Atypical Manifestations of Cat-Scratch Disease, United States, 2005–2014

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Release date: June 17, 2020; Expiration date: June 17, 2021

Learning Objectives

Upon completion of this activity, participants will be able to:

• Assess the epidemiology of atypical CSD in the United States, based on an analysis of data from the 2005 to 2014 MarketScan national health insurance claims databases
• Describe the clinical features of atypical CSD in the United States, based on an analysis of data from the 2005 to 2014 MarketScan national health insurance claims databases
• Describe the clinical and public health implications of the epidemiology and clinical features of atypical CSD in the United States, based on an analysis of data from the 2005 to 2014 MarketScan national health insurance claims databases

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Laurie Barclay, MD, freelance writer and reviewer, Medscape, LLC. Disclosure: Laurie Barclay, MD, has disclosed no relevant financial relationships.

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DOI: https://doi.org/10.3201/eid2607.200034
Atypical manifestations that can be severe and difficult to diagnosis develop in 5%–20% of patients with cat-scratch disease. To clarify the epidemiology of atypical cat-scratch disease in the United States, we analyzed data from the 2005–2014 MarketScan national health insurance claims databases by using the International Classification of Diseases, 9th Revision, Clinical Modification, codes for cat-scratch disease and selected atypical manifestations: retinitis/neuroretinitis, conjunctivitis, neuritis, encephalitis, hepatosplenic disease, osteomyelitis, erythema nodosum, and endocarditis. Atypical cat-scratch disease accounted for 1.5% of all cases, resulting in an average annual incidence of 0.7 cases/100,000 population. Atypical cat-scratch disease was associated with increased risk for hospitalization (risk ratios 8.77, 95% CI 6.56–11.72) and occurred most often in female patients 10–14 years of age. Ocular (48.7%), hepatosplenic (24.6%), and neurologic (13.8%) manifestations were most common among patients. A more comprehensive understanding of atypical cat-scratch disease can improve patient diagnosis and potentially elucidate pathophysiology of the disease.

Cat-scratch disease, a zoonotic bacterial infection, occurs worldwide and is caused by Bartonella henselae, a fastidious, intracellular gram-negative bacillus (1). Cats are the major reservoir of B. henselae and are infected by Ctenocephalides felis cat fleas. Although most cats infected with B. henselae are asymptomatic, signs such as fever and myocarditis might develop in some cats (2,3). Humans usually become infected through the scratches or bites of infected cats. B. henselae has also been shown to infect dogs (4), in some cases resulting in canine endocarditis (5–7). Although some human cases of cat-scratch disease have been linked to canine–human transmission (8–12), further research is needed to clarify the public health significance of B. henselae infection in dogs.

The true burden of cat-scratch disease in the United States is unknown because it is not a reportable condition; however, efforts have been made to estimate its incidence in the United States. In 1993, an analysis of hospital discharge data estimated a nationwide incidence of hospitalized cases of 0.77–0.86 cases/100,000 population annually (13). A subsequent study that examined a database of national health insurance claims during 2005–2013 found that incidence of cat-scratch disease in the United States was highest in southern states (6.4 cases/100,000 population) and in children 5–9 years of age (9.4 cases/100,000 population) (14).

Cat-scratch disease typically manifests as fever and an erythematous papule at the site of the cat scratch or bite, followed by lymphadenopathy in the regional lymph nodes that drain the area of inoculation (15). The papule usually appears 3–10 days after inoculation and can persist for several weeks, with regional lymphadenopathy developing 1–3 weeks postinoculation (1). From 80% to 95% of cases of cat-scratch disease are consistent with this typical presentation, and the remainder of cases manifest as atypical and more severe symptoms (16,17).

Atypical manifestations of cat-scratch disease can involve the eyes, nervous system, heart, liver, spleen, skin, or musculoskeletal system and might result in major illness (1,15). When cat-scratch disease involves the eye, the anterior compartment might be affected by Parinaud oculoglandular syndrome, and the posterior compartment might be affected by retinitis, retinochoroiditis, optic neuritis, uveitis, and vitritis (18–20). Nervous system involvement most commonly manifests as encephalitis, but seizures, nerve palsies, neuritis, myelitis, and cerebellar ataxia have also been reported (21,22).

Endocarditis is more often seen in adults with cat-scratch disease than in children, although preexisting valvular disease puts children at increased risk for this complication (1). Bartonella infection can also cause abdominal pain and microabscesses in the liver and spleen (23), and in immunocompromised hosts can result in bacillary peliosis hepatitis (24). In addition to the classic erythematous papule at the site of inoculation, erythema nodosum and bacillary angiomatosis are reported dermatologic manifestations of atypical infection (25,26). Osteolytic lesions, osteomyelitis, and arthritis have also been associated with cat-scratch disease (16,24,26). A study in 1998 found cat-scratch disease to be the third leading cause of prolonged fever of unknown origin in children, and a history of cat exposure was frequently absent (27).

Atypical manifestations of B. henselae infection can be severe, difficult to diagnose, and lead to lasting impairment. It is unclear why certain patients develop atypical cat-scratch disease, and little is known about its epidemiology. Improved understanding of atypical cat-scratch disease could lead to better recognition of cases by clinicians and inform efforts to understand the pathophysiology of this disease. The purpose of this study was to better characterize the rare and serious complications of this nonreportable zoonotic infection by using nationwide insurance claims data.

