Clinical Review: management of Lower respiratory tract infection in pediatric population

Ghasna Mohiuddin¹, Syed Wasif Gillani², Anam Azhar¹, Rukhsar A. Majeed¹

¹Pharm D candidate, College of Pharmacy Gulf Medical University, Ajman, UAE
²Assoc. Prof. Department of Pharmacy Practice, College of Pharmacy Gulf Medical University, Ajman, UAE

Received: 02 October, 2019
Accepted: 09 December, 2019

*Correspondence to:
Dr. Syed Wasif Gillani, PhD
Email: dr.syedwasif@gmu.ac.ae

INTRODUCTION

Lower respiratory tract infections are usually caused by bacteria and virus. The pediatric population is observed to be the most susceptible to it as the antibiotics are not used properly thus leading to resistance (1) LRTIs can be divided into pneumonia, chronic bronchitis and acute exacerbations of chronic obstructive pulmonary disease (2,3). These infections can affect people of all age, but mostly its children who are likely to suffer (3). It has also been statistically proven that around 13% of all infectious illnesses are in infants younger than 2 years (4).

The respiratory rate thresholds displayed by World Health Organization for children with this infection are as follows: Children younger than 2 months have greater than or equal to 60 beats per minute, Children aged between 2-11 months show greater than or equal to 50 breaths per minute, and children aged around 1-3 years have greater than or equal to 40 breaths per minute (5).

Congestion, fever and irritability are a few signs and symptoms of LRTI (4) Various tests were taken into consideration to diagnose this particular infection, for example: Auscultation by stethoscope, cultures, complete blood cell count (CBC) and chest radiography & Ultrasonography (3).

A broad spectrum of antibiotics was used to treat lower respiratory tract infections. High-dose amoxicillin is used as a first line (5). Combination therapy (ampicillin and either gentamicin or cefotaxime) is typically used in the initial treatment of newborns and young infants (4). Carbapenems (B-lactams), are one of the safest as it has the least side-effects, however the frequent use of it can lead to resistance (6,7). The time period, drug which is administered, and the strength of
the dose can vary depending on different factors such as the patient’s age, etc. (8). Cephalosporin’s are the other class of drugs used in children, which includes 4 generations (9). This class of the drug is one of the most effective when it comes to the bacteria which has been exposed for long as it inhibits bacterial cell wall synthesis. (10) Third generation cephalosporin’s are less effective against gram positive cocoa when compared to gram negative rods. However, they are the most used at they are effective in treating bacterial meningitis in children (2).

First generation of cephalosporins: Cephalexin

Cephalexin belongs to the first class of cephalosporins with the chemical name, 7-(D-a-amino -a-phenylacetamido)-3-methyl-3-cephem-4-carboxylic acid, and it is related to cephaloglycin. Cephalexin is a known bactericidal antibiotic (11). It is a white crystalline compound with high solubility hence orally administered (12). It gets easily dissolved in 2mg/ml distilled water at room temperature (14). It’s not effected by plasma protein significantly hence it is excreted without any metabolical changes.

The in vitro antimicrobial functioning of cephalexin is not equivalent to the in vitro activity of cephaloglycin. (11) Cephalexin is very quickly solubilized but it’s binding sites available are not sufficient hence, balances out low in vitro antimicrobial activity. (13) When comparing the lab activity of cephaloglycin, tetracyclin and chloramphenicol with cephalexin, cephalexin gave much better results when the same procedure was carried out on mice with bacterial infection (11,14)

A clinical study was carried out on 12 people, 6 males and 6 females, all of them being healthy individuals. The age of these samples units was between 20 and 41 years, weight of these individuals between 52.5 and 76 kg where mean weight = 67.8kg. All of these volunteers did not have any medication for past 14 days. The results obtained were as such: there was a rapid increase to the average peak concentration of 38.8 ± 8.1 mg / liter (averaging 55.5 + 21.8 min). And also relatively a very rapid reduction to just 1.2 mg / liter after 6 hours, culminating in a biological half-life of 62.5 ± 5.5 min (13)

Second generation of cephalosporins: Cefoxitin

Cefoxitin is a 8-lactam cephamycin antibiotic, injectable. It’s administration is either intravenously or intramuscularly(16). It is usually preferred for pediatric population as it has high antibacterial activity(16). In vitro studies show it’s high effectiveness on many common pathogens usually found in pediatrics(17). Cefoxitin is used for bacteriae like: Streptococcus pneumonia, Strepococcus aureus, Haemophilus influenzae etc(17). it’s acts by absorbing onto the proteins therefore inhibiting the cell wall functioning such as synthesis but its activity is highly dependent on the kind of organism and where it is present because that determines how well it can be absorbed hence the dose strength is determined(18).

