Pharmacokinetics, Pharmacodynamics, and Pharmacogenomics

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LEARNING OBJECTIVES

1. Evaluate the physiologic changes that affect drug absorption and distribution in critically ill patients.
2. Distinguish the effects of changing hemodynamics on drug metabolism and elimination and predict potential pharmacokinetic changes in critically ill patients.
3. Apply knowledge of how different drug properties affect the influence of physiologic changes on pharmacokinetics in critically ill patients.
4. Justify pharmacotherapy on the basis of pharmacodynamic parameters that correspond with the maximal efficacy of commonly used antimicrobials.
5. Demonstrate the value of pharmacogenomic applications with commonly used drugs in critically ill patients.

ABBREVIATIONS IN THIS CHAPTER

<table>
<thead>
<tr>
<th>Abbreviation</th>
<th>Definition</th>
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<tbody>
<tr>
<td>AKI</td>
<td>Acute kidney injury</td>
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<tr>
<td>Clh</td>
<td>Hepatic clearance</td>
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<td>Clint</td>
<td>Intrinsic clearance</td>
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<tr>
<td>Cmax</td>
<td>Maximum concentration</td>
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<td>CYP450</td>
<td>Cytochrome P450</td>
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<tr>
<td>Ef</td>
<td>Hepatic extraction efficiency or ratio</td>
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<tr>
<td>Fu</td>
<td>Fraction unbound</td>
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<tr>
<td>GFR</td>
<td>Glomerular filtration rate</td>
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<tr>
<td>KDIGO</td>
<td>Kidney Disease: Improving Global Outcomes</td>
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<td>LNPEP</td>
<td>Leucyl/cystinyl aminopeptidase</td>
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<td>PCI</td>
<td>Percutaneous coronary intervention</td>
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<td>PD</td>
<td>Pharmacodynamics</td>
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<td>PGx</td>
<td>Pharmacogenomics</td>
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<td>PK</td>
<td>Pharmacokinetics</td>
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<tr>
<td>Q</td>
<td>Hepatic blood flow</td>
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<tr>
<td>TDM</td>
<td>Therapeutic drug monitoring</td>
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<tr>
<td>Vd</td>
<td>Volume of distribution</td>
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Table of other common abbreviations.

INTRODUCTION

Pharmacotherapy of critically ill patients is challenging. Given the relative dearth of data available specifically for patients in ICUs, critical care pharmacists must use a combination of data extrapolation and clinical intuition to design optimal therapeutic dosing plans. Furthermore, each critically ill patient is distinct, with a differing pathogenesis and a rapidly changing physiology, which may further complicate treatment choices. The development of an individualized therapeutic plan requires the selection of a dosing and monitoring regimen that balances available information with the highest likelihood of positive outcome and minimal adverse effects. To successfully develop individualized dosing regimens and administer them to critically ill patients, an understanding of pharmacokinetics (PK), pharmacodynamics (PD), and pharmacogenomics (PGx) is paramount.

Every drug entering the body follows an identical process of absorption, distribution, metabolism, and elimination—but one that is unique to that specific medication. That process ultimately determines how much drug is available at the targeted site of action. Pharmacokinetics refers to the sum of the processes the body is conducting on the drug. In contrast, pharmacodynamics refers to the physiologic and biochemical effects of the drug on the body. The intended effects of the drug, at a concentration that minimizes potential adverse effects, are determined by the intricate balance between PK and PD. Pharmacogenomics refers to a patient’s possible shift in the balance of PK and PD through innate genetic polymorphisms, which can alter the way the body and the drug (or its metabolites)
interact with each other. The relationship between PK, PD, and PGx is further complicated by the rapid physiologic changes seen in critically ill patients. An understanding of the general principles of PK, PD, and PGx can assist clinicians in making the best choices when designing a drug regimen for critically ill patients. This chapter describes the general concepts behind PK, PD, and PGx and their use as principles in clinical examples seen in the critical care setting.

**PK CONCEPTS AND DERANGEMENTS IN CRITICALLY ILL PATIENTS**

**Absorption**

*Absorption* refers to the ability of a drug to migrate from the site of administration into the bloodstream. The extent of absorption is typically measured in terms of bioavailability, defined as the fraction of an administered dose that reaches the systemic circulation. All drugs administered outside the intravenous route are affected by absorption. However, few studies have directly evaluated the effects of critical illness on absorption. A number of factors may affect enteral absorption in critically ill patients (e.g., gastric pH, GI motility, bowel wall edema, splanchnic perfusion, first-pass metabolism, feeding tube or enteral feed interactions). Furthermore, the physiologic and therapeutic changes that occur during critical illness are unpredictable, and summation of the results may lead to an increased, decreased, or net unchanged enteral absorption. Theoretically, in states of decreased perfusion, the body’s natural physiologic response is to shunt blood toward vital organs and away from the GI tract, which may decrease the enteral absorption of drugs (Beale 2004). In patients requiring vasopressors for hemodynamic support, many studies have documented differing degrees of alterations in splanchnic perfusion (Hollenberg 2004).

Among the many reasons for decreased gastric motility in critically ill patients are abdominal surgery, opioid use, electrolyte abnormalities, ileus, immobility, and traumatic brain injury (Nguyen 2007). The delay in gastric emptying may alter absorption based on the drug’s characteristics. Absorption from the stomach is usually slow and favors acidic drugs, whereas absorption from the small intestines is usually fast and favors basic drugs. Delayed gastric emptying may decrease the absorption of drugs normally absorbed in the small intestines, because increased transit time from the stomach can lead to degradation. On the other hand, for drugs absorbed in the stomach, delayed emptying may increase absorption because of more time spent at the site of absorption (Prescott 1974).

Another common factor that may affect drug absorption is the use of enteral feeding tubes and formulas. Drugs administered through enteral feeding tubes may adhere to the lumen of the tube or have physical incompatibilities that limit bioavailability. In critically ill patients, decreased absorption from feeding tube drug administration has occurred with amiodarone, carbamazepine, ciprofloxacin, phenytoin, theophylline, digoxin, and warfarin (Lourenco 2001, Zhu 2013). In general, many drug-nutrient interactions can be circumvented by withholding enteral nutrition for 1 to 2 hours both before and after drug administration, although using that practice for drugs frequently administered may lead to suboptimal nutritional support.

The conclusion that critical illness has unpredictable effects on enteral absorption is further corroborated by discrepant clinical PK evaluations of drugs administered enterally in critically ill patients. In an open-label, single-dose study of healthy volunteers, hospitalized general ward patients, and critically ill patients, atorvastatin maximal concentration (C$_{max}$) and AUC 0 to 24 hours after administration were about 18- and 15-fold higher, respectively, in critically ill patients with sepsis (p<0.01 for both C$_{max}$ and AUC) compared with healthy volunteers. Notably, all doses were administered orally or, for those who could not swallow (23 of 25 critically ill patients), by way of a nasogastric tube. Although the results may have been influenced by the use of CYP inhibitors in some critically ill patients, similar results were observed in selected patients not receiving CYP inhibitors, wherein the mean AUC and C$_{max}$ were about 10-fold higher. Notably, despite the 10-fold-higher C$_{max}$ and AUC, those differences did
not reach statistical significance, most likely as a result of lack of study power (Kruger 2009).

The higher observed \( C_{\text{max}} \) and AUC in that study are in contrast with an evaluation of the effects of various clinical characteristics likely to affect paracetamol (acetaminophen) absorption. That study enrolled 27 ICU patients with normal hepatic and renal functions and no contraindications to nasogastric feeding. Of this group, 9 patients required the use of dopamine; those patients had significant decreases in paracetamol AUC\(_{0-60\text{min}}\) compared with those who were hemodynamically stable (mean 337.4 vs. 728.2 mg·L\(^{-1}\)·minute, \( p=0.005 \)). Notably, the usual AUC\(_{0-60\text{min}}\) in healthy volunteers is about 600 mg·L\(^{-1}\)·minute. This study also observed a considerable range in AUC\(_{0-60\text{min}}\) among all of the ICU patients (61.2 to 1400.3 mg·L\(^{-1}\)·minute) (Tarling 1997).

