Population History and Altitude-Related Adaptation in the Sherpa

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The first ascent of Mount Everest by Tenzing Norgay and Sir Edmund Hillary in 1953 brought global attention to the Sherpa people and human performance at altitude. The Sherpa inhabit the Khumbu Valley of Nepal, and are descendants of a population that has resided continuously on the Tibetan plateau for the past ∼25,000 to 40,000 years. The long exposure of the Sherpa to an inhospitable environment has driven genetic selection and produced distinct adaptive phenotypes. This review summarizes the population history of the Sherpa and their physiological and genetic adaptation to hypoxia. Genomic studies have identified robust signals of positive selection across EPAS1, EGLN1, and PPARA, that are associated with hemoglobin levels, which likely protect the Sherpa from altitude sickness. However, the biological underpinnings of other adaptive phenotypes such as birth weight and the increased reproductive success of Sherpa women are unknown. Further studies are required to identify additional signatures of selection and refine existing Sherpa-specific adaptive phenotypes to understand how genetic factors have underpinned adaptation in this population. By correlating known and emerging signals of genetic selection with adaptive phenotypes, we can further reveal hypoxia-related biological mechanisms of adaptation. Ultimately this work could provide valuable information regarding treatments of hypoxia-related illnesses including stroke, heart failure, lung disease and cancer.

Keywords: Sherpa, Tibetan, Sherpa physiology, hypoxia adaptation, genetic selection, high altitude adaptation, natural selection

INTRODUCTION

The term “sher-pa” is the Tibetan for “eastern-people”. The Sherpa reside primarily in the Solukhumbu district of Nepal but there are also smaller settlements in the Tibet Autonomous Region of China. The Sherpa speak a Tibetan dialect, and they share similar cultural and religious practices with Tibetans. They are traditionally engaged in farming; cultivating barley, potatoes and rearing yak and sheep. Starting with the first Everest expeditions in the 1920's, the Sherpa have become renowned for their ability as mountaineers and today they often aid and lead climbing expeditions in the Himalayas. Examples of their exceptional climbing feats include the first ascent of Mount Everest by Tenzing Norgay Sherpa, who accompanied Sir Edmund Hillary in the final stage of the 1953 expedition and Ang Rita Sherpa (known as “The Snow Leopard”) who, between 1983 and 1996, summited Everest ten times without the use of supplemental oxygen. The remarkable tolerance of the Sherpa to hypoxia has, over the last 60 years, been a focus of attention for the scientific community, in particular physiologists (Gilbert-Kawai et al., 2014).
The Sherpa are direct descendants of an ancestral population that has resided continuously on the Tibetan plateau for the past 25,000 to 40,000 years (Aldenderfer, 2011; Zhang et al., 2018). This long exposure to the evolutionary pressure presented by high altitude has driven physiological adaptation, which in turn has allowed the Sherpa to thrive. The adaptive physiological makeup of the Sherpa can inform on treatments for hypoxia-related illness including pulmonary, cardiac, neurological and renal disorders (Martin et al., 2013; Luks and Hackett, 2014; Gilbert-Kawai et al., 2015). Thus, studying the Sherpa at altitude offers a unique, “natural laboratory” that can provide insight to the molecular mechanisms of hypoxia.

An early paper on Sherpa physiology, published in 1965, suggested that the Sherpa have an efficient mechanism of oxygen utilization at the cellular level, allowing them to perform well under hypoxia (Lahiri and Milledge, 1965). Since then, our knowledge of Sherpa adaptation has grown, largely by comparing different physiological parameters between the Sherpa and people of lowland origin. With the development of high throughput DNA genotyping and sequencing platforms, genomics studies of indigenous high-altitude populations, including the Sherpa, have begun to emerge. These have provided insight into population history and genetic signatures of altitude-driven natural selection. In this review, we (1) summarize the population history of, (2) describe distinct adaptive phenotypes and (3) discuss signatures of selection, in the Sherpa. We highlight the need for further research connecting genetic factors to physiological adaptation in the Sherpa at extreme altitude.

