**Klebsiella pneumoniae** producing extended spectrum β-lactamase in Regional Military University Hospital of Oran, Algeria: antibiotic resistance, biofilm formation, and detection of *bla*_{CTX-M} and *bla*_{TEM} genes

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**Abstract:**

**Background:** *Klebsiella pneumoniae* is a bacterial pathogen commonly associated with severe nosocomial and community acquired infections especially through the acquisition of extended spectrum β-lactamases (ESBL) and biofilm formation capacity. The objectives of this study are to determine the prevalence of *K. pneumoniae* ESBL (KP-ESBL)-producing isolates in the Regional Military University Hospital of Oran (HMRUO) Algeria, characterize their antibiotic resistance profile, genetically detect *bla*EM and *bla*CTX-M genes, and evaluate their biofilm formation capacity.

**Methodology:** Different clinical specimens including blood, cerebrospinal fluids, urine and catheter, pus, perirectal abscess, and surgical wounds were collected from patients with suspected clinical infections in different units and departments of the hospital. The specimens were cultured on Blood, MacConkey and CLED agar (for urine only) plates and incubated aerobically for 24 hours at 37°C for preliminary identification of bacteria using conventional colony morphology, Gram stain reaction, and disk diffusion test for antibiotic susceptibility testing (AST). Confirmation of isolates, antibiogram, minimum inhibitory concentration (MIC) and detection of resistance phenotypes, were carried out by the automated Vitek 2 (BioMérieux) identification and susceptibility method. ESBL production was confirmed by the synergy and combination disk tests. ESBL genes were detected by conventional simplex PCR and biofilm formation was detected by the tissue culture plate (TCP) method.

**Results:** A total of 630 patients’ clinical samples (one sample per patient) were processed. *Klebsiella pneumoniae* was isolated in 40 (6.3%) samples, and 15 of these (37.5%) produced ESBL. In the disk diffusion AST assay, all 40 *K. pneumoniae* isolates were resistant to ampicillin and ticarcillin while all 40 isolates were sensitive to cefoxitin, imipenem and ertapenem. KP-ESBL producing isolates were more frequently recovered from intensive care unit (33.3%) and from urine (46.7%) samples. Group 1 *bla*CTX-M genes were detected in 13 of the 15 (86.7%) KP-ESBL isolates, and 46.7% of these isolates were moderate biofilm producers.

**Conclusion:** There is need for health departments to put in place preventive measures through regular surveillance of these resistant pathogens and initiating appropriate infection prevention and control strategies to limit their spread in Algerian hospitals and worldwide.

**Keywords:** *Klebsiella pneumoniae*, ESBL, biofilm, PCR, antibacterial resistance

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**Klebsiella pneumoniae** producitrice de-lactamase spectre tendu dans l’hôpital universitaire militaire régional d’Oran, Algérie: résistance aux antibiotiques, formation de biofilm et détection des gènes *bla*_{CTX-M} et *bla*_{TEM}

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**Abstrait:**

**Contexte:** Klebsiella pneumoniae est un pathogène bactérien communément associé aux infections nosocomiales et communautaires sévères, en particulier par l’acquisition de β-lactamases à spectre étendu (ESBL) et la capacité de formation de biofilm. Les objectifs de cette étude sont de déterminer la prévalence des isolats de K. pneumoniae producteurs de βLSE (KP-βLSE) au CHU d’Oran (HMRUO) Algérie, caractériser leur profil de résistance aux antibiotiques, déceler génétiquement les gènes blatem et blactX-M, et évaluer leur capacité de formation de biofilm.

**Méthodologie:** Différents échantillons cliniques, y compris du sang, des liquides céphalo-rachidiens, de l’urine mictionnelle et du cathéter, du pus, des abcès périrectaux, et des plaies chirurgicales, ont été prélevés des patients suspects d’infections cliniques dans différentes unités et départements de l’hôpital. Les échantillons ont été cultivés sur des milieu de culture: déglise au sang, MacConkey et CLED (pour l’urine uniquement) et incubés en aérobie pendant 24heures à 37°C pour l’identification préliminaire des bactéries en utilisant la morphologie conventionnelle des colonies, la coloration de Gram et le test de diffusion sur disque pour les tests de sensibilité aux antibiotiques (AST). La confirmation des isolats, l’antibiogramme, la concentration minimale inhibitrice (CMI) et la détection des phénomètes de résistance ont été réalisés par la méthode automatisée d’identification et de sensibilité sur Vitek 2 (BioMérieux). La production de βLSE a été confirmée par les tests de synergie et de double disques. Les gènes de βLSE ont été détectés par PCR simplex conventionnelle et la formation de biofilm a été détectée par la méthode de la plaque de culture tube à air (TCP).