**Methods**

To identify potential cases of atypical cat-scratch disease, we conducted a retrospective analysis of
persons enrolled in the Truven Health MarketScan Commercial Claims and Encounters Database (Truven Health Analytics, https://www.ibm.com) during 2005–2014. The MarketScan Commercial Claims and Encounters Database includes persons <65 years of age covered by select employer-sponsored health insurance plans in all 50 states and contains administrative claims data on outpatient visits, inpatient admissions, and emergency department visits. Demographically, the MarketScan population generally mirrors the US population, with a slight overrepresentation of persons 50–59 years of age and a slight underrepresentation of persons 20–29 years of age (28).

Billing codes from outpatient, inpatient, and emergency department visits are assigned by either a clinician or billing specialist according to the International Classification of Diseases, 9th Revision, Clinical Modification (ICD-9-CM), and procedures are captured as either ICD-9-CM codes, Current Procedure Terminology codes, or Healthcare Common Procedure Coding System codes. Because the International Classification of Diseases, 10th Revision, Clinical Modification, was not officially adopted in the United States until 2015, those codes were not included.

We identified cat-scratch disease cases by extracting all enrollee visit records during the study period with an ICD-9-CM code for cat-scratch disease (078.3). The first instance of a 078.3 diagnosis code in a patient record was considered the index event. Atypical manifestations of interest were selected for analysis if they had recorded precedent in the literature as a complication of cat-scratch disease and distinct, clearly discernable ICD-9-CM codes associated with the specific manifestation. Based on these criteria, the known complications of cat-scratch diseases included for analysis were endocarditis, osteomyelitis, erythema nodosum, conjunctivitis, retinitis/neuroretinitis, encephalitis, neuritis, and hepatosplenic disease. We included ICD-9-CM codes associated with optic neuritis in the retinitis/neuroretinitis category. Encounters with an ICD-9-CM code for cat-scratch disease and an accompanying diagnostic code to indicate the anatomic location of a wound or inoculation site for B. henselae were also flagged for analysis and were categorized as either head or neck region, arm or shoulder region, leg or hip region, or torso region. We compiled a detailed list of all ICD-9-CM codes used to identify atypical manifestations of cat-scratch disease (Appendix Table, https://wwwnc.cdc.gov/EID/article/26/7/20-0034-App1.pdf).

We extracted insurance billing records of enrollees with ICD-9-CM codes for cat-scratch disease and selected manifestations at either the same encounter or within a 30-day window of one another. These records were evaluated along with previous and subsequent records by 2 independent reviewers (R.J.M. and C.A.N.) to ensure that the clinical picture was consistent with the coded atypical manifestation based on diagnosis codes, procedure codes, and provider types. If plausible alternative causes of the selected manifestation or likely coding errors were identified, we did not include the enrollee record as an atypical case. In cases of discordance, a third reviewer (Paul Mead) determined final categorization based on record review.

We included persons with an ICD-9-CM code for cat-scratch disease but without accompanying atypical manifestation as typical cases for comparison. Residence in a rural area was assigned if an enrollee did not reside in a metropolitan statistical area, as designated by the US Office of Management and Budget. Because previous research has identified increases in cat-scratch disease in late summer, fall, and January (13,14,29), we categorized month of onset as either late summer and fall, January, or all other months for analysis.

We performed descriptive and comparative statistical analyses by using JMP version 13.2.1 (https://www.jmp.com) and SAS version 9.3 (https://www.sas.com). We used Pearson χ² tests or Fisher exact tests for comparisons of categorical variables. To compare the conditional probability of having atypical cat-scratch disease across strata of potential variables of interest (e.g., sex, age category), we calculated risk ratios (RRs) and associated 95% CIs. Human subjects review at the Centers for Disease Control and Prevention determined that institutional review board approval was not required for this study.

**Results**

**Study Population and Incidence**

During 2005–2014, the MarketScan database contained a median of 44,488,485 (range 16,159,068–53,131,420) enrollees each year. Of 14,824 cat-scratch disease cases identified from MarketScan during this period, 224 (1.5%) cases were classified as atypical (Table 1). The average annual incidence of atypical cat-scratch disease diagnoses during the study period was 0.7 cases/100,000 population.

Atypical cat-scratch disease was most common among adults 15–49 years of age (47.3%), and patients with atypical cat-scratch disease were more likely to be hospitalized than those with typical manifestations (p<0.0001).
Atypical Manifestations of Cat-Scratch Disease

**Table 1.** Characteristics of patients with cat-scratch disease and risk factors for development of atypical cat-scratch disease, United States, 2005–2014

| Characteristic                        | Typical disease, no. (%) | Atypical disease, no. (%) | Risk ratio (95% CI)* |
|--------------------------------------|--------------------------|---------------------------|---------------------|
| Sex                                  |                          |                           |                     |
| M                                    | 5,583 (38.2)             | 94 (42.0)                 | 1.17 (0.90–1.52)    |
| F                                    | 9,017 (61.8)             | 130 (58.0)                | Referent            |
| Age, y                               |                          |                           |                     |
| Child <14                            | 4,678 (32.0)             | 81 (36.2)                 | 1.20 (0.91–1.57)    |
| Adult, 15–49                         | 6,421 (44.0)             | 106 (47.3)                | Referent            |
| Adult, 50–64                         | 3,501 (24.0)             | 37 (16.5)                 | 0.63 (0.44–0.90)    |
| Month of onset                       |                          |                           |                     |
| Late summer and fall†                | 5,470 (37.5)             | 93 (41.5)                 | 1.18 (0.90–1.56)    |
| January                              | 1,490 (10.2)             | 22 (9.8)                  | 1.03 (0.65–1.64)    |
| All other months‡                    | 7,640 (52.3)             | 109 (48.7)                | Referent            |
| Hospitalized                         | 487 (3.3)                | 56 (25)                   | 8.77 (6.56–11.72)   |
| Residence in southern state          | 7,732 (53.0)             | 129 (57.6)                | 1.20 (0.93–1.57)    |
| Residence in rural area              | 3,235 (22.1)             | 51 (22.8)                 | 1.06 (0.78–1.45)    |
*Outcome for risk ratio calculations was development of atypical cat-scratch disease.
†August, September, October, and November.
‡February, March, April, May, June, July, and December.

