An 82-year-old man was admitted to hospital in late June with progressive fever and confusion. The patient was native to Calgary, Alberta and had not travelled outside the city for many years. While previously well, in the five days preceding hospitalization progressive confusion and an increasing frequency of falls were noted. On admission he had a temperature of 39.1°C, and was hypoxic on room air (85%). A chest x-ray showed evidence of pulmonary edema and a focal infiltrate consistent with rickettsial disease as etiology of fever in individuals presenting with leukopenia/lymphopenia, thrombocytopenia and elevated transaminases during periods of tick activity. Prompt recognition and treatment are important in minimizing resultant morbidity and mortality.

Key Words: Anaplasma phagocytophilum; Ehrlichia; Ixodes; Leukopenia; Thrombocytopenia; Ticks

CASE PRESENTATION

An 82-year-old man was admitted to hospital in late June with progressive fever and confusion. The patient was native to Calgary, Alberta and had not travelled outside the city for many years. While previously well, in the five days preceding hospitalization progressive confusion and an increasing frequency of falls were noted. On admission he had a temperature of 39.1°C, and was hypoxic on room air (85%). A chest x-ray showed evidence of pulmonary edema and a focal infiltrate could not be ruled out. He had normal computed tomography and magnetic imaging brain scans. Initial blood work was remarkable for leukopenia; total 3.7×10^9/L (normal 4.0×10^9/L to 11.0×10^9/L), lymphocytes; 0.1×10^9/L (normal 0.7×10^9/L to 11.0×10^9/L), neutrophils; 3.3×10^9/L (normal 2.0×10^9/L-8.0×10^9/L), thrombocytopenia; 114×10^9/L (normal 150×10^9/L-400×10^9/L), aminotransferase of 102 U/L (normal 8 U/L to 40 U/L) with otherwise normal liver enzymes and a progressive thrombocytopenia to a nadir of 34×10^9/L. A peripheral smear revealed evidence of intracytoplasmic inclusions within granulocytes consistent with rickettsial morulae (Figure 1), suggesting the diagnosis of human granulocytic anaplasmosis (HGA). The patient was immediately started on doxycycline 100 mg by mouth twice daily and made a complete recovery within 48 h. Polymerase chain reaction of whole blood performed at Mayo Medical Laboratories (Rochester, Minnesota) was positive for Anaplasma phagocytophilum. He was believed to have acquired the tick(s) during one of many repeated forays into local urban wooded recreational areas foraging for golf balls. Interestingly, polymerase chain reaction analysis of the removed tick performed at National Microbiology Laboratory of the Public Health Agency of Canada for HGA was negative.

Human granulocytic anaplasmosis (HGA) is a tick-borne rickettsial infection of peripheral blood neutrophils caused by *Anaplasma phagocytophilum*. While this infection is increasingly recognized as endemic throughout much of the United States, no Canadian cases have been previously described, despite the agent being identified in Canadian ticks. Herein we present a case of HGA acquired in an urban Alberta centre. Canadian physicians must be aware of the possibility of tick-borne rickettsial diseases as etiology of fever in individuals presenting with leukopenia/lymphopenia, thrombocytopenia and elevated transaminases during periods of tick activity. Prompt recognition and treatment are important in minimizing resultant morbidity and mortality.

L’anaplasmose granulocytaire humaine (AGH) est une rickettsiose, plus particulièrement une infection des neutrophiles du sang périphérique, transmise par les tiques et causée par *Anaplasma phagocytophilum*. Bien que la maladie soit de plus en plus reconnue comme endémique dans une grande partie des États-Unis, aucun cas n’avait été signalé jusqu’à maintenant, au Canada, malgré le fait que l’agent causal, lui, ait été détecté dans des tiques au pays. Il sera donc question, dans le présent article, d’un cas d’AGH apparu dans un centre urbain, en Alberta. Les médecins, au Canada, doivent penser à une rickettsiose transmise par les tiques comme cause possible de fièvre chez les personnes présentant une leucopénie ou une lymphocytopenie, une thrombocytopenie et un taux élevé de transaminases durant la période d’activité des tiques. Il est important de reconnaître la maladie et de la traiter rapidement afin de diminuer la morbidité et la mortalité.