**Résultats:** Un total de 630 échantillons cliniques de patients (un échantillon par patient) ont été traités. Klebsiella pneumoniae a été isolé dans 40 échantillons (6,3%) et 15 d’entre eux (37,5%) ont produit des βLSE. Dans le test AST à diffusion sur disque, tous les 40 isolats de K. pneumoniae étaient résistants à l’ampicilline et à la ticarcilline, tandis que les 40 isolats étaient sensibles à la céfoxitine, à l’imipénème et à l’ertapénème. Les isolats producteurs de KP-βLSE ont été plus fréquemment récupérés dans les unités de soins intensifs (33,3%) et dans les échantillons d’urine (46,7%). Les gènes blactX-M du groupe 1 ont été détectés dans 13 des 15 isolats de KP-βLSE (86,7%), et 46,7% de ces isolats étaient des producteurs de biofilm modérés.

**Conclusion:** Il est nécessaire que les services de santé mettent en place des mesures préventives grâce à une surveillance régulière de ces pathogènes résistants et à la mise en place de stratégies appropriées de prévention et de contrôle des infections pour limiter leur propagation dans les hôpitaux algériens et dans le monde.

**Mots clés:** Klebsiella pneumoniae, βLSE, biofilm, PCR, résistance antibactérienne

**Introduction:**

Bacterial antibiotic resistance (ATB) is constantly evolving. For over 30 years, antibiotic resistance among enterobacteria to the third-generation cephalosporins (3GC) has been steadily increasing, notably through the production of extended-spectrum beta lactamases (ESBLs). These enzymes including TEM, SHV, CTX-M and their derivatives confer resistance on enterobacteria to all β-lactam agents with exception of cephemycins and carbapenems (1,2). While ESBL-producing enterobacteria were mostly observed in hospitals with varying isolation frequencies from hospital to hospital, and even from department to department within the same hospital, the diffusion of these multiresistant pathogens into community is of increasing concern. The transmission of genes encoding ESBL, mainly through plasmids, is responsible of their rapid dissemination and thus account for the increased prevalence of ESBL-producing bacteria worldwide, which constitute a major public health problem (1,3).

Klebsiella pneumoniae is an important opportunist pathogen causing nosocomial and community associated infections (4). The organism is part of the normal microflora of the intestine and commonly responsible for severe infections of the respiratory tracts (hospital acquired and ventilator associated pneumonia), catheter-related urinary tract infection, meningitis, blood stream infections (bacteremia and sepsicaemia), infections of surgical and non-surgical wounds, diarrhea, diseases, prosthetic valve endocarditis, peri-tonitis, and osteomyelitis (5-11).

The first ESBL-producing K. pneumoniae strains were first reported in Europe in 1982 where a new resistance to ceftazidime and aztreonam from plasmid-transmitted β-lactamase enzyme was quickly and easily disseminated to other Gram-negative bacteria including Escherichia coli. Since the discovery of these enzymes, they have not stopped growing, and today there are over 200 different ESBL enzymes. The impact of the ESBL strains is very significant, especially in the intensive care units with a high propensity for epidemic outbreaks. It has been shown in previous European studies that K. pneumoniae and E. coli were the two most common bacterial species frequently involved in ESBL production (12).

There is need for active surveillance
for ESBL-producing pathogens in high-risk populations using appropriate antimicrobial techniques because these pathogens are generally multiresistant (12,13). ESBL genes are typically carried by large transferable plasmids (85–275kb) on which they are often other associated genes coding for resistance to aminoglycosides, chloramphenicol, sulfonamides, trimethoprim, tetracyclines and fluoroquinolones (12). Although, the β-lactamase inhibitors are used to treat serious infection caused by ESBL-producing K. pneumoniae, this should be done with caution because in vitro sensitivity does not necessarily predict in vivo efficacy (13). Surveys conducted in different countries during specific time periods revealed that once a CTX-M β-lactamase enters a specific geographic area, it becomes predominant, and displace or superimpose itself over TEM and SHV ESBL variants (14).

