Divergent and parallel routes of biochemical adaptation in high-altitude passerine birds from the Qinghai-Tibet Plateau

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When different species experience similar selection pressures, the probability of evolving similar adaptive solutions may be influenced by legacies of evolutionary history, such as lineage-specific changes in genetic background. Here we test for adaptive convergence in hemoglobin (Hb) function among high-altitude passerine birds that are native to the Qinghai-Tibet Plateau, and we examine whether convergent increases in Hb-\textsubscript{O\textsuperscript{2}} affinity have a similar molecular basis in different species. We documented that high-altitude parid and aegeithalid species from the Qinghai-Tibet Plateau have evolved derived increases in Hb-\textsubscript{O\textsuperscript{2}} affinity in comparison with their closest lowland relatives in East Asia. However, convergent increases in Hb-\textsubscript{O\textsuperscript{2}} affinity and convergence in underlying functional mechanisms were seldom attributable to the same amino acid substitutions in different species. Using ancestral protein resurrection and site-directed mutagenesis, we experimentally confirmed two cases in which parallel substitutions contributed to convergent increases in Hb-\textsubscript{O\textsuperscript{2}} affinity in codistributed high-altitude species. In one case involving the ground tit (Parus umilis) and gray-crested tit (Lophophanes dichrous), parallel amino acid replacements with affinity-enhancing effects were attributable to nonsynonymous substitutions at a CpG dinucleotide, suggesting a possible role for mutation bias in promoting recurrent changes at the same site. Overall, most altitude-related changes in Hb function were caused by divergent amino acid substitutions, and a select few were caused by parallel substitutions that produced similar phenotypic effects on the divergent genetic backgrounds of different species.

When different species experience similar selection pressures in a shared environment, the probability that they will evolve similar adaptations may be influenced by differences in population size (which determines levels of standing genetic variation and the rate of input of new mutations) and/or differences in the duration of residency in that environment (which determines the time available for new mutations to arise). The probability of evolving similar adaptive solutions may also be influenced by legacies of evolutionary history. Prior genetic changes may preclude or potentiate future changes in a particular trait, in which case the “happenstance of a realized beginning” (1) may play an outsize role in channeling subsequent pathways of evolutionary change. Due to lineage-specific changes in genetic background, different species may hit upon idiosyncratic solutions to the same problem simply because they evolved from different ancestral starting points at the onset of selection.

Mountain ranges and highland plateaus in different parts of the world provide an opportunity to investigate the extent to which native species have followed similar or different routes of high-altitude adaptation. The two highest elevation plateaus in the world, the Andean Altiplano in South America and the Qinghai-Tibet Plateau in Asia, have very different physiographic and biogeographic histories and are inhabited by members of phylogenetically distinct faunas. In the Andes, the passerine birds that inhabit the highest elevations include representatives of the globally distributed Passerida clade (oscines) as well as representatives of the exclusively Neotropical Tyranni clade (suboscines). The avifauna of the Qinghai-Tibet Plateau has a very different phylogenetic composition, and the passerine birds that inhabit the highest elevations include a highly disproportionate number of tits in the family Paridae and long-tailed tits in the family Aegithalidae (3–7). Tits are widely distributed throughout the northern hemisphere and tropical Africa, whereas long-tailed tits are mainly restricted to Eurasia; both groups have their center of diversity in East Asia (7, 8). At high altitude, the challenge of matching reduced \textsubscript{O\textsuperscript{2}} availability with an undiminished cellular \textsubscript{O\textsuperscript{2}} demand is especially acute.

Significance

Mountain ranges and highland plateaus in different parts of the world provide an opportunity to investigate the extent to which native species have followed similar or different routes of adaptation to the challenges of life at high altitude. Here we demonstrate that high-altitude songbirds from the Qinghai-Tibet Plateau independently evolved derived increases in hemoglobin-\textsubscript{O\textsuperscript{2}} affinity in comparison with their closest lowland relatives in East Asia. In comparisons that also included more distantly related high-altitude avian taxa, site-directed mutagenesis experiments revealed two cases in which convergent increases in hemoglobin-\textsubscript{O\textsuperscript{2}} affinity were caused by identical amino acid substitutions at the same sites. However, most adaptive convergence in protein function was attributable to different amino acid substitutions in different species.


