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Epistatic Adaptive Evolution of Human Color Vision

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Abstract

Establishing genotype-phenotype relationship is the key to understand the molecular mechanism of phenotypic adaptation. This initial step may be untangled by analyzing appropriate ancestral molecules, but it is a daunting task to recapitulate the evolution of non-additive (epistatic) interactions of amino acids and function of a protein separately. To adapt to the ultraviolet (UV)-free retinal environment, the short wavelength-sensitive (SWS1) visual pigment in human (human S1) switched from detecting UV to absorbing blue light during the last 90 million years. Mutagenesis experiments of the UV-sensitive pigment in the Boreoeutherian ancestor show that the blue-sensitivity was achieved by seven mutations. The experimental and quantum chemical analyses show that 4,008 of all 5,040 possible evolutionary trajectories are terminated prematurely by containing a dehydrated nonfunctional pigment. Phylogenetic analysis further suggests that human ancestors achieved the blue-sensitivity gradually and almost exclusively by epistasis. When the final stage of spectral tuning of human S1 was underway 45–30 million years ago, the middle and long wavelength-sensitive (MWS/LWS) pigments appeared and so-called trichromatic color vision was established by interprotein epistasis. The adaptive evolution of human S1 differs dramatically from classical originological pigments with a major mutational effect used in achieving blue-sensitivity in a fish and several mammalian species and in regaining UV vision in birds. These observations imply that the mechanisms of epistatic interactions must be understood by studying various orthologues in different species that have adapted to various ecological and physiological environments.

Introduction

The chance of survival of novel mutations is affected strongly by the molecular background in which they appear [1–3]. To understand the evolutionary dynamics of these mutations, it is necessary to characterize the phenotypic variation they generate. For vertebrates, however, understanding the genotype-phenotype relationship remains challenging because of technical difficulties in connecting the structure and function of evolving proteins and in evaluating non-additive (epistatic) interactions among amino acids unambiguously [3–7].

About 90% of human populations detect the entire range of visible color using three types of cone pigments: short wavelength-sensitive (SWS1) pigment (human S1), middle wavelength-sensitive (MWS) pigment (human M) and long wavelength-sensitive (LWS) pigment (human L), which detect light maximally ($\lambda_{\text{max}}$) at 414, 530 and 560 nm, respectively [8,9]. Human S1 can be made UV-sensitive ($\lambda_{\text{max}} = 360$ nm) by introducing seven mutations T16F, L49F, F52T, L66F, P93T, G114A and T118S, whereas the UV-sensitive pigment in mouse (Mus musculus) (mouse S1, $\lambda_{\text{max}} = 359$ nm) can be made blue-sensitive ($\lambda_{\text{max}} = 411$ nm) by the seven reverse mutations; however, when the seven mutations are introduced into mouse S1 individually, none of the individual changes produce any $\lambda_{\text{max}}$-shift [10], showing an extreme case of epistatic interactions. Largely free from the technical difficulties in evaluating the genotype-phenotype relationships as well as their strong associations to variable ecological and physiological environments, visual pigments make vertebrate vision a powerful model to directly study the dynamics of genotype-phenotype relationship during phenotypic adaptation [11–13]. The crystal structure of the visual pigments in bovine rod photoreceptors [14,15] and a large dataset on ecology of vision [16-18] also allow us to link the chemistry, genetics, organismal biology and ecology of vertebrate vision.

Regulated primarily by the lens [19], photons with wavelengths shorter than 400 nm do not reach the retinas of some primates, including human, and sciurid rodents, whereas UV light does reach the retinas of mouse, rat and other mammals [19,20]. As it may be suspected from these observations, SWS1 pigments in most vertebrate ancestors, including the mammalian ancestor, had $\lambda_{\text{max}}$ values of 360 nm and were UV-sensitive [21]. Subsequently, in the lineage leading to humans, the UV-transmitting lens evolved into a UV-absorbing lens [20]. To adapt to the UV-free retinal environment [19,20,22], human S1 switched from detecting UV to absorbing blue light during the last 90 million years (My) [21].

