Monitoring of Nesting Songbirds Detects Established Population of Blacklegged Ticks and Associated Lyme Disease Endemic Area in Canada

by
John D Scott, Emily L Pascoe,
Muhammad S Sajid and Janet E Foley

March 18, 2020
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John D. Scott,1,* Emily L. Pascoe 2, Muhammad S. Sajid 2 and Janet E. Foley 2

1 International Lyme and Associated Diseases Society, 2 Wisconsin Circle, Suite 700, Chevy Chase 20815-7007, MD, USA
2 School of Veterinary Medicine, Department of Medicine and Epidemiology, University of California Davis, One Shields Avenue, Davis-95616-8734, CA, USA; jefoley@ucdavis.edu (J.E.F.); mssajid@ucdavis.edu (M.S.S.); elpascoe@ucdavis.edu (E.L.P.)

* Correspondence: jkscott@bserv.com; Tel.: +1-519-843-3646

Received: 20 January 2020; Accepted: 03 March; Published: date

Abstract: This study provides a novel method of documenting established populations of bird-feeding ticks. Single populations of the blacklegged tick, *Ixodes scapularis*, and the rabbit tick, *Haemaphysalis leporispalustris*, were revealed in southwestern Québec, Canada. Blacklegged tick nymphs and, similarly, larval and nymphal rabbit ticks were tested for the Lyme disease bacterium, *Borrelia burgdorferi sensu lato* (Bbsl), using PCR and the flagellin (*flaB*) gene, and 14 (42%) of 33 of blacklegged tick nymphs tested were positive. In contrast, larval and nymphal *H. leporispalustris* ticks were negative for Bbsl. The occurrence of Bbsl in *I. scapularis* nymphs brings to light the presence of a Lyme disease endemic area at this songbird nesting locality. Because our findings denote that this area is a Lyme disease endemic area, and *I. scapularis* is a human-biting tick, local residents and outdoor workers must take preventive measures to avoid tick bites. Furthermore, local healthcare practitioners must include Lyme disease in their differential diagnosis.

Keywords: songbirds; nesting; fledgling; ticks; established population; *Ixodes scapularis*; *Haemaphysalis leporispalustris*; Canada

1. Introduction

Blacklegged ticks, *Ixodes scapularis* (Acari: Ixodidae), have a native range east of the Rocky Mountains, and are vectors of multiple enzootic pathogens. This human-biting ectoparasite is known to harbor and transmit at least nine different tick-borne, zoonotic pathogens [1]. Songbirds (Order: Passeriformes) are well known to widely disperse *I. scapularis* larvae and nymphs in Canada extending as far west and as far north as northern Alberta [2–8]. During northbound migratory flights, certain avifauna have transcontinental or trans-equatorial flights [9,10]. These long-distance passerines fly hundreds of kilometers, and transport *Amblyomma* ticks, including *A. americanum* [2], *A. dissimile* [11], *A. humerale* [12], *A. imitator* [4], *A. maculatum* [2,3], *A. rotundatum* [13], *A. sabanerae* [14], and *Ixodes* ticks, namely *I. affinis* [15], *I. minor* [16], *I. scapularis* [2,4–8] into Canada from the southern United States and the Neotropics. During the nesting and fledgling period, passerines make multiple flights back and forth to the nest. When these ground-foraging songbirds are parasitized by a number of co-
infesting blacklegged ticks, they have the potential to initiate new geographical foci of ticks in outlying areas [17–19].