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for small, active endotherms like passerine birds that cannot rely on metabolic suppression as a general strategy of hypoxia tolerance. To compensate for the reduced partial pressure of O₂ (P0₂) in inspired air, physiological adjustments involving numerous steps in the O₂-transport pathway can help sustain O₂ flux to the tissue mitochondria in support of aerobic ATP synthesis (9–11). In combination with changes in the cardiorespiratory system and microcirculation, changes in the oxygenation properties of hemoglobin (Hb) can enhance the O₂ capacitance of the blood (the total amount of O₂ unloaded for a given arterio-venous difference in O₂ tension). Under severe hypoxia, an increased Hb-O₂ affinity safeguards arterial O₂ saturation, thereby securing tissue oxygenation, albeit at a lower pressure gradient for O₂ diffusion from the peripheral capillaries to the cells of respiring tissues (12). Evolutionary increases in Hb-O₂ affinity may be caused by amino acid mutations that increase the intrinsic O₂ affinity of the Hb tetramer and/or mutations that suppress the sensitivity of Hb to the affinity-reducing effects of allosteric cofactors (nonheme ligands such as Cl⁻ ions and organic phosphates) (12, 13).

In the Andes, birds that are high-altitude natives have generally evolved derived increases in Hb-O₂ affinity in comparison with their closest lowland relatives (14–17). However, convergent increases in Hb-O₂ affinity in different high-altitude species are seldom attributable to convergent or parallel changes at the amino acid level (17). Here we test whether different high-altitude parids and aegithalid species from the Qinghai-Tibet Plateau have independently evolved increased Hb-O₂ affinities in comparison with their closest lowland relatives in East Asia, and we examine whether convergent changes in Hb function have a similar molecular basis in different phylogenetic lineages. In this context we make comparisons among different high-altitude parid and aegithalid species in the Sino-Himalayan region, and, using previously published data (14–17), we also make comparisons with a phylogenetically diverse set of high-altitude Andean birds. If lineage-specific changes in genetic background or happenstances of biogeographic history predisposed the Andean and Sino-Himalayan passerines to follow different evolutionary paths, then representatives of the two distinct avifaunas may exhibit qualitatively distinct adaptations to hypoxia. If such contingencies influence the adaptive evolution of Hb function, such that highland representatives of different lineages hit upon different solutions to the same problem, this may be reflected in clade-specific or region-specific patterns of molecular parallelism.

Results and Discussion
We conducted a survey of sequence variation in the adult-expressed α- and β-type globin genes in a set of 162 bird specimens representing 13 tit species in the family Paridae (n = 135 specimens) and 4 long-tailed tit species in Aegithalidae (n = 27 specimens). We collected these specimens from localities spanning a broad range of elevations on the Qinghai-Tibet Plateau, the mountains of Southwest China, and in eastern China. Phylogenetic relationships and elevational ranges of focal species are summarized in Fig. S1.

For a subset of six tit species, we experimentally examined functional properties of native Hbs purified from red blood cells. The set of species used in these experiments included extreme high-alpine specialists such as the ground tit (Parus humilis, elevational range = 3,100–5,500 m above sea level), which is endemic to the Qinghai-Tibet Plateau (6, 7), as well as predominantly highland species such as the rufous-vent ed tit (Periparus rubidiventris, 2,400–4,300 m) and gray crested tit (Lophophanes dichrous, 2,300–4,600 m), and predominantly lowland species such as the oriental tit (Parus minor, sea level–2,000 m) and the marsh tit (Poecile palustris, sea level–2,100 m). We also sampled multiple specimens from high- and low-altitude subspecies of the broadly distributed coal tit [Periparus ater aemodius (2,100–4,600 m) and Periparus ater pekinensis (sea level–1,800 m)]. In addition to analyzing the native Hbs of these seven taxa (six distinct species, including geographically distinct subspecies of Periparus ater), we also functionally tested recombinantly expressed Hbs (rHbs) from an additional pair of high- and low-altitude sister species in the family Aegithalidae: the black-browed bush tit (Aegithalos bonvaloti) and the sooty bush tit (Aegithalos fuliginosus), respectively. We performed experiments on rHbs for these two species because we did not have adequate quantities of blood from wild-caught birds to purify native Hbs.