Making the UV-sensitive mouse S1 blue-sensitive may give an impression that human S1 evolved from the SWS1 pigment of the


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Author Summary

Mapping the genotype-phenotype relationship is necessary to understand how variable phenotypes have evolved in nature. The blue-sensitive visual pigment in human (human S1) evolved from the UV-sensitive pigment in the Boreoeutherian (or Boreotherian) ancestor (AncBoreotheria S1) by seven mutations. Mutagenesis experiments reveal that 4,008 (~80%) of all 5,040 possible evolutionary trajectories connecting from AncBoreotheria S1 to human S1 are terminated prematurely. Quantum chemical analyses suggest that the premature termination of trajectories was caused by containing a dehydrated nonfunctional pigment. Phylogenetic analysis further suggests that the blue-sensitivity was achieved only gradually and almost exclusively by the seven non-additively interacting amino acids. During the period between 45 and 30 My ago, human S1 was in the final stage of developing its blue-sensitivity. This was the time when two red-sensitive pigments appeared by gene duplication and one of them became green-sensitive. Trichromatic color vision in the human lineage was fully developed by 30 My ago by interprotein epistasis among the three visual pigments. Manipulation of the genetically engineered ancestral molecule is the key to recapitulate the evolution of phenotypic adaptation and that of epistatic interaction separately.

Boreoeutherian (or Boreotherian) ancestor (AncBoreotheria S1) by F46T, F49L, T52F, F86L, T93P, A114G and S118T. To prove this, however, we have to show that these mutations actually switched the $\lambda_{\text{max}}$ of AncBoreotheria S1 into that of human S1. Even if such critical mutations are identified, it is still unclear how they have accumulated mutations and modified the $\lambda_{\text{max}}$ of AncBoreotheria S1 during evolution. To address these questions, we engineer AncBoreotheria S1, construct all possible evolutionary trajectories that connect AncBoreotheria S1 and human S1, recapitulate the evolutionary changes in the epistatic interaction and the $\lambda_{\text{max}}$ separately; in this way, we establish the genotype-phenotype relationship unambiguously during the entire process of human S1 evolution. We then consider how human ancestors acquired the present trichromatic color vision based on the three cone pigments.

Results

Evolutionary trajectories

Based on the phylogenetic tree of 33 representative SWS1 pigments sampled from a wide range of vertebrate species (Fig. 1), the amino acid sequences at different nodes have been inferred using PAML [23] and AncBoreotheria S1 was reconstructed (for details, see Methods). The in vitro assay shows that AncBoreotheria S1 has a $\lambda_{\text{max}}$ of 357 nm (Fig. 2A, in black spectrum) and its mutant with F46T, F49L, T52F, F86L, T93P, A114G and S118T has a $\lambda_{\text{max}}$ of 344 nm (S1 Table). These results, indeed, demonstrate that human S1 evolved from AncBoreotheria S1 (or AncBoreotheria (P357)) by accumulating the seven mutations.

To accommodate seven sequential amino acid replacements, there are 7! ( = 5,040) possible evolutionary trajectories. We traced all these trajectories by introducing seven single and all 120 possible combinations of the seven mutations into AncBoreotheria S1 and evaluating their effects on the $\lambda_{\text{max}}$-shift (or $\Delta\lambda_{\text{max}}$). The in vitro assays revealed that 60 out of 127 mutants could not make functional visual pigments, making 4,008 possible evolutionary paths prematurely terminated (S1 Fig.). These incomplete trajectories are caused most often by T52F primarily because if this mutation occurs first, the evolutionary path is immediately terminated. The phenotypic difference between AncBoreotheria S1 and the T52F mutant is obvious; that is, compared with the absorption spectrum of AncBoreotheria S1 with a clear peak at 357 nm, the T52F mutant does not have any absorption peak for the entire region of visible light, showing that the mutant is structurally unstable (Fig. 2A, in red spectrum). If F46T, F49L, F86L, T93P, A114G or S118T occur first, the numbers of trajectories that can accumulate the remaining 6 mutations are 134, 74, 252, 14, 348, 102 and 122, respectively, with various magnitudes of $\Lambda_{\text{max}}$ values during evolution (S2 Fig.). Therefore, a total of 1,032 (20.47%) out of the 5,040 possible trajectories are evolutionarily accessible.