Key Words: *Anaplasma phagocytophilum*; *Ehrlichia*; *Ixodes*; Leukopenia; Thrombocytopenia; Ticks
TABLE 1
Tick-borne diseases endemic in North America that are transmissible to humans

<table>
<thead>
<tr>
<th>Disease</th>
<th>Causative agent</th>
<th>Vector</th>
<th>Area of endemicity</th>
<th>Common clinical features</th>
<th>Preferred treatment</th>
</tr>
</thead>
<tbody>
<tr>
<td>Tick paralysis (12)</td>
<td>Adherent tick – tick toxin mediated</td>
<td><em>Dermacentor species &gt; Amblyomma species &gt; Ixodes species</em></td>
<td>W Canada and W-central USA</td>
<td>Ascending symmetrical paralysis. Absence of fever</td>
<td>Tick removal, supportive care</td>
</tr>
<tr>
<td>Human granulocytic anaplasmosis (5,6)</td>
<td><em>Anaplasma phagocytophilum</em></td>
<td><em>Ixodes scapularis</em></td>
<td>NE, mid-Mid Atlantic, mid-W and Pacific NW USA, S Ontario, S Nova Scotia</td>
<td>Fever, headache, absence of rash</td>
<td>Doxycycline</td>
</tr>
<tr>
<td>Lyme disease (1)</td>
<td><em>Borrelia burgdorferi</em></td>
<td><em>I scapularis</em></td>
<td>NE, Mid-Atlantic, mid-W and Pacific NW USA, S Ontario, S Nova Scotia, S British Columbia</td>
<td>Early: erythema migrans rash; Late: polyarthritis, neurological abnormalities</td>
<td>Amoxicillin, doxycycline, ceftriaxone</td>
</tr>
<tr>
<td>Babesiosis (13)</td>
<td><em>Babesia microti, Babesia species</em></td>
<td><em>I scapularis</em></td>
<td>NE and mid-W USA</td>
<td>Isolated fever, fulminant disease possible in asplenics</td>
<td>Atovaquone + azithromycin OR clindamycin + quinine</td>
</tr>
<tr>
<td>Rocky Mountain spotted fever (6)</td>
<td><em>Rickettsia rickettsii</em></td>
<td><em>Dermacentor species</em></td>
<td>SE and central USA</td>
<td>Fever, generalized macular-papular rash. High mortality</td>
<td>Doxycycline</td>
</tr>
<tr>
<td>Human monocytic ehrlichiosis (5,6)</td>
<td><em>Ehrlichia chaffeensis</em></td>
<td><em>Amblyomma species &gt; Dermacentor species</em></td>
<td>South-central, SE and mid-Mid Atlantic USA</td>
<td>Fever, headache, absence of rash</td>
<td>Doxycycline</td>
</tr>
<tr>
<td>Tularemia (14)</td>
<td><em>Francisella tularensis</em></td>
<td><em>Amblyomma species/Dermacentor species</em></td>
<td>Central, S-central, Pacific NW USA</td>
<td>Fever, ulcer at inoculation site and lymphadenitis</td>
<td>Gentamicin or tobramycin ± chloramphenicol or doxycycline</td>
</tr>
<tr>
<td>Q fever (14)</td>
<td><em>Coxiella burnetii</em></td>
<td><em>Dermacentor species</em></td>
<td>Endemic in North America</td>
<td>Febrile illness, pneumonia, hepatitis</td>
<td>Doxycycline ± hydroxychloroquine</td>
</tr>
<tr>
<td>Relapsing fever (15)</td>
<td><em>Borrelia species including</em> <em>Borrelia hermsii</em></td>
<td><em>Ornithodoros species</em></td>
<td>W, S-central USA and W Canada</td>
<td>Remittent bouts of high-grade fevers</td>
<td>Doxycycline or chloramphenicol</td>
</tr>
<tr>
<td>Powassan encephalitis (16)</td>
<td>Powassan virus</td>
<td><em>Ixodes species</em></td>
<td>NE USA and E Canada</td>
<td>Fever, weakness, and encephalitis ± meningitis</td>
<td>Supportive care</td>
</tr>
<tr>
<td>Colorado tick fever (17)</td>
<td><em>Colorado tick fever virus</em></td>
<td><em>Dermacentor species</em></td>
<td>W USA and SW Canada</td>
<td>Fever, myalgia, headache</td>
<td>Supportive care</td>
</tr>
</tbody>
</table>

This list is not all inclusive. E East; N North; S South; W West; USA United States

DISCUSSION

There are a number of medically important ticks capable of transmitting zoonoses to humans found in North America (Table 1). In particular, *Ixodes* species ticks are present in many parts of Canada. Self sustaining endemic populations of *I scapularis* are known to exist in eastern Canada (northeast shores of Lakes Ontario and Erie, and southeast Nova Scotia, and possibly southern Quebec, Manitoba and New Brunswick), whereas *Ixodes pacificus*, a vector less commonly implicated in tick-borne disease, is found in many areas of British Columbia (1). However, a recent passive surveillance study of *I scapularis* ticks collected from humans and animals in Canada found that more than 98% of ticks were identified in areas outside of those distinct geographic locations with replicating populations of ticks (representing provinces from Saskatchewan to Newfoundland) (2). Indeed, the majority of ticks in Canada are suspected of being adventitiously introduced by northerly migrating birds. In fact, Ogden et al (3) have extrapolated data collected from more than 39,000 northward migratory birds in the spring of 2005 and 2006 to estimate that between 50 and 175 million *I scapularis* ticks are imported into Canada each year.