**Distribution by Age and Sex**
Children ≤14 years of age accounted for 36.2% of atypical cat-scratch disease diagnoses overall; 26 cases (11.4%) were in female patients 10–14 years of age (Figure 1). Among female patients 10–14 years of age, 16 patients (61.5%) had ocular manifestations (13 retinitis/neuroretinitis and 3 conjunctivitis), and 6 patients (23.1%) had hepatosplenic disease.

Nearly half of all patients with atypical cat-scratch disease were younger adults (15–49 years of age). When we compared older adults with younger adults, older adults (50–64 years of age) had a decreased risk for having atypical cat-scratch disease (RR 0.63, 95% CI 0.44–0.90) (Table 1).

**Seasonality**
Atypical cat-scratch disease diagnoses increased from August through March, and diagnoses were concentrated during August–October (33.5% of diagnoses) and January–March (29.5% of diagnoses) (Figure 2), although neither diagnosis in late summer and fall or diagnosis in January was found to be risk factors for development of atypical cat-scratch disease (Table 1). Trends in atypical cat-scratch disease diagnoses were similar to trends in typical cat-scratch disease diagnoses. However, typical cat-scratch disease had less defined peak periods, and diagnoses decreased sharply after January.

**Geographic Distribution and Residence in Rural Area**
Most (57.6%) cases of atypical cat-scratch disease occurred in the southern region of the United States (57.6%), followed by the midwest (16.5%) and northeast (12.5%) regions (Figure 3). The geographic distribution of atypical cases did not differ significantly from cases of typical cat-scratch disease.

Residence in a rural area was not a risk factor for development of atypical cat-scratch disease (RR 1.06, 95% CI 0.78–1.45). Also, the proportion of patients with atypical cat-scratch disease living in a rural area did not differ from the proportion of patients with typical cat-scratch disease living in a rural area (p = 0.70).
Atypical Manifestations by Type

Among 224 patients with atypical cat-scratch disease, 109 (48.7%) had ocular manifestations (retinitis/neuroretinitis or conjunctivitis), 55 (24.6%) had hepatosplenic disease, and 31 (13.8%) had neurologic manifestations (neuritis or encephalitis). The remaining 33 (14.7%) case-patients had osteomyelitis, erythema nodosum, or endocarditis (Table 2).

Among patients with ocular manifestations, 82.6% had retinitis/neuroretinitis; most (64.5%) patients with neurologic manifestations had encephalitis. Three (1.3%) patients with atypical cat-scratch disease had >1 manifestation: 1 patient with osteomyelitis and hepatosplenic disease; 1 patient with endocarditis and hepatosplenic disease; and 1 patient with osteomyelitis, encephalitis, and hepatosplenic disease.

Children ≤14 years of age were at increased risk for hepatosplenic disease (RR 1.76, 95% CI 1.04–2.99) and osteomyelitis (RR 3.81, 95% CI 1.28–11.37) compared with persons ≥15 years of age. Older adults (50–64 years of age) were less likely to show development of ocular manifestations (retinitis/neuroretinitis and conjunctivitis) than younger adults (15–49 years of age) (RR 0.49, 95% CI 0.28–0.85).

Among persons with ocular (retinitis/neuroretinitis and conjunctivitis) manifestations, most diagnoses were made during August–October (31.2%) and January–March (35.8%). Among persons with neurologic (neuritis and encephalitis) manifestations, diagnoses were concentrated during October (22.6%). We observed no notable trends in seasonality of diagnoses for other manifestations of atypical cat-scratch disease (Figure 4). We also observed no differences in geographic distribution or rurality by manifestation of atypical cat-scratch disease.

Hospitalization of Atypical Case-Patients

Patients with atypical cat-scratch disease were more likely to be hospitalized than patients with typical cat-scratch disease (RR 8.77, 95% CI 6.56–11.72) (Table 1). Among patients with atypical cat-scratch disease, children ≤14 years of age accounted for 60.7% of hospitalizations and had an increased risk for hospitalization compared with adults 15–49 years of age (RR 2.34, 95% CI 1.44–3.79). A total of 57.1% of the hospitalizations occurred during August–November, and we found an overall increased risk for hospitalization during this period when compared with all other months, except for January (RR 1.88, 95% CI 1.15–3.05) (Table 3).

Increased risks of hospitalization were found for neurologic manifestations (neuritis and encephalitis) (RR 1.88, 95% CI 1.15–3.08), hepatosplenic disease (RR 2.30, 95% CI 1.49–3.55), and osteomyelitis (RR 2.13, 95% CI 1.20–3.82). Ocular manifestations (retinitis/neuroretinitis and conjunctivitis) were associated with decreased risk for hospitalization (RR 0.23, 95% CI 0.12–0.43).

