

EXPLORATIONS: AN OPEN INVITATION TO BIOLOGICAL ANTHROPOLOGY

Editors: Beth Shook, Katie Nelson, Kelsie Aguilera and Lara Braff

American Anthropological Association
Arlington, VA

2019



Explorations: An Open Invitation to Biological Anthropology is licensed under a Creative Commons Attribution-NonCommercial 4.0 International License, except where otherwise noted.

ISBN – 978-1-931303-63-7

www.explorations.americananthro.org

9. Early Hominins

Coauthors:

Kerryn Warren, Ph.D., University of Cape Town

Lindsay Hunter, Ph.D., University of Witwatersrand

Navashni Naidoo, M.Sc., University of Cape Town

Silindokuhle Mavuso, M.Sc., University of Witwatersrand

Kimberleigh Tommy, M.Sc., University of Witwatersrand

Rosa Moll, M.Sc., University of Witwatersrand

Nomawethu Hlazo, M.Sc., University of Cape Town

Learning Objectives

- Define what is meant by “hominin”
- Understand what is meant by “derived” and “primitive” traits and why this is relevant for understanding early hominin evolution
- Understand changing paleoclimates and paleoenvironments during early human evolution, and contextualize them as potential factors influencing adaptations during this time
- Describe the anatomical changes associated with bipedalism in early hominins, and the implications for changes in locomotion
- Describe the anatomical changes associated with dentition in early hominins, and its implication for diet in the Plio-Pleistocene
- Describe several early hominin genera and species: their currently understood date and geographic expanse and (briefly) what we know about them
- Describe the earliest stone tool techno-complex and what this implies about the transition from early hominins to our genus

DEFINING HOMININS

It is through our study of our hominin ancestors and relatives that we are exposed to a world of “might have beens”: of other paths not taken by our species, other ways of being human. But in order to better understand these different evolutionary trajectories, we must first define the terms we are using. If an imaginary line were drawn between ourselves and our closest relatives, the great apes, **bipedalism** (or habitually walking upright on two feet) is where that

line would be. **Hominin**, then, means everyone on “our” side of the line: humans and all of our extinct bipedal ancestors and relatives since our divergence from the **last common ancestor (LCA)** with chimpanzees.

Historic interpretations of our evolution, prior to our finding of early hominin **fossils**, varied. Debates in the mid-1800s regarding hominin origins focused on two key issues:

1. Where did we evolve?
2. Which traits evolved first?

Charles Darwin hypothesized that we evolved in Africa, convinced that we shared greater commonality with chimpanzees and gorillas on the continent. Others, such as Ernst Haeckel and Eugene Dubois, insisted that we evolved in Eurasia, closer in affinity to orangutans, and where, until the discovery of the Taung Child in South Africa in 1924, all humanlike fossils (of Neanderthals and *Homo erectus*) had been found. Adding to this debate was the discovery of the Piltdown Man in England, which turned out later to be a forgery of a modified orangutan mandible and medieval human skull.

Within this conversation, naturalists and early **paleoanthropologists** (people who study human evolution) speculated as to which human traits came first. These included the evolution of a big brain (**encephalization**), the evolution of the strange way in which we move about on two legs (**bipedalism**), and the evolution of our strange flat faces and small teeth (indications of dietary change). Original hypotheses suggested that in order to be motivated to change diet and move about in a bipedal fashion, the large brain needed to have evolved first. And, until research picked up in Africa, fossil finds of species mentioned above predominantly had larger heads.

We now know that bipedal locomotion is one of the first things that evolved in our lineage, with early relatives having small brains and more apelike dentition. In this chapter, we will tease out the details of what this looks like **morphologically** (i.e., what physical traits researchers look for on the skeleton and teeth).

We also know that early human evolution occurred in a very complicated fashion. We have multiple species (multiple genera), diverse in the extent to which they move like us and the diets they subsist on. Specimen finds have been made all along the **East African Rift System (EARS)** (in Ethiopia, Kenya, Tanzania, and Malawi), in limestone caves in South Africa, and in Chad. Dates of these early relatives range from around 7 million years ago (mya) to around 1 mya, overlapping temporally with members of our genus, *Homo* (Figure 9.1).



Figure 9.1 East Africa Rift System.

Yet there is still so much to understand. Modern debates now look at the relatedness of these species to us and to one another. Discussions as to which of these species were able to make and use tools continue. Every **site** discovery in the patchy hominin fossil record tells us more about our evolution. New scientific techniques provide us with insight into the diets, environments, and lifestyles of these ancient relatives that were not available to researchers even 10 years ago.

A Note on Brain Size

It is worth noting that while brain size expansion is seen primarily in our genus, *Homo*, earlier hominin brain sizes were highly variable between and within taxa, from 300 cc (cranial capacity, cm^3), estimated in *Ardipithecus*, to 550 cc, estimated in *Paranthropus boisei*. The lower estimates are well within the range of variation of nonhuman extant Great

Apes, and body size variability also plays a role in the interpretation of whether brain size could be considered large or small for a particular species or specimen.

Increases in brain size do not necessarily correlate with an increase in intelligence in animals, especially if body size is not taken into consideration. However the brain is an expensive tissue to build and maintain. Researchers therefore argue that the cost of maintenance must yield some evolutionary benefit. This is more easily understood in hominins where the stone tool record (an indication of behavior and intelligence) is well associated with a species.

Ancestral and Derived Traits

In Chapter 5, you were introduced to ways of organizing living **taxa**. In the past, **taxonomies** were primarily based on morphology (i.e., the physical features of organisms). Today they are tied to known relationships based on molecular **phylogeny** (such as based on DNA), or a combination of the two. This technique is complicated when applied to living taxa, but it becomes immensely more difficult when we seek to categorize ancestor-descendant relationships in long-extinct forms, where molecular information is no longer preserved. In many ways we find ourselves falling back on morphological comparisons (often on teeth and partially fossilized skeletal material) in the absence of genetic material.

It is here that we turn to the related concepts of **cladistics** and **phylogenetics**. Cladistics groups organisms according to their last common ancestors based on shared **derived** traits. These are traits (in the case of early hominins, morphological) that are evolved, differing from those seen in earlier populations or forms. These new or modified traits or characteristics provide evidence of evolutionary relationships, and organisms with the same derived traits are grouped in the same **clade** (Figure 9.2). For example, using feathers as a trait which groups pigeons and ostriches into the clade “birds.” A good example we will see in this chapter is the grouping of what is known as the “Robust Australopithecines,” whose cranial and dental features differ from those of other hominins of a similar or earlier time period, and can therefore be considered derived.

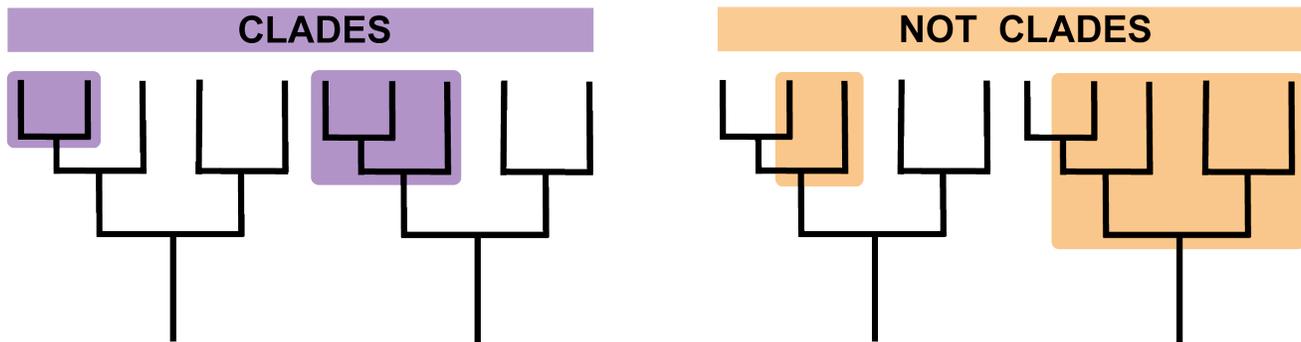


Figure 9.2 Clades refer to a group of species or taxa that share a common ancestor.

It is also worth noting that species designations for early hominin specimens are often highly contested. This is, in part, due to the fragmentary nature of the fossil record, the large timescale (in millions of years) with which paleoanthropologists need to work, and because of the difficulty in evaluating whether morphological differences and similarities are due to meaningful phylogenetic or biological differences or subtle differences/variation in niche occupation or time. In other words, do morphological differences indicate different species? How would classifying species in the paleoanthropological record compare with classifying living species today, where we can sequence their genomes and observe their lifestyles?

There are also broader philosophical differences among researchers when it comes to paleo-species designations. Some scientists, known as “**lumpers**,” argue that large variability is expected among multiple populations in a given species over time. These researchers will therefore prefer to “lump” specimens of subtle differences into single taxa. Others, known as “**splitters**,” argue that species variability can be measured and that even subtle differences can imply differences in niche occupation that is extreme enough to mirror modern species differences. In general, splitters would consider geographic differences among populations as meaning that a species is **polytypic** (i.e., capable of interacting and breeding biologically, but with morphological population differences). This is worth keeping in mind when learning about why species designations may be contested.

This further plays a role in evaluating ancestry. Debates over which species “gave rise” to which continue to this day. It is common to try to create “lineages” of species: in other words, where one species evolves into another over time. We refer to these as **chronospecies**. Constructed hominin phylogenetic trees are routinely variable, changing with new specimen discoveries, new techniques in evaluating and comparing species, and, some have argued, nationalist or biased interpretations of the record. More recently, some researchers have shifted away from “treelike” models of ancestry toward more nuanced metaphors such as the “braided stream,” where some levels of interbreeding among species and populations are seen as natural processes of evolution.

Finally, it is worth considering the process of fossil discovery and publication. Some fossils are easily diagnostic to a species level and allow for easy and accurate interpretation. Some, however, are more controversial. This could be because they do not easily preserve or are incomplete, making it difficult to compare and place within a specific species (e.g., the patella or knee bone). Researchers often need to make several important claims when announcing or publishing a find: a secure date (if possible), clear association with other finds, and an adequate comparison among multiple species (both extant and fossil). It is therefore not uncommon for the scientific community to know that an important find was made years before it is scientifically published.

PALEOENVIRONMENT AND HOMININ EVOLUTION

There is more to paleoanthropology than digging up and grouping fossil hominins: the discipline seeks to explain and understand the evolution of our ancestors’ behavior and morphology. There is no doubt that one of the major drivers (selective pressures) in hominin evolution is the environment. Large-scale changes in global and regional climate, as well as the environment, are all linked to hominin diversification, dispersal, and extinction.

Environmental reconstructions often use modern analogues. Let us take, for instance, the hippopotamus. It is an animal that thrives in environments which have abundant water to keep its skin cool and moist. If the environment for some reason becomes drier, it is expected that hippopotamus populations will reduce. If a drier environment becomes wetter, it is possible that hippopotamus populations may be attracted to the new environment and thrive. Such instances have occurred multiple times in the past, and the bones of some animals (like the hippopotamus) that are sensitive to these changes give us insights into these events.

Reconstructing **paleoenvironments** relies on a range of techniques, which vary depending on whether research interests focus on local changes or more global environmental changes/reconstructions. For local environments (reconstructing those of a single site or region), looking at the **faunal assemblages** (collections of fossils of other animals found at a site) and comparing them to animals found in certain modern environments allow us to determine if the environments in the past mirror those seen today in the region. Changes in the faunal assemblages, when they occur and how they occur, tell us about past environmental changes. Other techniques are also useful in this regard. **Isotopes** of these fauna, for instance, tell us about the relative diets of individual fauna (e.g., differentiating between eating

more grassland-heavy or bushland/tree-heavy diets using carbon isotopes) and whether the environment of individual animals was wetter or drier than the present day (e.g., nitrogen isotopes).

Global climatic changes in the distant past, which fluctuated between being colder and drier and warmer and wetter on average, would have global implications for environmental change (Figure 9.3). These can be studied using marine core and terrestrial soil data, and by comparing these lines of evidence across multiple localities/sites/regions. These techniques allow us to use chemistry (such as nitrogen and oxygen isotopes in shells and sediments) or pollen grains (which show directly the kinds of **flora** surviving in an environment at a specific time period). This means that there are multiple lines of evidence which allow us to visualize global trends over millions of years. Although it should be mentioned that the direction and extent of these changes could differ by geographic region.

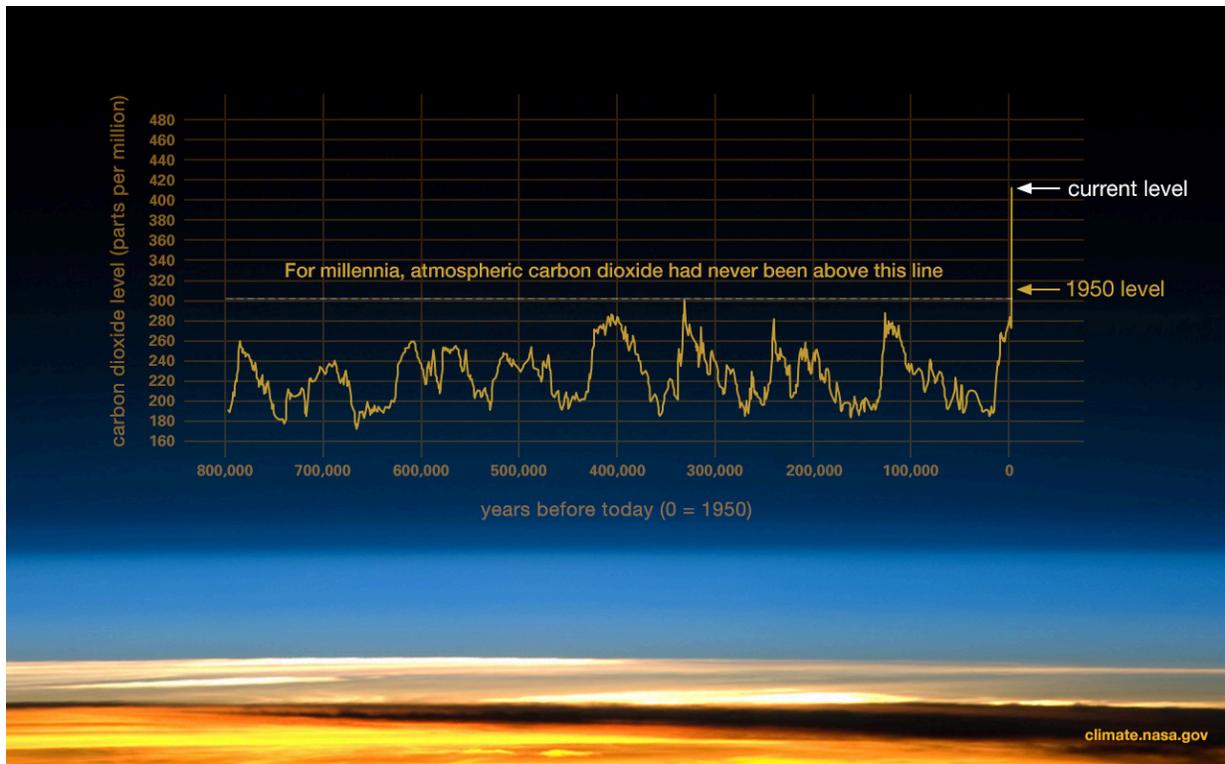


Figure 9.3 This graph, based on the comparison of atmospheric samples contained in ice cores and more recent direct measurements, illustrates how atmospheric CO₂ has fluctuated over time and increased sharply since the Industrial Revolution.

Both local and global climatic/environmental changes have been used to understand parameters affecting our evolution. There are numerous hypotheses regarding how climate has driven and continues to drive human evolution. Environmental change acts as an important keystone in hypotheses regarding the onset of several important hominin traits which are seen in early hominins and which are discussed in this chapter. Namely, the environment has been interpreted as:

1. the driving force behind the evolution of bipedalism (terrestrial locomotion on two legs),
2. the changing and diversifying of early hominin diets, and
3. the diversification of multiple early hominin species.

Here, we will explore the five most popular hypotheses.

Savannah Hypothesis (or Aridity Hypothesis)

The hypothesis: This popular theory was first penned by Charles Darwin and supported by anthropologists like Raymond Dart. It suggests that the expansion of the savannah (or less densely forested, drier environments) forced early hominins from an **arboreal** lifestyle (one living in trees) to a terrestrial one, where bipedalism was a more efficient form of locomotion (Figure 9.4). This hypothesis stems from the idea that the Last Common Ancestor (LCA) between us and chimpanzees was a knuckle-walking quadruped like chimpanzees and gorillas. However, this idea was supported by little fossil or paleoenvironmental evidence and was later refined as the **Aridity Hypothesis**. The Aridity Hypothesis states that the long-term **aridification** and, thereby, expansion of savannah biomes were drivers in diversification in early hominin evolution. It advocates particularly for periods of accelerated aridification leading to early hominin speciation events.

