

*flab* gene was targeted by using multiplex qPCR according to a previous described protocol (1). For quality control of qPCRs, we included positive and negative controls. Sequences of qPCR products were analyzed and compared with sequences available in GenBank.

*B. miyamotoi* was detected in 7 ticks: 2 (1.59%) of 126 males, 2 (0.68%) of 296 females, and 3 (6.52%) of 46 nymphs. *A. phagocytophilum* was detected in 16 ticks: 1 (0.79%) of 126 males, 11 (3.72%) of 296 females, and 4 (8.70%) of 46 nymphs. *Candidatus* N. mikurensis was detected in 25 ticks: 5 (3.97%) of 126 males, 18 (6.08%) of 296 females, and 2 (4.35%) of 46 nymphs. Overall prevalences were 1.50% for *B. miyamotoi*, 3.42% for *A. phagocytophilum*, and 5.34% for *Candidatus* N. mikurensis. Prevalences of each pathogen in specific varied by locality (Table). No co-infections were detected.

We analyzed *flab*, *msp2*, and *groEL* gene sequences obtained by qPCR. These sequences showed 99%–100% identities with gene sequences of *B. miyamotoi* (GenBank accession no. KJ847050), *A. phagocytophilum* (accession no. KP164415), and *Candidatus* N. mikurensis (accession no. FJ966365).

In Romania, the density of *Ix. ricinus* ticks is high and their host diversity is extensive (7). However, data for effects of tickborne pathogens on public health are scarce in this country. In this study, we detected *B. miyamotoi*, *A. phagocytophilum*, and *Candidatus* N. mikurensis in questing *Ix. ricinus* ticks in Romania, which confirms the emerging trend of these pathogens in Europe. Because of the scarcity of information on human infections with these pathogens in Romania, serologic and molecular investigations and their implementation are needed for diagnosis, which might help in assessing the effect of these pathogens on public health.

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## References

- Hansford KM, Fonville M, Jahfari S, Sprong H, Medlock JM. *Borrelia miyamotoi* in host-seeking *Ixodes ricinus* ticks in England. *Epidemiol Infect.* 2015;143:1079–87. <http://dx.doi.org/10.1017/S0950268814001691>
- Jongejan F, Uilenberg G. The global importance of ticks. *Parasitology.* 2004;129(Suppl):S3–14. <http://dx.doi.org/10.1017/S0031182004005967>
- Platonov AE, Karan LS, Kolyasnikova NM, Makhneva NA, Toporkova MG, Maleev VV, et al. Humans infected with relapsing fever spirochete *Borrelia miyamotoi*, Russia. *Emerg Infect Dis.* 2011;17:1816–23. <http://dx.doi.org/10.3201/eid1710.101474>
- Crowder CD, Carolan HE, Rounds MA, Honig V, Mothes B, Haag H, et al. Prevalence of *Borrelia miyamotoi* in Ixodes ticks in Europe and the United States. *Emerg Infect Dis.* 2014;20:1678–82. <http://dx.doi.org/10.3201/eid2010.131583>
- Fonville M, Friesema IH, Hengeveld PD, Docters van Leeuwen A, Jahfari S, Harms MG, et al. Human exposure to tickborne relapsing fever spirochete *Borrelia miyamotoi*, the Netherlands. *Emerg Infect Dis.* 2014;20:1244–5. <http://dx.doi.org/10.3201/eid2007.131525>
- Kalmár Z, Cozma V, Sprong H, Jahfari S, D'Amico G, Mărcuțan DI, et al. Transstadial transmission of *Borrelia turcica* in *Hyalomma aegyptium* ticks. *PLoS ONE.* 2015;10:e0115520. <http://dx.doi.org/10.1371/journal.pone.0115520>
- Kalmár Z, Mihalca AD, Dumitrache MO, Gherman CM, Magdaş C, Mircean V, et al. Geographical distribution and prevalence of *Borrelia burgdorferi* genospecies in questing *Ixodes ricinus* from Romania: a countrywide study. *Ticks Tick Borne Dis.* 2013;4:403–8. <http://dx.doi.org/10.1016/j.ttbdis.2013.04.007>
- Jahfari S, Fonville M, Hengeveld P, Reusken C, Scholte EJ, Takken W, et al. Prevalence of *Neoehrlichia mikurensis* in ticks and rodents from north-west Europe. *Parasit Vectors.* 2012;5:74. <http://dx.doi.org/10.1186/1756-3305-5-74>
- Matei IA, Kalmár Z, Magdaş C, Magdaş V, Toray H, Dumitrache MO, et al. *Anaplasma phagocytophilum* in questing *Ixodes ricinus* ticks from Romania. *Ticks Tick Borne Dis.* 2015;6:408–13. <http://dx.doi.org/10.1016/j.ttbdis.2015.03.010>
- Andersson M, Zaghoudi-Allan N, Tamba P, Stefanache M, Chitimia L. Co-infection with 'Candidatus Neoehrlichia mikurensis' and *Borrelia afzelii* in an *Ixodes ricinus* tick that has bitten a human in Romania. *Ticks Tick Borne Dis.* 2014;5:706–8. <http://dx.doi.org/10.1016/j.ttbdis.2014.05.013>