THE SHERPA, A RECENTLY DERIVED TIBETAN POPULATION

Stone tools used by early humans have been found at Nwya Devu in central Tibet at an altitude of 4,600 m. Dating to 30,000 to 40,000 years before present (YBP), these findings represent the earliest archeological record of human colonization of the Tibetan plateau (Zhang et al., 2018). Genetic studies have suggested that the ancestors of both the Sherpa and Tibetans diverged from a Han Chinese population and arrived on the Tibetan plateau from lowland East Asia around 40,000 years ago (Qi et al., 2013; Jeong et al., 2014).

The prevailing hypothesis is that, during the 16th century, the ancestors of the Sherpa migrated from Tibet to the Khumbu Valley of Nepal, driven by political and religious turmoil resulting from a Mongol invasion (Oppitz, 1974). The presence of Sherpa-specific mitochondrial DNA (mtDNA) lineages (Kang et al., 2013) in a Nepalese context, with an estimated age of less than 1,500 years and derived from Tibetans, further supports this hypothesis of a recent migration of the Sherpa to the Khumbu valley (Bhandari et al., 2015).

There is a long history of migration from the Tibetan plateau to Nepal. To illustrate, genomic analysis of human dental samples (dating to between 1,700 and 3,000 YBP) from a northern region of Nepal show strong affinity for contemporary Tibetans (Jeong et al., 2016). Analysis of both autosomal data (Lu et al., 2016; Gncechi-Ruscone et al., 2017) and uniparental mtDNA and Y-chromosome markers (Bhandari et al., 2015) have shown the Sherpa and Tibetans to share relatively recent common ancestry. Tibetans also share recent common ancestry with other Nepalese populations including the Rai, Magar, Tamang, and Gurung (Cole et al., 2017). The Sherpa share more genetic affinity with these Tibeto-Burman speaking populations than with other Indo-Aryan populations of Nepal. However, the Sherpa are distinct from other Nepalese populations in that the Sherpa have elevated levels of runs of homozygosity (Cole et al., 2017), and illustrate very little or no admixture with Nepalese or South Asian populations (Cole et al., 2017). Thus, the Khumbu Valley Sherpa can be considered from the perspective of population genetics as a “bottlenecked” population recently derived from Tibetans.

COMPARATIVE PHYSIOLOGICAL STUDIES BETWEEN SHERPA AND LOWLANDERS

In 1952, Griffith Pugh conducted a series of pioneering physiological experiments on Mount Cho Oyo (at 8,188 m, 20 km west of Mount Everest) that suggested a superior work capacity of the Sherpa at high altitude (Pugh, 1962; Pugh et al., 1964). They also provided the scientific rationale for the hydration, nutrition and oxygen requirements for the first Everest summiting in 1953 (Milledge, 2002). Although the physiology of the Sherpa has been studied over the intervening 60 years, the scientific literature is limited in number, and most of the studies are based on small sample sizes. There are obvious challenges to studying the Sherpa; they reside in a remote region, at an altitude over 2,800 m, where altitude sickness is common for sojourners. Despite this, several remarkable findings have emerged and below we discuss specific phenotypes that may be linked to hypoxia-related genetic signals of selection reported to date. For a discussion of other hypoxia-related physiological parameters studied in Sherpa, such as ventilation, lung volume, exercise capacity and cerebral function (see Gilbert-Kawai et al., 2014; Table 1).

Hemoglobin Concentration

The hypoxic challenge presented by high altitude drives changes in hemoglobin concentration. Elevated hemoglobin levels (≥19 g/dl in females; and ≥21 g/dl in males) resulting from hypoxia can lead to chronic mountain sickness (Leon-Velarde et al., 2005). Relative to lowland controls, the literature suggests the Sherpa display lower hemoglobin concentrations at high altitude (Beall and Reichisman, 1984; Wu et al., 2013; Bhandari et al., 2016). Sherpa women with lower hemoglobin concentrations (13.8 g/dl ± 1.3 g/dl) are reported to have better reproductive outcomes (Beall et al., 1997, 2004; Cho et al., 2017). Increased exercise capacity has been reported in Tibetan males with a low erythropoietic response (Simonson et al., 2015). It is yet to be determined whether the lower hemoglobin concentration observed in Sherpa is due to a blunted erythropoietic response or to some other physiological parameters that impact hemoglobin concentration.
### TABLE 1 | Physiological parameters studied in Sherpa and lowlanders at altitude.