The evidence: While early bipedal hominins are often associated with wetter, more closed environments (i.e., not supporting the Savannah Hypothesis), both marine and terrestrial records seem to support general cooling, drying conditions, with isotopic records indicating an increase in grasslands (i.e., colder and wetter climatic conditions) between 8 mya and 6 mya across the African continent. This can be contrasted with later climatic changes derived from aeolian dust records (sediments transported to the site of interest by wind), which demonstrate increases in seasonal rainfall between 3 mya and 2.6 mya, 1.8 mya and 1.6 mya, and 1.2 mya and 0.8 mya.



Figure 9.4 The African savannah grew during early hominin evolution. This may have forced early hominins from an arboreal lifestyle to a terrestrial one, where bipedalism was a more efficient form of locomotion.

Interpretation(s): Despite a relatively scarce early hominin record, it is clear that two important factors occur around the time period in which we see increasing aridity. Firstly, diversification of taxa, where high morphological variation between specimens has led to the naming of multiple hominin genera and species. Secondly, the earliest hominin fossils appear to have traits associated with bipedalism and are dated to around the drying period (as based on isotopic records). Some have argued that it is more accurately a combination of bipedalism and arboreal locomotion, which will be discussed later. However, the local environments in which these early specimens are found (as based on the faunal assemblages) do not appear to have been dry.

Turnover Pulse Hypothesis

The hypothesis: In 1985, paleontologist Elisabeth Vrba noticed that in periods of extreme and rapid climate change, **ungulates** (hoofed mammals of various kinds) that had generalized diets fared better than those with specialized diets. **Specialist** eaters (eating and relying primarily on specific food types) faced extinction at greater rates than their **generalist** (able to eat more varied and variable diets) counterparts because they were unable to adapt to new environments. This meant that specialist eater species were often confined to isolated areas with dwindling resources, whereas generalist eaters were able to move out across the landscape in search of new food sources. Thus, periods with extreme climate change would be associated with high **faunal turnover**: that is, the extinction of many species and the speciation, diversification, and migration of many others to occupy various niches.

The evidence: The onset of the **Quaternary Ice Age**, between 2.5 mya and 3 mya, brought extreme global, cyclical **interglacial** and **glacial** periods (warmer, wetter periods with less ice at the poles, and colder, drier periods with more ice near the poles). Faunal evidence from the Turkana basin in East Africa indicates multiple instances of faunal turnover and extinction events, in which global climatic change resulted in changes from closed/forested to open/grassier habitats at single sites. Similarly, work in the Cape Floristic Belt of South Africa shows that extreme changes in climate play a role in extinction and migration in ungulates. While this theory was originally developed for ungulates, its proponents have argued that it can be applied to hominins as well. However, the link between climate and speciation is only vaguely understood.

Interpretation(s): While the evidence of rapid faunal turnover among ungulates during this time period appears clear, there is still some debate around its usefulness as applied to the paleoanthropological record. Specialist hominin species do appear to exist for long periods of time during this time period, yet it is also true that *Homo*, a generalist genus with a varied and adaptable diet, ultimately survives the majority of these fluctuations, and the specialists appear to go extinct.

Forest Hypothesis

The hypothesis: Based on contrasting environmental evidence to the Savannah hypothesis, Rayner et al. (1993) hypothesized that forested environments, rather than savannahs, were a key influence on the development of bipedalism in hominins. Unlike the Savannah Hypothesis, one criterion for this may be that the last common ancestor (LCA) between chimpanzees and us used an arboreal form of bipedal locomotion (i.e., walking along branches using the arms for stability), similar to orangutans, and was not a knuckle-walker like contemporary chimpanzees.

The evidence: Pollen evidence from the site of Makapansgat in South Africa indicated that around the time early hominins occupied the area, it was a closed, wooded environment. Similarly, the earliest evidence for bipedalism occurs in specimens (associated with taxa such as *Orrorin* and *Ardipithecus* spp. as well as *Australopithecus anamensis*) found in sites with evidence of **closed habitats**. Furthermore, evidence of knuckle-walking in older hominin species is sorely lacking or highly contested.

Interpretation(s): This hypothesis can be considered in contrast to the Savannah Hypothesis, and appears to be evidence-based. However, it is worth noting that preservation and resulting fossilization might be better in these kinds of environments, biasing this interpretation of the fossil record. Evidence for knuckle-walking in our more distant ancestors is also highly contested.

Variability Selection Hypothesis

The hypothesis: This hypothesis was first articulated by paleoanthropologist Richard Potts (1988). It links the high amount of climatic variability over the last 7 million years to both behavioral and morphological changes. Unlike previous notions, this hypothesis states that hominin evolution does not respond to habitat-specific changes or to specific aridity or moisture trends. Instead, long-term environmental unpredictability over time and space influenced morphological and behavioral adaptations that would help hominins survive, regardless of environmental context. The Variability Selection Hypothesis states that hominin groups would experience varying degrees of natural selection due to continually changing environments and potential group isolation. This would allow certain groups to develop genetic combinations that would increase their ability to survive in shifting environments. These populations would then have a genetic advantage over others that were forced into habitat-specific adaptations.

The evidence: Similar evidence as for the Turnover Pulse Hypothesis: large climatic variability, and higher survivability of generalists versus specialists. However, this hypothesis accommodates for larger time-scales of extinction and survival events.

Interpretation(s): In this way, the Variability Selection Hypothesis allows for a more flexible interpretation of the evolution of bipedalism in hominins, accommodating the discrepancies in evidence between the conflicting Savannah and Forest Hypotheses. This also allows for a more fluid interpretation of the Turnover Pulse Hypothesis, where species-turnover is meant to be more rapid. In some ways, this hypothesis accommodates both environmental data and our interpretations of an evolution toward greater variability among species and the survivability of generalists.

Pulsed Variability Selection Hypothesis

The hypothesis: This hypothesis proposes that the East African Rift System (EARS) and changes in deep lakes are key drivers of diversification during early human evolution. EARS first developed about 10 mya and is responsible for the creation of large **super lakes** (e.g., Lake Baringo and Lake Turkana) within East Africa. The water contents of these lakes were primarily affected by both monsoonal rains and **solar precessional cycles** (cyclical changes in earth's axis rotation that have global climatic effects). According to the Pulsed Variability Selection Hypothesis, human evolution was impacted by 200,000-year cyclical changes in aridity and humidity caused by precessional cycles in this region.

The evidence: Proponents of this hypothesis name three extreme humid phases in East Africa at 2.7 mya to 2.5 mya, 1.8 mya to 1.6 mya, and 1 mya to 0.7 mya. During these periods, changes in precessional cycles increased the monsoonal system, causing more rain in East Africa, thereby increasing lake sizes. This is documented by the increase of diatomaceous lake sediments during these times. These expanded lakes would act as geographic barriers to hominin populations, allowing for changes and diversification in diet and adaptive behavior to the variable regions, even resulting in (allopatric) speciation.

Interpretation(s): High levels of species diversity during these time periods as well as environmental indications of these barriers may allow for an interpretation of allopatric-speciation (i.e., speciation due to geographic barriers) events. However, the degree of inter-specific variability and the extent to which these barriers acted as drivers of speciation are still debated.

Paleoenvironment Consolidated Summary

Some of the hypotheses presented in this section pay specific attention to habitat (Savannah and Forest Hypotheses) while others point to large-scale climatic forces (Pulsed Variability and Variability Selection Hypotheses). Some are complementary (Pulsed Variability and Turnover Pulse Hypotheses), whereas others are directly opposed (Savannah and Forest Hypotheses). Some may be interpreted to describe the evolution of traits such as bipedalism (Savannah and Forest Hypotheses) and others more generally explain the diversification of early hominins (Turnover Pulse and Variability Selection Hypotheses). While there is no consensus as to how environment drove our evolution, it is clear that the environment shaped both habitat and resource availability in ways that would have influenced our early ancestors physically and behaviorally.

DERIVED ADAPTATIONS: BIPEDALISM

The unique form of locomotion exhibited by modern humans, called **obligate bipedalism**, is important in distinguishing our species from the **extant** (living) Great Apes. The ability to walk habitually upright is thus considered one of the defining attributes of the hominin lineage. We also differ from other animals that walk bipedally (such as kangaroos) in that we do not have a tail to balance us as we move.

The origin of bipedalism in hominins has been debated in paleoanthropology, but at present there are two main ideas:

1. that early hominins descended from trees, and so we were a product of an arboreal last common ancestor (LCA); or
2. that our LCA was a terrestrial quadrupedal knuckle-walking species, more similar to extant chimpanzees.

Most research supports the theory of an arboreal LCA (i.e., idea 1) based on skeletal morphology of early hominin genera that demonstrate adaptations for climbing but not for knuckle-walking. This would mean that both humans and chimpanzees can be considered “derived” in terms of locomotion since chimpanzees would have independently evolved knuckle-walking.

There are many current ideas regarding selective pressures that would lead to early hominins adapting upright posture and locomotion. Many of these selective pressures, as we have seen in the previous section, coincide with a shift in environmental conditions, supported by paleoenvironmental data. In general, however, it appears as though early hominins thrived in forested regions, similar to extant great apes, with dense tree coverage, which would indicate an arboreal lifestyle. As the environmental conditions changed and a savannah/grassland environment became more widespread, the tree cover would become less dense, scattered and sparse and bipedalism would become more important.

There are several proposed selective pressures for bipedalism.

1. **Energy conservation:** modern bipedal humans conserve more energy than extant chimpanzees, which are predominantly knuckle-walking quadrupeds when walking over land. While chimpanzees, for instance, are faster than humans terrestrially, they expend large amounts of energy being so. Adaptations to bipedalism include “stacking” the majority of the weight of the body over a small area around the centre of gravity (i.e., the head is above the chest, which is above the pelvis, which is over the knees, which is above the feet). This reduces the amount of muscle needed to be engaged during locomotion to “pull us up,” and allows us to travel longer distances expending far less energy.
2. **Thermoregulation:** less surface area (i.e., only the head and shoulders) is exposed to direct sunlight during the

hottest parts of the day (i.e., midday). This means that the body is exposed to less heat and has less need to employ additional “cooling” through mechanisms such as sweating, which additionally means less water loss.

3. Bipedalism freed up our ancestors’ hands such that they could more easily gather food and carry tools or infants. This further frees the hands for more specialized adaptations associated with the manufacturing and use of tools.

These selective pressures are not mutually exclusive, and bipedality could have evolved from a combination of these selective pressures, in ways that increased the chances of early hominin survival.

Skeletal Adaptations for Bipedalism

Humans, as the only obligate bipedal species among primates, have highly specialized adaptations to facilitate this kind of locomotion (Figure 9.5). Many of these adaptations occur within the soft tissue of the body (e.g., muscles and tendons). However, when analyzing the paleoanthropological record for evidence of the emergence of bipedalism, all that remains is the fossilized bone. Interpretations of locomotion are therefore often based on comparative analyses between fossil remains and the skeletons of extant primates with known locomotor behaviors. These adaptations occur throughout the skeleton and are summarized in Table 9.1.

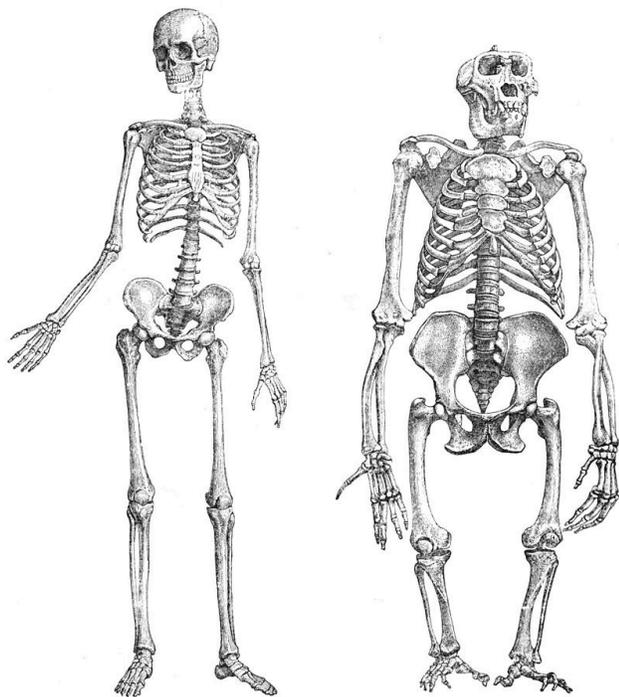


Figure 9.5 Compared to gorillas (right) and other apes, humans (left) have highly specialized adaptations to facilitate bipedal locomotion.

The majority of these adaptations occur in the **postcranium** (the skeleton from below the head). In general, these adaptations allow for greater stability and strength in the lower limb, by allowing for more shock absorption, a larger surface area for muscle attachment, and for the “stacking” of the skeleton directly over the center of gravity to reduce energy needed to be kept upright. These adaptations often mean less flexibility in areas such as the knee and foot.

However, these adaptations come at a cost. Evolving from a non-obligate bipedal ancestor means that the adaptations we have are evolutionary compromises. For instance, the valgus knee (angle at the knee) is an essential adaptation to balance the body weight above the ankle during bipedal locomotion. However, the strain and shock absorption at an angled knee eventually takes its toll, with runners often experiencing joint pain. Similarly, the long neck of the femur absorbs stress and accommodates for a larger pelvis, but is a weak point, resulting in hip replacements being commonplace among the elderly, especially in cases where the bone additionally weakens through osteoporosis. Finally, the S-shaped curve in our spine allows us to stand upright,

relative to the more curved, C-shaped spine of an LCA. Yet the weaknesses in the curves lead to pinching of nerves and back pain. Since many of these problems primarily are only seen in old age, they can potentially be seen as an evolutionary compromise.

Region	Feature	Obligate Biped (<i>H. sapiens</i>)	Non-obligate Biped
Cranium	Position of the foramen magnum	Positioned inferiorly (immediately under the cranium) so that the head rests on top of the vertebral column for balance and support (head is perpendicular to the ground)	Posteriorly positioned (to the back of the cranium). Head is positioned parallel to the ground
Postcranium	Body proportions	Shorter upper limb (not used for locomotion)	Nonhuman apes: Longer upper limbs (used for locomotion)
Postcranium	Spinal curvature	S-curve due to pressure exerted on the spine from bipedalism (lumbar lordosis)	C-curve
Postcranium	Vertebrae	Robust lumbar (lower-back) vertebrae (for shock absorbance and weight bearing). Lower back is more flexible than that of apes as the hips and trunk swivel when walking (weight transmission).	Gracile lumbar vertebrae compared to those of modern humans
Postcranium	Pelvis	Shorter, broader bowl-shaped pelvis (for support); very robust. Broad sacrum with large sacroiliac joint surfaces	Longer, flatter, elongated ilia, more narrow and gracile, narrower sacrum, relatively smaller sacroiliac joint surface
Postcranium	Lower limb	In general, longer, more robust lower limbs and more stable, larger joints <ul style="list-style-type: none"> • Large femoral head and longer neck (absorbs more stress and increases the mechanical advantage). • Valgus angle of knee, positions knee over the ankle and keeps the center of gravity balanced over stance leg during stride cycle (shock absorbance). • Distal tibia (lower leg) of humans has a large medial malleolus for stability. 	In general, smaller, more gracile limbs with more flexible joints <ul style="list-style-type: none"> • Femoral neck is smaller in comparison to modern humans and has a shorter neck. • The legs bow outward, there is no valgus angle of the knee (no "knock-knees"). • The distal tibia in chimpanzees is trapezoid (wider anteriorly) for climbing and allows more flexibility.
Postcranium	Foot	Rigid, robust foot, without a midtarsal break. Non-opposable and large, robust big toe (for push off while walking) and large heel for shock absorbance.	Flexible foot, midtarsal break present (which allows primates to lift their heels independently from their feet), opposable big toe for grasping.

Figure 9.6 Skeletal comparisons between modern humans (obligate bipeds) and non-obligate bipeds (e.g., chimpanzees).

Despite relatively few postcranial fragments, the fossil record in early hominins indicates a complex pattern of emergence of bipedalism. Key features, such as a more anteriorly placed foramen magnum, are argued to be seen even in the earliest discovered hominins, indicating an upright posture. Some early species appear to have a mix of primitive (arboreal) and derived (bipedal) traits, which indicate a mixed locomotion, and a more **mosaic evolution** of the trait. Some early hominins appear to, for instance, have bowl-shaped pelvises (hip bones) and angled femurs suitable for bipedalism, but would retain an opposable **hallux** (big toe) or curved fingers and longer arms (for arboreal locomotion). These mixed morphologies may indicate that earlier hominins were not fully obligate bipeds, thriving in mosaic environments.

