

Tick-borne Disease Diagnostic Reference

When to Consider Tick-Borne Diseases (TBDs)

- TBD signs and symptoms may be nonspecific and include fever, headache, myalgia and gastrointestinal manifestations. Rash is not associated with all TBDs and may not be an early indicator.
- New Yorkers can be exposed to TBDs throughout all of NYS.
- Diagnostic testing can help guide clinical management, but do not delay therapy if a TBD is suspected. Prompt treatment can prevent severe disease.
- Coinfection is uncommon but more likely associated with TBDs caused by blacklegged ticks.
- Ticks emerge when snow melts and stay active until temperatures fall below freezing.
- Ticks found crawling and unattached on skin are not considered a risk for TBD transmission.

Ticks Found in NYS



Blacklegged ticks: Lyme disease, babesiosis, anaplasmosis, Powassan virus and *Borrelia miyamotoi*



Lone star ticks: Ehrlichiosis, Heartland and Bourbon viruses, and tularemia



American dog ticks: Tularemia and Rocky Mountain Spotted Fever (*Rickettsia rickettsii*)



Woodchuck ticks: Powassan virus



Asian longhorned ticks: Not been found to carry pathogens that cause TBDs in the U.S.

TBDs to Consider

	Lyme Disease	Babesiosis	Anaplasmosis and Ehrlichiosis	RMSF	Powassan, Heartland and Bourbon Viruses, <i>B.miyamotoi</i> , and Tularemia
Manifestations	<ul style="list-style-type: none"> • EM • Cranial neuritis (usually facial palsy) • Acute oligoarthritis • Carditis (usually atrioventricular block) 	<ul style="list-style-type: none"> • Hemolytic anemia • Thrombocytopenia • Illness is more severe if asplenic, immunocompromised or an older adult 	<ul style="list-style-type: none"> • Thrombocytopenia • Leukopenia • Anemia • Mildly to moderately elevated hepatic transaminases <p>Some rickettsial diseases, especially RMSF, can be life-threatening if untreated</p>		<ul style="list-style-type: none"> • Powassan meningitis or encephalitis • Travel to the Midwestern U.S. (Bourbon virus)
Rash or Eschar	<ul style="list-style-type: none"> • EM and occasional multiple secondary annular rashes 	<ul style="list-style-type: none"> • Not applicable 	<ul style="list-style-type: none"> • Rare in anaplasmosis • Uncommon in ehrlichiosis in adults and typically maculopapular. Occurs in up to 60% of pediatric cases 	<ul style="list-style-type: none"> • Maculopapular (initially on wrists, forearms and ankles, then trunk and sometimes palms and soles, followed by a petechial rash) • Less than 50% of patients have a rash in the first three days of illness 	<ul style="list-style-type: none"> • Rash is uncommon (<i>B. miyamotoi</i>)
Testing	<ul style="list-style-type: none"> • EM alone is diagnostic and should be treated empirically • A two-tiered serologic test (most sensitive two weeks after illness onset) where the initial EIA or IFA is positive or equivocal, followed by a western blot or other FDA-cleared EIA test that is also positive or equivocal • If symptoms began more than 30 days from the test date, use IgG serologic results only. Disregard IgM results, even if positive • PCR is typically insensitive for most specimens 	<ul style="list-style-type: none"> • PCR • Intraerythrocytic <i>Babesia</i> parasite on blood smear • Serology (IFA for IgG offers evidence of infection but cannot distinguish between active and prior infection) 	<ul style="list-style-type: none"> • PCR on whole blood (most sensitive during the first week of illness) • Serology • A negative acute test does not rule out infection <p>• Confirm a serologic diagnosis with a demonstration of a fourfold rise in IgG titers by IFA in serum samples collected two to four weeks apart. Single antibody results cannot be independently relied on for confirmation</p> <p>• IgM antibodies are less specific than IgG antibodies and more likely to generate false positives</p> <p>• IgM results alone should not be used for lab diagnoses. Antibody titers are frequently negative in the first seven to ten days of illness</p>	<ul style="list-style-type: none"> • Serology • PCR on whole blood (less sensitive in early stages of disease) • A negative acute test does not rule out infection • PCR of skin biopsy of rash available for detection of rickettsial DNA 	<ul style="list-style-type: none"> • Testing for rare or emerging TBDs, particularly viral diseases, may not be available at commercial labs. For assistance, please contact the local health department where the patient resides • <i>B. miyamotoi</i> testing is available at several commercial diagnostic labs • RMSF antibody tests often cross-react with <i>R. akari</i> and <i>R. parkeri</i>

Treatment	<ul style="list-style-type: none"> • For post-exposure prophylaxis and TBD treatment guidance, visit cdc.gov/ticks/tickbornediseases • Consult with an infectious disease specialist • Antibodies may normally persist in the blood for months or years after infection, so testing cannot be used to determine a cure
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EIA: Enzyme immunoassay
IgG: Immunoglobulin G

EM: Erythema migrans
IgM: Immunoglobulin M

FDA: Food and Drug Administration
PCR: Polymerase chain reaction

IFA: Immunofluorescence assay
RMSF: Rocky Mountain spotted fever

For more information, visit health.ny.gov/tickfree

Images on this poster were adapted from the Centers for Disease Control and Prevention.

