Recent Loss of Vitamin C Biosynthesis Ability in Bats

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Abstract

The traditional assumption that bats cannot synthesize vitamin C (Vc) has been challenged recently. We have previously shown that two Old World bat species (Rousettus leschenaultii and Hipposideros armiger) have functional L-gulonolactone oxidase (GULO), an enzyme that catalyzes the last step of Vc biosynthesis de novo. Given the uncertainties surrounding when and how bats lost GULO function, exploration of gene evolutionary patterns is needed. We therefore sequenced GULO genes from 16 bat species in 5 families, aiming to establish their evolutionary histories. In five cases we identified pseudogenes for the first time, including two cases in the genus Pteropus (P. pumilus and P. conspicillatus) and three in family Hipposideridae (Coelops frithi, Hipposideros speoris, and H. bicolor). Evolutionary analysis shows that the Pteropus clade has the highest ω ratio and has been subjected to relaxed selection for less than 3 million years. Purifying selection acting on the pseudogenized GULO genes of roundleaf bats (family Hipposideridae) suggests they have lost the ability to synthesize Vc recently. Limited mutations in the reconstructed GULO sequence of the ancestor of all bats contrasts with the many mutations in the ancestral sequence of recently emerged Pteropus bats. We identified at least five mutational steps that were then related to clad origination times. Together, our results suggest that bats lost the ability to biosynthesize vitamin C recently by exhibiting stepwise mutation patterns during GULO evolution that can ultimately lead to pseudogenization.

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Introduction

Vitamin C (Vc), or L-ascorbic acid is a water-soluble vitamin that is an essential nutrient important in animal metabolism. Vc is involved in tissue growth and repair, and also functions as an antioxidant to block damage caused by free radicals. It is also a cofactor in enzymatic reactions that are catalyzed by Cu²⁺-dependent monooxygenases and Fe²⁺-dependent dioxygenases [1]. Vc is required in the diet of all vertebrates in order to sustain good health [2], and Vc deficiency can lead to potentially fatal scurvy in humans. Most vertebrates can satisfy their Vc requirements by synthesizing it de novo with glucose [3]. However, some mammals, including haplorhine primates and guinea pigs, have lost this ability, and thus have to obtain Vc from their diet [4]. The ability to synthesize Vc has been reported in many ancestral vertebrate lineages [5], [6], suggesting the ability for de novo synthesis is ancient. Moreover, there is an apparent transition of the organs used for the biosynthesis of Vc during evolution, from the kidney of reptiles to the liver of mammals [7].

The ability to synthesize Vc has been lost independently several times in vertebrates e.g. in some fishes [5], in some passeriform birds [7], in some bats [8], in guinea pigs [10] and in primates of the suborder Haplorrhini (e.g. monkeys, apes and humans) [7], [9–10]. All of these species lack activity of L-Gulonolactone oxidase (GULO) in their livers (or kidneys) to catalyze the last step of the Vc synthesis pathway so that they need to compensate by obtaining Vc from their food [8–10]. The gene encoding GULO in guinea pigs and humans has become a pseudogene [11], [12].

Our recent research has challenged the traditional opinion that bats cannot synthesize Vc [8], [13] by showing that GULO genes in two species (Rousettus leschenaultii and Hipposideros armiger) are still in their intact forms and can produce functional proteins [14]. Bats are perhaps in the process of large-scale loss of Vc biosynthesis ability [14], and show varying degrees of lack of GULO function. For example, the genera Pteropus and Rousettus belong to the same chiropteran family (Pteropodidae), and although the former has lost the ability to synthesize Vc, the latter retains it [14].

Our previous study on Vc synthesis in bats raises the question-what is the evolutionary pattern that shapes bat GULO evolution in bats? Given the uncertainty of when and how bats lost GULO gene function, it is important to sequence GULO genes of more bat to explore patterns of GULO evolution. In this study, we therefore sequenced the GULO genes of 16 bat species and aimed to reconstruct bat GULO evolutionary history. Using ancestral reconstructions, we infer stepwise mutation patterns showing how bats may have lost GULO function.

