





# Subsequent Antituberculous Treatment May Not Be Mandatory Among Surgically Resected Culture-Negative Pulmonary Granulomas: A Retrospective Nationwide Multicenter Cohort Study

Che-Liang Chung, Wei-Chang Huang, 3.4,5.6,7 Hung-Ling Huang, 3.9,10 Chun-Shih Chin, Meng-Hsuan Cheng, Meng-Rui Lee, 11,12 Sheng-Hao Lin, Ann-Yuan Wang, Ching-Hsiung Lin, 13,14,15 Inn-Wen Chong, 11,10-11,10 Shih, 12 and Chong-Jen Yu<sup>11,12</sup>

<sup>1</sup>Department of Internal Medicine, Yuanlin Christian Hospital, Changhua, Taiwan, <sup>2</sup>Division of Chest Medicine, Department of Internal Medicine, Taichung Veterans General Hospital, Taichung, Taiwan, <sup>3</sup>College of Medicine, National Chung Hsing University, Taichung, Taiwan, <sup>5</sup>School of Medicine, Chung Shan Medical University, Taichung, Taiwan, <sup>6</sup>Master Program for Health Administration, Department of Industrial Engineering and Enterprise Information, Tunghai University, Taichung, Taiwan, <sup>7</sup>Department of Medical Technology, Jen-Teh Junior College of Medicine, Nursing and Management, Miaoli, Taiwan, <sup>8</sup>Department of Internal Medicine, Kaohsiung, Municipal Ta-Tung Hospital, Kaohsiung, Taiwan, <sup>9</sup>Division of Pulmonary and Critical Care Medicine, Department of Internal Medicine, Kaohsiung Medical University Hospital, Kaohsiung, Taiwan, <sup>10</sup>Graduate Institute of Medicine, College of Medicine, National Taiwan University Hospital, Hsin-Chu, Taiwan, <sup>12</sup>Department of Internal Medicine, National Taiwan University Hospital, Hsin-Chu, Taiwan, <sup>13</sup>Division of Chest Medicine, Department of Internal Medicine, Changhua Christian Hospital, Changhua, Taiwan, <sup>14</sup>Institute of Genomics and Bioinformatics, National Chung Hsing University, Taichung, Taiwan, <sup>15</sup>Department of Recreation and Holistic Wellness, MingDao University, Changhua, Taiwan, <sup>16</sup>Department of Biological Science and Technology, National Yang Ming Chiao Tung University, Hsin-chu, Taiwan

*Background.* Histologic diagnosis of granuloma is often considered clinically equivalent to a definite diagnosis of pulmonary tuberculosis (TB) in endemic areas. Optimal management of surgically resected granulomatous inflammation in lung with negative mycobacterial culture results, however, remains unclear.

*Methods.* From 7 medical institutions in northern, middle, and southern Taiwan between January 2010 and December 2018, patients whose surgically resected pulmonary nodule(s) had histological features suggestive of TB but negative microbiological study results and who received no subsequent anti-TB treatment were identified retrospectively. All patients were followed up for 2 years until death or active TB disease was diagnosed.

**Results.** A total of 116 patients were enrolled during the study period. Among them, 61 patients (52.6%) were clinically asymptomatic, and 36 (31.0%) patients were immunocompromised. Solitary pulmonary nodule accounted for 44 (39.6%) of all cases. The lung nodules were removed by wedge resection in 95 (81.9%), lobectomy in 17 (14.7%), and segmentectomy in 4 (3.4%) patients. The most common histological feature was granulomatous inflammation (n = 116 [100%]), followed by caseous necrosis (n = 39 [33.6%]). During follow-up (218.4 patient-years), none of the patients developed active TB.

**Conclusions.** In patients with surgically resected culture-negative pulmonary granulomas, the incidence rate of subsequent active TB is low. Watchful monitoring along with regular clinical, radiological, and microbiological follow-up, instead of routine anti-TB treatment, may also be a reasonable option.

Keywords. acid-fast stain; caseous necrosis; granulomatous inflammation; pulmonary nodule; tuberculosis.

With the widespread use of chest computed tomography (CT), solitary pulmonary nodules (SPNs) and multiple pulmonary nodules are now frequently encountered in clinical setting. The underlying etiologies can be diverse and may vary with

Received 6 August 2021; editorial decision 1 November 2021; accepted 4 November 2021; published online 9 November 2021.

Correspondence: Meng-Rui Lee, MD, PhD, Department of Internal Medicine, National Taiwan University Hospital, #7 Chung-Shan South Road, Zhongzheng District, Taipei 10002, Taiwan (Ieemr@ntu.edu.tw).

