Biochemical paedomorphosis and genetic assimilation in the hypoxia adaptation of Tibetan antelope

Anthony V. Signore and Jay F. Storz*

University of Nebraska, School of Biological Sciences, Lincoln, Nebraska, 68588

*Correspondence to: jstorz2@unl.edu
Abstract

Developmental shifts in stage-specific gene expression can provide a ready mechanism of phenotypic change by altering the rate or timing of ontogenetic events. We discovered that the high-altitude Tibetan antelope (*Panthelops hodgsonii*) has evolved an adaptive increase in blood–O₂ affinity by truncating the ancestral ontogeny of globin gene expression such that a high-affinity juvenile hemoglobin isoform (isoHb) completely supplants the lower-affinity isoHb that is expressed in the adult red blood cells of other bovids. This juvenilization of blood properties represents a canalization of an acclimatization response to hypoxia that has been well-documented in adult goats and sheep. We also discovered the genomic mechanism underlying this regulatory isoHb switch, revealing how a reversible acclimatization response became genetically assimilated as an irreversible adaptation to chronic hypoxia.
**Introduction**

When members of multigene families are developmentally regulated, shifts in the stage-specific expression of individual genes can provide a ready mechanism of phenotypic change by altering the rate or timing of ontogenetic events (heterochrony). For example, retained activity of early-expressed genes in later stages of ontogeny can result in the retention of juvenile characters into adulthood, a well-documented developmental mechanism of phenotypic evolution (1, 2). In extreme cases, deceleration of development can produce a truncation of the ancestral ontogeny, resulting in the juvenilization of the adult-expressed phenotype, a phenomenon known as paedomorphosis.

In jawed vertebrates, the subfamilies of genes that encode the α- and β-type subunits of tetrameric hemoglobin (Hb) are developmentally regulated such that structurally and functionally distinct αβ2 Hb isoforms (isoHbs) are expressed during different ontogenetic stages. During mammalian development, different pre- and postnatally expressed isoHbs have evolved different oxygenation properties and perform distinct O2–scavenging/O2–transport tasks during different ontogenetic stages (3–6). Genetically based shifts in stage-specific isoHb expression could therefore provide a heterochronic mechanism of evolutionary change in respiratory gas transport and aerobic metabolism. Similarly, in humans, hereditary persistence of fetal Hb alleviates the severity of thalassemias and other pathologies affecting the synthesis or stability of adult Hb (7).

Given that prenatally expressed isoHbs of eutherian mammals exhibit substantially higher O2–affinities than adult-expressed isoHbs, and given that an increased Hb–O2 affinity is generally beneficial under conditions of severe hypoxia due to the importance of safeguarding arterial O2 saturation (6, 8-11), the retention of early isoHb expression into adulthood could provide an effective mechanism of adaptation to chronic O2 deprivation. Consistent with this
hypothesis, when adult goats and sheep are exposed to acute hypoxia, they upregulate a juvenile isoHb at the expense of the normal adult isoHb (12, 13). Here we report the discovery of a canalized version of this response in the high-altitude Tibetan antelope, *Panthelops hodgsonii* (Artiodactyla: Bovidae), a champion among mammals in aerobic exercise performance under hypoxia. This species is endemic to the Tibetan Plateau and lives at altitudes 3600-5500 m above sea level. At an altitude of 5500 m, the partial pressure of O2 ($P_{O_2}$) is roughly half the value at sea level, a level of hypoxia that severely compromises aerobic exercise performance in humans and most other mammals (14-15). Remarkably, however, at these altitudes Tibetan antelope can sustain running speeds of >70 km/h over distances of >100 km (16).

In addition to documenting the phenotypic consequences of developmentally displacing the low-affinity adult isoHb with a higher-affinity juvenile isoHb – a form of biochemical paedomorphosis – we also discovered the genomic mechanism by which the upregulation of the juvenile isoHb became canalized in Tibetan antelope. Specifically, we document how a reversible acclimatization response to acute hypoxia – as observed in modern-day sheep and goats – became genetically assimilated as an irreversible adaptation to chronic hypoxia.