| Parameter(s) | Sherpa at high altitude | Lowlander at altitude (meter) | Reference(s) |
|--------------|-------------------------|-------------------------------|---------------|
|              | Sample size | Parameter value | Altitude | Sample size | Duration (days) | Parameter value |
| Heart rate while working at 900 kg-m/min-beats/min | 1 | 162 | 5,800 | 2 | 240 | 122 | Pugh, 1962; Pugh et al., 1964 |
| Lung diffusion capacity for oxygen-ml/min | 1 | 97 | 5,800 | 2 | 240 | 52.5 | Pugh, 1962 |
| Basal metabolic rate, kcal/m² h | 3 | 46.1 ± 1.0 | 5,800 | 8 | 240 | 41.1 ± 3.6 | Gill and Pugh, 1964 |
| 10 different physiological parameters; measured, to test oxygen utilization at the cellular level | 4 | efficiently used O₂ | 4,880 | 3 | 60 | less efficient to use O₂ | Lahiri and Milledge, 1965 |
| Heart rate (while work rate at 1,265 kg-m/min) | 4 | 198 | 4,880 | 2 | 63 | 146 | Lahiri et al., 1967 |
| Lung diffusion capacity for oxygen-ml/min | 4 | 28.6 | 4,880 | 5 | 0 | 25.9 | Lahiri and Milledge, 1967 |
| Hemoglobin level in Tibetans living at 3658 m in Nepal; g/100 ml | 5 | Male: 16.8 ± 1.4; Female: 14.5 ± 0.7 | | | | | Adams and Shresta, 1974 |
| Hemoglobin level in Tibetans living at 4000 m in Nepal; g/100 ml | 5 | Male: 17.0 ± 1.25; Female: 15.3 ± 0.8 | | | | | Adams and Strang, 1975 |
| Ratio of 2, 3 diphosphoglycerate and hemoglobin | 7 | 0.9 | 3,900 | 2 | 30 | 1.26 | Morpurgo et al., 1976 |
| Mean oxygen half saturation of hemoglobin | 7 | 27.3 ± 1.8 | 3,500 | 7 | 120 | 28.2 ± 1.3 | Samaja et al., 1979 |
| Arterial oxygen saturation (SaO₂) | 10 | 88 ± 0.74 | 4,243 | 25 | 12 | 85.6 ± 1.0 | Hackett et al., 1980 |
| Body weight changes- Mean weight loss (kg) | 4 | constant | 5,400 | 13 | 25 | 1.9 to 4 | Boyer and Blume, 1984 |
| Partial pressure of oxygen in arterial blood (Torr) | 6 | 34.5 ± 3.2 | 5,400 | 9 | – | 41.0 ± 3.3 | Santolaya et al., 1989 |
| Partial pressure of carboxyhemoglobin in arterial blood (Torr) | 6 | 27.5 ± 2.2 | 5,400 | 9 | – | 20.0 ± 2.8 | Sutton et al., 1988 |
| Resting glucose appearance rate at sea level (1.79 ± 0.02) mg.kg⁻¹.l.min⁻¹ | 5 | 15 ± 1 | 22 | | | 28 ± 2 | Groves et al., 1993 |
| Glucose metabolic rates of myocardial regions | 6 | 0.32 ± 0.05 | 226 | 6 | | 0.20 ± 0.04 | Holden et al., 1995 |
| Brain glucose metabolic rates | 6 | 0.71 | 6 | 19 | | 0.73 | Hochachka et al., 1996b |