Most established populations of *I. scapularis* are infected with the Lyme disease bacterium, *Borrelia burgdorferi* sensu lato (Bbsl) [20]. Worldwide, the Bbsl complex consists of at least 24 genospecies (*Borrelia maritima* being the latest entry); the most common borrelial genospecies in the Temperate Zone of North America is *Borrelia burgdorferi* sensu stricto (Bbss), which is pathogenic to humans and several domestic animals [21]. At least six Bbsl genospecies in North America are known to be pathogenic to humans. Epidemiologically, at least seven human-biting *Ixodes* species act as vectors of Lyme disease spirochetes in Canada, namely *Ixodes angustus*, *Ixodes cookei* (groundhog tick), *Ixodes muris* (mouse tick), *Ixodes pacificus* (western blacklegged tick), *Ixodes scapularis* (blacklegged tick), and *Ixodes spinipalpis* [8,22–24].

When the tick salivary glands are infected with Lyme disease spirochetes, *Ixodes* ticks can transmit Bbsl to the host in less than a day [25], especially during nymphal blood meals. Based on North American studies, Bbsl is normally transmitted in 24–48 h; however, when the salivary glands are infected with Lyme disease spirochetes, transmission to a host can occur in less than 16 hours [26,27]. Using European Bbsl strains and *Ixodes ricinus* ticks, Bbsl has been found disseminated in mice within 12 h [26].

In order for a location to meet the criteria for an established population of *I. scapularis*, an area must have at least six individual ticks or at least two of the three motile life stages identified within a single collection period [28]. Normally the collection period is one year; however, in the present study, it was two months.

The aim of this study was: 1) to determine whether tick-infested songbirds monitored during the nesting period are a means of pinpointing unrecognized populations of ticks and 2) to ascertain if these bird-feeding ticks are infected with tick-borne, zoonotic pathogens.

2. Materials and Methods

2.1. Tick Collection

Ticks were collected from songbirds during the period 8th June to 2nd August 2019 at Montée Biggar (45.088226° N, 74.215678° W) in southwestern Québec, Canada (Figure 1). Live ticks were collected from passerines using hardened stainless steel, #5, superfine-tipped forceps (BioQuip Products, Rancho Dominguez, CA). Live ticks were put into a transparent 8.5 mL, 15.7 mm × 75 mm, round-bottomed polypropylene tube (Sarstedt, Montréal, Québec, CA); each tube contained ticks from a single host. In order to keep live ticks from escaping, a piece of tulle netting (3-cm diameter) was placed over the mouth of the tube, and a push cap, which had a 7-mm hole, was promptly inserted into tube opening. This technique prevented live ticks from escaping and, at the same time, provided ventilation for live ticks. For mailing, polypropylene tubes were put in a double-zipper plastic bag with slightly moistened paper towel to sustain high humidity. All ticks were sent directly to the lab (J.D.S.) for identification. Live, partially and fully engorged larval and nymphal ticks were held to molt (metamorphose to the next developmental life stage) to confirm the species identification and, in certain cases, reveal transstadial passage of tick-borne, zoonotic pathogens.
Figure 1. Map shows the songbird monitoring site at Montée Biggar, Québec. The red dot indicates the location of an established population of blacklegged ticks and associated Lyme disease endemic area.

Taxonomic keys and scientific articles were employed to confirm ixodid tick identifications, namely *I. scapularis* larvae [29,30] and nymphs [30,31] and, likewise, *Haemaphysalis leporispalustris* (rabbit tick) larvae [29] and nymphs [32].

Engorged ticks were exposed to a long-day photoperiod of 16:8 (L:D) h. Background and processing information (i.e., host species, leg band number, geographic location, date collected, collector’s name, developmental life stage, degree of engorgement) were logged for ticks from each avian host. After identification, ticks were stored in 2-mL microtubes (Sarstedt, Montréal, Québec, Canada) containing 95% ethyl alcohol to preserve them and stabilize DNA.