Location of Wound

Information on wound location was available for only 10 (4.5%) patients with atypical cat-scratch disease. Among these persons, 2 with conjunctivitis, 1 with encephalitis, and 2 with hepatosplenic disease had a wound on the head or neck; 1 with osteomyelitis and 3 with hepatosplenic manifestations had a wound on the arm or shoulder; 1 with endocarditis had a wound on the leg or hip;
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Discussion

Using US nationwide insurance claims data, we identified and characterized 224 atypical cases of cat-scratch disease during 2005–2014 and estimated an average annual incidence of 0.7 cases/100,000 population. Nearly half of all atypical cat-scratch disease cases had ocular manifestations, most of which were retinitis/neuroretinitis. Atypical cat-scratch disease was most prevalent among female patients 10–14 years of age, who most commonly had ocular manifestations.

Trends in hospitalizations of patients with cat-scratch disease highlight the severity of atypical cat-scratch disease compared with typical cat-scratch disease. Atypical cat-scratch disease appears to be particularly severe among children ≤14 years of age, who had an increased risk for hospitalization. Adults 50–64 years of age had the lowest risk for development of atypical cat-scratch disease and specifically ocular manifestations. Reasons that older adults might have complications associated with cat-scratch disease less often than other age groups are unclear and require further study.

Severity of cat-scratch disease in children has been previously documented. In a study conducted by Reynolds et al., ≈25% of hospitalizations of children for cat-scratch disease were caused by complications associated with atypical cat-scratch disease; neurologic and hepatosplenic complications were most common (30). Although children in our study were also particularly at risk for hepatosplenic disease, neurologic and hepatosplenic complications were associated with increased risk for hospitalization in our overall study population, indicating that these manifestations are particularly severe for all age groups. Encephalitis was the most common neurologic manifestation in our population, which was also consistent with findings of Reynolds et al., in which most hospitalizations of children for neurologic complications of cat-scratch disease were caused by encephalitis or encephalopathy (30). Thus, physicians should consider cat-scratch disease in patients who have encephalitis or new onset hepatosplenic abnormalities, especially children.

Table 2. Demographic characteristics for patients by manifestation of atypical cat-scratch disease, United States, 2005–2014

| Characteristic | No. (%) | Sex, no. (%) | Age category, y, no. (%) | Hospitalized, no. (%) |
|---------------|---------|--------------|--------------------------|-----------------------|
| Atypical disease | 224* | 94 (42.0) | 130 (58.0) | 81 (36.2) | 106 (47.3) | 37 (16.5) | 56 (25.0) |
| Ocular disease | 109 (48.7) | 46 (42.0) | 63 (58.5) | 33 (30.7) | 60 (56.6) | 16 (43.2) | 10 (17.9) |
| Retinitis/neuroretinitis | 90 (40.2) | 36 (40.0) | 54 (45.1) | 23 (28.4) | 53 (50.0) | 14 (37.8) | 8 (14.3) |
| Conjunctivitis | 19 (8.5) | 10 (52.6) | 9 (47.4) | 10 (52.6) | 7 (36.8) | 2 (5.2) | 2 (10.5) |
| Hepatosplenic disease | 55 (24.6) | 24 (43.6) | 31 (57.4) | 25 (45.5) | 21 (39.1) | 9 (16.3) | 24 (42.9) |
| Neurologic disease | 31 (13.8) | 13 (41.9) | 18 (57.1) | 12 (38.7) | 13 (41.9) | 6 (19.4) | 13 (23.2) |
| Encephalitis | 20 (8.9) | 12 (60.0) | 8 (40.0) | 12 (60.0) | 8 (40.0) | 0 (0.0) | 13 (23.2) |
| Neuritis | 11 (4.9) | 1 (9.1) | 10 (90.9) | 0 (0.0) | 5 (45.5) | 6 (54.5) | 0 (0.0) |
| Osteomyelitis | 14 (6.3) | 6 (42.9) | 8 (57.1) | 9 (64.3) | 4 (30.8) | 1 (7.1) | 7 (50.0) |
| Erythema nodosum | 11 (4.9) | 2 (18.2) | 9 (81.8) | 4 (36.4) | 5 (45.5) | 2 (18.2) | 4 (36.4) |
| Endocarditis | 8 (3.6) | 4 (50.0) | 4 (50.0) | 1 (12.5) | 4 (50.0) | 3 (37.5) | 2 (25.0) |

*A total of 228 manifestations of atypical cat-scratch disease were seen among 224 patients. Three patients had >1 manifestation; 1 patient had osteomyelitis and hepatosplenic disease; 1 patient had endocarditis and hepatosplenic disease; and 1 patient had osteomyelitis, encephalitis, and hepatosplenic disease.

and 1 with endocarditis had a wound on an unspecified limb.

Figure 4. Seasonal variation of atypical cat-scratch disease manifestations by month of diagnosis, United States, 2005–2014.
Previous studies have documented the highest rates of cat-scratch disease in late summer and fall and a separate peak often seen in January (13,14,29). One such study found that rates of B. henselae seropositivity among samples submitted to Mayo Clinic Laboratories over a 10-year period were highest during September–January, with the highest annual rates in January (29). Typical cat-scratch disease diagnoses in our study followed similar seasonal patterns to those previously reported. However, atypical cat-scratch disease appeared more concentrated during August–October and January–March. The reasons for this finding are unclear but might include delays in diagnosis of atypical cat-scratch disease. For example, patients who contract cat-scratch disease and had complications during January might not be given a diagnosis of atypical cat-scratch disease at that time because they do not show classic symptoms or their symptoms take time to develop and care-seeking is delayed.

Furthermore, a recent case series of ocular manifestations of atypical cat-scratch disease reported that 9 of 10 patients had symptoms ≤3 months before showing development of ocular complications and that 3 patients had been originally given misdiagnoses of etiologies other than cat-scratch disease (31). Given that ocular manifestations of cat-scratch disease were most common in our study, increased diagnoses of atypical cat-scratch disease through March could be a sign of delayed diagnoses, particularly for manifestations that are less severe, such as those involving the eye.

