

1  
2  
3  
4 DOES THE OCCURRENCE AND  
5 DURATION OF HEALTH INSULTS  
6 AMONG SHIWIAR  
7 FORAGER-HORTICULTURALISTS  
8 INDICATE THAT HEALTH CARE  
9 PROVISIONING REDUCES JUVENILE  
10 MORTALITY?  
11  
12  
13  
14  
15  
16  
17  
18

19 Lawrence S. Sugiyama  
20  
21

22  
23 **ABSTRACT**  
24

25 *Humans lifespan is characterized by delayed maturation. Delayed maturation*  
26 *may arise when juvenile mortality is reduced. Recent research suggests that*  
27 *juvenile mortality reduction could be achieved via provisioning to weaned*  
28 *juveniles, particularly during health crises. Here I test this idea with data*  
29 *on the causes, distribution, and duration of injuries and illnesses suffered by*  
30 *Shiwiar forager-horticulturalists during the juvenile period. Health insults*  
31 *for which prolonged care is necessary for survival are a recurrent feature of*  
32 *the juvenile lifespan. About half the individuals for whom data on disability*  
33 *duration were gathered suffered health insults likely to be lethal without*  
34 *extended aid; over 80% were born after a parent suffered such an event; and*  
35

36  
37 **Socioeconomic Aspects of Human Behavioral Ecology**  
38 **Research in Economic Anthropology, Volume 23, 377–400**  
39 **Copyright © 2004 by Elsevier Ltd.**  
40 **All rights of reproduction in any form reserved**  
**ISSN: 0190-1281/doi:10.1016/S0190-1281(04)23016-5**

1     *over 90% were born after a direct ancestor in the two ascending generations*  
2     *suffered such an event. The data indicate that health-care provisioning*  
3     *reduces juvenile mortality, and that provisioning of sick and injured juveniles*  
4     *has important fitness consequences in this population.*  
5  
6

## 7                                       **INTRODUCTION**

8  
9     Evolutionary life history theory examines how natural selection produces age-  
10    related resource allocation between somatic (growth and maintenance) and  
11    reproductive (mating and parental) effort (Charnov & Schaffer, 1973; Hill &  
12    Hurtado, 1996; Kaplan et al., 2000; MacArthur & Wilson, 1967; Schaffer, 1974;  
13    Williams, 1966). Most species delay maturity and hence reproduction for a length  
14    of time during which growth and development takes place. The evolution of  
15    delayed maturity is a central question for life history theory because, all else equal,  
16    delayed maturity decreases the probability that juveniles will live to reproductive  
17    age (Charnov, 1991; Hill & Hurtado, 1996; Kaplan et al., 2000; Pagel & Harvey,  
18    1993).

19    Primates in general have delayed reproduction (Pagel & Harvey, 1993; Pereira,  
20    1993), but even controlling for body size, humans have later reproduction and  
21    longer lifespans than other primates do, including our closest living relatives  
22    the common chimpanzee *Pan troglodyte* (e.g. Hill et al., 2001; Kaplan et al.,  
23    2000). For instance, in natural fertility foraging societies human females begin  
24    reproduction at about 17–20 years of age (Hawkes et al., 1998; Hill & Hurtado,  
25    1996; Kaplan et al., 2000), whereas chimpanzee females begin reproducing at  
26    13–15 years (Boesch & Boesch, 2000; Nishida et al., 1990; Pusey, 1990). While  
27    42% of precontact Aché foragers reached 50 years of age, only about 9% of wild  
28    chimpanzees live to be 50 years old (Hill et al., 2001). This difference is not just  
29    the outcome of higher chimpanzee mortality. Chimpanzees senesce more rapidly  
30    than do humans, indicating that the difference in lifespan is the result of selection  
31    acting differentially on the life history of each species (Hill et al., 2001).

32    Human life history also includes subsistence and other support of females and  
33    their offspring by post-reproductive and other females (e.g. Hawkes et al., 1998,  
34    2000; Hrdy, 1999, n.d.), while males contribute resources to mates, other adults, and  
35    juveniles (Hawkes et al., 2001; Hewlett, 1992; Hill & Hurtado, 1996; Kaplan et al.,  
36    2000; Marlowe, 1999, 2001; Winterhalder, 1996). Humans also exhibit exceptional  
37    intelligence, complex social skills, and a large capacity for developmental learning  
38    (e.g. Bogin, 1999; Byrne, 1997; Flinn et al., manuscript; Geary & Flinn, 2001;  
39    Hill & Kaplan, 1999; Kaplan et al., 2000; Tooby & DeVore, 1987). Kaplan et al.  
40    (2000) and others have argued that human adult foraging competence requires a

1 long period of skill and knowledge acquisition (e.g. Bock, 2002; Hill & Kaplan,  
2 1999; Walker et al., 2001). Although this claim is currently debated and the degree  
3 to which juvenile foragers contribute to their own subsistence varies (e.g. Bird &  
4 Bliege Bird, 2002; Bliege Bird & Bird, 2002; Blurton Jones & Marlowe, 2002;  
5 Blurton Jones et al., 1994; Sugiyama & Chacon, n.d.; Tucker et al., n.d.), humans  
6 are nevertheless distinguished by a long period of juvenile dependence (e.g. Bliege  
7 Bird & Bird, 2002; Bogin, 1999; Hawkes et al., 2001; Hewlett, 1992; Hill & Kaplan,  
8 1999; Kaplan et al., 2000).

9 Three basic inter-related factors are used to explain delayed maturity and  
10 long lifespan: demographic factors, skill and/or knowledge acquisition, and  
11 invariant patterns of growth (Pagel & Harvey, 1993). The demographic perspective  
12 emphasizes that reproductive age and lifespan are largely the function of extrinsic  
13 mortality rates. Species with high juvenile mortality tend to mature faster and be  
14 shorter-lived, because they cannot afford the high risk of dying before reproducing  
15 (e.g. Horn, 1978; Rose, 1983; Williams, 1957). Species with lower juvenile  
16 mortality can afford to mature later.

17 In addition, all else equal, delayed maturity can evolve if the pre-reproductive  
18 mortality risk it entails is offset by fitness-enhancing benefits acquired during the  
19 juvenile period. Longer adult lifespan allows more time for these payoffs to be  
20 realized. Thus, a second factor in explanations of long juvenile lifespan is that it  
21 is a time during which juveniles acquire “embodied capital,” not only via growth,  
22 but also via knowledge or skills – social, parental, fighting, or foraging skills  
23 – that enhance later fertility and/or reduce mortality (e.g. Harvey & Zammuto,  
24 1983; Kaplan et al., 2000; Pereira & Altmann, 1985; Promislow & Harvey, 1990;  
25 Sutherland et al., 1986). Rates of adult mortality are therefore expected to covary  
26 with fertility and age at maturity such that investments in the juvenile period  
27 – defined here as the period between weaning and first reproduction (Pagel &  
28 Harvey, 1993) – are compensated by higher lifetime fitness. Low adult mortality  
29 allows a longer lifespan, and consequently an increased age at maturity, decreased  
30 fecundity, and higher parental investment. Conversely, high adult mortality is  
31 expected to be compensated for by increased fecundity and/or rapid maturation  
32 (Pereira, 1993; Sutherland et al., 1986).

33 The final factor in explanations of long lifespan and delayed maturity focuses on  
34 invariant patterns of growth. The larger an animal’s adult body size, the greater its  
35 ability to produce energy, but the later the animal will begin reproduction because  
36 of the time it takes to grow to adult size (e.g. Bonner, 1965; Charnov, 1993;  
37 Charnov & Berrigan, 1993; Lindstedt & Swain, 1988). This approach assumes a  
38 tradeoff between resources spent growing and resources spent reproducing. Within  
39 a lifespan, the longer it takes to reach reproductive age, the shorter the period in  
40 which reproduction can take place (Charnov, 1993). This brings us back to rates of

1 extrinsic mortality, which set the length of lifespan and the probability of reaching  
2 reproductive age. Human maturational timing has been argued to be consistent  
3 with the pattern expected based on the length of the human lifespan (e.g. Alvarez,  
4 2000; Hawkes et al., 1998), and in this view evolution of delayed maturation  
5 requires “no special explanation” (Blurton Jones & Marlowe, 2002, p. 201). It  
6 is a byproduct of long lifespan. However, across primates, there is unexplained  
7 variance in life history traits even when the effects of body size are controlled (e.g.  
8 Harvey & Clutton-Brock, 1985; Harvey & Zammuto, 1985; Lindstedt & Swain,  
9 1988; Pereiara, 1993; Watts & Pusey, 1993). Further, explanation of long human  
10 lifespan is still needed. Several theories seek to explain important features of this  
11 uniquely human pattern (e.g. Bogin, 1999; Flinn et al., manuscript; Hawkes et al.,  
12 1998; Kaplan et al., 2000), here I focus on just one of those theories.