### Onen Forum Infectious Diseases®2021

© The Author(s) 2021. Published by Oxford University Press on behalf of Infectious Diseases Society of America. This is an Open Access article distributed under the terms of the Creative Commons Attribution-NonCommercial-NoDerivs licence (https://creativecommons.org/licenses/by-nc-nd/4.0/), which permits non-commercial reproduction and distribution of the work, in any medium, provided the original work is not altered or transformed in any way, and that the work is properly cited. For commercial re-use, please contact journals.permissions@oup.com https://doi.org/10.1093/ofid/ofab565

the nodular sizes, pattern, and local epidemiology [1]. In early reports, malignancy accounted for 10%–70% of SPNs [2, 3]. Nonetheless, recent lung cancer screening studies of smokers suggested that most pulmonary nodules detected on CT were benign [4, 5]. In 1 recent study of 2 separate low-dose-CT screening cohorts, for instance, the proportion of benign diseases among patients with nodules was 94.5% and 96.3%, respectively [4]. Surgical resection, however, was still performed among many patients to obtain a definite diagnosis of lung nodules [1, 6]. In approximately 25% of the surgical procedures, the resected pulmonary nodules were benign in diagnosis [4]. Furthermore, about 80% of all benign nodules were infectious granulomas [1, 6].

Granulomatous inflammation, a unique type of chronic inflammation, results from tissue reaction following cell injury due to various conditions [7]. Histologic diagnosis of granuloma is often considered as equivalent to a definite diagnosis of tuberculosis (TB) in endemic areas even when there is lack of TB bacteriological evidence [8–10]. Nevertheless, granulomatous inflammation is not specific to active TB. Atypical mycobacteria, coccidioidomycosis, and histoplasmosis can also be the culprit pathogen [1], and granulomatous inflammation can also result from various conditions such as autoimmune diseases, toxins, drugs, and neoplasms [7, 11]. Although necrotizing granulomatous inflammation is more likely to contain organisms, the etiology still could not be identified in 25%–40% of resected necrotizing granulomas after clinical, serological, and microbiological surveillance [12–14]. On the other hand, nonnecrotizing granuloma also does not readily exclude infectious disease as etiology [7].

Currently, most patients with surgically resected solitary pulmonary granuloma would receive anti-TB treatment after operation [10]. Adverse effects, however, could develop in as much as 53%-61% of patients who received treatment [10, 15]. Furthermore, the TB reactivation rate was low among patients who did not receive anti-TB treatment [9, 10, 15-18], and risk of active TB was similar regardless of receiving anti-TB treatment or not [15]. Nevertheless, in the above studies, extensive surveillance for TB including respiratory specimens (sputum, bronchoalveolar lavage, and tissue) for acid-fast stain (AFS) and mycobacterial culture was not performed universally among included patients. This would lead to difficulty in interpretation of the findings and clinical implication [15, 17, 18]. With the increasing use of chest CT and identification of SPNs [19], reliable evidence regarding optimal management and the indication of anti-TB treatment after surgical resection is in great need.

To evaluate the outcome of untreated pulmonary nodules after surgical resection with histological evidence of TB infection and negative culture results, we conducted this nationwide multicenter retrospective cohort study to investigate the incidence for the development of active TB in this population.

# **METHODS**

#### **Study Population**

Between 1 January 2010 and 31 December 2018, patients with pulmonary nodule(s) who received surgical resection at 7 medical institutions with a total of >7000 beds in northern, middle, and southern Taiwan (National Taiwan University Hospital [NTUH], Taichung Veterans General Hospital [VGHTC], Kaohsiung Medical University Hospital [KMUH], Changhua Christian Hospital [CCH], and their 3 branch hospitals) and met the following criteria (1) caseous necrosis, or granulomatous inflammation in the histological samples; (2) not receiving anti-TB treatment during or within 3 weeks after the surgery were included. The institutional review boards (IRBs) of each

of the above institutions approved the study (NTUH IRB No. 202001021RINB; KMUH IRB-E(I)-20200117; CCH IRB No. 200126; VGHTC IRB No. CE20127A). IRBs waived the need for informed consent as data utilized in this retrospective study have been de-identified.

Patients were excluded if (1) *Mycobacterium tuberculosis* or nontuberculous mycobacteria (NTM) were isolated within 60 days before or after surgical resection; (2) tissue culture of the pulmonary nodule was not performed; or (3) there was histological or culture evidence (such as sputum or bronchoalveolar lavage) of fungal, viral, or parasitic infection.

Per the guidelines of the World Health Organization, the standard regimen used for treating new TB cases in Taiwan consists of isoniazid, rifampicin, pyrazinamide, and ethambutol for 2 months, followed by isoniazid and rifampicin plus ethambutol for 4 months if susceptibility testing is not available [20–23].

#### **Data Collection**

Information regarding patient age, sex, underlying diseases, past medical history of TB, surgical procedures (wedge resection, segmentectomy, or lobectomy), histology, laboratory test results at the beginning of tissue diagnosis, and other associated symptoms was retrieved retrospectively from medical records.