Materials and Methods

Bat taxonomic coverage

Bat wing membrane biopsy specimens were taken from a collection at the Institute of Molecular Ecology and Evolution, East China Normal University. All experiments were conducted under permission to use these specimens granted by the by Animal Care Ethics Committee, East China Normal University (approval ID 20091225). Our screening included 16 bat species from 5
families: Pteropodidae, Hipposideridae, Rhinolophidae, Mega-

dermatidae, and Rhinopomatidae, and sampling locations include
China, Cameroon, Australia, India, and Vietnam (supplementary

table S1).

Bat GULO cloning and sequencing

Genomic DNA was extracted by using DNeasy Blood and Tissue

Kit (Qiagen) according to the manufacturer’s protocols. We aimed
to amplify exon-3 to exon-8, the six exons encoding the functional

region of the GULO enzyme (there are 12 exons in the gene in
total), of 16 bat species (supplementary table S1) by using a series
of primer pairs (supplementary table S2) designed according to
the genomic sequences of

P. vampyrus

(GeneScaffold_1205), dog

(Canis familiaris, ENSCFAFT00000013370), cow

(Bos taurus, ENSBATAT00000052052) and pig

(Sus scrofa, ENSSSCT00000010600) that contained
GULO
genes from the Ensembl database (http://

www.ensembl.org/). Polymerase chain reactions (PCR) were

performed using Ex Taq™ polymerase (TaKaRa) with the reaction

conditions as follows: 94°C for 5 min followed by 30 cycles
consisting of 94°C for 30 sec, 57–62°C for 15–30 sec, 72°C for
1 min, and a final extension of 72°C for 10 min. All PCR products

were ligated into pGEM-T Easy vectors (Promega) and trans-

formed. The universal T7 (5’-TAC GAC TCA CTA TAG G-3’) and SP6 (5’-ATT TAG GTG ACA CTA TAG-3’)

sequencing primers were used to sequence all positive clones on

an ABI 3730 DNA sequencer (Applied Biosystems). All new

sequences are submitted to GenBank (supplementary table S1).

Phylogenetic construction

To reconstruct the bat GULO phylogeny, we first retrieved non-bat
orthologous genes in GenBank (www.ncbi.nlm.nih.gov/genbank/):

horse

(Equus caballus, XM_001492727), dog

(Canis familiaris, XM_543226), pig

(Sus scrofa, XM_543226), cow

(Bos taurus, NM_001034043), panda

(Ailuropoda melanoleuca, XM_002914414), rabbit

(Oryctolagus cuniculus, XM_002709304), rat

(Rattus norvegicus, NM_022220), mouse

(Mus musculus, NM_178747), opossum

(Monodelphis domestica, XM_001380006) and platypus

(Ornithorhynchus anatinus, XM_001521551). We then aligned all these sequences with
the bat sequences using ClustalW [15] implemented in MEGA4 [16].
Indels (deletions or insertions) and premature stop codons were
excluded from the sequences before alignment. Because GULO genes
are highly conserved in most mammals, evolutionary history was
inferred using the Neighbor-joining [NJ] method, which was perfectly
used in many studies [17]. All nucleotide positions were included.
The evolutionary distances were computed using the Maximum
Composite Likelihood model [18] with branch lengths represent
genetic distances. All positions containing gaps and missing data were
eliminated. Bootstrapping with 2,000 replicates [19] was used to test
phylogenetic robustness and nodes with bootstrap values lower than
50% were collapsed. Evolutionary analyses were conducted in
MEGA4 [16].

Evolutionary analyses

A widespread method used to infer selection pressures acting on
specific genes is to estimate non-synonymous (dN) and synony-

Figure 1. Alignments of bat GULO nucleotide gene sequences.

