Table S1 QC details in PA medical records

Serial number	Diagnostic classification	Defect type	Defect detail
1	CPA	Examination	Chest CT not done
2	CPA	Diagnosis	History of less than 3 months
3	CPA	Treatment	No antifungal therapy
4	CPA	Treatment	Itraconazole was not used as the antifungal drug of first choice
5	ABPA	Treatment	Unreasonable treatment of choice, no oral corticosteroids
6	ABPA	Treatment	No antifungal prescribed
7	ABPA	Treatment	Unreasonable treatment of choice, without itraconazole as antifungal drug of first choice
8	CCPA	Examination	Chest CT not done
9	CCPA	Treatment	No antifungal prescribed
10	CCPA	Treatment	Itraconazole and Voriconazole were not used as antifungal drugs of first choice
11	IPA	Treatment	No antifungal prescribed
12	IPA	Treatment	Voriconazole not used as antifungal drug of Itraconazole was not used as the antifungal drug of first choice
13	IPA	Treatment	Echinocandins cannot be used as the first choice of treatment
14	IA	Treatment	When there is no contraindication, voriconazole is not selected as the drug of choice in patients with acquired immunodeficiency (HIV) and IA
15	IA	Treatment	Did not switch to caspofungin when resistance to voriconazole developed in patients with parenchymal organ transplantation and IA
16	IA	Treatment	In patients with IA treated with voriconazole, trough serum concentrations were not measured after 2 to 5 days of treatment
17	IA	Treatment	Patients with parenchymal organ transplantation and IA do not discontinue voriconazole and switch to caspofungin when they experience liver function impairment
18	CFPA	Examination	Chest CT not done
19	CFPA	Diagnosis	Lack of diagnostic evidence, severe fibrosis in at least two lobes
20	SA	Examination	Chest CT not done
21	AN	Examination	Chest CT not done
22	AN	Diagnosis	CT chest without visible nodule
23	SAIA	Examination	Chest CT not done
24	SAIA	Examination	Tissue microbiological test was not performed
25	CNPA	Treatment	No antifungal prescribed
26	CNPA	Treatment	voriconazole was not used as the antifungal drug of first choice

QC, quality control; PA, pulmonary aspergillosis; CPA, chronic pulmonary aspergillosis; ABPA, allergic bronchopulmonary aspergillosis; CCPA, chronic cavitary pulmonary aspergillosis; IPA, invasive pulmonary aspergillosis; IA, invasive aspergillosis; CFPA, chronic fiberoptic pulmonary aspergillosis; SA, simple pulmonary aspergillosis; AN, Aspergillus nodule; SAIA, subacute invasive aspergillosis; CNPA, chronic necrotizing pulmonary aspergillosis; CT, computed tomography.