Microbiological data included AFS results from respiratory specimens and histology; mycobacterial culture results of sputum, bronchial washing, bronchoalveolar lavage, and tissue specimens were also retrieved.

Chest images (chest CT preferred) before surgical procedure were recorded, including presence of solitary or multiple pulmonary nodules, fibrocalcified lesions, bronchiectasis, cavitation, ground glass opacities, and maximum size of lesions.

## **Mycobacterial Culture Methods**

All respiratory specimens sent for mycobacterial culture were processed as previously described [24, 25]. In brief, the specimens for acid-fast bacilli smears were processed with auraminerhodamine fluorochrome and examined using standard procedures. Kinyoun stain method was used to confirm fluorochrome stain—positive smears. The standard *N*-acetyl-L-cysteine and sodium hydroxide method was used for the process of mycobacterial cultures. Cultures were performed by inoculating 0.5 mL of the processed specimens into Middlebrook 7H11 selective agar (Remel, Lexena, Kansas) and MGIT 960 culture tubes (BACTEC Mycobacteria Growth Indicator Tube 960 System, Becton-Dickinson Diagnostic Instrument Systems, Sparks, Maryland) [24, 25].

# Follow-up and Outcome

All patients were followed up for 2 years, or until death or a diagnosis of active TB. According to the policy and regulation of the National Tuberculosis Program of Taiwan, report of all culture-confirmed TB or clinically suspicious TB cases to the

Taiwan Centers for Disease Control (CDC) is mandatory [20]. TB culture results and follow-up outcomes were obtained from the database of the 7 institutions and confirmed by obtaining information on the TB registration databases of the Taiwan CDC.

#### Statistical Analysis

All statistical analyses were performed using IBM SPSS software (version 23.0; SPSS Inc, Chicago, Illinois).

#### **RESULTS**

## **Selection of Study Participants**

Figure 1 shows the process of patient identification and enrollment process. From January 2010 to December 2018, cases with histological evidence of caseous necrosis or granulomatous inflammation and negative tissue culture results who did not receive anti-TB therapy were retrieved for further analysis. All cases with positive tissue culture results for mycobacteria, fungi, or positive culture from other samples within 3 months were excluded.

#### **Clinical Characteristics of Included Patients**

In total, 116 patients were included for analysis (Figure 1). The median follow-up duration was 730 days (interquartile range: 0 day, quartile 1: 730 days, quartile 3: 730 days) since resection. Among them, 87.1% (n=101) of subjects were followed for 2 years.

Table 1 summarizes the clinical characteristics of the included 116 cases. The median age was 58 years, with a female:male ratio of 1:1. The most common comorbidity was malignancy (n = 29 [25.0%]), followed by diabetes mellitus (n = 27 [23.3%]). Twenty-one (18.1%) cases had a medical history of TB. Among patients with malignancy, 17 of 116 (14.7%) had lung cancer and 16 of 29 (55.1%) received systemic therapy after pulmonary nodule resection. There were neither human immunodeficiency virus–positive cases nor patients receiving biologic agent in our cohort.

Fifteen (12.9%) patients did not have complete clinical follow-up for 2 years in our hospitals and among them, no active TB development was found in the following 2 years based on national TB registry.

Sixty-one patients (52.6%) were clinically asymptomatic, and they received CT either due to lung cancer screening or incidental abnormal chest radiographic findings. In the other 55 symptomatic patients, 44 (80%) patients had cough symptoms and 15 (27.2%) had dyspnea. Because the symptoms were considered due to underlying comorbidities, the 55 cases did not receive anti-TB treatment. Additionally, none of our included patients received anti-TB treatment within 6 months prior to surgery or during the follow-up period.

### Surgical Procedures and Histological Characteristics of SPN

Of the 116 cases, 95 (81.9%) received wedge resection, 17 (14.7%) received lobectomy, and 4 (3.4%) received segmentectomy. Among patients with multiple pulmonary

```
1 Jan 2010-31 Dec 2018, histological evidence of TB (caseous necrosis or
     granulomatous inflammation) (n = 3792) in 5 medical centers in Taiwan
Histological evidence of pulmonary fungal infection (n = 65)
                    Lung tissue specimens available (n = 828)
          Tissue culture for mycobacteria or fungus performed (n = 560)
Positive tissue culture
 MTB (n = 113), MAC (n = 22), M. kansasii (n = 13), M. abscessus (n = 6),
 M. gordonae (n = 1), nonchromogens (n = 1), photochromogens (n = 1),
 NTM not otherwise specified (n = 34), Cryptococcus neoformans (n = 2)
                         Tissue culture negative (n = 367)
Positive culture from other samples within 3 months
 MTB (n = 24), MAC (n = 16), M. fortuitum (n = 9), M. abscessus (n = 5), M. kansasii (n = 3),
 M. gordonae (n = 2), M. chelonae (n = 1), nonchromogens (n = 1), M. lentiflavum (n = 1),
 M. parascrofulaceum (n = 2), NTM not otherwise specified (n = 41), fungus (n = 2)
       Culture negative for both lung tissue and other specimens (n = 260)
Receiving anti-TB treatment (n = 144)
                    Not receiving anti-TB treatment (n = 116)
```

Figure 1. Case selection process. Abbreviations: MAC, Mycobacterium avium-intracellulare complex; MTB, Mycobacterium tuberculosis; NTM, nontuberculous mycobacteria; TB, tuberculosis.