(A) The intact bat GULO nucleotide gene sequences; (B) the bat GULO

pseudogenized nucleotide gene sequences. The nucleotide position

numbers are denoted according to the nucleotide sequence of
Rousettus leschenaultii GULO (HQ415789). The boxes denote insertions,
deletions, or premature stop codons that break the gene reading
frames.
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mous (dS) nucleotide substitution rates and their ratio dN/dS (ω) [20]. We used the CODEML program with the likelihood method implemented in PAML4 [20] to evaluate selection pressures acting on each lineage of bat \textit{GULO} genes mapped onto the published species tree [21], [22]. Two models were employed: the \textit{free-ratio} model that allows the ω ratios to vary for each branch, and the \textit{two-ratio} model that compares two different \textit{ω} ratios between specified branches (e.g. \textit{Pteropus} bats) and other branches. To test for the significance of each model used, likelihood ratio tests (LRTs) [23] were implemented, conducted by comparing twice the difference in likelihood between nested statistical models, i.e. the free-ratio versus the one-ratio model (which assumes an average ω for all lineages), and we also compared two-ratio versus one-ratio models.

To trace the amino acid changes during \textit{GULO} gene evolution, ancestral reconstruction was employed [24]. The program CODEML using the empirical Bayes method in PAML4 [20,24] was used for reconstructing amino acids in extinct ancestors on the species tree [21], [22]. Different amino acid changes were recorded after alignment using reconstructed ancestral sequences.

### Results

#### Bat \textit{GULO} cloning

Bats showed lineage-specific gene pseudogenization including premature stop codons, insertions and deletions. Basically, our molecular cloning revealed two major patterns: 1) an intact \textit{GULO} form: \textit{Rousettus leschenaultii}, \textit{R. aegyptiacus}, \textit{Pteropus rodricensis}, \textit{P. vampyrus}, \textit{Eonycteris spelaea} (Pteropodidae), \textit{Rhinolophus ferrumequinum}, \textit{Hipposideros armiger}, \textit{H. ater}, \textit{H. pratti} (Hipposideridae), \textit{Megaderma lyra} (Megadermatidae), and \textit{Rhinopoma hardwickii} (Rhinopomatidae) (figure 1A); 2) pseudogenized form: \textit{Pteropus conspicillatus}, \textit{P. pumilus} (Pteropodidae), \textit{Coelops frithii}, \textit{Hipposideros bicolor}, and \textit{H. speoris} (Hipposideridae) (figure 1B).

#### Patterns of selection in bat \textit{GULO} gene evolution

The phylogenetic gene tree for \textit{GULO} (figure 2) closely resembled the published species tree based on large-scale gene sequencing [21], [22]. Our selection tests showed high \textit{ω} ratios (P-value = 5.0×10^{-6}, LRT) in all main clades of bats, giving a 12–54x higher \textit{ω} ratio than in the ancestor of Laurasiatheria species.

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**Figure 2. Phylogenetic tree based on \textit{GULO} genes.** The evolutionary history was reconstructed using the NJ method in MEGA4 [16]. The bootstrap consensus tree inferred from 2,000 replicates is shown to represent \textit{GULO} evolution for each taxon. Bootstrap values lower than 50% are not shown. The scale bar represents genetic distance. The evolutionary distances were computed using the Maximum Composite Likelihood model. Lineages for pseudogenes are marked with ψ.

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The Pteropus clade has the highest $\omega$ ratio of 0.648. A pair of two-ratio models (with the Pteropus clade was set as the foreground) was then constructed by having the $\omega$ ratio fixed to 1 (neutral evolution) in one model and a $\omega$ ratio not fixing the other. LRT showed no significance when comparing the two models ($P$-value = 0.677), which suggests that the evolution of Pteropus clade GULO genes is close to being neutral.

The higher $\omega$ ratios of Pteropus suggest their GULO genes have already been subjected to relaxed selection (over a period of less than 3 million years ago [mya], the origination time of this clade) [21], [22]. GULO genes of Coelops frithii, Hipposideros armiger, and H. pratti may have been at the early stages of pseudogenization because these genes have relatively low $\omega$ ratios (suggesting they are still under purifying selection). Several two-ratio models were also established with different bat clades as the foreground. LRTs (two-ratio model versus one-ratio model) showed significance only in the Pteropus clade (data not shown), supporting the above conclusion that this clade has subjected to relaxed selection.