Address for correspondence: Zsuzsa Kalmár, University of Agricultural Sciences and Veterinary Medicine, Calea Mănăştur 3–5, Cluj-Napoca 400372, Romania; email: [zsuzsa\\_kalmar@yahoo.com](mailto:zsuzsa_kalmar@yahoo.com)

## Suspected Rabies in Humans and Animals, Laikipia County, Kenya

Mark Obonyo, James M. Akoko, Austine B. Orinde, Eric Osoro, Waqo Gufu Boru, Ian Njeru, Eric M. Fèvre

Author affiliations: Ministry of Agriculture, Livestock and Fisheries, Nairobi, Kenya (M. Obonyo, A.B. Orinde); Field Epidemiology and Laboratory Training Program, Nairobi (M. Obonyo, W.G. Boru); International Livestock Research Institute, Nairobi (J.M. Akoko, E.M. Fèvre); Kenya Zoonotic Disease Unit, Nairobi (A.B. Orinde, E. Osoro); Ministry of Health, Kenya (W.G. Boru, I. Njeru); University of Liverpool Institute of Infection and Global Health, Liverpool, UK (E.M. Fèvre)

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**To the Editor:** Dog bites are a serious public health problem because of the associated risk for rabies virus exposure in countries to which the virus is endemic (1,2).

Human rabies can be prevented by administration of post-exposure prophylaxis (PEP). However, PEP rabies vaccine may be unavailable or prohibitively expensive (3). Delay in or failure to receive PEP after possible rabies virus exposure contributes to increased incidence of human rabies deaths (3).

We performed a retrospective investigation of animal bites and postbite treatment in Laikipia North sub-county, Kenya, during January 2013–February 2014. Laikipia North is 1 of 3 sub-counties in Laikipia County and has a population of 32,726 (4). Our investigation was instigated by 3 suspected human rabies deaths that were informally reported to the Kenya Government Zoonotic Disease Unit (ZDU) during early 2014. We reviewed animal bite records from sub-county health facilities and veterinary offices and administered a structured household questionnaire to determine outcomes, knowledge of rabies, bite management, healthcare-seeking behavior, and economic costs. This public health response was government coordinated and approved; no personal identifiers were retained.

During January 1, 2013–February 10, 2014, a total of 106 bites were recorded by 6 government-run health facilities in Laikipia North. Median reported bite incidence per month was 24 bites/100,000 persons (range 6–45 bites/100,000 persons). The median age of bite victims was 13 years (range 1–81 years); 61 (58%) bites occurred in males. Of all bites recorded, 94 (88%) were by dogs, 8 (8%) by scorpions, and 4 (4%) by humans.

The deaths of 3 humans reported to the ZDU occurred in November and December 2013. To assess whether these cases were part of an exposure cluster, we followed up on bite cases during November 1–December 31, 2013. During this period, 17 additional animal bite cases were recorded. Of these 20 bite cases, we successfully traced the households of 11 (55%) case-patients, including 2 of the 3 who died from rabies. Bites were predominantly received from owned pets (82%), and most bites (82%) were reported to be unprovoked. All bites were inflicted on extremities, and almost all (91%) were single-bite injuries (Table 1).

Of 11 animals that bit case-patients, 7 had unknown histories of rabies vaccination and 4 were not vaccinated (Table 1). Four of the 11 animals were suspected to be rabid, including 1 cat and 3 dogs. All the suspected rabid animals were reported to exhibit aggressive or abnormal behavior, drooling or salivation, vocalization, and roaming tendencies (5; online Technical Appendix Table 1, <http://wwwnc.cdc.gov/EID/article/22/3/15-1118-Techapp1.pdf>). Three of the animals reportedly died; status was unknown for 1.