13 Kaplan et al. (2000) integrate insights from previous models, but note that  
14 theirs diverges from others in key ways. They propose that our extended juvenile  
15 dependency and lifespan are the product of the co-evolution of: (1) the dietary  
16 transition to high-quality, difficult-to-acquire foods (primarily game); (2) increased  
17 investment in the learning of complex subsistence strategies to exploit such foods;  
18 (3) increased food sharing and provisioning of conspecifics; and (4) the extension  
19 of this food provisioning to others when they are sick, which assisted recovery  
20 and reduced mortality risk. According to this model, as hominid dietary reliance  
21 on high quality, difficult-to-acquire game resources increased, fitness benefits  
22 could be realized from lengthening the pre-reproductive period and, hence, the  
23 period of foraging skill acquisition. This led to the co-evolution of an increased  
24 flow of resources from older individuals to juveniles in order to support this  
25 period of learning. This development was problematic because the fitness benefits  
26 of an extended learning period would be reduced by the increased probability of  
27 mortality during this lengthened pre-reproductive stage (e.g. Horn, 1978; Kaplan  
28 et al., 2000; Pagel & Harvey, 1993; Pereira, 1993). However, if provisioning  
29 to weaned juveniles was extended across periods of illness and injury, it could  
30 have effectively reduced mortality risk during the pre-reproductive lifespan, and  
31 extended juvenile period and delayed reproduction could co-evolve (Kaplan et al.,  
32 2000).

33 To date, a number of studies have presented data on food-sharing, productivity,  
34 and life-history (Bird & Bliege Bird, 2002; Bliege Bird & Bird, 2002; Blurton Jones  
35 & Marlowe, 2002; Bock, 2002; Hill & Kaplan, 1999; Hill et al., 2001; Sugiyama,  
36 2004; Walker et al., 2001), but systematic evidence regarding the occurrence and  
37 duration of injury, illness, and disability from this perspective is limited for forager  
38 and forager-horticulturalist populations living without regular access to Western  
39 medicine. Where there are reports, potentially disabling health insults are common  
40 (e.g. Bailey, 1991; Baksh & Johnson, 1990; Gurven et al., 2000; Hill & Hurtado,

1 1996; Howell, 1979; Kaplan et al., 2000; Sugiyama, 2004; Sugiyama & Chacon,  
2 2000), and sometimes result in disability that lasts for prolonged periods (e.g.  
3 Gurven et al., 2000; Sugiyama, 2004; Sugiyama & Chacon, 2000). Besides the  
4 immediate problem of surviving a health crisis once it occurs, juveniles also face  
5 fitness costs from lowered nutritional intake when adult providers suffer disabling  
6 incidents that prevent them from foraging (Sugiyama & Chacon, 2000). This is  
7 particularly likely where high quality but difficult to acquire items such as game  
8 are key components of the diet, as they are for many human foragers, and where  
9 juveniles are dependent on adult provisioning for a significant portion of their  
10 diet (e.g. Kaplan et al., 2000; Kelly, 1995; Sugiyama & Chacon, 2000; Tooby &  
11 DeVore, 1987).

12 In a variety of primates (including humans) and other mammals, nutritional  
13 deficits are associated with a number of fitness costs, including shorter reproductive  
14 lifespan, fewer offspring, delay of menarche and onset of puberty, lower proportion  
15 of live births, lower infant body weight, and increased juvenile mortality (see e.g.  
16 Altmann, 1991; Fritch & McArthur, 1974; Green et al., 1986; Hill & Hurtado,  
17 1996; Kohrs et al., 1976; Manocha & Long, 1977; Prentice et al., 1987; Riley et al.,  
18 1993; Schwartz et al., 1988). Among Yora foragers of Peru, prolonged recovery  
19 from injury to an average hunter resulted in an 18% reduction in average daily per  
20 capita protein intake in the village. Injury to the highest producing hunter would  
21 result in reduction of average daily per capita protein intake by approximately  
22 37% (Sugiyama, 1996; Sugiyama & Chacon, 2000). Thus, in foraging contexts  
23 with high levels of food transfers, disability among adults may severely impact  
24 nutrition of not only those juveniles whose primary caretaker may be temporarily  
25 disabled, but also all juveniles within the sharing network (Sugiyama & Chacon,  
26 2000).

27 Although these findings show clear costs to being sick and injured, and  
28 suggest that provisioning to sick and injured individuals could reduce juvenile  
29 mortality among people in small-scale, subsistence-based societies with little  
30 access to Western medical attention, systematic data on the occurrence and  
31 duration of illness and injury during the pre-reproductive lifespan are limited. If  
32 human delayed maturity and human longevity co-evolved with juvenile mortality  
33 reduction resulting from the extension of provisioning to times of health crisis, then  
34 the following are predicted to have characterized our recent evolutionary past:  
35 (1) health risk was a recurrent force during the pre-reproductive lifespan (even  
36 though risk-taking behavior may show facultative lifetime variation); (2) juveniles  
37 experienced health insults that would have been lethal without provisioning; (3)  
38 the probability and distribution of their occurrence was sufficiently great; that (4)  
39 provisioning during health crises effectively reduced juvenile mortality rates. Of  
40 course, paleopathological evidence can be used to test these assumptions, but doing

1 so entails problems (e.g. Steckel et al., 2002; Sugiyama, 2004; Wood et al., 1992)  
2 that ethnobiological evidence can help alleviate (Sugiyama, 2004; Walker et al.,  
3 1998).

4 Here I report on the types, causes, and duration of injuries and illnesses  
5 suffered during the pre-reproductive lifespan of Shiwiar forager-horticulturalists  
6 of the Ecuadorian Amazon, based on physical evidence and reported occurrence  
7 of pathologies. I address the following questions: (1) what health insults do  
8 individuals in this population suffer from during the pre-reproductive lifespan?  
9 (2) With what frequency do these episodes occur? (3) With what frequency and  
10 duration do these cases cause disability severe enough to necessitate survival  
11 assistance during the pre-reproductive portion of the lifespan? and (4) What  
12 are the demographic and fitness effects of individuals having received long-term  
13 provisioning without which they are likely to have died?  
14  
15

## 16 **STUDY POPULATION**

17

18 The Shiwiar are a Jivaroan-speaking people who live in the southern Oriente  
19 (tropical forest) of Ecuador and northeastern Peru. Approximately 2000 Shiwiar  
20 occupy a region along the Corrientes River and its tributaries in the upper Amazon.  
21 Long-term direct contact with non-indigenous populations began in the later  
22 1970s, when Shiwiar actively solicited missionary contact to reduce mortality  
23 from raiding and warfare. Mission contact also provided greater access to Western  
24 goods than previously used trade networks. Prior to this, Shiwiar lived in scattered  
25 households linked by marriage ties and the influence of powerful individuals  
26 (Descola, 1988). Unnavigable rivers, hostility toward outsiders, and border conflict  
27 between Ecuador and Peru have limited colonial incursions. Most Shiwiar villages  
28 have now cut small dirt airstrips in the forest, around which houses form loose  
29 clusters. These airstrips provide some access to medical and other facilities outside  
30 of Shiwiar territory via missionary light aircraft, although access is neither regular  
31 nor consistent. Shiwiar subsistence is based on foraging and horticulture, and  
32 internal politics are governed by traditional big-man, consanguine- and affine-  
33 based alliances.

