

ORIGINAL ARTICLE

Community-acquired pneumonia in Chile: the clinical relevance in the detection of viruses and atypical bacteria

Vivian Luchsinger,¹ Mauricio Ruiz,² Enna Zunino,³ María Angélica Martínez,⁴ Clarisse Machado,⁵ Pedro A Piedra,⁶ Rodrigo Fasce,⁷ María Teresa Ulloa,⁴ Maria Cristina Fink,⁵ Pamela Lara,¹ Mónica Gebauer,³ Fernando Chávez,³ Luis F Avendaño¹

¹Programa de Virología, Facultad de Medicina, Universidad de Chile, Santiago, Chile

²Facultad de Medicina, Hospital Clínico Universidad de Chile, Santiago, Chile

³Facultad de Medicina, Hospital de Infecciosos Dr. Lucio Córdova, Santiago, Chile

⁴Programa de Microbiología, Facultad de Medicina, Universidad de Chile, Santiago, Chile

⁵Laboratorio de Virología, Instituto de Medicina Tropical de São Paulo, Universidade de São Paulo, São Paulo, Brazil

⁶Department of Molecular Virology and Microbiology, and Pediatrics, Baylor College of Medicine, Houston, Texas, USA

⁷Sección Virus Respiratorios y Exantemáticos, Subdepartamento Enfermedades Virales, Instituto de Salud Pública de Chile, Santiago, Chile

Correspondence to

Luis F Avendaño, Programa de Virología, Facultad de Medicina, Universidad de Chile, Av. Independencia 1027, Independencia, Santiago 8380453, Chile; lavendan@med.uchile.cl

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ABSTRACT

Background Adult community-acquired pneumonia (CAP) is a relevant worldwide cause of morbidity and mortality, however the aetiology often remains uncertain and the therapy is empirical. We applied conventional and molecular diagnostics to identify viruses and atypical bacteria associated with CAP in Chile.

Methods We used sputum and blood cultures, IgG/IgM serology and molecular diagnostic techniques (PCR, reverse transcriptase PCR) for detection of classical and atypical bacteria (*Mycoplasma pneumoniae*, *Chlamydia pneumoniae*, *Legionella pneumoniae*) and respiratory viruses (adenovirus, respiratory syncytial virus (RSV), human metapneumovirus, influenza virus, parainfluenzavirus, rhinovirus, coronavirus) in adults >18 years old presenting with CAP in Santiago from February 2005 to September 2007. Severity was qualified at admission by Fine's pneumonia severity index.

Results Overall detection in 356 enrolled adults were 92 (26%) cases of a single bacterial pathogen, 80 (22%) cases of a single viral pathogen, 60 (17%) cases with mixed bacterial and viral infection and 124 (35%) cases with no identified pathogen. *Streptococcus pneumoniae* and RSV were the most common bacterial and viral pathogens identified. Infectious agent detection by PCR provided greater sensitivity than conventional techniques. To our surprise, no relationship was observed between clinical severity and sole or coinfections.

Conclusions The use of molecular diagnostics expanded the detection of viruses and atypical bacteria in adults with CAP, as unique or coinfections. Clinical severity and outcome were independent of the aetiological agents detected.

INTRODUCTION

Community-acquired pneumonia (CAP) is a worldwide cause of morbidity and mortality.^{1–3} In spite of the advances in diagnostics, the aetiology of CAP often remains uncertain and therapy is empirical.^{4–5} The low sensitivity of blood and sputum cultures and the presence of a heterogeneous bacterial flora in the upper respiratory tract confound the interpretation of the tests.⁶ Respiratory viruses are the worldwide cause of acute lower respiratory tract infections (ALRIs) in children and their

Key messages

What is the key question?

- ▶ Viruses and atypical bacteria were frequently detected as single and mixed agents in adults with community-acquired pneumonia (CAP).

What is the bottom line?

- ▶ Although agent detection increased notably by PCR diagnosis, clinical severity and outcome were independent of the pathogenic agents detected.

Why read on?