It is also worth noting that, while not directly related to bipedalism per se, other postcranial adaptations are evident in the hominin fossil record from some of the earlier hominins. For instance, the hand and finger morphologies of many of the earliest hominins indicate adaptations consistent with arboreality. These include longer hands, more curved metacarpals and phalanges (long bones in the hand and fingers), and a shorter, relatively weaker thumb. This allows for gripping onto curved surfaces during locomotion. The earliest hominins appear to have mixed morphologies for both

bipedalism and arborealism. However, among Australopiths, there are indications for greater reliance on bipedalism as the primary form of locomotion. Similarly, adaptations consistent with tool manufacture (shorter fingers and a longer, more robust thumb, in contrast to the features associated with arborealism) have been argued to appear before the genus *Homo*.

Earliest Hominins: Sahelanthropus and Orrorin

We see evidence for bipedalism in some of the earliest fossil hominins, dated from within our estimates of our divergence from chimpanzees. These hominins, however, also indicate evidence for arboreal locomotion.

The earliest dated hominin find (between 6 mya and 7 mya, based on radiometric dating of volcanic tufts) has been argued to come from Chad and named *Sahelanthropus tchadensis* (Figure 9.7). The initial discovery was made in 2001 by Ahounta Djimdoumbaye and announced in *Nature* in 2002 by a team led by French paleontologist Michel Brunet. The find has a small cranial capacity (360 cc) and has canines smaller than those in extant Great Apes, yet still larger and pointier than those in humans. This implies strongly that, over evolutionary time, the need for display and dominance among males has reduced, as has our sexual dimorphism. A short cranial base and a foramen magnum (the hole through which the spinal cord enters the cranium) that is more humanlike in positioning have been argued to indicate upright walking. However, the inclusion of *Sahelanthropus* in the hominin family has been debated by researchers, since the evidence for bipedalism is based on cranial evidence alone. Researchers have suggested that in order to conclude if it is a truly bipedal species, postcranial remains such as a pelvis or a leg bone would support the idea of upright walking. An unpublished femur (thigh bone) thought to belong to *Sahelanthropus* was discovered in 2001 and could potentially shed light on this topic once it is fully studied. However, the extent to which this femur is hominin-like is currently unknown.

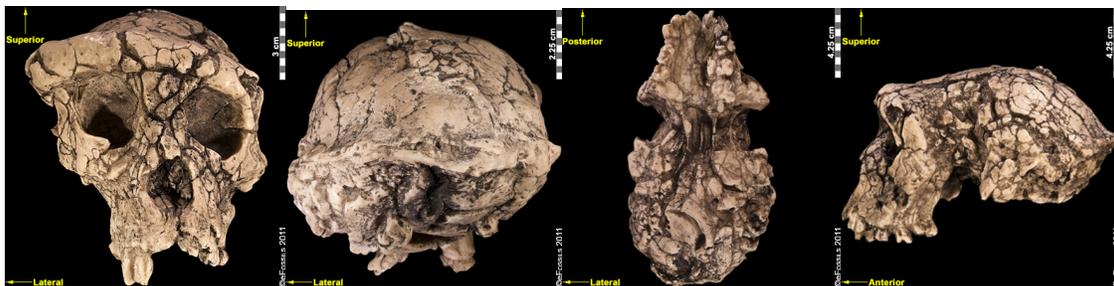


Figure 9.7 *Sahelanthropus tchadensis* exhibits a set of a set of derived features, including a long low cranium, a small ape-sized braincase, and relatively reduced prognathism.

Orrorin tugenensis (Orrorin meaning “original man”; dated to between 6 mya and 5.7 mya) was discovered near Tugen Hills in Kenya in 2000. Smaller **cheek teeth** (molars and premolars) than those in even more recent hominins (i.e., derived), thick enamel, and reduced, but apelike, canines characterize this species. This is the first species which clearly indicates adaptations for bipedal locomotion, with fragmentary leg, arm, and finger bones having been found but few cranial remains. One of the most important elements discovered was a proximal femur, BAR 1002'00. The femur is the thigh bone and the proximal part is that which articulates with the pelvis—it is very important when studying posture and locomotion. This femur indicates that *Orrorin* was bipedal, and recent studies suggest that it walked in a similar way to later **Pliocene** hominins. Some have argued that features of the finger bones suggest potential toolmaking capabilities, although many researchers argue that these features are also consistent with climbing.

Earliest Hominins: The Genus *Ardipithecus*

Another genus, *Ardipithecus*, is argued to be represented by at least two species: *Ardipithecus ramidus* and *Ar. kadabba*.

Ardipithecus ramidus (“ramid” means root in the Afar language) is currently the best known of the earliest hominins (Figure 9.8). Unlike *Sahelanthropus* and *Orrorin*, this species has a large sample size of over 110 specimens from Aramis alone. Dated to 4.4 mya, *Ar. ramidus* was found in Ethiopia (in the Middle Awash region and in Gona). This species was announced in 1994 by American palaeoanthropologist Tim White, based on a partial female skeleton nicknamed “Ardi” (ARA-VP-6/500). Ardi demonstrates a mosaic of ancestral and derived characteristics in the postcrania. For instance, an opposable big toe (hallux), similar to chimpanzees (i.e., “primitive” or more ancestral), which could have aided in climbing trees effectively. However, the pelvis and hip show that she could walk upright (i.e., it is derived), supporting her hominin status. A small brain (300 cc to 350 cc), midfacial projection, and slight prognathism show retained primitive cranial features, but the cheek bones are less flared and robust than in later hominins.



Figure 9.8 Researchers believe that *Ardipithecus ramidus* was able to walk upright, although not as efficiently as later humans. It possessed the musculature required for tree-climbing and while moving quadrupedally, it likely placed weight on the palms of the hands rather than on the knuckles.

Ardipithecus kadabba (the species name means “oldest ancestor” in the Afar language) is known from localities on the western margin of the Middle Awash region, the same locality where *Ar. ramidus* has been found. Specimens include mandibular fragments and isolated teeth as well as a few postcranial elements from the Asa Koma (5.5 mya to 5.77 mya) and Kuseralee (5.2 mya) Members. This species was discovered in 1997 by paleoanthropologist Dr. Yohannes Haile-

Selassie. Originally these specimens were referred to as a subspecies of *Ar. ramidus*. In 2002, six teeth were discovered at Asa Koma and the dental wear patterns confirmed that this was a distinct species, named *Ar. kadabba*, in 2004. One of the postcranial remains recovered included a 5.2 million-year-old toe bone that demonstrated features that are associated with toeing off (pushing off the ground with the big toe leaving last) during walking, a characteristic unique to bipedal walkers. However, the toe bone was found in the Kuseralee Member, and therefore some doubt has been cast by researchers about its association with the teeth from the Asa Koma Member.

Bipedal Trends in Early Hominins

Trends toward bipedalism are seen in our earliest hominin finds. However, many specimens also indicate retained capabilities for climbing. Trends include a larger, more robust hallux; a more compact foot, with an arch; a robust, long femur, angled at the knee; a robust tibia; a bowl-shaped pelvis; and a more anterior foramen magnum. While the level of bipedality in *Salehanthropus tchadenisis* is debated since there are few fossils and no postcranial evidence, *Orrorin tugenensis* and *Ardipithecus* show clear indications of some of these bipedal trends. However, some retained primitive traits, such as an opposable hallux in *Ardipithecus*, indicate some retention in climbing ability.

DERIVED ADAPTATIONS: EARLY HOMININ DENTITION

The Importance of Teeth

Teeth are abundant in the fossil record, primarily because they are already highly mineralized as they are forming, far more so than even bone. Because of this, teeth preserve readily. And, because they preserve readily, they are well-studied and better understood than many skeletal elements. Even in the sparse hominin (and primate) fossil record, teeth are, in some cases, all we have.

Teeth also reveal a lot about the individual from whom they came. We can tell what they evolved to eat, which other species they may be more closely related to, and even, to some extent, the level of sexual dimorphism, or general variability, within a given species. This is powerful information that can be contained in a single tooth. With a little more observation, the wearing patterns on a tooth can tell us about the diet of the individual in the weeks leading up to its death. Furthermore, the way in which a tooth is formed, and the timing of formation, can reveal information about changes in diet (or even mobility) over infancy and childhood, using isotopic analyses. When it comes to our earliest hominin relatives, this information is vital for understanding how they lived.

The purpose of comparing different hominin species is to better understand the functional morphology as it applies to dentition. In this, we mean that something about the morphology of the teeth or masticatory system (which includes jaws), can tell us something about the way in which they were used, and therefore the kinds of foods these hominins ate. When comparing the features of hominin groups, it is worth considering modern analogues (i.e., animals with which to compare) in order to make more appropriate assumptions about diet. In this way, hominin dentition is often compared with that of chimpanzees, gorillas (our two closest relatives), and/or modern humans.

The most divergent group, however, is humans. Humans around the world have incredibly varied diets. Among hunter-gatherers, it can vary from a honey- and plant-rich diet, as seen in the Hadza in Tanzania, to a diet almost entirely reliant

on animal fat and protein, as seen in Inuits in polar regions of the world. We are therefore considered **generalists**, more general than the largely **frugivorous** (fruit-eating) chimpanzee or the **folivorous** (foliage-eating) gorilla.

One way in which all humans are similar is our reliance on the processing of our food. We cut up and tear meat with tools using our hands, instead of using our front teeth (incisors and canines). We smash and grind up hard seeds, instead of crushing them with our hind teeth: molars. This means that, unlike our ape relatives, we can rely more on developing tools to navigate our complex and varied diets. Our brain, therefore, is our primary masticatory organ. Evolutionarily, partially in response to our increased reliance on our hands and brain, our teeth have reduced in size and our faces are flatter, or more **orthognathic**. Similarly, a reduction in teeth and a more generalist dental morphology could also indicate an increase in softer and more variable foods, such as the inclusion of more meat. These trends begin early on in our evolution. The link has been made between some of the earliest evidence for stone tool manufacture, the earliest members of our genus, and the features that we associate with these specimens.

General Dental Trends in Early Hominins

Several trends are visible in the dentition of early hominins. However, worth noting is that all tend to have the same **dental formula**. The dental formula is a method to characterize how many of the different kinds of teeth are present in the mouth. Going from the most anterior (front) of the mouth, this includes the square, flat **incisors**; the pointy **canines**; the small, flatter **premolars**; and the larger hind **molars**. In many primates, from Old World Monkeys to Great Apes, the typical dental formula is 2:1:2:3. This means that if we divide the mouth into quadrants, each should have two incisors, one canine, two premolars and three molars. In total that is eight teeth a quadrant, for a total of 32 teeth. In humans, this number can be variable. Unlike in other apes, it is not uncommon for people to have only two molars in one or more of their quadrants. One explanation for this is that, because of our processed foods, there are fewer dietary constraints, i.e., less pressure to have many teeth for additional processing. Furthermore, with smaller mouths and faces, fewer teeth may be advantageous. All early hominins have the primitive condition shared with other Great Apes.

The morphology of the individual teeth is where we see the most change. Among primates, large incisors are associated with food procurement or preparation (such as biting small fruits), while small incisors indicate a diet which may contain small seeds or leaves (where the preparation is primarily in the back of the mouth). Most hominins have relatively large flat, vertically aligned incisors which **occlude** (touch) relatively well, forming a “bite.” This differs from, for instance, the orangutan, whose teeth stick out (i.e., are **procumbent**).

While the teeth are often sensitive, evolutionarily speaking, with diet, the canines may be misleading in that regard. We tend to associate pointy, large canines with the ripping required for meat, and the reduction (or, in some animals, the absence) of canines as indicative of more herbivorous diets. In humans, our canines are often a similar size to our incisors, and are therefore considered **incisiform** (Figure 9.9). However, among our closest relatives, all have very long, pointy canines, particularly on their upper dentition. This is true even for the Gorilla, which lives almost exclusively on plants, as you have seen in previous chapters. The canines, in these instances, possibly indicate more about social structure and sexual dimorphism than diet.

Early on in human evolution, we see a reduction in canine size. *Sahelanthropus tchadensis* and *Orrorin tugenensis* both have smaller canines than those in extant Great Apes yet still larger and pointier than those in humans or more recent hominins. This implies strongly that, over evolutionary time, the need for display and dominance among males has reduced, as has our sexual dimorphism. In *Ardipithecus ramidus*, there is no obvious difference between male and female canine size, yet they are still slightly larger and pointier than in humans. This implies a less sexually dimorphic social structure in the earlier hominins relative to modern-day chimpanzees and gorillas.

Along with a reduction in canine size is the reduction or elimination of a canine **diastema**: a gap between the teeth on the mandible which allows room for elongated teeth on the maxilla to “fit” in the mouth. Absence of a diastema is an excellent indication of a reduction in canine size. In animals with large canines (such as baboons), there is also often a **honing P3**, where the first premolar (also known as P3 for evolutionary reasons) is triangular in shape, “sharpened” by the extended canine from the upper dentition. Evidence for this is also seen in some of the early hominins such as *Ardipithecus*, where, even though the canines are much smaller and almost the same height as the incisors, they are larger than those in more recent hominins.



Figure 9.9 In humans, our canines are often a similar size to our incisors.

The hind dentition, such as the bicuspid (two cusped) premolars or the much larger molars, are also highly indicative of a generalist diet in hominins. Among the earliest hominins, the molars are larger than we see in our genus, increasing in size to the back of the mouth, and angled in such a way from the much smaller anterior dentition as to give these hominins a **parabolic** (V-shaped) dental arch. This is opposed to our living relatives, and some of the earliest hominins, such as *Sahelanthropus*, whose molars and premolars are relatively parallel between the left and right sides of the mouth, creating a U-shape.

Among more recent early hominins, the molars are relatively large, larger than those in the earliest hominins, and far larger than those in our own genus, *Homo*. Large, short molars with thick **enamel** allow these early cousins of ours to grind away at fibrous, coarse foods, such as sedges which require plenty of chewing. This is further evidenced in the low **cusps**, or ridges, on the teeth, which are ideal for chewing. In our genus, the hind dentition is far smaller than in these early hominins. Our teeth also have medium-size cusps, which allow for both efficient grinding and for tearing/shearing meats.

Understanding the dental morphology has allowed researchers to extrapolate very specific behaviours of early hominins. It is worth noting that while teeth preserve well and are abundant, a slew of other morphological traits additionally provide evidence for many of these hypotheses. Yet there are some traits which are ambiguous. For instance, while there are definitely high levels of sexual dimorphism in *Au. afarensis*, which we will introduce in the next section, the canine teeth are reduced in size, implying that while canines may be useful indicators for sexual dimorphism, it is worth noting other lines of evidence.

Dental Trends in Early Hominins

Trends among early hominins include a reduction in procumbency, reduced hind dentition (molars and premolars), a reduction in canine size (more incisiform with a lack of canine diastema and honing P3), flatter molar cusps, and thicker dental enamel. All early hominins have the primitive dental formula of 2:1:2:3. These trends are all consistent with a generalist diet, incorporating more fibrous foods.

THE GENUS *AUSTRALOPITHECUS*

The Australopithecines are a diverse group of hominins comprised of various species. *Australopithecus* is the given group or genus name. It stems from the Latin word “Australo” meaning “southern” and “Pithecus” a Greek word meaning “ape.”

Within this section, we will outline these differing species' geological and temporal distributions across Africa, unique derived and/or shared traits, and importance in the fossil record.

Between 3 mya and 1 mya, there seems to be differences in dietary strategy between species of hominins designated as Australopithecines, which is evident from the peculiar size of the molars in one of the groups. This pattern of larger posterior dentition (even relative to the incisors and canines) with thick enamel, and cranial evidence for large chewing muscles, is far more pronounced in a group known as the **robust australopithecines**, as opposed to their earlier contemporaries or predecessors, the **gracile australopithecines** and certainly larger than those seen in early *Homo*, which emerges during this time. This pattern of incredibly large hind dentition (and very small anterior dentition), has led people to refer to robust australopithecines as **megadont** hominins (Figure 9.10).



Figure 9.10: Robust Australopithecines (left) had large molars and chewing muscles compared to modern humans (right).

This section has been categorized into “gracile” and “robust” Australopithecines, highlighting the morphological differences between the two groups (which many researchers have designated as separate genera: *Australopithecus* and *Paranthropus*, respectively) and then focusing on the individual species. It is worth noting, however, that not all researchers accept these clades as biologically or genetically disparate, with some researchers insisting that the relative gracile and robust features found in these species are due to parallel evolutionary events, toward similar dietary niches.

Despite this genus's ancestral traits and small cranial capacity, all members show evidence of bipedal locomotion. It is generally accepted that *Australopithecus* species display varying degrees of arborealism and bipedality: These individuals were walking on the ground on two legs but were probably still comfortable with climbing trees.

Gracile Australopithecines

The section below describes individual species from across Africa. These species have coined the term “**gracile**” **australopithecines** because of the less exaggerated, smaller, and less robust features seen in the divergent “robust” group. Numerous Australopithecine species have been named, but some are still only based on a handful of fossil finds, whose designations are controversial.