Trends related to geographic distribution of cases did not differ between atypical and typical cat-scratch disease. Similar to findings from 3 previous studies that reported the highest incidences of cat-scratch disease in the southern United States (13,14,30), in our study, most typical (53.0%) and atypical (57.6%) cat-scratch disease cases occurred in patients residing in this region. In addition, a national survey of US healthcare providers found that those in the Pacific and southern regions of the United States were more likely to have been given a diagnosis of cat-scratch disease than in other regions (32). These findings are further supported by studies that have found higher average B. henselae seroprevalences and active bacteremia in pet cats from warmer, more humid climates, including the southern United States (33,34). Thus, healthcare providers in regions with climates that support flea abundance should be aware of the risk for cat-scratch disease and be able to recognize its atypical manifestations.

This study had several limitations. First, although MarketScan is a large database of insurance claims data from persons covered by employer-sponsored insurance, it is a convenience sample and may not accurately represent the characteristics of all persons in the United States. For example, trends we see in atypical cat-scratch disease by geographic region and rural residence might be biased by differences in coverage and access to care that are not accounted for here. Furthermore, MarketScan does not include data for adults >65 years of age, military personnel, uninsured persons, or Medicaid/Medicare enrollees. These specific populations might show varying degrees of cat-scratch disease severity or risk that are not captured in our results. In addition, because only persons <65 years of age are included in the database, the proportion of children who have cat-scratch disease might be artificially inflated. The number of patients who had atypical cat-scratch disease was small, especially when broken down by manifestation. Thus, it is difficult to draw conclusions regarding risk factors for specific manifestations of atypical cat-scratch disease and hospitalization within these groups.

### Table 3. Demographic characteristics for patients hospitalized with atypical cat-scratch disease and associated risk factors for hospitalization, United States, 2005–2014

| Characteristic | Hospitalized, no. (%) | Not hospitalized, no. (%) | Risk ratio (95% CI) |
|---------------|-----------------------|---------------------------|--------------------|
| Sex | | | |
| M | 26 (46.4) | 68 (40.5) | 1.19 (0.76–1.89) |
| F | 30 (53.6) | 100 (59.5) | Referent |
| Age, y | | | |
| Child <14 | 34 (60.7) | 47 (28.0) | 2.34 (1.44–3.79) |
| Younger adult, 15–49 | 19 (33.9) | 87 (51.8) | Referent |
| Older adult, 50–64 | 3 (5.4) | 34 (20.2) | 0.45 (0.14–1.44) |
| Month of onset | | | |
| Late summer and fall | 32 (57.1) | 61 (36.3) | 1.88 (1.15–3.05) |
| January | 4 (7.1) | 18 (10.7) | 0.99 (0.38–2.62) |
| All other months† | 20 (35.7) | 89 (53.0) | Referent |
| Residence in southern state | 36 (64.3) | 93 (55.4) | 1.33 (0.82–2.14) |
| Residence in rural area | 11 (19.6) | 40 (23.8) | 1.26 (0.71–2.26) |

*August, September, October, and November.
†February, March, April, May, June, July, and December.
Furthermore, misclassification could have occurred when ICD-9-CM codes were used to classify atypical cat-scratch disease for several reasons. ICD-9-CM codes are subject to error from the clinicians and billing specialists who enter them. In addition, we excluded records that fit our criteria for manifestations of atypical cat-scratch disease but lacked additional supporting information, which could have caused us to underestimate the true burden of atypical cat-scratch disease. Last, codes for some known atypical cat-scratch disease manifestations, such as pulmonary complications and thrombocytopenia, were excluded because of etiologic ambiguity in enrollee records.

In conclusion, our findings indicate that atypical cat-scratch disease in the United States follows trends similar to those for typical cat-scratch disease but is more prevalent and severe among children <14 years of age and is least likely to occur in older adults (50–64 years of age). In addition, differences in seasonality of diagnoses were seen, which might be an indication that diagnosis of atypical cat-scratch disease is often delayed. Ocular (retinitis/neuroretinitis and conjunctivitis) and hepatosplenic complications were the most common manifestations of atypical cat-scratch disease. Improved understanding of atypical cat-scratch disease might lead to better recognition of cases by clinicians, as well as inform efforts to clarify the pathophysiology of this disease.

Acknowledgments
We thank Paul Mead for serving as the third reviewer of insurance billing records of enrollees with ICD-9-CM codes in cases of discordance between the 2 reviewers and Dan Pastula for providing assistance with defining neurologic manifestations of cat-scratch disease.

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Appendix Table. ICD-9-CM codes and descriptions used to identify atypical manifestations of cat-scratch disease*

