1 Supplementary Appendix

- 2 Title: Cohort profile: Community Burden of Acute Respiratory Infections in
- 3 Shanghai, a longitudinal cohort study in respiratory pathogens, China, 2024-2027
- 4 **Running head:** Community burden of ARIs

5 Tables & Forms

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14 Supplementary Table 1. Baseline Characteristics at Enrollment

15 Survey (CRF01)

Date of signing informed consent:
Participant ID: DDDDDD Family ID: DDDDD
1. Basic Characteristics
1.1 Name of participants:
1.2 Sex: 1=male; 2=female
1.3 Your identification number:
1.4 Your date of birth: DDDD/DD/DD(YYYY/mm/dd)
1.5 Your home address (to street) :
1.6 Place where your residence is registered? 1=Shanghai; 2=Other provinces
1.7 Your occupation:
1=school student; 2=housewife and unemployment; 3=retired;
4= service workers/food delivery; 5=security guards; 6=house keeping;
7=medical staff; 8=office clerk; 9=others
1.8 Your education attainment:
1=primary school level and under; 2=Junior high school; 3= high school/technical secondary school; 4= university/college level or above
1.9 Are you covered by the following insurance (multiple choice allowed)?
1= basic medical insurance for urban workers; 2=basic medical insurance for urban
residents; 3= new rural cooperative medical care; 4= commercial medical insurance; 5= uninsured;
6= others
2. Overall health status
2.1 Your height: cm
2.2 Your weight: kg
2.3 Are you pregnant? 1=Yes; 0=No; 9=Unknown
If yes, your gestational age isweeks.
2.4 Are your children a premature baby (for children 6 years and under)? 1=Yes; 0=No;
9=Unknown
(A premature baby is defined as a child born at less than 37 weeks of gestational age.)
(11 promutare buby is defined as a child born at ress than 57 weeks of gestational age.)
2.5 Are your children born a low-birth-weight baby (for children 6 years and under) ? 1=Yes;
0=No; 9=Unknown
(A low-birth-weight baby is defined as a baby weighing less than 2500 grams within 1 hour of
birth.)
2.6 Do you have or have had any of the following underlying conditions (multiple choice allowed):
1= diabetes mellitus; 2= hypertension; 3= heart disease; 4=asthma; 5= chronic
bronchitis/bronchitis; 6=COPD; 7= chronic kidney disease; 8= myocardial infarction; 9= cerebral
stroke; 10=cancer; 11=immunocompromised (defined as having received a solid organ or
hematopoietic stem cell transplant, undergoing cancer chemotherapy, having a history of HIV or
AIDS, or using steroids for >30 days); 12=others; 13=no medical underlying conditions
2.7 Do you smoke?
1=current smoking; 2= used to smoke, but not smoke now; 3=never smoke; 4= exposure to
second-hand smoke
2.8 Do you drink alcohol?
1=never; 2=occasionally; 3=drink often (once per week); 4=drink everyday
2.9 During the past three months, have you experienced a common cold or any of the following
symptoms, e.g., fever, cough, runny nose, sore throat, stuffy nose, and body aches? 1=Yes; 0=No
If yes, the nearest date of symptom onset: $\Box\Box\Box\Box\Box\Box\Box\Box$ (YYYY/mm/dd);
If yes, how many episodes have you experienced? times.

