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Interstitial lung disease following COVID-19 vaccination: a disproportionality analysis using the Global Scale Pharmacovigilance Database (VigiBase)

Database (VigiBase) Min-Taek Lee,^{1,2} Ju Won Lee,^{1,2} Hyeon Ji Lee,^{1,2} Jong-Min Lee,^{1,2} Jae Chol Choi,^{3,4} Kang-Mo Gu,^{4,5} Sun-Young Jung ¹

ABSTRACT Background and objective Despite several case reports,

analvsis.

vaccines.

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Dr Sun-Young Jung; jsyoung@cau.ac.kr reporting of ILD was observed for COVID-19 vaccines compared with all other vaccines. Moreover, when compared with the influenza vaccines that are known

to cause ILD, no signal was observed. This study results might help decision-making on the subsequent COVID-19 vaccination strategy of ILD. Further large and prospective studies are required for more conclusive evidence.

population-based studies on interstitial lung disease (ILD)

following COVID-19 vaccination are lacking. Given the

unprecedented safety issue of COVID-19 vaccination,

it is important to assess the worldwide patterns of ILD

investigate the signals of COVID-19 vaccine-associated ILD

compared with other vaccinations using disproportionality

Methods We analysed the VigiBase database during the

We adopted the case/non-case approach to assess the

disproportionality signal of ILD for COVID-19 vaccines via

1:10 matching by age and sex. We compared COVID-19

vaccines with all other vaccines as the reference group.

Results Among 1 233 969 vaccine-related reports, 679

were reported for ILD. The majority of ILD cases were

related to tozinameran (376 reports, 55.4%), Vaxzevria

The reporting OR of ILD following COVID-19 vaccination

was 0.86 (95% CI 0.64 to 1.15) compared with all other

Conclusion No significant signal of disproportionate

(129 reports, 19.0%) and elasomeran (78 reports, 11.5%).

period between 13 December 2020 and 26 January 2023.

following COVID-19 vaccination. This study aimed to

INTRODUCTION

As of September 2023, a total of 13.5 billion doses of COVID-19 vaccines have been administered, which averted millions of death worldwide.¹² The World Health Organization (WHO) has recently declared the expiration of COVID-19 public health emergency and revise the long-term COVID-19 disease management strategies.³ It is expected that the COVID-19 vaccines will be included in

WHAT IS ALREADY KNOWN ON THIS TOPIC

⇒ Since the first case report of interstitial lung disease (ILD) following COVID-19 vaccination was published on 9 June 2021, several ILD cases have been reported. However, population-based studies on ILD following COVID-19 vaccination are lacking.

WHAT THIS STUDY ADDS

⇒ No significant signal of disproportionate reporting of ILD was observed for COVID-19 vaccines compared with other vaccines (reporting OR 0.86, 95% CI 0.64 to 1.15). These findings were consistent across several analyses conducted after considering potential biases.

HOW THIS STUDY MIGHT AFFECT RESEARCH, PRACTICE OR POLICY

⇒ This study may provide information that can be useful for making decisions on subsequence COVID-19 vaccine strategies. Moreover, further studies using patient-level information such as disease history and diagnostic test results are required for more conclusive evidence.

the regular immunisation schedule similar to other seasonal influenza vaccines.^{4 5} To ensure a successful vaccination programme, safety information is important, especially for concerns that are not fully addressed. WHO encourages countries to perform research on vaccines with respect to unknown critical information.³

Since August 2021, cases of interstitial lung disease (ILD) following COVID-19 vaccination have been reported, although the underlying aetiology remained poorly understood. ILD is a heterogeneous group of diseases characterised by progressive inflammation and injury to the interstitium and alveoli.^{6 7} The incidence of ILD varies according to age, sex, region and race and the prevalence is



approximately 6.3–76.0 cases per 100 000 people.⁸⁹ The causes of ILD are not clearly known, but several potential risk factors have been suggested, including systemic autoimmune disease and drug exposure.⁷ The incidence and prevalence of drug-induced ILD are not well known; however, approximately 2.5%-5.0% of all prevalent ILD cases are estimated to be drug induced.^{10 11} Amiodarone and methotrexate are known to cause drug-induced ILD, and the use of these medications has been reported in over 10% of cases with mortality.¹⁰ According to previous reports, vaccination, especially for influenza, is likely to cause ILD.^{12–14} Conversely, there have been some case reports suggesting an association between COVID-19 vaccination and the development and progression of ILD.¹⁵⁻²² It remains unknown whether COVID-19 vaccination-associated ILD has distinct characteristics compared with the disease induced by other vaccines, such as the influenza vaccine.

Spontaneous reports are a useful source to assess signals of rare but serious adverse events (AEs), including COVID-19 vaccine-induced ILD. Studies suggest a temporary increase in reporting rate after product approval and safety alerts due to safety concerns. Therefore, it is necessary to verify ILD cases as the COVID-19 vaccination rate increases. Given that the unprecedented safety issues of COVID-19 vaccines have been raised, it is important to study the identifying characteristics of ILD following COVID-19 vaccination and investigate factors affecting the risk of ILD or reporting rate. Therefore, this study aimed to assess the disproportionality of reporting of ILD associated with COVID-19 vaccines using the case/ non-case approach by analysing the WHO global pharmacovigilance database.

METHODS

Data source

We used VigiBase, the largest global pharmacovigilance database with over 30 million reports of suspected AEs of medicines since 1968.²³ It was developed and maintained by WHO-Uppsala Monitoring Centre (UMC). The WHO-UMC receives individual case safety reports (ICSRs) from over 150 countries participating in the WHO programme for international drug monitoring.²³ VigiBase is composed of several medical and drug classification elements, such as the medical dictionary for regulatory activities (MedDRA) and WHODrug. The AEs analysed in our study were investigated using MedDRA version 26.0 (released March 2023) with preferred terms (PTs) and lowest level terms (LLTs), and drugs were coded using WHODrug Global B3/C3-format 1 March 2023.

Variables

We extracted the ICSRs with vaccines as suspected drugs between 13 December 2020 and 26 January 2023.²⁴ ILD was defined using MedDRA-standardised MedDRA queries (SMQ) (SMQ code=20000042) narrow terms to provide a clear definition and to account for specificity (cases highly likely to be of interest). There were 79 PTs and 132 LLTs in the ILD defined by MedDRA SMQ (online supplemental table 1). The COVID-19 vaccines tozinameran, elasomeran, Vaxzevria, Ad26. COV2.S, Gam-COVID-Vac, NVX-CoV2373 and GBP510 were included in our study and were defined using drug record numbers in WHODrug (online supplemental table 2) and the Anatomical Therapeutic Chemical (ATC) classification (J07BN). The other vaccines were classified according to ATC classification (J07; vaccines). We included physicians, pharmacists and other health professionals as notifier types, and excluded reports with missing values, including age and sex. The dates on which the reports were entered into the VigiBase were arranged by quarters. The geographical regions were divided into five groups: Africa, the Americas, South-East Asia, Europe, Eastern Mediterranean and Western Pacific. Using ICSRs, we calculated the time-to-onset, which is the time interval between vaccine administration and initiation of the event. In VigiBase, ICSRs contain multiple AEs with several different time-to-onset. Therefore, we selected the vaccine-AE pairs with the most information, including dechallenge action/outcome and rechallenge action/outcome as representatives. Moreover, if vaccine-AE pairs have an equal number of outcome information, we chose the ICSR with the shortest time-to-onset. Furthermore, we regarded time-to-onset as an outlier by individual pairs (coded as missing values) if it was outside the study period.

Statistical analysis

We performed a descriptive analysis of ILD cases and non-cases. Continuous variables, including time-to-onset, were presented as the mean±SD and were compared using the Student's t-test. The categorical variables, including reported quarter, age groups, sex, type of report, type of COVID-19 vaccines, seriousness, region and type of notifier, were reported as numbers (percentage) and compared using the χ^2 test and Fisher's exact test.

The association between COVID-19 vaccines and ILD was evaluated using case/non-case analysis.²⁵ The case/non-case analysis is a disproportionality approach performed in the pharmacovigilance databases developed during the early 1980s.²⁶ Briefly, it is similar to casecontrol analysis but uses non-case instead of control. In the spontaneous AE report database, the ICSRs indicate reports of exposure to the drug of interest at least once and any AE experienced any AE at least once.²⁵ In our study, cases were defined as ICSRs of ILD while the remaining ICSRs were considered non-cases. The primary analysis compared the COVID-19 vaccines with all other vaccines. We compared ILD cases and non-cases by 1:10 matching according to age and sex as matching variables. The logistic regression model was used for calculating reporting ORs (RORs) and 95% CI.²⁷ We determined the detection of a signal according to the three criteria: the

ROR is greater than 1, the lower bound 95% CI is greater than 1 and the number of cases is greater than $3.^{25}$

We performed subgroup analyses using stratification by age groups, sex and region. We determined factors reported in previous case reports that may affect the occurrence of ILD. Individuals were categorised into two groups based on (1) age (<65 and \geq 65 years), (2) sex (male and female) and (3) region (Western Pacific region and the other regions).

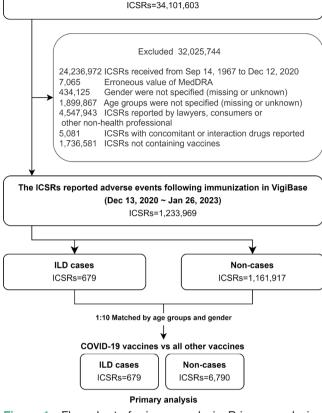
Moreover, sensitivity analyses were performed to identify diverse AE definitions and the extent of contribution of potential biases as follows. First, we designed sensitivity analysis 1 to define ICSRs with vaccines as suspected, concomitant and interaction drugs, as opposed to the primary analysis performed with suspected drugs. Second, we defined ILD using both MedDRA SMQ narrow and broad terms (broad search), thereby including all possible cases. Third, we excluded ICSRs that included drugs known to cause ILD, such as those used to treat cancer, rheumatic diseases, infection and cardiac diseases (online supplemental table 2) as these can influence the likelihood of detecting a signal between COVID-19 vaccines (drug competition bias).²⁸ Fourth, since events known as scientific and specific medical concerns about COVID-19 vaccines could affect other signal events (competition bias), we excluded reports containing 14 AEs of special interest of COVID-19 vaccine, including myocarditis, pericarditis and thrombosis, as suggested by Brighton collaboration.²⁹ Fifth, ICSRs reported as serious AEs were restricted in sensitivity analysis 5. The Weber effect could arise due to the market authorisation of new COVID-19 vaccines. Sixth, we included ICSRs with a reporting date before 9 August 2021, which may have influenced reporting in sensitivity analysis 6 (notoriety bias).¹⁵ Finally, in sensitivity analysis 7, we compared the COVID-19 vaccines with the influenza vaccines (ATC: J07BB, influenza vaccines) as a positive control. This choice was based on previous reports¹²⁻¹⁴ indicating that influenza vaccines have been known to cause ILD. Additionally, there is a report suggesting that the mechanism of ILD following COVID-19 vaccination may be similar to the mechanism of ILD following influenza vaccination.¹³ We have organised the overall analysis strategies in online supplemental table 3.