- ▶ Although adult CAP is a relevant worldwide cause of morbidity and mortality, the aetiology often remains uncertain and the therapy is empirical.

specific diagnosis is often made by immunodiagnostic techniques. In children, unlike in adults, an aetiological diagnosis is easily assessed by many diagnostic techniques because viral shedding is high and prolonged.^{7–8} The development of molecular methods with improved sensitivity and specificity has paved the way for the detection of new viruses, for the identification of pathogens that are difficult to culture, and for the detection of pathogens later in the disease process. Molecular diagnostics have improved our understanding of ALRIs in children and in adults.^{9–12} Seasonality of ALRIs is similar in paediatric and adult respiratory illnesses, suggesting a similar aetiology. In prior reports we presented the epidemiology and clinical impact of *Mycoplasma pneumoniae* and respiratory syncytial virus (RSV) in Chilean adults with CAP.^{13–15} Here we present a comprehensive evaluation of the bacterial and viral aetiology of CAP in adults. We postulate that viruses and atypical bacteria either as sole or coinfections play a relevant role in adult CAP and that molecular diagnostics will define the specific aetiology in a majority of cases. We also explore the relationship between respiratory pathogen and clinical severity with a validated severity score index.¹⁶

METHODS

Patients and study design

A prospective study was conducted in patients ≥ 18 years of age presenting with CAP in two hospitals in Santiago, Chile from February 2005 to December 2007. The study was approved by the University of Chile and Health Institutional Ethics Committee and all subjects at enrolment gave written informed consent. CAP was defined by the presence of acute respiratory symptoms for less than 1 week and chest radiography showing new pulmonary infiltrates. Exclusion criteria included immunocompromising conditions (ie, HIV, active treatment for cancer, organ transplant, immunosuppressive therapy) and hospitalisations within 30 days preceding enrolment. Information on age, gender, smoking status, prior antibiotic treatment, comorbidities and clinical presentation, vital signs, laboratory parameters, antimicrobial therapy and hospital course were recorded for all patients in standardised files. Chest radiographic patterns were described by independent radiologists as alveolar, interstitial or mixed infiltrates; the extension was categorised according to the number of lobes involved and the presence of pleural effusion. Patient severity was assessed during the first 48 h after enrolment by the pneumonia severity index described by Fine.¹⁶ The illness outcome and the presence of complications like renal failure, shock, mechanical ventilation and others were recorded. Patients who died between admission and 30 days after discharge were recorded.

Sample collection

In addition to routine laboratory tests, such as complete blood cell count, biochemistry panel, oxygen saturation at admission, patients had collection of urine and blood for bacterial culture and acute serum sample for serology. Induced sputum and nasopharyngeal aspirate were obtained and immediately transported on ice to the laboratory for bacterial and viral diagnostics. Aliquots were prepared and stored at -80°C for later testing. Participants were contacted at 4–6 weeks for follow-up and collection of convalescent sera. Serum samples were processed immediately and stored at -20°C until tested.

Viral study

Specimen processing

Sample aliquots of respiratory secretions were prepared for immunofluorescent assay, viral isolation in cell culture and real time PCR.

Viral isolation

Each specimen was processed as previously described.¹⁰ Briefly, samples were inoculated onto HEp-2 and Madin–Darby canine kidney (MDCK) cell cultures and observed for development of cytopathic effect (CPE) for 1 week, after which confirmatory immunofluorescence assays (IFAs) for RSV, adenovirus, influenza and parainfluenza viruses were performed in both cell cultures with and without CPE.

Indirect immunofluorescence assay

Smears were prepared in triplicate and IFA was performed for RSV, adenovirus, influenza A and B, parainfluenza virus 1–3 as described elsewhere, using monoclonal antibodies kindly provided by L Anderson (CDC, Atlanta) and P Pothier (Dijon, France). We used commercial conjugate (Sigma) and virus-specific monoclonal antibodies (Chemicon) for influenza and parainfluenza virus typing.^{7, 8}

Serology

Serum haemagglutination inhibition (HI) test for antibodies to influenza (H3N2, H1N1 and B) was performed as described elsewhere.¹⁷ Paired sera were inactivated with receptor-destroying enzyme (Denka Seiken Corporation, Tokyo, Japan), incubated overnight at 37°C , and heated at 56°C for 30 min. A 0.5% turkey red blood cell suspension, control antigens and reference antisera were used. Cases displaying at least a fourfold rise between the acute and convalescent sera were recorded as acute infection.