Hb Isoform Composition. During adulthood, most bird species express two structurally distinct Hb isoforms that incorporate different α-chain subunits but share identical β-chains. The major HbA isoform (α²β²) incorporates products of the α²-globin gene, and the minor HbD isoform (α²β¹) incorporates products of the α²-globin gene (18–20). We used isoelectric focusing (IEF) analysis to characterize Hb isoform composition in the red blood cells of each of the focal species. These analyses revealed that the minor HbD isoform accounted for 20–30% of total Hb (Table S1), consistent with data from the majority of other passerine taxa examined to date (15, 17, 19–21).

Oxygenation Properties of HbA and HbD Isoforms. We purified HbA and HbD isoforms from red cell lysates of select specimens with known α²-, α¹-, and β¹-globin genotypes. We then measured the oxygenation properties of purified HbA and HbD solutions in the presence and absence of two main allosteric cofactors that regulate Hb-O₂ affinity: Cl⁻ ions (added as 0.1 M KCl) and inositol hexaphosphate (IHP, a chemical analog of inositol pentaphosphate). Experimental measurements on purified Hb samples from all species revealed that the HbD isoform exhibited a uniformly higher O₂ affinity than HbA, both in the absence (“stripped”) and presence of allosteric cofactors (Fig. 1A–C and Table S2). This is indicated by the lower values of P50 (the P0₂ at which heme is 50% saturated) for HbD relative to HbA. This consistent isoform differentiation in Hb–O₂ affinity suggests that the up-regulation of HbD could provide a ready means of optimizing blood oxygenation properties in response to changes in O₂ availability. However, the absence of altitude-related differences in the HbA/HbD ratio among species (Table S1) indicates that regulatory adjustments in red cell Hb isoform composition do not play an important role in hypoxia adaptation. Both HbA...
and HbD exhibited cooperative O₂ binding, as indicated by Hill coefficients at half-saturation \((n_H) > 2\) in the presence of anions (Table S2).

**Evolved Changes in Hb-\(O_2\) Affinity and Its Structural Basis.** Experimental results for the nine focal taxa revealed that highland species generally have higher Hb-\(O_2\) affinities in comparison with their close lowland relatives. Within the genus *Parus*, the high-altitude *Parus humilis* and the low-altitude *Parus minor* provide the basis for an especially informative comparison because the two closely related species have completely nonoverlapping elevational ranges (Fig. S1). The experiments revealed that the HbA and HbD isoforms of *Parus humilis* exhibited significantly higher O₂ affinities than the corresponding isoforms of *Parus minor*, both in the absence and presence of allosteric cofactors (Fig. L4 and Table S2). In the presence of Cl⁻ and IHP, \(P_{50}\) values for HbA and HbD of *Parus humilis* were lower than those of *Parus minor* by factors of 2.0-fold and 1.6-fold, respectively (Table S2). The difference in HbA-\(O_2\) affinity is attributable to the independent or combined effects of 17 substitutions: 12 in \(\alpha\) and 5 in \(\beta\) (Fig. 2). The qualitatively similar species difference in HbD-\(O_2\) affinity (Fig. L4) is associated with eight substitutions: three in \(\alpha\) in addition to the above-mentioned \(\beta\) substitutions (Fig. 2).

Within the Poecile/Lophophanes clade, the high-altitude *Lophophanes dichrous* and the low-altitude *Poecile palustris* also have nonoverlapping elevational ranges (Fig. S1). Our experiments revealed that HbA and HbD of *Lophophanes dichrous* have significantly higher O₂ affinities than the corresponding isoforms of the low-altitude *Poecile palustris* (Fig. 1B and Table S2). Species differences in HbA-\(O_2\) affinity are attributable to the independent or combined effects of five substitutions (three in \(\alpha\), two in \(\beta\)) (Fig. 2). A similar difference in HbD-\(O_2\) affinity is associated with five substitutions (three in \(\alpha\) in addition to the above-mentioned \(\beta\) substitutions) (Fig. 2). Among the \(\alpha\) substitutions, it is notable that *Lophophanes dichrous* shares the same \(\alpha^A34T\) substitution as *Parus humilis* (Fig. 2). This site is located at an intradimer \(\alpha_1\beta_2/\alpha_2\beta_1\) contact surface. Mutations at such sites are known to affect Hb-\(O_2\) affinity and quaternary structural stability (22–26). Homology modeling analysis suggests that the replacement of Ala with Thr at \(\alpha^A34\) introduces four new hydrogen bonds between opposing \(\alpha\) and \(\beta\) chain subunits of the oxygenated R conformation; the same intersubunit \(\alpha_1\beta_2/\alpha_2\beta_1\) interface is not altered appreciably in the deoxygenated T conformation (Fig. 3 and Table S3). The allosteric equilibrium between the oxygenated (R) and deoxygenated (T) conformations of the Hb tetramer is a primary determinant of Hb-\(O_2\) affinity. The differential stabilization of oxygenated Hb therefore increases \(O_2\) affinity by shifting the T=R allosteric equilibrium in favor of the high-affinity R state.