Patient and public involvement

As this is a secondary database study, the database is anonymised and served without identifiers of the study participants. The patients were not involved in the design, conduct or dissemination of this study.

RESULTS

A total of 1 233 969 reports with AEs following COVID-19 vaccination were identified from 12 December 2020 to 26 January 2023. After 1:10 matching by age group and sex, 7469 reports were determined to be ILD cases (679 ICSRs) and non-cases (6790 ICSRs) (figure 1). The



All ICSRs in VigiBase (Sep 14, 1967 ~ Jan 26, 2023)

Figure 1 Flow chart of primary analysis. Primary analysis compared COVID-19 vaccines with all other vaccines. ICSR, individual case safety report; ILD, interstitial lung disease; MedDRA, Medical Dictionary for Regulatory Activities.

characteristics of ILD cases/non-cases of the primary analysis, including reported quarter, age groups, sex, type of report, type of vacines, seriousness, region and timeto-onset, are shown in table 1. This study analysed six COVID-19 vaccines, including tozinameran, elasomeran, Vaxzevria, Ad26.COV2.S., Gam-COVID-Vac and NVX-CoV2373. GBP510 was not included in our study. Most of the reports were received in the third quarter of 2021 (104 ICSRs, 17.1%) with ILD cases following COVID-19 vaccination being first reported (table 1). A significant proportion of ILD cases received tozinameran (376 ICSRs, 55.4%) and were Europeans (577 reports, 85.0%). Serious AEs including death, life-threatening conditions and hospitalisation/prolonged hospitalisation were more likely in ILD cases (625 ICSRs, 92.1%) compared with non-cases (2343 ICSRs, 34.5%). ILD cases had the highest proportion of reports received by physicians (527 ICSR, 77.6%), followed by other health professionals (102 ICSRs, 105.8%) and pharmacists (50 ICSRs, 7.4%). The median time-to-onset was 7 (IQR 1-36) days for ILD cases and 1 (IQR 0-15) day for non-cases (p=0.0077).

The number of monthly ILD cases following COVID-19 vaccination is shown in figure 2. Most of the cases of ILD following COVID-19 vaccination (40 cases) were

 Table 1
 Characteristics of interstitial lung disease (ILD) and non-ILD cases from VigiBase database: primary analysis

 (compared COVID-19 vaccines with all other vaccines)

	ILD cases (N=679)	Non-cases (N=6790)	
	N (%)	N (%)	P value
Reported quarter (Q)			0.036
2020.4Q (13 December 2020–31 December 2020)	0 (0.0)	19 (0.3)	
2021.1Q (1 January 2021–31 March 2021)	49 (7.2)	703 (10.4)	
2021.2Q (1 April 2021–30 June 2021)	104 (15.3)	1193 (17.6)	
2021.3Q (1 July 2021–30 September 2021)	116 (17.1)	1066 (15.7)	
2021.4Q (1 October 2021–31 December 2021)	104 (15.3)	1002 (14.8)	
2022.1Q (1 January 2022–31 March 2022	102 (15.0)	883 (13.0)	
2022.2Q (1 April 2022–30 June 2022)	80 (11.8)	820 (12.1)	
2022.3Q (1 July 2022–30 September 2022)	44 (6.5)	463 (6.8)	
2022.4Q (1 October 2022–31 December 2022)	66 (9.7)	560 (8.3)	
2023.1Q (1 January 2023–26 January 2023)	14 (2.1)	81 (1.2)	
Age groups			1
0–27 days	2 (0.3)	20 (0.3)	
28 days to 23 months	26 (3.8)	260 (3.8)	
2-11 years	4 (0.6)	40 (0.6)	
12–17 years	4 (0.6)	40 (0.6)	
18–44 years	92 (13.6)	920 (13.6)	
45–64 years	187 (27.5)	1870 (27.5)	
65–74 years	144 (21.2)	1440 (21.2)	
≥75 years	220 (32.4)	2200 (32.4)	
Sex			1
Male	356 (52.4)	3560 (52.4)	
Female	323 (47.6)	3230 (47.6)	
Report type			<0.0001
Spontaneous	618 (91.0)	6205 (91.4)	
Report from study	54 (8.0)	317 (4.7)	
Other	7 (1.0)	267 (3.9)	
Not available to sender (unknown)	0 (0.0)	1 (0.0)	
/accines type			
COVID-19 vaccines	626 (92.2)	6331 (93.2)	0.3039
Tozinameran	376 (55.4)	3324 (49.0)	0.0014
Elasomeran	78 (11.5)	642 (9.5)	0.0871
Vaxzevria	129 (19.0)	1492 (22.0)	0.073
Ad26.COV2.S	8 (1.2)	224 (3.3)	0.0024
Gam-COVID-Vac	0 (0.0)	2 (0.0)	0.6547
NVX-CoV2373	0 (0.0)	3 (0.0)	1.000
Influenza vaccines	34 (5.0)	123 (1.8)	<0.0001
Pneumococcal vaccines	11 (1.6)	105 (1.6)	0.8824
Drug known to cause ILD included in the ICSRs*	. ,	. ,	
Cancer therapy	17 (2.5)	10 (0.2)	<0.0001
	27 (4.0)	25 (0.4)	< 0.0001
Rheumatology therapy			
Rheumatology therapy Anti-infection agent	3 (0.4)	3 (0.0)	0.0121

Continued

Table 1 Continued

	ILD cases (N=679)	Non-cases (N=6790)	
	N (%)	N (%)	P value
Serious			<0.0001
Yes	625 (92.1)	2323 (34.5)	
Seriousness			<0.0001
Death	116 (17.1)	287 (4.2)	
Life threatening	97 (14.3)	162 (2.4)	
Caused/Prolonged hospitalisation	285 (42.0)	582 (8.6)	
Disabling/incapacitating	13 (1.9)	91 (1.3)	
Congenital anomaly/birth defect	0 (0.0)	3 (0.0)	
Other	114 (16.8)	1218 (17.9)	
Region			<0.0001
African	4 (0.6)	364 (5.4)	
Americas	54 (8.0)	632 (9.3)	
South-East Asia	6 (0.9)	121 (1.8)	
European	577 (85.0)	4398 (64.8)	
Eastern Mediterranean	8 (1.2)	342 (5.0)	
Western Pacific	30 (4.4)	933 (13.7)	
Notifier type			< 0.0001
Physician	527 (77.6)	3675 (54.1)	
Pharmacist	50 (7.4)	1055 (15.5)	
Other health professional	102 (15.0)	2060 (30.3)	
Time to onset (case=574, non-case=6064)			0.0077
Mean±SD	32.7±64.7	26.6±57.3	
Median (Q1–Q3)	7 (1–35)	1 (0–15)	

*Cancer therapy: bleomycin; gemcitabine; epidermal growth factor receptor-targeted agent (erlotinib, gefitinib, panitumumab, cetuximab); mammalian target of rapamycin-inhibitor (everolimus, temsirolimus, sirolimus); immune checkpoint inhibitor (nivolumab, pembrolizumab, avelumab, durvalumab, ipilimumab), rheumatology drugs: methotrexate; leflunomide, biological disease-modifying anti-rheumatic drugs (tumour necrosis factor) agent (infliximab, etanercept, adalimumab), tocilizumab, rituximab), anti-infection agents (nitrofurantoin, daptomycin, interferon), cardiology drugs (amiodarone, bepridil, statin (lovastatin, simvastatin, pravastatin, atorvastatin, fluvastatin, rosuvastatin, cerivastatin)).

ICSRs, individual case safety reports.

reported in September 2021, while the first report was from January 2021. The number of reports decreased steadily until the end of the study. Serious AE reports accounted for 84.4% to 100% of all ILD cases following COVID-19 vaccination (figure 2).

We identified characteristics of ILD cases by COVID-19 vaccines, influenza vaccines and other vaccines. The COVID-19 vaccines contained reports from European and had the longest median time-to-onset (1 (IQR 1–36) day) than the influenza vaccine (5 (IQR 2–23)) and others (6 (IQR 1–21)) (online supplemental table 4). The most frequently reported AEs with regard to PT or LLT were pneumonitis (134 ICSRs, 19.7%), ILD (70 ICSRs, 10.3%) and interstitial pneumonia (46 ICSRs, 6.8%) in ILD cases (online supplemental table 5). AEs that included both narrow and broad terms were aligned with the AEs from narrow terms. Details of ILD cases are provided in online supplemental tables 4 and 5.

Case/non-case analysis

The results of ILD cases/non-cases, including those of primary, secondary and subgroup analyses, are shown in table 2. The ROR of ILD following COVID-19 vaccination was 0.86 (95% CI 0.64 to 1.15) compared with other vaccines. The ROR of mRNA vaccines was 0.99 (95% CI 0.73 to 1.34), tozinameran was 0.98 (95% CI 0.73 to 1.33) and elasomeran was 1.04 (95% CI 0.71 to 1.50). Moreover, no signal of disproportionate reporting was observed in viral vector COVID-19 vaccines (viral vector COVID-19 vaccines ROR 0.69 (95% CI 0.50 to 0.96); Vaxzevria ROR 0.75 (95% CI 0.53 to 1.05) and Ad26.COV2.S ROR 0.32 (95% CI 0.15 to 0.67)) (table 2).