Quantum chemistry of T52F

Interestingly, mouse S1 has a $\lambda_{\text{max}}$ of 359 nm and is UV-sensitive like AncBoreotheria S1 (Fig. 2B, in black spectrum), but its T52F mutant is functional (Fig. 2B, in red spectrum) [10]. At the chemical level, each visual pigment consists of a mixture of pigments with protonated Schiff base nitrogen-linked 11-cis-retinals (PSBR) and those with unprotonated Schiff base nitrogen-linked 11-cis-retinals (SBR) [24]. An SWS1 pigment is UV-sensitive when SBR is more stable than PSBR; otherwise it is blue-sensitive. Moreover, the relative stability of a pigment with SBR and PSBR depends strongly on the water molecules around the 11-cis-retinal [24–26]. The cause for the contrasting roles of T52F in AncBoreotheria S1 and mouse S1 can be seen in two steps.

First, when they are hydrated, AncBoreotheria S1, mouse S1, and their T52F mutants with SBR are all 7 kcal/mol more stable than their PSBR counterparts, which show that all of these pigments are UV-sensitive. In the dehydrated states, the pigments with PSBR achieve similar H-bond interactions; much to our surprise, however, the pigments with SBR become nonfunctional because E113 moves to the protein surface (Fig. 3A). To attain a functional pigment, therefore, water molecules that keep E113 near the 11-cis-retinal are required (Fig. 3B). Second, mouse S1 and AncBoreotheria S1 consist of different types of water channels that allow water molecules to flow from the surface to the interior of the pigments; mouse S1 has two surface openings at T52 and V79 (Figs. 3C, D), but AncBoreotheria S1 has basically one opening at site 52 because of bulky I79. When T52F is introduced into AncBoreotheria S1, the bulky F52 blocks its only water channel; however, in mouse S1 with T52F, the channel opening through V79 is still operational. Hence, the mouse S1 mutant can still be hydrated, but the AncBoreotheria S1 mutant becomes dehydrated and is nonfunctional. In general, such disruptions in water trafficking affecting the H-bond network [24] near the 11-cis-retinal seem to be a major cause for generating nonfunctional pigments.

The mode of phenotypic adaptation of human S1

It would be ideal if we could identify the evolutionary trajectory that actually took place in the evolution of human S1. At present, we can consider the SWS1 pigments of nine primate species for this purpose. A composite evolutionary tree of these pigments and those of six other mammalian species reveals that 1) T93P and A114G, 2) F86L, 3) F49L and S118T and 4) F46T and T52F occurred in that order. Hence, we can identify eight most likely trajectories for describing human S1 evolution (Fig. 4). Then, going back to our mutagenesis results and using the divergence times estimated by others [27,28], we can see that the ancestral
human S1 remained UV-sensitive until about 80 My ago and its $\lambda_{\text{max}}$ increased 20 nm in the next 5 My and another 20 nm in the next 30 My, thus reaching 400 nm by 45 My ago and, finally, the current $\lambda_{\text{max}}$ value was achieved by 30 My ago (Fig. 5, in black and broken trajectories).

When the seven critical amino acid replacements are started with T93P or A114G, the number of trajectories total 450. The eight most likely trajectories are clustered in the center of the 450 trajectories and are characterized by the gradual increase in their $\lambda_{\text{max}}$ values, each step consisting with $|\Delta\lambda_{\text{max}}|$, 25 nm. Further support for this conclusion comes from an analysis of the relatively smaller variance for the amount of change for pathways predicted by the phylogeny as compared with the distribution of all possible variances (S3 Fig.). The gradual $\lambda_{\text{max}}$-shifts might have been necessary because our ancestors switched from their nocturnal life to a diurnal-life style by adjusting their vision slowly to various twilight conditions. The slowly evolving human S1 implies that the evolutionarily acceptable 1,032 trajectories can be subdivided further depending on whether they are characterized by $|\Delta\lambda_{\text{max}}|<25$ nm at every evolutionary step (335 paths, 32.5%) or not (67.5%). In this sense, the evolutionary trajectory of human S1 might have been any one of the 335 evolutionary trajectories (6.65% of 5,040 possible trajectories).

