

# Spread of endogenous retrovirus K is similar in the DNA of humans and rhesus monkeys

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According to paleontologic and molecular studies, the chimpanzee (*Pan troglodytes*) is the closer relative to the humans (*Homo sapiens*) and that both lineages had a common ancestor at 5 to 7 million years ago. Moreover, the human-chimp lineage split from that of the rhesus monkey (*Macaca mulatta*) around 25 million years ago.

However, by studying the population dynamics of complete copies of primate endogenous retrovirus family K (ERV-K) in the genomes of humans, chimpanzee and rhesus monkey, a surprising pattern was observed.

The study by Romano and colleagues being published this week on PLoS ONE revealed that human ERV-K had a similar demographic signature to that of the rhesus monkey, both differing greatly from that of the chimpanzee. The data suggested that the humans and rhesus have been purging ERV-K copies from their genomes while the chimpanzee ERV-K population kept the signature of increasing numbers of ERV-K amplification in the genome of ancestral primates during the last 20 million years.

Hominins have been moving out of Africa for the last 2 million years and the modern humans (*Homo sapiens*) spread around almost the entire globe during the last 100 thousand years. Moreover, *Macaca* is the most specious primate genus and it is believed to have originated around 2.5 million years ago and became widely dispersed within a short period of time, from the West in Afghanistan to the Eastern coast of China. It is also known that speciation events partition and restrict flow among genetic pools.

As a consequence, both *Homo* and *Macaca* by colonizing new environments and undergoing successive population fluctuations that caused

severe genetic bottlenecks, possible purged ERV-K from their genomes in a similar fashion. On the other hand, populations chimpanzee have been restricted to Eastern and Central Sub-Saharan Africa, ever since and crucially, are also known to have a greater genetic diversity than humans (due to a greater effective population number  $N_e$ ), even when humans have a far greater census population.

While the population size fluctuations due to dispersal or speciation may have had impact on genome architecture, the several expansion and bottlenecks experienced by *Homo* and *Macaca* may have played an important role in shaping ERV-K dynamics.

Because *Pan* did not suffer severe bottlenecks since their separation from the *Pan* – *Homo* (human) common ancestor, they not only show a greater genetic diversity but also they preserved a greater number of complete ERV-K copies in their genomes.

The most remarkable result was that for the first time we could observe that genetic fluctuations caused by bottlenecks and expansion in host species play a fundamental role not only in their genetic diversity but also in the interaction with latent parasites that leave their genome copies in our DNA.

Source: Public Library of Science

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