Last, in a study of the enteric absorption and PK of oseltamivir in 41 critically ill patients with suspected H1N1 influenza, the median and trough plasma concentrations of oseltamivir and its metabolite were similar to those observed in other clinical trials involving ambulatory patients (Ariano 2010). The aforementioned data are representative of the degree of variability in enteric absorption that could be expected on the parts of critically patients. Taken together, the data further confirm that enteral absorption of drugs in critically ill patients is highly erratic. If possible, enteral routes of administration should be avoided in acutely ill patients until the clinical status has been stabilized. Using alternative routes of administration may be especially important with drugs that have narrow therapeutic indexes. Regarding drugs without parenteral alternatives, clinicians should be aware of unpredictable absorption and extra vigilant in monitoring efficacy and safety.

Critical illness can also affect absorption in the subcutaneous route. Given the physiologic shunting of blood to vital organs during shock, perfusion to subcutaneous tissue may be affected. In a study evaluating the antifactor Xa activity of a subcutaneously administered low-molecular-weight heparin for deep-vein thromboembolic prophylaxis, the use of a vasopressor was an independent predictor of having an undetectable peak anti-factor Xa level (Jochberger 2005). In a separate study, the antifactor Xa activities in patients on prophylactic doses of subcutaneously administered low-molecular-weight heparin while on vasopressors were compared with antifactor Xa activities in hemodynamically stable, critically ill patients and hospitalized non-ICU patients. When compared with the two other groups, patients who received vasopressors had significantly lower concentrations of antifactor Xa activity that was not within the recommended therapeutic range (Dorffler-Melly 2002). In addition to physiologic shunting of peripheral blood volume to vital organs, aggressive fluid resuscitation, vasodilatory states, and third-space shifts of intravascular volume may cause significant peripheral edema, further decreasing subcutaneous absorption.

The effects of edema in critically ill patients have been evaluated in several studies. In a study of 20 multiple-trauma patients, a PK evaluation on subcutaneously administered enoxaparin was performed on patients classified as nonedematous or edematous (presence of edema and increase in admitting body weight of 10 kg or more). That investigation demonstrated highly variable AUCs in the entire cohort. In addition, a significant decrease in antifactor Xa activity was seen in edematous patients compared with nonedematous patients. In fact, in 7 of the 10 edematous patients, the majority of the sampled antifactor Xa assays were below the lower limit of detection (Haas 2005).

That finding was similar to finding in other investigations, which corroborated the relatively low plasma antifactor Xa activity in critically ill patients receiving standard doses of subcutaneous enoxaparin for deep-vein thrombosis prophylaxis (Dorffler-Melly 2002, Mayr 2002, Priglinger 2003). However, a study of 14 patients in a mixed medical-surgical ICU revealed that the presence of edema did not affect the \( C_{\text{max}} \) or AUC of dalteparin (Rommers 2006). Notably, the definition of peripheral edema as used in this study was slightly more conservative than the one used in the study mentioned earlier; hence the lack of difference in dalteparin concentrations may reflect the relatively lower magnitude of edema and/or the small sample size.

In summary, the subcutaneous absorption in critically ill patients—especially those on vasopressors or with more than 10 kg of fluid weight gains during their ICU admissions—is highly variable and most likely suboptimal.

**Distribution and Protein Binding**

The distribution of a drug depends largely on the drug’s hydrophilicity and its acid dissociation constant, which affects its binding to proteins and other macromolecules. Only a free drug, unbound by protein, can diffuse across tissue and have physiologic effect. Many acidic drugs are bound to albumin and many basic drugs are bound to \( \alpha \)-acid glycoprotein. During states of critical illness, albumin concentration decreases as a result of increased vascular permeability, decreased production, and catabolism (Fleck 1985). In contrast, \( \alpha \)-acid glycoprotein is an acute-phase reactant and tends to increase during states of stress (Edwards 1982). The changes in protein concentrations could affect both the amount of free drug available and the overall volume of distribution \( (V_d) \).

Decreased protein binding leads to an increased free fraction and overall \( V_d \) whereas increased protein binding has the opposite effect. Although decreased binding would lower the total serum concentration, increased free fraction may lead to better therapeutic effects than normally observed. In one study of the effect of albumin concentration on total and free phenytoin concentrations, free fraction of phenytoin was inversely related to plasma albumin concentration in 10 critically ill trauma patients (Boucher 1988). Hence,
Comparing the subcutaneous and skeletal muscle concentrations of piperacillin to plasma concentrations in six pairs of age- and sex-matched patients with septic shock and healthy patients. Using microdialysis methods, the investigators determined that in patients with septic shock, the piperacillin concentrations were 5- to 10-fold lower in the subcutaneous and skeletal muscle tissues compared with plasma concentrations. In the healthy volunteers, the free piperacillin concentrations in peripheral tissues were about 95% of those seen in plasma concentrations. Furthermore, the plasma concentrations were also significantly higher in healthy volunteers, which may reflect the increased Vd seen in critically ill patients (Joukhadar 2001). For hydrophilic drugs, aggressive fluid resuscitation may lead to increases in total body water, which results in a larger Vd and lower serum concentrations. That effect of aggressive fluid management has also been demonstrated among aminoglycosides and other β-lactam antibiotics (Dasta 1988, Triginer 1990).

In summary, critical illness may lead to changes in endogenous protein concentrations, tissue perfusion, and total plasma volume. The extent of those changes on each drug depends ultimately on the drug’s properties. Table 1-1 describes drugs commonly used in critical care, their characteristics that may affect volume of distribution, and the potential effects caused by critical illness.

<table>
<thead>
<tr>
<th>Drug Characteristic</th>
<th>Pharmacokinetic Characteristics</th>
<th>Examples</th>
<th>Critical-Illness Changes</th>
<th>Effect on Vd</th>
</tr>
</thead>
<tbody>
<tr>
<td>Basic charge</td>
<td>Bound to α1-acid glycoprotein</td>
<td>Azithromycin, carvedilol, fentanyl, lidocaine, meperidine, mirtaline, nicardipine, olanzapine, phenobarbital, verapamil</td>
<td>α1-acid glycoprotein is an acute phase reactant, which would increase in states of stress</td>
<td>Decreased volume of distribution, Decreased free fraction</td>
</tr>
<tr>
<td>Acidic or neutral charge</td>
<td>Bound to albumin</td>
<td>Amiodarone, ceftriaxone, diazepam, midazolam, morphine, phenytoin, prednisolone, propofol, valproic acid, warfarin</td>
<td>Decreased production and increased vascular permeability and catabolism lead to hypoalbuminemic states</td>
<td>Increased volume of distribution, Increased free fraction</td>
</tr>
<tr>
<td>Lipophilic</td>
<td>Distributes widely in tissue</td>
<td>Diazepam, fentanyl, fluoroquinolones, lorazepam, macrolides, midazolam, propofol, rifampin, tetracycline</td>
<td>Minimal changes</td>
<td>Minimal changes</td>
</tr>
<tr>
<td>Hydrophilic</td>
<td>Distribution primarily into serum</td>
<td>Aminoglycosides, β-lactams, daptomycin, hydromorphone, morphine, vancomycin</td>
<td>Fluid resuscitation, Third spacing, Reduced tissue perfusion</td>
<td>Increased volume of distribution, Decreased serum concentration</td>
</tr>
</tbody>
</table>

Metabolism

The liver plays a major role in the metabolism of many drugs, and critical illness and drug properties can affect hepatic clearance. Intrinsic patient factors include volume status, perfusion, gut motility, and liver function. The major drug properties that affect the quantity of elimination by the liver include extraction ratio and protein binding (Rodighiero 1999). To fully understand the impact of critical illness on a medication’s hepatic metabolism, an appreciation of the underlying determinants of hepatic clearance is necessary. Hepatic clearance ($Cl_h$) of a medication is a function of hepatic blood flow ($Q$) and the liver’s extraction efficiency (ratio) for the particular drug ($E_h$) (Verbeeck 2008); this can be represented by the formula:

$$Cl_h = Q \times E_h$$

A particular drug’s extraction efficiency depends on liver blood flow, intrinsic clearance of unbound drug ($Cl_{int}$), and the fraction of unbound ($f_u$) drug in the blood (Verbeeck 2008); it can be represented by the formula:

$$E_h = (f_u \times Cl_{int}) / (Q + (f_u \times Cl_{int}))$$

Together the equation for hepatic clearance is:

$$Cl_h = (Q \times f_u \times Cl_{int}) / (Q + (f_u \times Cl_{int}))$$

The equation contains the three primary components of hepatic drug elimination: blood flow, drug protein binding, and intrinsic clearance. Intrinsic clearance is based on the sum of all of the hepatic enzyme and transport activity involved in the removal of drug from the blood.