Fig. 1. A phylogenetic tree of 33 representative SWS1 pigments. The numbers after P indicate the $\lambda_{\text{max}}$ values. Divergence times inferred using through the “TimeTree of Life” web server (www.Timetree.org) are shown at the bottom. Black and blue rectangles indicate UV- and blue-sensitive pigments, respectively.

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The molecular basis of the phenotypic adaptation of human S1

Knowing the $\lambda_{\text{max}}$ of AncBoreotheria S1 and those of all 127 mutant pigments allows us to evaluate the $\Delta\lambda_{\text{max}}$ values caused by F46T ($\theta_{46}$)… S118T ($\theta_{118}$) as well as those of two-way ($\theta_{46}\times49$… $\theta_{114}\times118$)… and seven-way ($\theta_{46}\times49\times52\times86\times93\times114\times118$) interactions. To follow this method, it becomes necessary to infer the $\lambda_{\text{max}}$ values of 60 nonfunctional pigments. This is possible because when a new mutation prevents the formation of a functional pigment or when it actually does not shift the $\lambda_{\text{max}}$, the highest level of an epistatic effect can be regarded as zero and the $\lambda_{\text{max}}$ of the mutant pigment can be estimated (see Statistical Analyses, Methods, S1 Table; unstable in bold italics). One special example of this is $\theta_{52} = 0$, which was established using the mouse T52F mutant (Fig. 2B). This procedure is required if we want to obtain all 120 epistatic interaction terms; but, as we will see below,
the $\lambda_{\text{max}}$-shift of each evolutionary path can be recapitulated without obtaining all individual $\theta$ values. The 127 $\theta$ values determined from 127 linear equations show two things. First, the individual mutational effects on the $\Delta \lambda_{\text{max}}$ are very close to zero, i.e. $\theta_{46} = -2, \theta_{105} = -3, \theta_{112} = 0, \theta_{113} = 2, \theta_{114} = 1$ (S2 Table). Second, 21 out of 120 interaction terms show that $|\theta|$ is less than 5 nm, most of which (20 out of 21) reflect significant influences of the interaction between F86L and T93P; in particular, F86L is always involved in generating measurable epistatic interactions (Fig. 6). Consequently, the $\lambda_{\text{max}}$ values for the eight most likely trajectories are explained mostly by the epistatic interactions alone (Fig. 5, in red trajectories). As it was indicated earlier, the evolution of the $\lambda_{\text{max}}$ of human S1 during the last 90 My can be recapitulated without estimating all $\theta$ values individually. For example, when we consider the trajectory with the most conservative functional change, i.e. the smallest $|\Delta \lambda_{\text{max}}|$ values, each step of the observed phenotypic change can be explained solely by epistatic effects (S3 Table).

Discussion

Human S1 evolved gradually from AncBoreotheria S1. The blue-sensitive pigment in African clawed frog (Xenopus laevis) (frog S1, $\lambda_{\text{max}} = 423$ nm) also evolved from the UV-sensitive pigment of the Amphibian ancestor (AncAmphibian S1, $\lambda_{\text{max}} = 359$ nm; Fig. 1) by accumulating F86L, V91I, T93P, V109A, E113D, L116V and S118T [29]. Again, because of their small individual mutational effects on the $\lambda_{\text{max}}$-shift [29], the phenotypic evolution of frog S1 occurred mostly by epistatic interactions. Living in shallow waters, the light environments of the frog ancestors must have been similar to those of human ancestors. Hence, it is likely that the transition from the $\lambda_{\text{max}}$ of AncAmphibian S1 to that of frog S1 occurred gradually.