2.10 During the past one year, have you ever visited a doctor? 1=Yes; 0=No
If yes, how many visits? visits.
2.11 During the past one year, have you ever been hospitalized? 1=Yes; 0=No
If yes, how many days have you been hospitalized? days.
3. Household information
3.1 Total number of members living in your family:(persons)
3.2 The number of children aged <5 years in your family:(persons)
3.3 The number of people aged≥65 years in your family:(persons)
3.4 Total living area of your family:m ²
3.5 Per capita living area of your family:m ²
3.6 What is the average monthly income of your family?
1=less than 5000 Chinese yuan; 2=5000-9999 Chinese yuan; 3=10000-19999 Chinese yuan;
4=≥20000 Chinese yuan; 9= Unknown
3.1 Total number of members living in your family:(persons)
3.2 The number of children aged <5 years in your family:(persons)
3.3 The number of people aged≥65 years in your family:(persons)
4. Vaccination history (self-reported)
4.1 Have you ever received the flu vaccine since October 2023?
1=Yes; 0=No; 9=Unknown
4.2 Have you ever received a Covid-19 vaccine? 1=Yes; 0=No; 9=Unknown
If yes, how many doses have been administered cumulatively?
1=1 dose; 2=2 doses; 3=3 doses; 4=4 doses and more
4.3 Have you received the 23 valent pneumococcal polysaccharide vaccine?
1=Yes; 0=No; 9=Unknown
4.4 Have you received the 13 valent pneumococcal conjugate vaccine?
1=Yes; 0=No; 9=Unknown
4.5 Have you received the Haemophilus influenzae type b (Hib) conjugate vaccine?
1=Yes; 0=No; 9=Unknown
Time of survey completion:/ (YYY/mm/dd:HH/MM)
Name of investigator:

17 Supplementary Table 2. Symptoms of ARIs Monitoring Form

18 (CRF02)

Participant ID: Family I	D:				
1. Occurrence of ARIs					
1.1 Since our last contact, have you experie congestion or discharge, sore throat, body or If "no", survey ends. If "yes", please fill in the 2 APL Symptome/Sing (multiple choices)	muscle aches and pain he following information	, etc.)? 1=Yes; 0=No			
2. ARI Symptoms/Signs (multiple choices allowed)					
Symptoms/Signs	Symptoms/Signs	If "yes", onset time (days ago)			
Fever	1=Yes 0=No	days ago			
Chills	1=Yes 0=No	days ago			
Headache	1=Yes 0=No	days ago			
Body or muscle aches	1=Yes 0=No	days ago			
Sore throat	1=Yes 0=No	days ago			
Fatigue	1=Yes 0=No	days ago			
Nasal congestion or discharge	1=Yes 0=No	days ago			
Wheezing, or dyspnea	1=Yes 0=No	days ago			
Cough	1=Yes 0=No	days ago			
Sputum production	1=Yes 0=No	days ago			
Chest Pain	1=Yes 0=No	days ago			
Other	Please specify	days ago			
For children under 2 years old only	·				
Chest wall indrawing	1=Yes 0=No	days ago			
Head nodding	1=Yes 0=No	days ago			
Central cyanosis	1=Yes 0=No	days ago			
Apnea or difficulty in breathing	1=Yes 0=No	days ago			
Crying can't be eased by parents	1=Yes 0=No	days ago			
Unable to feed or choked while breastfeeding	1=Yes 0=No	days ago			
Lethargy or difficulty to wake up	1=Yes 0=No	days ago			
2.1 Does the subject meet the ARIs' case de date DDD/DD/DD (Format: YYYY/mm/dd).		If yes, please provide the onset			
3. Sampling Information					
3.1 Is a swab sampling scheduled? 1=Yes; 3.2 Name of the Community Healthcare Cen 3.3 Scheduled swab sampling time: Time of survey completion://	ter for scheduled swab	sampling: mm/dd)			
Name of investigator:					