34 Shiwiar access to Western medicine and medical supplies varies considerably  
35 across communities and through time. Emergency air flights can sometimes be  
36 arranged via a two-way radio. Villages also have health promoters trained to  
37 dispense first aid and medicine as best they can given a chronic shortage in supplies.

38 Nevertheless, in day-to-day life, the Shiwiar lifeway mirrors many aspects  
39 of our evolutionary past relevant to adaptations related to mortality reduction  
40 via provisioning during health crises. The Shiwiar live in small kin-based

1 communities in which some foods, particularly game, are shared; they rely on  
2 hunting and fishing for most of their dietary fat and protein, and on gathered  
3 plant products for fruits, starch, construction, and tool material. They are closely  
4 related to most of the people with whom they have daily interaction, and come  
5 into repeated contact with relatively few people. They lack access to modern  
6 contraceptives, baby formula, and have limited access to Western medicine, but  
7 a detailed system of indigenous medical knowledge.

8 Shiwiar also grow at least 26 horticultural products (Descola, 1988). The  
9 predominant starch in the diet is manioc; other root crops (e.g. sweet potatoes,  
10 yams), and plantains are also important. Each female head of household has  
11 between two and four gardens at different stages of production, and on most  
12 mornings women go to the garden to harvest, replant, and weed. Gardens are larger  
13 than necessary for household consumption, even when the need to buffer the risk  
14 of losses due to animal and other pests is considered (Descola, 1988). Horticultural  
15 production provides a highly productive, reliable, and spatially concentrated patch  
16 of carbohydrate resources. Large-scale clearing is done via *mingas*, village-wide  
17 parties of cooperative labor exchange involving both men and women.

18 Both blowguns and muzzle-loading shotguns are used in hunting, although  
19 single-shot cartridge shotguns are increasingly used when cash is available for  
20 the relatively costly shells. A wide variety of small, medium and large game are  
21 taken. Hunting returns are relatively high, and the day-to-day risk of returning  
22 from a hunting trip without game is low (Sugiyama, 2000; Sugiyama & Chacon,  
23 2000). Fishing is done either with hooks and line or by using *barbasco* fish poison  
24 (Sugiyama, 2000; Sugiyama & Chacon, 2000; Walker et al., 1998). Finally, the  
25 industrially produced machete is used ubiquitously as an all-purpose cutting, chop-  
26 ping, and digging tool. Metal pots, hooks, axes, and knives are also widely used.

27 Shiwiar life includes a number of health risks including lacerations from a  
28 variety of objects such as tools, branches, sticks, spines, logs, and falling trees.  
29 Local and parasitic infections, as well as diseases such as chicken pox, measles, and  
30 malaria are common. Animal and insect bites and stings are a ubiquitous problem.  
31 The threat from several species of venomous snake is serious (Sugiyama, 2004).  
32 Further, although jaguar predation is no longer a major concern because of shotgun  
33 use in hunting, informants state that in the past, when shot and powder were less  
34 available, jaguars were a real threat.

35  
36  
37  
38

## METHODS

39 Participant observation and interviews that recorded injuries and illness during  
40 the study periods provide the ethnographic context for Shiwiar reactions to injury.

1 Formal and informal interviews were conducted between 1993 and 1998 in four  
2 Shiwiar villages to gather genealogical and life history data. Physical examination  
3 of 19 male and 20 female Shiwiar individuals ranging in age from three to 50 years  
4 was conducted to document scars, broken bones, or other observable signs of past  
5 pathological events. An additional male was not examined during the study because  
6 he was not present in the village at the time. However, all critical information  
7 regarding a near-lethal snakebite was available for him from previous interviews,  
8 so it was included in the analysis (Sugiyama, 2004).

9 The examination for evidence of injury or illness proceeded in a standardized  
10 way. Beginning with the right foot, the examination proceeded up the right leg as  
11 far as was comfortable for the informant, and then down the left leg. The left and  
12 right arm were then examined, followed by the front and then rear of the torso  
13 and neck, followed by the face and head. Scars and evidence of broken bones  
14 were noted on standardized forms depicting front and rear line drawing views of a  
15 human form, and enlarged views of the hands and feet. Each health insult recorded  
16 was coded as visible, reported (by the informant), current, or some combination  
17 of these in order to specify the evidence upon which the recording of each  
18 observation was based. For each scar or evidence of a broken bone observed, the  
19 subject was asked to provide information about the cause, activity engaged in at  
20 the time, and age at which the event occurred. Informants from one of the sample  
21 villages ( $n = 17$ ) were also asked the duration of disability if applicable, and  
22 this information crosschecked with other informants. A standard set of questions  
23 about past illnesses, injuries, and treatment received (either from a shaman or  
24 Western medical practitioner) was then administered (Sugiyama, 2004). Disability  
25 duration gives a clearer indication of the proportion of health insults that would  
26 have been lethal without extended provisioning, and thus serves as a means by  
27 which juvenile mortality reduction via provisioning during health crises can be  
28 examined.

29 While the methods used provide systematic data collection on past and present  
30 illness and injury, they do entail problems. All scars on young individuals are  
31 easily observed, but older individuals have been subject to so many lacerations,  
32 abrasions, and infections that only the most prominent or most recent can be  
33 accurately recorded. Numbers of injuries recorded for older adults reported here  
34 thus primarily reflect the most recent or most serious injuries. For individuals  
35 under about 25 years of age, all health insults leaving visible evidence could be  
36 recorded. Methods used were time consuming however, and informants no doubt  
37 differed in levels of patience and quantity of details they were willing to supply.  
38 Although these problems could not be entirely solved, they were more likely to  
39 result in under-reporting than over-reporting. Independent means of crosschecking  
40



1 information suggest that overestimation or false reporting was highly likely to be  
2 exposed. The principle source of evidence was physical observation; reports of  
3 broken bones were verified both by tactile examination for evidence of a healed  
4 fracture and by corroboration by other informants. For information on both  
5 disability and health insults leaving no direct physical evidence, informant reports  
6 were crosschecked with others who were present at the time of the injury/illness  
7 or knowledgeable about the incident in question. Incidents causing significant  
8 disability were known by multiple informants, and the most significant were  
9 common knowledge.

10 On the other hand, infectious disease, pathogen load, and common illness are  
11 certainly underestimated by the methods used. Few infectious diseases leave  
12 visible traces, and intestinal parasitic infection is endemic, but is not visibly  
13 assessable. Only the most recent or severe are likely to have been recalled or  
14 recorded. I did not attempt to ask subjects about common colds or influenza,  
15 because while frequency was expected to be high, reported frequency was not  
16 expected to be accurate. Conversely, informants were asked if they had suffered  
17 from the following infectious diseases or parasitic infections known to occur in  
18 Shiwiar territory: measles, whooping cough (pertusis), tuberculosis, chicken pox  
19 (varicella), leishmaniasis, and malaria.

20 Finally, some health insults do not fit neatly into one category. As a guiding  
21 principle, I categorized illness or injury type according to the proximate cause  
22 of the condition with regard to its potential for causing disability. For instance, a  
23 health insult was recorded as laceration if the wound did not result in infection,  
24 but as infection if serious infection occurred, because for most lacerations  
25 observed, subsequent infection had greater potential to cause disability. That said  
26 however, severe lacerations were noted as such, and, subsequent infection was  
27 limited compared to what I expected based on previous observations (Sugiyama  
28 & Chacon, 2000). This may be because Shiwiar usually pack severe lacerations  
29 with masticated plant matter that appears to be an effective antibiotic. Similarly,  
30 insect bites that resulted in subsequent minor skin infection were recorded as  
31 infections, because such infections may become severe as other ectoparasites  
32 infect the wound. Finally, an incident was only recorded as pain when it occurred  
33 beyond simultaneous occurrence of another condition. For example, although  
34 snakebite causes extreme pain, at the time of the bite it is the venom and not  
35 the pain that poses the greatest threat of disability and death. Thus, pain of this  
36 type was recorded as snakebite. However, subsequent nerve or tissue damage  
37 can cause pain and limitations on mobility that can last long after the threat  
38 from venom is past. Accordingly, pain of this type was reported as pain, not  
39 snakebite.