East African Australopithecines

East African Australopithecines are found throughout the EARS, a system running from Malawi to Ethiopia, and include the earliest species associated with this genus. Numerous fossil-yielding sites, such as Olduvai, Turkana, and Laetoli, have excellent, datable stratigraphy, owing to the layers of **volcanic tufts** which have accumulated over millions of years. These tufts may be dated using absolute dating techniques, such as Potassium-Argon dating. This means that it is possible to know a relatively refined date for any fossil if the **context** of that find is known. Similarly, comparisons between the faunal assemblages of these stratigraphic layers have allowed researchers to chronologically identify environmental changes.

The earliest known Australopithecine is dated to 4.2 mya to 3.8 mya. ***Australopithecus anamensis*** (after “Anam,” meaning “lake” from the Turkana region in Kenya) is currently found from sites in the Turkana region (Kenya) and Middle Awash (Ethiopia) (Figure 9.11). Recently, a 2019 find from Ethiopia, named MRD, after Miro Dora where it was found, was discovered by an Ethiopian herder named Ali Bereino. It is one of the most complete cranial finds of this species. A small brain size (370 cc), relatively large canines, projecting cheekbones, and primitive earholes show more primitive features as compared to those of more recent Australopithecines. The most important element discovered associated with this species that indicates bipedalism is a fragment of a tibia (shinbone), which demonstrates features associated with weight transfer during bipedal walking. Similarly, the earliest found hominin femur belongs to this species. Primitive traits in the upper limb (such as the humerus) indicate some retained arboreal locomotion. Some researchers suggest that *Au. anamensis* is an intermediate form of the chronospecies which becomes *Au. afarensis*, evolving from *Ar. ramidus*. However, this is debated, with other researchers suggesting morphological similarities and affinities with more recent species instead. Almost 100 specimens, representing over 20 individuals, have been found to date.



Figure 9.11 As seen in this mandible, *Australopithecus anamensis* had relatively large canine teeth.

Au. afarensis is one of the oldest and most well-known australopithecine species and consists of a large number of fossil remains. *Au. afarensis* (which means “from the Afar region”) is dated to between 2.9 mya and 3.9 mya and found in sites all along the EARS system, in Tanzania, Kenya, and Ethiopia (Figure 9.12). The most famous individual stemming from this species is a partial female skeleton discovered in Hadar (Ethiopia), later nicknamed “Lucy,” after the psychedelic Beatles song “Lucy in the Sky With Diamonds,” which was played in celebration of the find. This skeleton was found in 1974 by Donald Johanson and dates to approximately 3.2 mya (Figure 9.13). In addition, in 2002 a juvenile of the species was found by Zeresenay Alemseged and given the name “Selam” (meaning “peace,” DIK 1-1), though it is popularly known as “Lucy’s Child” or as the “Dikika Child.” Similarly, the “Laetoli Footprints” (discussed in Chapter 7) have drawn much attention.



Figure 9.12 Artistic reconstructions of *Australopithecus afarensis* by artist John Gurche. Female “Lucy” is left and a male is on the right.

Au. afarensis has reduced canines and molars relative to Great Apes, but larger than in modern humans (indicative of a generalist diet), a prognathic face, and robust facial features which indicate relatively strong chewing musculature

(compared with *Homo*), but less extreme than in *Paranthropus*. Despite a reduction in canine size in this species, large overall size variation indicates high levels of sexual dimorphism in this species.

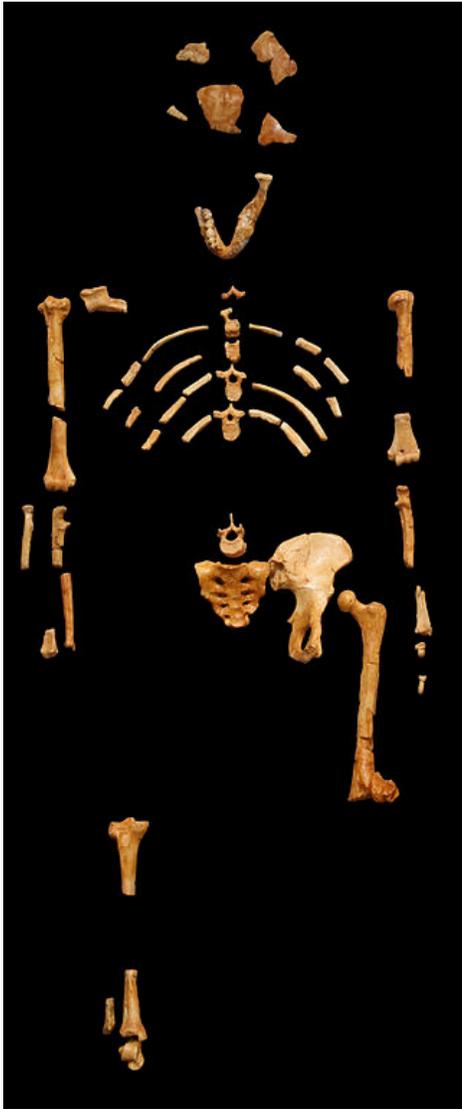


Figure 9.13 The humanlike femoral neck, valgus knee, and bowl-shaped hip seen in the “Lucy” skeleton indicates that *Australopithecus afarensis* was bipedal.

Skeletal evidence indicates that this species was bipedal, primarily through examining the pelvis and lower limb which demonstrate a humanlike femoral neck, a valgus knee, and bowl-shaped hip. More evidence of bipedalism is found not in the skeleton but in the footprints of this species. *Au. afarensis* is associated with the Laetoli Footprints, a 24-meter trackway of hominin fossil footprints preserved in volcanic ash discovered by Mary Leakey in Tanzania and dated to 3.5 mya to 3 mya. This set of prints is thought to have been produced by three bipedal individuals as there are no knuckle imprints, no opposable big toes, and a clear arch is present. The infants of this species are thought to have been more arboreal than the adults as discovered through analyses of the foot bones of the Dikika Child dated to 3.32 mya.

Although not found in direct association with stone tools, potential evidence for cut marks on bones, found at Dikika, and dated to 3.39 mya indicates a potential temporal/geographic overlap between meat-eating, tool use, and this species. However, this evidence is fiercely debated. Others have associated the earliest tool finds from Lomekwi, Kenya, temporally (3.3 mya) and in close geographic proximity to this species.

Contested Species

Many named species are highly debated and/or newly described. Often these species are argued to have specimens associated with a more variable *Au. afarensis* or *Au. anamensis* species. Sometimes these specimens from which these species are named are dated to times, or found in places, when there are “gaps” in the palaeoanthropological record. Often these are argued to represent chronospecies or variants of *Au. afarensis*. However, it is possible that, with more discoveries, these species definitions will hold.

Australopithecus bahrelghazali (named after the Bahr el Ghazal river valley in the southern region of Borkou-Ennedi Tibesti in Chad; and not within the

EARS system) is dated to within the time period of *Au. afarensis* (3.6 mya). Yet this discovery holds such significance in the field as this was the first Australopithecine to be discovered in Chad in central Africa (unlike those in the usual east Africa and South Africa). Researchers argue that the **holotype**, whom discoverers have named “Abel,” falls under the range of variation of *Au. afarensis*, and therefore that *A. bahrelghazali* does not fall into a new species. If a member of *Au. afarensis*, this would exceed the geographic range of the species even further.

On a different note, ***Australopithecus deyiremada*** (meaning “close relative” in the Ethiopian language of Afar) is dated to 3.5 mya to 3.3 mya, and is based on fossil mandible bones discovered in 2011 in Woranso-Mille (in the Afar region of Ethiopia) by Yohannes Haile-Selassie, an Ethiopian paleoanthropologist. The discovery indicated smaller teeth with thicker enamel than seen in *Au. afarensis*, with a potentially harder diet and a larger mandible and more projecting cheekbones than in *Au. afarensis*. The discoverers believe that this was a pivotal find in the palaeoanthropology field as it shows evidence of more than one closely related hominin species occupying the same region at the same temporal

period. If so, it may also imply that other *Au. afarensis* specimens have been incorrectly designated and researchers should attempt to better understand the variability among these species. However, others have argued that this species has been prematurely identified, and more evidence is needed before splitting the taxa, since the variation appears subtle and may be due to slightly different niche occupations between populations over time.

Australopithecus garhi is another species found in the Middle Awash region of Ethiopia and currently dated to 2.5 mya, and is therefore younger than *Au. afarensis*. Researchers have suggested it fills in a much needed temporal “gap” between hominin finds in the region. It has a relatively large cranial capacity (450 cc) and larger hind dentition than seen in other gracile Australopithecines. The discoverers argue that the postcrania is similarly different, and the femur of a fragmentary partial skeleton (argued to belong to *Au. garhi*) indicates that this species may be longer limbed than *Au. afarensis*, although still able to move arboreally. However, this species is not well documented or understood, and is based on only several fossil specimens. More astonishingly, crude/primitive stone tools resembling Oldowan (which will be described later) have been found in association with *Au. garhi*. While lacking some of the features of the Oldowan, this is one of the earliest technologies found in direct association with a hominin.

Kenyanthropus platyops (the name “platyops” refers to its flatter-faced appearance) is a highly contested genus/species designation, of a specimen (KNM-WT 40000) from Lake Turkana in Kenya, discovered by Maeve Leakey in 1999 (Figure 9.14). Dated to between 3.5 mya and 3.2 mya, some have suggested this specimen is an *Australopithecus*, perhaps even *Au. afarensis* (with a brain size which is difficult to determine, yet appears small), while still others have placed this specimen in *Homo* (small dentition, and flat-orthognathic face). While taxonomic placing of this species is quite divided, the discoverers have argued that this species is ancestral to *Homo*, in particular to *Homo rudolfensis*. Some have placed the species itself into the genus *Homo*, although the cranial capacity and general cranial features are not as derived. Some researchers have additionally associated the earliest tool finds from Lomekwi, Kenya, temporally (3.3 mya) and in close geographic proximity to this species/specimen.

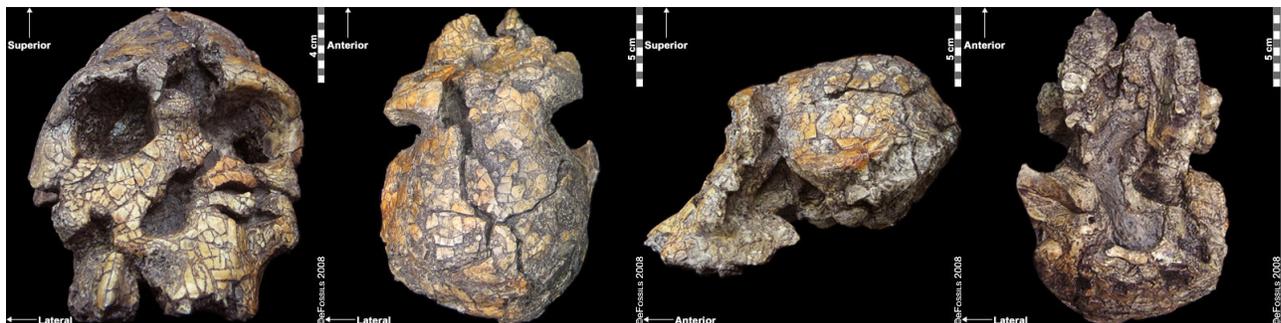


Figure 9.14 This specimen (*Kenyanthropus platyops*) has small dentition, a small brain case and a relatively flat face. Its genus/species designation remains contested.

South African Australopithecines

Since the discovery of the Taung Child there have been numerous Australopithecine discoveries from the region known as “**The Cradle of Humankind**,” recently given UNESCO World Heritage Site status as “The Fossil Hominid Sites of South Africa.” The limestone caves found in the Cradle allow for the excellent preservation of fossils. Animals navigating the landscape in the past falling into cave openings, or caves used as dens by carnivores, led to the accumulation of deposits over millions of years. Many of the hominin fossils we have, encased in **breccia** (hard calcareous sedimentary rock), are recently exposed from limestone quarries mined in the previous century. This means that extracting fossils requires excellent and detailed exposed work, often by a team of skilled technicians.

While these sites have historically been difficult to date, with mixed assemblages accumulated over large time periods, advances in techniques such as Uranium-series dating have allowed for greater accuracy. Historically, the excellent faunal record from East Africa has traditionally been used to compare sites based on **relative dating**. In this, the knowledge of environmental/faunal changes and extinction events allows us to know which hominin finds are relatively younger or older than others.

Research Highlight: The Taung Child:

The well-known fossil of a juvenile *Australopithecine*, the “Taung Child,” was the first early hominin evidence ever discovered and was the first to demonstrate our common human heritage in Africa (Figure 9.15). The tiny facial skeleton and natural endocrast were discovered in 1924 by a local quarryman in the North West Province in South Africa and painstakingly removed from the surrounding cement-like breccia by Raymond Dart using his wife’s knitting needles. When first shared with the scientific community in 1925, it was discounted as being nothing more than a young monkey of some kind. Prevailing biases of the time made it too difficult to contemplate that this small-brained hominin could have anything to do with our own history. The fact that it was discovered in Africa simply served to strengthen this bias.

It was not until adult specimens of the species began to be discovered at Sterkfontein Caves that scientific and public opinion began to be swayed. The most complete of these remains is best known as “Mrs. Ples” and was discovered in 1947.

The Taung Child is also an excellent example of how understanding the morphology and physiology of teeth can reveal information about our evolution. While many introductory lectures will showcase how the foramen magnum is more anteriorly placed (indicating a head that sits centrally on the body, ideal for bipedalism), Taung Child also shows very humanlike dentition. For one, the canine teeth were relatively small compared to other apes. And there was little to no diastema (or gap) between the canines and incisors. The incisors themselves were vertical and close together, and not at an angle to the mouth (procumbent), as seen in other apes. Furthermore, the overall shape of the dental arcade was more rounded, or parabolic. Even though these features were related to deciduous teeth, they were overwhelmingly more humanlike than those seen in other apes.

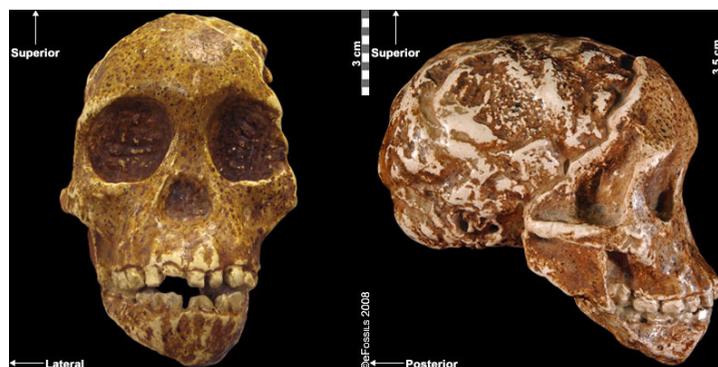


Figure 9.15 The Taung Child has a nearly complete face, mandible, and partial endocranial cast.

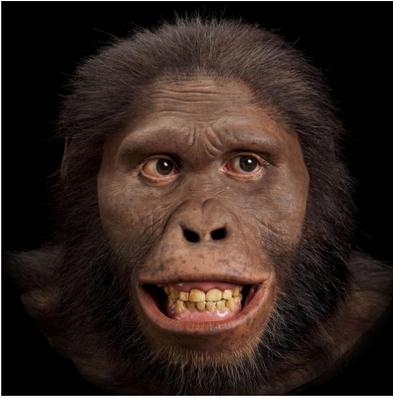


Figure 9.17 An artistic reconstruction of *Australopithecus africanus* by John Gurche.

The discovery of the Taung Child in 1924 shifted the focus of palaeoanthropological research from Europe to Africa (although acceptance was slow). The species with which it is assigned, ***Australopithecus africanus*** (name meaning “Southern Ape of Africa”), is currently dated to between 3.3 mya and 2.1 mya, with discoveries from Sterkfontein, Taung, Makapansgat, and Gladysvale in South Africa (Figure 9.17). A relatively large brain (400 cc to 500 cc), small canines without an associated diastema, and more rounded cranium and smaller teeth than *Au. afarensis* indicate some derived traits. Similarly, the postcranial remains (in particular, the pelvis) indicate bipedalism. However, the sloping face and curved phalanges (indicative of retained arboreal locomotor abilities) show some primitive features. Although not in direct association with stone tools, a 2015 study noted that the trabecular bone morphology of the hand was consistent with forceful tool manufacture and use, suggesting potential early tool abilities.

Another famous *Au. africanus* skull, the skull of “Mrs. Ples,” previously attributed to *Plesianthropus transvaalensis* (meaning “near human from the Transvaal,” the old name for Gauteng Province, South Africa), was promptly shortened by contemporary journalists to “Ples”(Figure 9.16). Due to the prevailing mores of the time, the assumed female found herself married, at least in name, and has become widely known as “Mrs. Ples.” It was later reassigned to *Au. africanus* and is now argued to be a young male rather than an adult female cranium.