| ICD-9-CM code | Description                                      |
|---------------|--------------------------------------------------|
| 3689          | Unspecified visual disturbance                   |
| 22803         | Hemangioma of retina                             |
| 36011         | Sympathetic uveitis                               |
| 36100         | Retinal detachment with retinal defect, unspecified |
| 36101         | Recent retinal detachment, partial, with single defect |
| 36102         | Recent retinal detachment, partial, with multiple defects |
| 36103         | Recent retinal detachment, partial, with giant tear |
| 36104         | Recent retinal detachment, partial, with retinal dialysis |
| 36105         | Recent retinal detachment, total or subtotal      |
| 3612          | Serous retinal detachment                        |
| 36189         | Other forms of retinal detachment                |
| 3619          | Unspecified retinal detachment                   |
| 36213         | Changes in vascular appearance of retina          |
| 36229         | Other nondiabetic proliferative retinopathy       |
| 36230         | Retinal vascular occlusion, unspecified           |
| 36254         | Macular cyst, hole, or pseudohole                 |
| 36281         | Retinal hemorrhage                                |
| 36282         | Retinal exudates and deposits                     |
| 36283         | Retinal edema                                     |
| 36289         | Other retinal disorders                           |
| 3629          | Unspecified retinal disorder                     |
| 36321         | Pars planitis                                     |
| 36330         | Choroidal detachment, unspecified                 |
| 36370         | Serous choroidal detachment                       |
| 36371         | Hemorrhagic choroidal detachment                  |
| 3638          | Other disorders of choroid                        |
| 3639          | Unspecified disorder of choroid                   |
| 36405         | Hypopyon                                          |
| 36901         | Better eye: total vision impairment; lesser eye: total vision impairment |
| 36902         | Better eye: near-total vision impairment; lesser eye: not further specified |
| 36903         | Better eye: near-total vision impairment; lesser eye: total vision impairment |
| 36904         | Better eye: near-total vision impairment; lesser eye: near-total vision impairment |
| 36905         | Better eye: profound vision impairment; lesser eye: not further specified |
| 36906         | Better eye: profound vision impairment; lesser eye: total vision impairment |
| 36907         | Better eye: profound vision impairment; lesser eye: near-total vision impairment |
| 36908         | Better eye: profound vision impairment; lesser eye: profound vision impairment |
| 36909         | Moderate or severe impairment, better eye, impairment level not further specified |
| 36910         | Better eye: severe vision impairment; lesser eye: blind, not further specified |
| 36911         | Better eye: severe vision impairment; lesser eye: total vision impairment |
| 36912         | Better eye: severe vision impairment; lesser eye: near-total vision impairment |
| 36913         | Better eye: severe vision impairment; lesser eye: not further specified |
| 36914         | Better eye: severe vision impairment; lesser eye: profound vision impairment |
| 36915         | Better eye: moderate vision impairment; lesser eye: blind, not further specified |
| 36916         | Better eye: moderate vision impairment; lesser eye: total vision impairment |
| 36917         | Better eye: moderate vision impairment; lesser eye: near-total vision impairment |
| 36918         | Better eye: moderate vision impairment; lesser eye: profound vision impairment |
| 36919         | Better eye: moderate or severe impairment, both eyes, impairment level not further specified |
| 36921         | Better eye: severe vision impairment; lesser eye: impairment not further specified |
| 36922         | Better eye: severe vision impairment; lesser eye: severe vision impairment |
| 36923         | Better eye: moderate vision impairment; lesser eye: severe vision impairment |
| 36924         | Better eye: moderate vision impairment; lesser eye: severe vision impairment |
| ICD-9-CM code | Description |
|--------------|-------------|
| 36925 | Better eye: moderate vision impairment; lesser eye: moderate vision impairment |
| 36926 | Unqualified visual loss, both eyes |
| 36961 | Profound impairment, one eye, impairment level not further specified |
| 36962 | One eye: total vision impairment; other eye: not specified |
| 36963 | One eye: total vision impairment; other eye: normal vision |
| 36964 | One eye: near-total vision impairment; other eye: vision not specified |
| 36965 | One eye: near-total vision impairment; other eye: near-normal vision |
| 36966 | One eye: near-total vision impairment; other eye: normal vision |
| 36967 | One eye: profound vision impairment; other eye: vision not specified |
| 36968 | One eye: profound vision impairment; other eye: near-normal vision |
| 36969 | One eye: profound vision impairment; other eye: normal vision |
| 36970 | Moderate or severe impairment, one eye, impairment level not further specified |
| 36971 | One eye: severe vision impairment; other eye: vision not specified |
| 36972 | One eye: severe vision impairment; other eye: near-normal vision |
| 36973 | One eye: severe vision impairment; other eye: normal vision |
| 36974 | One eye: moderate vision impairment; other eye: vision not specified |
| 36975 | One eye: moderate vision impairment; other eye: near-normal vision |
| 36976 | One eye: moderate vision impairment; other eye: near-normal vision |
| 36977 | One eye: moderate vision impairment; other eye: normal vision |
| 36978 | Unqualified visual loss, one eye |
| 36979 | Unspecified visual loss |
| 37221 | Angular blepharoconjunctivitis |
| 37222 | Contact blepharoconjunctivitis |
| 37289 | Other disorders of conjunctiva |
| 37289 | Unspecified disorder of conjunctiva |
| 37700 | Papilledema, unspecified |
| 37702 | Papilledema associated with decreased ocular pressure |
| 37703 | Papilledema associated with optic nerve sheaths |
| 37732 | Retrolubar neuritis (acute) |
| 37742 | Hemorrhage in optic nerve sheaths |
| 37749 | Other disorders of optic nerve |
| 3688 | Other specified visual disturbances |