SWS1 pigments can also take dramatically different modes of evolution. For example, the blue-sensitive pigment in scabbardfish (Lepidopus fitchi) (scabbardfish S1, $\lambda_{\text{max}} = 423$ nm) evolved from the UV-sensitive pigment in the euteleost ancestor (AncEuteleost S1, Fig. 1) by the deletion of F86 [26]. Similarly, the blue-sensitive pigments in bovine (Bos taurus) and wallaby (Macropus eugenii) also evolved essentially in one step by single mutations F86Y [5,30,31]. In addition, C90S makes the UV-sensitive pigments of zebra finch (Taeniopygia guttata) and budgerigar (Melopsittacus undulatus) blue-sensitive [32,33], whereas S90C transforms the blue-sensitive pigments of chicken (Gallus gallus) and pigeon (Columba livia) into UV-sensitive pigment [32].

These observations seem to suggest that there are two distinct modes of evolution among SWS1 pigments. However, even when major $\lambda_{\text{max}}$-shifts are caused by single mutations, epistatic interactions cannot be ignored. For example, the deletion of F86 in AncEuteleost S1 seems to increase the $\lambda_{\text{max}}$ by 59 nm, but the same mutation in the UV-sensitive pigment in a vertebrate ancestor (AncGnathostome S1, $\lambda_{\text{max}} = 360$ nm; Fig. 1) increases the $\lambda_{\text{max}}$ only by 19 nm [26]. Hence, despite the same UV-sensitivity of the two ancestral pigments, their molecular backgrounds are critically different, causing different epistatic effects on the $\lambda_{\text{max}}$-shift. Similarly, S90C causes a wide range of $\lambda_{\text{max}}$-shifts, between -46 nm and 0 nm, depending on SWS1 pigments manipulated [10,21,31–34].

During the period between 45 and 30 My ago, the last two amino acid replacements (F46T and T52F) were in progress in the ancestral human S1 pigment (Fig. 5). This was the time when the LWS pigment in the Boreotherian ancestor (or AncBoreotheria L) achieved two critical changes. First, two AncBoreotheria L copies were generated by a gene duplication event and, second, one of them retained the ancestral $\lambda_{\text{max}}$ (560 nm) and became modern human L and the other decreased its $\lambda_{\text{max}}$ to 350 nm by accumulating three mutations (S180A, Y277F and T285A) and became the modern human M [35–37]. At present, the order of S180A, Y277F and T285A in AncBoreotheria L is not known. However, mutagenesis analyses of AncBoreotheria L show that $\theta_{180} = -5, \theta_{277} = -10, \theta_{285} = -17, \theta_{180 \times 277} = 0, \theta_{180 \times 285} = -2,$
h277 × 625 = 1 and h180 × 277 × 285 = 4 and the sum of these values is −29 nm, which fully explains the evolution of human M [36]. Hence, if S180A and Y277F occurred before T285A, the χmax of the mutant pigment was 545 nm; on the other hand, if T285A was included in the first two mutations, the χmax values of the mutants were 532–538 nm, much similar to that of human M.

Eventually, we are interested in another phenotype that is synthesized by human S1, human M and human L – color vision. Color vision may be best characterized by wavelength discrimination, which describes the minimum wavelength difference (Δλ) along a wide range of wavelengths that human, or any experimental animals, can discriminate [13,17,38]. A typical trichromat having the three cone pigments exhibits Δλ<3 nm, sometimes Δλ<1 nm, along the wavelength between 450 and 625 nm (Fig. 7, Δλ with white circles); in contrast, deuteranopes or protanopes, who are missing functional human M or human L, respectively, have Δλ<5 nm only at around 500 nm (Fig. 7, Δλ with black circles for a deuteranope) [17,39]. Incorporating one or two of the three critical amino acids into AncBoreotheria L, our ancestors have achieved different levels of anomalous trichromatic color vision, conditions known as deuteranomaly, who achieve intermediate wavelength discrimination functions between those of deuteranopes and trichromats. That is, deuteranomalous people can discriminate the wavelength outside of 500 nm, particularly at around 600 nm, much better than deuteranopes (e.g. [40]). Therefore, as the three critical mutations accumulate in one of the duplicated AncBoreotheria L pigments, the U-shaped discrimination function of the human ancestor started to become more flat and eventually reached the more flat discrimination function of modern trichromats (Fig. 7).