A study was carried out on 32 infants ranging from 3 month old to 26 month old and the recorded results were shown in figure 1 (19):

![Figure 1: Results of administration of Cefoxitin to 32 infants.](image)

For checking how efficient cefoxitin is, regular tests are done on patients as well as healthy individuals as references and bacterias are extracted before they are administered with any kind of medications, antibiotics specifically, and cultured and examined(19).
### Generic Name | Therapeutic Use | Dose | Absorption | Distribution | Metabolism | Elimination | Excretion
--- | --- | --- | --- | --- | --- | --- | ---
Cefpodoxime | Acute Community-Acquired Pneumonia | >12 years 200 mg PO q12hr for 14 days. | BA: 50%, acid stable. Peak plasma time: ≤1 hr | Distributed well into tissues, also lungs and tonsils; penetrates into pleural fluid. Protein bound: 18-23% | Metabolized in liver to active metabolite | Half-life: 2-3 hr; prolonged with renal impairment | Urine (80%) as unchanged drug in 24 hr

Cefotaxime (25,35) | Pneumonia | ≤12 years or <50 kg: 200 mg/kg/day IV divided q8hr >12 years or >50 kg: 1-2 g IV/IM q8hr. | peak plasma time: IM, 30 min | widely distributed to body tissues and fluids, including aqueous humor, ascitic and prostatic fluids, bone, penetrates CSF when meninges inflamed | Partially metabolized in liver | Metabolite: Desacetylcefotaxime (active) | Half-life: Parent drug, 1-1.5 hr; active metabolite, 1-1.9 hr | Urine.

Ceftazidime (27) | Hospital acquired pneumonia (HAP) and Community acquired pneumonia (CAP) | If child is 4 weeks - 30mg/kg every 12Hours ≤If child is 1 month - 12 y/o – 30 to 50mg/kg every 8 hours (28,30) | IV Peak concentration after IV administration was 109mg | Peak concentration after IM administration was 53mg | aqueous humour, bile, blister fluid, bone, bronchial secretions, CSF, abcess, myometrium etc | Protein binding: 5-24%. (31) | not metabolized by the liver and as a result its clearance and elimination from the body is solely dependent on the RENAL SYSTEM. (29) | IV T1/2 of 4.17 | IM T1/2 of 3.8 | A 24 hour urianalysis showed that 89% of drugs was recovered unchanged.

Ceftibuten (32,33,34) | Bronchitis | 12 years or older: 400 mg orally once a day for 10 days | Rapid; food decreases peak concentrations, delays Tmax, and lowers AUC. peak time: 2 to 2.6 hours | Distributes into middle ear fluid, bronchial secretions, and sputum. Protein binding: 63% | predominantly eliminated through kidney in the pediatric population. | Half life elimination: Children: 2 hours; | Urine (56%); feces (39%) |

### Figure 2: Graphical presentation of comparative study of two successful antibiotics for LRTIs in pediatrics. Pathogens separated from 59 cases included H. influenzae (47.5%), S. pneumoniae (23.7%), M. catarrhalis (11.9%) and million deaths due to these infections are otherwise preventable hence, physicians are now progressively using other substitutes such as third generation cephalosporins. (23) A third-generation cephalosporin, cefpodoxime proxetil has a broader spectrum and it is also active against the bacteria responsible for LRTIs in children. A study was done to assess its clinical effectiveness in children with acute febrile LRTIs as shown in figure 2. (23)

*Haemophilus parainfluenzae* (6.8%). There was no difference in bacteriologic efficacy of the two treatments, and both drugs were well-tolerated. Thus, cefpodoxime proxetil is highly effective and well-tolerated in childhood pneumonia and is at least as effective as the comparator drug. (23).
Fourth generation of cephalosporins: Cephalosporins, Cefepime, Maxipime

Maxipime is a semi-synthetic cephalosporin antibacterial whose route of administration is usually parenteral (20). Maxipime is used to treat moderate Pneumonia triggered by Streptococcus Pneumonia, Klebsiella Pneumonia, Pseudomonas aeruginosa or Enterobacter species (20). Maxipime can slow down the progression of resistance of the bacteria towards the drugs and also maintain the effect of maxipime and also other drugs used to treat (20). When treating with maxipime, epidemiological conditions and also the patterns (if available) should be looked into (20).

