

Lyme Disease, Human Granulocytic Anaplasmosis & Babesiosis

Thomas Novicki Ph.D. D(ABMM)
Marshfield Labs (A division of Marshfield Clinic)
Marshfield WI



Major Tick-Borne Diseases of the USA and Their Tick Vectors

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|--|--|
| <input type="checkbox"/> Lyme Disease (LD) | <input type="checkbox"/> <i>Ixodes</i> |
| <input type="checkbox"/> Human Granulocytic Anaplasmosis (HGA) | <input type="checkbox"/> <i>Dermacentor</i> |
| <input type="checkbox"/> Tularemia | <input type="checkbox"/> <i>Amblyomma</i> |
| <input type="checkbox"/> Ehrlichiosis | <input type="checkbox"/> <i>Ornithodoros</i> |
| <input type="checkbox"/> Relapsing Fever | |
| <input type="checkbox"/> Rocky Mtn Spotted Fever | |
| <input type="checkbox"/> Colorado Tick Fever | |
| <input type="checkbox"/> Babesiosis | |
| <input type="checkbox"/> Tick Paralysis | |



Tick Borne Diseases of the Upper Midwest

- ☐ LD
 - ☐ *Borrelia burgdorferi* sensu stricto
- ☐ HGA
 - ☐ *Anaplasma phagocytophilum*
- ☐ Babesiosis
 - ☐ *Babesia microti*
- ☐ Tick paralysis
 - ☐ Tick neurotoxins
- ☐ Tularemia
 - ☐ *Francisella tularensis*



The Vector - *Ixodes scapularis*

- Primary Hosts
- ☐ Larva: white footed mouse, other small mammals
 - ☐ Nymph: small rodents, humans *i.e. The principle vector for human LD*
 - ☐ Adult: white tail deer, and sometimes humans
 - ☐ Wild animals remain asymptomatic



Images courtesy of CDC



Epidemiology

- ☐ LD, HGA, Babesiosis are zoonotic diseases
 - ☐ Cycle between small and large mammal populations
 - ☐ Birds, reptiles also play roles
- ☐ Humans are effectively incidental, dead end hosts



I. *Scapularis* - Feeding

- ☐ Female is the predominant feeder, source of disease
- ☐ Tick remains attached 3-7 days if not disrupted
- ☐ Main blood meal, "the big sip" occurs in final 4hrs
- ☐ Transmission occurs



The Etiological Agents and their Diseases

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LD

- *Borrelia burgdorferi*, a spiral bacterium related to *Treponema* (syphilis, yaws, pinta)
- Three species of *B. burgdorferi* sensu lato
 - *B. burgdorferi* sensu stricto
 - *B. afzelii*
 - *B. garinii*
- Geographic differences
 - N. America: *B. burgdorferi* sensu stricto only
 - Europe: all three species
 - Disease spectrum in Europe differs

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LD

- Early, Local (days-weeks post tick bite)
 - Primary erythema migrans (EM) at site of bite
 - Papule, expanding in annular rings
 - 80-90% of patients exhibit EM
 - Early signs of dissemination may also occur
 - Fever
 - Malaise/myalgia
 - Headache/stiff neck
 - Migratory arthralgias
 - Local lymphadenopathy

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1° EM

- Classic EM form, but may be more diffuse, less annular
- Has central *punctum*, site of tick bite
- And, 10-20% have no EM lesion
- Bottom Line: not always easy to diagnose!

Image courtesy of CDC



LD

- Early, Disseminated (weeks-months post bite)
 - Multiple 2° EM lesions (no punctum)
 - Lyme carditis
 - Neuroborreliosis
 - Meningitis
 - Cranial neuritis
 - Myelitis
 - Encephalitis
 - Lymphocytoma (cutaneous B-cell pseudolymphoma),
acrodermatitis chronica atrophicans
 - Primarily seen in Europe, is rare in here

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LD

- Late, Disseminated (months-years post bite)
 - Migrating arthritis, esp. knees
 - Various chronic neuropathies

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“Chronic” LD

- Is there disease beyond late LD?
 - Post-Lyme Disease Syndrome
 - Small proportion of patients living in endemic areas, who are diagnosed by validated lab methods, and complete approved treatment continue to show some residual symptoms
 - Symptoms usually mild, abate over time
 - Immune-related?