During acute illness, many pathophysiologic changes may affect drug metabolism and reduce $Cl_{int}$. During states of vasodilatory shock, cardiac output may be either normal, increased, or decreased depending on the progression of shock and the adequacy of fluid resuscitation. However, hepatic flow is likely to be impaired in states of low cardiac output—such as hypovolemia—or a state of cardiogenic or hemorrhagic shock. Furthermore, mechanical ventilation has decreased venous return, leading to decreased cardiac output and hepatic blood flow (Bonnet 1982). A reduction in liver perfusion may lead to a decrease in enzymatic activity. However, an increase in cardiac output may further increase the hepatic extraction of medications.

Two different types of reactions are primarily responsible for the liver’s metabolizing capabilities: phase I oxidative metabolism and phase II glycosylation and glucuronidation. Phase I reactions, which are usually mediated by the CYP isoenzymes, require the presence of oxygen molecules and are therefore more susceptible to functional deficiencies caused by lack of oxygenation from decreased hepatic perfusion. Being less reliant on the presence of oxygen, conjugation phase II reactions such as glucuronidation are less susceptible to the effects of liver cirrhosis and hypoxia (Westphal 1997).

Each drug’s extraction efficiency determines the extent to which its hepatic metabolism is affected by critical illness. Medications can be categorized according to the $E_h$: high ($E_h >0.6$), low ($E_h <0.3$), or intermediate ($0.3 < E_h <0.6$). Drugs with high extraction ratios depend on blood flow and are usually relatively insensitive to changes in protein binding or enzyme activity ($Cl_h = Q$). Drugs in this category are most influenced by cardiac output and hepatic blood flow. In fact, studies that evaluated drugs with intermediate to high extraction ratios have found increases in serum concentrations ranging from 2- to 12-fold from decreases in hepatic blood flow (Rodighiero 1999). On the other hand, drugs with low extraction efficiency are affected by changes in protein binding and intrinsic hepatic clearance ($Cl_h = f_u \times Cl_{int}$) (Verbeeck 2008). The hepatic elimination of those drugs depends more on hepatic enzymatic reactions and is more sensitive to decreases in metabolism from states of cirrhosis or acute liver dysfunction. Table 1-2 illustrates drugs commonly used in critical care and their respective extraction ratios.

### Table 1-2. Hepatic Extraction of Drugs Commonly Used in Critical Care

<table>
<thead>
<tr>
<th>Extraction Characteristic</th>
<th>Hepatic Extraction</th>
<th>Examples</th>
</tr>
</thead>
<tbody>
<tr>
<td>Low extraction</td>
<td>&lt;0.3</td>
<td>Amoxicillin, ceftriaxone, chlor Diazepoxide, clarithromycin, clindamycin, diazepam, doxycycline, fluconazole, lansoprazole, lorazepam, methadone, methylprednisolone, metronidazole, mycophenolate, oxazepam, phenobarbital, phenytoin, prednisolone, prednisone, primidone, rifampin, theophylline, valproic acid</td>
</tr>
<tr>
<td>Intermediate extraction</td>
<td>0.3–0.6</td>
<td>Alfentanil, amiodarone, atorvastatin, azathioprine, carvedilol, codeine, diltiazem, erythromycin, itraconazole, lidocaine, meperidine, nifedipine, omeprazole, ranitidine</td>
</tr>
<tr>
<td>High extraction</td>
<td>&gt;0.6</td>
<td>Fentanyl, isosorbide dinitrate, morphine, nitroglycerin, propofol, sufentanil</td>
</tr>
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</table>

Elimination

Although drugs can be excreted through the biliary tract, feces, and respiration, the kidneys eliminate most drugs and metabolites in the critical care setting, and they are the ones most likely to be affected by physiologic changes during acute illness. Drugs are eliminated by the kidney through glomerular filtration and tubular secretion. Although tubular secretion function is not directly measurable, glomerular filtration rate (GFR) is a good marker of overall renal drug clearance.

Although much focus is on acute kidney injury (AKI) and reduction of renal function, critical illness could also lead to augmented renal clearance. There is no universally accepted definition of augmented renal clearance, although a value of 10% above the upper limit of normal (GFR >160 mL/minute/1.73m² in men and >150 mL/minute/1.73m² in women) has been proposed (Udy 2010). Critical care conditions that are likely to lead to augmented renal clearance include those that induce low systemic vascular resistance and high cardiac output. Although there are no robust epidemiologic study data on the incidence of augmented renal clearance, it has been reported as high as 85% of all patients at some point during their ICU stay (Udy 2010). Augmented renal clearances have generally been observed in patients who are younger (<55 years), are post-trauma (particularly head trauma), are postsurgery, have sepsis, are diagnosed with hematologic malignancies, or have significant burn injuries (Albanese 2004, Bennmalek 1999, Brater 1986, Brown 1980, Conil 2007, Fuster-Lluch 2008, Lamoth 2009). Clinicians caring for patients with adequate urine output, no laboratory indications of AKI, and conditions in which renal blood flow may be increased should be aware of the possibility of augmented renal clearances and decreased serum concentrations of renally eliminated drugs. Of interest, direct correlations between augmented renal clearance and lower serum drug levels have been observed for many antimicrobial agents including β-lactams, aminoglycosides, and glycopeptides (Udy 2010).

Little is known about optimal adjustments to drug dosing in patients with AKI. The Kidney Disease: Improving Global Outcomes (KDIGO) society provides guidelines on drug-dosing considerations, but few details are offered for the management of patients with AKI (Matzke 2011). Box 1.1 lists recommended steps for assessing and adjusting drug regimens in patients with AKI or chronic kidney disease. Specifically for patients with chronic kidney disease, the KDIGO guidelines recommend that drug dosages be adjusted according to FDA-approved labeling. For patients with AKI, the guidelines acknowledge that there are few or no specific data. However, because of PK changes seen in AKI patients— including the potential for increases in Vț—and because of the inability to accurately estimate renal function, the guidelines recommend a liberal dosing approach. That approach entails, for hydrophilic drugs, administering an aggressive 25%–50% greater than normal loading dose, followed by a normal or near normal maintenance dosage (Matzke 2011).

### Box 1.1. Assessing and Adjusting Drug Regimens in Patients with Acute Kidney Injury or Chronic Kidney Disease

**Step 1 – Assess the following**
- Demographic information
- Medical history
- Current clinical information
- Current laboratory information
- DNA polymorphisms

**Step 2 – Estimate eGFR or creatinine clearance by using:**
- Age
- Body size
- Ethnicity
- Concomitant diseases

**Step 3 – Review current medications**
- Identify drugs needing individualized dosing

**Step 4 – Calculate individualized treatment regimen**
- Determine treatment goals (PK or PD values)
- Calculate dosage regimen (based on drug PK and changes noted in the patient)

**Step 5 – Monitor regimen**
- Drug response
- Signs or symptoms of toxicity
- Therapeutic drug monitoring if applicable

**Step 6 – Revise regimen**
- Adjust regimen according to patient response

eGFR = estimated glomerular filtration rate; PD = pharmacodynamics; PK = pharmacokinetics.

Adapted with permission from Swanson LM. Pharmacokinetics/Pharmacodynamics. ACCP Updates in Therapeutics 2015: Critical Care Pharmacy Preparatory Review Course; 2:335-66.

The rationale for such an aggressive approach is the importance of administering an adequate dose to quickly optimize pharmacologic activity, particularly for antimicrobials. In addition, in the setting of AKI, increased nonrenal clearances of some agents have been observed. The clinical applicability of that recommendation would clearly depend on the patient’s clinical status and the assessment of the drug’s therapeutic index with the potential for adverse reactions.