These drugs are used for pediatric populations, so a study was carried out on 88 patients who were treated with cefepime (14). The age of these patients was between 2 months old to 26 years old (14). Cefepime was administered through IV route and IM route (15). Single IV dose, multiple IV doses and IM doses (15). In all the patients, after steady dosages, 2 days later urine or CSF samples were collected (15). In all the results collected cefepime was observed to be decreasing sometime after and when multiple doses were given, no significant improvement in the results hence proving that the dose is independent of any parameter (15). The results were as listed in table 2 (15):

**Table 2:** Results from urine or CSF sample collected from patients administered cefepime.

For moderate to severe pneumonia, the dose prescribed should be 50mg/kg every 8 hours (17). Cefepime was well absorbed when given through IM route and it also penetrated into CSF in patients suffering from bacterial meningitis, differentiated with beta lactams (14,17).

Fifth generation of cephalosporins: Ceftaroline

Ceftaroline is a fifth-generation cephalosporin antibiotic. It acts by preventing the production of cell wall (B-lactamase) (21). They adsorb on the surface of PBP-2a, manufactured by MRSA and PBP-2a which is synthesized by penicillin resistant Streptococcus pneumonia (22). It has high activity on gram positive and gram negative (22). It is used to treat skin infections (also bacterial infections on the skin) triggered by MRSA and many common pathogens (23).

Ceftaroline is a prodruk and it has to be metabolised to convert it into its active form (21). It is metabolised by phosphate enzymes, the drug is later hydrolysed to inactive form (21). It is usually administered by injections, it is highly effected by the strength of the drug, as stronger the dose higher the Cmax concentration as well as AUC (21). The drug is recommended for regular use for fourteen day but not more than 14 because if used for fourteen days at regular intervals the drug shows no complications in any of the body compartments, metabolism and excretion is controlled well as the drug as low affinity towards proteins hence less binding (22). The approved doses are as shown in table 3 (22):

**Table 3:** Ceftaroline doses based on age and weight of the child.

<table>
<thead>
<tr>
<th>Age/weight</th>
<th>Dose</th>
</tr>
</thead>
<tbody>
<tr>
<td>Neonates and infants younger than 2 months</td>
<td>6 mg/kg every 8 hr</td>
</tr>
<tr>
<td>2 months to less than 2 years old</td>
<td>8 mg/kg every 8 hr</td>
</tr>
<tr>
<td>2 years old and above</td>
<td>12 mg/kg every 8 hr</td>
</tr>
<tr>
<td>Weight less than 33 kg</td>
<td>400 mg every 8 hr</td>
</tr>
<tr>
<td>Weight more than 33 kg</td>
<td>600 mg every 12 hr</td>
</tr>
</tbody>
</table>

If the patient has cystic fibrosis, it will have an impact on ceftaroline as its pharmacokinetics in the body as it enhances the mean clearance level and also elivated the total distribution volume throughout the body, reason being malnutrition and less deposition in the adipose tissues hence decreasing the total drug concentration in the body (20). A study was carried on a six year old child suffering from cystic fibrosis and another six year old healthy child who does not have any other health issues (21). The child suffering from cystic fibrosis has shown some improvement clinically but
there is a risk of child gaining the resistance towards the drug hence the drug cannot be used for a long term (21).

Ceftaroline was tested randomly as well as controlled experiments were done on children from the age of six months to 24 months and then it was approved by the ABSSI and CABP for pediatric population (under the age of two months) (22). Ceftaroline is not considered to be the first choice of drug but later when FDA approved it to be a bactericidal it is considered to be a good alternative option (24). Some common Side effects of ceftaroline are rashes, fever, gastrointestinal irritation and also the effects it causes (25).