2.2. Spirochete Detection

DNA was extracted from ixodid ticks preserved in 95% ethyl alcohol using the Qiagen DNeasy Blood and Tissue Kit (Qiagen, Valencia, CA, USA) following the manufacturer’s instructions. Quantitative real-time PCR (qPCR) was used in a combined thermocycler/fluorometer (ABI Prism 7700; Applied Biosystems, Foster City, CA) to screen for *B. burgdorferi* s.l. DNA. A probe (6FAM-TTC-GGT-ACT-AAC-TTT-TTA-A) with a 3′ end modified with minor groove binding protein and labelled with corresponding dye, forward (5′-GCT-GTA-AAC-GAT-GCA-GTT-CAA-TC-3′) and reverse (5′-GGC-GGC-ACA-CTT-AAC-ACG-TTA-G-3′) primers targeting a region of the 16S rRNA gene were used [33,34]. Water negative controls were included in each run, while positive controls consisted of DNA from cultured Bbsl. Ticks were considered positive if the cycle threshold was <40, and there was a characteristic amplification curve. To identify *B. burgdorferi* s.l. genospecies in qPCR-positive samples, conventional PCR using primers (5′-AAR-GAA-TTG-GCA-GTT-CAA-TC-3′ and 5′-GCA-ATT-TTA-GCA-AGT-GAT-G-3′) that target the flagellin (*flaB*) gene were performed as previously described [35]. Modifications were made to use GoTaq Green Master Mix (Promega, Madison, WI). Gel electrophoresis was performed on conventional PCR products using 1% agarose gel stained with Gelstar (Lonza, Rockland, ME). Amplicons of approximately 497 bases were excised from the gel and purified using the QIAquick Gel Extraction Kit (Qiagen, Valencia, USA), and sequenced using an ABI 3730 sequencer (Davis Sequencing, Davis, CA). Results were examined for accuracy of base determination, and end-read errors were trimmed [36]. We compared sequences to those in the GenBank database using the Basic Local Alignment Search Tool (BLAST), National Center for Biotechnology Information (NCBI), Bethesda, MD [37]. The MUSCLE algorithm was used to perform sequence alignments for *fla* gene sequences from ticks and from reference *Borrelia* genospecies obtained from the NCBI GenBank database [38].

3. Results

3.1. Tick Collection

In total, 68 ticks were collected from 23 songbirds representing five families within the Order Passeriformes (Table 1). These songbird-derived ticks include 64 blacklegged ticks (nymphs, \( n = 63 \); larva, \( n = 1 \)) and four *H. leporispalustris* (nymphs, \( n = 3 \), larva, \( n = 1 \)). It is noteworthy that all tick collections were made before southward migration started and, therefore, all ticks were locally acquired.

In one particular bird parasitism, a Veery was parasitized by three engorged *I. scapularis* nymphs, and one fully engorged nymph molted to a male in 31 d. As well, another Veery was parasitized by two engorged *I. scapularis* nymphs, and a partially engorged nymph molted to a male in 33 d. In addition, a partially engorged *I. scapularis* nymph was collected from a Rose-breasted Grosbeak, and it molted to a
female in 40 d; this female was positive for Bbsl. In the present study, Veeries were the most frequently occurring songbirds followed by Common Yellowthroats (Table 1).

3.2. Spirochete Detection

All ticks testing positive for Bbsl belonged to the genospecies *B. burgdorferi* sensu stricto. Significantly, 14 (42%) of 33 *I. scapularis* nymphs tested were positive (Table 1). A single *I. scapularis* larva was collected, but not tested for Bbsl. All larval and nymphal *H. leporispalustris* were negative for Bbsl. Four *I. scapularis* nymphs were collected from an American Robin on 22<sup>nd</sup> July 2019, and each was positive for Bbsl. Of epidemiological merit, five (100%) of five *I. scapularis* nymphs collected from American Robins were positive for Bbsl. As well, seven *I. scapularis* nymphs collected from an American Robin were pooled for Bbsl testing; the pool was positive for Bbsl. This pool was not included in the Bbsl prevalence calculations.

A total of 23 ticks were not tested for Bbsl because they got hot in transit, and died and spoiled.

Bbsl DNA sequences from this study were submitted to GenBank, and the accession numbers are listed in Table 2.

