. Three- and twelve-month infection-free survival are 55.7% and 48.4%, which is significantly higher than a previous Chinese study focusing on AAV patients with renal involvement[40]. Regarding the infectious location, pulmonary accounts for more than 90% of the infection. Fungal and viral infections are not uncommon due to active immunosuppressive therapy. Multivariate analysis reveals that AH and Al are both independently related to higher risk of infection. While Al patients in this study are less prone to relapse. Therefore, immunosuppressive therapy strategies must keep into account not only AAV activity, but patient’s radiological characteristics and their risk of infection as well.
There were some limitations in our study. On the one hand, ours is a retrospective study from a single center and therefore may be affected by referral bias and missing data. On the other hand, it was not possible to determine whether pre-existing AI was associated with AAV, we thus excluded these patients. Nevertheless, a recent study reported that the prognosis of patients with bronchiectasis prior to AAV diagnosis is poorer, which needs further investigation[33].

5. Conclusion

This study demonstrates that AI is also one of the common radiological manifestations of AAV, and AAV-related pulmonary involvement with diverse radiological features have different clinical characteristics and outcome. Specifically, the ILD group tends to have a poor long-term prognosis, the PG group is prone to relapse, and the AI group is apt to infection. The AH group has a high risk of both early infection and relapse, thus a poor short-term prognosis. Therefore, the intensity of immunosuppressive therapy must be carefully valued by considering the baseline CT findings among AAV patients with pulmonary involvement.

Abbreviations

AAV: antineutrophil cytoplasm antibodies-associated vasculitis
AI: airway involvement
ANCA: antineutrophil cytoplasm antibodies
BVAS: Birmingham Vasculitis Activity Scores
CNS: central nervous system
CT: computed tomography
DLCO: diffusing capacity for carbon monoxide
EGPA: eosinophilic granulomatosis with polyangiitis
ENT: ear-nose-throat
FEF\textsubscript{25-75} = forced expiratory flow at 25–75% of FVC
FEF\textsubscript{50} = forced expiratory flow at 50% of FVC
FEF\textsubscript{25} = forced expiratory flow at 50% of FVC
FEV1: forced expiratory volume
FFS: five factors score

FVC: forced vital capacity

GPA: granulomatosis with polyangiitis

MPA: microscopic polyangiitis; no: number

MPO: myeloperoxidase

NSIP: non-specific interstitial pneumonia

OP: organizing pneumonia

PR3: proteinase-3

TLC: total lung capacity

pred: predicted

UIP: usual interstitial pneumonia

**Declarations**

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Figures
AAV: ANCA-associated vasculitis; AH: alveolar hemorrhage; AI: airway involvement; CT: computed tomography; CTD: connective tissue disease; ILD: interstitial lung disease; PG: pulmonary granuloma.

Figure 1

Flow chart of the study population.
Figure 2

Kaplan-Meier survival curves for ANCA-associated vasculitis patients with pulmonary involvement.
Figure 3

Kaplan-Meier analysis comparing ANCA-associated vasculitis patients with different patterns of pulmonary involvements. (A) overall survival; (B) relapse-free survival; (C) infection-free survival.

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