**Carbapenams:**

Carbapenams are of the traditional antibiotics which are given by IV and have a time duration of 15-60 minutes (36). They are used to treat a variety of diseases, for example, nosocomial pneumonia, community acquired and *pseudomonas* infections in cystic fibrosis which are very severe and needs treatment before the condition worsens. (38) When you compare the structures of penicillin and carbapenams, they are quite similar except the fact that the sulfur atom is replaced by a carbon atom at position one and are therefore classified as beta lactam antibiotics, and play a role against both gram positive and negative bacteria. (7) Carbapenams are time-dependent antibiotics as they kill the bacteria when the drug concentration is above the minimum concentration required to cause an inhibitory effect (MIC). (41)

Carbapenams mode of action initiates as they move through the bacterial wall and binds to penicillin binding proteins (enzyme) (37) Transpeptidase is the main target enzyme of carbapenams during bacterial cell wall synthesis. (40) A covalent bond is formed by penicillin-binding proteins which consists of carboxypeptidase and transpeptidase enzyme. (7) This helps in preventing the cross-linkage of peptides during peptidoglycan biosynthesis, therefore resulting in cell apoptosis within the bacterial cell (41). Table 4 gives us detailed knowledge of the dose, excretion and the distribution of various type of carbapenems (6)

### Table 4: Recommended dose according to the child’s age and their respective pharmacokinetic data,

<table>
<thead>
<tr>
<th>Drug</th>
<th>Pediatric Age</th>
<th>Dose</th>
<th>Excretion</th>
<th>Distribution</th>
</tr>
</thead>
<tbody>
<tr>
<td>Imipenem/cilastatin</td>
<td>Newborns (&lt;1 week)</td>
<td>25mg/kg every 12 hours</td>
<td>Clearance rate: 2.3-6ml/kg/min</td>
<td>Vd = 0.36-0.67L/kg</td>
</tr>
<tr>
<td></td>
<td>1-4 weeks</td>
<td>25mg/kg every 8 hours</td>
<td>Half-life: 1-3 hours</td>
<td></td>
</tr>
<tr>
<td></td>
<td>1-3 months</td>
<td>25mg/kg every 6 hours</td>
<td></td>
<td></td>
</tr>
<tr>
<td></td>
<td>3 months and above</td>
<td>15-25mg/kg every 6 hours</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Meropenem</td>
<td>Neonates</td>
<td>20mg/kg every 12 hours</td>
<td>Clearance rate: 1.8-5.63ml/kg/min</td>
<td>Vd = 0.43-0.48L/kg</td>
</tr>
<tr>
<td></td>
<td>3 months and less than 50kgs</td>
<td>20mg/kg every 8 hours</td>
<td>Half-life: 1-3 hours</td>
<td></td>
</tr>
<tr>
<td></td>
<td>More than or equal to 50kgs</td>
<td>1g every 8 hours</td>
<td></td>
<td></td>
</tr>
<tr>
<td></td>
<td></td>
<td>2g/dose for meningitis</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Ertapenem</td>
<td>3 months-12 years</td>
<td>15mg/kg every 12 hours</td>
<td>Clearance rate: 2.1-3.3ml/kg/min</td>
<td>Vd= 0.2L/kg</td>
</tr>
<tr>
<td></td>
<td>More than or equal to 1 year</td>
<td>1 day every day</td>
<td>Half-life: 2-5.4 hours</td>
<td></td>
</tr>
<tr>
<td>Doripenem</td>
<td>3 months to less than 2 years</td>
<td>10mg/kg every 8 hours</td>
<td>Clearance: 3.1-6.5ml/kg/min</td>
<td>Vd= 0.22-0.46L/kg</td>
</tr>
<tr>
<td></td>
<td>More than or equal to 2 years</td>
<td>15mg/kg every 8 hours</td>
<td>Half-life: 0.9-1.1 hours</td>
<td></td>
</tr>
<tr>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