Table 1. Clinical Characteristics of Study Participants (N = 116)

Variable	No. (%)
Male sex	58 (50.0)
Age, y, median (min–max)	58 (22–82)
Follow-up duration, d, median (min–max)	730 (18–730
Surgical procedure	
Wedge	95 (81.9)
Segmentectomy	4 (3.4)
Lobectomy	17 (14.7)
Lymph node sampling	58 (50.0)
Histology	
Granulomatous inflammation	116 (100.0)
Caseous necrosis	39 (33.6)
Acid-fast staining	
Not performed	37 (31.9)
Negative	72 (62.1)
Positive	7 (6.0)
Lymph node involvement <sup>a</sup>	15 (12.9)
PAS or GMS stain	
Not performed	24 (20.7)
Negative	92 (79.3)
Concomitant malignancy	14 (12.1)
Underlying disease	
Malignancy	29 (25.0)
Lung cancer	17 (14.7)
Others	12 (10.3)
Status	
Remission	13 (11.2)
Under systemic treatment	16 (13.8)
Diabetes mellitus	27 (23.3)
History of tuberculosis	21 (18.1)
Chronic kidney disease stage ≥3	14 (12.1)
Cirrhosis of liver	2 (1.7)
Transplant recipients	2 (1.7)
Alcoholism	3 (2.6)
Autoimmune disease	11 (9.5)
Symptoms	
Cough	44 (37.9)
Sputum	24 (20.7)
Dyspnea	15 (12.9)
Hemoptysis	7 (6.0)
Fever	6 (5.2)
Weight loss	16 (13.8)
Clinically asymptomatic	61 (52.6)
Development of active TB within 2 y	0

Data are expressed as No. (%) unless otherwise indicated.

Abbreviations: GMS, Gomori methenamine silver; PAS, periodic acid-Schiff; TB, tuberculosis.

nodules, the granulomatous nodules in the same lung segment were removed at the same time during surgery. Otherwise, whether all lung nodules were removed during surgery depended on lung function reserve and complexity of surgical procedure. Fourteen cases (12.1%) had concomitant malignancy and granulomatous inflammation in histological specimen. The most common histological feature was granulomatous

Table 2. Radiographic Pattern and Laboratory Data of All Participants (N = 116)

Variable	No. (%)
Findings on chest CT	
Multiple nodules	64 (57.7)
Solitary nodule	44 (39.6)
Lesion size >3 cm	10 (9.0)
Associated findings	
Ground glass opacity	36 (32.4)
Fibrosis	24 (21.6)
Calcification	17 (15.3)
Bronchiectasis	13 (11.7)
Cavitation	7 (6.3)
Laboratory data, mean ± SD	
Albumin, g/dL	4.1 ± 0.6
Hemoglobin, mg/dL	13.1 ± 1.9
Leukocyte count, K/µL	$6.8 \pm 2.2$
Segment, %	63.3 ± 11.6
Lymphocytes, %	24.7 ± 10.8
C-reactive protein, mg/dL	3.2 ± 7.8

Data are expressed as No. (%) unless otherwise indicated. Abbreviations: CT, computed tomography; SD, standard deviation.

inflammation (n = 116 [100%]), followed by caseous necrosis (n = 39 [33.6%]). In 21 cases with past history of TB, 9 had histological finding of caseous necrosis. Fifteen cases (12.9%) had concomitant histological evidence of caseous necrosis or granulomatous inflammation of lymph nodes in addition to pulmonary nodule(s). Tissue AFS showed positive results in 7 (6%) patients. Of them, 5 had past medical history of TB. Fungal stain (periodic acid-Schiff or GMS stain) was performed in 92 (79.3%) of all cases, all of which yielded negative results.

