

## CITY AND COUNTY OF SAN FRANCISCO PUBLIC HEALTH LABORATORY

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## **Test Order**

## Malaria Molecular Identification (Malaria Real-Time PCR)

Synonym(s)	Plasmodium for Identification, Plasmodium Species Identification
Methodology	Real-time PCR (for <i>Plasmodium</i> /malaria identification)
Acceptable Specimen	Whole blood with EDTA. Also required: thick and thin blood smears
Type(s) for Testing	(stained or unstained* with Giemsa stain) on glass microscopy slides
	per CCR, Title 17, Section 2505. *Note: for unstained slides, thin smears
	must be methanol-fixed prior to submission.
Transport / Collection	Collect a 1-5 ml blood sample in Vacutainer® EDTA tubes prior to anti-
Medium	parasitic therapy, shipped on cold packs (not frozen); smears in slide
	transport container.
Storage and	Store whole blood refrigerated (2-8°C) until transport, and submit sample
Preservation of	within 5 days of collection; smears can be refrigerated (2-8°C) or stored at
Specimen	ambient temperature.
Minimum Volume	500 μl whole blood for malaria PCR
Required	NI/A
Additional Collection Instructions	N/A
Additional Required	Patient travel history, patient clinical history, physician's full name and
Information	contact number, onset of symptoms, county of residence.
Send Out?	No
Turnaround Time	1-3 business days
Testing Restrictions	1 0 basinoss days
Requisition Form(s)	http://sfcdcp.org/document.html?id=1036
Limitations / Notes /	This PCR test is to be used in conjunction with microscopic examination of
Disclaimers	blood smears, as it does not determine the level of malarial parasitemia. In
Biodiaiiioio	addition, this PCR test does not test for the presence of <i>Plasmodium knowlesi</i>
	or <i>Babesia</i> species DNA. Whole blood negative for <i>Plasmodium</i> species on
	this assay does not rule out the possibility of infection with <i>Plasmodium</i>
	knowlesi or Babesia species. Thick and thin smear examination will be used
	in conjunction with this assay to detect the presence of other blood parasites.
	This PCR test was developed and its performance characteristics were
	determined by the San Francisco Public Health Laboratory. It has not been
	cleared or approved by the U.S. Food and Drug Administration. The FDA has determined that such approval is not necessary and results should not be
	regarded as investigational or for research.
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