

## In this issue . . .

### Genomic history of Italian brown bears

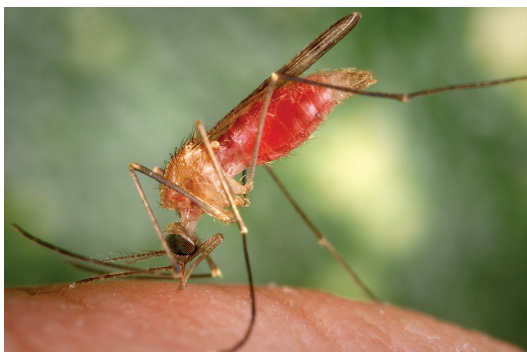
Apennine bears are a critically endangered population of approximately 50 Italian brown bears that live in the central Apennine Mountains. Apennine bears (*Ursus arctos marsicanus*) are separated from other bear populations by several hundred kilometers. Andrea Benazzo et al. (pp. E9589–E9597) explored genomic variation and evolutionary divergence in Apennine bears. The authors sequenced the full genomes of six Apennine bears and six other European brown bears. The authors estimate that Apennine bears were separated from other brown bears during the Neolithic Period, resulting in a 40-fold decline in the Apennine population; Apennine bears may have become isolated due to human expansion and land clearing, the authors suggest. The genetic analysis indicates that Apennine bears are highly inbred with a complete loss of variation in the mitochondrial genome and along long stretches of the nuclear genome. Random drift appears to have fixed several deleterious mutations, but also led to unique features such as small size, distinctive cranial morphology, and reduced aggressiveness, compared with other bears. In combination with a lack of competitors, a nearly vegetarian diet, and low aggressiveness, peaks of high variation in genes associated with the immune and olfactory systems may have helped Apennine bears avoid extinction, according to the authors. — L.C.



Apennine bear. Photo by Valentino Mastrella and image courtesy of Archive from the Abruzzo, Lazio e Molise National Park.

### Synergistic malaria vaccines

In recent years, progress has been made in reducing malaria mortality. However, malaria remains a significant global public health challenge in part due to the emergence of malaria parasites resistant to current frontline antimalarial drugs. A highly effective, strain-transcending vaccine would be a useful tool for malaria elimination. Leyla Bustamante et al. (pp. 12045–12050) evaluated a panel of 29 blood-stage antigens from *Plasmodium falciparum*, a deadly human malaria parasite. The authors used antibodies to systematically screen the antigens for vaccine candidates in two genetically diverse parasite strains in vitro. The screen identified several antigen targets that produced a synergistic effect in blocking erythrocyte invasion, a critical stage of malaria parasite development when the parasite is vulnerable to antibody-directed inhibition. Related immunoepidemiological data from



*Anopheles gambiae*. Image courtesy of the CDC.

an endemic population in Mali mirrored the in vitro findings and revealed that protection from febrile malaria as well as reduced malaria risk were associated with antibodies specific to certain combinations of the antigens. In vitro video microscopy studies of