| Signs of mild cortical atrophy | 7 | Seen in 1 | | | | | Garrits et al., 1996 |
| Partial pressure of carbon dioxide, mm Hg | 5 | 28.8 ± 1.2 | 3,400 | 4 | 40 | 22.0 ± 0.4 | Samaja et al., 1997 |
| Hypoxic ventilatory response (HVR)-end-tidal PO₂, 40 Torr | 6 | 34.5 ± 2.2 | 1,300 | 11 | 14 | 281 | Droma et al., 2008 |
| Partial pressure of oxygen in arterial blood (Torr) | 6 | 34.5 ± 3.2 | 226 | 5 | | 0.20 ± 0.04 | Holden et al., 1995 |
| Partial pressure of carboxyhemoglobin in arterial blood (Torr) | 6 | 27.5 ± 2.2 | 5,400 | 9 | – | 20.0 ± 2.8 | Sutton et al., 1988 |
| Resting glucose appearance rate at sea level (1.79 ± 0.02) mg.kg⁻¹.l.min⁻¹ | 5 | 15 ± 1 | 22 | | | 28 ± 2 | Groves et al., 1993 |
| Glucose metabolic rates of myocardial regions | 6 | 0.32 ± 0.05 | 226 | 6 | | 0.20 ± 0.04 | Holden et al., 1995 |
| Brain glucose metabolic rates | 6 | 0.71 | 6 | 19 | | 0.73 | Hochachka et al., 1996b |
| Signs of mild cortical atrophy | 7 | Seen in 1 | | | | | Garrits et al., 1996 |
| Partial pressure of carbon dioxide, mm Hg | 5 | 28.8 ± 1.2 | 3,400 | 4 | 40 | 22.0 ± 0.4 | Samaja et al., 1997 |
| Mean arterial blood pressure, mm Hg | 9 | 83 ± 6 | 4,243 | 10 | 7 | 94 ± 7 | Jansen et al., 2000 |
| Forced expiratory volume of adult male (%) | 146 | 110(107–114) | 3,840 | 103 | | 103.8 | Havryk et al., 2002 |
| Heart Rate (beats min⁻¹) means ± S.D. | 7 | 167 ± 10 | 5,050 | 10 | 28 | 149 ± 7 | Marconi et al., 2004 |
| Carried loads of their body weight (mean ± SD) | 96 | 93 ± 36% | 2,880 | 10 | | 75% | Bastien et al., 2005a,b, 2016 |
| Arterial oxygen saturation (SaO₂) or (SpO₂) lower than Tibetans | – | – | 8,400 | 4 | | 145.8 ml per L | Wu, 1990; Wu and Kayser, 2006 |
| Arterial oxygen saturation, % | 10 | 88 ± 3 | 40 | 10 | | 97 ± 2 | Jansen et al., 2007 |
| Statistically significant gender specific differences in SpO₂ | | | Adult Tibetan female show higher SpO₂ value than male | | | | Weitz and Garruto, 2007 |
| Serum angiotension-converting enzyme activity, IU/L/37°C | 105 | 14.5 ± 0.4 | 1,300 | 111 | | 14.7 ± 0.4 | Droma et al., 2008 |
| Mean arterial oxygen content at 8,400 m (26% lower than at 7100 m) | | | | | | | |
| Muscle phosphocreatine recovery halftime-PCr₁/₂ (s) | 7 | 22.2 ± 1.6 | 50 | 7 | | 16.1 ± 1.1 | Edwards et al., 2010 |
TABLE 1 | Continued