In the subgroup analysis of primary analysis, we did not find an increased reporting of ILD according to age groups, sex and region (table 3, online supplemental table 6). The ILD following COVID-19 vaccination was not associated with a disproportionality signal regardless

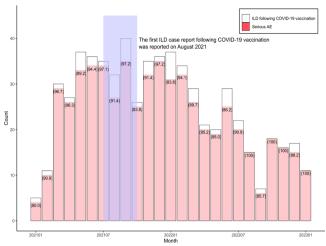


Figure 2 The number of ILD cases following COVID-19 vaccination during the study period. The brackets () present the proportion of serious AE among ICSRs reported ILD following COVID-19 vaccination. AE, adverse event; ICSR, individual case safety report; ILD, interstitial lung disease.

of age groups (under 65 years; ROR 0.94 (95% CI 0.65 to 1.35), 65 years and older; ROR 0.70 (95% CI 0.41 to 1.17). The COVID-19 vaccination emerged with no signal in both males and females (males ROR 0.81 (95% CI 0.55 to 1.19), females ROR 0.93 (95% CI 0.59 to 1.46)). There was no signal when stratifying Western Pacific region and the other regions (ROR 0.42 (95% CI 0.16 to 1.14) and ROR 0.88 (95% CI 0.65 to 1.21).

The results of the sensitivity analyses were similar to those of the primary analysis (figure 3, online supplemental table 7). There was no disproportionality signal when considering diverse AE definitions and potential biases (figure 3, online supplemental table 7). The ROR of influenza vaccines was 0.44 (95% CI 0.27 to 0.71). The results of sensitivity analysis 7 compared with influenza a vaccines were consistent with the primary analysis (online

DISCUSSION

supplemental tables 8 and 9)

The present study aimed to identify the characteristics of ILD following COVID-19 vaccination and the disproportionality between COVID-19 vaccines and ILD using the global pharmacovigilance database. We identified 679 ILD cases from VigiBase defined using MedDRA SMQ and performed disproportionality analysis. To the best of our knowledge, this is the first study to investigate the signals of disproportionate reporting of ILD associated with COVID-19 vaccines. Compared with other vaccines, no significant signal of disproportional reporting of ILD was observed for COVID-19 vaccines. These findings were consistent across several analyses conducted after considering potential biases. Moreover, the signal of disproportionality was not detected when compared with the influenza vaccine which is known to induce ILD.

In our study, reports received from European accounted for the majority of ILD cases (85.0%) following COVID-19 vaccination. In contrast to the present study, most ILD cases following COVID-19 vaccination have been reported in South-East Asia, including South Korea and Japan since Park et al reported the first ILD case following mRNA COVID-19 vaccination.^{15 16 18-20} Kono et al suggested that South-East Asian population should be carefully monitored since it is at a high risk of COVID-19 vaccine-related ILD.³⁰ Among 30 cases of ILD identified following COVID-19 vaccination in the Western Pacific, which is classified as Asia by WHO, and the signal of ILD was not detected when compared with other vaccines (ROR 1.68, 95% CI 0.68 to 4.16) (table 2). However, the result of subgroup analysis according to the region should be interpreted with caution because of the small number of cases and incomplete information on ICSRs.

be of analysis	ILD cases	Non-cases	ROR (95% CI)
mary analysis (cases: 679, non-cases: 679	0)		
The other vaccines	53 (7.8)	459 (6.8)	Reference
COVID-19 vaccines	626 (92.2)	6331 (93.2)	0.86 (0.64 to 1.15)
mRNA COVID-19 vaccines	448 (66.0)	3912 (57.6)	0.99 (0.73 to 1.34)
Tozinameran	373 (58.9)	3282 (54.4)	0.98 (0.73 to 1.33)
Elasomeran	73 (11.5)	611 (10.1)	1.04 (0.71 to 1.50)
Viral vector COVID-19 vaccines	137 (20.2)	1718 (25.3)	0.69 (0.50 to 0.96)
Vaxzevria	126 (19.9)	1461 (24.2)	0.75 (0.53 to 1.05)
Gam-COVID-Vac	0 (0.0)	2 (0.0)	NC
Ad26.COV2.S	8 (1.3)	220 (3.6)	0.32 (0.15 to 0.67)
Protein-based COVID-19 vaccines	0 (0.0)	3 (0.0)	NC
NVX-CoV2373	0 (0.0)	3 (0.1)	NC
Others	41 (6.0)	698 (10.3)	0.51 (0.33 to 0.78)

ype of analysis	ILD cases	Non-cases	ROR (95% CI)
Subgroup analysis			
Age			
Age <65 (case=315, non-case=	3150)		
The other vaccines	36 (11.4)	339 (10.8)	Reference
COVID-19 vaccines	268 (85.1)	2447 (77.7)	0.94 (0.65 to 1.35)
Age ≥65 (case=364, non-case=	3640)		
The other vaccines	17 (4.7)	120 (3.3)	Reference
COVID-19 vaccines	347 (95.3)	3520 (96.7)	0.70 (0.41 to 1.17)
Gender			
Male (case=356, non-case=356	60)		
The other vaccines	31 (8.7)	254 (7.1)	Reference
COVID-19 vaccines	325 (91.3)	3306 (92.9)	0.81 (0.55 to 1.19)
Female (case=323, non-case=3	3230)		
The other vaccines	22 (6.8)	205 (6.4)	Reference
COVID-19 vaccines	301 (93.2)	3025 (93.7)	0.93 (0.59 to 1.46)
Region			
Western Pacific region (case=3	0, non-case=933)		
The other vaccines	5 (16.7)	73 (7.8)	Reference
COVID-19 vaccines	25 (83.3)	860 (92.2)	0.42 (0.16 to 1.14)
The other regions (case=649, n	on-case=5857)		
The other vaccines	48 (7.4)	386 (6.6)	Reference
COVID-19 vaccines	601 (92.6)	5471 (93.4)	0.88 (0.65 to 1.21)

A previous systematic review of drug-induced ILD iden-

tified male has been as a risk factor for drug-induced ILD, especially in those treated with amiodarone, methotrexate, epidermal growth factor receptor tyrosine kinase inhibitor (EGFR-TKI) and premetrexed.³¹ Males were predominant in previous case reports of ILD related

Type of analysis	ILD cases	Non-ILD cases		ROR (95% CI)
Sensitivity analysis 1: ICS	SRs reported as	suspected, concomita	nt, and interaction dru	ıgs
The other vaccines	53 (7.8)	449 (6.6)	Reference	1
COVID-19 vaccine	626 (92.2)	6,338 (93.4)	0.84 (0.62-1.13)	
Sensitivity analysis 2: Ext	tended the ILD d	efinition (broad search	ר)	
The other vaccines	67 (6.1)	521 (4.8)	Reference	
COVID-19 vaccine	1,026 (93.9)	10,409 (95.2)	0.77 (0.59-1.00)	
Sensitivity analysis 3: IC	SRs excluding re	ports included drugs	known to cause drug-	induced ILD
The other vaccines	43 (7.1)	416 (7.2)	Reference	
COVID-19 vaccine	538 (92.9)	5,374 (92.8)	1.02 (0.73-1.42)	
Sensitivity analysis 4: ICS	SRs excluding re	ports included COVID	-19 vaccine AESIs	
The other vaccines	43 (7.2)	414 (6.9)	Reference	
COVID-19 vaccine	554 (92.8)	5,556 (93.1)	0.96 (0.69-1.33)	
Sensitivity analysis 5: Re	stricted to seriou	s AE reports		
The other vaccines	47 (7.5)	344 (5.5)	Reference	
COVID-19 vaccine	578 (92.5)	5,906 (94.5)	0.72 (0.52-0.98)	
Sensitivity analysis 6: Re	ported before Au	gust 6, 2021		
The other vaccines	10 (5.1)	76 (3.9)	Reference	
COVID-19 vaccine	187 (94.9)	1,894 (96.1)	0.75 (0.38-1.48)	
Sensitivity analysis 7: De	finition of Influen	za vaccine as a positi	ve control	
Influenza vaccines	20 (3.3)	89 (1.5)	Reference	
COVID-19 vaccine	576 (96.7)	5,971 (98.5)	0.44 (0.27-0.71)	_

Figure 3 Reporting OR (ROR) of sensitivity analysis. AE, adverse event; AESI, adverse event of special event; ICSR, individual case safety report; ILD, interstitial lung disease; MedDRA, medical dictionary for regulatory activities; SMQ, standardised MedDRA queries.

to COVID-19 vaccination.^{15 16 18–20} However, this study observed no signal of disproportionate reporting regardless of sex (males (ROR 0.81,95% CI 0.55 to 1.19), females (ROR 0.93,95% CI, 0.59 to 1.46)) (online supplemental table 6). Further studies are required to identify the risk according to demographic characteristics.

We analysed the data of spontaneous reporting systems to assess signals of AE of COVID-19 vaccination. The spontaneous reporting systems have several biases due to factors that could affect reporting, which results in incorrect signal detection. These biases can be notoriety bias, information bias, selection bias and competition bias.^{25 32 33} We implemented different minimisation strategies against these biases. First, we designed a primary analysis to address factors that could lead to information bias by considering to be suspected, healthcare professionals and complete information on age groups and sex. Moreover, in sensitivity analysis 2, we used MedDRA SMQ with narrow and broad terms. This result was in line with the primary analysis that showed no signal of disproportionate reporting (ROR 0.77, 95% CI 0.59 to 1.00). Second, for competition bias, it is necessary to eliminate factors associated with vaccines/AEs of interest (sensitivity analyses 3, 4). Results derived from sensitivity analysis considering competition biases showed that copyright.

COVID-19 vaccines had no disproportionality signal of ILD compared with the other vaccines (ROR 1.02, 95% CI 0.73 to 1.42 and ROR 0.96, 95% CI, 0.69 to 1.33, respectively). Third, in pharmacovigilance, temporal bias (the Weber effect or notoriety bias) refers to variation in the number of reports after a specific event, such as safety alerts and market authorisation. The signal of ILD was not observed when minimising temporal biases; the RORs of sensitivity analyses 5 and 6 were 0.72 (95%) CI 0.52 to 0.98) and 0.75 (95% CI 0.38 to 1.48), respectively. Fourth, we defined reference groups that received influenza and other vaccines instead of all other drugs to avoid selection bias. The analysis using influenza vaccines as a positive control in the secondary analysis showed that COVID-19 vaccines emerged with no signal when compared with influenza vaccines (ROR 0.44, 95% CI 0.27 to 0.71). However, the risk-benefits of COVID-19 vaccines should be carefully assessed.