### Table 1. Detection of *Borrelia burgdorferi* sensu stricto in ticks collected from nesting songbirds captured at Montée Biggar, Québec, 8th June to 2nd August, 2019.

<table>
<thead>
<tr>
<th>Bird species</th>
<th>No. of hosts</th>
<th>No. of ticks</th>
<th>No. of</th>
<th>Hlp</th>
<th>No. of nymphs</th>
<th>Isc</th>
<th>No. of nymphs</th>
<th>I. scapularis nymphs (%)</th>
</tr>
</thead>
<tbody>
<tr>
<td>American Robin, <em>Turdus migratorius</em> L.</td>
<td>3</td>
<td>14&lt;sup&gt;*&lt;/sup&gt;</td>
<td>0/0</td>
<td>0/0</td>
<td></td>
<td>0/0</td>
<td>5/5, 2NT</td>
<td>5/5 (100)</td>
</tr>
<tr>
<td>Common Yellowthroat, <em>Geothlypis trichas</em> (L.)</td>
<td>6</td>
<td>19</td>
<td>0/0</td>
<td>0/1</td>
<td></td>
<td>0/0</td>
<td>2/6, 12NT</td>
<td>2/6 (33)</td>
</tr>
<tr>
<td>Rose-breasted Grosbeak, <em>Pheucticus ludovicianus</em> (L.)</td>
<td>2</td>
<td>3</td>
<td>0/0</td>
<td>0/0</td>
<td></td>
<td>0/0</td>
<td>1/1, 2NT</td>
<td>1/1 (100)</td>
</tr>
<tr>
<td>Chestnut-sided Warbler, <em>Setophaga pensylvanica</em> (L.)</td>
<td>1</td>
<td>1</td>
<td>0/0</td>
<td>0/0</td>
<td></td>
<td>0/0</td>
<td>0/0, 1NT</td>
<td>0/0 (0)</td>
</tr>
<tr>
<td>Veery, <em>Catharus fuscescens</em> (Stephens)</td>
<td>8</td>
<td>26</td>
<td>0/1</td>
<td>0/2</td>
<td></td>
<td>1NT</td>
<td>6/19, 3NT</td>
<td>6/19 (32)</td>
</tr>
<tr>
<td>Song Sparrow, <em>Melospiza melodia</em> (Wilson)</td>
<td>1</td>
<td>3</td>
<td>0/0</td>
<td>0/0</td>
<td></td>
<td>0/0</td>
<td>0/0, 3NT</td>
<td>0/0 (0)</td>
</tr>
<tr>
<td>Gray Catbird, <em>Dumetella carolinensis</em> (L.)</td>
<td>1</td>
<td>1</td>
<td>0/0</td>
<td>0/0</td>
<td></td>
<td>0/0</td>
<td>0/1</td>
<td>0/1 (0)</td>
</tr>
<tr>
<td>Nashville Warbler, <em>Oreothlypis ruficapilla</em> (Wilson)</td>
<td>1</td>
<td>1</td>
<td>0/0</td>
<td>0/0</td>
<td></td>
<td>0/0</td>
<td>0/1</td>
<td>0/1 (0)</td>
</tr>
<tr>
<td>Total: 8 bird species</td>
<td>23</td>
<td>68</td>
<td>0/1</td>
<td>0/3</td>
<td></td>
<td>1NT</td>
<td>14/33, 23NT</td>
<td>14/33 (42)</td>
</tr>
</tbody>
</table>

* Bbsl, *Borrelia burgdorferi* sensu stricto. Hlp, *Haemaphysalis leporispalustris*; Isc, *Ixodes scapularis*. NT, not tested. * Seven *I. scapularis* nymphs collected from an American Robin were pooled for testing, but were not included in the infection prevalence calculations for Bbsl.