The evolution of human S1 could have taken any one of a small proportion of trajectories (335 out of 5,040 possible trajectories, or 6.65%). This observation is consistent with the results of epistatic adaptive evolutionary studies such as antibiotic resistance [2], drug resistance [41], coenzyme evolution [2,42] and coevolution of two ecotypes [43] in microbial systems as well as the evolution of hormone receptors [7], visual pigments [44] and hemoglobins [45] in vertebrates. However, one important implication of epistasis, often neglected, is that forward and reverse mutations can depict
dramatically different epistatic interactions, leading us to erroneous conclusions on the mechanisms of phenotypic adaptation [4,5]. To obtain the correct molecular mechanism of phenotypic adaptation, therefore, it is critical not only to identify appropriate ancestral molecules, engineer and manipulate them but also to recapitulate the evolution of epistatic interactions.

Fig. 5. Eight most likely evolutionary paths used during the evolution of human S1. Human S1 evolution starting either with T93P (in black lines) or with A114G (in broken lines) are shown separately. In the background, a total of 442 evolutionarily accessible trajectories are given. The eight trajectories are characterized by the gradual increase in their $\lambda_{\text{max}}$ values, consisting of each step with $|\Delta \lambda_{\text{max}}| < 25$ nm. The path in red shows the $\lambda_{\text{max}}$ values predicted by considering only epistatic interactions among the seven mutations. $\Delta \lambda_{\text{max}}$ values smaller (in blue) or larger (in orange) than 25 nm are distinguished. doi:10.1371/journal.pgen.1004884.g005

Fig. 6. The $\theta$ values ($|\theta| > 5$ nm) that are generated by epistasis in AncBoreotheria S1. The $\lambda_{\text{max}}$ values of the 127 SWS1 mutant pigments were expressed as that of AncBoreotheria S1 ($\lambda_{\text{max}}$) plus the effects of the appropriate single and multiple amino acid changes on the $\lambda_{\text{max}}$-shift (denoted by a sum of $\theta$ values). These $\theta$ values were estimated by solving a total of 128 simultaneous linear equations with the ancestral $\lambda$ and 127 mutant values. The individual effects of $\theta_{46}$, $\theta_{49}$, $\theta_{52}$, $\theta_{93}$, $\theta_{93}$, and $\theta_{118}$ are $-2$, $-3$, $0$, $0$, $2$, $1$ and $1$ nm, respectively, and their roles in human S1 evolution are negligible compared with those of epistatic interactions. The graph shows the $\theta \pm$ s.e. $^*P < 0.05$. $^{**}P < 0.01$. doi:10.1371/journal.pgen.1004884.g006
Materials and Methods

Ethics statement

The project does not involve any live animals and has been approved by the Institutional Animal Care and Use Committee of the Emory University, in compliance with the USA Public Health Service Policy on Human Care and Use of Laboratory Animals.