| Parameters                                                                 | Sherpa at high altitude | Lowlander at altitude (meter) | Reference(s)                  |
|---------------------------------------------------------------------------|-------------------------|--------------------------------|--------------------------------|
|                                                                           | Number | Parameters values | Altitude | Number | Duration (days) | Parameters values |
| Radial arterial plasma NO\(_2\) (nmol l\(^{-1}\))                        | –      | –                 | 4,559    | 26     | 4               | 263.6 ± 61.2       |
| Middle cerebral artery diameter [at 6,400 m = 6.66 mm]                     | –      | –                 | 7,950    | 5      | 71              | 9.34 mm            |
| Flow-mediated dilatation (FMD)-shear rate                                 | 12     | 24490 ± 7230      | 5,050    | 12     | 14              | 14802 ± 5306       |
| Arterial oxygen saturation (mean ± SE)                                    | 13     | 86 ± 1            | 5,050    | 13     | 9               | 83 ± 2             |
| Hemoglobin level ml. min\(^{-1}\), mmHg\(^{-1}\)                         | 13     | 61 ± 4            | 5,050    | 13     | 9               | 37 ± 2             |
| Lung diffusing capacities                                                 | 13     | 226 ± 18          | 5,050    | 13     | 9               | 153 ± 9            |
| Systolic pulmonary artery pressure                                        | 95     | 29.4 ± 5.5        | 13       | 64     | –               | 23.6 ± 4.8         |
| Left ventricular untwisting velocity, °/s                                  | 11     | –93 ± 31          | 5,050    | 9      | 13              | –153 ± 38          |
| Relative PPARα mRNA expression of muscles tissues                         | 15     | 0.5158            | 5,300    | 10     | 19              | 1.0045             |
| Post reproductive, Tibetan women (n=959)-Hemoglobin concentration, gm/dl | –      | 13.8              | –        | –      | –               | –                  |
| Increase in nocturnal time course of blood oxygen saturation level at rest | –      | 3,050             | 10      | 10     | 21              | 94.5% (91-97)      |
| FMD unchanged (in rest and maximal exercise), at low and high altitude    | –      | –                 | 3,800    | 9      | 7               | 6.3 ± 1.3%         |
| Brachial artery blood flow [at Sea level-(142.7 ± 30.6)], ml/min          | –      | –                 | 5,050    | 14     | 21              | 53.1 ± 11.1        |
| Number of circulating microparticles in blood (CD 66b+)/µl (21 ± 4) Sea level | –      | –                 | 3,800    | 10     | 3               | 74 ± 17            |
| Birth-weight (kg) in Tibetans & Han; at 3,000–4,000 m altitude             | 100    | 3.14 (3.06, 3.22) | <4,000   | 100    | 2.61 (2.34, 2.88) |
| Case report of a 32 week gestation Sherpa at 5160 m and her data after 10 month postpartum | –      | –                 | No apparent maternal, fetal or neonatal complications | – |
| Arterial oxygen pressure (PaO\(_2\); mm Hg)                               | 9      | 50.1 ± 11.3       | 5,050    | 9      | –               | 54 ± 1.2           |
| Prefatigue, maximal voluntary contraction torque, N. m                    | 10     | 44.3 ± 14.1       | 5,050    | 12     | 10              | 58.2 ± 8.1         |
| Brachial artery flow-mediated dilatation (FMD)                             | 12     | 5.8 ± 2.8%        | 5,050    | 22     | 10              | 3.8 ± 2.8%         |
| Resting posterior cerebral artery velocity                                 | –      | –                 | 4,240    | 10     | 13              | 43 cm/s            |
| Lowland origin; Female SpO\(_2\); Mean (SD), (%)95.2 (1.2); at 600 m      | –      | –                 | 3,500    | 20     | 1               | 76.7 (5.6)         |
| Partial pressure of arterial carbon dioxide, mmHg                           | 11     | 32.1 ± 2.5        | 5,050    | 21     | 21              | 30.0 ± 1.9         |
| Peripheral oxygen saturation in female [at 600 m; 96.9 (1.0) Mean (SD)]%  | –      | –                 | 3,840    | 20     | 1               | 86.5 (6.5)         |
| SpO\(_2\) (%) [at Sea Level (244 m) is 98 ± 1]                            | –      | –                 | 3,800    | 12     | 10              | 89.1 ± 3           |
| Free cysteine and plasma total free thiol concentrations                   | –      | –                 | 4,559    | 4      | Elevated at 4,559 m than at 50 m |
| Sublingual capillary total vessel density [at Sea Level; 18.81 ± 3.92 mm mm\(^{-2}\)] | –      | –                 | 7,042    | 10     | 21              | 21.25 ± 2.27       |
| Sympathetic nerve activity, burst frequency (bursts min\(^{-1}\))         | 8      | 22 ± 11           | 5,050    | 14     | 20              | 30 ± 9             |