**Results and Discussion**

We characterized the genomic organization of globin genes in Tibetan antelope and other bovid artiodactyls using published genome assemblies (17). Among mammals, bovid artiodactyls are unusual in that the entire $\beta$-globin gene cluster has undergone multiple rounds of *en bloc* duplication involving the same set of pre- and post-natally expressed $\beta$-type globin genes (Fig. 1) (18-20). Cows (*Bos taurus*) possess two duplicated gene blocks, each containing separate paralogs of the $\beta$-globin gene, $\beta^A$ and $\beta^F$, in the 5’ and 3’ blocks, respectively (Fig. 1). As with
other eutherian mammals, the product of $\beta^A$ is incorporated into an adult-expressed isoHb, HbA ($\alpha_2\beta^A_2$), whereas the $\beta^F$ gene has been recruited for prenatal expression, and is incorporated into a fetal isoHb, HbF ($\alpha_2\beta^F_2$) (21). Goats (Capra hircus) and sheep (Ovis aries) possess an additional gene block at the 5’ end of the cluster that contains a third $\beta$-globin paralog, $\beta^C$ (Fig. 1) (18-20). Whereas the $\beta^A$ and $\beta^F$ genes in goats and sheep have retained the same developmental expression profiles as their respective orthologs in cow, the $\beta^C$ gene has been recruited for a new ontogenetic stage of expression during the first few months of neonatal life, and its product is incorporated into a juvenile isoHb, HbC ($\alpha_2\beta^C_2$) (22).

The $\beta$-globin gene cluster of Tibetan antelope appears superficially similar to that of cow in terms of gene content (Fig. 1), suggesting that the Tibetan antelope inherited the same pair of $\beta^A$- and $\beta^F$-containing gene blocks. The alternative hypothesis is that Tibetan antelope inherited the additional *en bloc* duplication observed in goats and sheep, but one of the triplicated gene blocks was secondarily deleted, in which case the sole remaining pair of $\beta$-globin genes would be represented by one of three possible combinations: $\beta^A + \beta^F$ (a reversion to the ancestral gene complement observed in cow), $\beta^C + \beta^A$, or $\beta^C + \beta^F$ (Fig. 2A-C). Either of the latter two combinations would implicate a novel isoHb profile that is not observed in other bovid taxa. To distinguish among these three alternative scenarios, we estimated the phylogeny of bovid $\beta^C$, $\beta^A$, and $\beta^F$ genes and the pair of Tibetan antelope $\beta$-globin paralogs. Estimated phylogenies (Fig. 2D, supplementary Fig. S1) clearly demonstrate that the 5’ and 3’ $\beta$-globin genes of Tibetan antelope are orthologous to bovid $\beta^C$ and $\beta^F$, respectively, consistent with the scenario illustrated in Fig. 2B. This result indicates that Tibetan antelope inherited the triplicated set of $\beta^C$, $\beta^A$, and $\beta^F$-containing gene blocks observed in goats and sheep (Fig. 1), and that the 5’ gene block containing $\beta^A$ was secondarily deleted. This phylogenetic inference is unambiguously
corroborated by patterns of conserved synteny and pairwise sequence matches (Fig. 3), as the β^C- and β^F-containing gene blocks of goat and sheep match the 5’ and 3’ gene blocks in Tibetan antelope. This comparative genomic analysis revealed that a ~45 kb region of the Tibetan antelope β-globin gene cluster was deleted – a gene region that contained the ortholog of the β^A gene that encodes the β-chain of adult Hb in bovids and all other mammals. Deletion of the adult β^A gene in the ancestor of Tibetan antelope effectively truncated the ancestral ontogeny of globin gene expression, such that juvenile HbC completely supplanted HbA in adult red blood cells. Thus, blood–O_2 transport in Tibetan antelope has been juvenilized relative to the ancestral phenotype of adult bovids. To examine the effects of this paedomorphic change, we measured the oxygenation properties of purified recombinant Hb from Tibetan antelope and purified native Hbs from adult specimens of 10 other bovid species (Fig. 4; Table S1). The adult red cells of these other taxa contain HbA alone or in combination with HbC as a minor component. We measured the Hb–O_2 affinity of purified total Hb from each bovid species in both the absence (stripped) and presence of 100 mM Cl^- (in the form of KCl). The stripped treatment provides a measure of intrinsic Hb–O_2 affinity whereas the +KCl treatment provides a measure that is relevant to in vivo conditions in bovid red cells, as Cl^- ions are the principal allosteric regulators of Hb–O_2 affinity (i.e., heme reactivity is modulated oxygenation-linked binding of Cl^- ions at sites remote from the heme iron)(6, 21, 23). Results of our in vitro experiments revealed that Hb of Tibetan antelope has a substantially higher O_2–affinity than that of all other bovid taxa (Fig. 4; Table S1). Hbs of all taxa were similarly responsive to Cl^-, as the average P_{50} (the PO_2 yielding 50% Hb–O_2 saturation) was 27.1% higher (i.e., Hb–O_2 affinity was lower) in the +KCl treatment (Table S1).
As a follow-up experiment, we isolated and purified HbC and HbA from two of the bovid species expressing both components and we measured isoHb-specific O₂–binding properties to determine how blood–O₂ affinity would be affected by elimination of the major HbA isoHb (as would occur with the deletion of the β² gene, leaving juvenile HbC as the sole-remaining isoHb). There was very little among-species variation in the measured O₂–affinities of either juvenile HbC or adult HbA (Fig. S2; Table S1), but the O₂–affinity of HbC exceeded that of HbA by a consistent margin (average 10.6 torr) in all species (Fig. S2; Table S1). Moreover, O₂–affinity of HbC alone was always substantially higher than that of the composite HbA + HbC mixture (with the two isoHbs present in their naturally occurring relative abundance) (Fig. S2; Table S1), reflecting the fact that the lower affinity HbA is always present as the major isoHb in adult red cells (average HbA/HbC ratio = ~80:20).