Within the genus *Periparus*, experiments revealed no appreciable differentiation in overall Hb-\(O_2\) affinity between the high- and low-altitude subspecies of *Periparus ater* revealed no differentiation in overall Hb-\(O_2\) affinity, all species-level comparisons between close relatives were consistent in that the highland species always had higher Hb-\(O_2\) affinities than the lowland species. In each case, the evolved changes were exclusively attributable to an increase in intrinsic \(O_2\) affinity (as revealed by \(P_{50}\) values for stripped Hbs), as there were no appreciable differences in sensitivity to allosteric cofactors (Table S2).

In principle, altitude-related changes in Hb-\(O_2\) affinity could be attributable to derived increases in Hb-\(O_2\) affinity in high-altitude species, derived reductions in low-altitude species, or a combination of both. These alternatives can be distinguished by deciphering the character polarity of causative amino acid substitutions. For example, if a given high-altitude species has evolved a derived increase in Hb-\(O_2\) affinity relative to its low-altitude sister species, then the evolved change in protein function must be attributable to one or more substitutions where the derived amino acid state is fixed in the high-altitude species. As described below, we estimated globin gene trees to identify the phylogenetic intervals in which potentially causative amino acid substitutions occurred in the history of each high-altitude lineage.

**Genealogical Discordance of Globin Genes.** To infer the polarity of character state change at substituted sites in HbA and HbD, we estimated DNA-based phylogenies of the \(\alpha^A\), \(\alpha^B\), and \(\beta^A\)-globin genes from all 17 species. Estimated phylogenies revealed high levels of genealogical discordance among loci (Fig. S2), likely reflecting a history of incomplete lineage sorting and/or introgressive hybridization.
Molecular Convergence and Parallelism. It is critically important to account for genealogical discordance when testing for evidence of molecular parallelism and convergence (30, 31). If amino acid substitutions in the \( \alpha^A \)- and \( \alpha^D \)-globin genes were mapped onto the branches of the species tree (Fig. S1) rather than the appropriate gene trees (Fig. S2A and B), it would create the false appearance that multiple substitutions had occurred independently in different lineages.

Estimates of gene tree topologies for the \( \alpha^A \), \( \alpha^D \), and \( \beta^D \)-globin genes permit an assessment of the true prevalence of molecular convergence and parallelism in our set of focal taxa. Some degree of homoplasy (due to convergence, parallelism, or mutational reversion) is expected due to chance alone. The key question concerns the number of convergent or parallel substitutions that actually contributed to convergent increases in \( \text{Hb}_{O_2} \) affinity in different high-altitude taxa; this applies to substitutions in \( \text{HbA} \) and \( \text{HbD} \) of both \( \text{Parus humilis} \) and \( \text{Lophophanes dichrous} \), \( \text{HbD} \) of \( \text{Periparus rubidiventris} \), and \( \text{HbA} \) of \( \text{Aegithalos bonvaloti} \). The sequence data revealed that few derived amino acid states are shared between high-altitude taxa to the exclusion of lowland taxa. Within the set of high-altitude parid and aegithalid taxa that we examined, it appears that there is only one true parallel substitution that is associated with convergent increases in \( \text{Hb}_{O_2} \) affinity: \( \alpha^A \)-A34T in the \( \alpha^A \)-globin orthologs of \( \text{Parus humilis} \) and \( \text{Lophophanes dichrous} \) (Fig. 2).

