

8	4	2	1	8	4	2	1	8	4	2	1	8	4	2	1	8	4	2	1	8	4	2	1
<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>

Marquer les cases d'un trait noir:

juste faux



Service de Pharmacologie clinique
Service des Maladies Infectieuses
Réception des laboratoires BH 18-100
Rue Bugnon 46, 1011 Lausanne
www.chuv.ch/pcl

Contact:
Laboratoire de Pharmacologie clinique
Tél.: 021 31 44 271
Fax: 021 31 48 098 – PP439
Interprétation clinique:
Tél.: 021 31 42 500
Heures d'ouverture: lundi – vendredi 08h–17h

Patient

Last name:

First name:

Gender:

Full birth date:

Address:

ZIP/City:

Your reference:

Invoice to:

Patient Applicant

Etiquette code barres pour le CHUV



Date and time of sampling (required)

Day:

1	2	3	4	5	6	7	8	9	10	11	12	13	14	15	
16	17	18	19	20	21	22	23	24	25	26	27	28	29	30	31

Month:

1	2	3	4	5	6	7	8	9	10	11	12
---	---	---	---	---	---	---	---	---	----	----	----

Hour:

0	1	2	3	4	5	6	7	8	9	10	11
12	13	14	15	16	17	18	19	20	21	22	23

 Min.:

15	30	45
----	----	----

COPY OF THE RESULTS TO:

Physician name:

Address:

ZIP/City:

DEMANDEUR

Physician name:

Address:

Phone: Fax:

Etiquette déviation
ou
Etiquette de projet

CLINICAL JUSTIFICATION(S):

- | | |
|--|---|
| <input type="checkbox"/> Therapeutic Drug Monitoring | <input type="checkbox"/> Unclear compliance |
| <input type="checkbox"/> Infection not responding | <input type="checkbox"/> Pregnancy |
| <input type="checkbox"/> Suspected toxicity | <input type="checkbox"/> Study: |
| <input type="checkbox"/> Suspected drug interactions | <input type="checkbox"/> Other reason: |
| | <input type="checkbox"/> Urgent request (specify reason): |

INDICATION OF TREATMENT:

- | | | |
|--|---|---|
| <input type="checkbox"/> Prophylaxis | <input type="checkbox"/> 1 st line treatment | <input type="checkbox"/> Empirical therapy for persistent fever |
| <input type="checkbox"/> Pre-emptive treatment | <input type="checkbox"/> Salvage therapy | <input type="checkbox"/> Asymptomatic infection (CMV) |
| <input type="checkbox"/> Documented infection | <input type="checkbox"/> Follow-up treatment | <input type="checkbox"/> Other (specify): |

Request for clinical interpretation of drug level measurement

yes (by default) no

CLINICAL DATA:

Organ dysfunction:	<input type="checkbox"/> heart	<input type="checkbox"/> liver	Immunosuppression:	<input type="checkbox"/> neutropenia
	<input type="checkbox"/> lung	<input type="checkbox"/> none		<input type="checkbox"/> long-term corticosteroids
	<input type="checkbox"/> other (specify):			<input type="checkbox"/> other (specify):
Graft:	<input type="checkbox"/> heart	<input type="checkbox"/> hematopoietic allograft	Renal function:	<input type="checkbox"/> renal insufficiency
	<input type="checkbox"/> liver	<input type="checkbox"/> hematopoietic autograft		serum creatinine: μmol/L
	<input type="checkbox"/> kidney	<input type="checkbox"/> other (specify):		<input type="checkbox"/> hemodialysis
	<input type="checkbox"/> lung			<input type="checkbox"/> continuous renal replacement therapy
				<input type="checkbox"/> other (specify):
Adverse events:	<input type="checkbox"/> none		Weight: kg	Height: cm
	<input type="checkbox"/> hepatic (specify):		For premature baby, gestational age:	
	<input type="checkbox"/> renal (specify):		Co-medications:	
	<input type="checkbox"/> neurologic (specify):			
	<input type="checkbox"/> other (specify):			

Important information:

Blood collection:

- Do not collect blood sample from the line used for the infusion of antimicrobial treatment.
- Blood specimens are usually drawn JUST BEFORE the next dose (=trough level) (exception: rifampicin: 2h post-dose) and at steady state conditions (cf. Recommendations: http://www.chuv.ch/pcl/pcl_home/pcl-prestations/pcl-prestations-tdm.htm)
- **Ship without delay** the blood specimens to your centralized hospital laboratory within a maximum of 30 minutes after sampling
- Processing of blood samples: centrifugation at 2000g for 10 minutes and 4°C, freezing of plasma at -80°C (or at least -20°C) within 1h after sampling. If this timing cannot be respected, keep the blood sample at 4°C until centrifugation and freezing (max: 6h after sampling).