20 Supplementary Table 3. Weekly Illness Updates and Clinical

21 Recovery Follow-up (D07/D14/D21/D28) Data Form (CRF03)

Participant ID:	Family ID: DDDDD				
1. ARI Symptoms/Signs (multiple	choices allowed)				
1.1 Since our last contact, have yo	ou experienced any	of the following syn	nptoms?		
1=Yes, please specify the sympton	ms (multiple choices	allowed); 0=No			
ARI Symptoms/Signs	Day 7	Day 14	Day 21	Day 28	
Fever	1=Yes 0=No	1=Yes 0=No	1=Yes 0=No	1=Yes 0=No	
Chills	1=Yes 0=No	1=Yes 0=No	1=Yes 0=No	1=Yes 0=No	
Headache	1=Yes 0=No	1=Yes 0=No	1=Yes 0=No	1=Yes 0=No	
Body or muscle aches	1=Yes 0=No	1=Yes 0=No	1=Yes 0=No	1=Yes 0=No	
Sore throat	1=Yes 0=No	1=Yes 0=No	1=Yes 0=No	1=Yes 0=No	
Fatigue	1=Yes 0=No	1=Yes 0=No	1=Yes 0=No	1=Yes 0=No	
Nasal congestion or discharge	1=Yes 0=No	1=Yes 0=No	1=Yes 0=No	1=Yes 0=No	
Wheezing, or dyspnea	1=Yes 0=No	1=Yes 0=No	1=Yes 0=No	1=Yes 0=No	
Cough	1=Yes 0=No	1=Yes 0=No	1=Yes 0=No	1=Yes 0=No	
Sputum production	1=Yes 0=No	1=Yes 0=No	1=Yes 0=No	1=Yes 0=No	
Chest Pain	1=Yes 0=No	1=Yes 0=No	1=Yes 0=No	1=Yes 0=No	
Other	Please specify	Please specify	Please specify	Please specify	
For children aged under 2 years o	old only				
Chest wall indrawing	1=Yes 0=No	1=Yes 0=No	1=Yes 0=No	1=Yes 0=No	
Head nodding	1=Yes 0=No	1=Yes 0=No	1=Yes 0=No	1=Yes 0=No	
Central cyanosis	1=Yes 0=No	1=Yes 0=No	1=Yes 0=No	1=Yes 0=No	
Apnea or difficulty in breathing	1=Yes 0=No	1=Yes 0=No	1=Yes 0=No	1=Yes 0=No	
Crying can't be eased by parents	1=Yes 0=No	1=Yes 0=No	1=Yes 0=No	1=Yes 0=No	
Unable to feed or choked while	1=Yes 0=No		1=Yes 0=No	1=Yes 0=No	
breastfeeding		es 0=No 1=Yes 0=No 1=Yes		1-163 0-110	
Lethargy or difficulty to wake up	1=Yes 0=No	1=Yes 0=No	1=Yes 0=No	1=Yes 0=No	
2. Healthcare Utilization					
2.1 Since our last contact, have yo					
Healthcare Utilization	Day 7	Day 14	Day 21	Day 28	
Outpatient/clinic visit	1=Yes 0=No	1=Yes 0=No	1=Yes 0=No	1=Yes 0=No	
Emergency department visit	1=Yes 0=No	1=Yes 0=No	1=Yes 0=No	1=Yes 0=No	
Hospital admission	1=Yes 0=No	1=Yes 0=No	1=Yes 0=No	1=Yes 0=No	
Absent from school or work	1=Yes 0=No	1=Yes 0=No	1=Yes 0=No	1=Yes 0=No	
2.2 Outpatient and Emergency D					
2.2.1 Total number of visits to out		ency Department: _	times		
2.2.2 Name of the hospital or clin					
2.2.3 Date of the first visit:	•	YY/mm/dd)			
2.2.4 Diagnosis from the first visit		2 202 402	2 500 000	4 4000	
2.2.5 Total cumulative expenditu	re: 1=Below 200 yu	ian; 2=200-499 yua	in; 3=500-999 yuan	; 4=1000 yuan ar	
above					
2.3 Hospitalization					
2.3.1 Name of the Hospital:					
2.3.2 Admission Date:/	/🗆 (Format: YYYY/	mm/dd)			
2.4 Absence from Work/School					
2.4.1 Total number of days absen	t from work or scho	ol due to the illness	:		
3. Clinical Outcome					
3.1 By the end of the follow-up pe	eriod, the clinical ou	tcome for the subje	ect is:		

1=Clinical recovery; 2=Improvement or remission; 3=Worsening or Hospitalization; 4=Death.

