



**Delaware Division of Public Health
Bureau of Epidemiology**

Lyme Disease (LD) Case Report Form

PATIENT INFORMATION

Last Name:		First Name:		Telephone #:
Street Address:			Zip Code:	County: <input type="checkbox"/> New Castle <input type="checkbox"/> Kent <input type="checkbox"/> Sussex
Gender: <input type="checkbox"/> Male <input type="checkbox"/> Female	Date of birth:	Hispanic Ethnicity: <input type="checkbox"/> Yes <input type="checkbox"/> No <input type="checkbox"/> Unknown	Race: <input type="checkbox"/> American Indian or Alaskan Native <input type="checkbox"/> Asian <input type="checkbox"/> Black or African American <input type="checkbox"/> Other <input type="checkbox"/> Native Hawaiian or Pacific Islander <input type="checkbox"/> Unknown <input type="checkbox"/> White	

PHYSICIAN / PROVIDER INFORMATION

Physician:	Address:	Phone:
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LABORATORY RESULTS

EIA/IFA (IgM, IgG, Total) <input type="checkbox"/> Positive <input type="checkbox"/> Negative <input type="checkbox"/> Equivocal <input type="checkbox"/> Not done <input type="checkbox"/> Titer _____		Other tests (check all that apply) <input type="checkbox"/> <i>B. burgdorferi</i> culture positive? <input type="checkbox"/> CSF titer higher than serum titer*? <input type="checkbox"/> Other positive result? (PCR, etc.): Specify: _____
Specimen Source: <input type="checkbox"/> Serum <input type="checkbox"/> CSF <input type="checkbox"/> Synovial Fluid <input type="checkbox"/> Other		
Western Blot (WB) <i>Please indicate positive WB bands, if known. For IgM, 2 of 3 bands must be positive For IgG, 5 of 10 bands must be positive</i>		
IgM: <input type="checkbox"/> Positive <input type="checkbox"/> Negative <input type="checkbox"/> Not done <input type="checkbox"/> 41kDa (FlaB) <input type="checkbox"/> 39 kDa (BmpA) <input type="checkbox"/> 21-25 kDa (OspC)		
IgG: <input type="checkbox"/> Positive <input type="checkbox"/> Negative <input type="checkbox"/> Not done <input type="checkbox"/> 93 kDa <input type="checkbox"/> 66 kDa <input type="checkbox"/> 58 kDa <input type="checkbox"/> 45 kDa <input type="checkbox"/> 41 kDa <input type="checkbox"/> 39 kDa <input type="checkbox"/> 30 kDa <input type="checkbox"/> 28 kDa <input type="checkbox"/> 21 kDa <input type="checkbox"/> 18 kDa		

CLINICAL

Did you diagnose the patient with LD? <input type="checkbox"/> Yes <input type="checkbox"/> No Date of LD diagnosis: _____ Date of symptom onset: _____																																																					
Case definition signs and symptoms <table border="1" style="width:100%; border-collapse: collapse;"> <thead> <tr> <th></th> <th>Yes</th> <th>No</th> <th>Unknown</th> </tr> </thead> <tbody> <tr><td>Erythema Migrans (EM) rash</td><td><input type="checkbox"/></td><td><input type="checkbox"/></td><td><input type="checkbox"/></td></tr> <tr><td>Arthritis (<i>objective episodes of joint swelling</i>)</td><td><input type="checkbox"/></td><td><input type="checkbox"/></td><td><input type="checkbox"/></td></tr> <tr><td>Bells palsy or other cranial neuritis</td><td><input type="checkbox"/></td><td><input type="checkbox"/></td><td><input type="checkbox"/></td></tr> <tr><td>Radiculoneuropathy</td><td><input type="checkbox"/></td><td><input type="checkbox"/></td><td><input type="checkbox"/></td></tr> <tr><td>Lymphocytic meningitis</td><td><input type="checkbox"/></td><td><input type="checkbox"/></td><td><input type="checkbox"/></td></tr> <tr><td>Encephalomyelitis*</td><td><input type="checkbox"/></td><td><input type="checkbox"/></td><td><input type="checkbox"/></td></tr> <tr><td><i>*If encephalomyelitis is checked, CSF titer must be higher than serum titer</i></td><td><input type="checkbox"/></td><td><input type="checkbox"/></td><td><input type="checkbox"/></td></tr> <tr><td>2nd or 3rd degree atrioventricular block</td><td><input type="checkbox"/></td><td><input type="checkbox"/></td><td><input type="checkbox"/></td></tr> </tbody> </table>		Yes	No	Unknown	Erythema Migrans (EM) rash	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	Arthritis (<i>objective episodes of joint swelling</i>)	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	Bells palsy or other cranial neuritis	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	Radiculoneuropathy	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	Lymphocytic meningitis	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	Encephalomyelitis*	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<i>*If encephalomyelitis is checked, CSF titer must be higher than serum titer</i>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	2 nd or 3 rd degree atrioventricular block	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	Other signs and symptoms (check all that apply): <table border="1" style="width:100%; border-collapse: collapse;"> <tbody> <tr><td><input type="checkbox"/> Arthralgias</td><td><input type="checkbox"/> Myocarditis</td></tr> <tr><td><input type="checkbox"/> Bundle branch block</td><td><input type="checkbox"/> Neck pain</td></tr> <tr><td><input type="checkbox"/> Cognitive impairment</td><td><input type="checkbox"/> Other rash</td></tr> <tr><td><input type="checkbox"/> Encephalopathy</td><td><input type="checkbox"/> Palpitations</td></tr> <tr><td><input type="checkbox"/> Fatigue</td><td><input type="checkbox"/> Paresthesias</td></tr> <tr><td><input type="checkbox"/> Fever/Sweats/Chills</td><td><input type="checkbox"/> Peripheral neuropathy</td></tr> <tr><td><input type="checkbox"/> Headache</td><td><input type="checkbox"/> Visual/auditory impairment</td></tr> <tr><td><input type="checkbox"/> Myalgias</td><td><input type="checkbox"/> Symptom(s) not listed</td></tr> </tbody> </table>	<input type="checkbox"/> Arthralgias	<input type="checkbox"/> Myocarditis	<input type="checkbox"/> Bundle branch block	<input type="checkbox"/> Neck pain	<input type="checkbox"/> Cognitive impairment	<input type="checkbox"/> Other rash	<input type="checkbox"/> Encephalopathy	<input type="checkbox"/> Palpitations	<input type="checkbox"/> Fatigue	<input type="checkbox"/> Paresthesias	<input type="checkbox"/> Fever/Sweats/Chills	<input type="checkbox"/> Peripheral neuropathy	<input type="checkbox"/> Headache	<input type="checkbox"/> Visual/auditory impairment	<input type="checkbox"/> Myalgias	<input type="checkbox"/> Symptom(s) not listed
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SUPPLEMENTAL INFORMATION