Sera were tested for antibodies to RSV and human metapneumovirus (hMPV) by microneutralisation assays to RSV/A/Tracy (A2 like virus) and RSV/B/18537 (prototype B virus) to measure RSV/A and RSV/B specific neutralising antibodies as previously described.¹⁸ hMPV antibodies were also detected by ELISA assays.¹⁹ Seroconversion was defined by at least a fourfold rise of antibody titre between acute and convalescent sera by one or more of the serological tests.

Reverse transcriptase PCR for RSV and hMPV

Samples were treated with guanidinium thiocyanate-phenol-chloroform method for RNA extraction.²⁰ cDNA was synthesised with $5\ \mu\text{L}$ RNA (sample) and $0.52\ \mu\text{M}$ F gene primer (F844: 5'-TGTCTAACTATTGAACA-3') for RSV and $10\ \mu\text{L}$ RNA and $0.52\ \mu\text{M}$ random hexamer primer (Amersham Bioscience) for hMPV, during 1 h at 37°C , followed by 5 min at 95°C in a AmpPCR System 2400. For RSV, a sequence of the N gene was amplified with $10\ \mu\text{M}$ (each) of N1 and N2 primers,²¹ $1\times$ Master SYBR Green I (Roche) and $2\ \mu\text{L}$ of cDNA. For hMPV, a sequence of the N gene were amplified with $0.5\ \mu\text{M}$ (each) of N2 and N3 primers.²² Cycling conditions included a denaturation step of 10 min at 94°C , followed by 50 (RSV) or 40 cycles (hMPV) of 10 s at 94°C , 5 s at 58°C (RSV) or 52°C (hMPV), 30 s at 72°C in a Light Cycler 1.5 instrument (Roche). Amplicons were analysed by melting curve analysis. Appropriated controls were included in each assay.

Reverse transcriptase PCR for rhinovirus and coronavirus detection

Human coronavirus (HCoV), and picornavirus (PV) were detected by real time reverse transcriptase PCR (RT-PCR) by amplifying the conserved region of the replicase 1a gene and the 5'NCR conserved region, respectively, as previously described.^{23, 24} The 390 bp fragment of PV was purified and sequenced using the kit ABI PRISM Dye Terminator Cycle Sequencing ready Reaction in the automated sequencer ABI model 377 (Applied Biosystems, Inc, USA). Sequences were aligned to other sequences in the Gene Bank using the programme Blast to differentiate between rhinovirus and enterovirus.

Bacterial study

Cultures

Sputum samples displaying >25 leukocytes and <10 epithelial cells per $100\times$ power field after Gram staining were cultured and processed according to standard techniques.^{6, 25} Urinary antigens for *Streptococcus pneumoniae* were detected using immunochromatographic tests (Binax NOW, Portland, Oregon, USA).²⁶ For *Legionella* species isolation, a subset of 256 samples were inoculated onto BCYE and GVPC media (Oxoid) and incubated at 37.8°C for up to 10 days; 125 of these cases were also studied for urinary antigen using immunochromatographic tests.²⁷ Confirmatory PCR was performed on positive samples as described elsewhere.²⁸

Polymerase chain reaction

Chlamydomphila pneumoniae was detected using a nested PCR directed to a chromosomal DNA segment (*PstI* 474) using the reference strains AR-39, TW-183 and CM-1 as positive controls. Primers HL and HR were used for the first PCR and nested PCR was performed as described elsewhere.^{29 30} For *M pneumoniae* a PCR directed to a 277 bp fragment of the 16S rRNA gene was used, including a *M pneumoniae* FH strain as positive control.^{14 31}

Serology

Serum samples were tested for IgM and IgG antibodies by indirect IFA using commercial kits. Sera were absorbed with an anti-human IgG reagent (Zorba, Zeus, Inc, USA) prior to IgM testing. The *Chlamydia* kit (SeroFIA, Sayvon, Israel) was used for *C pneumoniae* and IgM $\geq 1:16$ or a fourfold rise in IgG titres between paired sera were regarded as acute infection; single or standing IgG titres $\geq 1:512$ were considered as past infections. For *M pneumoniae* another IFA test was used (Zeus, USA) and acute infection was assigned with IgM $\geq 1:32$ or seroconversion in paired sera.³² Sera determinations were done blindly with respect to other viral or bacterial tests.

Statistical analysis

Analysis was performed using the Z-test for categorical data and the t test, Mann–Whitney rank sum or Kruskal–Wallis one-way analysis of variance for continuous variables. The level of significance was set at $p < 0.05$. Data were analysed using SigmaStat software.