Bronchitis and pneumonia are the two major types of acute lower respiratory tract infections which increase the death rate in children under 5 years of age. (1) The most common pathogen was *Staphylococcus aureus* which had affected infants with lower respiratory tract infections *streptococcus pneumonia and E.coli* which was treated using Methicillin. *E.coli* showed less resistance to cephalosporin’s compared to other antibiotics such as piperacillin and ampicillin. Aminoglycosides were however contraindication to be used for children as they caused major side-effects such as hearing loss and achondroplasia (40,41). *Staphylococcus aureus* causes 25% of community acquired pneumonia cases. Mostly, the cases are related to skin and soft tissue infection. In addition to this, socio-economic factors such as low income, poor lifestyle, limited health care due to insufficient breast feeding and other reasons such as air pollution therefore causing these
respiratory tract infections to become acute (8) However, more research has been conducted which suggests that antenatal care and routine immunization can help prevent diseases and save many lives (39)

Beta-lactamase are the enzymes which are produced by bacteria that causes resistance to some antimicrobials and can degrade β-lactam antimicrobials. (38) However, carbapenems which contains β-lactamase inhibitors cannot be degraded so easily (40). Carbapenems can cause hepatotoxicity when jaundice occurs as they have an insignificant hepatic metabolism effect (37). This is a very rare condition caused by carbapenems but should be taken into consideration when giving it to children. (41)

An enzyme, called dehydropeptidase-1 (DHP-1) is located in the renal tubules which allows most of the carbapenems to be broken down. (36) Due to this reason, it is advised for carbapenems to be administered with cilastatin which is a DHP-1 inhibitor. (38) The potency of such antibiotics is reduced due to increased resistance and other cost containment issues. (41) If the dose carbapenems is not taken with complete cautious, it gives rise to other several issues leading to more resistant organisms. (37)

Penicillins and Aminoglycosides

Table 5 summarises the mechanism of action, dose, absorption, distribution, metabolism and excretion of penicillins and aminoglycosides.

| Table 5: Pharmacokinetic data of Penicillin and its derivatives, and Aminoglycosides. |
|-----------------|-----------------|-----------------|
| **Mechanism of Action** | **Amoxicillin (43)** | **Ampicillin (42)** | **Penicillin (42)** |
| **Dose** | <3 months: 30 mg/kg/day PO divided q12hr for 48-72 hours; > 10 days for S. pyogenes infections. | <20 kg: 50mg/kg/day PO divided q6-8hr >20kg: 250mg PO q6hr | 250 mg PO q8-12hr for 10 days |
| **Absorption** | Bioavailability: 74-92% Peak plasma time: 2hr(capsule) 3.1hr (extended release tab) 1hr (suspension). | Bioavailability: 30-40% Peak plasma time:1-2hr (oral) | Bioavailability: 60-73% Peak plasma time: 0.5-1hr |
| **Distribution** | Protein bound: 15-25% Blister and tissue fluids, bile, and CSF with inflamed meninges | Protein bound: 80% Crosses placenta; poor Blood-Brain Barrier diffusion. | |
| **Metabolism** | Hepatic | Hepatic | Hepatic |
| **Excretion** | Excretion in urine Half-life: 3.7hr(full-term neonates); 1-2hr (infants and children). | Excretion in urine (90% within 24 hr). Half-life: 1.1-1.8hr (normal renal function); 7-20hr (anuria/end-stage renal disease). | Excretion in urine Half-life: 0.5-0.6hr |
| **Gentamycin (3)** | Amikacin (3) | Tobramycin (5) |
| **Mechanism of Action** | It irreversibly binds to specific 30S-subunit proteins and 16S rRNA (specifically binds to four nucleotides of 16S rRNA and a single amino acid of protein S12). | Hinders mRNA binding and rRNA acceptor sites by binding to bacterial 30S ribosomal subunits which results in disrupted normal protein synthesis and non-functional or toxic peptides are produced. As a result, it interferes with bacterial growth. | Aminoglycoside has two binding sites on the 30S ribosomal subunit, to one of which tobramycin binds and inhibits bacterial protein synthesis. Additionally, tobramycin may also bind to 16S rRNA and destabilize bacterial membrane. |
| **Dose** | (Children, 1 year or older) 6 to 7.5mg/kg/day IV/IM in equally divided doses every 8 hours for 7 to 10 days. (Infants and neonates, 1 year or younger) 7.5 mg/kg/day | Neonates: 10mg/kg IV/IM loading dose followed with 7.5 mg/kg IV/IM every 12 hours; MAX 15 mg/kg/day total daily dose. Infants and children: 15 mg/kg/day IV/IM divided | Premature or full-term neonates, 1 week or younger: up to 4mg/kg/day IV in 2 equally divided doses every 12 hours. Greater than 1 week of age: 6-7.5mg/kg/day IV in 3-4 equally divided doses every 6 to 8 hours. |
**Figure 3**: Comparison between penicillin and aminoglycosides’ safety and toxicity.