**Nitric Oxide Concentration**

Nitric oxide acts as a vasodilator and is believed to protect against pulmonary hypertension at high altitude (Busch et al., 2001). It also plays a role in haematocrit regulation by controlling blood viscosity (Ashmore et al., 2014). Serum nitric oxide levels have been reported as reduced in the Sherpa relative to lowlanders.
content, but their muscle is somehow maximizing the oxygen (Kayser et al., 1991). Sherpa also display a reduced mitochondrial capillaries per cross-sectional area, in comparison to lowlanders. Sherpa muscle contains a significantly greater number of capillaries per cross-sectional area, in comparison to lowlanders (Hochachka et al., 1996a). This ratio remains steady in the Sherpa (Hochachka et al., 1996a). However, the mechanism by which this reduced myocardial relaxation impacts on the exercise capacity of the Sherpa is unclear (Stembridge et al., 2015). These are presumably the result of exposure over many generations to the hypoxia-related selective pressure presented by the Tibetan plateau. Indeed, some examples have already emerged of specific genetic signatures of selection associating with distinct adaptive traits (Simonson, 2015; Moore, 2017).

Skeletal Muscle
Sherpa muscle contains a significantly greater number of capillaries per cross-sectional area, in comparison to lowlanders (Kayser et al., 1991). Sherpa also display a reduced mitochondrial content, but their muscle is somehow maximizing the oxygen consumption to mitochondrial volume ratio (Kayser et al., 1991; Horscroft et al., 2017). Under hypoxia, Sherpa skeletal muscle prefers carbohydrate over fatty acids as a metabolic substrate (Murray, 2009). Sherpa muscle maintains fatty acid oxidation relative to lowlanders at high altitude. Incomplete fatty acid oxidation results in production of byproducts such as acylcarnitines and reactive oxygen species. Acylcarnitines and markers of oxidative stress (e.g., reduced/oxidized glutathione and methionine sulfoxide) are increased in lowlander muscle relative to the Sherpa (Gelfi et al., 2004; Horscroft et al., 2017). However, oxidative damage in lowlanders was reduced to levels comparable with the Sherpa, where acclimatization has taken place (Janocha et al., 2017). Lactate dehydrogenase activity is elevated in Sherpa muscle (Allen et al., 1997; Horscroft et al., 2017), indicating greater capacity for anaerobic lactate production. With increasing altitude, lowlanders experience a gradual reduction in phosphocreatine (PCr) and ATP levels (Levett et al., 2015). But the Sherpa maintain PCr and ATP levels at altitude (Horscroft et al., 2017). Thus, the superior muscle energetics displayed by the Sherpa is probably the result of adaptation at the metabolic level.

Birth Weight
Women of European and Han Chinese ancestry exhibit reduced birth weights following gestation at high altitude, quantified at 100 g reduction for every 1,000 m elevation (Moore, 2003; Julian et al., 2009; Moore et al., 2011). The Sherpa (and Tibetans), however, maintain normal birth weight at both low (1,330 m) and high (3,930 m) altitude (Smith, 1997; Moore et al., 2001). Genes including PPARα are expressed in the placenta (Barak et al., 2008) and have been shown to influence female reproductive function (Bogacka et al., 2015). HIFs play a critical role in mammalian embryo and placental development (Dunwoodie, 2009; Pringle et al., 2009). EPAS1 expression appears reduced in umbilical endothelial cells and placentas of Tibetan women (Peng et al., 2017). Intrinsic variants in CCDC14I have been shown in Tibetan and Sherpa women to associate with the number of live births, and the same locus also shows evidence of positive selection (Jeong et al., 2018). The increased reproductive success of the Sherpa is therefore likely to be, at least in part, due to cardiac-related traits (Jeong et al., 2018) and placental adaptation (Burton et al., 2016). Further studies are required to understand the molecular mechanisms by which the Sherpa maintain normal intrauterine growth at altitude.

In summary, the Sherpa display distinct physiological responses to hypoxia that contrast to lowlanders at high altitude (Table 1). These are presumably the result of exposure over many generations to the hypoxia-related selective pressure presented by the Tibetan plateau. Indeed, some examples have already emerged of specific genetic signatures of selection associating with distinct adaptive traits (Simonson, 2015; Moore, 2017).
population-specific signatures of selection for adaptation across the human genome. There are now several complementary genomic tests available for detecting genetic selection (Scheinfeldt and Tishkoff, 2013) and the application of these tests to data from indigenous high-altitude people including the Sherpa have identified numerous and remarkable genetic signals of selection. Here, we focus on the three most robust signals of selection detected to date in the Sherpa: EPAS1, EGLN1, and PPARA (Table 2).