Critically ill patients with renal failure may require renal replacement therapy, which provides an extracorporeal mechanism for drug removal. The choice of dialytic technique depends on institutional practice and the patient’s hemodynamic status. There are considerable institutional differences in the use of continuous renal replacement therapy and of extended daily dialysis as regards dialysis machines, flow rates, filter sizes, and indications. As such, the medical literature provides little guidance about selecting the most appropriate dosing regimens for each specific dialytic mechanism. To select a dosing regimen, the KDIGO guidelines recommend a strategy that accounts for extracorporeal clearance of the drug. One such method is to use the formula:

\[
Dose = Dose_n \times \left[ \frac{Cl_{n\text{onrenal}} + (Q_{\text{eff}} \times SC)}{Cl_{\text{norm}}} \right]
\]
Where \( \text{dose}_n \) = normal dose; \( \text{Cl}_{\text{nonrenal}} \) = nonrenal clearance; \( Q_{\text{eff}} \) = effluent rate; \( SC \) = saturation coefficient (or sieving coefficient if using hemofiltration modes of dialysis); and \( \text{Cl}_{\text{norm}} \) = normal clearance (Matzke 2011). Normal clearance in this case can also be interpreted as the total body clearance seen in usual population without organ dysfunction.

In the foregoing equation, the saturation coefficient (or sieving coefficient) is, in general, similar among different continuous renal replacement therapy machines and membranes. In situations when the saturation coefficient is not available, the fraction unbound provides a reasonable estimate. A nephrology team usually chooses an effluent rate based on a patient’s weight and needs for solute removal. The extent of nonrenal clearance varies with the medication. In some instances, patients may also have residual renal function, which should be accounted for.

Given the complexity of the calculations, institutions should attempt to quantify their most common methods and practices associated with continuous dialytic methods and derive appropriate dosing guidelines for commonly administered drugs. The literature contains several continuous renal replacement therapy dosing reviews (Churchwell 2009, Heintz 2009, Kuang 2007, Trotman 2005), but most of these studies used older dialysis machines or filters in an era when dialysis and replacement fluid flow rates were relatively low. In other words, the doses described in clinical reviews may not be adequate for current dialytic modes and practices.

**PK Summary**

The effects of critical illness on PK are vast and unpredictable. Ultimately, serum concentrations of medications administered to critically ill patients depend on the interaction between physiologic derangements and drug properties. Figure 1-1 summarizes the physiologic effects of critical illness and the corresponding effects on drug concentrations. If medications are to exert their intended effects, the establishment of adequate serum concentrations in critically ill patients is important—especially when, for certain agents, the drug’s PD rely on achieving a particular concentration target.

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**Figure 1-1.** Effect of critical illness on pharmacokinetics of medications.

- Drugs excreted by kidneys are susceptible to augmented renal clearance.
- Applicable for hydrophilic medications.
- Acidic or neutral drugs usually bound to albumin increase their free fractions; basic drugs decrease their free fractions.
- Likely to affect hydrophilic drugs with a relatively small volume of distribution (less than plasma blood volume).

ECMO = extracorporeal membrane oxygenation; RRT = renal replacement therapy.
PHARMACODYNAMICS

Clinical Pharmacodynamic Applications in Antimicrobials

The majority of PD evaluations and applications for critically ill patients involve the management of antimicrobials, which is the focus of this section. Antimicrobials function through the inhibition of vital processes seen in microbiologic pathogens. For antimicrobials, the best PD responses, defined as improvements in microorganism clearance or in clinical outcomes, are more likely when certain targets are achieved. Table 1-3 summarizes known PD targets for some of the most commonly used antimicrobials in the ICU. The desired antimicrobial PD targets correspond to reaching a time, concentration, or AUC ratio above the organism’s MIC. As such, the probability of antimicrobial PD target attainment and desired antimicrobial effect depends on the interplay between pathogen resistance and each patient’s PK parameters.

Antibiotics such as aminoglycosides induce bacterial killing through concentration-dependent PD, wherein the rate and extent of the kill increases as the ratio between the maximum concentration and MIC increases. In an analysis of 236 patients treated with aminoglycosides for gram-negative infections, high Cmax/MIC ratios were associated with improved outcomes.

### Table 1-3. Pharmacodynamic Targets of Antimicrobials

<table>
<thead>
<tr>
<th>Antimicrobial PD Category and Corresponding Antibiotics</th>
<th>PD Targets</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>Concentration Dependent</strong></td>
<td></td>
</tr>
<tr>
<td>Aminoglycosides</td>
<td>Cmax/MIC 8–10 (Drusano 2007); AUC/MIC &gt;70 (Highet 1999)</td>
</tr>
<tr>
<td>Metronidazole</td>
<td>N/A</td>
</tr>
<tr>
<td><strong>Time Dependent</strong></td>
<td></td>
</tr>
<tr>
<td>Penicillins</td>
<td>40–50% T &gt; MIC (Craig 1996)</td>
</tr>
<tr>
<td>Cephalosporins</td>
<td>60–100% T &gt; MIC (Crandon 2010; McKinnon 2008)</td>
</tr>
<tr>
<td>Carbapenems</td>
<td>50–75% T &gt; MIC (Ariano 2005; Li 2007)</td>
</tr>
<tr>
<td><strong>Concentration Dependent and Time Dependent</strong></td>
<td></td>
</tr>
<tr>
<td>Fluoroquinolones</td>
<td>AUC/MIC &gt;125 (Ariano 2005; Li 2007)</td>
</tr>
<tr>
<td>Vancomycin</td>
<td>AUC/MIC &gt;400 (Moise-Broder 2004)</td>
</tr>
<tr>
<td>Linezolid</td>
<td>AUC/MIC &gt;80–120 (Rayner 2003)</td>
</tr>
<tr>
<td>Tigecycline</td>
<td>AUC/MIC 13–18 (Bhavnani 2012; Rubino 2012)</td>
</tr>
<tr>
<td>Daptomycin</td>
<td>AUC/MIC 38–171 (Dandekar 2003)</td>
</tr>
<tr>
<td>Colistin</td>
<td>AUC / MIC 7-23 (Dudhani 2010)</td>
</tr>
<tr>
<td>Fluconazole</td>
<td>AUC / MIC &gt;12 (Baddley 2008)</td>
</tr>
</tbody>
</table>

Cmax = maximal concentration; PD = pharmacodynamics; T > MIC = time above MIC during dosing interval.

infections, improvements in clinical response were seen in more than 90% of patients with peak:MIC ratios of 10 or more (Moore 1987). Similar findings were observed in patients with gram-negative nosocomial pneumonia (Kashuba 1999). Given the availability of aminoglycoside serum concentration monitoring and the variability in Vd observed in critically ill patients, clinicians should perform first-dose pharmacokinetic evaluations with a view to aggressive dosing of these antibiotics so as to target known PD parameters.

For antibiotics such as fluoroquinolones, when serum concentration monitoring is not readily available, clinicians should still be familiar with the targeted PD parameters and understand the importance of hitting those targets. Using modeling data, clinicians should be able to assess clinical scenarios in which the probability of PD goal target attainment is lower than ideal. For example, with fluoroquinolones, as the MIC approaches 1 mcg/mL (even with aggressive dosing regimens), the probability of PD target attainment drops off considerably (Drusano 2004b).

As demonstrated in the previous section, critically ill patients have a number of physiologic changes that may alter the PK of antimicrobials. The changed PK has the propensity to decrease the probability of antimicrobial PD target attainment. Indeed, using microbiologic data generated from 14 ICUs and Monte Carlo simulation of PK parameters garnered from published reports of infected patients, one team demonstrated that routine β-lactam dosing had a probability of target attainment as low as 45% (DeRyke 2007). Notably, simulations of high-dose ceftazidime and cefepime (2 g every 8 hours) against Pseudomonas aeruginosa exceeded the PD target more than 90% of the time. An open-label multicenter study evaluated the PK profiles of 80 critically ill patients treated with aggressively dosed β-lactam (e.g., piperacillin 4 g every 6 hours, cefepime 2 g every 8 hours, ceftazidime 2 g every 8 hours, meropenem 1 g every 8 hours); the proportion of patients who met prespecified PD targets for meropenem, ceftazidime, cefepime, and piperacillin-tazobactam were 75%, 28%, 16%, and 44%, respectively (Taccone 2010). Notably, that study’s PD target was time greater than four times the MIC, which is considerably more aggressive than the target used in the DeRyke study.

