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cattle (2,8). Taken together, the prevalence of EHEC serotype O145:H– in cats, humans, and cattle might indicate that the girl was probably more likely the infection source for the cat than vice versa. Third, a cycle of mutual infection and reinfection between the girl and her pet cat cannot be ruled out. Although the excretion rate for EHEC changes over time and EHEC can therefore remain undetected in stool samples while still present within the patient, the child tested EHEC negative for a short period. Despite all the precautions taken, the girl may have been reinfected by the cat.

This case illustrates several issues: 1) domestic animals such as cats (3), dogs (3,4), and rabbits (9) may serve as reservoirs for EHEC, irrespective of whether they are the primary or secondary source for these bacteria; 2) domestic cats as carriers may excrete EHEC for a prolonged period; 3) autovaccination may be effective for treating EHEC-infected animals; and 4) fondness for pets may be problematic: although EHEC O145:H– is among the 4 most often isolated EHEC serotypes associated with severe colitis or life-threatening hemolytic uremic syndrome (10), the girl’s parents, after weighing the infectious risks against the psychological benefits for both their daughter and her feline companion, decided not to send the cat to an animal shelter until its EHEC infection disappeared.

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Misdiagnosing Melioidosis

To the Editor: Melioidosis is endemic in southern and Southeast Asia and northern Australia. Although relatively few indigenous cases are recognized in the Indian subcontinent, a substantial proportion of cases imported into the United Kingdom originate there, probably reflecting patterns of immigration and travel, and underdiagnosis within the Indian subcontinent (1–3).

A 33-year-old woman spent 3 months in India. Shortly after arriving there, fever, myalgia, rigors, pharyngitis, and tender cervical lymphadenopathy developed. After she received antimicrobial agents, her symptoms initially improved, but in September 2005, 1 week after returning to the United Kingdom, she visited her general practitioner with recurrent fever and increasingly painful cervical lymphadenopathy. She was given a course of oral co-amoxiclav 625 mg 3× daily. However, the following week she visited the emergency department of her local hospital with left-sided suppurative cervical lymphadenitis. Pus aspirated from the lymph node grew an aminoglycoside-resistant "pseudomonad" identified as Pseudomonas fluorescens (API20NE profile 1056554), assumed to be a contaminant. She was discharged home to complete a further 10-day course of co-amoxiclav.

One month later, the patient again went to the emergency department, this time with a submental abscess. An otolaryngology consultation was sought, and the abscess was incised and drained. Although tuberculosis was suspected, no acid-fast bacilli were identified, and cultures were negative for mycobacteria; histologic examination showed noncaseating granulomata. Culture of fluid from the submental collection again yielded an aminoglycoside-resistant pseudomonad, however. At this point misidentifi-
cation was suspected, and the isolate, which had a characteristic colony form on Ashdown’s Medium, microscopic appearance (Figure, panel A), API20NE profile (1556574), and fatty acid profile, was identified as *Burkholderia pseudomallei*, the etiologic agent of melioidosis. The patient had no relevant past medical history. Before immigrating to the United Kingdom 3 years earlier, she had lived in Tanjore, a rice-farming area of Tamil Nadu. She had stayed with family there during her recent trip, which coincided with the monsoon season, but she denied rural travel, fresh water contact, or skin abrasions. On examination, she was obese with acanthosis nigricans and tender cervical lymphadenopathy. Blood tests showed a mild microcytosis, low ferritin level, and erythrocyte sedimentation rate 40 mm/h; serum biochemistry and levels of C-reactive protein, fasting glucose, and hemoglobin by electrophoresis were normal. Two blood cultures were negative. Results of chest and abdominal imaging were normal. The patient was treated with intravenous ceftazidime for 10 days and oral co-trimoxazole for 4 months. She remains well.

*B. pseudomallei* serologic tests, performed subsequently, showed negative results by ELISA against the standard laboratory strain (204). However, when the assay was repeated using the patient’s own isolate, the result was positive (immunoglobulin G titer 4,000). Comparison of lipopolysaccharide (LPS) antigens from the 2 strains by sodium dodecyl sulfate–polyacrylamide gel electrophoresis and immunoblotting showed that they differed in O-repeating units (Figure, panel B).

*B. pseudomallei* is an aerobic, gram-negative, environmental saprophyte ubiquitous in soil and surface water (e.g., paddy fields) in disease-endemic areas. Acquisition may occur through skin abrasions, aspiration of fresh water, inhalation, and possibly ingestion and may occasionally occur in the laboratory. An association between severe respiratory melioidosis and heavy monsoonal rains suggests that inhalation has previously been underrecognized as a route of infection (4); this is the likely mode in this case.