The main virulence factors contributing to the pathogenesis of K. pneumoniae are the capsular polysaccharides and pili type 1 and 3 (6,7). Biofilms formation is another virulence factor. Biofilms are bacterial populations linked by exopolysaccharide matrices at the surface. This matrix of extracellular polymeric substances (EPS) consists mainly of polysaccharides, proteins, lipids and nucleic acids in varying amounts (11). The first strain of K. pneumoniae producing biofilms was described in the late 1988 (7). Biofilm-forming bacteria are often observed on the urinary catheter and on the surface of tissues and biomaterials usually at sites of persistent infection, where biofilm formation is a major cause of implant failure, and often limits the lifetime of many permanent medical devices (7,11).

The ability of K. pneumoniae to form biofilm provides protection against the host immune system as well as to antibiotics, therefore, biofilms are a major cause of resistance to antimicrobial agents, with resultant high costs of treatment of infections caused by these strains (15). For these reasons, information on ESBL and biofilm producing organisms are required for implementation of preventive procedures and application of infection control measures. The objectives of this study are; to determine the prevalence of ESBL-producing K. pneumoniae, isolates and their antibiotic resistance profile, genetically detect blat://TEM and blat://CTX-M genes, and evaluate their capacity to form biofilms.

Materials and method:

Study setting and design
This descriptive cross-sectional study was conducted over a period of 5 months (from October 29, 2017 to March 29, 2018) on 630 patients from different units and departments of the Regional Military University Hospital of Oran, Algeria, whose clinical samples were collected and processed at the Microbiology Laboratory of the hospital.

Sample and data collection, culture, bacteria isolation and identification
Clinical specimens comprising voided urine (n=333), pus (n=173), blood for culture (n=98), cerebrospinal fluids (n=12), catheter urine (n=8), perirectal abscess (n=4), and surgical wound (n=2) were collected by standard procedures from patients in different departments/units (paediatrics, infectious disease, general surgery, oncology, pneumology, haematology, intensive care unit, urology, re-education, nephrology, child surgery, and psychiatry) of the hospital, and from some external patients. From each patient, socio-demographic information including age and gender, specimen types, service units, and sampling date, were collected into a designed collection form.

Samples were cultured on Blood and MacConkey agar plates (Fluka), with the exception of urine samples which were cultured on Cysteine Lactose Electrolytes Deficient (CLED) agar medium. All culture plates were incubated aerobically for 24 hours at 37°C, and bacteria were preliminarily identified by conventional microbiological methods of colony morphology and Gram reactions, and then purified by successive subcultures on MacConkey agar purity plates. Confirmation of K. pneumoniae isolates, antibiogram, minimum inhibitory concentrations (MICs) determination and detection of resistance pheno-
types were performed with automated VITEK 2 (BioMérieux) ID and susceptibility platform, using GN and AST-N 233 tapes. The susceptibility results were interpreted according to the recommendations of CLSI (16).

Synergy test for ESBL production
Synergy test was performed under the same conditions of the antibiogram after detection of resistance phenotypes by VITEK 2 (BioMérieux) 1D. Amoxicillin/clavulanic acid (AMC) disk (20/10μg) was placed at 30 mm center to center of ceftazidime (CAZ) disk (30μg), cefotaxime (CTX) disk (30μg), ceftriaxone (CRO) disk (30μg) and aztreonam (ATM) disk (30 μg) on MH agar plate that has been inoculated with 0.5 McFarland standards suspension of K. pneumoniae isolates. Klebsiella pneumoniae ATCC 700603 was used as control. ESBL production results in the appearance of an image of synergy or champagne plug (17).

Combination disk test for ESBL production
The combination disk test was performed by placing clavulanic acid disk (10μg) and a third generation cephalosporin (3GC) disk at a distance of 30 mm on a MH agar
plate that has been inoculated with 0.5 McFarland standards suspension of *K. pneumoniae* isolates (test), and *K. pneumoniae* ATCC 700603 as control. ESBL production was phenotypically confirmed when the inhibition zone diameter of the 3GC disk tower applied after diffusion of the clavulanic acid disk is ≥ 5 mm in relation to the inhibition zone diameter of the 3GC disk tower (17).