LID

Norm
Prénom

Matériel adressé: K-EDTA whole blood (2.6 ml S-Monovette) sample received frozen
 (Also accepted: 2.6 ml serum tube (without gel) and 3 ml citrate tube; Neonatology: serum or EDTA-K 0.3 ml Microvette®)

ANTI-INFECTIOUS DOSE AGENT

- Routine TDM is currently recommended for the antimicrobials **voriconazole**, **cefepime** (underscored) (along with aminoglycosides and vancomycin).
- For the other antimicrobial agents in bold, **TDM interpretation is only recommended in special situations** (i.e. difficulties in dosing adaptation (i.e. renal replacement therapy, dialysis), drug interactions problems, toxicity, drug resistances, etc.)
- For all the other antimicrobial agents of the list, the TDM is not yet advised due to the current lack of clinical evidence supporting concentration measurements interpretation.

ANTIBIOTICS:

Carbapenems:

- Ertapenem
 Imipenem
 Meropenem

Cephalosporins:

- Cefazoline
 Cefepime
 Ceftaroline
 Ceftazidime
 Ceftobiprole
 Ceftolozane/tazobactam
 Ceftriaxone
 Cefuroxime

Penicillins:

- Amoxicillin
 Benzylpenicillin
 Flucloxacillin
 Piperacillin/tazobactam

Quinolones:

- Ciprofloxacin
 Levofloxacin

Tetracyclines:

- Doxycycline
 Tygecycline

Other antibiotics:

- Clindamycin
 Colistin/Colistimethate
 Daptomycin
 Fosfomycin
 Hydroxychloroquine
 Linezolid
 Rifampicin
 Trimethoprim/sulfamethoxazole

ANTIFUNGALS

Triazole derivates:

- Fluconazole
 Isavuconazole
 Itraconazole / hydroxy-itraconazole
 Posaconazole
 Voriconazole

Echinocandins:

- Anidulafungin
 Caspofungin

ANTIVIRALS:

- Aciclovir
 Ganciclovir
 Valaciclovir
 Valganciclovir

OTHER ANTI-INFECTIOUS AGENTS

INFORMATION ON LAST DOSE ADMINISTRATION (REQUIRED)

Exact date of last dose administration:

Day: 1 2 3 4 5 6 7 8 9 10 11 12 13 14 15 16 17 18 19 20 21 22 23 24 25 26 27 28 29 30 31
 Month: 1 2 3 4 5 6 7 8 9 10 11 12

Exact time of last dose administration:

Hour: 0 1 2 3 4 5 6 7 8 9 10 11 12 13 14 15 16 17 18 19 20 21 22 23
 Min.: 15 30 45

Date of onset of treatment or last change in drug dose:

Dosing schedule:

Dose (mg): 0.1 0.2 0.3 0.4 0.5 0.6 0.7 0.8 0.9 1 2 3 4 5 6 7 8 9 10 20 30 40 50 60 70 80 90
 100 200 300 400 500 600 700 800 900 1000 2000 3000 4000 5000 6000 7000 8000 9000
 Number of dose(s) /24h: 1 2 3 4 5 6

(example : for a 1500 mg dose, tick the box 1000 and the box 500)

Administration route: oral i.v. i.v. continuous infusion Perfusion time:

Other dosage:

ESSENTIAL INFORMATION FOR THE INTERPRETATION OF THE RESULTS

Bacterial infection:

- clinically documented (specify):
 microbiologically documented (specify):
 pathogen(s):

Fungal infection:

- Invasive aspergillosis
 Invasive candidiasis
 Other mycosis (specify):

Viral infection:

- Cytomegalovirus disease (CMV)
 Influenza *Viral subtype identified:*
 Post-exposure prophylaxis (Influenza)
 Other viral infection (specify):

Generalities:

Localization of infection (specify):

In vitro sensitivity of pathogen to the current treatment:

- susceptible intermediate resistant

Minimal inhibitory concentration (MIC) of the antibiotic/antifungal agent: mg/L

Inhibitory concentration 50 (IC₅₀) of the antiviral: mg/L

- Severity of infection: sepsis
 severe sepsis
 septic shock

Response of infection to the antibiotic/antifungal agent or to the antiviral drug: complete partial
 stable deterioration