Of the 11 traced bite case-patients, 9 washed their wound before going to a healthcare facility and 8 were prescribed PEP. The median time from bite to reporting to a health facility was 1 day (range 0–3 days). Four respondents delayed in starting PEP: 3 after 3 days, and 1 after 2 days.

Reasons given for delay included the high cost of PEP by 3 (including 1 who died); a health facility being too far away by 1, who died; and vaccine unavailable at nearest health facility by 2, 1 of whom died. Of 8 respondents who received PEP, 7 traveled >10 km to reach the nearest health facility. PEP availability was inconsistent at the sub-county hospital and local dispensaries; 6 of 8 respondents seeking PEP visited multiple facilities to receive PEP, including a county referral facility that was >100 km away. The World Health Organization's 5-dose PEP regimen is recommended in Kenya (1). However, only 3 case-patients were prescribed and

**Table.** Responses to questionnaire interview of 11 animal bite victims assessed for rabies, Laikipia County, Kenya, 2014\*

Variables/categories	No. (%) case-patients
Time of bite	
Evening	6 (55)
Morning	4 (36)
Afternoon	1 (9)
Part of body bitten	
Legs	8 (73)
Arms	3 (27)
Circumstances of bite	
Unprovoked	9 (82)
Animal provoked	2 (18)
Type of animal	
Dog	10 (91)
Cat	1 (9)
Ownership of biting animal	
Owned	9 (82)
Stray	2 (18)
Rabies vaccination history of biting animal	
Unknown	7 (64)
Not vaccinated	4 (36)
Outcome of biting animal	
Alive and normal	7 (64)
Deceased	4 (36)
Wound washed after bite	
Yes	9 (82)
No	2 (18)
Treatment at healthcare facility	
Anti-tetanus	9 (82)
PEP rabies vaccination	8 (72)
Pain killers	5 (46)
Distance from nearest PEP facility, km	
>10	7 (64)
5–10	3 (27)
0–5	1 (9)
Source of PEP	
Government facility	5 (63)
Chemist	2 (25)
Private hospital	1 (13)
Costs of PEP, US\$†	
No. doses of PEP administered	23
Cost categories	Average (range)
Cost/dose of PEP	≈8 (2–15)
Total cost of PEP doses	≈23 (8–50)
Direct medical cost	≈65 (2–500)
Indirect medical cost	≈34 (4–100)
Average cost for obtaining 1 dose of PEP	≈45 (8–120)

\*PEP, postexposure prophylaxis.

†Average annual exchange rate during 2013 was 1 Kenya shilling/\$0.011586 US.

received 5 doses. Five respondents were prescribed 3, 4, or 6 doses (online Technical Appendix Table 2). This finding indicates large inconsistencies in the PEP prescribing practices in this region of Kenya, a pattern that is similar in other parts of East Africa (6).

Respondents bore all medical costs without subsidy. Direct medical costs were ≈\$2 \$500 (US) per bite victim, and indirect medical costs were ≈\$4 \$100. The average cost of obtaining a single dose of PEP ranged from \$8 to \$120 (Table; online Technical Appendix Table 2).

All respondents had heard of rabies. Nine (82%) knew it was transmitted to humans through a bite from a rabid dog, and 4 (36%) knew that rabies among dogs could be prevented through vaccination.

During 2014, at least 3 suspected human rabies deaths and 4 domestic animal deaths were associated with this cluster. Postbite care, including PEP, is a heavy economic burden on this community, more so because rabies vaccine is not always locally accessible. Dog vaccination rates are low in this region and rabies in suspected animals is rarely definitively diagnosed, increasing risks for human rabies virus exposures and the economic burden of PEP administration. We recommend implementation of regular and comprehensive mass dog vaccination campaigns, in line with Kenya's National Rabies Elimination Strategy (7), and further detailed studies on the epidemiology of rabies in this ecosystem, which supports human, wildlife, and domestic dog populations.