**Molecular detection of ESBL genes**

Conventional simplex PCR for the detection of *blaCTX-M* and *blaTEM* genes was performed in the laboratory of Department of Public Health in Naples, Italy, with specific primers (18-20) as shown in Table 1. The bacterial DNA was extracted by the boiling method (21). PCR was performed in a thermal cycler (Applied Biosystems 2720, California, USA) and the amplification conditions were: initial denaturation at 94°C for 5 mins followed by 30 cycles at 94°C for 25 seconds, annealing at 52°C for 40 seconds, elongation for 72°C for 50 seconds and final elongation at 72°C for 6 minutes. The PCR products were electrophoresed on 1.5% agarose gel, visualized in the UV transilluminator after staining with ethidium bromide, and then photographed. The molecular weight marker (100 bp) was used to determine the sizes of the expected bands (Table 1).

**Biofilm formation by Tissue Culture Plate method (TCP)**

Test for biofilm formation for the *K. pneumoniae* isolates was performed as previously described (22,23) with some modifications. For this purpose, 96-well polystyrene microplates were used. Isolates were grown on nutrient agar for 18-24 hours at 37°C. One colony of each strain was inoculated in 5 ml of Brain Heart Infusion Broth (BHIB) and incubated at 37°C for 24 hours, then diluted to 1:100 in BHIB + 2% sucrose. Each well in the microplate was filled with 200μL of this dilution (three independent cultures for each species). A sterile broth of BHIB+2% sucrose was used as a negative control. The microplates were incubated at 37°C for 24 to 48 hours, after which the well contents were gently removed and rinsed four times with TBS (pH 7.2) and allowed to dry at 60°C for 30 min. Then, the cells adhering to the polystyrene support in the wells were coloured with 200μL of 1% (w/v) crystal violet for 30 minutes incubation at room temperature. The extra purple crystal violet was poured out and rinse four times at the sterile physiological waters. 200μL of ethanol 95% (v/v) was added to each well and left for 15 minutes before reading the optical density (OD) at 550nm using a microplate reader (Bio-Rad ELISA, PR 5100).

The interpretation of the results was carried out according to Vuotto et al., (15). The OD of the strains was obtained by the average of the three wells and compared to the OD (mean absorbance) of the negative control (ODc). Non-biofilm producer had ODc<OD, weak biofilm producer had ODc<2xODc, moderate biofilm producer had 2ODc<ODc≤4xODc, and strong biofilm producer had 4xODc<OD.

**Statistical analysis**

Data were analysed and presented on Excel sheet as frequency distribution tables and simple graphs. The susceptibility data were analyzed and interpreted using the WHONet 5.6 antibiotic susceptibility surveillance software.

![Image](image_url)
Results:

During the study period, a total of 630 samples from 630 patients with clinical infections were collected and processed for isolation and identification of bacteria in the laboratory. *K. pneumoniae* was isolated in 40 (6.3%) samples. In the AST assay, all the *K. pneumoniae* isolates were resistant (100%) to ampicillin and ticarcillin, 10% to amoxicillin/clavulanic acid, and 2.5% to piperacillin/tazobactam. On the other hand, 100% of the isolates were sensitive to cefotixin, imipenem and ertapenem, while sensitivity to other antibiotics were amikacin (97.5%), ofloxacin (87.5%), ciprofloxacin (82.5%), nalidixic acid (67.5%), gentamicin (67.5%), tobramycin (65%), ceftazidime (62.5%), cefotaxime (61.5%), cephalothin (60%), nitrofurantoin (55%), and trimethoprim/sulfamethoxazole (50%) (Fig 1).

The MIC of cefotaxime (CTX) was ≥ 64μg/ml, MIC of ceftazidime (CAZ) ranged from 1–64μg/ml, MIC of cefoxitin (FOX) was in the range of ≤4–8 μg/ml and MIC of amoxycillin/clavulanic acid (AMC) was between 16 and ≥ 32μg/ml. Fifteen (15) of the 40 (37.5%) *K. pneumoniae* isolates were ESBL-producing strains (Fig 2a,b) but no carbapenemase producing strain was detected.
The ESβL-producing isolates were more frequently recovered from urine (46.7%), followed by pus, blood culture and perirectal abscess (13.3% each), and least from surgical wounds and catheters (6.7% each). There were no ESβL-producing strains in the cerebrospinal fluids (Table 2).