(Note: Clinical recovery is defined as a normal body temperature for two consecutive days and the complete disappearance of symptoms such as body or muscle aches and pain, fatigue, cough, nasal congestion or discharge, sore throat, and wheezing, or dyspnea. Improvement/Remission is defined as an improvement in systemic and/or respiratory symptoms by the 28-day follow-up, but without complete resolution. Worsening or Hospitalization is defined as being admitted to the hospital during the follow-up period) Time of survey completion: DOM (YYY/mm/dd:HH/MM)

22

Name of investigator:

23 Supplementary Table 4. Hospital Discharge Data Collection

24 Form (CRF04)

Participant N	No.				Family	No.			
1. Basic in	forn	nation	of admi	ission					
1.1 Hospital	name	:							
1.2 Date of a	1.2 Date of admission:								
1.3 Admittin	g dia	gnosis:							
1.4 Date of d	lischa	ırge:□□	100/00/01	□ (Form	at: YYY	Y/mm	/dd)		
1.5 Discharg	e dia	gnosis:							
_	-	Princip	al diagnos	is 1					
		Second	lary diagno	osis 1	;	2		_; 3	
2. Clinical 2.1 Signs/syr			ion resu		me				
	-	-						h	
_					beats/min	Hea	irt rate	e: <u>b</u> eats/min	
			<u>mmHg</u> on (oxygei		on) sPO-		%		
Pulse ox	gen s	saturati	on (withou	it oxygen	a) sPO ₂ :	~	10		
Pulmona	ry aus	scultati	on: 1=dry	rales; 2=	wet rales	; 3=no			
				ır; 2=dro	owsiness;	3=ir	ritabili	ity; 4=delirium;	5=convulsion;
		=norma		9	_			9	
			VBC	$_{\times 10^{\circ}/L}$; L	$_{\times 10^{\circ}}$	7L; N_	<u>×10⁹/L; Pl</u>	.t×
10 ⁹ /L; Hb	g/L	;	amination	CDD	ma/L	CLU		mmal/L DUN	тта 1/Г.
		ncal ex	amination	CRP	mg/L	; GLU		mmol/L; BUN	mmol/L;
PCTµg/L			e c	. ,	d		c) NI
2.4 Clinical 2.4.1 If yes, th								ormed. 1=Yes; (J=No
2.4.2 If yes, th	e met	hod of	laboratory	testing:	1=PCR: 2	2=antis		sting; 3=antibody	testing
2.4.3 If yes, th	e resu	ilt of la	boratory te	sting: 1=	positive;	0=neg	gative	, sing, s-untroody	lesting
2.5 Whether	chest	radiog	raphy or ot	her chest	imaging	was p		ned? 1=Yes; 0=No	D
			of pneumo	nia? 1=Y	es; 0=No)			
2.6 Complica			umonia	Pastaria	1	via 🗆	Dnoun	aothoray	
Septic shock Viral pneumonia Bacterial pneumonia Pneumothorax ARDS Bronchiolitis Respiratory failure Coagulopathy									
Anemia	Pleura	l effusio	on Acu	te kidney	injury	Myoly	sis	_	
							ngitis	pancreatitis	
Convulsion Hyperglycem	ia 🗌	Hypog	a Liver lvcemia	Congest	tive heart	failure	Пне	eart infection	
								()	
3. Treatme									
3.1 Was oxyge									
If yes, the method of treatment: 1=nasal cannula or mask oxygen; 2=high-flow nasal cannula;									
3=non-invasive mechanical ventilation; 4=invasive mechanical ventilation; 5= Other 3.2 Admission to the ICU? 1=Yes; 0=No									
If yes, the length of ICU admission (days)									
3.3 Were vasopressors administered? 1=Yes; 0=No									
3.4 Were extracorporeal membrane oxygenation (ECMO) administered? 1=Yes; 0=No									
3.5 Were Continuous renal replacement therapy(CRRT) administered? 1=Yes; 0=No									
4. Drugs a	dmi	nister	ed						
Drug name	Drug name Category Ro		Route	Daily	v dose	Frequ	iencv	Starting date	Stop date
2145 hume	Cut	-5017	muu	Dose	Unit	riequ	y	(YYYY/mm/dd)	(YYYY/mm/dd)