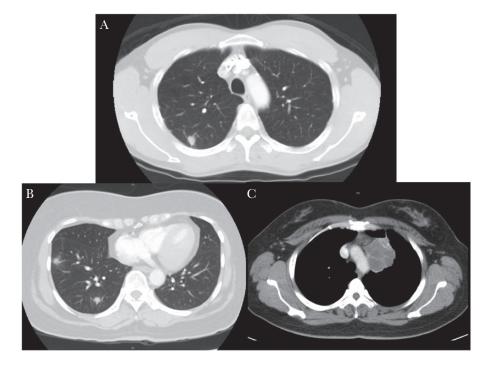
# **Radiographic Features of SPNs**

Table 2 shows laboratory data and radiographic features of the cohort. In summary, most patients had normal hemogram and albumin values. SPNs (Figure 2A) accounted for 44 (39.6%) of all cases, whereas multiple pulmonary nodules (Figure 2B) accounted for 64 (57.7%) cases. Ten (22.7%) cases had at least 1 lesion size >3 cm in diameter (Figure 2C). All 7 (6.3%) cases with cavitary lesions had multiple pulmonary nodules.

### **Outcome After Observation Without Anti-TB Treatment**

During a total of 218.4 patient-years of follow-up, none of the patients developed active TB according to the medical records and the data in the TB registration databases of Taiwan CDC. One 67-year-old male patient died at day 18 after surgery due to septic shock. The pathological report showed negative results of acid-fast staining and fungal staining. Cultures from sputum, bronchoalveolar lavage, and surgical specimens all showed no evidence of mycobacterial infection. In 36 cases with immunocompromised status (cirrhosis, diabetes mellitus, transplant recipients, autoimmune diseases), none developed active TB within a total follow-up of 66.6 patient-years.

<sup>&</sup>lt;sup>a</sup>Lymph node involvement refers to concomitant histological evidence of caseous necrosis or granulomatous inflammation of lymph nodes in addition to pulmonary nodule(s).



**Figure 2.** Chest computed tomography: a 35-year-old man with an ovoid 1.7-cm nodule with pleural tagging at the posterior aspect of the right upper lobe (*A*); a 52-year-old woman with 2 round nodules in the right lower lobe (*B*); and a 36-year-old woman with one 6.2-cm lobulated mass-like lesion in the left upper lobe (*C*).

#### **DISCUSSION**

This study investigated the incidence of active TB within 2 years in patients with culture-negative surgically resected pulmonary granulomas after thorough clinical, radiological, and microbiological workup. Mycobacterial culture was performed in all resected pulmonary granulomas with negative findings. While patients in this study may be considered to be not at high risk for TB by their physicians, the fact that no active TB developed in this cohort during a total of 218.4 patient-years follow-up implies that the risk of TB in this special group is likely to be lower than TB contacts who are test-positive by IGRA (5% in the first 2 years) [26]. Routine treatment with anti-TB drugs immediately for cases with histological evidence suggestive of TB but negative culture results should not be considered the only and standard option.

A literature review of patients with resected lung nodules having histological findings suggestive of TB yet subsequently not receiving anti-TB treatment was summarized in Table 3 [9, 10, 12, 15–18, 27]. Four hundred twenty-one cases with "tuberculoma," "granuloma," or "granulomatous inflammation" were identified in 7 studies. Of them, 212 (50.4%) cases received no anti-TB therapy. The duration of follow-up in these studies varied from 0.1 to 16.7 years. Although some data were unavailable in these studies, only 1 cancer patient who underwent regular chemotherapy developed TB reactivation [15]. In the other report, 2 cases with resected pulmonary necrotizing granulomas developed new lung nodules but did not receive additional treatment or develop new symptomatic disease [12].

Overall, the estimated incidence of active TB among patients who received no anti-TB treatment in the 7 studies was <1 per 647.2 patient-years (Table 3).

Benefit of empiric anti-TB treatment should be weighed with risk of treatment-related drug toxicity. In 1 retrospective study from Taiwan, 53% of patients with culture-negative granulomas developed adverse events after empiric anti-TB treatment and 18% of them had drug-induced hepatotoxicity [15]. In our cohort, 49.1% of enrolled cases were >60 years old and concern exists regarding the impact of age and comorbidity on management and follow-up of pulmonary nodules [28]. Considering higher risk of serious adverse effects among elderly patients [29] and uncertainty of TB diagnosis, the necessity of empiric anti-TB treatment should be carefully evaluated in culture-negative granuloma patients with low possibility of infectiousness.

In 1 large retrospective study, despite increasing use of chest CT and identifying of incidental SPNs, the incidence of lung cancer diagnosis within 2 years of SPN detection was not increased [19]. Therefore, more frequent SPN identification was presumed to be secondary to indolent infections, granulomas, and scar formation [19]. With the increasing encountering of pulmonary granulomas in clinical practice, skepticism regarding the need for treatment is warranted.