Table S2 Reference for quality control points

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Serial number	Guide details	Reference guide
1	The diagnosis of CPA requires a combination of characteristics: a consistent appearance in thoracic imaging (preferably by CT)	2015 ERS and ESCMID guideline [32]
2	The diagnosis of CPA requires a combination of characteristics: one or more cavities with or without a fungal ball present or nodules on thoracic imaging, either direct evidence of Aspergillus infection (culture or microscopy from biopsy) or an IgG antibody response to Aspergillus spp. and exclu_x0002_ sion of alternative diagnoses (especially mycobacterial infection), all present for at least 3 months	2017 ESCMID-ECMM-ERS guideline [6]
3	Table 38	2017 ESCMID-ECMM-ERS guideline [6]
4	Table 38	2017 ESCMID-ECMM-ERS guideline [6]
5	Corticosteroids are a cornerstone of therapy for exacerbations (Table 1)	2016 IDSA guideline [5]
6	Table 1	2016 IDSA guideline [5]
7	Table 1	2016 IDSA guideline [5]
8	84. Oral itraconazole and Voriconazole are the preferred oral antifungal agents (strong recommendation; high-quality evi_x0002_dence); posaconazole is a useful third-line agent for those with adverse events or clinical failure (strong recommendation; moderate-quality evidence)	2016 IDSA guideline [5]
9	84. Oral itraconazole and Voriconazole are the preferred oral antifungal agents (strong recommendation; high-quality evi_x0002_dence); posaconazole is a useful third-line agent for those with adverse events or clinical failure (strong recommendation; moderate-quality evidence)	2016 IDSA guideline [5]
10	81. The diagnosis of CCPA requires: (i) 3 months of chronic pul_x0002_ monary symptoms or chronic illness or progressive radiologic radiographic abnormalities, with cavitation, pleural thickening, pericavitary infiltrates, and sometimes a fungal ball; (ii) Aspergillus IgG antibody elevated or other microbiological data; and (iii) no or minimal immunocompromise, usually with one or more underlying pulmonary disorders. The Aspergillus IgG an tibody test is the most sensitive microbiological test (strong rec ommendation; moderate-quality evidence). Sputum Aspergillus PCR testing is more sensitive than culture (weak recommenda tion; moderate-quality evidence)	2016 IDSA guideline [5]
11	25. We recommend primary treatment with Voriconazole (strong recommendation; high-quality evidence)	2016 IDSA guideline [5]
12	25. We recommend primary treatment with Voriconazole (strong recommendation; high-quality evidence)	2016 IDSA guideline [5]
13	28. Combination antifungal therapy with Voriconazole and an echinocandin may be considered in select patients with docu_x005f mented IPA (weak recommendation; moderate-quality evidence)	2016 IDSA guideline [5]
	29. Primary therapy with an echinocandin is not recommend ed (strong recommendation; moderate-quality evidence). Echinocandins (micafungin or caspofungin) can be used in settings in which azole and polyene antifungals are contrain dicated (weak recommendation; moderate-quality evidence)	
14	Table 32	2017 ESCMID-ECMM-ERS guideline [6]
15	Table 32	2017 ESCMID-ECMM-ERS guideline [6]
16	All patients receiving Voriconazole prophylaxis for IA, Measure serum trough	2017 ESCMID-ECMM-ERS guideline [6]
	level after 2–5 days of therapy or soon after, and 4 days after change of dose	
17	Table 32	2017 ESCMID-ECMM-ERS guideline [6]
18	Severe fibrotic destruction of at least two lobes of lung complicating CCPA leading to a major loss of lung function. Severe fibrotic destruction of one lobe with a cavity is simply referred to as CCPA affecting that lobe. Usually the fibrosis is manifest as consolidation, but large cavities with surrounding fibrosis may be seen	2016 IDSA guideline [5]
19	Severe fibrotic destruction of at least two lobes of lung complicating CCPA leading to a major loss of lung function. Severe fibrotic destruction of one lobe with a cavity is simply referred to as CCPA affecting that lobe. Usually the fibrosis is manifest as consolidation, but large cavities with surrounding fibrosis may be seen	2016 IDSA guideline [5]
20	Single pulmonary cavity containing a fungal ball, with serological or microbiological evidence implicating Aspergillus spp. in a non- immunocompromised patient with minor or no symptoms and no radiological progression over at least 3 months of observation	2016 IDSA guideline [5]
21	Aspergillus nodules, which may be single or multiple, may mimic malignancy as well as nodules seen in rheumatoid arthritis, coccidioidomycosis, tuberculosis, non-tuberculous mycobacterial infection and, rarely, actinomycosis or rheumatoid arthritis. Typically, Aspergillus nodules appear rounded, some with low attenuation or cavitation within. Some are spiculated, a common feature of carcinoma	2017 ESCMID-ECMM-ERS guideline [6]
22	Aspergillus nodules, which may be single or multiple, may mimic malignancy as well as nodules seen in rheumatoid arthritis, coccidioidomycosis, tuberculosis, non-tuberculous mycobacterial infection and, rarely, actinomycosis or rheumatoid arthritis. Typically, Aspergillus nodules appear rounded, some with low attenuation or cavitation within. Some are spiculated, a common feature of carcinoma	2017 ESCMID-ECMM-ERS guideline [6]
23	SAIA should be diagnosed according to established definitions of invasive aspergillosis in immunocompromised patients (or highly debilitated patients), with a slower course than acute invasive aspergillosis (1–3 months), and commonly with both detectable Aspergillus antibody and antigen in the serum. Histological confirmation derives from seeing hyphae invading lung parenchyma	2016 IDSA guideline [5]
24	SAIA should be diagnosed according to established definitions of invasive aspergillosis in immunocompromised patients (or highly debilitated patients), with a slower course than acute invasive aspergillosis (1–3 months), and commonly with both detectable Aspergillus antibody and antigen in the serum. Histological confirmation derives from seeing hyphae invading lung parenchyma	2015 ERS and ESCMID guideline [32]
25	Voriconazole preferred for CNPA and patients with fungal balls to minimize risk of resistance; Voriconazole Start 150–200 mg bid, adjust with TDM	2017 ESCMID-ECMM-ERS guideline [6]
26	Voriconazole preferred for CNPA and patients with fungal balls to minimize risk of resistance; Voriconazole Start 150–200 mg bid, adjust with TDM	2017 ESCMID-ECMM-ERS guideline [6]

CPA, chronic pulmonary aspergillosis; CT, computed tomography; CCPA, chronic cavitary pulmonary aspergillosis; PCR, polymerase chain reaction; IPA, invasive pulmonary aspergillosis; IA, invasive aspergillosis; SAIA, subacute invasive aspergillosis; CNPA, chronic necrotizing pulmonary aspergillosis; TDM, therapeutic drug monitoring.

Lable 05 111 medical records diagnostic county reference

Diagnostic classification	Term coding	Code source
PA	B44	ICD10
CPA	733171006	SNOMED CT V20190131
ABPA	B44.101+	ICD10
CCPA	733171006&2483006	SNOMED CT V20190131
IPA	3214003&2704003, 3214003&89187006, B44.0	SNOMED CT V20190131, ICD10
IA	721798004	SNOMED CT V20190131
CFPA	733171006&112674009	SNOMED CT V20190131
SA	6042001&13673007	SNOMED CT V20190131
AN	6042001&27925004	SNOMED CT V20190131
SAIA	782761005	SNOMED CT V20190131
CNPA	782761005	SNOMED CT V20190131

PA, pulmonary aspergillosis; CPA, chronic pulmonary aspergillosis; ABPA, allergic bronchopulmonary aspergillosis; CCPA, chronic cavitary pulmonary aspergillosis; IPA, invasive pulmonary aspergillosis; IA, invasive aspergillosis; CFPA, chronic fiberoptic pulmonary aspergillosis; SA, simple pulmonary aspergillosis; AN, Aspergillus nodule; SAIA, subacute invasive aspergillosis; CNPA, chronic necrotizing pulmonary aspergillosis.