| 4387 | Late effects of cerebrovascular disease, disturbances of vision |
| 37239 | Other conjunctivitis |
| 3410 | Neuromyelitis optica |
| 3630 | Focal chorioretinitis and focal retinochoroiditis (fill) |
| 3631 | Disseminated chorioretinitis and disseminated retinochoroiditis (fill) |
| 3632 | Other and unspecified forms of chorioretinitis and retinochoroiditis (fill) |
| 3634 | Visual field defects (fill) |
| 3725 | Conjunctival degenerations and deposits (fill) |
| 3770 | Papilledema (fill) |
| 3771 | Optic atrophy (fill) |
| 3772 | Other disorders of optic disc (fill) |
| 3773 | Optic neuritis (fill) |
| 3790 | Scleritis and episcleritis (fill) |
| 3792 | Disorder of vitreous body (fill) |
| 3799 | Unspecified disorder of eye and adnexa (fill) |
| 36900 | Profound impairment, both eyes, impairment level not further specified |
| 36912 | Panuveitis |
| 36212 | Exudative retinopathy |
| 36231 | Central retinal artery occlusion |
| 36232 | Retinal arterial branch occlusion |
| 36233 | Partial retinal arterial occlusion |
| 36234 | Transient retinal arterial occlusion |
| 36235 | Central retinal vein occlusion |
| 36241 | Venous tributary (branch) occlusion |
| 36300 | Central serous retinopathy |
| 36301 | Focal chorioretinitis, unspecified |
| 36302 | Focal chorioretinitis and chorioretiitns, juxtapapillary |
| 36303 | Focal chorioretinitis and choriioretinitis of other posterior pole |
| 36304 | Focal choroiditis and chorioretinitis, peripheral |
| 36305 | Focal retinitis and retinochoroiditis, juxtapapillary |
| 36306 | Focal retinitis and retinochoroiditis, macular or paramacular |
| 36307 | Focal retinitis and retinochoroiditis of other posterior pole |
| 36308 | Focal retinitis and retinochoroiditis, peripheral |
| 36310 | Disseminated chorioretinitis, unspecified |
| 36311 | Disseminated chorioretinitis and chorioretinitis, posterior pole |
| 36312 | Disseminated chorioretinitis and chorioretinitis, peripheral |
| 36313 | Disseminated chorioretinitis and chorioretinitis, generalized |
| ICD-9-CM code | Description                                              |
|--------------|----------------------------------------------------------|
| 36314        | Disseminated retinitis and retinocchoroiditis, metastatic |
| 36315        | Disseminated retinitis and retinocchoroiditis, pigment epitheliopathy |
| 36403        | Secondary iridocyclitis, infectious                       |
| 37600        | Acute inflammation of orbit, unspecified                 |
| 37701        | Papilledema associated with increased intracranial pressure |
| 372          | Disorders of conjunctiva                                 |
| 3720         | Acute conjunctivitis (fill)                              |
| 37201        | Serous conjunctivitis, except viral                       |
| 37202        | Acute follicular conjunctivitis                           |
| 37203        | Other mucopurulent conjunctivitis                         |
| 37204        | Pseudomembranous conjunctivitis                           |
| 3722         | Blepharoconjunctivitis, unspecified                       |
| 37222        | Blepharoconjunctivitis, unspecified (fill)                |
| 3723         | Other and unspecified conjunctivitis                     |
| 37230        | Conjunctivitis, unspecified                              |
| 37261        | Granuloma of conjunctiva                                 |
| 3773         | Optic neuritis (fill)                                    |
| 37730        | Optic neuritis, unspecified                              |
| 37739        | Other optic neuritis                                     |
| 36320        | Chorioretinitis, unspecified                             |
| 7301         | Chronic osteomyelitis (fill)                             |
| 73010        | Chronic osteomyelitis, site unspecified                   |
| 73011        | Chronic osteomyelitis, shoulder region                    |
| 73012        | Chronic osteomyelitis, upper arm                          |
| 73013        | Chronic osteomyelitis, forearm                            |
| 73014        | Chronic osteomyelitis, hand                              |
| 73015        | Chronic osteomyelitis, pelvic region and thigh            |
| 73016        | Chronic osteomyelitis, lower leg                          |
| 73017        | Chronic osteomyelitis, ankle and foot                     |
| 73018        | Chronic osteomyelitis, other specified sites              |
| 73019        | Chronic osteomyelitis, multiple sites                     |
| 73030        | Periostitis, without mention of osteomyelitis, site unspecified |
| 73031        | Periostitis, without mention of osteomyelitis, shoulder region |
| 73032        | Periostitis, without mention of osteomyelitis, upper arm  |
| 73033        | Periostitis, without mention of osteomyelitis, forearm    |
| 73034        | Periostitis, without mention of osteomyelitis, hand       |
| 73035        | Periostitis, without mention of osteomyelitis, pelvic region and thigh |
| 73036        | Periostitis, without mention of osteomyelitis, lower leg  |
| 73037        | Periostitis, without mention of osteomyelitis, ankle and foot |
| 73038        | Periostitis, without mention of osteomyelitis, other specified sites |
| 73039        | Periostitis, without mention of osteomyelitis, multiple sites |
| 73000        | Acute osteomyelitis, site unspecified                     |
| 73001        | Acute osteomyelitis, shoulder region                      |
| 73002        | Acute osteomyelitis, upper arm                            |
| 73003        | Acute osteomyelitis, forearm                              |
| 73004        | Acute osteomyelitis, hand                                 |
| 73007        | Acute osteomyelitis, ankle and foot                       |
| 73009        | Acute osteomyelitis, multiple sites                       |
| 73022        | Unspecified osteomyelitis, upper arm                      |
| 73023        | Unspecified osteomyelitis, forearm                        |
| 73024        | Unspecified osteomyelitis, hand                           |
| 73027        | Unspecified osteomyelitis, ankle and foot                 |
| 73029        | Unspecified osteomyelitis, multiple sites                 |
| 73030        | Orbital osteomyelitis                                    |
| 7305         | Acute osteomyelitis, pelvic region and thigh              |