### Table 2. Results of sequence analysis of *Borrelia* species positive for *Borrelia burgdorferi* sensu lato and associated NCBI GenBank accession numbers detected in *Ixodes scapularis* nymphs collected from nesting songbirds at Montée Biggar, Québec, 2019.

<table>
<thead>
<tr>
<th>Bird species</th>
<th>Reference strain</th>
<th>flaB gene sequence</th>
</tr>
</thead>
<tbody>
<tr>
<td>American Robin*</td>
<td>CN19-71</td>
<td>MT039724</td>
</tr>
<tr>
<td>American Robin</td>
<td>CN19-75</td>
<td>MT039725</td>
</tr>
<tr>
<td>Common Yellowthroat</td>
<td>CN19-76B</td>
<td>MT039726</td>
</tr>
<tr>
<td>Rose-breasted Grosbeak</td>
<td>CN19-77</td>
<td>MT039727</td>
</tr>
<tr>
<td>Veery</td>
<td>CN19-109A</td>
<td>MT039728</td>
</tr>
<tr>
<td>American Robin†</td>
<td>CN19-112A-1</td>
<td>MT039729</td>
</tr>
<tr>
<td>American Robin†</td>
<td>CN19-112A-2</td>
<td>MT039730</td>
</tr>
<tr>
<td>American Robin†</td>
<td>CN19-112B-1</td>
<td>MT039731</td>
</tr>
<tr>
<td>American Robin†</td>
<td>CN19-112B-2</td>
<td>MT039732</td>
</tr>
<tr>
<td>Veery</td>
<td>CN19-113A</td>
<td>MT039733</td>
</tr>
<tr>
<td>Veery§</td>
<td>CN19-117A-2</td>
<td>MT039735</td>
</tr>
<tr>
<td>Veery§</td>
<td>CN19-117B-1</td>
<td>MT039736</td>
</tr>
<tr>
<td>Veery</td>
<td>CN19-118B</td>
<td>MT039737</td>
</tr>
<tr>
<td>Common Yellowthroat</td>
<td>CN19-121</td>
<td>MT093795</td>
</tr>
</tbody>
</table>
4. Discussion

In this study, we exposed *I. scapularis* and *H. leporispalustris* populations by monitoring a wooded area in southwestern Québec during the nesting and fledgling period. This deciduous woodland turned out to be a breeding colony of *I. scapularis* infected with Lyme disease spirochetes. The presence of Bbsl in these bird-feeding *I. scapularis* nymphs indicates that this nesting area is endemic for Lyme disease. Most importantly, these bird parasitisms during the nesting and fledgling periods provide a unique means of identifying established populations of ticks.

![Common Yellowthroat](image)

Figure 2. Common Yellowthroat, after-second year male, infested with 12 *I. scapularis* nymphs (eight not visible). This multi-tick bird parasitism brings to light the potential of this bird, and the attached nymphs, to establish a new population of blacklegged ticks. Photo credits: Ana Morales.

4.1. Nesting Area Marks Established Population of Blacklegged Ticks

Firstly, a Common Yellowthroat parasitized by 12 *I. scapularis* nymphs signifies an established population in this nesting environs (Figure 2) [28]. Secondly, a Veery was parasitized by immature stages of *I. scapularis* (two nymphs, one larva), and these two developmental stages on a single host within a single year, indicate that this locale is an established population [28]. When engorged *I. scapularis* nymphs molt, they become adults in late June to mid-August. Consistent with other bird-tick-Bbsl studies reporting Bbsl infection prevalence which range from 31% to 59% in *I. scapularis* nymphs in the Great Lakes area [5,7,8,14], we obtained an infection prevalence of 42%. Since these Bbsl-infected nymphs will molt to adults, we extrapolate that the Bbsl infection prevalence for *I. scapularis* adults will be approximately 42% at this locale. Thirdly, when we combine all of the three motile life stages (larvae, nymphs, adults), there is an established population in this woodland locality. In three different ways, we meet the surveillance criteria for an established population of *I. scapularis* ticks [28]. Based on the Bbsl infection prevalence, we show that this area is clearly endemic for Lyme disease.