Classification of PA	Number of QC points to be verified [®]	Number of medical records to be verified	Total number of QC points verification*
ABPA	3	37	111
SA	1	17	17
IPA	7	64	448
CCPA	3	19	57
CFPA	2	4	8
CPA	4	59	236
Overall calculation	20	200	877

Table S4 Results of manual verification of 200 medical records from QCSA

⁸, the number of quality control points to be verified included in QCSA for each diagnostic classification; *, the result of multiplying the number of quality control points for each classification by the number of medical records that fit that classification. QCSA, quality control system for pulmonary aspergillosis; PA, pulmonary aspergillosis; QC, quality control; ABPA, allergic bronchopulmonary aspergillosis; SA, simple pulmonary aspergillosis; IPA, invasive pulmonary aspergillosis; CCPA, chronic cavitary pulmonary aspergillosis; CFPA, chronic fiberoptic pulmonary aspergillosis; CPA, chronic pulmonary aspergillosis.



Figure S1 QCSA multi-role workflow. QCSA, quality control system for pulmonary aspergillosis.



Figure S2 Approach to forming QCSA using guidelines. There are four steps in the construction of the QCSA. Step 1: PA clinical experts sort out the guidelines and select key diagnosis and treatment suggestions to form quality control points. Step 2: extract the nouns of diagnosis, treatment and examination that appear in the quality control points, map them to the ontology library for standardization, and then form a standardized quality control logic. Step 3: connect the QCSA with the hospital business system, extract the information from the electronic medical records, carry out entity identification and standardization mapping, and then match the quality control logic in QCSA to judge whether there is a defect. Step 4: if a defect is found, transfer this information to the page end and give the corresponding prompt. QCSA, quality control system for pulmonary aspergillosis.

Characteristics of PA medical records	Total PA medical records	Defective PA medical records
Number	699	162
Sex (M/F)	460 (65.8%)/239 (34.2%)	112 (69.1%)/50 (30.9%)
Median age	56	56
Median hospital stay	8	7
PA (unspecified)	284	0
IPA	132	29
СРА	121	84
ABPA	78	34
CCPA	44	9
SA	36	10
CFPA	2	2
Others*	0	0

Table S5 Characteristics of PA medical records from 2015 to 2020 in the First Affiliated Hospital of Guangzhou Medical University

*, others include AN, SAIA and CNPA. PA, pulmonary aspergillosis; IPA, invasive pulmonary aspergillosis; CPA, chronic pulmonary aspergillosis; ABPA, allergic bronchopulmonary aspergillosis; CCPA, chronic cavitary pulmonary aspergillosis; SA, simple pulmonary aspergillosis; CFPA, chronic fiberoptic pulmonary aspergillosis; AN, Aspergillus nodule; SAIA, subacute invasive aspergillosis; CNPA, chronic necrotizing pulmonary aspergillosis.

	ion results of each	elussification					
Catalogue	True positive	False positive	True negative	False negative	Precision	Recall	F1
Classification of PA							
ABPA	19	0	90	2	1.00	0.90	0.95
SA	5	0	12	0	1.00	1.00	1.00
IPA	13	0	434	1	1.00	0.92	0.96
CCPA	5	0	35	17	1.00	0.23	0.37
CFPA	1	0	6	1	1.00	0.50	0.67
CPA	33	3	198	2	0.92	0.94	0.93
Overall calculation	76	3	775	23	0.96	0.77	0.85
QC Type							
Diagnosis	4	2	55	2	0.67	0.67	0.67
Treatment	38	0	659	18	1.00	0.68	0.81
Examination	34	1	61	3	0.97	0.92	0.94

Table S6 QCSA evaluation results of each classification

QCSA, quality control system for pulmonary aspergillosis; PA, pulmonary aspergillosis; ABPA, allergic bronchopulmonary aspergillosis; SA, simple pulmonary aspergillosis; IPA, invasive pulmonary aspergillosis; CCPA, chronic cavitary pulmonary aspergillosis; CFPA, chronic fiberoptic pulmonary aspergillosis; CPA, chronic pulmonary aspergillosis; QC, quality control.

Table S7 The overall evaluation results of QCSA

Test indicators	Values
Specificity	0.77
Sensitivity	0.99
Positive predictive value	0.96
Negative predictive value	0.97
Precision	0.96
Recall	0.77
F1	0.85

According to the results of confusion matrix in *Table 1*, we calculated the results of evaluation indicators. QCSA, quality control system for pulmonary aspergillosis.