The higher Hb–O₂ affinity of Tibetan antelope relative to that of other bovid species is entirely attributable to a difference in isoHb composition: they only express the high-affinity HbC instead of jointly expressing HbA and HbC (with the lower-affinity HbA present as the major isoHb). To infer the direction of evolutionary change in isoHb-specific O₂–affinities, and to reconstruct the phenotypic effect of deleting β²-globin (thereby leaving HbC as the solo-expressed isoHb in adult red cells), we reconstructed the ancestral bovid β² and β⁵ genes as well as their single-copy, pre-duplication progenitor (β²⁵) (Fig. 5; Fig. S3). Triangulated comparison of O₂–affinities of the three recombinantly expressed ancestral isoHbs, AncHb-β², AncHb-β⁵, and AncHb-β²⁵ (all of which had identical α-chains), revealed that the juvenile AncHb-β⁵ evolved a slight increase in O₂–affinity relative to the estimated ancestral state (represented by AncHb-β²⁵), whereas adult HbA evolved a slight reduction in O₂–affinity (Fig. 5). These data indicate that if HbA and HbC were present in a 80:20 ratio in the red cells of the Tibetan...
antelope ancestor (as in extant bovids), the deletion of βA-globin and the consequent elimination of HbA from the ‘HbA + HbC’ composite mixture would result in a 13.5% increase in Hb–O₂ affinity in the presence of 100 mM Cl⁻ (P₅₀ decreased from 18.5 to 16.0 torr).

The derived blood phenotype of Tibetan antelope is consistent with the theoretical expectation that an increased Hb–O₂ affinity is adaptive under conditions of severe hypoxia (especially in highly athletic species), and is consistent with patterns observed in other high-altitude mammals and birds that maintain especially high rates of aerobic metabolism (24-32). In other case studies of high-altitude vertebrates, evolved increases in Hb–O₂ affinity have been traced to one or more amino acid substitutions in the α- and/or β-chain subunits of the α₂β₂ Hb tetramer (6, 30-31). Here we document a unique case in which an evolved change in Hb–O₂ affinity has been accomplished via a heterochronic shift in globin gene expression, such that a high-affinity, juvenile isoHb supplants the lower-affinity, adult isoHb. This juvenilization of blood properties represents a novel mode of biochemical adaptation and highlights the utility of heterochrony as an adaptive mechanism, whereby “…the existing channel of ordinary ontogeny already holds the raw material in a particularly effective state for evolutionary change” (2).

There has been debate in the literature regarding the relative importance of regulatory vs. coding changes in genetic adaptation (33) and phenotypic evolution in general (34-35). In the case of Tibetan antelope, the evolved increase in Hb–O₂ affinity was caused by an unusual combination of regulatory and structural changes. Specifically, a drastic regulatory switch in protein isoform expression (via truncation of the ancestral ontogeny of globin gene regulation) was caused by a large-scale chromosomal deletion, highlighting the surprising diversity of genetic mechanisms and substrates of phenotypic evolution.
Materials and Methods

Collection of blood samples

Frozen erythrocytes from 10 bovid species were provided by the San Diego Zoo Institute for Conservation Research (Uniform Biological Material Transfer Agreement BR2017063). This included six species in the subfamily Caprinae (*Capra aegagrus cretica, Capra caucasica caucasica, Ovis nivicola, Capra nubiana, Ovis orientalis musimon* and *Ovis canadensis nelsoni*), two species in the subfamily Alcelaphinae (*Damaliscus pygargus phillipsi* and *Connochaetes gnou*), and one species of each from Hippotraginae (*Addax nasomaculatus*) and Bovinae (*Bos gaurus*).