In a recent prospective, multinational, PK point-prevalence study of 248 ICU patients treated with β-lactams, 16% of patients did not achieve the PD target of 50% of free drug concentration T > MIC. Patients who did not achieve the PD target had a 32% lower chance of achieving a positive clinical outcome (Roberts 2014c). In that point-prevalent study, the antibiotic dosing and administration methods were dictated by the treating physician and varied among the different sites.

Aside from the decreased propensity for critically ill patients to reach PD targets because of PK changes, evidence suggests that PD targets in critical care should be different and higher than those seen in non-critically ill patients. Indeed, few of the targets proposed in Table 1-3 have been validated in critically ill patients. In fact, many of the targets were derived from animal models, in vitro kill studies, or non-critically ill patients.

Although most clinicians assume that the PD targets should be the same in critically ill patients, the data to support that hypothesis is sparse; some studies suggest that higher PD targets are necessary. In a study of 76 patients with serious infections treated with either cefepime or ceftazidime, both clinical cure and bacteriologic cure were significantly higher if drug concentrations were above the MIC throughout the entire dosing interval (100% T > MIC) (McKinnon 2008). This begs the question of whether, for severe infections, a higher proportion of time spent above the MIC for β-lactams is warranted.

Traditional in vitro studies also suggest that, for β-lactams, further increases in concentration above the MIC do not produce additional bacteriologic killing (Drusano 2004a). However, in a PK study of 101 patients with lower respiratory tract infections being treated with meropenem, the only PD target significantly correlated with clinical success was 100% time spent above five times the MIC (Li 2007). Taken together, it appears that the optimal PD parameter for patients with severe infections might not be fully elucidated. In a recent survey of nine ICUs in which therapeutic drug monitoring (TDM) of β-lactams is routine, the PD targets ranged from 100% T > MIC to 100% T > 4xMIC (Wong 2014), which is considerably higher than traditionally recommended.

The potential need for higher PD targets poses additional challenges for critically ill patients. Aside from the high inter-patient variability because of unpredictable PK, critically ill patients also have a higher risk of acquiring resistant organisms with higher MICs. In an epidemiologic study of resistance among gram-negative bacilli in ICU patients over a decade, consistent increases in resistance rates were observed (Lockhart 2007). The constellation of higher baseline MICs, the potential need for higher PD targets, PK changes that may lower serum concentrations, and dosing recommendations that are largely one size fits all based on healthy volunteers makes selection of an adequate antimicrobial dosing regimen for critical care patients a considerable challenge.

Two strategies have emerged as potential options for the application of antimicrobial PD principles in critically ill patients: TDM of antimicrobials (including ones not traditionally considered for TDM) and the use of alternative-dosing-administration techniques (e.g., continuous or extended infusions for β-lactams and once-daily dosing for aminoglycosides).

Evidence from pharmacokinetic modeling and Monte Carlo simulation suggests that the probability of PD target attainment for β-lactams is enhanced through continuous or extended infusion methods (Lodise 2006). However, whether the use of continuous- or extended-infusion β-lactams improves clinical outcomes remains a controversial topic, with meta-analyses demonstrating conflicting results (Falagas 2013, Roberts 2009). To address this controversy, a prospective, randomized, controlled, double-dummy trial compared β-lactams...
(piperacillin-tazobactam, ticarcillin-clavulanate, or meropenem) with continuously administered and intermittent 30-minute β-lactam infusions in 432 patients with severe sepsis in 25 ICUs. The study attempted to address many of the study design issues that had precluded clinical interpretation from previous studies. The issues included administration of a loading dose, double-blind design, selection of severely ill patients, administration of the same total daily dose in each arm, and aggressive dosing regimens. The study did not perform pharmacokinetic evaluations to determine whether continuous infusions of β-lactams were in fact associated with improvements in target attainment. The study found no significant differences in any primary or secondary outcomes, which included ICU-free days, 90-day mortality, 14-day clinical cure, duration of bactemia, and organ-failure-free days. In addition, no differences in adverse effects were observed (Dulhunty 2015).

Although this study found a lack of clinical outcome benefits with continuous infusions of β-lactams, several considerations remain. Of the 432 patients, only 83 actually had a pathogen isolated from blood. Furthermore, MIC information and differences in proportion of patients with PD target attainment were not evaluated. The median duration of study drug treatment was only 3.2 days in the study arm and 3.7 days in the intermittent arm. Furthermore, because the study did not evaluate serum concentrations, it is difficult to ascertain whether the lack of benefit was caused by the inability of continuous infusion to improve further on PD target attainment over aggressively dosed bolus administration of β-lactams. Although this study found evidence that continuous infusions of β-lactams may not be helpful for all patients with severe sepsis, it does not negate that such an administration technique has PK/PD benefits and may be of benefit in a more selective subpopulation.

Traditionally, TDM has been reserved for medications with narrow therapeutic indexes. Recently, TDM use has been suggested to ensure the optimization of the dosing of β-lactams in critically ill patients (Roberts 2014a). Several reports recently illustrated the feasibility of routine TDM use for dose optimization of β-lactams (Hites 2013, Patel 2012, Roberts 2014b). However, to date, no prospective studies have evaluated the clinical benefit associated with that practice. Furthermore, wide adoption of routine antimicrobial TDM is unlikely to occur until a quick turnaround and a reliable method for obtaining antimicrobial concentrations are available. Last, although dosing nomograms are available for certain antimicrobials (e.g., aminoglycosides, vancomycin), there are currently no standards for interpretation and dosing adjustments of other antimicrobials based on serum concentration. Until some of those barriers to TDM topple, it is unlikely that antimicrobial TDM will become routine at most institutions.

**Patient Care Scenario**

A 72-year-old man is admitted to a medical ICU with pneumonia and septic shock. Upon admission, he received 8 L of lactated Ringers and 1 L of 5% albumin. He has acute kidney injury requiring continuous renal replacement therapy and is currently receiving norepinephrine 45 mcg/minute, vasopressin 0.03 units/minute, cefepime, and levofloxacin. What physiologic changes are observed in this patient that could affect the pharmacokinetics and pharmacodynamics of the antimicrobials he is receiving?

**ANSWER**

This is a complicated patient with many physiologic changes that may affect antimicrobial PK and PD. Full ascertainment of the effects of the physiologic changes requires an assessment of the patient and the antimicrobial properties. Presumed changes, based on patient presentation, are larger intravascular volume from fluid resuscitation; decreased serum albumin concentration and increased α-acid glycoprotein, affecting protein binding; hemodynamic instability, decreasing perfusion and distribution of hydrophilic antimicrobials to intended site of action; and extracorporeal removal of drugs by way of continuous renal replacement therapy. The extent to which those physiologic changes would affect the PK and PD depends on the drugs’ properties: Cefepime is a time-dependent antimicrobial with a small volume of distribution and low protein binding and is eliminated through the kidneys. For cefepime, the patient’s physiologic changes are likely to lower serum concentration (increased volume of distribution), decrease distribution to active site of infection (hemodynamic instability), and decrease clearance (acute kidney injury). Regarding levofloxacin, this drug has a large volume of distribution, has low protein binding, and is also eliminated by the kidneys. Clearance is affected by acute kidney injury; however, serum concentration and delivery to site of infection should not be affected by the increase in intravascular volume. To effectively design a therapeutic regimen for this patient, a clinician must first weigh the potential effects of deranged PK and PD and then balance the safety and efficacy of potentially underdosing or overdosing the patient on any particular medication. An understanding of the antimicrobials’ PD targets and local MIC patterns would also assist in designing a regimen.

**PHARMACOGENOMICS**

Pharmacogenomics is the study of how genes affect a person’s response to drugs. The application of PGx information is an opportunity to further provide personalized medicine for critically ill patients. Currently, a large number of drugs have known pharmacogenomic variations in PK or PD. However, given the current structure of evidence-based medicine and FDA drug development regulations, the majority of drug dosing and monitoring remains a one-size-fits-all paradigm. Furthermore, the application of PGx to critically ill patients will necessitate the availability of a rapid genetic test panel that targets known genes that may affect critical care pharmacotherapy. Unfortunately, no such commercially available genetic test panel currently exists, and most PGx tests (unless available through in-house assays) have turnaround times of 5–7 days, thus precluding rapid bedside decisions. As such, application of the PGx tests and clinical scenarios discussed in the following rely on information from rapid genetic testing or incidental findings from previously administered PGx test panels. The discussion also pinpoints specific areas where further PGx testing development could enhance clinical application.