## References

1. World Health Organization. Expert consultation on rabies 2013. Second report. World Health Organization technical report series, (982), 1 [cited 2015 Nov 10]. <http://www.who.int/iris/handle/10665/85346>
2. Charkazi A, Naser B, Mehri F, Abdollatif E, Hossein S, Hashem H. Epidemiology of animal bite in Aq Qala city, northern of Iran. *J Educ Health Promot.* 2013;2:13 <http://dx.doi.org/10.4103/2277-9531.112682>.
3. Sambo M, Cleaveland S, Ferguson H, Lembo T, Simon C, Urassa H, et al. The burden of rabies in Tanzania and its impact on local communities. *PLoS Negl Trop Dis.* 2013;7:e2510 <http://dx.doi.org/10.1371/journal.pntd.0002510>.
4. Kenya National Bureau of Statistics. Kenya population and housing census 2009 [cited 2015 Nov 10]. [http://www.knbs.or.ke/index.php?option=com\\_phocadownload&view=category&id=109:population-and-housing-census-2009&Itemid=599](http://www.knbs.or.ke/index.php?option=com_phocadownload&view=category&id=109:population-and-housing-census-2009&Itemid=599)
5. Tepsunmethanon V, Wilde H, Meslin FX. Six criteria for rabies diagnosis in living dogs. *J Med Assoc Thai.* 2005;88:419–22.
6. Fèvre EM, Kaboyo RW, Persson V, Edelsten M, Coleman PG, Cleaveland S. The epidemiology of animal bite injuries in Uganda and projections of the burden of rabies. *Trop Med Int Health.* 2005;10:790–8 <http://dx.doi.org/10.1111/j.1365-3156.2005.01447.x>.
7. Republic of Kenya Zoonotic Disease Unit. Strategic plan for the elimination of human rabies in Kenya 2014–2030. Nairobi: Ministry of Health and Ministry of Agriculture, Livestock and Fisheries; 2014 [cited 2015 Nov 10]. <http://www.rr-africa.oie.int/docspdf/en/2015/Kenya-National-Rabies-Elimination-Strategy.pdf>

Address for correspondence: Mark O. Obonyo, Ministry of Agriculture, Livestock and Fisheries, Cathedral Road, Nairobi, PO Box 34188-00100, Kenya; email: [mobonyo@feltp.or.ke](mailto:mobonyo@feltp.or.ke)

## Generalized Cowpox Virus Infection in a Patient with HIV, Germany, 2012

Philipp Fassbender, Sabine Zange, Sofi Ibrahim, Gudrun Zoeller, Frank Herbstreit, Hermann Meyer

Author affiliations: Universitaet Duisburg-Essen, Essen, Germany (P. Fassbender, F. Herbstreit); Bundeswehr Institute of Microbiology, Munich, Germany (S. Zange, S. Ibrahim, G. Zoeller, H. Meyer); Edgewood Chemical Biological Center, Aberdeen Proving Ground, Maryland, USA (S. Ibrahim)

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**To the Editor:** In October 2012, a 35-year-old man with clinical category C HIV infection was admitted to the intensive care unit at the University of Duisburg–Essen, Essen, Germany. The man had severe respiratory distress syndrome with septic shock, and he was infected with hepatitis B and C viruses and Epstein-Barr virus. Standard infection-control procedures were followed: the patient was placed in a single room; healthcare providers wore personal protective equipment (gown, face shield, mask, and gloves); and a closed system was used for endotracheal suctioning.

Physical examination of the patient revealed multiple skin lesions on his right forearm and right leg. In the following days, more skin lesions appeared on his abdomen and head. The skin lesions were inflamed macules with central livid, hemorrhagic ulceration (1–2 cm in diameter) and raised edges. Kaposi sarcoma was suspected initially, but on hospital day 5, a skin biopsy showed large intracellular eosinophilic inclusion bodies pathognomonic for infection with cowpox virus (family *Poxviridae*, genus *Orthopoxvirus*). To confirm the diagnosis of cowpox virus infection, we conducted biopsies of 3 skin lesions on hospital day 7. Despite antimicrobial drug and supportive therapy, the patient died that day from septic shock.

The 3 biopsy samples obtained on hospital day 7 were cultured on African green monkey kidney (MA104) cells, and within 2 days, many plaques were observed. DNA extracted from homogenates and virus isolated from the biopsy material were tested by orthopoxvirus real-time PCR (*I*); results were positive for all 6 samples. We confirmed the presence of cowpox virus DNA in all samples by sequencing the hemagglutinin gene.

Serum obtained from the patient on day 2 after admission, when the first lesions were noted, was also positive for