**Ancestral reconstruction reveals stepwise mutation patterns**

Interestingly, ancestral sequence reconstruction exhibits a stepwise mutation pattern (figure 4) that starts around the time when the tested bat species first evolved from a common ancestor around 58 mya [21]. The ancestor of all bats maintains most of the original Laurasiatheria gene form (with only two mutations) after divergence with non-bat Laurasiatheria species; the ancestor of Hipposideridae, Rhinolophidae, and Megadermatidae (origin around 52 mya) has 3 mutations; the ancestor of Hipposideridae and Rhinolophidae (origin around 39 mya) has 4 mutations; the ancestor of Pteropodidae (origin around 23 mya) has 7 mutations; and the ancestor of the recently emerged Pteropus bats (around 3
Discussion

As GULO is present in all major vertebrate lineages except some bats (most of these being New World species) [8], anthropoid primates [12], [26], guinea pigs [11], some passerine birds [27], and some fishes [5], such loss-of-function is neither related to broad phylogenetic affiliations nor to diet [28]. Some researchers have even proposed that the loss of Vc synthesis is associated with higher speciation rates because of higher mutation rates [29], which seems unlikely and which has not been tested formally.

Having successfully cloned bat GULO genes from 16 species, we carried out detailed evolutionary analyses. Our results show a range of forms of the GULO gene in bats. Combined with our earlier functional studies [14], we identify the following conditions: 1) pseudogenes that will have lost function, as seen in some Pteropus and hipposiderid species, 2) intact genes that functional studies showed loss of function in Vc synthesis (e.g. Pteropus vampyrus), 3) intact genes that maintain some ability to synthesize Vc (Rousettus leschenaultii and Hipposideros armiger). We found that strong purifying selection has shaped the evolution of bat GULO genes, leading to a diversity of functional outcomes.

Figure 4. Stepwise mutation patterns during bat GULO evolution. Amino acid (abbreviation) changes of bat GULO are shown after the divergence of bats with non-bat Laurasiatheria species. The topology was retrieved from published species trees [21], [22]. Five major steps were identified according to the mutation pattern, and they are highlighted in Roman characters (I, II, III, IV, and V). The time of origin for each node was collected from published data [21], [25]. The positions of mutations are recorded according to the protein sequence of the Rousettus leschenaultii GULO gene (ADP88813). For example, G119A means that the amino acid G evolved from A at amino acid position 119. The outgroup of non-bat Laurasiatheria species include pig, cow, horse, cat, and panda.

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selection has shaped non-*Pteropus* bat pseudogenes, suggesting these bats are in early stages of loss in their ability to synthesize Vc. In the family Hipposideridae some species possess pseudogenes that show only small changes from the intact and functional genes of their close relatives. Together with the evidence for purifying selection our results suggest that Vc function has been lost recently in hipposiderid species showing pseudogenized *GULO*. Relaxed selection acting on *Pteropus* bat *GULO* suggests that bats in this genus lost the ability to synthesize Vc within the past 3 mya [25]. Thus we infer that pseudogenization of bat *GULO* evolved recently.

Ancestral reconstruction clearly shows a stepwise accumulating mutation pattern during bat *GULO* evolution. By mapping each mutation step with the origination times of each clade (figure 4), we surprisingly found that the more ancient the species are, the less mutations they had accumulated; conversely, more recently evolved bats often accumulated many mutations, which supports our hypothesis that Vc synthesis involving *GULO* is gradually becoming less important in bats. The ancestral bats were therefore presumably able to biosynthesize Vc, and during evolution, *GULO* gene function is gradually becoming redundant.

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In conclusion, our study shows that bats are beginning to lose their ability to biosynthesize vitamin C and some have lost this ability in no more than 3 mya. During gene degeneration, stepwise mutation patterns are evident and these are important mechanisms leading eventually to pseudogenization.

Supporting Information

Table S1 Bat GULO genes with taxa information. (XLS)

Table S2 Primers used for amplification of exon-3–exon-8 of bat GULO genes. (DOC)

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Author Contributions

Conceived and designed the experiments: JC SZ. Performed the experiments: JC XY LW. Analyzed the data: JC. Contributed reagents/materials/analysis tools: JC SZ. Wrote the paper: JC GJ.