The mechanisms of COVID-19 vaccine-induced ILD are unclear. To date, both cytotoxicity and immunemediated lung injury are considered as main mechanisms that initiate drug-induced ILD. Although it is rare, the event can be fatal and patients might require hospitalisation.³⁴ According to previous reports, the influenza vaccination can induce ILD by increasing the levels of inflammatory cytokines.¹² Several cases of COVID-19 mRNA vaccine associated ILD have been reported.¹⁵⁻²¹ Given the similar clinical characteristics with influenza vaccine-induced ILD, including onset time, chest CT findings and responsiveness to corticosteroids, it can be speculated that ILD following COVID-19 vaccination might also be due to immune-mediated pulmonary injury. These studies suggest that COVID-19 vaccination induces immune-mediated injury to the lungs through T-cells, which adopt a predominant type one phenotype in susceptible patients.^{17–21 35} However, further studies with a large number of ILD patients who received the COVID-19 vaccines are needed.^{12 15}

This study has several limitations. First, selective reporting of AEs might have been compromised in the spontaneous reporting database although we strived to minimise biases. During the pandemic, the number of ICSRs following COVID-19 vaccination increased rapidly, which might have resulted in differential reporting rates and influenced parameters. We applied 1:10 exact matching to reduce the imbalance between case and non-case and performed various analyses. The results of our study did not provide exhaustivity of COVID-19 vaccine-induced ILD although it suggests focusing on the risk. Second, we analysed ICSRs without causality assessment. However, VigiBase contains essential information required for causality assessment, including age, sex, primary reporter and time-to-onset. Third, concerns on the validity of ILD in spontaneous reporting database might be raised. To overcome this limitation, we restricted physicians (77.6%), pharmacists (7.4%) and other health professionals (14.8%) as notifier types and defined ILD using MedDRA SMQs, which are validated by expert

discussion. Fourth, in the present study, the majority of ILD cases following vaccination were predominantly in the European population, which may introduce bias due to population heterogeneity. Fifth, previous studies have suggested that COVID-19 infection can lead to the occurrence or exacerbation of ILD, referred to as post-COVID-19 ILD. Notably, the VigiBase we used cannot ascertain the COVID-19 infection status. Therefore, the study findings should be interpreted with caution. Finally, this study did not assess the risk of specific molecular components of vaccines. The excipients such as adjuvants, stabilisers, preservatives and trace components can cause AE following immunisation. Therefore, besides vaccines, the safety of excipients should also be evaluated. Despite these limitations, our study used a global pharmacovigilance database with over 30 million ICSRs and could offer additional hypotheses for AEs. In addition, the case/non-case approach allowed us to study rare AEs and could represent the use of drugs in real world settings.²⁵ Since there were no population-based studies and previous case reports have included exacerbation of pre-existing ILD with death, additional safety studies are needed.

CONCLUSION

In conclusion, we identified no significant disproportionality signal of ILD associated with COVID-19 vaccines using global pharmacovigilance database. This finding is consistent regardless of the subpopulation. Furthermore, the disproportional analysis compared with the influenza vaccines that are known to cause ILD emerged with no signal. However, serious AE accounted for the majority of ILD cases following COVID-19 vaccination and events, including hospitalisations, have been reported. We suggest careful monitoring of COVID-19 vaccine-induced ILD. This study may provide information that can be useful for making decisions on subsequence COVID-19 vaccine strategies. However, further studies using patientlevel information such as disease history and diagnostic test results are required for more conclusive evidence.

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Contributors M-TL and S-YJ participated in the conception and design of study. M-TL and S-YJ participated in the data acquisition and data analysis. M-TL, JWL, JCC, K-MG and S-YJ participated in the data interpretation. M-TL participated in the draft of the manuscript. JWL, HJL, J-ML, JCC and K-MG helped to revise the manuscript for intellectual content. All authors read and approved the final manuscript. S-YJ is reponsible for the overall content as guarantor.

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Competing interests None declared.

Patient and public involvement Patients and/or the public were not involved in the design, or conduct, or reporting, or dissemination plans of this research.

Patient consent for publication Not applicable.

Ethics approval This study was conducted in accordance with the Declaration of Helsinki. The study protocol was approved for exemption from review by the Institutional Review Board of Chung-Ang University (IRB number: 1041078-201903-HR-071-01), because this study analyaed a secondary database. Informed consent from subjects was waived due to the database containing anonymised data that cannot identify study subjects.

Provenance and peer review Not commissioned; externally peer reviewed.

Data availability statement Data may be obtained from a third party and are not publicly available. The data analysed in this study are available from VigiBase upon formal request to the Uppsala Monitoring Centre at the WHO Collaborating Centre for International Drug Monitoring.

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Supplementary Table 1. The adverse events and MedDRA code used in the study.

Adverse event	MedDRA code
ILD – narrow terms	SMQ code: 20000042
	(including MedDRA code: 10000001, 10000686, 10001028,
	10001713, 10001714, 10001881, 10001889, 10001890, 10001891,
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	10004942, 10006448, 10007701, 10008832, 10009164, 10011495,
	10014574, 10014952, 10014957, 10014962, 10015887, 10016221,
	10016222, 10016640, 10016641, 10016649, 10016656, 10016659,
	10016660, 10019107, 10020463, 10021225, 10021240, 10022611,
	10022612, 10022617, 10022618, 10022619, 10025088, 10025089,
	10025102, 10025103, 10026724, 10026725, 10026816, 10026819,
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	10035743, 10035750, 10035751, 10035754, 10035755, 10036028,
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	10045623, 10047130, 10047263, 10048016, 10048594, 10049202,
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	10060779, 10060902, 10061473, 10061924, 10062997, 10063725,
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	10082838, 10082999, 10083303, 10084211, 10084305, 10084309,
	10084872, 10085058, 10085188, 10085189, 10085190, 10085191,
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	10085668, 10085686, 10086041, 10086072, 10085117, 10085122,
	10086597, 10086598, 10086601, 10086926, 10086935, 10087294)
ILD – broad terms	SMQ code: 20000042
ILD – broad terms	
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Allaphylaxis	10002394, 10002395, 10002218, 10042930, 10042931, 10009320, 10002394, 10002395, 10002424, 10002425, 10002426, 10018257,
	10018259, 10020198, 10046744, 10001734, 10001718, 10001719,
	10001738
Thrombocytopenia	10043551, 10043554, 10043555, 10043560, 10043569, 10039884,
	10024922, 10035528, 10035529, 10035545, 10038213, 10043546,
	10074667, 10021243, 10021245, 10023095, 10051064, 10043552,
	10043558
Generalized convulsion	10010916, 10010917, 10018079, 10010914, 10010904, 10010906,
	10010922, 10039906, 10039910, 10016731, 10016735, 10039910,
	10009234, 10010915, 10010923, 10010926, 10016733, 10003628,
	10071377
Guillain Barré Syndrome	10018767, 10018766, 10042812, 10000813, 10067898, 10033803,
	10049567
Aseptic meningitis	10003458, 10027201, 10029669, 10057724, 10047469, 10046236,
	10027260, 10027262, 10028263, 10028273, 10027250, 10027220,
	10027213, 10027214, 10027222, 10027215, 10027229, 10027211,
	10011253
Encephalitis / Encephalomyelitis /	10014581, 10014601, 10014588, 10056198, 10014602, 10054373,
Acute disseminated encephalomyelitis	10028524, 10028526, 10028527, 10000709
Bell's Palsy	10004223, 10033559, 10016060, 10016062, 10033808
Vaccine associated enhanced disease	10085491, 10085001

Supplementary Table 2. The types and WHODrug recode number codes used in the study.

Type of drugs	WHODrug record number							
COVID-19 vaccines								
Tozinameran	15268625001, 15268625002, 15268625003, 15268625004, 15268625005, 15268625006, 15268625007, 15268625008, 15268625009							
Vaxzevria	15268603001, 15268603002, 15268603003, 15268603004, 15268603005,							
	15268603006, 15268603007, 15268603008, 15268603009, 15268603010,							
	15268603012, 15268603013, 15268603014							
Elasomeran	15268604001, 15268604002, 15268604003, 15268604004, 15268604005,							
2100011101011	15268604006, 15268604007, 15268604008							
Ad26.COV2.S	15268640001, 15268640002, 15268640003, 15268640004, 15268640005,							
	15268640006, 15268640007							
Gam-COVID-Vac	15268619005, 15268684002, 15268619007, 15268620004							
NVX-CoV2373	15268611001, 15268611002, 15268611003, 15268611004, 15268611005							
GBP510	15268665003							
Cancer therapy								
Bleomycin	001839xxxxx (94)							
Gemcitabine	012157xxxxx (326)							
Erlotinib	016114xxxxx (93)							
Gefitinib	015601xxxxx (92)							
Panitumumab	017773xxxxx (5)							
Cetuximab	014905xxxxx (5)							
Everolimus	015069xxxxx (61)							
Temsirolimus	017165xxxxx (6)							
Sirolimus	014591xxxxx (35)							
Nivolumab	078729xxxx (4)							
Pembrolizumab	083428xxxxx (6)							
Avelumab	089186xxxxx (3)							
Durvalumab	087572xxxxx (4)							
Ipilimumab	057456xxxxx (7)							
Rhematology therapy								
methotrexate	001138xxxxx (416)							
Leflunomide	014148xxxxx (214)							
Infliximab	014456xxxx (24)							
Etanercept	014906xxxxx (25)							
Adalimumab	016129xxxxx (40)							
Tocilizumab	017591xxxxx (11)							
Rituximab Anti-infection	014025xxxxx (60)							
	000244/mmm (220)							
Nitrofurantoin	000244xxxxx (230)							
Daptomycin	016772xxxx (57)							
Interferon	005968xxxxx (4)							
Cardiology drugs	001221							
Amiodarone	001331xxxxx (293)							
Bepridil	007889xxxx (11)							
Lovastatin	008967xxxx (249)							
Simvastatin	008481xxxxx (946)							
Pravastatin	008804xxxxx (381)							
Atorvastatin	013261xxxxx (1225)							
Fluvastatin	012245xxxxx (146)							

Rosuvastatin	015886xxxxx (999)			
Cerivastatin	013413xxxxx (28)			
The bracket () indicates the numbers of drugs.				