The determination of whether blacklegged ticks form an established population in an area is often determined by active surveillance. However, in this study, we have shown that ticks collected from songbirds during the nestling and fledgling period are, indeed, an established population. The June–July period is the time of year when songbirds stay in close proximity to the nest. Ecologically, the present bird-tick-pathogen study was conducted during the nesting and fledgling period, which coincides with the peak questing period of nymphal blacklegged ticks [39]. As a result, ground-foraging songbirds are frequently parasitized by host-seeking *I. scapularis* nymphs.

Cricetid rodents, such as deer mice (*Peromyscus maniculatus*) and eastern chipmunks (*Tamias striatus*), act as reservoir hosts for Lyme disease spirochetes, and maintain an enzootic transmission cycle of Bbsl, year-round [40–42]. In an established breeding colony of bird-feeding ticks, overwintering songbirds act as...
maintenance hosts, whereas migratory songbirds are incidental hosts [17]. Because migratory songbirds widely disperse ticks during bidirectional migration, heavily-infested passerine migrants can initiate Bbsl-endemic foci in new areas hundreds of kilometers away from their original source [17–19].

4.2. Short-distance Flights Signify Tick Populations

During the nesting and fledgling period, songbirds forage contiguous to the nest, and maintain short-distance trips back and forth for food. Short runs ensure that the eggs in the nest stay warm and protected from predators. Additionally, during the fledgling period, ground-foraging songbirds make hasty flights near the nest, and promptly return to feed their young. As the ground-frequenting songbirds forage through the leaf litter or grass refuge searching for arthropods, they may become parasitized by host-seeking (questing) ticks. In the present study, bird banders, who were conducting demographic monitoring and collecting data related to songbird activity, had the opportunity to collected bird-feeding ticks. Notably, in late July, songbirds are in moult (shed and replace feathers), and only have localized flight activity. Based on the criteria for an established population [28], we were able to ascertain that there were solo populations of *I. scapularis* and *H. leporispalustris* ticks in the nesting and fledgling area at Montée Biggar, Québec. We were also able to determine that this nesting area is endemic for Lyme disease.

![American Robin](image)

**Figure 3.** American Robin, second year female, parasitized by four *I. scapularis* nymphs (two not visible). On 22nd July 2019, this bird was in moult in preparation for the upcoming, southbound, fall migration. Photo credits: Simon Duval.

4.3. Songbirds as Reservoir Hosts

Certain bird species are reservoir hosts of Bbsl. An American Robin (female, second year) was parasitized with four *I. scapularis* nymphs, and each nymph was infected with Bbsl (Figure 3). Anderson et al. isolated Lyme disease spirochetes from the liver of a Veery [43] and, also, from *I. scapularis* larvae feeding on this avian host. These acarologists also isolated Bbsl from larval *I. scapularis* feeding on Rose-breasted Grosbeaks and Common Yellowthroats suggesting that these songbirds are also reservoir-competent hosts. Other researchers provide substantive evidence that the Veery, Song Sparrow, Common Yellowthroat are competent reservoirs of Bbsl [17,44,45]. Using xenodiagnostic methods, Richter et al. placed unfed spirochete-free *I. scapularis* larvae on Bbsl-inoculated American Robins [46] and, when replete, let these fully engorged larvae molt to nymphs. These unfed nymphs were then placed on spirochete-free American Robins and, subsequently, became infected with Lyme disease spirochetes. These findings reveal that American Robins are competent reservoirs. Similarly, Anderson et al. found that American Robins are reservoir hosts of Bbsl [18]. In the present study, five (100%) of five *I. scapularis* nymphs parasitizing American Robins were infected with Bbsl. In addition, six (32%) of 19 *I. scapularis* nymphs collected from Veeries were infected with Bbsl. Since we did not take any blood samples from host birds, we do not know whether any songbirds were spirochetemic for Bbsl.
Of enzootic significance, a partially engorged *I. scapularis* nymph molted to an unfed female, and this female was positive for Bbsl; this molt demonstrates transstadial passage of Bbsl. This enzootic finding show that Bbsl can move to the next developmental life stage at this site.