RESULTS

Patients and samples

Respiratory secretions and urine were obtained from 356 patients from 330 hospitalised and 26 ambulatory cases. Good-quality sputum samples were available for 233 (65%) cases. Blood culture was performed in 241 (75%) patients and paired sera were available for 211 (68%) patients. Rhinovirus and coronavirus were studied in a subset of 268 cases from August 2005 to September 2007. Serology for influenza was performed only in cases with paired sera, which covered the three annual seasonal influenza epidemics. A respiratory pathogen was detected in 232 (65.2%) of 356 cases, corresponding to bacteria in 92 (26%) and viruses in 80 (22%) cases. Mixed viral/bacterial infections were detected in 60 (17%) cases. In 124 (35%) patients no agent was detected (table 1). Death occurred in 28 (7.8%) cases, without significant difference between those with or without a detected infectious agent.

Table 1 Distribution of 356 adults with community-acquired pneumonia by infection classification

Infection classification	N	%
Negative cases	124	34.8
Only bacteria*	92	25.8
Only viruses*	80	22.5
Mixed†	60	16.9

Santiago, Chile, February 2005–December 2007.

*One or more agents per case.

†Mixed virus and bacteria, with one or more agents per case.

Clinical characteristics of population

Demographic risk factors, comorbidities and outcomes were not significantly different between those with or without a detectable pathogen (table 2). Also laboratory parameters like haemogram, glucose, electrolytes, urea and liver function enzymes did not exhibit significant differences (data not showed). Only C-reactive protein was significantly higher in patients with a detectable pathogen (median=211 mg/dL) compared with patients without an identifiable pathogen (median=158 mg/dL) ($p=0.01$), although with a broad range in values (table 2).

The clinical outcome related to the presence of complications, like radiologic progression, respiratory failure, need for mechanical ventilation or presence of shock, failed to show any significant difference between both groups. Only liver damage (14/221=6.3% vs 1/119=0.8%; $p=0.03$) was significantly

Table 2 General characteristics of 356 patients with community-acquired pneumonia, according to detection of an infectious agent

Characteristic	With agent	Without agent	P Value*
N	232	124	
Age (years), median (range)	61 (18–94)	66 (18–93)	0.1
Men, n (%)	128 (55.2)	63 (50.8)	0.4
Death, n (%)	17 (7.3)	11 (8.9)	0.6
Hospital stay (days), median/n (range)	7/190 (1–71)	7/104 (1–82)	0.3
Hospitalisation, n (%)	211 (90.9)	119 (96.0)	0.1
ICU admission, n (%)	53 (22.8)	30 (24.2)	0.7
Outpatients, n (%)	21 (9.1)	5 (4.0)	0.1
Smokers, n (%)	90/220 (40.1)	37/119 (31.1)	0.1
Alcohol use >80 g/day, n/N (%)	31/221 (14)	11/119 (9.2)	0.2
Comorbidity, n (%)	N=221	N=119	
Any	112 (50.7)	66 (55.5)	0.4
Diabetes mellitus	36 (16.3)	19 (15.9)	0.9
COPD	36 (16.3)	23 (19.3)	0.5
Asthma	12 (5.4)	4 (3.4)	0.5
Cardiac failure	38 (17.2)	23 (19.3)	0.7
Liver damage	14 (6.3)	1 (0.8)	0.03
Renal disease	4 (1.8)	7 (5.9)	0.08
Antibiotics prior to admission, n/N (%)	48/220 (21.8)	24/119 (20.2)	0.8
Mental confusion, n/N (%)	55/219 (25.1)	22/115 (19.1)	0.2
Hypotension, n/N (%)	36/218 (16.5)	26/119 (21.9)	0.2
Shock, n/N (%)	21/217 (9.7)	9/119 (7.6)	0.6
Respiratory rate, median/n (range)	24/172 (18–48)	24/99 (15–46)	0.8
Pulse, median/n (range)	95/214 (17–153)	91/119 (10–180)	0.1
Systolic blood pressure (mm Hg), median/n (range)	120/178 (68–213)	123/178 (70–195)	0.6
Temperature at admission (°C), median/n (range)	37.2/219 (35–41)	37.4/117 (35–40.5)	0.9
C-reactive protein (mg/dL), median/n (range)	211/182 (2–569)	158/101 (3–454)	0.02
Chest radiograph, n (%)	N=217	N=115	
Interstitial patterns	31 (14.3)	12 (10.4)	0.4
Alveolar patterns	173 (79.7)	95 (83.3)	0.5
Both	13 (5.9)	8 (6.9)	0.9
Multilobar involvement	70 (32.6)	42/113 (37.2)	0.4

Santiago, Chile, February 2005–December 2007.