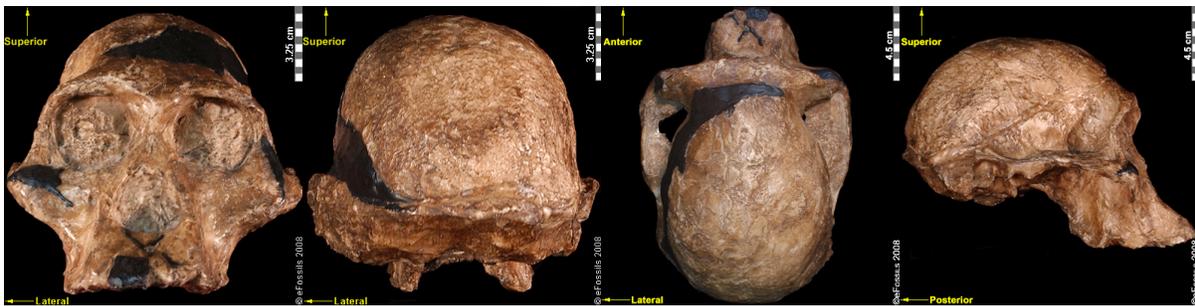


Figure 9.16 The “Mrs. Ples” brain case is small in size (like apes), but has a less prognathic face and its foramen magnum is positioned more like modern humans than in African apes.

In 2008, nine-year-old Matthew Berger, son of paleoanthropologist Lee Berger, noted a clavicle bone in some leftover mining breccia in the Malapa Fossil Site (South Africa). After rigorous studies, the species, ***Australopithecus sediba*** (meaning “fountain” or “wellspring” in the South African language of Sesotho), was named in 2010 (Figure 9.18). The first type specimen belongs to a juvenile male, Karabo (MH1), but the species is known from at least six partial skeletons of infants through adult. These specimens are currently dated to 1.97 mya. The discoverers have argued that *Au. sediba* shows **mosaic** features between *Au. africanus* and *Homo*, which potentially indicates a transitional species, although this is heavily debated. A small brain size (*Australopithecus*-like; 420 cc to 450 cc), but gracile mandible and small teeth (*Homo*-like). Similarly, the postcranial skeletons are also said to have mosaic features: scientists have interpreted this mixture of traits (such as a robust ankle, but evidence for an arch in the foot) as a transitional phase between a body previously adapted to arborealism (tree climbing, particularly in evidence from the bones of the wrist) to one that adapted to bipedal ground walking. Some researchers have argued

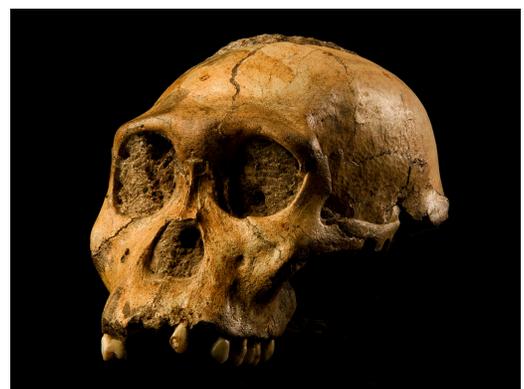


Figure 9.18 *Australopithecus sediba* shows mosaic features between *Au. africanus* and *Homo*.

that *Au. sediba* shows a modern hand morphology (shorter fingers and a longer thumb), indicating that adaptations to tool manufacture and use may be present in this species.

Another famous Australopithecine find from South Africa is that of the nearly complete skeleton now known as “Little Foot.” Little Foot (StW 573) is potentially the earliest dated South African hominin fossil (3.7 mya, based on radiostopic techniques, although some argue that it is younger than 3 mya). The name is jokingly in contrast to the crypto-species, “bigfoot,” and named because the initial discovery of four ankle bones indicated bipedality. Little Foot was discovered by Ron Clarke in 1994, when he came across the ankle bones while sorting through monkey fossils in the University of Witwatersrand collections. He asked Stephen Motsumi and Nkwane Molefe to identify the known records of the fossils, which allowed them to find the rest of the specimen within just days of searching the Sterkfontein Caves’ Silberberg Grotto. The discoverers of Little Foot insist that other fossil finds, previously identified as *Au. Africanus*, be placed in this new species based on shared primitive traits with older East African Australopithecines. These include features such as a relatively large brain size (408 cc), robust zygomatic arch, and a flatter midface. Furthermore, the discoverers have argued that the heavy anterior dental wear patterns, relatively large anterior dentition, and smaller hind dentition of this specimen more closely resemble that of *Au. anamensis* or *Au. afarensis*.

It has thus been placed in the species ***Australopithecus prometheus***. This species name refers to a previously defunct taxon named by Raymond Dart. The name “prometheus” refers to the Greek titan who stole fire from the gods to give to humanity. Raymond Dart believed that the bones he discovered at Makapansgat in South Africa reflected burning when it was later determined that they were in fact *Au. africanus* remains that had been stained by manganese during fossilization. The species designation was, through analyzing Little Foot, revived by Ron Clarke, who insists that many other fossil hominin specimens have prematurely been placed into *Au. africanus*. Others say that it is more likely that *Au. africanus* is a more variable species and not representative of two distinct species.

Paranthropus “Robust” Australopithecines

In the robust australopithecines, the specialized nature of the teeth and masticatory system, such as flaring zygomatic arches (cheekbone) to accommodate the large temporalis (chewing) muscle, indicated a shift in diet in these taxa. Some argued that the diet of the robust australopithecines was so specific that any change in environment would have accelerated their extinction. The generalist nature of the teeth of the gracile australopithecines, and certainly early *Homo*, would have made these hominins more resistant to environmental change. However, some have suggested that these features might have developed more in response to effectively eating **fall-back foods** in hard times, and might not indicate a lack of adaptability.

Paranthropus is usually referred to by scholars as the “robust” australopithecine, because of its defining distinct morphological features. Features that are closer to those of the assumed ancestral type are referred to as *P. aethiopicus*, and those that have become derived are referred to as both *P. boisei* and *P. robustus*. These features include a large, broad, dish-shaped face, zygomatic arches that are forward facing, including a large mandible with extremely large posterior dentition. These three species have been grouped together by a majority of scholars as a genus as they share more derived features (are more closely related to each other, or in other words, are **monophyletic**) than the other australopithecines. Much of the phylogenetic debate (and research, more generally) has revolved around the shared adaptations of these “robust” australopithecines linked to a diet of hard and/or tough foods, including their:

- large posterior dentition (they are referred to as **megadonts**),
- hyper-thick enamel,
- thick robust jaws, and
- flared zygomatic arches.

In terms of diet, the tougher, chewing diets of the robust australopithecines are supported by the extreme morphology of their face and cranium. Similarly, the **prognathic** faces, which jut out under the eyes in the earlier hominins, are less pronounced than in those of living Great Apes, yet are more pronounced than in early *Homo*. In contrast, the **orthognathic** (flat) face of our genus is well suited to accommodate our relatively small generalized teeth and processed diets.

Researchers have reached a consensus to the umbrella term *Paranthropus*. However there are those who remain cautious/disagree. As a collective, this genus spans 2.7 mya to 1.0 mya, although the dates of the individual species differ. The genus was first discovered in Kromdraai B, South Africa, by Robert Broom, who first attributed the **holotype** of specimen TM 1517.

The earliest of the *Paranthropus* species, ***Paranthropus aethiopicus***, is dated to between 2.7 mya and 2.3 mya, and is currently found in Tanzania, Kenya, and Ethiopia in the EARS system (Figure 9.19). It is well known because of the “Black Skull” (KNM-WT 17000), so called because of the mineral manganese which stained it black during fossilization. As with all robust Australopithecines, *P. aethiopicus* has the shared derived traits of large flat premolars and molars, large flaring zygomatic arches for accommodating large chewing muscles (the temporalis muscle), a sagittal crest for increased muscle attachment of the chewing muscles to the skull, and a robust mandible and supraorbital torus (brow ridge). However, only a few teeth have been found. A proximal tibia indicates bipedality and similar body size to *Au. afarensis*. In recent years, researchers have discovered and assigned a proximal tibia and juvenile cranium (L.338y-6) to the species.



Figure 9.19 The “Black Skull” (*Paranthropus aethiopicus*) had a large sagittal crest and large flared zygomatic arches which indicate it had large chewing muscles and a powerful biting force.



Figure 9.20 Artistic reconstruction of a *Paranthropus boisei*, male, by John Gurche.

First attributed as *Zinjanthropus boisei* (with the first discovery going by the nickname “Zinj” or sometimes “Nutcracker Man”), ***Paranthropus boisei*** was discovered in 1959 by Mary Leakey (Figure 9.20). This “robust” australopithecine species is distributed across countries in East Africa at sites such as Kenya (Koobi Fora, West Turkana, and Chesowanja), Malawi (Malema-Chiwondo), Tanzania (Olduvai Gorge and Peninj), and Ethiopia (Omo River Basin and Konso). The **hypodigm** has been found by researchers to date to roughly 2.4 mya to 1.4 mya. Due to the nature of its exaggerated larger and more robust features, *P. boisei* has been termed **hyper-robust**, that is, even more heavily built than other robust forms, with very large, flat posterior dentition (largest of all hominins currently known). Richard Leakey and Bernard Wood have both suggested that *P. boisei* could have made and used stone tools. Tools dated to 2.5 mya in Ethiopia have been argued to possibly belong to this species. Despite the cranial features of *P. boisei* indicating a tough diet of tubers, nuts, and seeds, isotopes indicate a diet high in C4 foods (e.g., grasses, such as sedges). This differs from what is seen in *P. robustus*. Another famous specimen from this species is the Peninj mandible from Tanzania, found in 1964 by Kimoya Kimeu.



Figure 9.21 “Nutcracker Man” (*Paranthropus boisei*) had hyper-robust features including very large dentition, flaring zygomatic arches, a broad concave face. It had a powerful and extremely efficient chewing force.

Paranthropus robustus was the first taxon to be discovered within the genus in Kromdraai B, by a school boy named Gert Terblanche, and subsequent fossil discoveries were made by researcher Robert Broom in 1938 (Figure 9.22). *Paranthropus robustus* dates approximately to 2.0 mya to 1 mya and is the only taxon from the genus to be discovered in South Africa. It has been found in sites all over the Cradle, such as Kromdraai B, Swartkrans, Gondolin, Drimolen, and Coopers Cave. Several of these fossils are fragmentary in nature, distorted, and not well preserved, because they have been recovered from quarry breccia using explosives. *P. robustus* features are neither as “hyper-robust” as *P. boisei* nor as primitive as *P. aethiopicus*, but have been described as having less derived, more general features that are shared with both East African species (e.g., the sagittal crest and zygomatic flaring). Enamel hypoplasia is also common in this species, possibly because of instability in the development of large, thick-enameled dentition.



Figure 9.22 *Paranthropus robustus* had less derived, more general features; it was not as robust as *P. boisei* and not as primitive as *P. aethiopicus*. It also appeared to be prone to a tooth defect (enamel hypoplasia) making the enamel thinner and more prone to decay.

Comparisons Between Gracile and Robust Australopiths

Comparisons between gracile and robust australopithecines may indicate different phylogenetic groupings but may also indicate parallel evolution in several species. In general, the robust australopithecines have large temporalis (chewing) muscles, as indicated by flaring zygomatic arches, sagittal crests, and robust mandibles (jawbones). Their hind dentition is large (megadont), with low cusps and thick enamel. Within the gracile australopithecines, researchers have debated the relatedness of the species, or even whether these species should be lumped together to represent more variable or polytypic species. Often researchers will attempt to draw chronospecific trajectories, with one taxon evolving into another over time.

EARLY TOOL USE AND TECHNOLOGY

Early Stone Age Technology (ESA)

The Early Stone Age (ESA) marks the beginning of recognizable technology as made by our human ancestors. Stone-tool (or **lithic**) technology is defined by the fracturing of rocks and the manufacture of tools through a process called **knapping**. The Stone Age lasted for more than 3 million years and is broken up into chronological periods called the **Early (ESA)**, **Middle (MSA)**, and **Later Stone Ages (LSA)**. Each period is further broken up in different techno-complexes, as explained below. This section focuses on the earliest tools associated with ESA. The ESA spanned the largest technological time period of human innovation from over 3 million years ago to around 300,000 years ago and is associated almost entirely with hominin species prior to modern *Homo sapiens*. As the ESA advanced, stone tool makers (known as **knappers**) began to change the ways they detached **flakes** and eventually were able to shape artifacts into functional tools. These advances in technology go together with the developments in human evolution and cognition, dispersal of populations across the African continent and the world, and climatic changes.

In order to understand the ESA, it is important to consider some definitions. A **techno-complex** is a term encompassing multiple **assemblages** (collections of artifacts) that share similar traits in terms of artifact production and morphology. Not all assemblages are exactly the same as within each techno-complex: one can have multiple phases and traditions at different sites. However, there is an overarching commonality between them. Within stone tool assemblages, both flakes or **cores** (the rocks from which flakes are removed) are used as tools. **Large Cutting Tools (LCTs)** are tools that are shaped to have functional edges. It is important to note that the information presented here is a small fraction of what is known about the ESA and there are many ongoing debates and discoveries within the archaeological discipline.

Currently, the oldest known stone tools, which form the techno-complex the Lomekwian, date to 3.3 mya. They were found at a site called Lomekwi 3 in Kenya. This techno-complex is the most recently defined and pushed back the oldest known date for lithic technology. There is only one known site thus far and, due to the age of the site, it is associated with species prior to *Homo*, such as *Kenyanthropus platyops*. Flakes were produced through indirect percussion, whereby the knappers held a rock and hit it against another rock resting on the ground. The pieces are very chunky and do not display the same fracture patterns as seen in later techno-complexes. Lomekwian knappers likely aimed to get a sharp-edged piece on a flake which would have been functional, although the specific function is currently unknown.

Stone tool use, however, is not only understood through the direct discovery of the tools. Cut marks on fossilized animal bones may illuminate the functionality of stone tools. In one controversial study in 2010, researchers argued that cut marks on a pair of animal bones from Dikika (Ethiopia), dated to 3.4 mya, were from stone tools. The discoverers suggested that they be more securely associated, temporally, with *Au. afarensis*. However, others have noted that these marks are consistent with teeth marks from crocodiles and other carnivores.

The Oldowan techno-complex is far more established in the scientific literature. It is called the **Oldowan** because it was originally discovered in Olduvai Gorge, Tanzania, but the oldest assemblage is from Gona in Ethiopia, dated to 2.6 mya. The techno-complex is defined as a core and flake industry. Like the Lomekwian, there was an aim to get sharp-edged flakes, but through a different production method. Knappers were able to actively hold or manipulate the core being knapped which they could directly hit using a hammerstone. This technique is known as free-hand percussion which demonstrates an understanding of fracture mechanics. It has long been argued that the Oldowan hominins were skillful in tool manufacture.

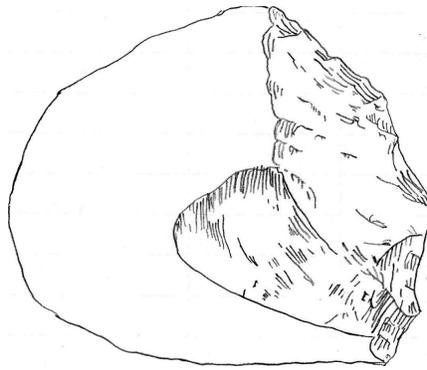


Figure 9.23 Some scholars believe that some *genuses* explored in this chapter were capable of producing more complex stone tools (Oldowan).

Because Oldowan knapping requires skill, earlier researchers have attributed these tools to members of our genus, *Homo*. However, some have argued that these tools are in more direct association with hominins in the *genuses* described in this chapter (Figure 9.23).

Invisible Tool Manufacture and Use

The vast majority of our understanding of these early hominins comes from fossils and reconstructed paleoenvironments. It is only from 3 mya when we can start “looking into their minds” and lifestyles by analyzing their manufacture and use of stone tools. However, the vast majority of tool use in primates (and, one can argue, in humans) is not with durable materials like stone. All of our extant Great Ape relatives have been observed using sticks, or leaves, or other materials for some secondary purpose (to wade across rivers, to “fish” for termites, or to absorb water for drinking). It is possible that the majority of early hominin tool use and manufacture may be invisible to us because of this preservation bias.

HOMININ CHARTS

Hominin	<i>Sahelanthropus tchadensis</i>
Dates	7 mya to 6 mya
Region(s)	Chad
Famous discoveries	The initial discovery, made in 2001
Brain size	360 cc average
Dentition	Smaller than in extant Great Apes, larger and pointier than in humans. Canines worn at the tips
Cranial features	A short cranial base and a foramen magnum (hole in which the spinal cord enters the cranium) that is more humanlike in positioning, has been argued to indicate upright walking.
Postcranial features	Currently little published postcranial material.
Culture	NA
Other	The extent to which this hominin was bipedal is currently heavily debated. If so, it would indicate an arboreal bipedal ancestor of hominins, not a knuckle-walker like chimpanzees.