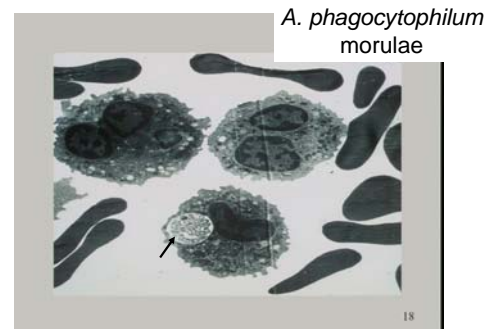
Chronic LD

- Many other cases do not fit these criteria - “Chronic LD”
 - No lab evidence of infection, or “evidence” by poorly validated methods
 - Live outside of endemic areas
 - No Hx of tick bite, EM
 - Vague symptoms - fatigue, aches, night sweats, etc
 - Can result in much money spent, questionable treatments



HGA

- *Anaplasma (Ehrlichia) phagocytophilum*
- Order *Rickettsiales*
- 1st described in 1994 (Chen et al JCM 32:589)
 - 6 pts. MN & WI
 - 33% mortality
 - Granulocytes of one pt. had cytoplasmic inclusions reminiscent of *Ehrlichia chaffeensis* **monocytic** inclusions (morulae)



Photos courtesy of Jim Kazmierczak DVM
WI DPH

HGA

- 16S rRNA sequence analysis:
 - > 99.8% homologous to the animal pathogens *E. phagocytophila* and *E. equi*
 - only 92.5% homologous to *E. chaffeensis*
- “Agent of human granulocytic ehrlichiosis”
- After several reclassifications, now known as “*Anaplasma phagocytophilum*”



HGA

- Common Symptoms
 - Fever
 - Headache
 - Malaise/myalgia
 - Thrombocytopenia, neutropenia
 - ↑ hepatic transaminases
 - Rash is rare: compare to Rocky Mountain Spotted Fever (*Rickettsia rickettsii*) where rash is common
- Usually self-limited, but fatalities occur (<1%)



Babesiosis

- Approximately 100 species of *Babesia*
- Human agents
 - Upper Midwest, Northeast USA: *B. microti*
 - West Coast: CA1, WA1
 - Europe: *B. divergens*



Babesiosis

- Unicellular protozoan that parasitizes RBCs
- Distant relative of *Plasmodium* (malaria)
 - *Babesia* differs from *Plasmodium*:
 - Vector = tick, not mosquito
 - No hepatic forms
 - No schizonts
 - Has extracellular forms



Babesiosis

- 1st human case reported Nantucket RI 1969
- Similarities to *Plasmodium* extend to symptoms:
 - Relapsing fever
 - Hemolytic anemia/jaundice
 - Fatigue
 - Chills, sweats
 - Headache
 - Myalgia/arthritis
 - Anorexia



Babesiosis

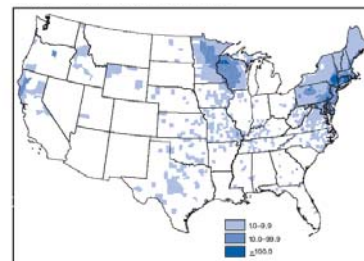
- Most cases asymptomatic
- Disease ranges from mild to fulminant (rare)
- Immunosuppression, advanced age, asplenia are predisposing factors
- Mortality $\leq 5\%$ if untreated