*p by Kruskal–Wallis test.

COPD, chronic obstructive pulmonary disease; ICU, intensive care unit.

higher in patients with a detectable pathogen (table 2). However, clinical severity of cases correlated with parameters classically described, like age, comorbidity, hypotension, hepatic failure and others (table 3).

Viral and bacterial agent detection

The most commonly identified bacteria was *S pneumoniae* (75=21.1%), although it was the sole pathogen in 38 cases (table 4). Of these 75 patients, 39 were tested by sputum culture, 61 by blood culture and 73 by urine antigen detection, and *S pneumoniae* was identified in 12 (30.8%), 15 (24.6%) and 60 (82.2%) cases, respectively.

M pneumoniae and *C pneumoniae* were detected in 32 (9%) and 28 (7.9%) cases, respectively, but only in 13 cases for each agent were they the sole pathogen. For *M pneumoniae*, PCR and serology yielded similar results (69.7% and 72.7%, respectively; 42.4% of cases were positive by both tests). *C pneumoniae* was detected by PCR (19/28) and serology (15/28), with 6 of 28

cases (21.4%) detected by both tests. *Legionella pneumophila* was detected in 13/256 cases (5.07%), 3 by culture, 10 by antigen detection (one by both); all cases were confirmed by PCR. The most commonly detected viruses were RSV (n=48), PV (n=41) and hMPV (n=41), but single infections occurred in only 18, 17 and 8 cases, respectively (table 4).

Yield of different techniques for viral diagnosis

Detection of a viral pathogen by conventional IFA and cell culture techniques was infrequent, with only 2 of 48 RSV infections and 6 of 27 influenza cases. The RT-PCR technique identified 32 RSV cases. The microneutralisation serology for RSV and HI for influenza detected seroconversion in 20 of 184 (10.8%) and 26 of 211 (12.8%) adults with paired sera, respectively. RT-PCR was the only test applied for hMPV, PV and coronavirus diagnosis, resulting in the detection of 41 (11.5%), 41 (11.5%) and 20 (4.5%) cases, respectively (table 4).

Table 3 Clinical characteristics of 330 patients with community-acquired pneumonia admitted to ICUs or general wards

Clinical parameters	ICU	General Wards	p Value*
N	83	247	
Age (years), median (range)	68 (20–93)	63 (18–94)	0.06
Men, n (%)	39 (47.0)	138 (55.9)	0.2
Hospital stay (days), median/n (range)	116/80 (24–196)	6/195 (1–61)	<0.001
Comorbidity, n (%)	N=81	N=233	
Any	56 (69.1)	118 (50.6)	0.006
Diabetes mellitus	19 (23.5)	66 (28.3)	0.4
Chronic obstructive pulmonary dis.	14 (17.3)	76 (32.6)	0.01
Asthma	1 (1.2)	26 (11.2)	0.01
Cardiac failure	22 (27.2)	55 (23.6)	0.6
Liver damage	7 (8.7)	13 (5.6)	0.7
Renal disease	5 (6.2)	8 (3.4)	0.7
Antibiotics prior to admission, n/N (%)	17/82 (20.7)	47/233 (20.2)	0.9
Infectious agent detected, n (%)			
Virus	17 (20.5)	58 (23.5)	0.6
Bacteria	19 (22.9)	63 (25.5)	0.7
Virus–bacteria coinfection	17 (20.5)	37 (14.9)	0.3
Negative	30 (36.1)	89 (36.0)	0.9
Mental confusion, n/N (%)	45/82 (54.9)	32/230 (13.9)	<0.001
Pleural effusion, n/N (%)	25/81 (30.9)	37/228 (16.2)	0.008
Hypotension, n/N (%)	40/80 (50.0)	19/231 (8.2)	<0.001
Respiratory rate, median/n (range)	28/79 (18–36)	24/168 (16–40)	<0.001
Pulse, median/n (range)	105/81 (73–142)	93/226 (41–180)	<0.001
Systolic pressure (mm Hg), median/n (range)	110/79 (68–213)	122/174 (75–195)	0.006
Diastolic pressure (mm Hg), median/n (range)	60/78 (26–133)	70/174 (43–97)	<0.001
Chest Rx multilobar involvement, n/N (%)	41/81 (50.6)	65/229 (28.9)	<0.001
Fine score, n (%)	N=83	N=247	
1	5 (6.0)	64 (25.9)	<0.001
2	6 (7.2)	59 (23.9)	<0.001
3	15 (18.1)	53 (21.5)	0.6
4	24 (28.9)	62 (25.1)	0.6
5	33 (39.8)	9 (3.6)	<0.001
Blood urea (mg/dL), median/n (range)	21/79 (8–116)	18/219 (6–75)	<0.001
Chest Rx progression, n/N (%)	13/80 (16.3)	7/231 (3.0)	<0.001
Hepatic failure, n/N (%)	8/80 (10.0)	1/231 (0.4)	<0.001
Respiratory failure, n/N (%)	29/80 (36.3)	24/232 (10.3)	<0.001
Shock, n/N (%)	27/80 (33.8)	3/231 (1.3)	<0.001
Death during hospitalisation and until 30 days post discharge, n/N (%)	17/83 (20.5)	9/247 (3.6)	<0.001