40

## RESULTS

### *Age/Sex Distribution of Sample*

As of 1998, 410 Shiwiar living in six villages were in the core study area, with an additional 87 siblings or offspring of core area individuals living in surrounding villages. Usual resident population in the two villages from which data on health insults were collected was 63 and 103 persons, respectively. At the time of study in 1994–1995, long-term visitors were present in one village, while many residents from the other village were away visiting, such that there were 67 and 87 persons present in each village respectively. The sample of 40 individuals represents 24% of the usual resident population in these villages, and about 9.76% of the population of the core study area. The 17-person disability sample represents 16.5% of the population of the second village, and includes individuals from 10 of the 12 village households. Residents of the other households were not present when data were collected. A Kruskal-Wallis test shows no significant differences between the ages of males and females in the sample overall ( $\chi^2 = 28.6$ ,  $df = 28$ ,  $p = 0.42$ ) or by 10-year age cohorts ( $\chi^2 = 1.514$ ,  $df = 4$ ,  $p = 0.82$ ).

### *Health Insults Suffered*

A total of 678 injuries and illnesses were recorded for the 40 individuals examined. The most commonly observed incidents were lacerations, followed by infections (including infectious disease), bites and stings, puncture wounds, abrasions, pain (either chronic or periodic), broken bones, and burns. Sugiyama (2004) provides detailed analysis of occurrence, definition, and distribution of pathologies by type, cause, and sex of victim across the lifespan. Data on health insults suffered during the pre-reproductive lifespan were extracted from this sample in two ways. Twenty-two sample individuals, 10 females and 12 males (aged 3–25), were pre-reproductive and all health insults they suffered were therefore included. Eighteen individuals, nine females and nine males, had begun reproduction by the time of the study. For these adults, health insults for which an age of occurrence or phase of lifetime (e.g. childhood or after first child) were reported, were examined for cases occurring prior to the individuals' age at first reproduction. Together, these yielded 456 health insults occurring prior to first reproduction. Data analysis presented below is based on these incidents. The Kruskal-Wallis test indicates no significant age difference between the sexes within this sub-sample ( $\chi^2 = 15.546$ ,  $df = 14$ ;  $p = 0.34$ ).

Table 1. Number of Shiwiar Health Insults Prior to First Reproduction.

Health Insult Type	Observed	Residual	Percent
Laceration	152 <sup>a</sup>	114.0	33.3
Bite/Sting	118 <sup>a,b</sup>	80.0	25.9
Infection	99 <sup>b,c</sup>	61.0	21.7
Puncture	30 <sup>c</sup>	-8.0	6.6
Abrasion	22	-16.0	4.8
Scars unknown	10	-28.0	2.2
Fracture	9	-29.0	2.0
Burn	6	-32.0	1.3
Contusion	4	-34.0	0.9
Pain	4	-34.0	0.9
Blisters	1	-37.0	0.2
Irritation	1	-37.0	0.2
Total	456		100.0

Pl. provide note for a, b and c.

As expected, significant differences were observed in the relative frequencies with which different types of health insults were observed (Table 1). Lacerations were the most frequently observed health insults, and were observed significantly more often than bites and stings, the second leading cause. Although the number of bites/stings and infections (the third leading cause of health insults) observed do not significantly differ, both were significantly more common than puncture wounds. Thus, infections were observed significantly more often than all remaining types of health insults: i.e. abrasions, scars of unknown type, fractures, burns, contusions, pain, blisters, and irritated skin in descending order observed (Table 1).

Sex Differences in Health Insults Suffered

Males generally suffer disproportionately more illness and injury than females, a fact attributable both to higher male behavioral risk-taking, aggression, and susceptibility to health problems (e.g. Dabbs & Dabbs, 2000; Daly & Wilson, 1988; Kraemer, 2000). As expected, Shiwiar males suffer significantly more health insults than females in the pre-reproductive years (Table 2). Analysis by type indicates that prior to age at first reproduction males were more frequent victims of laceration, infection, bites/stings, and puncture wounds than females. Males and females did not differ in number of abrasions or scars of unknown cause. Contusions, fractures, and pain did not occur with sufficient frequency for statistical tests to be meaningfully performed. In addition, only males suffered burns.

**Table 2.** Sex Differences in Shiwiar Health Insults Prior to First Reproduction.

Health Insult	Male		Female		Total	$\chi^2$	df	p-Value
	Observed	Expected	Observed	Expected				
Abrasion	15	11.0	7	11.0	22	2.91	1	0.088
Bite/Sting	82	59.0	36	59.0	118	17.93	1	0.000
Blisters	1		0		1	a,b		
Burn	6	6.0	0	0	6	a,b		
Contusion	3		1		4	b		
Fracture	5		4			b		
Infection	67	49.5	32	49.5	99	12.37	1	0.000
Irritation	0		1		1	a		
Laceration	111	76.0	41	76.0	152	32.24	1	0.000
Pain	2	2.0	2	2.0	4	b		
Puncture	23	15.0	7	15.0	30	8.53	1	0.003
Scars	5	5.0	5	5.0	10	0.00	1	1.00
Total	319	228.0	135	228.0	456	44.24	1	0.000

<sup>a</sup>This variable is constant. Chi-Square test cannot be performed.

<sup>b</sup>Expected cell frequency less than 5. Chi-square test not performed.

### *Causes of Health Insults Suffered in the Pre-reproductive Years*

The specific cause of each type of health insult was recorded at the time of data collection. Table 3 shows the frequency with which the 37 specific causes of observed health insult occurred during the pre-reproductive lifespan. There are significant differences in the frequency with which different kinds of health insults occurred. The four most frequently observed were bites from vampire bats ( $n = 98$ ), followed by machete wounds ( $n = 75$ ), lacerations, contusions, abrasions, and broken limbs from being hit by, or running into spines, branches, or logs ( $n = 70$ ), and insects (mostly ectoparasites [ $n = 60$ ]). Together these account for 66.4% of all health insults observed. Although each of these causes occur significantly more often than expected by chance, they also differ significantly from one another in relative frequency with which they are observed due to the difference between the highest frequency cause – vampire bat bites – and the least frequent – insects. Falls, lance wounds, chicken pox, snake bites, malaria, axe wounds, leishmaniasis (a tropical protozoic infection), and burns from cooking fires occur repeatedly but significantly less frequently than the first set of causes (Table 3).

Of particular interest when considering health risks in evolutionary perspective is the comparison of incidents caused by introduced technology and those that arise from non-introduced causes. Study results would be problematic if most health insults were due to the direct influence of Western technology. This does not

**Table 3.** Causes of Shiwiar Health Insults Prior to First Reproduction.<sup>a</sup>

Cause	Observed	Residual	Percent
Bat	98 <sup>a,b</sup>	86.0	21.5
Machete	75 <sup>a</sup>	63.0	16.4
Plant	70 <sup>a</sup>	57.7	15.3
Insect	60 <sup>a,b,c</sup>	48.0	13.2
Unknown	49	37.0	10.7
Fall	13 <sup>c</sup>	1.0	2.9
Lance	11	-1.0	2.4
Varicella	11	-1.0	2.4
Snake	8	-4.0	1.8
Malaria	7	-5.0	1.5
Axe	6	-6.0	1.3
Leishmaniasis	6	-6.0	1.3
Fire	5	-7.0	1.1
Injection	4	-8.0	0.9
Hit	3	-9.0	0.7
Knife	3	-9.0	0.7
Measles	3	-9.0	0.7
Assault	2	-10.0	0.4
Fingernail	2	-10.0	0.4
Pertusis	2	-10.0	0.4
Scorpion	2	-10.0	0.4
Amoebae	1	-11.0	0.2
Ant	1	-11.0	0.2
Bee	1	-11.0	0.2
Blisters	1	-11.0	0.2
Bot fly	1	-11.0	0.2
Collision	1	-11.0	0.2
Dog	1	-11.0	0.2
Ear piercing	1	-11.0	0.2
Fight	1	-11.0	0.2
Fish spine	1	-11.0	0.2
Harpoon	1	-11.0	0.2
Howler monkey	1	-11.0	0.2
Oncherosiasis	1	-11.0	0.2
Person	1	-11.0	0.2
Pirana	1	-11.0	0.2
Shotgun	1	-11.0	0.2
Total	456		100.0

Pl. provide note for b and c.