Figure 9.24

Hominin	<i>Orrorin tugenensis</i>
Dates	6 mya to 5.7 mya
Region(s)	Tugen Hills (Kenya)
Famous discoveries	Original discovery in 2000
Brain size	NA
Dentition	Smaller cheek teeth (molars and premolars) than even more recent hominins (i.e., derived), thick enamel, and reduced, but apelike, canines
Cranial features	Not many found
Postcranial features	Fragmentary leg, arm, and finger bones have been found. Indicates bipedal locomotion.
Culture	Potential toolmaking capability based on hand morphology, but nothing found directly
Other	This is the earliest species which clearly indicates adaptations for bipedal locomotion.

Figure 9.25

Hominin	<i>Ardipithecus ramidus</i>
Dates	4.4 mya
Region(s)	Middle Awash region and Gona (Ethiopia)
Famous discoveries	A partial female skeleton nicknamed "Ardi" (ARA-VP-6/500).
Brain size	300 cc to 350 cc
Dentition	Little differences between the canines of males and females (small sexual dimorphism).
Cranial features	Midfacial projection, slightly prognathic. Cheekbones less flared and robust than in later hominins.
Postcranial features	Ardi demonstrates a mosaic of ancestral and derived characteristics in the postcrania. For instance, an opposable big toe similar to chimpanzees (i.e., "primitive" or more ancestral), which could have aided in climbing trees effectively. However, the pelvis and hip show that she could walk upright (i.e., it is derived), supporting her hominin status.
Culture	None directly associated
Other	Over 110 specimens from Aramis

Figure 9.26

Hominin	<i>Ardipithecus kadabba</i>
Dates	5.2 mya to 5.8 mya
Region(s)	Middle Awash (Ethiopia)
Famous discoveries	This species discovery in 1997 by Yohannes Haile-Selassie
Brain size	NA
Dentition	Larger hind dentition than in modern chimpanzees. Thick enamel and larger canines than in later hominins.
Cranial features	NA
Postcranial features	A large hallux (big toe) bone indicates a bipedal “push off.”
Culture	NA
Other	Faunal evidence indicates a mixed grassland/woodland environment.

Figure 9.27

Hominin	<i>Australopithecus anamensis</i>
Dates	4.2 mya to 3.8 mya
Region(s)	Turkana region (Kenya), Middle Awash (Ethiopia)
Famous discoveries	A 2019 find from Ethiopia, named MRD
Brain size	370 cc
Dentition	Relatively large canines compared with more recent Australopithecines
Cranial features	Projecting cheekbones and primitive earholes.
Postcranial features	Lower limb bones (tibia and femur) indicate bipedality; arboreal features in upper limb bones (humerus) found.
Culture	NA
Other	Almost 100 specimens, representing over 20 individuals, have been found to date.

Figure 9.28

Hominin	<i>Australopithecus afarensis</i>
Dates	2.9 mya to 3.9 mya
Region(s)	Afar Region, Omo, Maka, Fejej, and Belohdelie (Ethiopia); Laetoli (Tanzania); Koobi Fora (Kenya).
Famous discoveries	Lucy, Selam (Dikika Child), Laetoli Footprints
Brain size	380 cc to 430 cc
Dentition	Reduced canines and molars relative to Great Apes, but larger than in modern humans
Cranial features	Prognathic face, facial features indicate relatively strong chewing musculature (compared with <i>Homo</i>), but less extreme than in <i>Paranthropus</i> .
Postcranial features	Clear evidence for bipedalism from lower limb postcranial bones. Laetoli Footprints indicate humanlike walking. Dikika Child bones indicate retained primitive arboreal traits in the postcrania.
Culture	None directly; but close in age and proximity to controversial cut marks at Dikika and early tools in Lomekwi
Other	<i>Au. afarensis</i> is one of the oldest and most well-known australopithecine species and consists of a large number of fossil remains.

Figure 9.29

Hominin	<i>Australopithecus bahrelghazali</i>
Dates	3.6 mya
Region(s)	Chad
Famous discoveries	“Abel,” the holotype
Brain size	NA
Dentition	NA
Cranial features	NA
Postcranial features	NA
Culture	NA
Other	Arguably within range of variation of <i>Au. afarensis</i>

Figure 9.30

Hominin	<i>Australopithecus deyiremada</i>
Dates	3.5 mya to 3.3 mya
Region(s)	Woranso-Mille (Afar region, Ethiopia)
Famous discoveries	First fossil mandible bones were discovered in 2011 in the Afar region of Ethiopia by Yohannes Haile-Selassie
Brain size	NA
Dentition	Smaller teeth with thicker enamel than seen in <i>Au. afarensis</i> , with a potentially harder diet
Cranial features	Larger mandible and more projecting cheekbones than in <i>Au. afarensis</i>
Postcranial features	NA
Culture	NA
Other	Contested species designation; arguably a member of <i>Au. afarensis</i>

Figure 9.31

Hominin	<i>Australopithecus garhi</i>
Dates	2.5 mya
Region(s)	Middle Awash (Ethiopia)
Famous discoveries	NA
Brain size	450 cc
Dentition	Larger hind dentition than seen in other gracile Australopithecines
Cranial features	NA
Postcranial features	A femur of a fragmentary partial skeleton, argued to belong to <i>Au. garhi</i> , indicates this species may be longer-limbed than <i>Au. afarensis</i> , although still able to move arboreally.
Culture	Crude/primitive stone tools resembling Oldowan (described later) have been found in association with <i>Au. garhi</i>
Other	This species is not well documented or understood, and is based on only several fossil specimens.

Figure 9.32

Hominin	<i>Australopithecus africanus</i>
Dates	3.3 mya to 2.1 mya
Region(s)	Sterkfontein, Taung, Makapansgat, Gladysvale (South Africa)
Famous discoveries	Taung Child, "Mrs. Ples," Little Foot (?)
Brain size	400 cc to 500 cc
Dentition	Smaller teeth (derived) relative to <i>Au. afarensis</i> . Small canines with no diastema.
Cranial features	A rounder skull compared with <i>Au. afarensis</i> in East Africa. A sloping face (primitive).
Postcranial features	Similar postcranial evidence for bipedal locomotion (derived pelvis) with retained arboreal locomotion (e.g., curved phalanges—fingers), as seen in <i>Au. afarensis</i>
Culture	None with direct evidence
Other	A 2015 study noted that the trabecular bone morphology of the hand was consistent with forceful tool manufacture and use, suggesting potential early tool abilities.

Figure 9.33

Hominin	<i>Australopithecus sediba</i>
Dates	1.97 mya
Region(s)	Malapa Fossil Site (South Africa)
Famous discoveries	Karabo (MH1)
Brain size	420 cc to 450 cc
Dentition	Small dentition with Australopithecine cusp-spacing
Cranial features	Small brain size (<i>Australopithecus</i> -like), but gracile mandible (<i>Homo</i> -like)
Postcranial features	Scientists have interpreted this mixture of traits (such as a robust ankle, but evidence for an arch in the foot) as a transitional phase between a body previously adapted to arborealism (tree climbing, particularly in evidence from the bones of the wrist) to one that adapted to bipedal ground walking.
Culture	None of direct association, but some have argued that a modern hand morphology (shorter fingers and a longer thumb) means that adaptations to tool manufacture and use may be present in this species.
Other	It was first discovered through a clavicle bone in 2008 by nine-year-old Matthew Berger, son of paleoanthropologist Lee Berger.

Figure 9.34

Hominin	<i>Australopithecus prometheus</i>
Dates	3.7 mya (debated)
Region(s)	Sterkfontein (South Africa)
Famous discoveries	“Little Foot” (StW 573)
Brain size	408 cc (Little Foot estimate)
Dentition	Heavy anterior dental wear patterns, relatively large anterior dentition and smaller hind dentition, similar to <i>Au. afarensis</i>
Cranial features	Relatively larger brain size, robust zygomatic arch, and a flatter midface.
Postcranial features	The initial discovery of four ankle bones indicated bipedality.
Culture	NA
Other	Highly debated new species designation

Figure 9.35

Hominin	<i>Paranthropus aethiopicus</i>
Dates	2.7 mya to 2.3 mya
Region(s)	West Turkana (Kenya), Laetoli (Tanzania), Omo River Basin (Ethiopia)
Famous discoveries	The 'Black Skull' (KNM-WT 17000)
Brain Size	410 cc
Dentition	<i>P. aethiopicus</i> has the shared derived traits of large flat premolars and molars, although few teeth have been found.
Cranial features	Large flaring zygomatic arches for accommodating large chewing muscles (the temporalis muscle), a sagittal crest for increased muscle attachment of the chewing muscles to the skull, and a robust mandible and supraorbital torus (brow ridge).
Postcranial features	A proximal tibia indicates bipedality, and similar size to <i>Au. afarensis</i> .
Culture	None
Other	The "Black Skull" is so called because of the mineral manganese which stained it black during fossilization.

Figure 9.36

Hominin	<i>Paranthropus boisei</i>
Dates	2.4 mya to 1.4 mya
Region(s)	Koobi Fora, West Turkana, and Chesowanja (Kenya), Malema-Chiwondo (Malawi), Olduvai Gorge and Peninj (Tanzania), and Omo River basin and Konso (Ethiopia)
Famous discoveries	“Zinj”; or sometimes “Nutcracker Man” (OH5), in 1959 by Mary Leakey. The Peninj mandible from Tanzania, found in 1964 by Kimoya Kimeu
Brain size	500 cc to 550 cc
Dentition	Very large, flat posterior dentition (largest of all hominins currently known). Much smaller anterior dentition. Very thick dental enamel.
Cranial features	Indications of very large chewing muscles (e.g., flaring zygomatic arches and a large sagittal crest).
Postcranial features	Evidence for high variability and sexual dimorphism, with estimates of males at 1.37 meters tall and females at 1.24 meters.
Culture	Richard Leakey and Bernard Wood have both suggested that <i>P. boisei</i> could have made and used stone tools. Tools dated to 2.5 mya in Ethiopia have been argued to possibly belong to this species.
Other	Despite the cranial features of <i>P. boisei</i> indicating a tough diet of tubers, nuts and seeds, isotopes indicate a diet high in C4 foods (e.g., grasses, such as sedges). This differs from what is seen in <i>P. robustus</i> .

Figure 9.37

Hominin	<i>Paranthropus robustus</i>
Dates	2.3 mya to 1 mya
Region(s)	Kromdraai B, Swartkrans, Gondolin, Drimolen, and Coopers Cave (South Africa)
Famous discoveries	SK48 (original skull)
Brain size	410 cc to 530 cc
Dentition	Large posterior teeth with thick enamel, consistent with other Robust Australopithecines. Enamel hypoplasia is also common in this species, possibly because of instability in the development of large, thick enameled dentition.
Cranial features	<i>P. robustus</i> features are neither as “hyper-robust” as <i>P. boisei</i> or as primitive as <i>P. aethiopicus</i> , but have been described as less derived more general features that are shared with both East African species, e.g., the sagittal crest and zygomatic flaring.
Postcranial features	Reconstructions indicate sexual dimorphism.
Culture	None
Other	Several of these fossils are fragmentary in nature, distorted and not well preserved, because they have been recovered from quarry breccia using explosives.

Figure 9.38

Hominin	<i>Kenyanthropus platyops</i>
Dates	3.5 mya to 3.2 mya
Region(s)	Lake Turkana (Kenya)
Famous discoveries	KNM-WT 40000
Brain size	Difficult to determine, but appears within the range of <i>Australopithecus afarensis</i>
Dentition	Small molars/dentition (<i>Homo</i> -like characteristic)
Cranial features	Flatter (i.e., orthognathic) face
Postcranial features	NA
Culture	Some have associated the earliest tool finds from Lomekwi, Kenya, temporally (3.3 mya) and in close geographic proximity to this species/specimen.
Other	Taxonomic placing of this species is quite divided. The discoverers have argued that this species is ancestral to <i>Homo</i> , in particular to <i>Homo rudolfensis</i> .

Figure 9.39

Review Questions

- What is the difference between a “derived” versus a “primitive” trait? Give an example of both, seen in *Au. afarensis*.
- Which of the paleoenvironment hypotheses have been used to describe early hominin diversity, and which have been used to describe bipedalism?
- Which anatomical features for bipedalism do we see in early hominins? Are these primarily obligate bipeds? Explain.
- Describe the dentition of gracile and robust australopithecines. What might these tell us about their relative diets?
- List the hominin species argued to be associated with stone tool technologies. Are you convinced of these associations? Why/why not?

Key Terms

Arboreal: Related to trees or woodland.

Aridification: Becoming increasingly arid or dry, as related to the climate or environment.

Assemblage: A collection demonstrating a pattern. Often pertaining to a site or region.

Bipedalism: The locomotor ability to walk on two legs.

Breccia: Sedimentary rock.

Canines: The pointy teeth just next to the incisors, in the front of the mouth.

Chronospecies: Species which are said to evolve into another species, in a linear fashion, over time.

Clade: A group of species or taxa with a shared common ancestor.

Cladistics: The field of grouping organisms into those of shared ancestry.

Closed habitat: A phrase typically referring to a woodland, or tree-filled, environment.

Context: As pertaining to palaeoanthropology: the place where an artifact or fossil is found.

Core: The remains of a rock which has been flaked or knapped.

Cusp: The “bumps” on the teeth.

Dental formula: A technique to describe the number of incisors, canines, premolars and molars in each quadrant of the mouth.

Derived: A newly evolved trait, or differing from that seen in the ancestor.

Diastema: A gap. In this chapter referring to a canine diastema, between the incisors and canines.

Early Stone Age (ESA): The earliest described archaeological period, whereby we start seeing stone tool technology.

East African Rift System (EARS): Often referring to the Rift Valley, expanding from Malawi to Ethiopia. This active geological structure is responsible for much of the visibility of the paleoanthropological record in East Africa.

Enamel: The highly mineralized outer layer of the tooth.

Encephalization: Expansion of the brain.

Extant: Currently living, i.e., not extinct.

Fallback foods: Food which may not be preferred by an animal (e.g., not nutritionally dense) but is essential for survival in times of stress.

Fauna: The animals of a particular region, habitat, or geological period.

Faunal turnover: The rate at which species go extinct and are replaced with new species.

Flake: The piece knocked off of a stone core during the manufacture of a tool and may be used as a stone tool.

Flora: The plants of a particular region, habitat, or geological period.

Folivorous: Foliage-eating.

Fossil: The remains or impression of an organism from the past.

Frugivorous: Fruit-eating.

Generalist: A species that can thrive in a wide variety of habitats and can have a varied diet.

Glacial: Colder, drier periods during an ice age when there is more ice trapped at the poles.

Gracile: Slender, less rugged or pronounced features.

Hallux: The big toe.

Holotype: A single specimen from which a species or taxon is described or named.

Honing P3: The mandibular premolar alongside the canine (in primates, the P3), which is angled to give space for (and sharpen) the upper canines.

Hominid: A primate which includes humans and our fossil relatives and, in more recent definitions, other Great Apes.

Hominin: A primate which includes humans and our fossil relatives since our divergence from extant Great Apes.

Incisors: The teeth in the front of the mouth, used to bite off.

Incisiform: An adjective referring to a canine which appears more incisor-like in morphology.

Interglacial: A period of milder climate in between two glacials.

Isotopes: Two or more forms of the same element that contain equal numbers of protons but different numbers of neutrons, giving them the same chemical properties but different atomic masses.

Knapping (knappers): The fracturing of rocks for the manufacture of tools (and the people who made them).

Large Cutting Tool (LCT): A tool that is shaped to have functional edges.

Last Common Ancestor (LCA): The hypothetical final ancestor (or ancestral population) of two or more taxa before their divergence.

Lithic: Relating to stone (here to stone tools).

Lumbar lordosis: The inward curving of the lower (lumbar) parts of the spine. The lower curve in the human S-shaped spine.

Lumpers: Researchers who prefer to lump variable specimens into a single species or taxon, and who feel high levels of variation is biologically real.

Megadont: An organism with extremely large dentition compared with body size.

Molars: The largest, most posterior of the hind dentition.

Monophyletic: A taxon or group of taxa descended from a common ancestor which is not shared with another taxon or group.

Morphology (morphological): The study of the form or size and shape of things (in this case, skeletal parts).

Mosaic evolution: The concept that evolutionary change does not occur homogeneously throughout the body in organisms.

Occlude: When the teeth from the maxilla come into contact with the teeth in the mandible.

Oldowan: Lower Palaeolithic, the earliest stone tool culture.

Orthognathic: The face below the eyes is relatively flat and does not jut out anteriorly.

Paleoanthropology (paleoanthropologist): The study of human evolution (a researcher in this field).

Paleoenvironment: An environment from a period in the Earth's geological past.

Paleolithic: The earliest part of the Stone Age.

Parabolic: Like a parabola (parabola-shaped).