Santiago, Chile, 2005–2007.
ICU, intensive care unit.

Table 4 Distribution of single and multiple viral and bacterial agents detected in 356 adults with community-acquired pneumonia

Bacterial agents						
	Only bacteria (n=92)		Mixed bacterial and viral infections (n=60)		Total	
	Only one	Plus other bacteria	Plus virus	Plus virus and other bacteria	N	%
<i>Streptococcus pneumoniae</i>	38	6	25	6	75	21.1
<i>Haemophilus influenzae</i>	3				3	0.8
<i>Moraxella catarrhalis</i>	1	3	2		6	1.7
Gram(-)bacillus	1	2		2	5	1.4
<i>Staphylococcus aureus</i>	3	1	2	2	8	2.2
<i>Streptococcus agalactiae</i>			1		1	0.3
<i>Mycoplasma pneumoniae</i>	13	5	13	1	32	9.0
<i>Chlamydia pneumoniae</i>	13	5	4	6	28	7.9
<i>Legionella pneumophila</i>	7	2	4		13	5.07*
Subtotal cases with bacteria	80				152†	42.7
Viral agents						
	Only viruses (n=80)		Mixed viral and bacterial infections (n=60)		Total	
	Only one	Plus other virus	Plus bacteria	Plus bacteria and other virus	N	%
RSV	18	8	12	10	48	13.5
hMPV	8	14	10	9	41	11.5
Flu	12	5	7	3	27	7.6
HCoV	3	9	3	5	20	5.6
Picornavirus	17	11	9	4	41	11.5
ADV	1		2		3	0.8
Subtotal cases with viruses	59				140‡	39.3
No agent					124	34.8

Santiago, Chile, February 2005–December 2007.

*13/256 cases=5.07%.

†Cases with unique bacteria or with bacteria plus other agent were counted as one individual case.

‡Cases with unique virus or with virus plus other agent were counted as one individual case.

ADV, adenovirus; HCoV, human coronavirus; hMPV, human metapneumovirus; RSV, respiratory syncytial virus.

Seasonality

During the 36-month survey, respiratory viruses were circulating with clear predominance of RSV during the three winter seasons (May–July) (figure 1); influenza viruses presented as an epidemic in autumn/winter and parainfluenza viruses had higher prevalence during the autumn months. This epidemiology is consistent with the patterns previously reported in Chile.¹⁰ CAP peaked during the winter months. It was influenced by the epidemiology observed with *S pneumoniae*, RSV and influenza virus. Other agents were endemic throughout the year (figure 1).

Clinical severity and infection classification

The analysis of demographic and clinical parameters failed to demonstrate a relationship between infection classification (bacteria, virus or mixed coinfection) and illness severity outcome. Furthermore, the presence of multiple pathogens did not contribute to more severe disease (data not shown). Fine's severity score applied at admission also did not display differences among the pathogenic groupings (table 5).