<sup>a</sup>For each cause, expected = 12.

37  
38  
39  
40

1 appear to be the case. Although machete cuts are highly prevalent, chi-square tests  
2 reveal that when introduced causes of injury (i.e. wounds by machete, axe, knife,  
3 injection, and shotgun) are compared to all other types of health insults, the latter  
4 occur significantly more often than the former ( $\chi^2 = 128.85$ ,  $df = 1$ ,  $p = 0.000$ ).  
5 It is possible that the significantly higher frequency of insults caused by natural  
6 or local versus introduced agents is a reflection of limited contact with modern  
7 products because 32 of the 37 observed causes of health insult were not directly  
8 related to introduced tools. Moreover, when expected frequencies are adjusted to  
9 account for this fact (86.5% of 407 = 352.06 for natural or local and 13.5% of  
10 407 = 54.95 for introduced, respectively), natural causes of injury and illness are  
11 still more frequently observed than ones attributable to introduced technologies  
12 ( $\chi^2 = 24.40$ ,  $df = 1$ ,  $p < 0.000$ ).

13 Within health insult type, some causes are more frequently observed than others  
14 are. Vampire bats are by far the most common of the nine types of animal bites and  
15 stings observed ( $\chi^2 = 621.68$ ,  $df = 8$ ,  $p < 0.000$ ), accounting for 98 of the total  
16 118 bites and stings. Fifteen different causes of local, parasitic, and contagious  
17 infections were recorded, again differing significantly in how often they caused  
18 infection ( $\chi^2 = 381.76$ ,  $df = 14$ ,  $p < 0.000$ ), with mild skin infections from  
19 insect bites that are scratched open most common. Chicken pox, leishmaniasis,  
20 and malaria were observed significantly less often. Sources of lacerations also  
21 differ significantly in how often they caused lacerations ( $\chi^2 = 339.86$ ,  $df = 10$ ,  
22  $p < 0.000$ ). Among these, machetes caused significantly more lacerations than the  
23 next most common, plants (i.e. running into logs, branches, sticks [ $\chi^2 = 16.96$ ,  
24  $df = 1$ ,  $p < 0.000$ ]), unknown causes or lance wounds ( $\chi^2 = 13.56$ ,  $df = 1$ ,  
25  $p < 0.000$ ). All other causes – axes, falls, knives, being hit, lacerated by a howler  
26 monkey, and assaults – occurred only a few times each. Puncture wounds, were  
27 most often caused by plants ( $\chi^2 = 29.27$ ,  $df = 6$ ,  $p < 0.000$ ).

### 28 29 30 *Disabling Health Insults* 31

32 Informants from one sample village were asked about the duration of disability  
33 associated with each reported pathological condition. This information was then  
34 crosschecked with other informants. Disability was defined as a condition that  
35 prevented the informant from leaving the house or engaging in foraging or garden  
36 work. The Kruskal-Wallis test indicates that the sex composition of this sub-  
37 sample, including eight males and nine females, does not differ significantly from  
38 the rest of the larger sample ( $\chi^2 = 0.100$ ,  $df = 1$ ,  $p = 0.725$ ) although it does  
39 contain an older age cohort ( $\chi^2 = 14.31$ ,  $df = 1$ ,  $p = 0.000$ ), ranging from seven  
40 to 50 years of age. This means that they have more pre-reproductive life experience

1 than the sample as a whole. It includes more individuals having survived their  
 2 entire juvenile lifespan, and thus provides a more complete picture of how likely  
 3 one is to encounter disability severe enough to demand provisioning for survival.  
 4 Shiwiar place high value on vigorous work and industriousness, therefore reports  
 5 of disability reflect conditions that physically precluded subsistence work or the  
 6 mobility necessary for subsistence work (Sugiyama, 2004). Informants usually  
 7 reported disability duration estimates in even units of days, weeks, and months;  
 8 these were converted to number of days for comparison.

9 I could determine either the age of occurrence, or whether an incident occurred  
 10 before or after first reproduction, for 131 of the 215 conditions recorded in the sub-  
 11 sample. Fifty-nine of these were cases in which disability was observed. Comparing  
 12 the age at which the disability occurred with genealogical data allows calculation of  
 13 the probable effects of disability on mortality and reproduction. Thirty-three of 59  
 14 (55.9%) cases of disability affected individuals prior to first reproduction (Table 4).  
 15 Duration of these disabilities ranged from one day to one year, with 12 of the 33  
 16 cases (22%) causing disability of one month or longer. These 12 incidents were  
 17 distributed among 10 of the 17 individuals (58.82%) for whom data were available.

18  
 19  
 20 **Table 4.** Frequency of Shiwiar Disability by Duration and Lifestage.

Disability Duration (Days)	Juvenile Period or Age During Adulthood														Total	
	Juvenile	21	22	23	24	25	26	27	28	30	35	37	39	41		43
1	4							1								5
2	1															1
3			1										1			2
4				1												1
6	1															1
7	6				1		1		1		1			1		11
10															1	1
14	2	2					1			1					1	7
15	3	1														4
17	1															1
20	1															1
21	2			1												3
30	7						1		1			1			1	11
45	2		1													3
60	1			1		1						1				4
90	1															1
180			1													1
365	1															1
Total	33	3	3	3	1	1	3	1	2	1	1	2	1	1	3	59

**Table 5.** Frequency of Disability by Individual, Duration, and Reproductive Success.

ID	Age	Sex	Disability Duration (Days)			RS (Descending Generations)			Descendants (% Population)			
			Total	7–13	14–29	30<	1st	2nd	Total	In Village	% Village	% Tot. Pop
8	16	f	2	1	1	0	0	0	0	0	0	
<b>5<sup>a</sup></b>	<b>18</b>	<b>f</b>	<b>8</b>	<b>2</b>	<b>3</b>	<b>2</b>	<b>0</b>	<b>0</b>	<b>0</b>	<b>0</b>	<b>0</b>	
<b>10<sup>a,b</sup></b>	<b>18</b>	<b>f</b>	<b>3</b>	<b>0</b>	<b>1</b>	<b>1</b>	<b>2</b>	<b>0</b>	<b>2</b>	<b>1.94</b>	<b>0.40</b>	
<b>6<sup>a</sup></b>	<b>25</b>	<b>f</b>	<b>1</b>	<b>0</b>	<b>0</b>	<b>1</b>	<b>0</b>	<b>0</b>	<b>0</b>	<b>0</b>	<b>0</b>	
<b>16<sup>a,b,c</sup></b>	<b>27</b>	<b>f</b>	<b>6</b>	<b>0</b>	<b>1</b>	<b>3</b>	<b>2</b>	<b>0</b>	<b>2</b>	<b>1.94</b>	<b>0.40</b>	
<b>12<sup>a,b,c</sup></b>	<b>29</b>	<b>f</b>	<b>14</b>	<b>1</b>	<b>1</b>	<b>4</b>	<b>8<sup>e</sup></b>	<b>0</b>	<b>8</b>	<b>8<sup>e</sup></b>	<b>7.78</b>	<b>1.61</b>
<b>11<sup>a,b</sup></b>	<b>37</b>	<b>f</b>	<b>4</b>	<b>1</b>	<b>0</b>	<b>3</b>	<b>9<sup>d</sup></b>	<b>5<sup>f</sup></b>	<b>14</b>	<b>7<sup>d</sup></b>	<b>6.80</b>	<b>2.82</b>
<b>2<sup>a,b,c</sup></b>	<b>43</b>	<b>f</b>	<b>10</b>	<b>2</b>	<b>3</b>	<b>5</b>	<b>10</b>	<b>8</b>	<b>18</b>	<b>15</b>	<b>14.56</b>	<b>3.6</b>
17	7	m	2	0	2	0	0	0	0	0	0	
<b>15<sup>a</sup></b>	<b>15</b>	<b>m</b>	<b>5</b>	<b>0</b>	<b>3</b>	<b>2</b>	<b>0</b>	<b>0</b>	<b>0</b>	<b>0</b>	<b>0</b>	
<b>1<sup>a,b</sup></b>	<b>22</b>	<b>m</b>	<b>4</b>	<b>2</b>	<b>0</b>	<b>2</b>	<b>2</b>	<b>0</b>	<b>2</b>	<b>2</b>	<b>1.94</b>	<b>0.40</b>
7 <sup>b</sup>	24	m	7	1	5	0	3	0	3	3	2.91	6.0
14 <sup>b</sup>	34	m	1	0	0	0	9 <sup>e</sup>	0	9	9 <sup>e</sup>	8.74	1.81
3 <sup>b,c</sup>	36	m	1	na	na	1	3	0	3	3	2.91	0.60
4 <sup>b</sup>	37	m	3	2	1	0	5	1	6	6	5.83	1.20
9 <sup>b</sup>	43	m	3	2	0	0	9 <sup>d</sup>	5 <sup>f</sup>	14	7 <sup>d</sup>	6.80	2.82
<b>13<sup>a,b</sup></b>	<b>50</b>	<b>m</b>	<b>8</b>	<b>1</b>	<b>1</b>	<b>3</b>	<b>11</b>	<b>14</b>	<b>25</b>	<b>24</b>	<b>23.30</b>	<b>5.03</b>
LLWPJ subtotal							44 <sup>g</sup>	27 <sup>g</sup>	71 <sup>g</sup>	60 <sup>g</sup>	58.25 <sup>g</sup>	14.63 <sup>g</sup>
Total			82	15	22	27	54 <sup>g</sup>	28 <sup>g</sup>	82 <sup>g</sup>	73 <sup>g</sup>	70.87 <sup>g</sup>	16.49 <sup>g</sup>