**Genetic Variations in CYP Metabolism**

The CYP enzymes are responsible for the hepatic metabolism of about 75% of all drugs (Guengerich 2008). Almost all CYP isoenzymes have genetic variations reported; however, a few enzymes have particular clinical applicability in critically ill patients. These include CYP2D6, CYP2C9, and CYP2C19, which are responsible for hepatic metabolism of 25%, 15%, and 10% of all drugs, respectively (Zhou 2009). Although involved in the hepatic metabolism of 50% of drugs, the CYP3A4 polymorphisms are less as common, have minimal clinical impact, and are not discussed here.

Phenotypic variations of CYP enzymes can be categorized into four categories: poor metabolizers, intermediate metabolizers, extensive metabolizers, and ultrarapid metabolizers (Samer 2013). Extensive metabolizers are generally considered wild-type representatives (typically most common) and have the usual amount of enzyme activity. Poor metabolizers result from either the presence of two nonfunctional (null) alleles or the deletion of the entire gene. Intermediate metabolizers are usually found in individuals carrying one null allele or one with reduced function. Ultrarapid metabolizers often carry one or more duplicate functional genes. Furthermore, the distribution of common variant alleles of CYP genes varies among different ethnic populations. Common phenotypes of the three most-studied isoenzyme genetic polymorphisms—with their corresponding frequencies in ethnic groups—and potentially affected drugs as seen in critically ill patients are listed in Table 1-4.

### Table 1-4. CYP Isoenzymes Commonly Seen in Critically Ill Patients

<table>
<thead>
<tr>
<th>CYP Isoenzyme</th>
<th>Common Metabolizer Types</th>
<th>Whites (%)</th>
<th>Africans (%)</th>
<th>Asians (%)</th>
<th>Affected Drugs</th>
</tr>
</thead>
<tbody>
<tr>
<td>CYP2D6</td>
<td>Poor</td>
<td>5–10</td>
<td>1–2</td>
<td>1–2</td>
<td>Antipsychotics, carvedilol, codeine, diltiazem, ethanol, hydrocortisone, metoprolol, oxycodone, propafenone, tramadol</td>
</tr>
<tr>
<td></td>
<td>Intermediate</td>
<td>10–15</td>
<td>50</td>
<td>30</td>
<td></td>
</tr>
<tr>
<td></td>
<td>Ultrarapid</td>
<td>1–5(^a)</td>
<td>0–2(^b)</td>
<td>2(^c)</td>
<td></td>
</tr>
<tr>
<td>CYP2C19</td>
<td>Poor</td>
<td>3–16</td>
<td>7–15</td>
<td>14–46</td>
<td>Clopidogrel, diazepam, phenobarbital, phenytoin, prasugrel, propranolol, proton pump inhibitors, valproic acid, voriconazole, warfarin</td>
</tr>
<tr>
<td></td>
<td>Ultrarapid</td>
<td>16–31</td>
<td>16–33</td>
<td>Rare</td>
<td></td>
</tr>
<tr>
<td>CYP2C9</td>
<td>Poor</td>
<td>35%</td>
<td>Rare</td>
<td>0–2</td>
<td>Amiodarone, clopidogrel, metronidazole, pantoprazole, sulfamethoxazole, valproic acid, voriconazole</td>
</tr>
</tbody>
</table>

\(^a\)Higher prevalence seen in southern European countries.

\(^b\)As high as 30% of Ethiopians.

\(^c\)As high as 30% of Saudi Arabians.

**Clinical Application of CYP Polymorphisms**

**Clopidogrel**

Clopidogrel is a thienopyridine prodrug that requires hepatic transformation to an active metabolite, which irreversibly inhibits the P2Y₁₂ receptor on the surface of blood platelets, causing decreased aggregation activity. Conversion of clopidogrel to its active metabolite requires two sequential oxidative steps involving several CYP enzymes (i.e., CYP1A2, CYP2B6, CYP2C9, CYP2C19, and CYP3A4). Patients with reduced CYP enzyme function may have decreased biotransformation of clopidogrel to its active metabolite.

Clopidogrel is often prescribed after percutaneous coronary intervention (PCI); however, patient response rates vary greatly, with up to 20% of patients being nonresponders or poor responders (Campos 2010). Several studies have linked CYP2C19 genetic polymorphisms with varying rates of clinical outcomes among post-PCI patients treated with clopidogrel (Collet 2009, Frere 2009, Giusti 2007, Mega 2009, Shuldiner 2009). Furthermore, meta-analysis has shown that patients treated with clopidogrel who are undergoing PCI and who are either homozygous (poor metabolizers) or heterozygous (intermediate metabolizers) with the CYP2C19 loss of functional allele(s) have an increased risk of major adverse cardiovascular events (Mega 2011). Furthermore, the relationship exhibits a stepwise increase in poor outcomes from intermediate metabolizers to poor metabolizers, which may further denote a physiologic or pharmacologic connection (HR for adverse cardiovascular event = 1.55; 95% CI, 1.11–2.17 for intermediate metabolizer; HR 1.76, 95% CI 1.24–2.50 for poor metabolizer).

Clinical Pharmacogenetics Implementation Consortium (CPIC) guidelines provide specific guidance in the use of clopidogrel in patients who are post-PCI and are intermediate or poor metabolizers of CYP2C19; for these patients, an alternative antiplatelet agent such as prasugrel or ticagrelor should be considered. Those recommendations may assist clinicians prior to the decision to initiate clopidogrel, but they may be of limited applicability for patients with confirmation of adequate antiplatelet activity, such as those with evidence of inhibition on platelet aggregation tests. Notably, the benefit of prasugrel and ticagrelor over clopidogrel in patients with post–acute coronary syndrome—with the majority undergoing PCI for the prevention of cardiovascular death, myocardial infarction, or stroke—was most pronounced in those with loss of functional CYP2C19 alleles (Mega 2009, Wallentin 2010). The American College of Cardiology guidelines on the management of ST-elevation myocardial infarction say genetic testing of clopidogrel metabolism may be considered on an individual patient basis (O’Gara 2014).

**Voriconazole**

The CYP2C19 polymorphisms may also have implications in the use of voriconazole for treatment of invasive fungal infections. Voriconazole is a triazole antifungal agent that is recommended as a first-line therapy for the treatment of acute invasive aspergillosis. It is metabolized primarily by hepatic enzyme CYP2C19, with some additional metabolism by CYP2C9 and CYP3A4. Voriconazole has a narrow therapeutic index and nonlinear pharmacokinetics, and it exhibits large, interpatient pharmacokinetic variability with a more than 100-fold difference in plasma concentrations observed with usual dosing (Meletiadis 2006). Those properties make it an ideal agent for the combination of pharmacogenetic evaluation and TDM.

Numerous studies have consistently found a relationship between voriconazole serum concentrations and CYP2C19 polymorphisms (Hassan 2011, Scholz 2009). In addition, the use of TDM to maintain voriconazole levels within therapeutic ranges has correlated with improvements in efficacy and adverse effects (Park 2012). However, to date, no studies have definitively proved that CYP2C19 polymorphisms are linked directly to differences in patient outcomes. That lack of proof may be due to increased use of TDM and dose adjustments, as well as the influence of other patient baseline characteristics on outcomes associated with invasive fungal infections. However, despite the lack of clinical evidence, the use of pharmacogenetic information to guide initial dosing of voriconazole seems prudent, particularly if the information is already available and would not delay therapy. That initial-dose guidance may be especially important for patients known to be ultrarapid metabolizers and to have active fungal infections. Subsequent dosing of voriconazole could be directed through TDM.

**Opioid Analgesics**

A number of opioid analgesics (e.g., codeine, hydrocodone, oxycodone, tramadol) rely on in vivo conversion to more active opioids for their analgesic properties. Codeine metabolism entails conversion to norcodeine by CYP3A4, glucuronidation conversion to codeine-6-glucuronide, and CYP2D6 conversion to morphine. Although the CYP2D6 pathway is responsible for only 5%–10% of codeine metabolism, CYP2D6 is responsible for the majority of analgesic effect because the parent drug has a 200-fold weaker affinity for the μ-opioid receptor compared with morphine.