Reconstruction of the ancestral SWS1 pigment

To infer the amino acid sequences at various ancestral pigments, we have constructed a composite phylogenetic tree of 33 representative SWS1 pigments (Fig. 1). Using these sequences and those of RH1 pigment of bovine (Bos Taurus; M21606), RH2 pigment of goldfish (Carassius auratus; L11865) and SWS2 pigment of zebrafish (Danio rerio; AB087809) as the outgroup, we inferred the amino acid sequences of SWS1 pigments at various nodes of the phylogenetic tree using PAML [23]. The SWS1 pigments of 33 vertebrate species have been considered as follows: lamprey (Lampetra marinus; U67123), goldfish (C. auratus; D85863), zebrafish (D. rerio; AB087810), scabbardfish (Lepidopus fitchi; FJ443126), tilapia (Oreochromis niloticus; AF191221), cichlid-Mzeb (Maylandia zebra; AF191219), bluefin killifish (Lucania goodei; AY296735), medaka (Oryzias latipes; AB223058), Pacific saury (Cololabis sauro; KP099197), toothfish (Dissostichus mawsoni; AY927651), lampfish (Stenobrachius leucopareus; FJ443127), frog (Xenopus laevis; U23463), salamander (Ambystoma tigrinum; AF038948), zebra finch (Taeniopygia guttata; AF222331), canary (Serinus canaria; AJ277922), budgerigar (Melopsittacus undulatus; Y11787), pigeon (Columba livia; AF149234), chicken (Gallus gallus; M92039), anole (Anolis carolinensis; AF134192), gecko (Gekko gecko; AY024356), sunbeam (Xenopeltes unicolor; FJ497234), human (Homo sapiens; M13295), macaque (Macaca fascicularis; AF158977), squirrel monkey (Saimiri sciureus; U53875), marmoset (Callithrix jacchus; L76201), Bovine (Bos taurus; U92557), mouse (Mus musculus; U63972), squirrel (Sciurus carolinensis; AB223058), Pacific saury (Cololabis saira; KP099197), toothfish (Dissostichus mawsoni; AY927651), lampfish (Stenobrachius leucopareus; FJ443127), frog (Xenopus laevis; U23463), salamander (Ambystoma tigrinum; AF038948), zebra finch (Taeniopygia guttata; AF222331), canary (Serinus canaria; AJ277922), budgerigar (Melopsittacus undulatus; Y11787), pigeon (Columba livia; AF149234), chicken (Gallus gallus; M92039), anole (Anolis carolinensis; AF134192), gecko (Gekko gecko; AY024356), sunbeam (Xenopeltes unicolor; FJ497234), human (Homo sapiens; M13295), macaque (Macaca fascicularis; AF158977), squirrel monkey (Saimiri sciureus; U53875), marmoset (Callithrix jacchus; L76201), Bovine (Bos taurus; U92557), mouse (Mus musculus; U63972), squirrel (Sciurus carolinensis; AB223058), Pacific saury (Cololabis saira; KP099197), toothfish (Dissostichus mawsoni; AY927651), lampfish (Stenobrachius leucopareus; FJ443127), frog (Xenopus laevis; U23463), salamander (Ambystoma tigrinum; AF038948), zebra finch (Taeniopygia guttata; AF222331), canary (Serinus canaria; AJ277922), budgerigar (Melopsittacus undulatus; Y11787), pigeon (Columba livia; AF149234), chicken (Gallus gallus; M92039), anole (Anolis carolinensis; AF134192), gecko (Gekko

Fig. 7. Evolutionary change in the wavelength discrimination by human ancestors. Forty-five My ago, our ancestor possessed almost final product of human S1 and human L and could have achieved color vision of deuteranopes who do not have functional MWS pigment (represented by a discrimination function with black circles). This has changed to the color vision of trichromats with the three cone pigments (represented by a discrimination function with white circles) in the following 15 My. The arrow indicates this evolutionary change. The wavelength discrimination functions of deuteranope and trichromat data are those of A. W. G. in [48] and W. D. W. in [49], respectively.

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forward and reverse primers. Reactions were run on a LI-COR (Lincoln, NE) 4300LD automated DNA sequencer.

To consider the phylogeny of 15 mammalian species, the SWS1 pigments of six additional species have been included: chimpanzee (Pan troglodytes; AF093433), gorilla (Gorilla gorilla gorilla; XM_004046176), orangutan (Pongo abelii; XM_002019421), gibbon (Nomascus leucogenys; XM_003261297), taxier (Tarsius bancanus; AB111463) and Aye-Aye (Daubentonia madagascariensis; EF667285). TimeTree of Life web server (www.timetree.org) shows that human diverged from macaque, marmoset, taxier, Aye-Aye and mouse about 30, 45, 75, 80 and 90 My ago, respectively [27,28]. Then, following the parsimony assumption that minimizes the number of amino acid replacements, the possible evolutionary pathways for the seven amino acid replacements have been determined.