Supplementary Table 3. Analysis strategies

Analysis types	Case	Non-case	Vaccine of interest	Other vaccines	Comment
Primary analysis	ILD narrow terms	Non-ILD	COVID-19 vaccines	The other vaccines	
Subgroup analysis	ILD narrow terms	Non-ILD	COVID-19 vaccines	The other vaccines	
1. Age	ILD narrow terms	Non-ILD	COVID-19 vaccines	The other vaccines	<65 and ≥65 years
2. Gender	ILD narrow terms	Non-ILD	COVID-19 vaccines	The other vaccines	Male and female
3. Region	ILD narrow terms	Non-ILD	COVID-19 vaccines	The other vaccines	Western Pacific region and the other regions
Sensitivity analysis					
1. ICSRs reported as suspected, concomitant, and interaction drugs	ILD narrow terms	Non-ILD	COVID-19 vaccines	The other vaccines	Diverse AE definition
2. Extended the ILD definition (broad search)	ILD narrow+broad	Non-ILD	COVID-19 vaccines	The other vaccines	Diverse AE definition (Information bias)
3. ICSRs excluding reports included either cancer therapy or rheumatology drugs or anti-infection agents, cardiology drugs [†]	ILD narrow terms	Non-ILD	COVID-19 vaccines	The other vaccines	Drug competition bias
4. ICSRs excluding reports included COVID-19 vaccine AESIs	ILD narrow terms	Non-ILD	COVID-19 vaccines	The other vaccines	Event competition bias
5. Restricted to serious AE reports	ILD narrow terms	Non-ILD	COVID-19 vaccines	The other vaccines	Weber effect
6. ICSRs before August 6, 2021	ILD narrow terms	Non-ILD	COVID-19 vaccines	The other vaccines	Notoriety bias
7. Definition of influenza vaccine as positive control	ILD narrow terms	Non-ILD	COVID-19 vaccines	Influenza vaccine	Positive control

⁺ Cancer therapy: bleomycin; gemcitabine; Epidermal Growth Factor Receptor (EGFR)-targeted agent (erlotinib, gefitinib, panitumumab, cetuximab); mammalian target of rapamycin (MTOR)-inhibitor (everolimus, temsirolimus, sirolimus); immune checkpoint inhibitor (nivolumab, pembrolizumab, avelumab, durvalumab, ipilimumab), rheumatology drugs: methotrexate; leflunomide, biological disease-modifying anti-rheumatic drugs (DMARDs) (tumor necrosis factor (TNF) agent (infliximab, etanercept, adalimumab), tocilizumab, rituximab), anti-infection agents (nitrofurantoin, daptomycin, interferon), cardiology drugs (amiodarone, bepridil, statin (lovastatin, simvastatin, pravastatin, atorvastatin, fluvastatin, rosuvastatin, cerivastatin))

ILD, interstitial lung disease; COVID-19, coronavirus disease 2019; AESI, adverse event of special interest

Supplementary Table 4. Characteristics of interstitial lung disease (ILD) by COVID-19 vaccines, influenza vaccines, and others.

	COVID-19	vaccines	Influenza vaccines		Others	
	Ν	%	Ν	%	Ν	%
Reported quarter (Q)						
2020.4Q (Dec. 13,2020~Dec. 31, 2020)	0	(0.0)	0	(0.0)	0	(0.0)
2021.1Q (Jan. 1, 2021 ~ Mar. 31, 2021)	46	(7.4)	3	(8.8)	1	(3.0)
2021.2Q (Apr. 1, 2021 ~ Jun. 30, 2021)	100	(16.0)	5	(14.7)	1	(3.0)
2021.3Q (Jul. 1, 2021 ~ Sep. 30, 2021)	107	(17.1)	8	(23.5)	2	(6.1)
2021.4Q (Oct. 1, 2021 ~ Dec. 31, 2021)	97	(15.5)	3	(8.8)	6	(18.2)
2022.1Q (Jan. 1, 2022 ~ Mar. 31, 2022	100	(16.0)	4	(11.8)	1	(3.0)
2022.2Q (Apr. 1, 2022 ~ Jun. 30, 2022)	70	(11.2)	4	(11.8)	8	(24.2)
2022.3Q (Jul. 1, 2022 ~ Sep. 30, 2022)	44	(7.0)				
2022.4Q (Oct. 1, 2022 ~ Dec. 31, 2022)	51	(8.2)	4	(11.8)	13	(39.4)
2023.1Q (Jan. 1, 2023 ~ Jan. 26, 2023)	11	(1.8)	3	(8.8)	1	(3.0)
Age groups				<i>、 ,</i>		. ,
0~27 days			2	(5.9)		
28 days to 23 months	2	(0.3)	3	(8.8)	21	(63.6)
2~11 years	2	(0.0)	1	(2.9)	3	(9.1)
12~17 years	4	(0.6)	-	()	5	(0.1)
18~44 years	91	(14.5)	1	(2.9)	1	(3.0)
45~64 years	182	(29.1)	6	(17.7)	2	(6.1)
65~74 years	134	(21.4)	11	(32.4)	4	(12.1)
\geq 75 years	213	(34.0)	10	(29.4)	2	(6.1)
Sex	215	(34.0)	10	(29.4)	Ζ	(0.1)
Male	325	(51.9)	18	(52.9)	20	(60.6)
Female	323	(48.1)	16	(32.9)	13	(39.4)
	301	(40.1)	10	(47.1)	13	(39.4)
Report type	E 70	(92.3)	20	(85.3)	22	(69.7)
Spontaneous	578	• •	29	. ,	23	• •
Report from study	41 7	(6.6)	5	(14.7)	10	(30.3)
Other Serious	/	(1.1)				
	E 70	(02.2)	21	(91.2)	20	(87.9)
Yes	578	(92.3)	31	(91.2)	29	(87.9)
Seriousness	107	(47.4)	-	(22.5)	-	(4 = 0)
Death	107	(17.1)	7	(20.6)	5	(15.2)
Life threatening	96	(15.3)	3	(8.8)		
Caused/Prolonged Hospitalization	260	(41.5)	14	(41.2)	16	(48.5)
Disabling/Incapacitating	13	(2.1)	1	(2.9)		
Congenital anomaly/Birth defect						
Other	102	(16.3)	6	(17.7)	8	(24.2)
Region						
African	4	(0.6)				
Americas	42	(6.7)	9	(26.5)	5	(15.2)
South-East Asia	5	(0.8)			1	(3.0)
European	543	(86.7)	20	(58.8)	25	(75.8)
Eastern Mediterranean	7	(1.1)	1	(2.9)		
Western Pacific	25	(4.0)	4	(11.8)	2	(6.1)
Notifier type				·		
Physician	493	(78.8)	24	(70.6)	21	(63.6)
Pharmacist	45	(7.2)	3	(8.8)	3	(9.1)
Pharmacist						

Mean±SD	33.5±61.7	17.1±24.7	40.9±128.7
Median (Q1-Q3)	8 (1-36)	5 (2-23)	6 (1-21)

SD; standard deviation

COVID-19 vaccines contained Tozinameran, Elasomeran, Vaxzevira, Ad26.COV2.S, Gam-COVID-Vac, and NVX-CoV2373. Influenza vaccines were defined using Anatomical Therapeutic Chemical (ATC) classification; J07BB.

Narrow + broad terms

Count

134

%

12.3

MedDRA terms

T	Pheumonitis	154	19.7	Pheumonius	154	12.5
2	Interstitial lung disease	70	10.3	Acute respiratory distress syndrome	128	11.7
3	Interstitial pneumonia	46	6.8	Sarcoidosis	83	7.6
4	Pulmonitis	45	6.6	Interstitial lung disease	70	6.4
5	Lung infiltration	37	5.4	Interstitial pneumonia	46	4.2
6	Interstitial pneumonitis	32	4.7	Pulmonitis	45	4.1
7	Bronchiolitis	30	4.4	Lung infiltration	37	3.4
8	Pneumonia interstitial	30	4.4	Interstitial pneumonitis	32	2.9
9	Ground glass opacity in thoracic CT	27	4.0	Pneumonia interstitial	30	2.7
10	Pulmonary fibrosis	19	2.8	Bronchiolitis	30	2.7
11	Lung inflammation	18	2.7	Cryptogenic organising pneumonia	29	2.7
12	Eosinophilic pneumonia	17	2.5	ARDS	27	2.5
13	Lung opacity	16	2.4	Ground glass opacity in thoracic CT	27	2.5
14	Eosinophilic granulomatosis with polyangiitis	16	2.4	Pulmonary alveolar haemorrhage	21	1.9
15	Pulmonary infiltration	15	2.2	Pulmonary fibrosis	19	1.7
16	Alveolitis	13	1.9	Lung inflammation	18	1.6
17	Hypersensitivity pneumonitis	12	1.8	Eosinophilic pneumonia	17	1.6
18	Bilateral pulmonary infiltrates	10	1.5	Eosinophilic granulomatosis with polyangiitis	16	1.5
19	Exacerbation of idiopathic pulmonary fibrosis	10	1.5	Lung opacity	16	1.5
20	Churg Strauss syndrome	9	1.3	Pulmonary infiltration	15	1.4
21	Lung fibrosis	8	1.2	Adult respiratory stress syndrome	14	1.3
22	Alveolitis allergic	7	1.0	Alveolitis	13	1.2
23	Idiopathic pulmonary fibrosis	6	0.9	Granulomatosis with polyangiitis	13	1.2
24	Pulmonary vasculitis	6	0.9	Hypersensitivity pneumonitis	12	1.1
25	Acute respiratory distress syndrome	6	0.9	Organizing pneumonia	11	1.0
26	ARDS	5	0.7	A.R.D.S.	11	1.0
27	Pneumonia eosinophilic	4	0.6	Antisynthetase syndrome	10	0.9
28	Interstitial pneumonia aggravated	4	0.6	Bilateral pulmonary infiltrates	10	0.9
29	Diffuse alveolar damage	4	0.6	Exacerbation of idiopathic pulmonary fibrosis	10	0.9
30	Rheumatoid arthritis- associated interstitial lung	4	0.6	Churg Strauss syndrome	9	0.8

Supplementary Table 5. The list of reported adverse events (AEs) in narrow and broad terms (>2 report).

%

19.7 Pneumonitis

Count

134

Narrow terms

MedDRA terms

Pneumonitis

disease

X-ray

Fibrosis lung

Lymphocytic alveolitis

Ground glass opacity on chest

31

32

33

No.