Many infections are initially subclinical but may result in latency and delayed manifestations, even after several decades. Clinical signs and symptoms include septicemia, cavitating pneumonia, bone and soft tissue infections, disseminated abscesses, mycotic aneurysms, lymphadenitis, and childhood parotitis. Most patients have an underlying predisposition to infection, especially diabetes, renal disease, alcoholism, and thalassemia, but in the largest Indian case series 50% patients had no traditional risk factors, as with our patient (5). *B. pseudomallei* is a category B potential bioterrorism agent.

Limited awareness of the disease, confusion with other conditions such as tuberculosis, and laboratory constraints all probably contribute to underdiagnosis of melioidosis in many areas (6). However, accurate diagnosis is important because septi- cemic melioidosis may be rapidly fatal. *B. pseudomallei* is intrinsically resistant to many antimicrobial agents, and prolonged treatment is usually required to minimize relapse. Diagnosis is usually by culture from sterile sites. Laboratory misidentification is not uncommon and occurred in this case because the diagnosis was not considered. Isolation of aminoglycoside-resistant pseudomonads in patients from disease-endemic areas should always prompt consideration of melioidosis and accurate identification. PCR is an emerging diagnostic tool not yet extensively validated (7).

The role of serology in diagnosis is limited by high background seropositivity rates in disease-endemic areas. No standardized serologic test is inter-

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**Figure.** A) Gram stain of pus from the patient’s submental collection, showing the characteristic safety-pin pattern (arrows) of bipolar staining. B) Sodium dodecyl sulfate–polyacrylamide gel electrophoresis of lipopolysaccharide (LPS) antigens from the patient and *Burkholderia pseudomallei* reference strain (204), showing different O-repeating units (bracket). A control isolate of *Pseudomonas aeruginosa* LPS (PA01) is shown for comparison.
Burkholderia pseudomallei and T.L. Pitt confirmed the isolate as
Handy and P.C. Matthews made the initial
responsibility for the patient. R.
mission to publish this case report.

This case illustrates potential pit-
falls in diagnosing melioidosis, which
requires clinical and laboratory aware-
ness and knowledge of its geographic
distribution. LPS-based serologic
assays should use a range of isolates
representative of known LPS types.

Acknowledgment
We are grateful to the patient for per-
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A. J. Brent and R. Handy had clin-
ical responsibility for the patient. R.
Handy and P.C. Matthews made the initial
microbiologic diagnosis of melioidosis,
and T.L. Pitt confirmed the isolate as
Burkholderia pseudomallei. T.L. Pitt per-
formed the serology and SDS-PAGE
analysis of lipopolysaccharide antigens.
All authors contributed to preparation of
the manuscript. A.J. Brent is guarantor
for the article, had full access to all the cli-
nical and microbiologic data, and had final
responsibility for the decision to submit
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Subclinical Plasmodium falciparum
Infection and HIV-1 Viral Load

To the Editor: Studies indicate that Plasmodium falciparum
infection increases HIV replication in adults
(1,2). Although malaria-related illness and
death are more common in chil-
dren, and HIV-1 generally progresses
faster in children than in adults (3,4),
to our knowledge the effect of inter-
mittent malaria on HIV-1 viral load
has not been directly explored in chil-
dren. To investigate this issue, we
monitored HIV-positive infants from
a 1996–2001 birth cohort study in
Kisumu, Kenya, a P. falciparum–
holoendemic area.

Study design and methods have
been described elsewhere (5,6).
Twenty-four children that were peri-
natally infected with HIV were
included in this substudy. During
monthly visits during the child’s first
2 years of life, malaria and HIV inci-
dence were recorded (5,6). Both chil-
dren with malaria-positive blood
smears and those with fever but no
smear result available were treated
with sulfadoxine-pyrimethamine
according to national guidelines. At
the time of this substudy, none of the
study participants were taking anti-
tretroviral drugs.

HIV and malaria diagnoses were
determined by using standard meth-
ods (5–7). To reduce the chance of
including infants infected through
breast-feeding, perinatal infection was
defined as >2 consecutive HIV posi-
tive tests, with the first positive PCR
result by 4 months of age (7). The so-
called baseline viral load was the pre-
malarial value measured 1 month
before the first observation in the
analysis. To be included in the analy-
sis, follow-up visits had to have data
available on the current and previous
months’ viral load and malaria status
and occur at roughly monthly inter-
vals at >4 months of age.

Malaria parasites were found at 53
of 146 visits in the month before viral
load measurement, although at 89%
of visits in which children were
malaria-positive, the children’s sam-
ple sizes had <1,000 parasites/µL, and in
only 13% of visits in which children
had parasitemia they also have fever (8).
Median number of observations
per child was 7 (range 2–18). No
significant demographic or clinical
differences were found between HIV-
positive children in this substudy and
those enrolled in the full cohort (data
not shown).

Clinical and demographic vari-
ables were evaluated in univariate
repeated measures analysis to deter-
mine associations with log-trans-
formed HIV-1 viral load. Age and