<sup>a</sup> Bold: individuals who suffered health insult likely to be lethal without provisioning during juvenile period (LLWPJ).

<sup>b</sup> Individuals who had started reproduction by time of the study.

<sup>c</sup> Individuals who suffered health insult likely to be lethal without provisioning after age of first reproduction.

<sup>d</sup> Represents same individuals: Offspring of a married couple both included in the sample.

<sup>e</sup> Eight of these nine are same individuals: offspring of a married couple both included in the sample.

<sup>f</sup> Represents same individuals.

<sup>g</sup> Totals calculated based on descendants of a couple both of whom are included in the subsample only once.

40  
39  
38  
37  
36  
35  
34  
33  
32  
31  
30  
29  
28  
27  
26  
25  
24  
23  
22  
21  
20  
19  
18  
17  
16  
15  
14  
13  
12  
11  
10  
9  
8  
7  
6  
5  
4  
3  
2  
1



1 For humans, death from dehydration occurs within about three–seven days in  
2 temperate environments (Egland, 2002) and, depending on a number of variables,  
3 hunger strikers have died from starvation after an average of 55 days if no food  
4 is taken, but medical attention, water, and salt are available (Beresford, 1987).  
5 Recent examples suggest that when sick or injured, a person can survive without  
6 care for about 30 days with water and a little food before death (Krakauer, 1996).  
7 Younger juveniles succumb more rapidly (Egland, 2002; see discussion below).  
8 I assume here that sick and injured individuals could acquire water and salt, but  
9 would be unable to forage. Because food deprivation during periods of illness  
10 or injury significantly hinder recovery, I therefore assume that 30 days without  
11 food under these circumstances would likely prove lethal. Ten of the 17 people in  
12 the second village sub-sample suffered an illness or injury of this severity during  
13 their juvenile years. Of these, seven had begun reproduction by the time of the  
14 study. If provisioning was required for individuals to survive disability of 30 days  
15 or longer, then without provisioning, over half of the sub-sample would have died  
16 sometime during their pre-reproductive years.

17 Fifty-four offspring and 28 grandchildren were born to individuals in the second  
18 village sample (children/grandchildren born to parents/grandparents who are both  
19 in the sample were counted only once), although only two members of the sample  
20 had completed or were approaching probable completed fertility. Provisioning  
21 during health crises can have large effects on individual fitness. Of the living  
22 descendants of individuals in the second village sample, 44 out of the 54 (81.5%)  
23 first descending generation, and 27 out of 28 (96.4%) of the second descending  
24 generation were born after a direct ancestor in the sample survived a juvenile health  
25 crisis likely to be fatal without provisioning. In other words, 71 of the 82 individuals  
26 born to members of the subsample, were either the child or grandchild of an indi-  
27 vidual who would not have survived to reproduce had s/he not received extended  
28 provisioning during a juvenile health crisis. Further, three individuals (ID numbers  
29 2, 11, and 13) who survived juvenile health crises likely to be lethal without provi-  
30 sioning, are either the parent or grandparent of 27 second-generation descendants.  
31 Together, these three individuals are either parent or grandparent of 11.5% of the  
32 Shiwiar population in the study area, and 45% of sub-sample village residents. One  
33 (ID number 13), is the parent or grandparent of 5% of the Shiwiar population block  
34 and 23% of the people in their village. Another (ID number 2) is the direct ancestor  
35 of 3.6% of the population and 14.5% of the people in their village (Table 5).

36  
37  
38  
39

## **DISCUSSION AND CONCLUSION**

40 To date, suggestive evidence indicated that provisioning during health-care crises  
could reduce juvenile mortality in forager and forager-horticulturalist societies.

1 No systematic data have been available on the overall degree to which individuals  
2 in such societies suffer injury/illness of sufficient duration to interfere with  
3 subsistence. Data presented in this paper address these questions more directly. As  
4 expected, health insults represent a significant hazard during the pre-reproductive  
5 lifespan in the sample population. Illness and injury are frequently encountered  
6 and widely distributed across individuals. Over half the sample individuals in the  
7 second study village suffered juvenile health crises expected to be lethal without  
8 extended care from others, indicating that provisioning effectively reduces juvenile  
9 mortality.

10 Further, it is clear that individual Shiwiari fitness is predicated on the existence  
11 of provisioning to temporarily disabled individuals. Of 54 offspring born to  
12 informants in the sub-sample, over 80% were born after a parent included in the  
13 sample suffered injury or illness likely to be fatal without extended provisioning.  
14 Of 28 individuals in the second descending generation of informants in the sub-  
15 sample, over 95% were born after a direct ancestor in the sample survived an  
16 incident likely to be fatal without provisioning. Additional work must be done to  
17 extend these findings, particularly through increasing sample size and constructing  
18 life history and mortality tables for the Shiwiari, so that the degree of mortality  
19 reduction via health-care provisioning can be estimated. Comparative studies  
20 among extant foraging and forager-horticulturalist societies living with little access  
21 to Western medical care are also critical.

22 My use of 30 days as an estimate of probable mortality without provisioning is  
23 rather crude, but I believe it is nevertheless reasonable. Outdoor survival guides  
24 report that adults can live for from three to seven days with no water, but here I  
25 conservatively assumed that even among those individuals who are temporarily  
26 disabled all but the most seriously sick or injured individuals could access water.  
27 However, chicken pox, measles, influenza, malaria, or other disease resulting in  
28 high fever, diarrhea and/or vomiting are such that dehydration, volume depletion,  
29 and poor nutrition can lead to rapid death. Food provisioning is critical in such  
30 cases, because even with access to water, volume depletion can lead to seizures,  
31 cerebral edema, or cardiovascular collapse when water but not sodium and other  
32 solutes are replaced (i.e. hyponatremic volume depletion; [Egland, 2002](#)). In the  
33 United States, about 5% of hospital admissions for children are the result of volume  
34 depletion. Worldwide, diarrheal illness is the third leading cause of death across all  
35 age groups (following respiratory and cardiac disease, and in developing countries  
36 volume depletion from diarrhea causes approximately 5 million deaths of children  
37 under the age of five each year ([Egland, 2002](#)). This means that the data presented  
38 here probably underestimate the effects of provisioning on mortality rates, and that  
39 the criteria for disability – defined here as a condition that prevented an informant  
40 from leaving the house or engaging in foraging or garden work – that would be

1 lethal without provisioning is conservative for such illnesses. Even though people  
2 can survive starvation alone for longer periods, and death from dehydration and  
3 volume depletion can occur much more quickly, particularly among juveniles,  
4 30 days seems a reasonable estimate of the time by which a condition causing  
5 disability severe enough to prevent foraging would be lethal without protection  
6 and provisioning. With large-bodied predators around, the time could well be  
7 shorter.