1

13

Polyarteritis nodosa

Cryptogenic organizing

Lung fibrosis

0.6

0.4

0.4

4

3

3

9

8

8

0.8

0.7

0.7

				pneumonia		
34	Lung infiltration NOS	3	0.4	Pulmonary sarcoidosis	7	0.6
35	Interstitial lung fibrosis	3	0.4	Goodpasture's syndrome	7	0.6
36	Acute interstitial pneumonitis	3	0.4	Organising pneumonia	7	0.6
37	Pulmonary alveolar haemorrhage	3	0.4	Wegener's granulomatosis	7	0.6
38				Alveolitis allergic	7	0.6
39				Loefgren syndrome	6	0.5
40				Pulmonary vasculitis	6	0.5
41				Idiopathic pulmonary fibrosis	6	0.5
42				Adult respiratory distress syndrome	6	0.5
43				Rheumatoid arthritis- associated interstitial lung disease	4	0.4
44				Ground glass opacity on chest X-ray	4	0.4
45				Interstitial pneumonia aggravated	4	0.4
46				Diffuse alveolar damage	4	0.4
47				Granulomatous polyangiitis	4	0.4
48				Pneumonia eosinophilic	4	0.4
49				Syndrome respiratory distress adult	4	0.4
50				Periarteritis nodosa	4	0.4
51				Lung injury	3	0.3
52				Lung infiltration NOS	3	0.3
53				Wegeners granulomatosis	3	0.3
54				Pulmonary renal syndrome	3	0.3
55				Acute interstitial pneumonitis	3	0.3
56				Pulmonary granuloma	3	0.3
57				Pulmonary alveolar hemorrhage	3	0.3
58				Interstitial lung fibrosis	3	0.3
59				Fibrosis lung	3	0.3
60				Lymphocytic alveolitis	3	0.3
61			1	Lung transplant rejection	3	0.3

* only included in narrow and broad terms

MedDRA, medical dictionary for regulatory activities

Type of analysis

Non-cases

ROR (95% CI)

Type of analysis	ILD-0	.ases	NOII-	Lases	KUK (95% CI)
Subgroup analysis 1, age					
Age<65 (case=315, non-case=3,150)					
The other vaccines	36	(11.4)	339	(10.8)	
COVID-19 vaccines	268	(85.1)	2,447	(77.7)	0.94 (0.65–1.35)
mRNA COVID-19 vaccines	198	(62.9)	1,611	(51.1)	1.16 (0.80–1.68)
Tozinameran	158	(52.2)	1,328	(48.1)	1.12 (0.77–1.64)
Elasomeran	40	(13.2)	272	(9.8)	1.39 (0.86–2.23)
Viral vector COVID-19 vaccines	70	(22.2)	837	(26.6)	0.79 (0.52-1.20)
Vaxzevria	64	(21.1)	676	(24.5)	0.89 (0.58–1.37)
Gam-COVID-Vac	0	(0.0)	1	(0.0)	NC
Ad26.COV2.S	5	(1.7)	147	(5.3)	0.32 (0.12-0.83)
Protein-based COVID-19 vaccines	0	(0.0)	1	(0.0)	NC
NVX-CoV2373	0	(0.0)	1	(0.0)	NC
Others	11	(3.5)	362	(11.5)	0.29 (0.14–0.57)
Age≥65 (case=364, non-case=3,640)					
The other vaccines	17	(4.7)	120	(3.3)	
COVID-19 vaccines	347	(95.3)	3,520	(96.7)	0.70(0.41-1.17)
mRNA COVID-19 vaccines	250	(68.7)	2,301	(63.2)	0.77(0.45-1.30)
Tozinameran	215	(65.2)	1,954	(59.7)	0.78(0.46-1.32)
Elasomeran	33	(10.0)	339	(10.4)	0.69(0.37-1.28)
Viral vector COVID-19 vaccines	67	(18.4)	881	(24.2)	0.54(0.31-0.95)
Vaxzevria	62	(18.8)	785	(24.0)	0.56(0.32-0.99)
Gam-COVID-Vac	0	(0.0)	1	(0.0)	NC
Ad26.COV2.S	3	(0.9)	73	(2.2)	0.29(0.08-1.02)
Protein-based COVID-19 vaccines	0	(0.0)	2	(0.1)	NC
NVX-CoV2373	0	(0.0)	2	(0.1)	NC
Others	30	(8.2)	336	(9.2)	0.63(0.34-1.18)
Subgroup analysis 2, gender					
Male (case=356, non-case=3,230)					
The other vaccines	31	(8.7)	254	(7.1)	
COVID-19 vaccines	325	(91.3)	3,306	(92.9)	0.81(0.55–1.19)
mRNA COVID-19 vaccines	235	(66.0)	2,009	(56.4)	0.96(0.65-1.43)
Tozinameran	194	(57.9)	1,697	(54.2)	0.94(0.63-1.40)
Elasomeran	40	(11.9)	304	(9.7)	1.08(0.66–1.77)
Viral vector COVID-19 vaccines	71	(19.9)	898	(25.2)	0.65(0.42-1.01)
Vaxzevria	67	(20.0)	739	(23.6)	0.74(0.47-1.16)
Gam-COVID-Vac	0	(0.0)	2	(0.1)	NC
Ad26.COV2.S	3	(0.9)	137	(4.4)	0.18(0.05–0.60)
Protein-based COVID-19 vaccines	0	(0.0)	0	(0.0)	NC
NVX-CoV2373	0	(0.0)	0	(0.0)	NC
Others	19	(5.3)	399	(11.2)	0.39(0.22-0.71)
Female (case=305, non-case=3,050)					
The other vaccines	22	(6.8)	205	(6.4)	-
COVID-19 vaccines	301	(93.2)	3,025	(93.7)	0.93(0.59–1.46)
mRNA COVID-19 vaccines	213	(65.9)	1,903	(58.9)	1.04(0.66–1.66)
Tozinameran	22	(7.4)	205	(7.1)	1.05(0.66–1.68)
Elasomeran	179	(60.1)	1,585	(54.6)	1.00(0.57–1.77)
Viral vector COVID-19 vaccines	66	(20.4)	820	(25.4)	0.75(0.45–1.24)
Vaxzevria	59	(19.8)	722	(24.9)	0.76(0.46–1.27)
Gam-COVID-Vac	0	(0.0)	0	(0.0)	NC

Supplementary Table 6. Reporting odds ratio (ROR) of subgroups analyses in primary analysis

ILD-cases

Ad26.COV2.S	5	(1.7)	83	(2.9)	0.56(0.21-1.53)
Protein-based COVID-19 vaccines	0	(0.0)	3	(0.1)	<.001 (<.001->999.9)
NVX-CoV2373	0	(0.0)	3	(0.1)	<.001 (<.001->999.9)
Others	22	(6.8)	299	(9.3)	0.69(0.37-1.27)
Subgroup analysis 3, region					
Western Pacific Region (case=30, non-case=	933)				
The other vaccines	5	(16.7)	73	(7.8)	
COVID-19 vaccines	25	(83.3)	860	(92.2)	0.42(0.16-1.14)
mRNA COVID-19 vaccines	14	(46.7)	359	(38.5)	0.57(0.20-1.63)
Tozinameran	13	(48.2)	316	(45.6)	0.60(0.21-1.74)
Elasomeran	1	(3.7)	43	(6.2)	0.34(0.04-3.00)
Viral vector COVID-19 vaccines	10	(33.3)	267	(28.6)	0.55(0.18-1.65)
Vaxzevria	8	(29.6)	219	(31.6)	0.53(0.17-1.68)
Gam-COVID-Vac	0	(0.0)	0	(0.0)	NC
Ad26.COV2.S	0	(0.0)	42	(6.1)	NC
Protein-based COVID-19 vaccines	0	(0.0)	0	(0.0)	NC
NVX-CoV2373	0	(0.0)	0	(0.0)	NC
Others	1	(3.3)	234	(25.1)	0.06(0.01-0.54)
The other regions (case=649, non-case=5,85	57)				
The other vaccines	48	(7.4)	386	(6.6)	
COVID-19 vaccines	601	(92.6)	5,471	(93.4)	0.88(0.65-1.21)
mRNA COVID-19 vaccines	434	(66.9)	3,553	(60.7)	0.98(0.72–1.35)
Tozinameran	360	(59.4)	2,966	(55.5)	0.98(0.71-1.34)
Elasomeran	72	(11.9)	568	(10.6)	1.02(0.69–1.50)
Viral vector COVID-19 vaccines	127	(19.6)	1,451	(24.8)	0.70(0.50-1.00)
Vaxzevria	118	(19.5)	1,242	(23.2)	0.76(0.54-1.09)
Gam-COVID-Vac	0	(0.0)	2	(0.0)	NC
Ad26.COV2.S	8	(1.3)	178	(3.3)	0.36(0.17-0.78)
Protein-based COVID-19 vaccines	0	(0.0)	3	(0.1)	NC
NVX-CoV2373	0	(0.0)	3	(0.1)	NC
Others	40	(6.2)	464	(7.9)	0.69(0.45-1.08)

COVID-19, coronavirus disease 2019; ILD, interstitial lung disease; ROR, reporting odds ratio; NC, not calculated