The association between genetic polymorphisms of CYP2D6 leading to phenotypic differences in metabolizing activity and PK and PD differences with codeine and morphine is well documented. In CYP2D6 ultrarapid metabolizers, PK studies have documented increased conversion of codeine to morphine (Kirchheiner 2007); in some cases, this has resulted in toxic systemic concentrations of morphine.
and life-threatening adverse effects after normal doses of codeine (Gasche 2004). In contrast, poor metabolizers of CYP2D6 have been associated with decreased formation of morphine and lack of efficacy. CYP2D6 is also responsible for the biotransformation of tramadol, hydrocodone, and oxycodone to more-potent opioids. Although the clinical evidence surrounding PK changes associated with various CYP2D6 polymorphisms is not as robust as that for codeine, the CPIC guidelines provide similar therapeutic recommendations for all of these agents (Crews 2014). The guidelines recommend that alternative opioids that don’t rely on CYP2D6 conversion (e.g., morphine, hydromorphone) should be considered for patients who are either ultrarapid metabolizers or poor metabolizers of CYP2D6 (Crews 2014).

Other Polymorphisms Relevant to Critical Care

Human Leukocyte Antigen

Human leukocyte antigen B (HLA-B) is a gene that is used for encoding a cell surface protein for antigen presentation to the immune system. It is responsible for the presentation of a variety of peptides for immune recognition. To date, there are more than 2000 identified HLA-B alleles. Of particular interest to critical care practitioners, HLA-B*15:02 is related to the development of severe cutaneous adverse drug reactions (e.g., Stevens-Johnson syndrome, toxic epidermal necrolysis) as caused by any of several antiepileptics, including carbamazepine, phenytoin, and oxcarbamazepine. Unlike polymorphisms of CYP metabolizing enzymes, whose varying degrees of metabolizing activity depend on the presence or absence of one or both alleles, HLA-B*15:02 genotyping is considered positive if one or two copies of the allele are present; it is considered negative only when no copies are present. HLA-B*15:02 appears to have a distinctive ethnic and regional distribution, with the highest rates among East Asians and South or Central Asians. Rates of HLA-B*15:02 positivity have been as high as 36% in some areas of China. In general, areas of Hong Kong, Singapore, Thailand, and Malaysia have rates of 6%–12%. Among other Asian, Koreans and Japanese have relatively low rates of HLA-B*15:02 positivity, at 0.5% and 1%, respectively. Among patients with known ancestry associated with high probability of having an HLA-B*15:02 allele, the negative predictive value for severe cutaneous reactions in those treated with carbamazepine is 100% (Hung 2006). However, the positive predictive value is only 7.7%, suggesting that not all patients who possess the allele will develop severe reactions.

Currently, the FDA recommends that patients with ancestries in at-risk populations (East Asians and South or Central Asians) be screened for the presence of the HLA-B*15:02 allele before initiation of carbamazepine and phenytoin. Patients positive for HLA-B*15:02 should avoid initiation of relevant antiepileptics.

Leucyl/cystinyl Aminopeptidase (LNPEP) Variants

Currently, the surviving sepsis campaign has an ungraded recommendation suggesting that low-dose vasopressin may be added to norepinephrine for the hemodynamic support of patients in septic shock (Dellinger 2013). Vasopressin is currently administered as a fixed-dose infusion; therefore, it is understandable that patient variations may lead to differing serum concentrations.

One particular genetic variation associated with vasopressin therapy has significant correlations with outcomes among patients with septic shock. The clearance of vasopressin depends primarily on actions by the LNPEP enzyme. A single nucleotide polymorphism on the section of gene that encodes LNPEP generates genotypic differences and phenotypic variations in enzymatic activity. Using data from the Vasopressin and Septic Shock Trial and a validation cohort, Nakada and colleagues were able to determine that the TT genotype of a specific single nucleotide within the LNPEP gene was associated with increased enzymatic activity, leading to decreased vasopressin concentrations (Nakada 2011). The TT genotype, when compared with normal genotype (AA or AT), was also associated with increased 28-day mortality (51% vs. 34%, p=<0.001). Although the screening of this particular single-nucleotide polymorphism is not currently commercially available for clinical care and no studies support that a clinical decision based on this genetic variant would yield outcomes benefit, the trial by Nakada and colleagues demonstrates the potential vastness of the applicability of PGx.

CONCLUSION

The application of PK/PD/PGx principles is at a crossroad, and it is unlikely that a large-scale study would clearly demonstrate clinical outcome benefits associated with the application of those principles—as became evident by a study comparing continuous infusion β-lactams with intermittent infusions. This is most likely because the stringent procedure called for by a randomized controlled trial is contradictory to the patient-specific approach endorsed by the application of PK, PD, and PGx principles. As critical care pharmacists, we must decide whether each patient should receive personalized dosing and monitoring schemes based on a balance between the importance of the efficacy of a drug and the potential for adverse reactions associated with under- or overdosing of the medication. Application of the individualized PK, PD, and PGx principles may constitute deviation from a traditional, evidence-based therapeutic approach. However, as clinicians, we should not—based only on lack of evidence—negate the known physiologic and pathophysiologic changes seen in critically ill patients and the potential for dosing optimization.
Practice Points

In applying pharmacokinetic, pharmacodynamic, and pharmacogenomic principles to critically ill patients, clinicians should consider following these steps.

- Quantify the patient’s current clinical physiologic changes and note how each change could alter the pharmacokinetics of medications.
- For medications with narrow therapeutic indexes or ones for which meeting pharmacodynamic targets is crucial, specifically evaluate the potential effects of the patient’s physiologic change on serum and tissue concentrations.
- Derive a pharmaceutical plan that applies a dosing strategy that optimizes the probability of pharmacodynamic target attainment without substantial increase in the risk of adverse effects.
- If using a dosing scheme that differs from the package insert’s recommended doses, ensure proper documentation of the scheme rationale and vigilantly monitor for both therapeutic and adverse effects.
- Adjust dosing and monitoring schemes accordingly based on observed effects and changes in the patient’s physiology.
- The clinician should also realize that until the advent of real-time genetic testing, pharmacogenomics applications are currently restricted to incidental findings from previously evaluated patients,

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Self-Assessment Questions

Questions 1–6 pertain to the following case.

S.W. is a 54-year-old woman (weight 45 kg) admitted to the ICU with signs and symptoms of pneumonia and septic shock requiring intubation, aggressive fluid resuscitation, broad-spectrum antimicrobials, and hemodynamic support with norepinephrine. Her medical history is significant for seizures, for which she is taking phenytoin 100 mg orally three times daily. S.W. received a total of 7 L of crystalloid fluids in the first 24 hours after ICU admission. She is currently sedated with midazolam and fentanyl, and an orogastric tube was placed to initiate tube feeds. Her laboratory test results include WBC count 18.2 x 10³ cells/mm³, hemoglobin 8.0 g/dL, albumin 2.0 g/dL, lactate 2.2 mmol/L, SCr 0.8 mg/dL, and total phenytoin concentration 6.0 mcg/mL. Her physical examination reveals anasarca, and chest radiography demonstrates the presence of pulmonary edema. S.W.’s other current drugs, aside from norepinephrine, include pantoprazole 40 mg by mouth once daily, enoxaparin 30 mg subcutaneously every 12 hours, ceftriaxone 1 g intravenously every 24 hours, and azithromycin 500 mg intravenously every 24 hours.

1. Which one of the following physiologic changes is most likely to decrease S.W.’s absorption of oral pantoprazole?
   A. Increased α-1 glycoprotein concentrations
   B. Decreased albumin concentrations
   C. Decreased gastric emptying
   D. Increased splanchnic perfusion

2. Based on her clinical status and pharmacokinetic changes, which of the following recommendations is the most appropriate regarding S.W.’s current enoxaparin regimen?
   A. Decrease enoxaparin to 40 mg once daily.
   B. Continue current enoxaparin regimen.
   C. Check 4-hour peak anti-factor Xa level.
   D. Change to subcutaneous unfractionated heparin.