Phylogeny: The study of the evolutionary relationships between groups of organisms.

Phylogenetics: The study of phylogeny.

Pliocene: A geological epoch between the Miocene and Pleistocene.

Polytypic: In reference to taxonomy, having two or more group variants.

Postcranium: The skeleton below the cranium (head).

Premolars: The smallest of the hind teeth, behind the canines.

Procumbent: In reference to incisors, tilting forward.

Prognathic: In reference to the face below the eyes jutting anteriorly.

Quaternary: The most recent geological time period, which includes the Pleistocene and Holocene Epochs, and is defined by the cyclicity of increasing and decreasing ice sheets at the poles.

Relative dating: Dating techniques which are in reference to a sequence (i.e., older or younger than others in the reference). No actual/absolute date is estimated.

Robust: Rugged or exaggerated features.

Site: A place in which evidence of past societies/species/activities may be observed through archaeological or palaeontological practice.

Specialist: A specialist species can thrive only in a narrow range of environmental conditions or has a limited diet.

Splitters: Researchers who prefer to split a highly variable taxon into multiple groups or species.

Taxa (taxonomy): Plural of taxon, a taxonomic group such as species, genus or family. Taxonomy is the science of grouping and classifying organisms.

Techno-complex: A term encompassing multiple assemblages that share similar traits in terms of artifact production and morphology.

Thermoregulation: Maintaining body temperature through physiologically cooling or warming the body.

Ungulates: Hoofed mammals e.g., cows and kudu.

Volcanic tuffs: Rock made from volcanic ash from volcanic eruptions in the past.

Valgus knee: The angle of the knee between the femur and tibia, which allows for weight distribution to be angled closer to the point above the center of gravity (i.e., between the feet) in bipeds.

About the Authors

Kerryn Warren, Ph.D.

University of Cape Town, kerryn.warren@gmail.com



Kerryn Warren

Kerryn Warren is a postdoctoral researcher at the University of Cape Town. She lectures on archaeology and human evolution. Her research interests include identifying hybridization in the hominin fossil record, stemming from research from her Ph.D., and understanding evolution education in South African schools. She is also currently one of the new “Underground Astronauts” selected to excavate *Homo naledi* remains from the Rising Star Cave System in the Cradle of Humankind. She is passionate about education and science communication.

Lindsay Hunter, Ph.D.

University of Witwatersrand



Lindsay Hunter

Lindsay Hunter is a trained paleoanthropologist who uses her more than 15 years of experience to make sense of the distant past of our species in ways that can help us to build a better future. She received her master’s degree in biological anthropology from the University of Iowa and is completing her Ph.D. in archaeology at the University of the Witwatersrand. She has studied fossil and human bone collections across five continents with major grant support from the National Science Foundation (United States) and the Wenner-Gren Foundation for Anthropological Research. More recently she developed and led the National Geographic “Umsuka” Public Palaeoanthropology Project in South Africa with support from the National Geographic Society and private donors. She now works as the Community Relations and Development Director for the Center for Academic Research and Training in Anthropogeny (CARTA) at UCSD.

Navashni Naidoo, M.Sc.

University of Cape Town



Navashni Naidoo

Navashni Naidoo is a researcher at Nelson Mandela University, lecturing on physical geology. Her research interests include developing paleoenvironmental proxies suited to the African continent, behavioral ecology, and engaging with community-driven archaeological projects. She has excavated at Stone Age sites across South Africa and East Africa.

Silindokuhle Mavuso, M.Sc.

University of Witwatersrand



Silindokuhle Mavuso

Silindokuhle has always been curious about the world around him and how it has been shaped. He is a Ph.D. Candidate at the University of Witwatersrand (Wits) conducting palaeoenvironmental reconstruction and change of the northeastern Turkana Basin's Pleistocene sequence. Silindokuhle begun his education with a B.Sc. (geology, archaeology, and environmental and geographical sciences) from the University of Cape Town before moving to Wits for a B.Sc. Honors (geology and palaeontology) and M.Sc. in geology. During this time he has gained more training as a Koobi Fora Fieldschool fellow (Kenya) as well as an Erasmus Mundus scholar (France). Silindokuhle is a Plio-Pleistocene geologist with a specific focus on identifying and explaining past environments that are associated with early human life and

development through time. He is interested in a wide range of disciplines such as micromorphology, sedimentology, geochemistry, geochronology, and stratigraphy. He has worked with teams from significant eastern and southern African hominid sites including Elandsfontein, Rising Star, Sterkfontein, Gondolin, Laetoli, Olduvai, and Koobi Fora. He plans to extend his knowledge from both parts of the continent to assist the better understanding of how we as humans came to being.

Kimberleigh Tommy, M.Sc.

University of Witwatersrand



Kimberleigh Tommy

Kimberleigh Tommy is currently a Ph.D. candidate in biological anthropology at the Human Variation and Identification Research Unit of the School of Anatomical Sciences at the University of Witwatersrand. Her current research focuses on the evolution and biomechanical implications of bipedal walking through analyses of trabecular bone structure in the joints of the lower limb. Kimberleigh was awarded her Master of Science (M.Sc.) degree with distinction (and no corrections) from the University of the Witwatersrand, specializing in palaeoanthropology and functional morphology in 2018. Her research interests include trabecular structure, functional morphology, primate locomotion, ontogenetic development of gait, biomechanics, and joint pathologies.

Rosa Moll, M.Sc.

University of Witwatersrand



Rosa Moll

Rosa Moll is an archaeology Ph.D. candidate at the University of Witwatersrand. She focuses on Earlier Stone Age core reduction strategies of east Africa and south Africa and received her M.Sc. with distinction in the same field. She is interested in how stool tool technological behaviors correspond with cognitive human evolution. In 2018 she was awarded the Baldwin Fellowship from the Leakey Foundation as part of her Ph.D.

Nomawethu Hlazo, M.Sc.

University of Cape Town



Nomawethu Hlazo

Nomawethu Hlazo is a student at the University of Cape Town currently undergoing her Doctoral Degree. She completed her undergraduate degree in biochemistry and archaeology. Since then her postgraduate studies have focused on the genus *Paranthropus* and the variation that exists between and within species. Following the fossil species, she has concentrated on the study of geometric morphometrics and will follow new techniques such as paleoproteomics to investigate not only shape change but contributions of evolutionary processes and ecological niches occupied by the genus *Paranthropus*. Since the start of her research with *Paranthropus* she has worked at several sites, not only in the Cradle of Humankind World Heritage Site in South Africa but also in Kenya. Her research has shown that this genus is highly diverse and more variable than we expected. After completion of her master's (with distinction), she has been able to show the contributions of both natural selection and genetic drift and their roles in shaping *Paranthropus* craniomandibular variation.

For Further Exploration

The Smithsonian website hosts descriptions of fossil species, an interactive timeline and much more! It is a highly recommended website. <http://humanorigins.si.edu/evidence>

The Maropeng Museum website hosts a wealth of information regarding South African Fossil Bearing sites in the Cradle of Humankind. <https://www.maropeng.co.za/content/page/human-evolution>

This quick comparison between *Homo naledi* and *Australopithecus sediba* from the Perot Museum: <https://perot-museum.imgix.net/2019-08-naledi-sediba-quick-comparison.pdf>

This explanation of the braided stream by the Perot Museum: <https://www.dropbox.com/s/11d2hv42psj21y9/Braided%20Stream-1920.mp4?dl=0>

A collation of 3-D files for visualizing (or even 3-D printing) for homes, schools, and universities: <https://www.hetmp.com/>

PBS learning materials, including videos and diagrams of the Laetoli footprints, bipedalism, and fossils: https://www.pbslearningmedia.org/resource/tdc02.sci.life.evo.lp_humanevo/human-evolution/

A wealth of information from the Australian Museum website, including species descriptions, family trees, and explanations of bipedalism and diet: <https://australianmuseum.net.au/learn/science/human-evolution/>

References

- Alemseged, Z., Spoor, F., Kimbel, W. H., Bobe, R., Geraads, D., Reed, D., and Wynn, J. G. 2006. A juvenile early hominin skeleton from Dikika, Ethiopia. *Nature* 443 (7109): 296–301.
- Asfaw, B.; White, T.; Lovejoy, O.; Latimer, B.; Simpson, S.; Suwa, G. 1999. “*Australopithecus garhi*: a new species of early hominid from Ethiopia.” *Science* 284 (5414): 629–635.
- Behrensmeyer, A. K., Todd, N. E., Potts, R., and McBrinn, G. E. 1997. Late Pliocene faunal turnover in the Turkana Basin, Kenya, and Ethiopia. *Science* 278: 637–640.
- Berger, L. R., De Ruiter, D. J., Churchill, S. E., Schmid, P., Carlson, K. J., Dirks, P. H., and Kibii, J. M. 2010. *Australopithecus sediba*: a new species of *Homo*-like australopith from South Africa. *Science* 328 (5975): 195–204.
- Beyene, Y., Zeleke, Y., and Uzawa, K. 1997. The Acheulean at Konso-Gardula: Results from locality KGA4-A2. In Fukui, K., Kurimoto, E., and Shigeta, M. (Eds.), *Ethiopia in Broader Perspective Vol 1*. Shokado: Kyoto, Japan: 376–381.
- Beyene, Y., Katoh, S., Wolde, Gabriel, G., Kart, W.K., Sudo, M., Kondo, M., Hyodo, M., Renne, P.R., Suwa, G., and Asfaw, B. 2013. The characteristics and chronology of the earliest Acheulean at Konso, Ethiopia. *Proceedings of the National Academy of Sciences* 110 (5): 1585–1591.
- Bobe, R. and Behrensmeyer, A.K. 2004. The expansion of grassland ecosystems in Africa in relation to mammalian evolution and the origin of the genus *Homo*. *Palaeogeography, Palaeoclimatology, Palaeoecology* 207 (3–4): 399–420.
- Brain, C.K. 1967. The Transvaal Museum’s Fossil Project at Swartkrans. *South African Journal of Science* 63 (9): 378–384.
- Broom, R. 1938a. More discoveries of *Australopithecus*. *Nature* 141 (1): 828–829.
- Broom, R. 1938b. The Pleistocene anthropoid apes of South Africa. *Nature* 142 (3591): 377–379.
- Broom, R. 1947. Discovery of a new skull of the South African ape-man, Plesianthropus. *Nature* 159 (4046): 672.
- Broom, R. 1950. The genera and species of the South African fossil ape-man. *American Journal of Physical Anthropology* 8 (1): 1–14.
- Brunet, M., Beauvilain, A., Coppens, Y., Heintz, E., Moutaye, A. H., and Pilbeam, D. 1995. The first australopithecine 2,500 kilometers west of the Rift Valley (Chad). *Nature* 378 (6554): 275–273.
- Cerling, T.E., Wynn, J.G., Andanje, S.A., Bird, M.I., Korir, D.K., Levin, N.E., Mace, W., Macharia, A.N., Quade, J., Remien, C.H. 2011. Woody cover and hominin environments in the past 6 million years. *Nature* 476: 51e56.
- Clarke, R. J. 1998. First ever discovery of a well-preserved skull and associated skeleton of *Australopithecus*. *South African Journal of Science* 94 (10): 460–463.
- Clarke, Ronald. 2013. “*Australopithecus* from Sterkfontein Caves, South Africa.” In: *The paleobiology of Australopithecus*. Springer 105–123.
- Clarke, R.J. and Kuman, K. 2019. The skull of StW 573, a 3.67 Ma *Australopithecus prometheus* skeleton from Sterkfontein Caves, South Africa. *Journal of Human Evolution* 134: 102634.

- Clarke, R. J., and Tobias, P. V. 1995. Sterkfontein Member 2 foot bones of the oldest South African hominid. *Science* 269 (5223): 521–524.
- Constantino, P.J., and Wood, B.A. 2004. Paranthropus paleobiology In: *Miscelanea en Homenae a Emiliano Aguirre*. Volumen III: Paleoantropologia. Alcala de Henares: Museo Arqueologico Regional.
- Constantino, P., and Wood, B. 2007. The evolution of Zinjanthropus boisei. *Evolutionary Anthropology. Issues, News, and Reviews* 16 (2): 49–62.
- Dart, R. A. 1925. Australopithecus africanus, the man-ape of South Africa. *Nature* 115: 195–199.
- Darwin, Charles, 1809–1882. *On the Origin of the Species by Means of Natural Selection, or Preservation of Favoured Races in the Struggle for Life*. London: John Murray, 1859.
- Darwin, Charles. 1871. *The Descent of Man: And Selection in Relation to Sex*. London: J. Murray.
- De Heinzelin, J., Clark, J.D., White, T., Hart, W., Renne, P., WoldeGabriel, G., Beyene, Y., and Vrba, E. 1999. Environment and behavior of 2.5-million-year-old Bouri hominids. *Science* 284 (5414): 625–629.
- De la Torre, I. 2016. The origins of the Acheulean: past and present perspectives on a major transition in human evolution. *Philosophical Transactions Royal Society B* 371: 20150245. DOI: 10.1098/rstb.2015.0245.
- De la Torre, I., and Mora, R. 2009. Remarks on the Current Theoretical and Methodological Approaches to the Study of Early Technological Strategies in Eastern Africa. In Hovers, E., and Braun, D.R. (Eds.), *Interdisciplinary Approaches to the Oldowan*. Dordrecht: Springer: 15–24.
- DeMenocal, Peter, B.D., 2004. African climate change and faunal evolution during the Pliocene–Pleistocene. *Earth and Planetary Science Letters* 220 (1–2): 3–24.
- deMenocal, P.B.D., and Bloemendal, J. 1995. Plio-Pleistocene climatic variability in subtropical Africa and the paleoenvironment of hominid evolution: a combined data-model approach. *Paleoclimate and evolution, with emphasis on human origins* 262–288.
- Dirks, P. H., Kibii, J. M., Kuhn, B. F., Steininger, C., Churchill, S. E., Kramers, J. D., ... and King, G. C. 2010. Geological setting and age of Australopithecus sediba from southern Africa. *Science* 328 (5,975): 205–208.
- Faith, J.T., and Behrensmeyer, A.K. 2013. Climate change and faunal turnover: testing the mechanics of the turnover-pulse hypothesis with South African fossil data. *Paleobiology* 39 (4): 609–627.
- Haile-Selassie, Y., Gibert, L., Melillo, S. M., Ryan, T. M., Alene, M., Deino, A., Scott, G., and Saylor, B. Z. 2015. New species from Ethiopia further expands Middle Pliocene hominin diversity. *Nature* 521 (7,553): 432–433.
- Haile-Selassie, Y., Melillo, S.M., Vazzana, A., Benazzi, S. and Ryan, T.M. 2019. A 3.8-million-year-old hominin cranium from Woranso-Mille, Ethiopia. *Nature*, pp.1–6.
- Hlazo, N. 2015. Paranthropus: Variation in Cranial Morphology. Honours Thesis: University of Cape Town, Cape Town.
- Hlazo, N. 2018. Variation and the Evolutionary drivers of diversity in the genus Paranthropus. Master's Thesis: University of Cape Town, Cape Town.
- Grine, F.E. 1988. New Craniodental fossils of Paranthropus from the Swartkrans Formation and their significance in “robust” australopithecine evolution. In Grine, F.E. (eds.). *Evolutionary History of the “Robust” Australopithecines*. New York: Aldine de Gruyter.