Functional Test of Adaptive Parallelism. The \( \alpha^A \)-globin gene tree shown in Fig. S2A indicates that the shared, derived Thr at \( \alpha^A \)34 was not present in the ancestral amino acid at that position of codon 34 (31, 32). This enabled us to test the effect of the \( \alpha^A \)34T mutation on an evolutionarily relevant genetic background. Ancestral amino acid states were estimated with a high level of statistical confidence, as site-specific posterior probabilities averaged 0.997 for the \( \alpha \)-chain and 1.000 for the \( \beta \)-chain. The reconstructed ancestral \( \text{Hb} \) was estimated to possess Ala at \( \alpha^A \)34 with a posterior probability of 0.988. Overall, the \( \text{AncParidae} \) \( \text{Hb} \) differed from the wild-type \( \text{Hb} \)s of \( \text{Parus humilis} \) and \( \text{Lophophanes dichrous} \) at 12 and 6 amino acid sites, respectively (Fig. S3).

In vitro functional tests on the purified \( \text{Hbs} \) confirmed that \( \text{AncParidae} \) \( \text{Hb} \) exhibited a significantly lower \( \text{Hb}_{O_2} \) affinity than the wild-type \( \text{Hbs} \) of both \( \text{Parus humilis} \) and \( \text{Lophophanes dichrous} \), indicating that each of the two high-altitude species independently evolved derived increases in \( \text{Hb}_{O_2} \) affinity due to the effects of one or more lineage-specific substitutions. Site-directed mutagenesis experiments revealed that \( \alpha^A \)34T produced a significant increase in intrinsic \( \text{Hb}_{O_2} \) affinity on the \( \text{AncParidae} \) background, and this affinity difference persisted in the presence of \( \text{Cl}^- \) and IHP (Fig. 4). This result indicates that the parallel \( \alpha^A \)34T substitutions in \( \text{Parus humilis} \) and \( \text{Lophophanes dichrous} \) contributed to convergent increases in \( \text{Hb}_{O_2} \) affinity in each of the two high-altitude species, but the effect of \( \alpha^A \)34T alone was not sufficient to completely recapitulate the evolved change in \( \text{O}_2 \) affinity in either species. Thus, additional amino acid substitutions in the \( \alpha \)- and/or \( \beta \)-globins of both \( \text{Parus humilis} \) and \( \text{Lophophanes dichrous} \) must have contributed to the derived increases in \( \text{Hb}_{O_2} \) affinity. Interestingly, although \( \alpha^A \)34T produced a significant increase in \( \text{O}_2 \) affinity on the \( \text{AncParidae} \) background, reversion of this mutation to the ancestral state (\( \alpha^A \)T34A) did not produce significant reductions in \( \text{Hb}_{O_2} \) affinity on the wild-type backgrounds of either \( \text{Parus humilis} \) or \( \text{Lophophanes dichrous} \) (Fig. 4). Asymmetry in the effects of forward mutations on ancestral backgrounds and reverse mutations on derived backgrounds, a phenomenon documented in numerous protein-engineering studies, is attributable to epistatic interactions between the focal mutations and residues at other substituted sites that distinguish the two backgrounds (31–36).

The Possible Role of Mutation Bias in Promoting Parallel Substitutions. Intriguingly, the parallel \( \alpha^A \)34T substitutions in both \( \text{Lophophanes dichrous} \) and \( \text{Parus humilis} \) are attributable to nonsynonymous substitutions at a \( \text{CpG} \) dinucleotide. In both species, replacement of the ancestral Ala for Thr at \( \alpha^A \)34 was caused by \( \text{CpG} \rightarrow \text{CpA} \) mutations in the first codon position. Depending on the methylation status of the cytosine, transition mutations at \( \text{CpG} \) sites are expected to occur at a much higher rate than non-\( \text{CpG} \) point mutations at the same nucleotide position (37–40). This suggests the hypothesis that the \( \alpha^A \)34T mutation may be especially likely to contribute to evolved increases in \( \text{Hb}_{O_2} \) affinity simply because the underlying \( \text{CpG} \rightarrow \text{CpA} \) transition mutation will occur at a higher rate than nonsynonymous mutations at non-\( \text{CpG} \) sites that produce similar affinity-enhancing effects. If adaptation is mutation-limited, an increase in the rate of mutation to a beneficial allele results in a commensurate increase in the allele’s probability of fixation (41–44). The extent to which evolution is mutation-limited in natural populations is not generally known, but a growing body of evidence suggests that mutation bias is an important cause of substitution bias and parallelism in adaptive protein evolution (15, 45, 46).