Sequencing of bovid globin genes

RNA was extracted from ~100 ul of flash frozen erythrocytes using an RNeasy Universal Plus Mini Kit (Qiagen). cDNA was synthesized from freshly prepared RNA using Superscript IV Reverse transcriptase (Invitrogen). Gene specific primers were used to amplify the a- and b-type globin transcripts. PCR reactions were conducted using 1 ml of cDNA template in 0.2 ml tubes containing 25 µl of reaction mixture (0.5 µl of each dNTP (2.5 mM), 2.5 µl of 10x Reaction Buffer (Invitrogen), 0.75 µl of 50 mM MgCl₂, 1.25 µl of each primer (10 pmol/µl), 1 µl of Taq polymerase (Invitrogen) and 16.75 µl of ddH₂O), using an Eppendorf Mastercycler® Gradient thermocycler. Following a 5-min denaturation period at 94°C, the desired products were amplified using a cycling profile of 94°C for 30 sec; 53-65°C for 30 sec; 72°C for 45 sec for 30 cycles followed by a final extension period of 5 min at 72°C. Amplified products were run on a 1.5% agarose gel and bands of the correct size were subsequently excised and purified using Zymoclean Gel DNA recovery columns (Zymo Research). Gel-purified PCR products were ligated into pCR™4-
TOPO® vectors using a TOPO™ TA Cloning™ Kit and were then transformed into One Shot™ TOP10 Chemically Competent *E. coli* (Thermo Fisher Scientific). Three to six transformed colonies were cultured in 5 ml of LB medium and plasmids were subsequently purified with a GeneJET Plasmid Miniprep kit (Thermo Fisher Scientific). Purified plasmids were sequenced by Eurofins Genomics (Fig. S3).

**Sequence analyses**

Genomic sequences containing the complete α- and β-globin gene clusters for the domestic goat (*Capra hircus*), sheep (*Ovis aries*), cow (*Bos taurus*), and Tibetan antelope (*Panthelops hodgsonii*) were mined from GenBank (Table S2). Sequence identity between bovid chromosomal regions containing the β-globin gene clusters was calculated using Blastn and conserved synteny was visualized using Easyfig 2.1 (36). Coding sequences of α- and β-globin genes were extracted from genomic and cDNA sequences mined from GenBank (Table S2) and combined with cDNA sequences collected above (Fig. S3). These sequences were aligned using MUSCLE (37) and were used for phylogenetic tree construction. The best fitting codon substitution model and initial tree search were estimated using IQ-TREE with the options -st CODON, -m TESTNEW, -allnni, and -bnni (38, 39). Initial trees were then subjected to 1000 ultrafast bootstrap replicates (40). Bootstrap consensus trees (Figs. S1A, S1B) were used to estimate ancestral globin sequences using IQ-TREE with the option -asr (Figs. S1C, S3). As bovid βC-globins are truncated by 9bp (relative to βA), the ancestral reconstruction of indels in the β-globin gene tree was performed by FastML (41).

**Protein purification**
Blood samples (~200µl) were added to a 5x volume of ice-cold water and incubated on ice for 30 minutes to lyse the red blood cells. Samples were centrifuged at 20,000 x g for 10 minutes to remove cell debris. Buffer was added to the supernatants to a final concentration of 0.01 M HEPES/0.2 M NaCl (pH 7.4) and passed through a PD-10 desalting column (GE Healthcare) equilibrated with 25 ml of 0.01 M HEPES/0.5mM EDTA (pH 7.2) to remove intracellular cofactors. Desalted lysates were loaded onto a HiTrap SP cation exchange column (GE Healthcare) and isoHbs were eluted using a linear pH gradient (0.01 M HEPES/0.5mM EDTA, pH 7.2 – 7.7). For each species, a subsample of each isoHb was pooled to create a “Total Hb” solution. Each Hb solution was then desalted using a PD-10 column (GE Healthcare) equilibrated with 0.01 M HEPES/0.5mM EDTA (pH 7.4) and eluates were concentrated using Amicon Ultra-4 Centrifugal Filter Units (Millipore).

**Measuring O₂–binding properties of purified Hbs**

O₂–equilibrium curves for purified Hb solutions (0.1 mM hemoglobin in 0.1 M HEPES/0.05 M EDTA buffer, pH 7.4) were measured at 37°C using a Blood Oxygen Binding System (Loligo Systems). O₂–equilibrium curves were measured in the absence (stripped) and presence of chloride ions (0.1 M KCl). Each Hb solution was sequentially equilibrated with three to five different O₂ tensions (PO₂) at saturation levels between 30 to 70% while the absorbance was continually monitored at 430 nm (deoxy peak) and 421 nm (oxy/deoxy isobestic point)(42-44). Hill plots (log[fractional saturation/[1-fractional saturation]] vs logPO₂) constructed from these measurements were used to estimate the PO₂ at half saturation (P₅₀) and the cooperativity coefficient (n₅₀) from the χ-intercept and slope of these plots, respectively. O₂–equilibrium
curves for each Hb solution were measured in triplicate and $P_{50}$ is reported as mean ± s.e.m. (Table S1).