- Hay, R. L., and Leakey, M. D. 1982. The fossil footprints of Laetoli. *Scientific American* 246 (2): 50–57.
- Hay, R.L. 1990. Olduvai Gorge: a case history in the interpretation of hominid paleoenvironments. In: L. Laporte (eds.). *East Africa. Establishment of a geologic framework for paleoanthropology*. Boulder: Geological Society of America Special Paper, 242 (2): 23–37.
- Harmand, S., Lewis, J.E., Feibel, C.S., Lepre, C.J., Prat, S., Lenoble, A., Boë, X., Quinn, R.L., Brenet, M., Arroyo, A., Taylor, N., Clément, S., Daver, G., Brugal, J.P., Leakey, L., Mortlocks, R.A., Wright, J.D., Lokorodi, S., Kirwa, C., Kent, D.V., and Roche, H. 2015. 3.3 million-year-old stone tools from Lomekwi3, West Turkana, Kenya. *Nature* 521: 310–316.
- Isaac G.L. 1969. Studies of early culture in East Africa. *World Archaeology* 1: 1–28.
- Johanson, D.C., White, T.D., and Coppens, Y. 1978. A new species of the genus *Australopithecus* (Primates: Hominidae) from the Pliocene of East Africa. *Kirtlandia* 28: 1–14.
- Kingston, J.D. and Harrison, T. 2007. Isotopic dietary reconstructions of Pliocene herbivores at Laetoli: Implications for early hominin paleoecology. *Palaeogeography, Palaeoclimatology, Palaeoecology* 243 (3–4): 272–306.
- Kingston, J.D. 2007. Shifting adaptive landscapes: progress and challenges in reconstructing early hominid environments. *American Journal of Physical Anthropology* 134 (S45): 20–58.
- Kimbel, W.H. 2006. The Species and Diversity of Australopiths. *Springer* 1(1): 1–35.
- Kimbel, W. H., and Deleuzene, L. K. 2009. “Lucy” redux: A review of research on *Australopithecus afarensis*. *American Journal of Physical Anthropology* 140 (S49): 2–48.
- Kuman, K. 2007. The Earlier Stone Age in South Africa: site context and the influence of cave studies. In: Pickering, T.R., Schick, K., and Toth, N. (eds) *Breathing Life Into Fossils: Taphonomic Studies in Honor of C.K. (Bob) Brain*: 181–198. Bloomington (Indiana): Stone Age Institute Press.
- Kuman, K., and Clarke, R.J. 2000. Stratigraphy, Artefact Industries and Hominid associations, Member 5. *Journal of Human Evolution* 38: 827–847.
- Leakey, L.S.B. 1959. A New Fossil Skull from Olduvai. *Nature* 184 (4685): 491–493.
- Leakey, M. 1971. *Olduvai Gorge, Vol. 3*. Cambridge: Cambridge University Press.
- Leakey, M. D., and Hay, R. L. 1979. Pliocene footprints in the Laetoli Beds at Laetoli, northern Tanzania. *Nature* 278 (5,702): 317–323.
- Leakey, M. G., Feibel, C. S., McDougall, I., and Walker, A. 1995. New four-million-year-old hominid species from Kanapoi and Allia Bay, Kenya. *Nature* 376 (6,541): 565–571.
- Leakey, M.G., Spoor, F., Brown, F.H., Gathogo, P.N., Kiarie, C., Leakey, L.N., and McDougall, I. 2001. New hominin genus from eastern Africa shows diverse middle Pliocene lineages. *Nature* 410 (6,827): 433.
- Lebatard, A. E., Bourlès, D. L., Durringer, P., Jolivet, M., Braucher, R., Carcaillet, J., ... and Likies, A. 2008. Cosmogenic nuclide dating of *Sahelanthropus tchadensis* and *Australopithecus bahrelghazali*: Mio-Pliocene hominids from Chad. *Proceedings of the National Academy of Sciences* 105 (9): 3226–3231.
- Lee-Thorp, J. 2011. The demise of “Nutcracker Man”. *Proceedings of the National Academy of Sciences* 108 (23): 9319–9320.
- Lepre, C.J., Roche, H., Vent, D.V., Harmand, S., Quinn, R. L., Brugal, J-P., Texier, P-J., Lenoble, A. and Feibel, C. S. 2011. An earlier origin for the Acheulian. *Nature* 477: 82–85.

- Lombard, M., Wadley, L., Deacon, J., Wurs, S., Parsons, I., Mohapi, M., Swart, J. and Mitchell, P. 2012. South African and Lesotho Stone Age Sequence Updated. *The South African Archaeological Bulletin* 67 (195): 123–144.
- Maslin, M.A., Brierley, C.M., Milner, A.M., Shultz, S., Trauth, M.H., and Wilson, K.E. 2014. East African climate pulses and early human evolution. *Quaternary Science Reviews* 101: 1–17.
- McHenry, H.M. 2009. “Human Evolution.” In Michael Ruse; Joseph Travis. *Evolution: The First Four Billion Years*. pp. 256–280.
- Patterson, B.; Howells, W. W. 1967. “Hominid Humeral Fragment from Early Pleistocene of Northwestern Kenya.” *Science* 156 (3771): 64–66.
- Pickering, R. and Kramers, J.D. 2010. Re-appraisal of the stratigraphy and determination of new U-Pb dates for the Sterkfontein hominin site. South Africa. *Journal of Human Evolution* 59 (1): 70–86.
- Potts, R. 1998. Environmental hypotheses of hominin evolution. *American Journal of Physical Anthropology* 107 (S27): 93–136.
- Potts, R. 2013. Hominin evolution in settings of strong environmental variability. *Quaternary Science Reviews* 73: 1–13.
- Rak, Y. 1983. *The australopithecine face*. New York: Academic Press.
- Rak, Y. 1988. On Variation in the Masticatory System of *Australopithecus boisei*. In Grine, F.E. (eds.). *Evolutionary History of the “Robust” Australopithecines*. New York: Aldine de Gruyter.
- Rayner, R.J., Moon, B.P., Masters, J.C. 1993. The Makapansgat australopithecine environments. *Journal of Human Evolution* 24: 219–231.
- Semaw, S. 2000. The World’s Oldest Stone Artefacts from Gona, Ethiopia: Their Implications for Understanding Stone Technology and Patterns of Human Evolution Between 2.6 Million Years Ago and 1.5 Million Years Ago. *Journal of Archaeological Science* 27: 1197–1214.
- Shipman, Pat. 2002. *The Man Who Found the Missing Link: Eugene Dubois and His Lifelong Quest to Prove Darwin Right*. Phoenix.
- Spoor, F. 2015. Palaeoanthropology: The middle Pliocene gets crowded. *Nature* 521(7553): 432– 433.
- Strait, D.S., Grine, F.E. and Moniz, M.A. 1997. A reappraisal of early hominid phylogeny. *Journal of Human Evolution* 32 (1): 17–82.
- Suwa, G., Asfaw, B., Kono, R.T., Kubo, D., Lovejoy, C.O. and White, T.D., 2009. The *Ardipithecus ramidus* skull and its implications for hominid origins. *Science* 326 (5949): 68–68e7.
- Thackeray, J. F. 2000. “Mrs. Ples” from Sterkfontein: Small male or large female? *The South African Archaeological Bulletin* 155–158.
- Thackeray, F., Braga, J., Treil, J., Niksch, N., and Labuschagne, J.H. 2002. “Mrs. Ples” (Sts 5) from Sterkfontein: an adolescent male? *South African Journal of Science* 98 (1–2): 21–22.
- Toth, N. 1985. The Oldowan reassessed. *Journal of Archaeological Science* 12: 101–120.
- Vrba, E. S. 1988. Late Pliocene climatic events and hominid evolution. Pp. 405–426 In Grine, F. E., ed. *The evolutionary history of the robust Australopithecines*. Aldine, New York.

- Vrba, E. S. 1998. Multiphasic growth models and the evolution of prolonged growth exemplified by human brain evolution. *Journal of Theoretical Biology* 190: 227–239.
- Vrba, E. S. 2000. Major features of Neogene mammalian evolution in Africa. Pp. 277–304 In Partridge, T. C. and Maud, R., eds. *Cenozoic geology of southern Africa*. Oxford University Press, Oxford.
- Walker, A.C., Leakey, R.E., Harris, J.M., and Brown, F.H. 1986. 2.5- my Australopithecus boisei from west of Lake Turkana, Kenya. *Nature* 322 (6079): 517–522.
- Walker, A.C. and Leakey, R.E. 1988. The Evolution of Australopithecus boisei. In Grine, F.E. (eds.). *Evolutionary History of the “Robust” Australopithecines*. New York: Aldine de Gruyter.
- Ward, C., Leakey, M., and Walker, A. 1999. The new hominid species *Australopithecus anamensis*. *Evolutionary Anthropology: Issues, News, and Reviews* 7 (6): 197–205.
- White, T.D. 1988. The Comparative Biology of “Robust” Australopithecus: Clues from Content. In Grine, F.E. (eds.). *Evolutionary History of the “Robust” Australopithecines*. New York: Aldine de Gruyter.
- White, T.D., Suwa, G., Asfaw, B., 1994. *Australopithecus ramidus*, a new species of early hominid from Aramis, Ethiopia. *Nature* 371: 306–312.
- Wood, B.A. 2010. Reconstructing human evolution: Achievements, challenges, and opportunities. *Proceedings of the National Academy of Sciences* 10 (2): 8902–8909.
- Wood, B. A. and Boyle, E.K. 2016. Hominin Taxic Diversity: Fact or Fantasy? *Yearbook of Physical Anthropology* 159 (S61): 37–78.
- Wood, B. and Schroer, K. 2017. Paranthropus: Where Do Things Stand? In Marom, A. and Hovers, E. (eds.). *Human Paleontology and Prehistory*. New York: Springer, Cham.

Acknowledgments

All of the authors in this section are students and early career researchers in paleoanthropology and related fields in South Africa (or at least have worked in South Africa). We wish to thank everyone who supports young and diverse talent in this field, and would love to further acknowledge black, African, and female academics who have helped pave the way for us.

Figure Attributions

Figure 9.1 [IMG_1696 Great Rift Valley](#) by [Ninara](#) is used under a CC BY 2.0 License.

Figure 9.2 Clades original to Explorations: An Open Invitation to Biological Anthropology by Katie Nelson is under a [CC BY-NC 4.0 License](#).

Figure 9.3 [CO₂ increase since the Industrial Revolution](#) by [NASA](#), original from Luthi, D., et al.. 2008; Etheridge, D.M., et al. 2010; Vostok ice core data/J.R. Petit et al.; NOAA Mauna Loa CO₂ record is in the [public domain](#) and used within [NASA guidelines on re-use](#).

Figure 9.4 [African savannah @ Masai Mara \(21308330314\)](#) by [Leo Li](#) from Hong Kong is used under a [CC BY 2.0 License](#).

Figure 9.5 [Skeleton of human \(1\) and gorilla \(2\), unnaturally sketched](#) by unknown from Brehms Tierleben, Small Edition 1927 is in the [public domain](#).

Figure 9.6 Skeletal comparisons between modern humans and non-obligate bipeds original to Explorations: An Open Invitation to Biological Anthropology is under a [CC BY-NC 4.0 License](#).

Figure 9.7a [Sahelanthropus tchadensis: TM 266-01-060-1 anterior view](#) by [eFossils](#) is [copyrighted and used for noncommercial purposes as outlined by eFossils](#).

Figure 9.7b [Sahelanthropus tchadensis: TM 266-01-060-1 posterior view](#) by [eFossils](#) is [copyrighted and used for noncommercial purposes as outlined by eFossils](#).

Figure 9.7c [Sahelanthropus tchadensis: TM 266-01-060-1 inferior view](#) by [eFossils](#) is [copyrighted and used for noncommercial purposes as outlined by eFossils](#).

Figure 9.7d [Sahelanthropus tchadensis: TM 266-01-060-1 lateral left view](#) by [eFossils](#) is [copyrighted and used for noncommercial purposes as outlined by eFossils](#).

Figure 9.8a [Ardipithecus ramidus Skull](#) by [@BoneClones](#) is used by permission and available here under a [CC BY-NC 4.0 License](#).

Figure 9.8b [Artist's rendition of "Ardi" skeleton](#) by [@BoneClones](#) is used by permission and available here under a [CC BY-NC 4.0 License](#).

Figure 9.9 [Adult human teeth](#) by [Genusfotografen](#) (Tomas Gunnarsson) through Wikimedia Sverige [Wikimedia Sverige](#) is used under a [CC BY-SA 4.0 License](#).

Figure 9.10 [Paranthropus bonsei compared to Homo sapiens](#) by Constantino, Paul J. is used under a [CC BY-SA 4.0 License](#).

Figure 9.11 [Australopithecus anamensis: KNM-KP 29281 occlusal view](#) by [eFossils](#) is [copyrighted and used for noncommercial purposes as outlined by eFossils](#).

Figure 9.12a [Australopithecus afarensis, "Lucy," adult female. Reconstruction based on AL-288-1 by artist John Gurche, front view close-up](#) by [Smithsonian](#) [exhibit: Reconstructed Faces, What does it mean to be human?] is [copyrighted and used for educational and noncommercial purposes as outlined by the Smithsonian](#).

Figure 9.12b [Australopithecus afarensis, adult male. Reconstruction based on AL444-2 by John Gurche](#) by [Smithsonian](#) [exhibit: Reconstructed Faces, What does it mean to be human?] is [copyrighted and used for educational and noncommercial purposes as outlined by the Smithsonian](#).

Figure 9.13 [Lucy blackbg](#) (AL 288-1, Australopithecus afarensis, cast from Museum national d'histoire naturelle, Paris) by 120 is used under a [CC BY-SA 3.0 License](#).

Figure 9.14a [Kenyanthropus platyops KNM WT 40000 anterior view](#) by [eFossils](#) is [copyrighted and used for noncommercial purposes as outlined by eFossils](#).

Figure 9.14b [Kenyanthropus platyops KNM WT 40000 superior view](#) by [eFossils](#) is [copyrighted and used for noncommercial purposes as outlined by eFossils](#).

Figure 9.14c [Kenyanthropus platyops KNM WT 40000 lateral left view](#) by [eFossils](#) is [copyrighted and used for noncommercial purposes as outlined by eFossils](#).

Figure 9.14d [Kenyanthropus platyops KNM WT 40000 inferior view](#) by [eFossils](#) is [copyrighted and used for noncommercial purposes as outlined by eFossils](#).

Figure 9.15a [australopithecus africanus: Taung 1 anterior view](#) by [eFossils](#) is [copyrighted and used for noncommercial purposes as outlined by eFossils](#).

Figure 9.15b [australopithecus africanus: Taung 1 lateral right view](#) by [eFossils](#) is [copyrighted and used for noncommercial purposes as outlined by eFossils](#).

Figure 9.16a [Australopithecus africanus Sts 5 anterior view](#) by [eFossils](#) is [copyrighted and used for noncommercial purposes as outlined by eFossils](#).

Figure 9.16b [Australopithecus africanus Sts 5 posterior view](#) by [eFossils](#) is [copyrighted and used for noncommercial purposes as outlined by eFossils](#).

Figure 9.16c [Australopithecus africanus Sts 5 superior view](#) by [eFossils](#) is [copyrighted and used for noncommercial purposes as outlined by eFossils](#).

Figure 9.16d [Australopithecus africanus Sts 5 lateral right view](#) by [eFossils](#) is [copyrighted and used for noncommercial purposes as outlined by eFossils](#).

Figure 9.17 [Australopithecus africanus. Reconstruction based on STS 5 by John Gurche by Smithsonian](#) [exhibit: Reconstructed Faces, What does it mean to be human?] is [copyrighted and used for educational and noncommercial purposes as outlined by the Smithsonian](#).

Figure 9.18 [Australopithecus sediba](#), photo by Brett Eloff, courtesy [Profberger](#) and [Wits University](#) is used under a [CC BY-SA 4.0 License](#).

Figure 9.19a [Paranthropus aethiopicus: KNM-WT 17000 anterior view](#) by [eFossils](#) is [copyrighted and used for noncommercial purposes as outlined by eFossils](#).

Figure 9.19b [Paranthropus aethiopicus: KNM-WT 17000 lateral right view](#) by [eFossils](#) is [copyrighted and used for noncommercial purposes as outlined by eFossils](#).

Figure 9.19c [Paranthropus aethiopicus: KNM-WT 17000 superior view](#) by [eFossils](#) is [copyrighted and used for noncommercial purposes as outlined by eFossils](#).

Figure 9.19d [Paranthropus aethiopicus: KNM-WT 17000 posterior view](#) by [eFossils](#) is [copyrighted and used for noncommercial purposes as outlined by eFossils](#).

Figure 9.19e [Paranthropus aethiopicus: KNM-WT 17000 inferior view](#) by [eFossils](#) is [copyrighted and used for noncommercial purposes as outlined by eFossils](#).

Figure 9.20 [Paranthropus boisei, male. Reconstruction based on OH 5 and KNM-ER 406 by John Gurche by Smithsonian](#) [exhibit: Reconstructed Faces, What does it mean to be human?] is [copyrighted and used for educational and noncommercial purposes as outlined by the Smithsonian](#).

Figure 9.21a [Paranthropus boisei: OH 5 anterior view](#) by [eFossils](#) is [copyrighted and used for noncommercial purposes as outlined by eFossils](#).

Figure 9.21b [Paranthropus boisei: OH 5 inferior view](#) by [eFossils](#) is [copyrighted and used for noncommercial purposes as outlined by eFossils](#).

Figure 9.21c [Paranthropus boisei: OH 5 posterior view](#) by [eFossils](#) is [copyrighted and used for noncommercial purposes as outlined by eFossils](#).

Figure 9.22a [Paranthropus robustus: SK 48 anterior view](#) by [eFossils](#) is [copyrighted and used for noncommercial purposes as outlined by eFossils](#).

Figure 9.22b [Paranthropus robustus: SK 48 superior view](#) by [eFossils](#) is [copyrighted and used for noncommercial purposes as outlined by eFossils](#).

Figure 9.22c [Paranthropus robustus: SK 48 inferior view](#) by [eFossils](#) is [copyrighted and used for noncommercial purposes as outlined by eFossils](#).

Figure 9.22d [Paranthropus robustus: SK 48 lateral left view](#) by [eFossils](#) is [copyrighted and used for noncommercial purposes as outlined by eFossils](#).

Figure 9.23 [Olduwan Industry Chopper 2](#) by [Emmyanne29](#) is used under a [CC0 1.0 License](#).