 4.1 Drug Name: (Please use the name of the drug. If it is a fixed compound preparation, please use the trade name.) 4.2 Category: A=antibiotics; B=antiviral drugs; C=steroid hormone drugs; D=angiotensin-converting enzyme inhibitors (ACE-Is) or angiotensin-receptor blockers (ARBs); 					
E=Statins 4.3 Route of medication: 1=oral administration, 2=intravenous injection, 3=intravenous drip,					
4=intramuscular injection, 5=inhalation, 6=others 4.4 Frequency: 1= continuous, 2= intermittent					
5. Patient prognosis					
cured improved and be discharged transferred to the other hospital					
Reasons for transfer: community rehabilitation/other ()					
reasons for giving-up: economic reasons/illness exacerbation/other ()					
death date of death:/(YYYY/mm/dd) death diagnosis:					
6. The total expenditure of your hospitalization:RMB yuan Time of survey completion:(UTYY/mm/dd:HH/MM)					
Name of investigator:					

26 Supplementary Table 5. Semi-annual Survey Data Form

27 (CRF05)

Participant No. DDDDDD Family No. DDDDDD
1. Update of family information
1.1 Total number of members living in your family:(persons)
1.2 The number of children aged <5 years in your family:(persons)
1.3 The number of people aged≥65 years in your family:(persons)
2. Update of vaccination information during the study
2.1 Have you received the flu vaccine during your participation in the study? 1=Yes; 0=No;
9=Unknown
2.2 Have you received the Covid-19 vaccine during your participation in the study? 1=Yes;
0=No; 9=Unknown
If yes, how many doses have been administered cumulatively? 1=1 dose; 2=2 doses; 3=3 doses; 4=4 doses and more
2.3 Have you received the 23 valent pneumococcal polysaccharide vaccine during your participation in the study? 1=Yes; 0=No; 9=Unknown
2.4 Have you received the 13 valent pneumococcal conjugate vaccine during your participation in
the study? 1=Yes; 0=No; 9=Unknown
2.5 Have you received the Haemophilus influenzae type b (Hib) conjugate vaccine during your
participation in the study? 1=Yes; 0=No; 9=Unknown
Time of survey completion: DDD/DD/DD:DD/DD (YYYY/mm/dd:HH/MM)
Name of investigator:

Supplementary Table 6. Lists of respiratory pathogens tested for 29

in the study 30

no.	Viruses	no.	Bacteria
1	Influenza A	27	Bordetella holme
2	Influenza B	28	Bordetella pertus
3	Respiratory syncytial virus subtype A/B	29	Chlamydophila p
4	SARS-Cov-2	30	Haemophilus infl
5	Human Coronavirus-229E	31	Klebsiella pneum
6	Human Coronavirus- HKU1	32	Legionella pneun
7	Human Coronavirus- NL63	33	Moraxella catarr
8	Human Coronavirus- OC43	34	Mycoplasma pne
9	MERS-CoV	35	Staphylococcus a
10	SARS-CoV	36	Streptococcus pn
11	Adenovirus		Fungus
12	Human parainfluenza virus serotype 1	37	Pneumocystis j
13	Human parainfluenza virus serotype 2		
14	Human parainfluenza virus serotype 3		
15	Human parainfluenza virus serotype 4		
16	Human metapneumovirus		
17	Rhinovirus		
18	Enterovirus		
19	Bocavirus		
20	varicella-zoster virus		
21	Epstein-Barr virus		
22	Cytomegalovirus		
23	Human herpesvirus 6		
24	Measles virus		
25	Mumps virus		
26	Parechovirus		

no.	Bacteria
27	Bordetella holmesii
28	Bordetella pertussis
29	Chlamydophila pneumoniae
30	Haemophilus influenzae
31	Klebsiella pneumoniae
32	Legionella pneumophila
33	Moraxella catarrhalis
34	Mycoplasma pneumoniae
35	Staphylococcus aureus
36	Streptococcus pneumoniae
	Fungus
37	Pneumocystis jirovecii