**Construction of Hb expression vector**

Globin sequences for domestic goat, Tibetan antelope, as well as the reconstructed ancestral globins were synthesized by GeneArt Gene Synthesis (Thermo Fisher Scientific) after optimizing the nucleotide sequences in accordance with *E. coli* codon preferences. The synthesized globin gene cassette was cloned into a custom pGM vector system along with the *methionine aminopeptidase* (MAP) gene, as described previously (45).

**Expression and purification of recombinant Hbs**

Recombinant Hb expression was carried out in the *E. coli* JM109 (DE3) strain as described previously (45-47). Bacterial cell lysates were then loaded onto a HiTrap Q HP anion exchange column (GE Healthcare) and were then equilibrated with 20 mM Tris/0.5 mM EDTA (pH 8.3) and eluted with a linear gradient of 0 - 0.25 M NaCl. Hb-containing fractions were then loaded on to a HiTrap SP HP cation exchange column (GE Healthcare) and eluted with a linear pH gradient (pH 6.8 to 8.4). Eluted Hb fractions were concentrated using Amicon Ultra-4 Centrifugal Filter Units (Millipore) and oxygenation properties were measured as described above.

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**Acknowledgments:** We thank C. Natarajan for assistance in the laboratory and K. Campbell for comments on this manuscript. **Funding:** This study was funded by grants to JFS from the National Institutes of Health (HL087216) and the National Science Foundation (MCB-1517636 and OIA-1736249). **Author contributions:** AVS and JFS designed the study, AVS acquired the data, AVS and JFS analyzed the data and wrote the manuscript. **Competing interests:** Authors declare no competing interests. **Data and materials availability:** Sequence data collected as part of this study are deposited to GenBank under accession numbers XXXXXXXX-YYYYYY.

**Supplementary Materials:**

Figures S1-S3

Tables S1-S2
**Fig. 1.** Genomic organization of bovid $\beta$-globin gene clusters. Colored boxes represent individual genes. Labels denote previously annotated $\beta^C$, $\beta^A$, and $\beta^F$-globin genes.
**Fig. 2.** Phylogenetic analyses of bovid $\beta^C$-, $\beta^A$-, and $\beta^F$-globin genes. Alternative histories of gene deletion in Tibetan antelope yield testable phylogenetic hypotheses: (A) deletion of $\beta^C$, (B) deletion of $\beta^A$, and (C) deletion of $\beta^F$. (D) Estimated maximum likelihood phylogeny of bovid $\beta$-type globin genes indicates that Tibetan antelope has retained copies of $\beta^C$ and $\beta^F$, and that $\beta^A$ has been secondarily lost. Bootstrap support values are shown for relevant nodes.
**Fig. 3.** Analysis of pairwise sequence matches in chromosomal regions containing bovid β-globin gene clusters reveal a large-scale deletion in Tibetan antelope. Purple, green and blue colored boxes represent genes within the βC-, βA-, and βF-containing gene blocks, respectively. (A) Gray shading denotes percent sequence identity between homologous β-globin gene clusters. (B) A ~45kb chromosomal deletion in the β-globin cluster of the Tibetan antelope lineage resulted in secondary loss of the βA-containing gene block.
**Fig. 4.** O₂-affinities of total purified Hb from select bovid species. O₂ tensions at half saturation ($P_{50}$) for total Hb in the absence (stripped; gray hatched bars) and presence of 0.1 M KCl (black bars) at 37°C, pH 7.4 (0.1 mM Hb₄). Values are shown as mean $P_{50} \pm$ s.e.m (n = 3).
Fig. 5. O2-affinities of reconstructed ancestral bovid isoHbs. (A) Reconstructed ancestral β-globin genes (βC, βA, and βAC) of bovids. (B) O2 tensions at half saturation (mean $P_{50} \pm$ s.e.m., n = 3) for recombinant ancestral isoHbs in the absence (stripped) and presence of 0.1 M KCl at 37°C, pH 7.4 (0.1 mM Hb4). Schematic diagrams show the subunit composition of the three ancestral isoHbs (which have identical α-chains and structurally distinct β-chains).