| 73028        | Unspecified osteomyelitis, other specified sites          |
| 73025        | Unspecified osteomyelitis, pelvic region and thigh        |
| 73026        | Unspecified osteomyelitis, other specified sites          |
| 73027        | Unspecified osteomyelitis, shoulder region                |
| 73028        | Unspecified osteomyelitis, pelvic region and thigh        |
| 73029        | Unspecified osteomyelitis, other specified sites          |
| ICD-9-CM code | Description |
|--------------|-------------|
| 73020        | Unspecified osteomyelitis, site unspecified |
| 4378         | Other ill-defined cerebrovascular disease |
| 4379         | Unspecified cerebrovascular disease |
| 2930         | Delirium due to conditions classified elsewhere |
| 2931         | Subacute delirium |
| 29389        | Other specified transient mental disorders due to conditions classified elsewhere, other |
| 2939         | Unspecified transient mental disorder in conditions classified elsewhere |
| 2948         | Other persistent mental disorders due to conditions classified elsewhere |
| 2949         | Unspecified persistent mental disorders due to conditions classified elsewhere |
| 3129         | Unspecified disturbance of conduct |
| 3207         | Meningitis in other bacterial diseases classified elsewhere |
| 32082        | Meningitis due to gram-negative bacteria, not elsewhere classified |
| 32089        | Meningitis due to other specified bacteria |
| 3209         | Meningitis due to unspecified bacterium |
| 3220         | Nonpyogenic meningitis |
| 32342        | Other myelitis due to other infections classified elsewhere |
| 32361        | Infectious acute disseminated encephalomyelitis |
| 32362        | Other postinfectious encephalitis and encephalomyelitis |
| 32363        | Postinfectious myelitis |
| 32382        | Other causes of myelitis |
| 3239         | Unspecified causes of encephalitis, myelitis, and encephalomyelitis |
| 32710        | Organic hypersomnia, unspecified |
| 32711        | Idiopathic hypersomnia with long sleep time |
| 32712        | Idiopathic hypersomnia without long sleep time |
| 32713        | Recurrent hypersomnia |
| 32714        | Hypersomnia due to medical condition classified elsewhere |
| 32719        | Other organic hypersomnia |
| 3332         | Myoclonus |
| 3449         | Paralysis, unspecified |
| 3568         | Other specified idiopathic peripheral neuropathy |
| 3570         | Acute infective polyneuritis |
| 78093        | Memory loss |
| 7811         | Disturbances of sensation of smell and taste |
| 7812         | Abnormality of gait |
| 7813         | Lack of coordination |
| 7814         | Transient paralysis of limb |
| 7861         | Abnormal reflex |
| 78039        | Other convulsions |
| 342          | Flaccid hemiplegia |
| 4374         | Cerebral arteritis |
| 345          | Infantile spasms |
| 436          | Acute, but ill-defined, cerebrovascular disease |
| 32341        | Other encephalitis and encephalomyelitis due to other infections classified elsewhere |
| 32381        | Other causes of encephalitis and encephalomyelitis |
| 37701        | Papilledema associated with increased intracranial pressure |
| 3483         | Encephalopathy, not elsewhere classified (fill) |
| 34830        | Encephalopathy, unspecified |
| 34839        | Other encephalopathy |
| 78097        | Altered mental status |
| 78002        | Transient alteration of awareness |
| 78009        | Other alteration of consciousness |
| 78040        | Alteration of consciousness (fill) |
| 780          | General symptoms |
| 7244         | Thoracic or lumbosacral neuritis or radiculitis, unspecified |
| 7292         | Neuralgia, neuritis, and radiculitis, unspecified |
| 7234         | Brachial neuritis or radiculitis NOS |
| 3343         | Other cerebellar ataxia |
| 3344         | Cerebellar ataxia in diseases classified elsewhere |
| 3412         | Acute (transverse) myelitis |
| 78931        | Abdominal or pelvic swelling, mass, or lump, right upper quadrant |
| 78941        | Abdominal rigidity, right upper quadrant |
| 78961        | Abdominal tenderness, right upper quadrant |
| 2895         | Other diseases of spleen (fill) |
| 28950        | Disease of spleen, unspecified |
| 28951        | Chronic congestive splenomegaly |
| 28953        | Neutropenic splenomegaly |
| 28959        | Other diseases of spleen |
| 570          | Acute and subacute necrosis of liver |
| 5719         | Unspecified chronic liver disease without mention of alcohol |
| ICD-9-CM code | Description                                           |
|---------------|-------------------------------------------------------|
| 5732          | Hepatitis in other infectious diseases classified elsewhere |
| 5733          | Hepatitis, unspecified                                |
| 5734          | Hepatic infarction                                   |
| 5735          | Hepatopulmonary syndrome                             |
| 5738          | Other specified disorders of liver                    |
| 5739          | Unspecified disorder of liver                         |
| 7590          | Anomalies of spleen                                  |
| 7904          | Nonspecific elevation of levels of transaminase or lactic acid dehydrogenase |
| 7905          | Other nonspecific abnormal serum enzyme levels        |
| 7948          | Nonspecific abnormal results of function study of liver |
| 78901         | Abdominal pain, right upper quadrant                  |
| 28952         | Splenic sequestration                                 |
| 7891          | Hepatomegaly                                          |
| 7892          | Splenomegaly                                          |
| 2894          | Hypersplenism                                         |
| 572           | Liver abscess and sequelae of chronic liver disease    |
| 5720          | Abscess of liver                                      |
| 421           | Acute and subacute endocarditis                       |
| 4249          | Acute and subacute endocarditis                       |
| 6951          | Erythema multiforme (fill)                            |
| 6952          | Erythema nodosum                                      |
| 2870          | Allergic purpura                                      |

*ICD-9-CM, International Classification of Diseases, 9th revision, clinical modification; NOS, not otherwise specified.