The in vitro assay
The opsin cDNA clones were expressed in COS1 cells by transient transfection [46]. The pigments were regenerated by incubating the opsins with 11-cis-retinal [a gift from Dr. Rosalie K. Crouch at Storm Eye Institute, Medical University of South Carolina and National Eye Institutes] and were purified using immobilized ID4 (The Culture Center, Minneapolis, MN) in buffer W1 (50 mM N-2-hydroxyethyl) piperazine-N’-2-ethanesulfonic acid (HEPES) (pH 6.6), 140 mM NaCl, 3mM MgCl₂, 20% (w/v) glycerol and 0.1% dodecyl maltoside). UV visible methods are identical.

h

hh

hh

\[ Y = CX \]

And the estimates of the effects and interactions are obtained by solving the equation for X, i.e., \( \hat{X} = (C^T C)^{-1}(C^T Y) \)

Since the estimates are linear functions of the observed \( \lambda_{\text{max}} \) values then the variance of each estimate is a function of the variances of the \( \lambda_{\text{max}} \) values and the coefficient associated with each parameter. For this analysis the variances for \( \lambda_{\text{max}} \) are assumed to be 1 and covariance 0. Given the variance-covariance matrix for \( Y = V \) then the standard errors of the estimates are on the diagonal of \( S_e \).

\[ Se = \sqrt{\left(C^T V^{-1} C\right)^{-1}} \]

Modeling of protein structures
The structures of AncBoreotheria S1 and mouse S1 pigments were obtained by 1) applying homology modeling (Modeller 9v7, www.salilab.org/modeller) to bovine rhodopsin (pdb code: 1U19), 2) adding the missing hydrogen atoms, water molecules, and 11-cis-retinal, and 3) optimizing them first at pure AMBER96 force field level [http://ambermd.org] and then using hybrid quantum mechanical/molecular mechanical (QM/MM) calculations in the ONIOM electronic embedding scheme (QM = B3LYP/6-31G*; MM = AMBER) [24]. The QM part of the calculations involves 11-cis-retinal with the covalently bound Schiff base (SB) nitrogen, or NH moiety, and E113 along with hydrogen link atoms, in which either the SB nitrogen or one of the carboxyl O atoms of the E113 is protonated, resulting retinal with protonated SB nitrogen (or PSBR) or retinal with unprotonated SB nitrogen (or SBR), respectively.

Supporting Information
S1 Fig Evolutionary trajectories with premature termination. Each termination event due to unstable protein structure is indicated by a filled red square. (EPS)

S2 Fig All accessible evolutionary trajectories with seven amino acid substitutions in AncBoreotheria S1. Six levels of \( \lambda_{\text{max}} \) values are color coded and the possibilities of whether or not a path includes a large phenotypic change (\( \Delta \lambda_{\text{max}}>25 \) nm) are also indicated. (EPS)

S3 Fig The variances of \( \Delta \lambda_{\text{max}} \) values along an evolutionary path based on whether or not it contains \( \Delta \lambda_{\text{max}}>30 \) nm. The green and blue columns indicate paths contain \( \Delta \lambda_{\text{max}} <30 \) nm and \( \Delta \lambda_{\text{max}} >30 \) nm, respectively. The two red bars indicate 8 paths derived from Fig. 4, four have variances that range from (46.2–49.2) and four range from (85.9–89.9). (EPS)
S4 Fig The amino acid sequences of AncBoreotheria S1, pigment g and human S1. Pigment g was reconstructed previously [21]. Dots indicate the identity of the amino acids with those of AncBoreotheria S1. The amino acids in red are taken from those of SWS1 pigments in mammals, whereas those indicated by orange shade and seven critical amino acid residues (star) are indicated.

S1 Table The amino acid changes, $\lambda_{\text{max}}$ values and $\lambda_{\text{shift}}$-shifts caused by the amino acid changes in AncBoreotheria S1.

(SPDF)

S2 Table The $\lambda_{\text{max}}$-shifts caused by the single and multiple amino acid changes.

(PDF)

References