Epidemiology

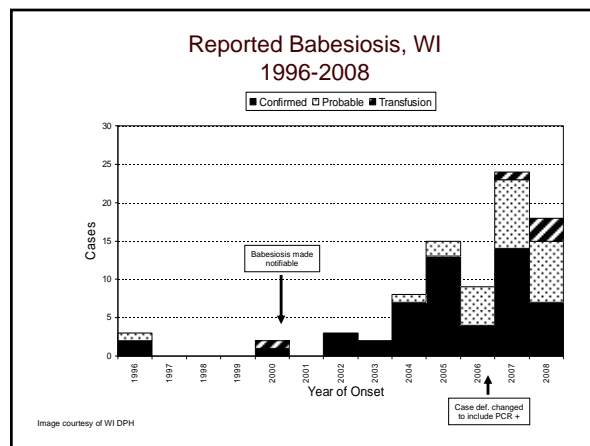
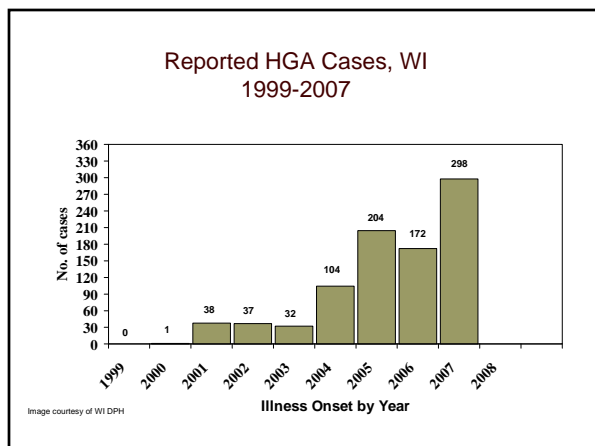
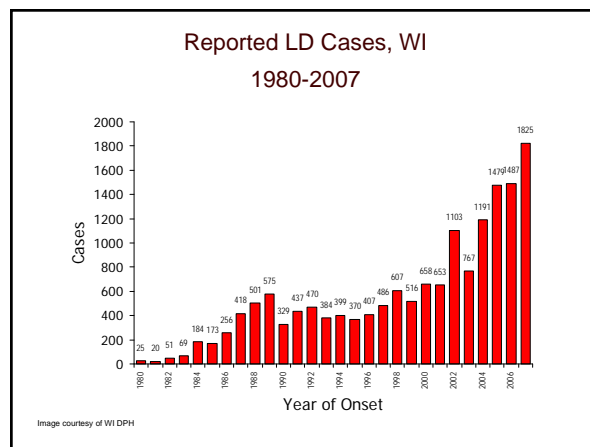
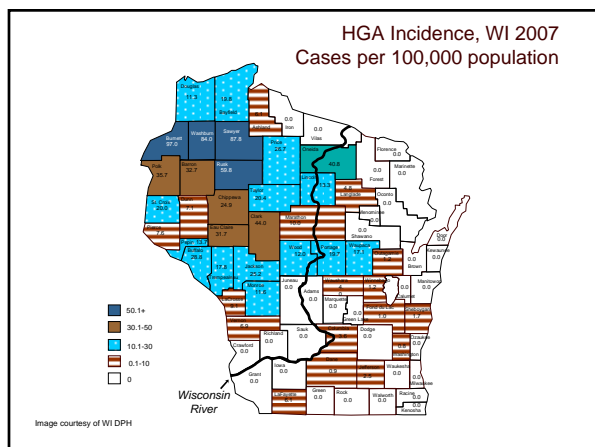
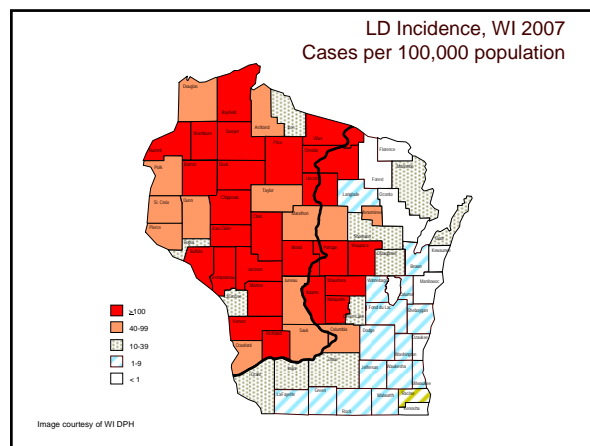
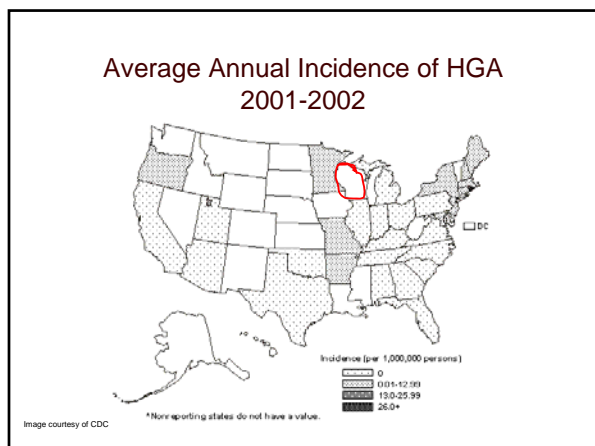