<table>
<thead>
<tr>
<th>Samples</th>
<th>Number positive for K. pneumonia (%)</th>
<th>No positive for ESβL (%)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Surgical wound (n=2)</td>
<td>1 (25)</td>
<td>1 (6.7)</td>
</tr>
<tr>
<td>Perirectal abscess (n=6)</td>
<td>2 (50)</td>
<td>2 (13.3)</td>
</tr>
<tr>
<td>Catheter urine (n=8)</td>
<td>2 (50)</td>
<td>1 (6.7)</td>
</tr>
<tr>
<td>Cerebrospinal fluid (n=12)</td>
<td>2 (50)</td>
<td>0</td>
</tr>
<tr>
<td>Blood (n=98)</td>
<td>2 (50)</td>
<td>2 (13.3)</td>
</tr>
<tr>
<td>Pus (n=173)</td>
<td>11 (27.5)</td>
<td>2 (13.3)</td>
</tr>
<tr>
<td>Urine (n=333)</td>
<td>20 (50.0)</td>
<td>7 (46.7)</td>
</tr>
<tr>
<td>Total (n=630)</td>
<td>40</td>
<td>15 (27.5)</td>
</tr>
</tbody>
</table>

Twenty-seven K. pneumoniae isolates (65.7%) were recovered from male patients, 13 (48.2%) of which were ESβL producers, while 13 isolates (32.5%) were recovered from female patients, 2 (15.4%) of which were ESβL producers. The age group distribution of the patients with KP-ESβL-producing isolates shows that 3 patients (20%) were in the age group 21-30 years, while 2 patients (13.3%) each were in age groups <10 years, 10-20 years, 31-40 years, 41-50 years, 51-60 years and >60 years. Most ESβL-producing strains were isolated in the intensive care unit (33.3%, n=5), followed by general surgery (20%, n=3), paediatrics and urology units (13.3%, n=2) each, and 6.7% (n=1) each from nephrology, re-education and psychiatry units (Table 3).

<table>
<thead>
<tr>
<th>Service</th>
<th>Number of K. pneumoniae isolates</th>
<th>No positive for ESβL (%)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Paediatrics</td>
<td>3</td>
<td>2 (33.3)</td>
</tr>
<tr>
<td>Infectious disease</td>
<td>6</td>
<td>0</td>
</tr>
<tr>
<td>General surgery</td>
<td>6</td>
<td>3 (50)</td>
</tr>
<tr>
<td>Oncology</td>
<td>1</td>
<td>0</td>
</tr>
<tr>
<td>Pneumology</td>
<td>3</td>
<td>0</td>
</tr>
<tr>
<td>Haematology</td>
<td>1</td>
<td>0</td>
</tr>
<tr>
<td>Intensive care unit</td>
<td>8</td>
<td>5 (33.3)</td>
</tr>
<tr>
<td>Urology</td>
<td>5</td>
<td>2 (33.3)</td>
</tr>
<tr>
<td>Re-education</td>
<td>1</td>
<td>1 (40)</td>
</tr>
<tr>
<td>Nephrology</td>
<td>4</td>
<td>1 (20)</td>
</tr>
<tr>
<td>Child surgery</td>
<td>1</td>
<td>0</td>
</tr>
<tr>
<td>Psychiatry</td>
<td>1</td>
<td>1 (40)</td>
</tr>
<tr>
<td>Total</td>
<td>40</td>
<td>15 (37.5)</td>
</tr>
</tbody>
</table>

Group 1 bla<sub>CTX-M</sub> gene was detected by PCR from 13 (86.7%) of the 15 KP-ESβL producing isolates, while 2 (13.3%) isolates did not contain the gene (Fig 3a). No bla<sub>TEM</sub> gene was detected in any of the isolates (Fig 3b). The results of the biofilm formation with TCP test for KP-ESβL producing isolates showed that 7 (46.7%) were moderate, 7 (46.7%) were weak biofilm producers, while 1 (6.7%) was non biofilm producer (Fig 4).