An examination of \( \alpha^A \)-globin nucleotide sequences in a phylogenetically diverse set of passerines revealed that the \( \text{CpG} \) dinucleotide involving the third position of codon 33 and the first position of codon 34 is clearly the ancestral state (and, hence, Ala is the ancestral amino acid at \( \alpha^A \)34) (Fig. S4). Thus, if the parallel \( \alpha^A \)34T substitutions in \( \text{Lophophanes dichrous} \) and \( \text{Parus humilis} \) were partly attributable to mutation bias, it is a bias that is not unique to parid species. On the basis of mutational accessibility alone, there...
is consistent with comparative data from other high-altitude birds (14–17, 21, 29) and demonstrates that there are multiple possible ways of evolving an increased Hb–O₂ affinity without sacrificing allosteric regulatory capacity. In mammals, by contrast, evolutionary changes in Hb–O₂ affinity often involve changes in responsiveness to CO and/or organic phosphates (47–52).

Despite the pervasive convergence in Hb–O₂ affinity (and convergence in the underlying functional mechanisms) among high-altitude avian taxa, the limited amount of convergence and parallelism at the amino acid level demonstrates that changes in Hb–O₂ affinity can be produced by numerous possible amino acid substitutions. The analysis of sequence variation in our set of parid and aegithalid species revealed several parallel amino acid substitutions, only one of which contributed to convergent increases in Hb–O₂ affinity in different high-altitude tit species: α²A34T in Parus humilis and Lophophanes dichrous. Most parallel substitutions were not uniquely associated with convergent increases in Hb–O₂ affinity in highland lineages. In studies of genetic convergence and parallelism, experimental testing of individual substitutions is necessary to distinguish between signal (replicated substitutions that contribute to adaptive convergence in phenotype) and noise (repli- cated substitutions that have no effect on the selected phenotype).

In broader comparisons involving other avian taxa, we documented one additional example in which parallel substitutions contributed to convergent increases in Hb–O₂ affinity in different high-altitude species: the α²P119A substitution in Aythya bonvaloti (Aegithalidae: Passeriformes) and the distantly related bar-headed goose, Anser indicus (Anatidae: Anseriformes). Results of site-directed mutagenesis experiments confirm that the parallel α²P119A substitutions have significant affinity-enhancing effects in the Hbs of both species (Fig. 1D and Table S2 (29)). The α²P119A mutation also produces a similar affinity-enhancing effect in human Hb (27). All else being equal with respect to effect sizes and mutational pleiotropy, allelic variants that have additive effects across a wide range of genetic backgrounds can be expected to make more consistent contributions to convergent adaptation than those with context-dependent effects. Overall, we did not observe any striking clade-specific patterns of parallelism that could be interpreted as evidence that the high-altitude parid or aegithalid species had hit upon adaptive solutions that were not equally accessible to other high-altitude avian taxa.

Methods

Sample Collection. We collected a total of 162 bird specimens (representing 17 species) from high-altitude localities on the Qinghai-Tibet Plateau and the mountains of Southwest China and in low-altitude localities throughout eastern China. The collection of all bird specimens was authorized by the Forestry Administrations of Qinghai Province and Xizang Autonomous Region and was in compliance with the National Wildlife Conservation Law of China. All birds were handled in accordance with regulations of the Animal Experimental and Medical Ethics Committee of the Institute of Zoology, Chinese Academy of Sciences. Details are provided in SI Methods.

Hb Isoform Composition and Sequence Variation. To characterize Hb isoform composition in the red blood cells of wild-caught birds, we separated native Hbs by means of IEF and used densitometric measurements to quantify isoform abundance (20). Details regarding cloning and sequencing protocols are provided in SI Methods. All sequences have been deposited in GenBank (accession nos. MG772099–MG772439).

Protein Purification and in Vitro Analysis of Hb Function. Using hemolysates of bird specimens with known globin genotypes, we isolated and purified the HbA and HbD isoforms by means of anion-exchange FPLC using a HiTrap Q HP column (GE Healthcare). Details regarding O₂-binding experiments are provided in SI Methods.

Vector Construction, Mutagenesis, and Recombinant Expression. The α²- and β²-globin sequences were synthesized in accordance with Escherichia coli codon preferences. The α²-globin gene cassette was cloned into a custom pGEM vector system, and recombinant Hb expression was carried out in the E. coli JM109 (DE3) strain (25, 53). Details regarding mutagenesis and protein purification are provided in SI Methods.