Endothelial PAS Domain-Containing Protein 1 (EPAS1)

One of the earliest signals for altitude-related adaptation to emerge from genomic selection studies was EPAS1. Initially discovered in Tibetans (Beall et al., 2010), the EPAS1 signal has been replicated in multiple other Tibetan populations (Bigham et al., 2010; Simonson et al., 2010; Yi et al., 2010; Peng et al., 2011; Wang et al., 2011; Xu et al., 2011) as well as the Sherpa (Hanaoka et al., 2012; Jeong et al., 2014; Bhandari et al., 2016). The selected EPAS1 haplotype is associated with lowered hemoglobin concentrations (Beall et al., 2010). Remarkably, it seems the adaptive EPAS1 haplotype likely descends from an introgression event with the Denisovan people, an extinct species of archaic humans (Huerta-Sánchez et al., 2014; Hu et al., 2017). A 3.4 kb copy number deletion, downstream of EPAS1, is elevated in frequency, in Tibetans and Sherpas relative to lowland controls (Lou et al., 2015). This deletion is in strong linkage disequilibrium with the previously reported (Beall et al., 2010) EPAS1 haplotype and has also been associated with lower hemoglobin levels. The actual functional EPAS1 variant(s) that are conferring advantage in relation to hypoxic adaptation remain unknown. However, the intronic and intergenic location of the selected variants would be consistent with a role in HIF-related transcriptional regulation.

### TABLE 2

A summary of genetic adaptations reported in the Sherpa, and replication in other population(s) or species.

| Genes name(s)                      | Sample Size | Sherga Reference(s) | Other population(s) or species | Reference(s) |
|-----------------------------------|-------------|---------------------|--------------------------------|--------------|
| ACE                               | 105         | Droma et al., 2008  | Elite European descent athletes | Montgomery et al., 1998; Jones et al., 2002 |
| HIF-la                            | 20          | Suzuki et al., 2003 | –                              | –            |
| eNOS                              | 105         | Droma et al., 2006  | Tibetan                        | Beall et al., 2010; Simonson et al., 2010; Yi et al., 2010; Bigham et al., 2010; Peng et al., 2011; Wang et al., 2011; Xu et al., 2011 |
| EPAS1                             | 105         | Hanaoka et al., 2012| Deedu Mongolian                | Xing et al., 2013 |
| 3.4 kb Copy Number Deletion-80 kb downstream of EPAS1 | 51          | Jeong et al., 2014  | Tibetan                        | Lou et al., 2015 |
| 3.4 kb Copy Number Deletion-80 kb downstream of EPAS1 | 582         | Bhandari et al., 2016 | Denisovan                      | Huerta-Sánchez et al., 2014 |
| EGLN1                             | 51          | Jeong et al., 2014  | Tibetan                        | Lorenzo, 2010; Simonson et al., 2010; Yi et al., 2010; Xiang et al., 2013; Lorenzo et al., 2014 |
| 3.4 kb Copy Number Deletion-80 kb downstream of EPAS1 | 582         | Bhandari et al., 2016 | Andean                         | Bigham et al., 2009; Bigham et al., 2010 |
| PPARA                             | 15          | Horscroft et al., 2017 | Tibetan                        | Pagani et al., 2012 |
| HYOU/HMB3                         | 51          | Jeong et al., 2014  | Tibetan                        | Lok et al., 2010; Simonson et al., 2010; Peng et al., 2011 |
| 3.4 kb Copy Number Deletion-80 kb downstream of EPAS1 | 111         | Zhang et al., 2017  | Andean                         | Scheinfeldt et al., 2012 |
| NOS1                              | 111         | Zhang et al., 2017  | Tibetan (GCH1), Andeans (NOS2) | Bigham et al., 2009; Bigham et al., 2010; He et al., 2018 |
| ANGPT1                            | 111         | Zhang et al., 2017  | Tibetan and grey wolves of TAR, China | Jeansson et al., 2011; Wang et al., 2011 |
| EPAS1, EGLN1, RP11-384F7.2 AC066833.1, ZNF53 2, HLA-DOB1/HLA-DBP1 | 10          | Arciero et al., 2018 | Pigs of TAR, China             | Ai et al., 2014 |
| ANKH                              | 10          | Arciero et al., 2018 | Pigs of TAR, China             | Li et al., 2016 |
| GRB2                              | 31          | Gnecci-Ruscone et al., 2017, 2018 | Pigs of TAR, China             | Gnecci-Ruscone et al., 2018 |
EPAS1 encodes the HIF2 alpha subunit of HIF2. The postnatal deletion of EPAS1 in adult mice causes anaemia (Gruber et al., 2007). Some cases of erythrocytosis are caused by missense mutations (e.g., G536W) in EPAS1 (Percy et al., 2008). Mice carrying the EPAS1 G536W mutation display excessive erythrocytosis and pulmonary hypertension (Tan et al., 2013). Another study in heterozygous EPAS1 knockout mice reported a blunted physiological response to chronic hypoxia (Peng et al., 2017). Further in-vivo and in-vitro studies are necessary to understand how the adaptive version of the EPAS1 gene is shaping human adaptation to altitude.