8 Although data supporting each component of the Kaplan et al. (2000) co-  
9 evolutionary model of human life-history evolution exist, other adaptationist  
10 explanations for elements of human life-history evolution have been proposed.  
11 These include Hawkes' and colleagues (1998, 2000) "grandmother" hypothesis,  
12 which posits that the dietary transition to high quality plant foods led to offspring  
13 provisioning, long lifespan, and with this extension the juvenile period was also  
14 drawn out (see though Kennedy, 2003; Peccei, 2001). The social intelligence  
15 hypothesis posits that the evolution of large brains and high intelligence is a  
16 consequence of an increasing intra-specific arms race for social intelligence (e.g.  
17 Alexander, 1989; Byrne & Whiten, 1997; Flinn et al., manuscript) which occurred  
18 after evolution of ecological foraging dominance (e.g. Flinn et al., manuscript).  
19 Finally, Miller's (2000) hypothesis is that increased human intelligence is largely  
20 the consequence of runaway sexual selection.

21 Blurton Jones and Marlowe (2002) challenge the assumption that adult foraging  
22 competence requires an extended period of learning, arguing that the reason  
23 Hadza children do not achieve adult levels of foraging competence is that they  
24 have insufficient size and/or strength to do so. Among Miriam Island foragers,  
25 children are reported to achieve adult foraging rates for some skill-intensive fishing  
26 techniques well before they achieve adult rates for techniques requiring less skill  
27 but greater size, speed or strength (Bird & Bliege Bird, 2002; Bliege Bird & Bird,  
28 2002; Bliege Bird et al., 1995). Although each of these hypotheses appears to  
29 have valid elements, uni-dimensional theories are unlikely to explain the complex  
30 complement of human traits ultimately at issue: (1) long pre-reproductive period  
31 and lifespan; (2) high levels of social and technical intelligence; (3) cultivation of a  
32 foraging niche based on acquisition of difficult-to-acquire, high-quality foods; (4)  
33 extended investment in offspring and in the learning of complex social and foraging  
34 strategies; and (5) conspecific provisioning of injured/ill individuals and of females  
35 and their offspring. Co-evolutionary models positing multiple sequential and/or  
36 simultaneous interacting evolutionary pressures appear necessary to explain these  
37 intersecting phenomena (e.g. Alexander, 1989; Flinn et al., manuscript; Kaplan  
38 et al., 2000).

39 Regardless of one's position with regard to these theories, the data reported  
40 here illustrate a clear point. If illness, injury and disability risk among the Shiwiar

1 are within the normal range of those associated with a foraging lifeway, then  
 2 whichever explanation of these phenomena one wishes to advance, the evolution  
 3 of mortality risk reduction via provisioning during health crises appears to be  
 4 a necessary feature that must be incorporated into one's model, either as a pre-  
 5 adaptation or co-evolutionary condition for the emergence of these features of the  
 6 human "cognitive niche" (Tooby & DeVore, 1987). Once juvenile provisioning  
 7 evolved and was provided during health crises, selection would be expected to  
 8 favor strategies for garnering strategic health-care provisioning during adulthood  
 9 as well (e.g. Gurven et al., 2000; Sugiyama, 1996, 2004; Sugiyama & Chacon,  
 10 2000). Without provisioning during health crises, features of the human lifeway  
 11 associated with long juvenile period, long lifespan, and long period of cognitive  
 12 development are unlikely to have evolved, because the fitness costs of extending the  
 13 pre-reproductive lifespan in the absence of extended juvenile provisioning during  
 14 health crises are overwhelmingly great.

## 15 16 17 REFERENCES

- 18  
19 Alexander, R. (1989). Evolution of the human psyche. In: P. Mellars & C. Stringer (Eds), *The*  
 20 *Human Revolution: Behavioural and Biological Perspectives on the Origins of Modern Humans*  
 21 (pp. 455–513). Princeton: Princeton University Press.  
 22 Altmann, S. (1991). Diets of yearling female primates predict lifetime fitness. *Proceedings of the*  
 23 *National Academy of Science*, 88, 420–423.  
 24 Alvarez, H. (2000). Grandmother hypothesis and primate life histories. *American Journal of Physical*  
 25 *Anthropology*, 113, 435–450.  
 26 Bailey, R. (1991). *The behavioral ecology of Efe pygmy men in the Ituri forest, Zaire*. Ann Arbor, MI:  
 27 University of Michigan Press.  
 28 Baksh, M., & Johnson, A. (1990). Insurance policies among the Machiguenga: An ethnographic analysis  
 29 of risk management in a non-western society. In: E. Cashdan (Ed.), *Risk and Uncertainty in*  
 30 *Tribal and Peasant Economies* (pp. 193–228). San Francisco, CA: Westview.  
 31 Beresford, D. (1987). *Ten men dead: The story of the 1981 Irish hunger strike*. New York: Harper  
 32 Collins.  
 33 Bird, D., & Bliege Bird, R. (2002). Children on the reef: Slow learning or strategic foraging? *Human*  
 34 *Nature*, 13, 269–298.  
 35 Bliege Bird, R., & Bird, D. (2002). Constraints of knowing or constraints of growing? Fishing and  
 36 collecting by the children of Mer. *Human Nature*, 13, 239–265.  
 37 Bliege Bird, R., Bird, D., & Beaton, J. (1995). Children and traditional subsistence on Mer, Torres  
 38 Strait. *Australian Aboriginal Studies*, 1, 2–17.  
 39 Blurton Jones, N., Hawkes, K., & Draper, P. (1994). Difference between Hadza and !Kung children's  
 40 foraging: Original Affluence or Practical Resason? In: E. Burch (Ed.), *Key Issues in Hunter-  
 Gatherer Research* (pp. 189–215). Oxford: Berg.  
 Blurton Jones, N., & Marlowe, F. (2002). Selection for delayed maturity: Does it take 20 years to learn  
 to hunt and gather? *Human Nature*, 13, 199–238.

