

Malaria

PATIENT DEMOGRAPHICS						
Name (last, first):		Birth date: / / Age:				
Address (mailing):		Sex: □Male □Female □Unk				
Address (physical):		Ethnicity: Not Hispanic or Latino				
City/State/Zip:		☐Hispanic or Latino ☐Unk				
Phone (home): Phone (work/	Race:					
Alternate contact: □Parent/Guardian □Spouse □Othe		□Asian □Am. Ind/AK Native				
Name:	Phone:	□Native HI/Other PI □Unk				
INVESTIGATION SUMMARY		·				
Local Health Department (Jurisdiction):		Entered in WVEDSS? □Yes □No □Unk				
Investigation Start Date://		Case Classification:				
Earliest date reported to LHD://	☐ Confirmed ☐ Probable ☐ Suspect					
Earliest date reported to DIDE://		□ Not a case □ Unknown				
REPORT SOURCE/HEALTHCARE PROVIDER (HCP)						
Report Source: □Laboratory □Hospital □HCP □Public H	ealth Agency 🗆 Other					
Reporter Name:	D					
Primary HCP Name:	Primary HCP Phone:					
CLINICAL	rimary rice rimone.					
	sis date: / /	Recovery date: / /				
Symptoms and Clinical Findings	Clinical Risk Factors					
Y N U	Y N U					
☐ ☐ ☐ Fever (Highest measured temperature: °F)	□ □ Underlying medical co	ndition				
□ □ □ Chills		previous 12 months (if yes, indicate species below)				
□ □ Sweats	☐ Vivax ☐ Falciparum	n 🗆 Ovale 🗆 Malariae 🗆 Unknown				
□ □ □ Headache						
□ □ □ Myalgia	Hospitalization					
□ □ Nausea	YNU					
□ □ □ Vomiting	□ □ □ Patient hospitalized fo	or this illness				
□ □ □ Fatigue	If yes, hospital name:					
Confusion	Admit date://	Discharge date://				
□ □ □ Neurologic focal signs						
	Death					
Complications	YNU	YNU				
YNU	☐ ☐ Patient died due to this illness If yes, date of death://					
☐ ☐ ☐ Acute respiratory distress syndrome (ARDS)	TREATMENT					
□ □ □ Coma	YNU					
☐ ☐ ☐ Cerebral malaria	☐ ☐ Patient received therapy for this attack (If yes, indicate type below)					
□ □ □ Kidney failure	-	Tetracycline				
□□□Liver failure	-	Exchange transfusion				
		tovaquone/proguanil				
LABORATORY (Please submit copies of all labs, including CBC	•					
Y N U	s, associated with this limess to bibl	7				
□ □ □ Anemia						
□ □ Demonstration of <i>Plasmodium</i> species in blood films (parasitemia:%)						
□ □ Demonstration of <i>Plasmodium</i> species by molecular testing (e.g. PCR)						
□ □ Detection of <i>Plasmodium</i> species by RDT without confirmation by microscopy or molecular testing (symptomatic or asymptomatic)						
□□□ Specimen(s) sent to CDC for testing (□ Smear □ Who	ole blood ⊔ Other:)					
If the species of <i>Plasmodium</i> has been identified from <u>any</u> of the above test methods, please specify:						
☐ Vivax ☐ Falciparum ☐ Ovale ☐ Malariae ☐ Unable to identify ☐ Other species (specify:)						

INFECTION TIMELINE	NFECTION TIMELINE								
Instructions: Enter onset	I	Ехро	Exposure period		1	Onset date	* Incubation period for infection from		
date in grey box. Count backward to determine	Days from onset	-30] [transfusion may be up		
probable exposure period	, ,	(Max Incubatio	on) (Min Incubation)		, ¹ ,		to 2 months. Some <i>P.</i> vivax strains have		
	Calendar dates:	_/_/_	/	/	`\;'	//	protracted incubation		
EDIDEMIOLOGIC EVDOS	LIDEC /bosed on the obs		novice wol		i	*~~I\	(8 to 10 months).		
Y N U	URES (based on the abo	ove exposure	perioa, unie	ess otne	rwise no	tea)			
☐ ☐ History of travel dur	ring exposure period (if yes, o	complete travel	history below):						
							1		
Destination (City,	County, State and Country)		Arrival Date	Departu	ire Date	Reason for travel			
				1					
YNU									
☐ ☐ Patient traveled (or the past 2 years	lived) outside of United State	s during							
☐ ☐ Patient resided in U	nited States prior to most rec	ent travel							
If no, please specify		ciit ti avei							
☐ ☐ Foreign arrival (e.g.									
If yes, country:									
□ □ Blood transfusion re	·	3							
If yes, date:/ _									
☐ ☐ Organ transplant red									
Where did exposure most like		State	: Co	untry:					
PUBLIC HEALTH ISSUES			PUBLIC HEA	LTH ACT	IONS				
YNU			YNU						
☐ ☐ Malaria chemoprop☐ ☐ Chloroquine ☐						k or other facility who	ere organs donated		
	Atovaquone/proguanil					vention information	provided to patient		
□ □ □ All chemoprophylax		ribed	and/or family/guardian						
	ed or not taken, please specify Didn't think needed	reason: e effects			•	ng of other symptoma	ntic persons who have		
•	Prematurely stopped taking once			red exposunt is lost to)			
□Unknown □	Other:		□ □ □ Other						
□ □ Case donated blood									
In the 30 days p Date://	orior to symptom onset								
Agency/location	n:								
Type of donation	on:								
☐ ☐ ☐ Case is pregnant (Du☐ ☐ ☐ Case knows someon		and is							
currently having sim		allu is							
☐ ☐ Epi link to another o	confirmed case of same condi	tion							
☐ ☐ Case is part of an ou	ıtbreak								
□ □ □ Other: WVEDSS									
Y N U									
□ □ □ Entered into WVEDS	SS (Entry date ://) Case S	Status: 🗆 Con	firmed \square	l Probable	☐ Suspect ☐ Not a	case 🛘 Unknown		
NOTES									