The collected data shown in figure 3 suggests that the best choice of antibiotic in the treatment of LRTI in pediatrics are penicillin’s since their efficacy to toxicity balance is better than in aminoglycosides.
SAFETY MENAMICS
Penicillin is one of the safest antibiotics which was used as it inhibits different type of bacteria without leading to severe side-effects which would make the condition more severe. Following this cephalosporin’s and carbapenems are considered to be beneficial as well. When the 5 generations of cephalosporins are compared, third, fourth and fifth generation agents achieve high concentration in the CSF as it can cross the blood brain barrier successfully and have a tremendous activity against gram negative aerobic bacteria including the pathogens responsible for bacterial meningitis against children compared to the first and the second. However, first generations are potentially useful against mild-moderate infections caused by gram positive bacteria. Aminoglycosides are potent but cause life-threatening side effects so they are the least safe class of antibiotics.

Penicillin > Cephalosporins (Third generation) > Fourth generation > Fifth generation > First generation > Second generation > Carbapenems > Aminoglycosides

CONCLUSION
This study analysed the effectiveness of different classes of antibiotics which could be used in the treatment of lower respiratory tract infections in the paediatric population. The efficacy, potency, safety and the pharmacokinetics of the antibiotics were the main factors which were taken into consideration when making the suitable choice as children especially infants who have a very sensitive immune system as it is still developing therefore is not completely function. Due to this reason, they are at a very high risk of infection. Aminoglycosides are the most potent hence they’re used for very specific infections (narrow spectrum) however, they cause toxicity so TDM is necessary (42). Penicillins on the other hand, are the safest in terms of toxicity since the only reported side effects are diarrhea and headache, but their frequent use, as it is a broad-spectrum antibiotic, has led to resistance (42). The best penicillin class antibiotic is Amoxicillin as it has the highest bioavailability compared to ampicillin and penicillin (43). When comparing the lab activity of cephaloglycin, tetracyclin and chloramphenicol with cephalixin, which is a first-generation cephalosporin gave much better results when the same procedure was carried out on mice with bacterial infection (11,14). Cefoxitin, which is a second-generation cephalosporin is usually preferred for pediatric population as it has high antibacterial activity (16) but dose should be carefully adjusted. For third generation Cephalosporins (Cefpodoxime, Cefotaxime, Cefazidime and Cefitiben), Cefazidime and Cefitiben are predominantly are eliminated through the renal system hence should be given with care in patients with renal problems (27, 33, 34). Cefepime; 4th generation cephalosporin was well absorbed when given through IM route and it also penetrated into CSF in patients suffering from bacterial meningitis, differentiated with beta lactams (14,17). Ceftaroline (5th generation cephalosporin) has high activity on gram positive and gram negative (22).

Carbapenems are another class of antibiotics that are used to treat a variety of diseases, for example, nosocomial pneumonia, community acquired and pseudomonas infections in cystic fibrosis which are very severe and needs treatment before the condition worsens however, their use is less because bacteria are becoming resistant to it. Additionally, Carbapenems may sometimes cause hepatotoxicity when jaundice occurs as they have an insignificant hepatic metabolism effect (37) so this should also be considered before using them in children.

REFERENCES
24. BNF. Chapter 18, Prescribing in children (2018); 76: 44-45.
34. Nix DE. Comparative pharmacokinetics of oral cefributen, cefixime, cefaclor, and cefuroxime axetil in healthy volunteers (1997); 17(1): 121-5.
37. Xiaoguang He. Antimicrobial resistance in bacterial pathogens among hospitalized children with community acquired lower respiratory tract infections in Dongguan, China. BMC infect Dis (2017); 17: 614.
42. Padari H. Pharmacokinetics of Penicillin G in preterm and term neonates. Antimicrobe agents Chemother (2018); 62(5).