HGA is a tick-borne rickettsial infection of neutrophils caused by *Anaplasma phagocytophilum*. While HGA has previously been described in many parts of the United States (northeastern, mid-Atlantic, upper-midwest and Pacific northwest states), Europe and Asia, it has not been described in Canada (although the agent has been found in Canadian ticks) (4). A reportable condition in the United States, documented infections tripled from 2000 to 2005. In 2005, there were 700 cases reported to the Center for Disease Control and Prevention (5). The true burden of infection may be much higher because seroprevalence studies in populations living in areas of high endemicity show 0.6% to 14.9% of individuals have reactive antibodies. However, prospective studies do not show this same burden of disease; therefore, it is unknown if this relates to mild or subacute infection or rather to cross-reacting antibodies to other agents.

Human infection occurs as a result of a bite from a tick infected with *Anaplasma phagocytophilum*. Patients often present with febrile illness associated with headache, fever and myalgias (4-6). Rash is not a common manifestation of disease. Most patients report tick exposure one to two weeks before the onset of infection. Laboratory findings often include thrombocytopenia, leukopenia/lymphopenia and elevated hepatic transaminases. Approximately 50% of patients with documented HGA require hospital admission and 7% require intensive care support. Unlike Rocky Mountain spotted fever and human monocytic ehrlichiosis, death is uncommon and occurs in only 0.5% of documented infections. Complications are uncommon but include sepsis-like syndromes, acute respiratory distress syndrome, heart failure, acute renal failure, rhabdomyolysis and
opportunistic infections. Risk factors for adverse outcomes include advanced age and host immune-suppression. Recently, secondary cases of human to human nosocomial transmission of HGA were documented in China, associated with large volume exposure to the index patient's blood and respiratory secretion (7).

Anaplasma species are uniformly susceptible to tetracyclines (4,5). Because of favourable pharmacokinetics, treatment with doxycycline 100 mg by mouth twice daily for five to 14 days is recommended in patients for whom the drug is not contraindicated. For those with history of allergy, pregnancy or young age, rifampin serves as alternate therapy. Patients typically exhibit full recovery within 24 h to 48 h of therapy.

The majority of I scapularis ticks are not infected with infectious disease agents that are transmissible to humans. Rates of infection vary greatly depending on region where ticks were collected but Canadian data suggests A phagocytophilum affects 0.4% of I scapularis (United States rates are much higher, at 4.7%) (3,8). Other common infections transmitted by I scapularis ticks include Lyme disease (caused by Borrelia burgdorferi), rickettsial species and babesiosis (caused by Babesia microti), although the latter two have not been described as endemic to Canada. There has, however, been one case of transfusion-associated babesiosis acquired in Canada where the blood donor had travelled to endemic zones in the United States (9). Multiply infected ticks are occasionally found, predominately in the eastern and mid-eastern parts of the United States, leading to coinfections with more than one individual tick-borne pathogen (5). There is increasing concern that through environmental changes, including global warming, there will be an associated expansion of tick populations to many other areas of Canada and with them, potentially tick-borne illnesses (10). High-density populations of suitable rodent/wildlife species could readily support their expansion.

While HGA has not previously been described in Canada, there have been occasions in which anaplasmosis has been described in animals. Most recently, three separate cases of canine anaplasmosis were described in Saskatchewan dogs presenting with fever, lethargy, vomiting and diarrhea (11). Furthermore, A phagocytophilum-infected I scapularis ticks have been infrequently identified from migratory birds in eastern Canada (3). Neither HGA nor any other tick-borne disease is currently a mandatory reportable illness in Canada.

In the case described herein, it is apparent that A phagocytophilum from a tick acquired in Canada has resulted in the first case of locally transmitted HGA. Locally reproducing populations of I scapularis are not known to exist in the province of Alberta and as such this likely represents an adventitiously introduced tick. The fact that the adherent tick was negative for A phagocytophilum when screened at the National Microbiology Laboratory should not be surprising, given that normal time from tick acquisition to disease is seven to 14 days (4,5), much longer then tick adherence. As such, it is likely that this gentleman unknowingly had repeated recent tick exposures and a previous tick was responsible for disease transmission.

The present case illustrates that Canadians are at risk for tick-borne illnesses not previously thought to occur in Canada. There is an urgent need for Canadian-specific data regarding the epidemiology of ticks and tick-borne agents in Canada. Patients presenting during times of tick activity (May to September) with fever, leukopenia/lymphopenia, thrombocytopenia and increased serum transaminases should have rickettsial diseases such as HGA included in the differential diagnosis. Lack of historical or epidemiological features for tick exposure should not exclude the diagnosis when the clinical features are present, as illustrated by the present case.

**DISCLOSURE:** All authors are free of potential conflicts. The patient described herein has provided written consent for the publication of this case report.

**REFERENCES**