Was the patient pregnant at the time of illness?	<input type="checkbox"/> Yes <input type="checkbox"/> No <input type="checkbox"/> Unknown <input type="checkbox"/> NA
Was the patient hospitalized for this illness?	<input type="checkbox"/> Yes <input type="checkbox"/> No <input type="checkbox"/> Unknown
Antibiotic(s) used for this illness:	_____
Antibiotic usage (days prescribed):	_____

-----FOR PUBLIC HEALTH SURVEILLANCE USE ONLY-----

CASE STATUS: <input type="checkbox"/> Confirmed: <input type="checkbox"/> Unconfirmed: <input type="checkbox"/> Not a Case:	DERSS ID#:	COMMENTS:
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Lyme Disease Surveillance Case Definition

Clinical Description:

A systemic, tick-borne disease with protein manifestations, including dermatologic, rheumatologic, neurologic, and cardiac abnormalities. The best clinical marker for the disease is the initial skin lesion, erythema migrans, which occurs among 60%-80% of patients.

Clinical Case Definition:

- Erythema migrans, OR
- At least one late manifestation, as defined below and laboratory confirmation of infection.

Laboratory Confirmation - For the purposes of surveillance, the definition of a qualified laboratory assay is:

- A positive culture for *B. Burgdorferi*
- Single-tier IgG immunoblot seropositivity
- A two-tiered approach using a sensitive enzyme immunoassay or immunofluorescence antibody followed by Western blot.

Case Classification – Confirmed: a case that meets one of the clinical case definitions above.

Definition of terms used in the clinical description and case definition:

A. Erythema migrans (EM)

- For purpose of surveillance, EM is defined as a skin lesion that typically begins as a red macule or papule and expands over a period of days to weeks to form a large round lesion, often with partial central clearing. A solitary lesion must reach at least 5cm in size. Secondary lesions may also occur. Annular Erythematous lesions occurring within several hours of tick bite represent hypersensitivity reactions and do not qualify as EM. For most patients, the expanding EM lesion is accompanied by other acute symptoms, particularly fatigue, fever, headache, mild stiff neck, arthralgia, and myalgia. These symptoms are typically intermittent. The diagnosis of EM must be made by a physician. Laboratory confirmation is recommended for persons with no known exposure.

B. Late manifestations – Include any of the following when an alternate explanation is not found:

- **Musculoskeletal system:** Recurrent, brief attacks (weeks or months) of objective joint swelling in one or few joints, sometimes followed by chronic arthritis in one or few joints. Manifestations not considered as criteria for diagnosis include chronic progressive arthritis not preceded by brief attacks and chronic symmetrical Polyarthritis. Additionally, arthralgia, myalgia, or fibromyalgia syndromes alone are not criteria for musculoskeletal involvement.
- **Nervous system:** (Any one the following alone, or in combination): lymphocytic meningitis; cranial neuritis, particularly facial palsy (may be bilateral); radiculoneuropathy; or rarely encephalomyelitis, which must be confirmed by showing antibody production against *B. Burgdorferi* in the cerebrospinal fluid (CSF), demonstrated by a higher titer of antibody in CSF than in serum. Headache, fatigue, paresthesias, or mild stiff neck alone, are not criteria for neurologic involvement.
- **Cardiovascular system:** Acute onset, high grade (2nd or 3rd degree) artioventricular conduction defects that resolve in days to weeks and are sometimes associated with myocarditis. Palpitations, bradycardia, bundle branch block, or myocarditis alone are not criteria for cardiovascular involvement.

C. Laboratory confirmation – As noted above, laboratory confirmation of infection with *B. Burgdorferi* is established when a laboratory isolates the spirochete from tissue or sterile body fluid or detects diagnostic levels of IgM or IgG antibodies to the spirochete in serum, CSF, or other sterile body fluid (i.e., synovial fluid). States may determine the criteria for laboratory confirmation and diagnostic levels of antibody. Syphilis and other known causes of biologic false-positive serologic test results should be excluded when laboratory confirmation has been based on serologic testing alone.

Note: *This surveillance case definition was developed for national reporting of Lyme disease; it is not intended to be used in clinical diagnosis.*

Source: Centers for Disease Control and Prevention (CDC)

For questions related to this form, please call the Bureau of Epidemiology at 1-888-295-5156

This form is available on our webpage: <http://www.dhss.delaware.gov/dhss/dph/epi/lyme.html>

Reviewed: 1/2008