![Fig 3a: Gel electrophoresis of the PCR products of bla<sub>CTX-M</sub> gene](image-url)
Clinical ESBL producing Klebsiella pneumoniae isolates in Algeria

Second lane, molecular weight marker; Gene Ruler 100 bp DNA ladder, lanes number 1, 2, 3, 4, 5, 6, 7, 8, 9, 10, 11, 12, 13, 14, 15 shows negative result. NC: Negative control

Fig 3b: Gel electrophoresis of the PCR products of blatan gene

Fig 4: Detection of biofilm formation by the TCP method

**Discussion:**

During the study period, K. pneumoniae was involved in 6.3% of clinical infections among the 630 patients, a rate that is similar to that reported by Khalifa and Khedher (12). All the K. pneumoniae isolates were resistant to ampicillin and ticarcillin (100%), which is consistent with the findings of Lagha et al., (24) and Muggeo et al., (25) in their previous studies. However, they were highly sensitive to amoxicillin/clavulanic acid (AMC) (90%) and piperacillin/tazobactam (97.5%), a finding that is also consistent with that of Rasa-miravaka et al., (26) especially for AMC, although Muggeo et al., (25) and Khalifa and Khedher (12) reported K. pneumoniae isolates with intermediate resistance to AMC in their studies, while higher AMC resistance rates were reported by Lagha et al., in 2014 (24) and Benaicha et al., in 2017 (27). However, for piperacillin/tazobactam combination, the resistance rate (2.5%) in our study is lower than the rates reported by Lagha et al., (24) and Muggeo et al., (25). The decrease in the resistance rate may be an indicator of the presence of CTX-M type β-lactamases and the association of penicillins with inhibitors, which permit recovery of the activity of the molecules. The β-lactamase inhibitors have structural similarity with penicillin, and are effective against many sensitive organisms expressing class A lactamases (17). In addition, the combination of penicillin and β-lactamase inhibitor has been actively used in the treatment of infections caused by ESBL-producing bacteria and could be proposed for outpatient treatment of urinary tract infections caused by ESBL-producing Escherichia coli strains (1,28).

All the K. pneumoniae isolates in the study were sensitive to cefoxitin, imipenem and ertapenem which agrees with the reports of Alibi et al., (20) and Lagha et al., (24), al-
though different susceptibility rates have been reported in many other studies (11, 15,25,26,29). Currently, carbapenems are the only class of antimicrobials historically effective against KP-ESBL producing strains (13), however, it is essential to ensure the rational use of carbapenems because there are no new antibiotics in the pipeline available for use in the near future for the treatment of infections caused by ESBL-producing Enterobacteriaceae (1). On the other hand, K. pneumoniae isolates in our study were highly sensitive to amikacin, ofloxacin and ciprofloxacin, and moderately sensitive to cephalothin, cefotaxime, ceftazidime, gentamicin, tobramycin, and nalidixic acid, which agrees with findings of some other studies (15,17,24,27,29).

However, Mugeeo et al., (25) and Rasamiravaka et al., (26) reported lower sensitivity (83%) to amikacin in their studies compared to 97.5% in our own study. Only half of the K. pneumoniae isolates were sensitive to trimethoprim/sulfamethoxazole and nitrofurantoin. Rasamiravaka et al., (26) and Vuotto et al., (15) have reported differing susceptibility rates of K. pneumoniae isolates to trimethoprim/sulfamethoxazole in their studies. The fluoroquinolones (FQ) are the potential drug of choice for treating infections caused by β-lactamase-producing enterobacteria that are usually FQ sensitive (25). However, Mugeeo et al., (25) reported 100% resistance of K. pneumoniae ST395 to fluoroquinolones in north-eastern France, which is contrary to the findings of high susceptibility in our study. Klebsiella species are naturally sensitive to FQ, but the misuse of these antibiotic in human and veterinary medicine practices have, over the past decades, resulted in evolution of resistance to this antibiotic family, reduced their effectiveness, and compromised the future use of this important class of antibacterial drugs (12).

In our study, 15 of the 40 (37.5%) K. pneumoniae isolates were ESBL-producing strains, which is proximate to 41.1% rate reported by Pirzaman et al., (2). Different prevalence rates for ESBL-producing strains have been reported in Tunisia (12), Algeria (24) and Russia (30), and ESBL rates are usually higher in Asian countries, with up to 75% (17). The phenomenon of ESBL production has been observed in various pathogenic bacteria, but more frequent in E. coli and K. pneumoniae. The KP-ESBL strains were most frequently recovered from urine in our study which is consistent with the findings of other studies (2,26,27,31,32,33), probably because urine is about the most commonly collected specimens for clinical investigation. Infections caused by ESBL-producing bacteria can occur in people of all ages, but distribution could be determined by the immunological status of patients and prevalence of antibiotic misuse. KP-ESBL isolates were recovered across all age groups in our study but most frequently in age group 21-30 years (20%), although the number of isolates from patients in ours study are too few to make any significant inference. However, Lagha et al., (24) reported that the most KP-ESBL were recovered from patients between the ages 27-85 years, Gravey et al., (34) reported 4.1% KP-ESBL rate in the age group 18–64 years and 4.2% in the age group >65 years. K. pneumoniae isolates as well as KP-ESBL strains were mostly recovered from male patients in our study. This is in close agreement with those of Bush et al., (33), Deng et al., (35) and Lagha et al., (24).