Fig. 4. O₂ affinities of recombinantly expressed Hbs representing wild-type genotypes of Parus humilis and Lophophanes dichrous and the reconstructed genotype of their most recent common ancestor AncParidae. Site-directed mutagenesis experiments revealed that α²A34T produces a significant affinity-enhancing effect on the ancestral background (AncParidae), as indicated by the reduction in P50 but does not completely recapitulate the evolved increases in Hb–O₂ affinity in Parus humilis or Lophophanes dichrous. Cross-hatched and solid bars show P50 values in the absence (stripped) and presence of anionic cofactors, respectively. Mutational reversions (α²A34T) on the wild-type backgrounds of Parus humilis and Lophophanes dichrous did not produce a symmetrical, affinity-reducing effect. The asterisk denotes a statistically significant difference (P < 0.05) in the presence of KCl + IHP.

is no reason to think that α²A34T substitutions were more likely to contribute to Hb adaptation in parid species than in other passerines.

Broader Patterns of Convergence and Parallelism Involving High-Altitude Birds from the Andes and the Qinghai-Tibet Plateau. An expansion of the comparative analysis to include other high-altitude birds revealed several convergent and parallel substitutions shared with parid and aegithalid species from the Qinghai-Tibet Plateau. The number of such substitutions is reduced significantly if we restrict our attention to shared, derived replacements that are associated with increases in Hb–O₂ affinity in high-altitude lineages. In the Andean birds, convergent and parallel substitutions were not uncommon, but site-directed mutagenesis experiments revealed that a small fraction of such substitutions (i.e., N/G83S, A86S, D94E, and A116S in β²-globin) actually contributed to convergent increases in Hb–O₂ affinity in two or more high-altitude species (16, 17). These same sites were all invariant in our sample of parid and aegithalid species.

Conclusions

We documented that high-altitude parid and aegithalid species from the Qinghai-Tibet Plateau have typically evolved increased Hb–O₂ affinities in comparison with lowland relatives. In broader comparisons involving other avian taxa, we documented one additional example in which parallel substitutions contributed to convergent increases in Hb–O₂ affinity in different high-altitude species: the α²P119A substitution in Aythya bonvaloti (Aegithalidae: Passeriformes) and the distantly related bar-headed goose, Anser indicus (Anatidae: Anseriformes). Results of site-directed mutagenesis experiments confirm that the parallel α²P119A substitutions have significant affinity-enhancing effects in the Hbs of both species (Fig. 1D and Table S2 (29)). The α²P119A mutation also produces a similar affinity-enhancing effect in human Hb (27). All else being equal with respect to effect sizes and mutational pleiotropy, allelic variants that have additive effects across a wide range of genetic backgrounds can be expected to make more consistent contributions to convergent adaptation than those with context-dependent effects. Overall, we did not observe any striking clade-specific patterns of parallelism that could be interpreted as evidence that the high-altitude parid or aegithalid species had hit upon adaptive solutions that were not equally accessible to other high-altitude avian taxa.

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Hb Isoform Composition and Sequence Variation. To characterize Hb isoform composition in the red blood cells of wild-caught birds, we separated native Hbs by means of IEF and used densitometric measurements to quantify isoform abundance (20). Details regarding cloning and sequencing protocols are provided in SI Methods. All sequences have been deposited in GenBank (accession nos. MG772099–MG772439).

Protein Purification and in Vitro Analysis of Hb Function. Using hemolysates of bird specimens with known globin genotypes, we isolated and purified the HbA and HbD isoforms by means of anion-exchange FPLC using a HiTrap Q HP column (GE Healthcare). Details regarding O₂-binding experiments are provided in SI Methods.

Vector Construction, Mutagenesis, and Recombinant Expression. The α²- and β²-globin sequences were synthesized in accordance with Escherichia coli codon preferences. The α²-globin gene cassette was cloned into a custom pGEM vector system, and recombinant Hb expression was carried out in the E. coli JM109 (DE3) strain (25, 53). Details regarding mutagenesis and protein purification are provided in SI Methods.
Phylogenetic Analysis. We used complete exon and intron sequences to estimate phylogenies of each globin gene. Details regarding alignments, sub-study models, and ancestral reconstructions are provided in SI Methods.

Structural Modeling. We modeled structures of avian Hbs using MODELER ver. 9.19 (54). We used Protein Data Bank (PDB) ID 1hhq and 3hhb as mate phylogenies of each globin gene. Details regarding alignments, sub-study Phylogenetic Analysis.