4.4. *H. leporispalustris* Vector Competency

In this study, we did not identify Bbsl in *H. leporispalustris* ticks. However, Banerjee et al. detected Lyme disease spirochetes in *H. leporispalustris* ticks collected from a snowshoe hare, *Lepus americanus*, in northern Alberta [47]. As well, Scott et al. discovered *Borrelia laneti*-like spirochetes and a *Babesia divergens*-like piroplasm concurrently in a *H. leporispalustris* (female) collected from an eastern cottontail, *Sylvilagus floridanus*, in southern Manitoba [48]. Scott & Durden provide the first record of a Bbsl-positive *H. leporispalustris* (nymph) collected from an avian host (Swainson’s Thrush) in Canada [14]. Previously, Scott et al. found Bbsl in a *H. leporispalustris* larva parasitizing a passerine (Canada Warbler) in Québec suggesting that this tick species is a reservoir-competent host [8]. During southbound, fall migration in Canada, larval and nymphal *H. leporispalustris* frequently parasitize passerines, and are widely dispersed in southern regions.

![Veery, after-second year male, infested with three *I. scapularis* nymphs. One fully engorged nymph moulted to a male in 31 d. On 22nd July 2019, this bird was moulting to prepare for the southward flight to its wintering range in Brazil. Photo credits: Simon Duval.](image)

4.5. Ticks Co-infest Songbirds

One Veery was co-infested with *H. leporispalustris* (one nymph and one larva), and *I. scapularis* (two nymphs). Not only is there a breeding colony of *I. scapularis* present in this Laurentian River basin, an established population of *H. leporispalustris* is also there. Since these ixodid ticks were collected during the nesting and fledgling period, this bird parasitism denotes a cohabitation of two tick species in this sylvan locale, and signifies that these two tick species are sympatric. The Veery has trans-border and trans-equatorial migration during its northward spring flight, and has a breeding range in southern Canada, including southwestern Québec and northern United States; the wintering range is in central and southeastern Brazil (Figure 4). During the breeding, nesting, and fledgling period, Veeries have localized activity in juxtaposition to the stationary nest. When the young have fledged the nest, these passerines replenish their fat reserves, and prepare for the southbound trek to wintering ranges in southern latitudes during August and September. In late July, they typically moult in preparation for the southbound marathon flight.
4.6. Definition of Lyme Disease Endemic Area

We put forth the following definition for a Lyme disease endemic area based on comprehensive, tick and Lyme disease research across Canada [8], and similarly, conducting field studies and monitoring multiple established tick populations [4,8,12,14,19,22,48–58]: A Lyme disease endemic area is defined as an established population of vector ticks infected with the Lyme disease bacterium, Borrelia burgdorferi sensu lato, at a geographic locality in a single collection period. The present bird–tick–pathogen study at Montée Biggar, Québec meets this definition for a Lyme disease endemic area.

4.7. Clinical Manifestations of Lyme Disease

Lyme disease is an insidious, multisystem spirochetosis that involves many body systems, including cardiac, cutaneous, endocrine, gastrointestinal, genitourinary, musculoskeletal, neurological, otologic, ophthalmological, encephalitic, and neuropsychiatric [59,60]. Only 14%–41% of Lyme disease patients remember a tick bite [61,62], and 9%–39% exhibit an erythema migrans (EM) rash; more than 50% of those with a rash have a homogenous rash [61,63,64]. Bbsl is pleomorphic with diverse forms (i.e., granules, blebs, spherocytes, spirochetes) and, combined together in a gelatinous matrix, these forms become biofilms [65–67]. Lyme disease spirochetes can sequester in protected niche reservoirs, including brain [68–71], bone [72], eye [73], collagenous tissues (ligaments, tendons) [74,75], heart [71,76], kidney [71], liver [71], muscle [77], synovial cells [78], central nervous system [79,80], glial and neuronal cells [81–83], and scar tissue [84].