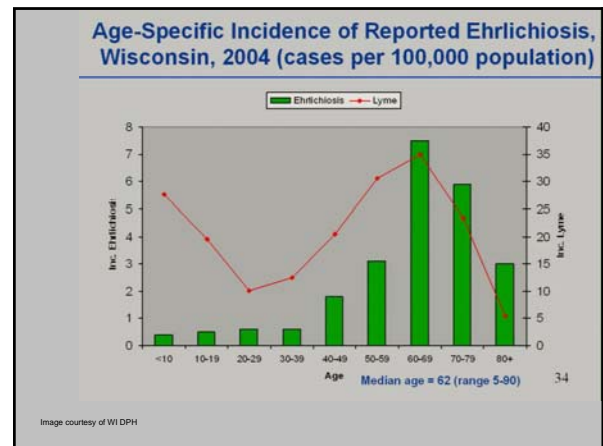
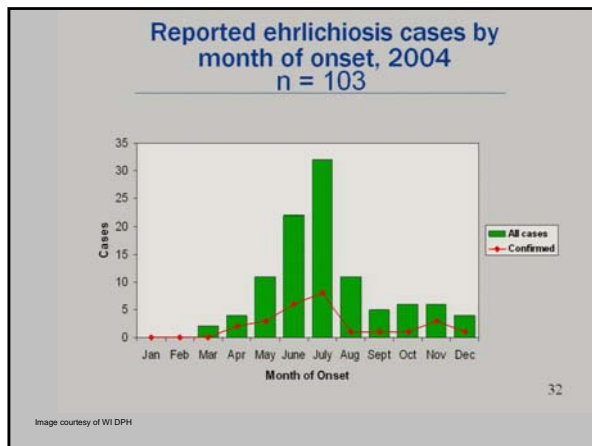
Average Annual Incidence of LD 1992-2006



* Per 100,000 population.

Image courtesy of CDC





Diagnosis

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- ### Diagnosis of Tick-borne Diseases
- Serology: the cornerstone of lab diagnosis
 - Fluorescent antibody staining
 - EIA
 - Western blot
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- ### Western Blot
1. Antigens of an organism are separated by size by gel electrophoresis, blotted onto a nitrocellulose strip
 2. Pt. serum is applied to the strip: any antibody(ies) present bind to the immobilized antigen(s); other serum components are then washed off
 3. Antigen/Antibody complex bands are visualized by immunochemistry
 4. Number and size of visible bands enumerated by eye or computerized blot scanner
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- ### Western Blot
- Newer technique "paints" various purified antigens onto strips at the "correct" positions
 - Also known as "immunoblots"
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Western Blot - LD

- IgM & IgG Blots add specificity over EIA and FA
- Must look at **intensity, size and number** of bands
- CDC interpretive criteria
 - Positive =
 - IgG \geq 5/10 **significant** bands
 - IgM \geq 2/3 **significant** bands



Diagnosis - LD

- CDC Two-tier algorithm
 - Screen with an EIA or IFA
 - Confirm Positive and Equivocal screens with immunoblot (IB)
 - IgM & IgG in 1st month of disease (i.e. 1^o EM present)
 - IgG only thereafter
 - 38%-100% Sensitive, 99% Specific (Bacon et al 2003 J Infect Dis 187:1187)
- CDC: "A clinical diagnosis" in the end