DISCUSSION

The aetiology of CAP in adults is not well understood. We undertook this prospective study over a 3-year period in Santiago to determine if comprehensive use of conventional and molecular diagnostics would improve the aetiological

identification of adult CAP and to determine if the aetiological agent impacted disease severity. An aetiological agent was identified in 65% of cases. *S pneumoniae* and RSV were the most common bacterial and viral pathogens identified. Single and coinfections occurred in 139 (39%) and 93 (26%) CAP cases, respectively. Potentially vaccine-preventable pathogens (*S pneumoniae* and influenza virus) were involved in 102 (28.7%) of the 356 CAP cases. The analysis was done globally, comparing groups with and without a detectable infectious agent and by infection classification (viral, bacterial and mixed viral and bacterial) because sole infection by a few major pathogens was uncommon.

Adult CAP was observed year round, with distinct winter seasonality. Outbreaks of *S pneumoniae*, RSV and influenza contributed to the increase in adult CAP during the winter months. Effective vaccines are available against influenza and *S pneumoniae*. Increased use of these vaccines could reduce the rates of adult CAP, in particular during the winter months. In this study information on vaccination history was not recorded. Historically vaccination against influenza and *S pneumoniae* in adults has not been well accepted by this population, even though there are existing national recommendations for vaccinating older adults and at-risk groups.

Comprehensive use of molecular diagnostics, serological assays, urinary antigen tests, and induced sputum and blood

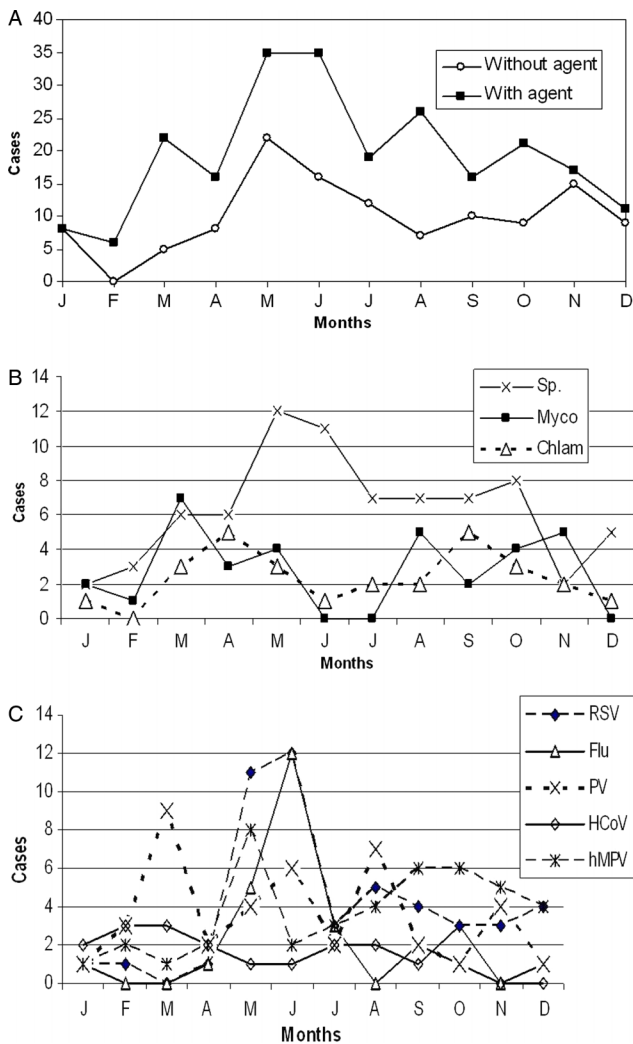


Figure 1 Monthly distribution of 356 adults presenting with community-acquired pneumonia, according to infectious agent identified in Santiago, Chile, February 2005 to December 2007. (A) Cases with and without a detectable infectious agent. (B) Cases associated with a bacterial pathogen (*Streptococcus pneumoniae* (Sp), *Mycoplasma pneumoniae* (Myc), *Chlamydia pneumoniae* (Chlam)). (C) Cases associated with a viral pathogen (respiratory syncytial virus (RSV), influenza (Flu), picornavirus (PV), coronavirus (HCoV), metapneumovirus (hMPV)).

cultures improved the overall detection of an infectious agent in adults presenting with CAP. Viral and the atypical bacteria detection were as frequent as the classical *S pneumoniae*. There were

no antecedent symptoms or signs that could be associated with one specific aetiological agent. In addition, chest radiograph findings could not define the aetiology.