Because Lyme disease patients often experience lengthy delays in diagnosis and treatment, these patients frequently become persistent cases that result in longstanding Bbsl infections (chronic Lyme disease) [69,72,74,75,78,85–100]. Chronic Lyme disease is defined as a persistent Bbsl infection of at least six months duration [59,60,71,100]. Persistent spirochetal infection has been demonstrated in many Lyme disease patients by PCR and culture [68,71,73,74,86,87,90,91]. Whenever there is continual spirochetemia, approximately 63% of patients infected with Bbsl develop chronic Lyme disease [101]. Patients with this spirochetosis are frequently seronegative especially when serological testing falls outside the peak immune response period, typically 6 to 8 wk after initial infection. Only 48% of patients with persistent Lyme disease test positive using the two-tier serological testing system [102]; dissimilar to AIDS testing, seronegativity is common in Lyme disease patients [103]. Even though Lyme disease patients are seronegative, they may still have active infection because Bbsl side-steps the immune response [71,77,86,88,93,96,104–106]. The Lyme disease spirochete is able to evade the host immune response using a mechanism called stealth pathogenesis. When biofilm busters are implemented prior to blood draw, they help to activate the immune response and provide more reliable serological testing [107]. Psychiatric illnesses, caused by Bbsl, may include violence, substance abuse, and developmental disabilities [108–110]. Chronic Lyme disease frequently causes severe disability, and potentially gives rise to central nervous system complications and cognitive impairment [49,50,61,90]. Under the dire duress of chronic Lyme disease, patients resort to suicide [108–110], and critically ill patients ultimately have fatal outcomes caused by advanced Bbsl spirochetosis [68,71,73,88,111,112]. Bbsl may be potentially transmitted to a partner via intimate relations [89,113]. Gestational Lyme disease takes place when pregnant mothers with Lyme disease pass Bbsl spirochetes across the placenta by vertical transmission from mother to the fetus in utero [114–119].

5. Conclusions

By monitoring nesting songbirds, we demonstrate a novel method of pinpointing tick populations. Nesting songbirds have short-distance flight, and reflect bird-feeding ticks acquired locally. By testing songbird-derived ticks during the nesting and fledgling period, we obtained an infection prevalence of 42% for I. scapularis nymphs, and discovered that this locality is a Lyme disease endemic area. This high level of Bbsl infectivity, reveals that outdoor people working in this newfound, Lyme disease endemic area during
snow-free days encounter a public health risk. Any individuals frequenting this area should take extra precautions to avoid tick bites, and do full-body tick checks at the end of the day. Because *Borrelia burgdorferi* sensu stricto is pathogenic to humans, anyone who is bitten by a tick or has Lyme disease symptoms should seek medical attention. Since chronic Lyme disease is a pernicious, debilitating infection, healthcare practitioners must take special steps to screen symptomatic patients for this incapacitating spirochetosis.

**Author Contributions:** J.D.S. was responsible for study design, coordinating this tick–host-microbe project, and writing the manuscript. J.E.F., E.L.P and M.S.S. conducted molecular testing of ticks and analysis on PCR amplicons. All authors read and approved the final manuscript.

**Funding:** Funding was provided by the Mary Alice Holmes Foundation, and by individual gifts from philanthropic donors Sharleine Haycock and Diane Kindree.

**Acknowledgments:** We thank bird banders Simon Duval and Ana Morales for collecting ticks. We are indebted to Amanda Green for computer graphics.

**Conflicts of Interest:** The authors declare no conflicts of interest.

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