According to previous reports, infectious granuloma, such as actinomycetes, mycobacteria, fungi, and helminths infection, accounted for about 80% of benign nodules [1, 6, 13, 30]. Nevertheless, no micro-organisms can be identified after

Table 3. Literature Review of Clinical Characteristics and Outcomes of Patients With Resected Lung Nodules Having Histological Findings Suggestive of Tuberculosis Yet Subsequently Not Receiving Antituberculosis

Study, First Author, Year [Ref.]	No. of Patients, Total/No ATT	Age, y <sup>a</sup>	Median FU Year (Range)	No. of Patients With AFS, Positive/Performed	Coexistence of Cancer, No. (%)	No. of Patients Developing Active TB (FU Duration <sup>b</sup> )
Ishida, 1992 [18]	36/8	53.5 (23–76)	(1–16)	16/NA	NA	0 (≥8 PY)
Mukhopadhyay, 2013 [12]	52/36	55 ± 14.9	7 (0.1–16.7)	0/52	NA	0 (217 PY) <sup>c</sup>
Yakar, 2016 [17]	48/37	63 (40–76)	≥2	0/25	48 (100%)	0 (≥74 PY)
Dagaonkar, 2017 [16]	19/18	63 (40-84)	1.3 (0.1-4.3)	0/19	19 (100%)	0 (23.4 PY)
Watanabe, 2017 [10]	8/3	59 (32-74)	NA	NA	1	0 (NA)
Chung, 2018 [15]	107/67	57 (21–91)	4 (0.5–4)	0/107	18 (17%)	1 (251.2 PY)
Wang, 2020 [9]	98/32	50.0 ± 13.2	2.3 (0.8-5.8)	59/98	NA	0 (73.6 PY)
Overall	421/212	NA	NA	59/301	NA	1 (≥647.2 PY)

Abbreviations: AFS, acid-fast staining; ATT, antituberculosis treatment; FU, follow-up; NA, not available; PY, person-years; TB, tuberculosis

Ziehl-Neelsen and GMS stains in more than one-fourth of patients who underwent resection of radiographically solitary pulmonary granuloma [14]. In a multinational study investigating 500 histological specimens containing pulmonary granulomas [31], 42% had no identifiable etiology. Of the 58% cases with identified etiology, sarcoidosis was the most common (27%), followed by mycobacterial or fungal infections (25%). Interestingly, mycobacteria were more commonly identified outside the United States, whereas fungi were more commonly in the United States [31]. Although incidence of tuberculomas and NTM pulmonary nodules had not been established, tuberculoma is still first considered for culture-negative granuloma in TB-endemic areas. In 2019, 8732 new TB cases were reported in in Taiwan and the overall incidence of TB was 37 cases per 100 000 population [32]. Of them, 20% were sputum smear-negative and culture-negative TB [33]. Tissue biopsy or resection may, therefore, be necessary for establishing diagnosis when noninvasive testing could not confirm the diagnosis.

Histological studies and microbiological cultures are complementary, although many cases can be tested positive for both modalities. In 1 report, cultures yielded more positive results in mycobacterial infections, yet fungi were identified in most histological specimens of fungal infections [31]. These results highlight the necessity to submit biopsied tissue for cultures as well as cultures of respiratory specimens whenever feasible. In our cohort, 191 cases with pulmonary granulomas were culture-positive for mycobacteria, while 65 cases with pulmonary granulomas were diagnosed as fungal infection histologically (Figure 1). This finding again emphasizes the importance of concomitant culture and pathology surveillance to increase diagnostic yield [31]. In a study of pulmonary necrotizing granulomas of unknown cause, a careful review of clinical features, radiographic studies, cultures, fungal serologies, and special stains could lead to identification of definite etiology in 60% of all histologically unexplained cases [12].

Our study still has limitations. First, since our study design was retrospective, whether anti-TB treatment could further lower the risk of TB reactivation in culture-negative granulomas remained unknown. The effect of anti-TB treatment on TB reactivation in this special clinical entity, however, can be trivial, as the results of our previous study suggested [15]. In addition, our untreated patients might not be considered high risk to have TB by their medical doctors, which might lead to selection bias. Second, this study did not have positron emission tomography (PET) scan results. Although PET scan is promising in helping differential diagnosis of lung nodules [34], it is still unable to reliably differentiate between active TB, inactive granuloma, and malignancy [10, 35]. Besides, workup for sarcoidosis was not universal and mandatory for all patients. None of our patients, however, were diagnosed with sarcoidosis during follow-up. Finally, nucleic acid amplification testing of histopathology was not routinely used in current study. Furthermore, neither the results of tuberculin skin testing nor interferon-y release assay (IGRA) were available since these tests are not recommended as diagnostic in this setting due to the high background rate of bacillus Calmette-Guérin (BCG, an attenuated Mycobacterium bovis) vaccine and the high cost of IGRAs in Taiwan [20, 36, 37].

In conclusion, after obtaining negative results through detailed clinical, radiological, microbiological, and histopathologic review, watchful monitoring instead of immediate anti-TB treatment may be a reasonable option. Regular radiographic examinations and microbiological study, however, are still warranted.