Supplementary Table 7. Reporting odds ratio (ROR) of sensitivity analyses

Type of analysis		ases	Non-	cases	ROR (95% CI)				
Sensitivity analysis 1 (case=644, non-cas	se=6,440)								
The other vaccines	53	(7.8)	449	(6.6)					
COVID-19 vaccines	626	(92.2)	6,338	(93.4)	0.84(0.62-1.13)				
mRNA COVID-19 vaccines	448	(66.0)	3,863	(56.9)	0.98(0.73–1.33)				
Tozinameran	373	(58.9)	3,265	(54.3)	0.97(0.71-1.31)				
Elasomeran	73	(11.5)	582	(9.7)	1.06(0.73–1.55)				
Viral vector COVID-19 vaccines	137	(20.2)	1,743	(25.7)	0.67(0.48–0.93)				
Vaxzevria	126	(19.9)	1,469	(24.4)	0.73(0.52-1.02)				
Gam-COVID-Vac	0	(0.0)	4	(0.1)	NC				
Ad26.COV2.S	8	(1.3)	241	(4.0)	0.28(0.13-0.60)				
Protein-based COVID-19 vaccines	0	(0.0)	2	(0.0)	NC				
NVX-CoV2373	0	(0.0)	2	(0.0)	NC				
Others	41	(6.0)	730	(10.8)	0.48(0.31–0.73)				
Sensitivity analysis 2 (case=548, non-case=5,480)									
The other vaccines	67	(6.1)	521	(4.8)					
COVID-19 vaccines	1,026	(93.9)	10,409	(95.2)	0.77(0.59–1.00)				
mRNA COVID-19 vaccines	704	(64.4)	6,340	(58.0)	0.86(0.66–1.13)				
Tozinameran	604	(60.0)	5,326	(55.0)	0.88(0.68–1.15)				
Elasomeran	95	(9.4)	983	(10.1)	0.75(0.54–1.05)				
Viral vector COVID-19 vaccines	251	(23.0)	2,906	(26.6)	0.67(0.51–0.89)				
Vaxzevria	221	(22.0)	2,483	(25.6)	0.69(0.52–0.92)				
Gam-COVID-Vac	1	(0.1)	4	(0.0)	1.94(0.21–17.65)				
Ad26.COV2.S	19	(1.9)	372	(3.8)	0.40(0.24–0.67)				
Protein-based COVID-19 vaccines	0	(0.0)	3	(0.0)	NC				
NVX-CoV2373	0	(0.0)	3	(0.0)	NC				
Others	71	(6.5)	1,160	(10.6)	0.48(0.34–0.68)				
Sensitivity analysis 3 (case=567, non-cas	se=6,131)								
The other vaccines	41	(7.1)	416	(7.2)					
COVID-19 vaccines	538	(92.9)	5,374	(92.8)	1.02(0.73–1.42)				
mRNA COVID-19 vaccines	391	(67.5)	3,264	(56.4)	1.22(0.87–1.70)				
Tozinameran	327	(59.9)	2,746	(53.5)	1.21(0.86–1.70)				
Elasomeran	62	(11.4)	498	(9.7)	1.26(0.83–1.91)				
Viral vector COVID-19 vaccines	117	(20.2)	1,497	(25.9)	0.79(0.55–1.15)				
Vaxzevria	109	(20.0)	1,283	(25.0)	0.86(0.59–1.26)				
Gam-COVID-Vac	0	(0.0)	3	(0.1)	NC				
Ad26.COV2.S	7	(1.3)	186	(3.6)	0.38(0.17–0.87)				
Protein-based COVID-19 vaccines	0	(0.0)	1	(0.0)	NC				
NVX-CoV2373	0	(0.0)	1	(0.0)	NC				
Others	30	(5.2)	612	(10.6)	0.50(0.31–0.81)				
Sensitivity analysis 4 (case=584, non-cas	se=5,840)								
The other vaccines	43	(7.2)	414	(6.9)					
COVID-19 vaccines	554	(92.8)	5,556	(93.1)	0.96(0.69–1.33)				
mRNA COVID-19 vaccines	398	(66.7)	3,461	(58.0)	1.11(0.80–1.54)				
Tozinameran	330	(59.4)	2,883	(54.5)	1.10(0.79–1.54)				
Elasomeran	66	(11.9)	559	(10.6)	1.14(0.76–1.70)				
Viral vector COVID-19 vaccines	119	(19.9)	1,474	(24.7)	0.78(0.54–1.12)				
Vaxzevria	110	(19.8)	1,241	(23.5)	0.85(0.59–1.24)				
Gam-COVID-Vac	0	(0.0)	1	(0.0)	<.001 (<.001->999.9)				
Ad26.COV2.S	7	(1.3)	193	(3.7)	0.35(0.15–0.79)				
Protein-based COVID-19 vaccines	0	(0.0)	0	(0.0)					

NVX-CoV2373	0	(0.0)	0	(0.0)	-
Others	37	(6.2)	621	(10.4)	0.57(0.36-0.91)
Sensitivity analysis 5 (case=591, non-cas	se=5,910)				
The other vaccines	47	(7.5)	344	(5.5)	
COVID-19 vaccines	578	(92.5)	5,906	(94.5)	0.72(0.52-0.98)
mRNA COVID-19 vaccines	417	(66.7)	4,078	(65.3)	0.75(0.54-1.03)
Tozinameran	347	(59.4)	3,582	(60.5)	0.71(0.51-0.98)
Elasomeran	68	(11.6)	477	(8.1)	1.04(0.70-1.55)
Viral vector COVID-19 vaccines	125	(20.0)	1,558	(24.9)	0.59(0.41-0.84)
Vaxzevria	115	(19.7)	1,323	(22.3)	0.64(0.44-0.91)
Gam-COVID-Vac	0	(0.0)	0	(0.0)	NC
Ad26.COV2.S	7	(1.2)	194	(3.3)	0.26(0.12-0.60)
Protein-based COVID-19 vaccines	0	(0.0)	1	(0.0)	NC
NVX-CoV2373	0	(0.0)	1	(0.0)	NC
Others	36	(5.8)	269	(4.3)	0.98(0.62-1.56)
Sensitivity analysis 6 (case=197, non-cas	se=1,970)				
The other vaccines	10	(5.1)	76	(3.9)	
COVID-19 vaccines	187	(94.9)	1,894	(96.1)	0.75(0.38-1.48)
mRNA COVID-19 vaccines	123	(62.4)	1,164	(59.1)	0.80(0.41-1.59)
Tozinameran	109	(56.8)	1,025	(56.8)	0.81(0.41-1.61)
Elasomeran	14	(7.3)	139	(7.7)	0.77(0.32-1.81)
Viral vector COVID-19 vaccines	59	(30.0)	567	(28.8)	0.79(0.39–1.61)
Vaxzevria	57	(29.7)	554	(30.7)	0.78(0.38-1.60)
Gam-COVID-Vac	0	(0.0)	1	(0.1)	NC
Ad26.COV2.S	2	(1.0)	11	(0.6)	1.38(0.27–7.16)
Protein-based COVID-19 vaccines	0	(0.0)	0	(0.0)	NC
NVX-CoV2373	0	(0.0)	0	(0.0)	NC
Others	5	(2.5)	163	(8.3)	0.23(0.08-0.71)
Sensitivity analysis 7 (case=606, non-cas	se=6,060)				
Influenza vaccines	20	(3.3)	89	(1.5)	Reference
COVID-19 vaccines	586	(96.7)	5,971	(98.5)	0.44(0.27-0.71)
mRNA COVID-19 vaccines	448	(73.9)	4,071	(67.2)	0.49(0.30-0.80)
Tozinameran	373	(62.2)	3,376	(56.5)	0.49(0.30-0.81)
Elasomeran	73	(12.2)	669	(11.2)	0.49(0.28-0.84)
Viral vector COVID-19 vaccines	137	(22.6)	1,893	(31.2)	0.32(0.19–0.54)
Vaxzevria	126	(21.0)	1,544	(25.8)	0.36(0.22–0.61)
Gam-COVID-Vac	0	(0.0)	6	(0.1)	NC
Ad26.COV2.S	8	(1.3)	292	(4.9)	0.12(0.05-0.29)
Protein-based COVID-19 vaccines	0	(0.0)	1	(0.0)	NC
NVX-CoV2373	0	(0.0)	1	(0.0)	NC
Others	1	(0.2)	6	(0.1)	0.74(0.09-6.51)

COVID-19, coronavirus disease 2019; ILD, interstitial lung disease; ROR, reporting odds ratio; NC, not calculated sensitivity analysis 1: included ICSRs of COVID-19 vaccines as suspected, concomitant, and interaction; sensitivity analysis 2: defined ILD including MedDRA SMQ narrow and broad terms (broad search); sensitivity analysis 4: excluded ICSRs that included drugs known to cause drug-induced ILD; sensitivity analysis 3: excluded reports containing adverse event special interest (AESI) of COVID-19 vaccines; sensitivity analysis 5: restricted to ICSRs reported serious AE following COVID-19 vaccination; sensitivity analysis 6: containing ICSRs with a reporting date before August 6, 2021; sensitivity analysis 7: compared the COVID-19 vaccines with the influenza vaccines as a positive control.

	ILD cases (N=606)		Non-cases (N=6,060)		
	N	%	(N=0 N	,000) %	<i>p</i> -value
Reported quarter (Q)		70		70	0.0216
2020.4Q (Dec. 13,2020~Dec. 31, 2020)	0	(0.0)	10	(0.2)	
2021.1Q (Jan. 1, 2021 ~ Mar. 31, 2021)	47	(7.8)	645	(10.6)	
2021.2Q (Apr. 1, 2021 ~ Jun. 30, 2021)	100	(16.5)	1,152	(19.0)	
2021.3Q (Jul. 1, 2021 ~ Sep. 30, 2021)	110	(18.2)	847	(14.0)	
2021.4Q (Oct. 1, 2021 ~ Dec. 31, 2021)	96	(15.8)	887	(14.6)	
2022.1Q (Jan. 1, 2022 ~ Mar. 31, 2022	98	(16.2)	843	(13.9)	
2022.2Q (Apr. 1, 2022 ~ Jun. 30, 2022)	66	(10.9)	773	(12.8)	
2022.3Q (Jul. 1, 2022 ~ Sep. 30, 2022)	39	(6.4)	408	(6.7)	
2022.4Q (Oct. 1, 2022 ~ Dec. 31, 2022)	40	(6.6)	426	(7.0)	
2023.1Q (Jan. 1, 2023 ~ Jan. 26, 2023)	10	(1.7)	69	(1.1)	
Age groups		(=)		()	1
0~27 days	2	(0.3)	20	(0.3)	-
28 days to 23 months	4	(0.7)	40	(0.7)	
2°11 years	1	(0.2)	10	(0.2)	
12~17 years	4	(0.7)	40	(0.7)	
18~44 years	90	(14.9)	900	(14.9)	
45~64 years	176	(29.0)	1,760	(29.0)	
65~74 years	127	(21.0)	1,270	(23.0)	
\geq 75 years	202	(33.3)	2,020	(33.3)	
Sex	202	(33.3)	2,020	(55.5)	1
Male	318	(52.5)	3,180	(52.5)	T
Female	288	(47.5)	2,880	(32.5)	
Report type	200	(47.5)	2,000	(47.5)	<.0001
Spontaneous	567	(93.6)	5,651	(93.3)	1.0001
Report from study	33	(5.5)	182	(3.0)	
Other	6	(1.0)	226	(3.7)	
Not available to sender (unknown)	0	(0.0)	220	(0.0)	
Vaccines type	0	(0.0)		(0.0)	
COVID-19 vaccines	586	(96.7)	5,971	(98.5)	0.0007
Tozinameran	376		3,437	(56.7)	
Elasomeran	78	(62.1)	5,437 715		0.0115 0.4368
Vaxzevria	129	(12.9)	1,589	(11.8) (26.2)	0.4368
		(21.3)			
Ad26.COV2.S	0	(0.0)	6	(0.1)	0.4384
Gam-COVID-Vac NVX-CoV2373	8 0	(1.3)	298 1	(4.9)	<.0001
Influenza vaccines	34	(0.0) (5.6)	133	(0.0)	1 <.0001
Pneumococcal vaccines	34 5	(5.6) (0.8)	133	(2.2) (0.2)	<.0001 0.0005
	5	(0.8)	9	(0.2)	0.0005
Drug known to cause ILD included in the ICSRs ⁺	1 -	(2 5)	n	(0,0)	< 0001
Cancer therapy	15	(2.5) (2.5)	2	(0.0)	<.0001
Rheumatology therapy	21	(3.5) (0.5)	25	(0.4)	<.0001
Anti-infection agent	3	(0.5)	6	(0.1)	0.0414
Cardiology drugs	55	(9.1)	244	(4.0)	<.0001
Serious		(02.5)	2.244	(20.1)	<.0001
Yes	561	(92.6)	2,311	(38.1)	
Seriousness				(15.0)	<.0001
Death	267	(4.4)	98	(16.2)	