Also, most KP-ESBL strains were isolated from the intensive care and general surgery units of the hospital, with 33.3% and 20% rates respectively. This agrees with the finding of Lagha et al., (24), but Khalifa and Khedher (12) reported lower rates than ours, with rates of 5% in paediatrics, 2.5% in medical resuscitation, and 0% in surgical unit. Numerous studies have reported isolation of KP-ESBL strains from hospitalized patients and nosocomial epidemics caused by these strains have been reported mainly in intensive care units (20). We did not isolate KP-ESBL strain from infectious, oncology, pneumology, haematology and child surgery departments of the hospital. Although, this might suggest good infection prevention and control practices in these service departments, the limited KP-ESBL strains in this study cannot allow us to generalize this finding.

PCR assays detected group 1 blaCTX-M genes in most (86.7%) of the KP-ESBL producing isolates, while few (13.3%) isolates did not carry the gene. This is similar to what Alibi et al., (20) reported in their study for blaCTX-M but in addition, they reported blaTEM in 56.8% of their isolates. However, Paterson et al., (36) reported 23.3% blaCTX-M and 87% blaTEM among their isolates. In Abidjan, Côte d’Ivoire, Guessennd et al., (31) reported that 63.4% of their strains carried blaTEM, 58.5% carried group 1 blaCTX-M, none carried groups 2 and 9 blaCTX-M, one strain carried group 8 blaCTX-M and three strains carried blaTEM, blaSHV and group1 blaCTX-M. In another study in Korea by Jung et al., (32), only group 1 blaCTX-M (75.9%) and/or group 9 blaCTX-M (20.5%) were reported. In an Egyptian study, blaCTX-M was reported in three cases and blaTEM was detected only in one case (37). KP-ESBL strains carrying blaCTX-M and/or blaTEM genes are usually resistant to third generation cephalosporins (ceftriaxone, cefotaxime, ceftazidime), and several studies have shown that the presence of these genes could confer resistance to the third-generation cephalosporins (38). With the wide-
spread use of cefotaxime and ceftriaxone (36), it is not surprising that blaCTX-M ESBL strains are now found in many countries and reported at high rates among K. pneumoniae and other bacteria pathogens (20). It has been widely reported in the literature that biofilm production provides a significant benefit by protecting pathogens against host immune system and reducing susceptibility to antibiotic therapy. In our study, 46.7% of KP-ESBL strains were moderate and 46.7% were mild biofilm producers, which agrees with the finding of Seifi et al., (7). Martino et al., (39) reported 48.5% of K. pneumoniae strains to be strong biofilm producers; Vuotto et al., (15) reported 67.5% of strains to be potent, and 25% to be moderate biofilm producers, and Khodadadian et al., (11) reported 91.2% of their isolates to be biofilm producers. Burgers et al., (40) have reported a close relationship between several virulence factors and the ability to produce biofilm. The ability of K. pneumoniae strains to adhere and colonize inert surfaces may be a general feature of this species, as high incidence of effective adhesion of K. pneumoniae strains to glass and polypropylene surfaces in clinical and water distribution systems, have been observed (39).

Conclusion:

Infections caused by ESBL-producing Gram-negative bacteria are increasing, particularly in immunocompromised patients and in high-risk units of hospitals. These infections are associated with higher costs of healthcare in Algeria and worldwide, as a result of prolonged hospitalization and the use of expensive drugs. The prevalence of KP-ESBL-producing strains reported in this study is a reflection of the level of infection prevention and control practices in our hospital.

Surveillance of these antibiotic resistant pathogens by detection of β-lactamases and molecular identification of prevalent ESBL genes, as well as good knowledge of biofilms formation, will be essential for reliable epidemiological characterization of these pathogens, in order to prevent the risk of transmission, and implement antibiotic stewardship and appropriate infection control measures.

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References:

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