3. Which of the following best describes ceftriaxone’s volume of distribution in S.W.?
   A. Decreased because of aggressive volume resuscitation
   B. Increased because of mechanical ventilation
   C. Decreased because of hypoalbuminemia
   D. Increased because of hypoalbuminemia

4. Which of the following statements best describes the total serum concentration of azithromycin in S.W.?
   A. Unaffected from aggressive fluid resuscitation
   B. Decreased because of aggressive fluid resuscitation
   C. Increased because of decreased protein binding
   D. Decreased because of increased protein binding

5. During daily patient rounds, you calculate S.W.’s adjusted phenytoin concentration, based on the current albumin concentration, to be about 12 mcg/mL. Which one of the following properties of phenytoin would best explain why an adjustment is necessary for S.W.?
   A. High protein binding, weakly acidic
   B. Moderate protein binding, weakly basic
   C. High protein binding, weakly basic
   D. Moderate protein binding, weakly acidic

6. Upon verifying the corrected phenytoin concentration, the team orders a home dose of oral phenytoin 100 mg three times daily for S.W. You recommended changing the drug to IV fosphenytoin. Which one of the following reasons best justifies this change to S.W.’s regimen?
   A. Concerns with diminished absorption because of orogastric tube interactions
   B. Vasodilatory shock leading to increased hepatic blood flow and first-pass metabolism
   C. Hypoalbuminemia leading to increased volume of distribution
   D. Mechanical ventilation leading to decreased hepatic blood flow and first-pass metabolism

7. Your coronary care ICU is gathering information in preparation for the possible development of routine pharmacogenomics evaluations. The ICU director wants your input on deciding which pharmacogenomics tests should be employed for post-percutaneous coronary intervention patients. Which of the following tests is most likely to have clinical applicability in your particular unit?
   A. CYP3A4 polymorphism
   B. LNPEP gene variants
   C. CYP2D6 polymorphisms
   D. HLA-B screening

8. A Chinese patient with a prolonged history of epilepsy is admitted to your neuro ICU with active seizures. The patient was chronically managed on carbamazepine and phenytoin as an outpatient, with stable doses for the past 2 years. On admission, the patient had a total phenytoin level of 6.2 mcg/mL. The patient received intravenous lorazepam, and the seizures have currently subsided. One of the neurology fellows recently read an article regarding the increased propensity for severe cutaneous reactions among certain individuals receiving carbamazepine and phenytoin and raises this concern during rounds while discussing whether this patient’s home anti-epileptics should be continued. Which one of the following recommendations is best regarding this patient’s home anti-epileptics?
L.H. is currently intubated and hypotensive requiring nor-epinephrine 15 mcg/min despite receiving large amounts of crystalloid, colloids, and blood products. She has no significant medical history. Upon admission, L.H. was started on piperacillin/tazobactam 3.375 g intravenously every 6 hours administered as 30-minute bolus infusions. Her current urine output is 150 mL/hour. Her relevant laboratory results are WBC count 16.7 x 10^3 cells/mm, hemoglobin 9.0 g/dL, albumin 2.4 g/dL, lactate 1.2 mmol/L, and Scr 0.3 mg/dL.

11. Which of the following is most likely to cause a lower than usual piperacillin/tazobactam serum concentration in L.H.?
A. Hemodynamic instability
B. Augmented renal clearance
C. Hypoalbuminemia
D. Decreased volume of distribution

12. Which one of the following interventions would best assist piperacillin/tazobactam pharmacodynamics target attainment in L.H.?
A. Decrease infusion time from 30 min per dose to 10 min per dose.
B. Administer total daily dose as one large bolus infusion.
C. Change dosing regimen to 2.25 g intravenously every 4 hours.
D. Change dosing regimen to 4.5 g intravenously every 8 hours.

13. Which one of the following statements best describes L.H.’s piperacillin/tazobactam pharmacodynamic target?
A. Peak concentration should be at least 10 times higher than the pathogen’s MIC.
B. Minimum pharmacodynamic target is serum concentration being above the MIC for 50% of each dosing interval.
C. Prolonged time of serum concentration above MIC beyond 50% is unnecessary.
D. A higher AUC to MIC ratio than what is traditionally recommended is likely to improve bacterial killing.

14. You recommend administering piperacillin/tazobactam to L.H. in a fashion that maximizes the probability of pharmacodynamics target attainment. Which one of the following reasons best justifies this recommendation for L.H.?
A. Proven mortality benefit
B. Decreased ventilator days
C. Decreased adverse effects
D. Improved in vitro killing

15. In discussing L.H. with your PGY-2 critical care resident, you expressed concern regarding the increased volume of distribution with piperacillin/tazobactam and the corresponding decreased serum and tissue concentrations.

Questions 11–15 pertain to the following case.
L.H. is a 25 year-old woman (weight 65 kg) admitted to the surgical-trauma ICU after a motor vehicle crash. She suffered blood loss, open bone fractures, and traumatic brain injury.
Which one of the following alternative regimens will be least likely to be affected by the pharmacokinetic changes seen in L.H.?

A. Cefepime
B. Levofloxacin
C. Gentamicin plus vancomycin
D. Aztreonam plus clindamycin

16. For the same dose, which of the following factors is the most important contributor to the probability of attaining pharmacodynamic (peak/MIC > 8) targets for aminoglycosides?

A. Serum protein concentrations
B. Volume of distribution
C. Pathogen resistance
D. Renal clearance

17. Which one of the following dosing regimens is best to recommend for M.M.?

A. 50 mg once daily
B. 100 mg once daily
C. 130 mg once daily
D. 175 mg once daily

18. M.M. has been receiving the new oral antiviral agent through a naso-gastric feeding tube. On day 3 of therapy, M.M. develops hemodynamic instability. An ECG demonstrates new left-ventricle wall motion abnormality and a decreased ejection fraction from baseline. New laboratory results demonstrate an INR of 2.0 and total bilirubin of 4.5 mg/dL. Based on its pharmacokinetic profile, which one of the following changes in serum concentrations of the antiviral agent is most likely to happen as a result of M.M.’s hemodynamic instability and altered hepatic metabolism?

A. Decreased because of decreased hepatic blood flow
B. Increased because of decreased hepatic blood flow
C. Decreased because of decreased intrinsic hepatic clearance
D. Increased because of decreased intrinsic hepatic clearance

19. A 54-year-old Asian man (weight 70 kg) with acute myeloid leukemia is admitted to your medical ICU with neutropenic fever and respiratory failure requiring mechanical ventilation. The patient’s chest CT scan demonstrates cavitary lesions suggestive of invasive fungal infection, and serum galactomannan assay is positive. The medical team decides to treat the patient’s presumed invasive fungal infection with voriconazole. Your institution has a mechanism to obtain both CYP2C19 polymorphism and voriconazole serum concentration results by a send-out test. Which of the following recommendations is best for this patient?

A. Test for CYP2C19 polymorphisms to determine initial dose; adjust dose based on therapeutic drug monitoring.
B. Administer standard intravenous doses, adjust dose based on therapeutic drug monitoring.
C. Administer standard intravenous doses, and monitor for radiologic signs of efficacy.
D. Administer standard intravenous dose, adjust dose based on CYP2C19 polymorphism.

20. A 40-year-old woman (height 66 in, weight 56 kg) is admitted to your medical ICU with signs and symptoms of pyelonephritis and septic shock. The patient has a history of chronic UTIs and kidney stones requiring the placement of ureteral stents. The patient was profoundly dehydrated upon admission and required 8 L of crystalloid fluid in the first 24 hours to maintain her blood pressure. The patient was anuric upon admission, but upon fluid resuscitation and removal of ureteral stents her urine output is now 150 mL/hr. She was initially started on ciprofloxacin for her infection, but urine cultures now demonstrate the presence of an extended β-lactamase producing K. pneumoniae, which is sensitive to meropenem. Relevant laboratory results are WBC count 26.7 x 10^3 cells/mm, lactate 4.2 mmol/L, and Scr 2.3 mg/dL (baseline 0.5 mg/dL). Which one of the following meropenem regimens would be best to recommend for this patient?

A. 500 mg once daily
B. 500 mg every 12 hours
C. 500 mg every 8 hours
D. 1000 mg every 8 hours