Supplementary Table 8. Characteristic of interstitial lung disease (ILD) and non-ILD of sensitivity analysis 7 (compared COVID-19 vaccines with influenza vaccines)

Life threatening	128	(2.1)	95	(15.7)	
Caused/Prolonged Hospitalization	576	(9.5)	254	(41.9)	
Disabling/Incapacitating	88	(1.5)	13	(2.2)	
Congenital anomaly/Birth defect	1	(0.0)	0	(0.0)	
Other	1,251	(20.6)	101	(16.7)	
Region					<.0001
African	4	(0.7)	333	(5.5)	
Americas	32	(5.3)	507	(8.4)	
South-East Asia	4	(0.7)	124	(2.1)	
European	533	(88.0)	4,276	(70.6)	
Eastern Mediterranean	6	(1.0)	127	(2.1)	
Western Pacific	27	(4.5)	693	(11.4)	
Notifier type					<.0001
Physician	484	(79.9)	3,555	(58.7)	
Pharmacist	43	(7.1)	929	(15.3)	
Other Health Professional	79	(13.0)	1,576	(26.0)	
Time to onset (case=549, non-case=5,743)					0.2784
Mean±SD	32.5±59.4		29.6±58.7		
Median (Q1-Q3)	8	(2-36)	1	(0-18)	

ILD; interstitial lung disease, SD; standard deviation

[†] Cancer therapy: bleomycin; gemcitabine; Epidermal Growth Factor Receptor (EGFR)-targeted agent (erlotinib, gefitinib, panitumumab, cetuximab); mammalian target of rapamycin (MTOR)-inhibitor (everolimus, temsirolimus, sirolimus); immune checkpoint inhibitor (nivolumab, pembrolizumab, avelumab, durvalumab, ipilimumab), rheumatology drugs: methotrexate; leflunomide, biological disease-modifying anti-rheumatic drugs (DMARDs) (tumor necrosis factor (TNF) agent (infliximab, etanercept, adalimumab), tocilizumab, rituximab), anti-infection agents (nitrofurantoin, daptomycin, interferon), cardiology drugs (amiodarone, bepridil, statin (lovastatin, simvastatin, pravastatin, atorvastatin, fluvastatin, rosuvastatin, cerivastatin))

Type of analysis	ILD-0	ILD-cases		cases	ROR (95% CI)	
Subgroup analysis 1, age						
Age<65 (case=277, non-case=2,770)						
The other vaccines	9	(3.3)	33	(1.2)		
COVID-19 vaccines	268	(96.8)	2,737	(98.8)	0.36(0.17–0.76)	
mRNA COVID-19 vaccines	198	(71.5)	1,783	(64.4)	0.41(0.19-0.86)	
Tozinameran	1,460	(53.4)	158	(57.3)	0.40(0.19-0.84)	
Elasomeran	308	(11.3)	40	(14.5)	0.48(0.21-1.07)	
Viral vector COVID-19 vaccines	70	(25.3)	953	(34.4)	0.27(0.12-0.59)	
Vaxzevria	743	(27.2)	64	(23.2)	0.32(0.15-0.69)	
Gam-COVID-Vac	4	(0.2)	0	(0.0)	NC	
Ad26.COV2.S	188	(6.9)	5	(1.8)	0.10(0.03-0.31)	
Protein-based COVID-19 vaccines	0	(0.0)	0	(0.0)	NC	
NVX-CoV2373	0	(0.0)	0	(0.0)	NC	
Others	0	(0.0)	1	(0.0)	NC	
Age≥65 (case=329, non-case=3,290)		()		()		
The other vaccines	11	(3.3)	56	(1.7)		
COVID-19 vaccines	318	(96.7)	3,234	(98.3)	0.50(0.26-0.97)	
mRNA COVID-19 vaccines	250	(76.0)	2,288	(69.5)	0.56(0.29–1.08)	
Tozinameran	215	(66.4)	1,916	(59.1)	0.57(0.30-1.11)	
Elasomeran	33	(10.2)	361	(11.1)	0.47(0.22-0.97	
Viral vector COVID-19 vaccines	67	(20.4)	940	(28.6)	0.36(0.18-0.73)	
Vaxzevria	62	(19.1)	801	(24.7)	0.39(0.20-0.79)	
Gam-COVID-Vac	0	(0.0)	2	(0.1)	NC	
Ad26.COV2.S	3	(0.9)	104	(3.2)	0.15(0.04-0.55)	
Protein-based COVID-19 vaccines	0	(0.0)	1	(0.0)	NC	
NVX-CoV2373	0	(0.0)	- 1	(0.0)	NC	
Others	1	(0.3)	- 5	(0.2)	1.02(0.11-9.59)	
Subgroup analysis 2, gender		(0.0)		(=-)		
Male (case=318, non-case=3,180)						
The other vaccines	11	(3.5)	47	(1.5)		
COVID-19 vaccines	307	(96.5)	3,133	(98.5)	0.42(0.22-0.82)	
mRNA COVID-19 vaccines	235	(73.9)	2,107	(66.3)	0.48(0.24-0.93)	
Tozinameran	194	(61.6)	1,739	(55.4)	0.48(0.24–0.93)	
Elasomeran	40	(12.7)	356	(11.3)	0.48(0.23-1.00)	
Viral vector COVID-19 vaccines	71	(22.3)	1,026	(32.3)	0.30(0.15-0.60)	
Vaxzevria	67	(21.3)	810	(25.8)	0.35(0.18-0.71)	
Gam-COVID-Vac	0	(0.0)	4	(0.1)	0.33(0.16 0.71) NC	
Ad26.COV2.S	3	(0.0)	182	(5.8)	0.07(0.02-0.26)	
Protein-based COVID-19 vaccines	0	(0.0)	102	(0.0)	0.07(0.02-0.20) NC	
NVX-CoV2373	0	(0.0)	0	(0.0)	NC	
Others	1	(0.0)	0	(0.0)	NC	
Female (case=288, non-case=2,880)	T	(0.5)	0	(0.0)		
The other vaccines	9	(3.1)	42	(1.5)		
COVID-19 vaccines	279		2,838			
mRNA COVID-19 vaccines	279	(96.9) (74.0)	2,838 1,964	(98.5) (68.2)	0.46(0.22–0.95) 0.51(0.24–1.05)	
	213 179	(74.0) (62.8)	1,964	(68.2)		
Tozinameran Elasomeran	33	(62.8)	•	(57.7)	0.51(0.24 - 1.07)	
		(11.6)	313	(11.0)	0.49(0.22-1.10)	
Viral vector COVID-19 vaccines	66 50	(22.9)	867 724	(30.1)	0.36(0.17-0.76)	
Vaxzevria Gam-COVID-Vac	59	(20.7) (0.0)	734	(25.9) (0.1)	0.38(0.17–0.81) NC	
	0	10.01	2	(0.1)	INC.	

Supplementary Table 9. Reporting odds ratio (ROR) of subgroups analyses in sensitivity analysis 7

Ad26.COV2.S	5	(1.8)	110	(3.9)	0.21(0.07-0.67)
Protein-based COVID-19 vaccines	0	(0.0)	110	(0.0)	0.21(0.07-0.07) NC
NVX-CoV2373	0	(0.0)	1	(0.0)	NC
Others	0	(0.0)	6	(0.2)	NC
Subgroup analysis 3, region	0	(0.0)	0	(0.2)	NC .
Western Pacific Region (case=27, non-cas	e=693)				
The other vaccines	3	(11.1)	17	(2.5)	
COVID-19 vaccines	24	(88.9)	676	(97.6)	0.20(0.06-0.73)
mRNA COVID-19 vaccines	14	(51.9)	376	(54.3)	0.21(0.06–0.80)
Tozinameran	13	(52.0)	330	(48.1)	0.22(0.06–0.86)
Elasomeran	1	(4.0)	46	(6.7)	0.12(0.01-1.27)
Viral vector COVID-19 vaccines	10	(37.0)	298	(43.0)	0.19(0.05-0.76)
Vaxzevria	8	(32.0)	240	(35.0)	0.19(0.05-0.78)
Gam-COVID-Vac	0	(0.0)	0	(0.0)	NC
Ad26.COV2.S	0	(0.0)	52	(7.6)	NC
Protein-based COVID-19 vaccines	0	(0.0)	1	(0.1)	NC
NVX-CoV2373	0	(0.0)	1	(0.2)	NC
Others	0	(0.0)	1	(0.1)	NC
The other regions (case=579, non-case=5	,367)				
The other vaccines	17	(2.9)	72	(1.3)	
COVID-19 vaccines	562	(97.1)	5,295	(98.7)	0.45(0.26-0.77)
mRNA COVID-19 vaccines	434	(75.0)	3,695	(68.9)	0.50(0.29–0.85)
Tozinameran	360	(62.6)	3,046	(57.6)	0.50(0.29–0.86)
Elasomeran	72	(12.5)	623	(11.8)	0.49(0.27-0.88)
Viral vector COVID-19 vaccines	127	(21.9)	1,595	(29.7)	0.34(0.19–0.59)
Vaxzevria	118	(20.5)	1,304	(24.7)	0.38(0.22–0.67)
Gam-COVID-Vac	0	(0.0)	6	(0.1)	NC
Ad26.COV2.S	8	(1.4)	240	(4.5)	0.14(0.06-0.34)
Protein-based COVID-19 vaccines	0	(0.0)	0	(0.0)	NC
NVX-CoV2373	0	(0.0)	0	(0.0)	NC
Others	1	(0.2)	5	(0.1)	0.85(0.09–7.73)

COVID-19, coronavirus disease 2019; ILD, interstitial lung disease; ROR, reporting odds ratio; NC, not calculated