#### Notes

Author contributions. M.-R. L. is the guarantor of the paper and takes responsibility for the content of the manuscript, including the data and analysis. C.-L. C., with M.-R. L. and J.-Y. W., drafted the manuscript and designed the study. C.-L. C., M.-R. L., W.-C. H., H.-L. H., and J.-Y. W. were involved in data processing. C.-L. C., C.-S. C., M.-H. C., S.-H. L., and J.-Y. W. performed statistical analysis. J.-Y. W., C.-H. L., I.-W. C., J.-Y. S., and

<sup>&</sup>lt;sup>a</sup>Data are expressed as median (min-max) or mean ± standard deviation.

bMedian number of follow-up years was assumed as mean number for calculation of incidence rate of active TB in References 9, 12, and 16. Minimum follow-up year was used for calculation in References 17 and 18. The exact follow-up period was available in the text in reference 15. Reference 10 was not included for calculation due to insufficient information.

<sup>&</sup>lt;sup>c</sup>Follow-up period was available in 31 of 36 no ATT patients.

C.-J. Y. supervised the research and provided critical revision of the article. All authors reviewed, provided input, and approved the final manuscript.

Acknowledgments. The authors would like to thank Dr Gwan-Han Shen, who supervised Laboratory No. 114 in Taichung Veterans General Hospital and passed away in 2014. We hold you dear in our memories. The authors also would like to thank members of Yuanlin Christian Hospital Center for Infection Control, Hui-Chen Yu and Chung-Ya Huang, for data collection.

*Financial support.* This research was supported by Taiwan Ministry of Science and Technology [109-2314-B-002-185-MY3]. The funder had no role in the study design, data analysis or manuscript writing.

Potential conflicts of interest. All authors: No reported conflicts of interest.

All authors have submitted the ICMJE Form for Disclosure of Potential Conflicts of Interest. Conflicts that the editors consider relevant to the content of the manuscript have been disclosed.

#### References

- Kikano GE, Fabien A, Schilz R. Evaluation of the solitary pulmonary nodule. Am Fam Physician 2015; 92:1084–91.
- Khouri NF, Meziane MA, Zerhouni EA, et al. The solitary pulmonary nodule. Assessment, diagnosis, and management. Chest 1987; 91:128–33.
- Siegelman SS, Khouri NF, Leo FP, et al. Solitary pulmonary nodules: CT assessment. Radiology 1986; 160:307–12.
- McWilliams A, Tammemagi MC, Mayo JR, et al. Probability of cancer in pulmonary nodules detected on first screening CT. N Engl J Med 2013; 369:910–9.
- Aberle DR, Adams AM, Berg CD, et al. Reduced lung-cancer mortality with lowdose computed tomographic screening. N Engl J Med 2011; 365:395–409.
- Ost D, Fein AM, Feinsilver SH. Clinical practice. The solitary pulmonary nodule. N Engl J Med 2003; 348:2535–42.
- 7. Shah KK, Pritt BS, Alexander MP. Histopathologic review of granulomatous inflammation. J Clin Tuberc Other Mycobact Dis 2017; 7:1–12.
- 8. Subotic D, Yablonskiy P, Sulis G, et al. Surgery and pleuro-pulmonary tuberculosis: a scientific literature review. J Thorac Dis 2016; 8:E474–85.
- Wang C, Liu Y, Lin H, et al. The necessity of anti-tuberculosis therapy after resection of pulmonary tuberculous nodules: a single center retrospective study. Ann Thorac Cardiovasc Surg 2020; 26:190–5.
- Watanabe H, Uruma T, Seita I, et al. Solitary pulmonary caseating granulomas: a 5-year retrospective single-center analysis. Mol Clin Oncol 2017; 6:839–45.
- Williams GT, Williams WJ. Granulomatous inflammation—a review. J Clin Pathol 1983; 36:723–33.
- Mukhopadhyay S, Wilcox BE, Myers JL, et al. Pulmonary necrotizing granulomas of unknown cause: clinical and pathologic analysis of 131 patients with completely resected nodules. Chest 2013; 144:813–24.
- Aubry MC. Necrotizing granulomatous inflammation: what does it mean if your special stains are negative? Mod Pathol 2012; 25(Suppl 1):S31-8.
- Ulbright TM, Katzenstein AL. Solitary necrotizing granulomas of the lung: differentiating features and etiology. Am J Surg Pathol 1980; 4:13–28.
- Chung CL, Chen YF, Lin YT, et al. Outcome of untreated lung nodules with histological but no microbiological evidence of tuberculosis. BMC Infect Dis 2018; 18:530.
- Dagaonkar RS, Choong CV, Asmat AB, et al. Significance of coexistent granulomatous inflammation and lung cancer. J Clin Pathol 2017; 70:337–41.
- Yakar F, Yakar A, Büyükpınarbaşılı N, Erelel M. Does every necrotizing granulomatous inflammation identified by NSCLC resection material require treatment? Med Sci Monit 2016; 22:1218–22.