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Diagnosis - LD

- Serological caveats
 - Sensitivity of two-tier serological algorithm increases with length of untreated disease
 - Early therapy blunts immune response
 - IgM persists for \geq 1 yr – do not test IgM after 1mo
- No data supports repeat sero-testing during treatment, or in suspected reoccurrence

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LD Diagnosis - New Fronts

- FDA cleared product scans blots, performs software analysis, and archives strip images
- Painted immunoblot strips may soon be available, allowing for more uniformity, ease of reading
- EIAs using purified VLSE and C6 antigens
 - promise better performance
 - may eliminate/reduce need for WB

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Diagnosis - LD

- Culture
 - Skin Bx: Reserve for very early, unusual EM
 - Recovery from other sources is poor
 - Takes 1-2 weeks or more
 - Not readily available
- PCR
 - Most sensitive on synovial fluid (83%) and CSF (73%)

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LD Diagnosis - Choosing a Reference Lab

- LD specialty labs have arisen in response to "chronic" LD. Can often be found on the Web
- Often do not follow the CDC two-tier serological method, do not use FDA-cleared lab tests, use FDA-cleared tests "off label", or use incompletely validated tests
- Your physicians or patients may ask you to use one of these labs

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LD Diagnosis - Choosing a Reference Lab

What can you do?

- When searching for a reference lab, ask:
 - Are they accredited? (Joint Commission, CAP, CLIA)
 - Does the lab use
 - FDA-cleared tests? If so, are they used "on label"?
 - The CDC 2-Tier LD algorithm?
 - Non-FDA cleared tests? If so, how validated? Data published in peer-reviewed journals?
 - Do the same physicians that run the lab also provide clinical services? (Potential conflict of interest)

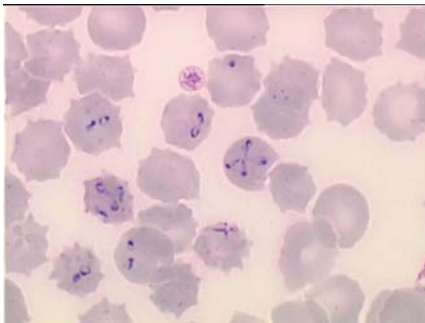


Diagnosis: HGA, Babesiosis

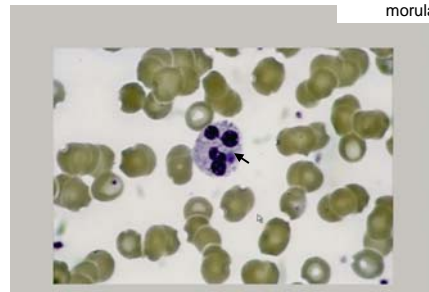
- Blood smear
 - Thin smear fresh whole blood stained with Wright or Giemsa
 - Carefully observe for characteristic forms
 - Ring and tetrad forms of *Babesia*
 - Multiply infected RBCs
 - Extracellular forms
 - Extreme size variation
 - Granulocyte morulae of *A. phagocytophilum*
 - Azure, stippled in appearance



B. microti



A. phagocytophilum
morulae



Photos courtesy of Jim Kazmierczak DVM
WI DPH

Diagnosis: HGA, Babesiosis

- Serology
 - Indirect Fluorescent Antibody (IFA)
 - IgG: 4X rise in acute & convalescent titers, or
 - $\geq 1:64$ HGA
 - $\geq 1:32$ babesiosis
 - IgM: any detectable level
 - Subject to challenges of all FA tests
 - Subjective, need FA 'scope and trained microscopist
- Blood PCR



Treatment



Treatment

- LD
 - **Doxycycline**, Ceftriaxone, Cefuroxime, Amoxicillin
- HGA
 - **Doxycycline**
- Babesiosis
 - Atovaquone+Azithromycin
 - Clindamycin+Quinine



Treatment – LD

- Treatment resistant/recurrent Lyme rarely occurs when appropriately treated
- Reinfection is now recognized, usually in patients previously treated in early disease
- Co-infection does occur: incidence is not clear



Questions?



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