Egl-9 Family Hypoxia Inducible Factor 1 (EGLN1)

Another high altitude genetic selection signal to emerge from early studies on Tibetans was EGLN1 (Simonson et al., 2010; Yi et al., 2010). Similar to EPAS1, this signal was later demonstrated in the Sherpa (Jeong et al., 2014). Two functional EGLN1 mutations (rs12097901, D4E, and rs186996510, S127C) appear to be driving the selection signal and are present in both Sherpa (Bhandari et al., 2016) and Tibetans (Lorenzo, 2010; Xiang et al., 2013; Lorenzo et al., 2014). Whether the mode of action of these two mutations is via gain of function (Lorenzo et al., 2014) or loss of function (Song et al., 2014) remains unclear.

EGLN1 encodes proline hydroxylase 2 (PHD2), an isoform of HIF prolyl-hydroxylase. Homozygous knockout PHD2 mice are unviable and die at the embryonic stage due to severe placental defects (Takeda et al., 2006). Knockout mice with PHD2 disruption targeted to specific organs including the liver, heart, kidney and lung develop excessive vascular growth (Takeda et al., 2007). Adult mice deficient for PHD2 display excessive erythrocytosis (Takeda et al., 2008) and heterozygous PHD2 mice have an increased ventilatory sensitivity to hypoxia and carotid body hyperplasia (Bishop et al., 2013).

Peroxisome Proliferator-Activated Nuclear Receptor A (PPARA)

PPARA encodes PPARα, a transcriptional regulator of fatty acid oxidation in liver, heart and muscle (Gilde and Van Bilsen, 2003). PPARα has tissue-specific expression and, under hypoxic conditions, is downregulated by HIFs (Narravula and Colgan, 2001). Positive selection across the PPARA gene has been reported in Tibetans (Simonson et al., 2010) and Sherpa (Horscroft et al., 2017), and the selected PPARA SNPs correlate with reduced hemoglobin levels (Simonson et al., 2010). Sherpa carriers of the positively selected PPARA alleles switch to more efficient fuels such as glucose and display decreased muscular fatty acid oxidation (Horscroft et al., 2017). Most of the PPARA SNPs reported to be under selection appear to be non-coding variants (Kinota et al., 2018). It is unclear if these variants directly affect transcriptional regulation or are linked with functional variants in other genes or nearby inter-genic regions.

CONCLUSION

The Sherpa show remarkable performance in the hypoxic environment presented by high altitude. Comparative physiological studies have suggested numerous distinct, adaptive phenotypes in the Sherpa including advantageous levels of hemoglobin, oxygen saturation and birth weight, and the elevated reproductive success of Sherpa women. Genomic studies have identified robust signals of positive selection across genes including EPAS1, EGLN1 and PPARA. All three of these signals of genetic selection have been shown to correlate with advantageous levels of hemoglobin. However, Sherpa-specific signals of genetic selection have also been reported, suggesting that whilst some of the genetic basis for adaptation in the Sherpa is shared with Tibetans, there may be features unique to the Sherpa, which could in turn explain distinct Sherpa phenotypes. Collectively, this illustrates how the outstanding physiological performance of the Sherpa at altitude is, at least in part, a result of hypoxia driven genetic selection spanning the ∼35,000 years of seasonal migration on the Himalayan plateau. Further comparative physiological studies are required to refine existing, and identify additional adaptive phenotypes, in particular those that are specific to the Sherpa. By correlating these phenotypes with known and emerging signals of genetic selection, we can shed light on biological mechanisms of Sherpa hypoxic adaptation. Ultimately this work can inform on treatments of hypoxia-related illness including pulmonary, cardiac, neurological and renal disorders.

AUTHOR CONTRIBUTIONS

Both authors drafted, edited, and approved the final version of the manuscript.

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**Conflict of Interest Statement:** The authors declare that the research was conducted in the absence of any commercial or financial relationships that could be construed as a potential conflict of interest.

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