- Ishida T, Yokoyama H, Kaneko S, et al. Pulmonary tuberculoma and indications for surgery: radiographic and clinicopathological analysis. Respir Med 1992; 86:431-6.
- Gould MK, Tang T, Liu IL, et al. Recent trends in the identification of incidental pulmonary nodules. Am J Respir Crit Care Med 2015; 192:1208–14.
- Centers for Disease Control, Ministry of Health and Welfare, Taiwan. Taiwan Guidelines for TB Diagnosis and Treatment [in Chinese]. 6th ed. Taipei: Taiwan Centers for Disease Control; 2017. https://www.cdc.gov.tw/En/File/Get/Ptat7PMqyrzuqYbiPlZ4VQ. Accessed 21 October 2021.
- World Health Organization. Guidelines for Treatment of Drug-Susceptible Tuberculosis and Patient Care, 2017 Update. 2017. https://apps.who.int/iris/bitstream/handle/10665/255052/9789241550000-eng.pdf. Accessed 21 October 2021.
- World Health Organization. Guidelines for Treatment of Tuberculosis. 4th ed. 2010. https://apps.who.int/iris/bitstream/handle/10665/44165/9789241547833\_eng.pdf. Accessed 21 October 2021.
- Nahid P, Dorman SE, Alipanah N, et al. Official American Thoracic Society/ Centers for Disease Control and Prevention/Infectious Diseases Society of America clinical practice guidelines: treatment of drug-susceptible tuberculosis. Clin Infect Dis 2016; 63:e147–95.
- Lee MR, Yang CY, Shu CC, et al. Factors associated with subsequent nontuberculous mycobacterial lung disease in patients with a single sputum isolate on initial examination. Clin Microbiol Infect 2015; 21:250.e1-7.
- Ruan SY, Chuang YC, Wang JY, et al. Revisiting tuberculous pleurisy: pleural fluid characteristics and diagnostic yield of mycobacterial culture in an endemic area. Thorax 2012; 67:822-7.
- Mack U, Migliori GB, Sester M, et al; TBNET. LTBI: latent tuberculosis infection or lasting immune responses to *M. tuberculosis*? A TBNET consensus statement. Eur Respir J 2009; 33:956–73.
- Hsu KY, Lee HC, Ou CC, Luh SP. Value of video-assisted thoracoscopic surgery in the diagnosis and treatment of pulmonary tuberculoma: 53 cases analysis and review of literature. J Zhejiang Univ Sci B 2009; 10:375–9.
- Wong ML, Shi Y, Fung KZ, et al. Age, comorbidity, life expectancy, and pulmonary nodule follow-up in older veterans. PLoS One 2018; 13:e0200496.
- Yee D, Valiquette C, Pelletier M, et al. Incidence of serious side effects from firstline antituberculosis drugs among patients treated for active tuberculosis. Am J Respir Crit Care Med 2003; 167:1472–7.
- Kradin RL, Mark EJ. Pulmonary infections. In: Kradin R, ed. Diagnostic Pathology of Infectious Disease. 1st ed. Philadelphia: Saunders Elsevier: 2010:125–88.
- Mukhopadhyay S, Farver CF, Vaszar LT, et al. Causes of pulmonary granulomas: a retrospective study of 500 cases from seven countries. J Clin Pathol 2012; 65:51.7
- Centers for Disease Control, Ministry of Health and Welfare, Taiwan. 2020 CDC Annual Report. Taipei: Taiwan Centers for Disease Control; 2020. https://www.cdc.gov.tw/File/Get/VzxSlitRN9UWzV7N9z6bNA. Accessed 21 October 2021.
- Centers for Disease Control, Ministry of Health and Welfare, Taiwan. Taiwan tuberculosis control report 2019 [in Chinese]. 2020. https://www.cdc.gov.tw/File/ Get/eohpjs5F-9obJG4sMlmHBw. Accessed 21 October 2021.
- Kim IJ, Lee JS, Kim SJ, et al. Double-phase 18F-FDG PET-CT for determination of pulmonary tuberculoma activity. Eur J Nucl Med Mol Imaging 2008; 35:808–14.
- Priftakis D, Riaz S, Zumla A, Bomanji J. Towards more accurate 18F-fluorodeoxyglucose positron emission tomography (18F-FDG PET) imaging in active and latent tuberculosis. Int J Infect Dis 2020; 92S:85–90.
- Jou R, Huang WL, Su WJ. Tokyo-172 BCG vaccination complications, Taiwan. Emerg Infect Dis 2009; 15:1525-6.
- Chiu WT, Scholl J, Li YJ, Wu J. So few COVID-19 cases in Taiwan: has population immune health played a role? Front Public Health 2021; 9:676750.