The identification of a bacterial agent using sputum and blood cultures was infrequent, and the detection of a bacterial pathogen was improved with the use of the urinary antigen assay for *S pneumoniae*. The implementation of molecular diagnostic technology has enhanced and broadened the spectrum of potential aetiological agents because of the improved identification of known agents and the recognition of new agents associated with respiratory illnesses.³³⁻³⁴ In virology, molecular techniques are extensively used and for some agents they constitute the principal method for identification (ie, hMPV, PV, coronavirus). Furthermore, the diminished load and shorter duration of viral shedding that occurs in adults compared with children makes the use of highly sensitive diagnostic tools necessary in adults. For atypical bacteria, PCR should be used as the primary diagnostic technique; unfortunately, its application is often restricted to major academic and diagnostic centres.

To our surprise, we were unable to find a relationship between illness severity and major pathogenic grouping or infection classification. Most of the agents detected occurred more frequently with other pathogens than as the sole pathogen. We detected many infectious agents using new diagnostic technology, but the translation of this finding was not clear. The detection of traditional pathogens, like *S pneumoniae* and atypical bacteria, could explain the clinical outcome, but the detection of RSV and hMPV, recognised viral respiratory pathogens in the paediatric population, must be carefully considered in adults. The identification of other viruses such as rhinoviruses and coronaviruses makes it more difficult to assign an aetiological interpretation because asymptomatic infections and prolonged viral shedding can also occur. The frequent detection of two or more infectious agents could be the more likely situation in CAP compared with the classical assignment of a single agent to a pathogenic outcome.³⁴ Clinical or radiological features are not characteristic of *S pneumoniae* CAP. In this study nearly 50% of *S pneumoniae* cases occurred as coinfections. This finding suggests that the classical presentation of pneumococcal CAP as the sole pathogen might not be correct and that other pathogens are likely to contribute to the disease. Today the use of more sensitive diagnostic technology is rendering it difficult to make a simple association between agent and illness.

In summary, factors such as age and comorbidities are relevant to the severity of CAP. Despite improvements in identifying an aetiological agent associated with CAP, we were unable to demonstrate a clinical significance during the acute process. The relevant detection of classic, atypical bacteria and viruses should be considered in the initial empirical antimicrobial intervention.

Table 5 Clinical severity according to Fine score and infection classification in 356 adults with community-acquired pneumonia

Fine class	Total cases		Without agent		With agent		Bacteria		Virus		Bacteria and virus	
	N	%	N	%	N	%	N	%	N	%	N	%
1	84	23.6	24	19.4	60	25.9	38	25.0	33	23.6	11	18.3
2	74	20.8	28	22.6	46	19.8	25	16.4	31	22.0	10	16.7
3	69	19.4	29	23.4	40	17.2	27	17.8	28	21.3	15	25.0
4	87	24.4	33	26.6	54	23.3	39	25.7	31	22.0	16	26.7
5	42	11.8	10	8.1	32	13.8	23	15.1	17	12.1	8	13.3
Total	356	100	124	34.8	232	65.2	152	42.7	140	39.3	60	16.9

Santiago, Chile, February 2005–December 2007. Statistical analyses were performed using Z test, $p > 0.1$.

Atypical bacteria were detected in approximately 20% of cases, justifying antimicrobial coverage. The seasonality of viral respiratory agents should also influence the recommendation for influenza antiviral therapy.

In conclusion:

1. Viruses and atypical bacteria are detected as frequently as *S pneumoniae* in patients with CAP if modern diagnostic technology is applied.
2. We were unable to demonstrate that either type of infection (bacterial vs viral) or number of pathogens (sole vs coinfections) adversely affected clinical outcome.
3. The detection of multiple infectious agents in adult CAP is common and should be considered in the selection of empirical antimicrobial therapy.
4. Vaccination against influenza and *S pneumoniae* can reduce adult CAP admission, especially during the winter season.

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Competing interests None.

Patient consent Obtained.

Ethics approval All cases included completed a duly signed consent form. The study was approved by the Universidad de Chile and Health Services Research and Ethics Committee. Also two National Research Institutions approved the study: FONIS SA04 I 2084 and FONDECYT 1050734.

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