- 1 Bock, J. (2002). Learning, life history, and productivity: Children's lives in the Okavango Delta,  
2 Botswana. *Human Nature*, *13*, 161–197.
- 3 Boesch, C., & Boesch, H. (2000). *The chimpanzees of the Tai forest: Behavioral ecology and evolution*.  
4 Oxford: Oxford University Press.
- 5 Bogin, B. (1999). *Patterns of human growth*. Cambridge: Cambridge University Press.
- 6 Bonner, J. (1965). *Size and cycle: An essay in the structure of biology*. Princeton: Princeton University  
7 Press.
- 8 Byrne, R. (1997). Machiavellian intelligence. *Evolutionary Anthropology*, *5*, 172–180.
- 9 Byrne, R., & Whiten, A. (1997). Machiavellian intelligence. In: A. Whiten & R. Byrne (Eds),  
10 *Machiavellian Intelligence II* (pp. 1–23). Cambridge: Cambridge University.
- 11 Charnov, E. (1991). Evolution of life history variation among female mammals. *Proceedings of the*  
12 *National Academy of Sciences*, *88*, 1134.
- 13 Charnov, E. (1993). *Life history invariants: Some explanations of symmetry in evolutionary ecology*.  
14 Oxford: Oxford University Press.
- 15 Charnov, E., & Berrigan, D. (1993). Why do female primates have such long lifespans and so few  
16 babies? Or life in the slow lane. *Evolutionary Anthropology*, *1*, 191–194.
- 17 Charnov, E., & Schaffer, W. (1973). Life-history consequences of natural selection: Cole's result  
18 revisited. *American Naturalist*, *107*, 791–793.
- 19 Dabbs, J., & Dabbs, M. (2000). *Heroes, rogues and lovers*. New York: McGraw-Hill.
- 20 Daly, M., & Wilson, M. (1988). *Homicide*. New York: Aldine de Gruyter.
- 21 Descola, P. (1988). *La selva culta: Simbolismo y praxis en la ecología de los Achuar*. Quito: Ediciones  
22 Abya-Yala.
- 23 Eglund, A. (2002). Pediatrics, dehydration. In: *Emedicine: Instant Access to the Minds of Medicine*.  
24 Address: <http://www.emedicine.com>.
- 25 Flinn, M., Geary, D., & Ward, C. (n.d.). Ecological dominance, social competition, and coalitionary  
26 arms races: Why humans evolved extraordinary intelligence. Manuscript.
- 27 Fritch, R., & McArthur, J. (1974). Menstrual cycles: Fatness as a determinant of minimum weight  
28 necessary for their maintenance and onset. *Science*, *185*, 949–951.
- 29 Geary, D., & Flinn, M. (2001). Evolution of human parental behavior and the human family. *Parent*  
30 *Science and Practice*, *1*, 5–61.
- 31 Green, B., Weiss, J., & Daling, J. (1986). Risk for ovulatory infertility in relation to body weight.  
32 *Fertility and Sterility*, *50*, 721–726.
- 33 Gurven, M., Allen-Arave, W., Hill, K., & Hurtado, A. (2000). It's a wonderful life: Signaling generosity  
34 among the Aché of Paraguay. *Evolution and Human Behavior*, *21*, 263–282.
- 35 Harvey, P., & Clutton-Brock, T. (1985). Life history variation in primates. *Evolution*, *39*, 559–581.
- 36 Harvey, P., & Zammuto, R. (1985). Patterns of mortality and age at first reproduction in natural  
37 populations of mammals. *Nature*, *315*, 318–329.
- 38 Hawkes, K., O'Connell, J., & Blurton Jones, N. (2001). Hunting and nuclear families: Some lessons  
39 from the Hadza about men's work. *Current Anthropology*, *42*, 631–695.
- 40 Hawkes, K., O'Connell, J., Blurton Jones, N., Alvarez, H., & Charnov, E. (1998). Grandmothering,  
menopause, and the evolution of human life histories. *Proceedings of the National Academy of*  
*Sciences*, *95*, 1336–1339.
- Hawkes, K., O'Connell, J., Blurton Jones, N., Alvarez, H., & Charnov, E. (2000). The grandmother  
hypothesis and human evolution. In: L. Cronk, N. Chagnon & W. Irons (Eds), *Human Behavior*  
*and Adaptation: An Anthropological Perspective* (pp. 371–395). New York: Aldine de Gruyter.
- Hewlett, B. (1992). *Father-child relations: Cultural and biosocial contexts*. New York: Aldine de  
Gruyter.

Pl. provide the  
complete  
information for  
ref. "Flinn, M.,  
Geary, D., &  
Ward, C. (n.d.)".

- 1 Hill, K., Boesch, C., Goodall, J., Pusey, A., Williams, J., & Wrangham, R. (2001). Mortality rates  
2 among wild chimpanzees. *Journal of Human Evolution*, 40, 437–450.
- 3 Hill, K., & Hurtado, A. (1996). *Aché life history: The ecology and demography of a foraging people*.  
4 New York: Aldine de Gruyter.
- 5 Hill, K., & Kaplan, H. (1999). Life history traits in humans: Theory and empirical studies. *Annual*  
6 *Review of Anthropology*, 28, 397–430.
- 7 Horn, H. (1978). Optimal tactics of reproduction and life history. In: J. Krebs & N. Davies (Eds),  
8 *Behavioral Ecology: An Evolutionary Approach* (pp. 411–429). Sunderland: Sinauer.
- 9 Howell N. (1979). *Demography of the Dobe !Kung*. New York: Academic.
- 10 Hrdy, S. (1999). *Mother nature: A history of mothers, infants and natural selection*. New York:  
11 Pantheon.
- 12 Hrdy, S. (n.d.). Cooperative breeding: An evolutionary and comparative perspective. In: B. Hewlett  
13 & M. Lamb (Eds), *Hunter-Gatherer Childhoods: Evolutionary, Developmental, and Cultural*  
14 *Perspectives*. New York: Aldine de Gruyter.
- 15 Kaplan, H., Hill, K., Lancaster, J., & Hurtado, A. (2000). A theory of human life history evolution:  
16 Diet, intelligence, and longevity. *Evolutionary Anthropology*, 9, 156–185.
- 17 Kelly, R. (1995). *The foraging spectrum*. Washington, DC: Smithsonian Press.
- 18 Kennedy, G. (2003). Palaeolithic grandmothers? Life history theory and early Homo. *Journal of the*  
19 *Royal Anthropological Institute*, 9, 549–572.
- 20 Kohrs, M., Harper, A., & Kerr, G. (1976). Effects of a low-protein diet during pregnancy of the rhesus  
21 monkey: 1. Reproductive efficiency. *American Journal of Clinical Nutrition*, 29, 136–145.
- 22 Kraemer, S. (2000). The fragile male. *British Medical Journal*, 321, 1609–1612.
- 23 Krakauer, J. (1996). *Into the wild*. New York: Villard Books.
- 24 Lindstedt, S., & Swain, S. (1988). Body size as a constraint of design and function. In: M. Boyce (Ed.),  
25 *Evolution of Life Histories of Mammals* (pp. 93–105). New Haven, CT: Yale University Press.
- 26 MacArthur, R., & Wilson, E. (1967). *The theory of island biogeography*. Princeton, NJ: Princeton  
27 University Press.
- 28 Manocha, S. L., & Long, J. (1977). Experimental protein malnutrition during gestation and breeding  
29 performance of squirrel monkeys. *Primates*, 18, 923–930.
- 30 Marlowe, F. (1999). Showoffs or providers? The parenting effort of Hadza men. *Evolution and Human*  
31 *Behavior*, 20, 391–404.
- 32 Marlowe, F. (2001). Male contribution to diet and female reproductive success among foragers. *Current*  
33 *Anthropology*, 42, 755–763.
- 34 Miller, G. (2000). *The mating mind: How sexual choice shaped the evolution of human nature*. New  
35 York: Doubleday.
- 36 Nishida, T., Takasaki, H., & Takahata, Y. (1990). Demography and reproductive profiles. In: T. Nishida  
37 (Ed.), *The Chimpanzees of the Mahale Mountains* (pp. 3–97). Tokyo: University of Tokyo Press.
- 38 Pagel, M., & Harvey, P. (1993). Evolution of the juvenile period in primates. In: M. Pereira & L.  
39 Fairbanks (Eds), *Juvenile Primates: Life History, Development, and Behavior* (pp. 28–37).  
40 New York: Oxford University Press.
- Peccei, J. (2001). A critique of the grandmother hypotheses: Old and new. *American Journal of Human*  
*Biology*, 13, 434–452.
- Pereira, M. (1993). Juvenility in animals. In: M. Pereira & L. Fairbanks (Eds), *Juvenile Primates: Life*  
*History, Development, and Behavior* (pp. 17–27). New York: Oxford University Press.
- Pereira, M., & Altmann, J. (1985). Development of social behavior in free-living nonhuman primates.  
In: E. Watts (Ed.), *Nonhuman Primate Models for Human Growth and Development* (pp.  
217–309). New York: Liss.

Pl. provide year  
for the ref. "Hrdy,  
S. (n.d.)."

- 1 Prentice, A., Cole, T., Foord, F., Lamb, W., & Whitehead, R. (1987). Increased birthweight after prenatal  
2 dietary supplementation of rural African women. *American Journal of Clinical Nutrition*, 46,  
3 912–925.
- 4 Promislow, D., & Harvey, P. (1990). Living fast and dying young: A comparative analysis of life-history  
5 variation among mammals. *Journal of Zoology*, 220, 417–437.
- 6 Pusey, A. (1990). Behavioral changes at adolescence in chimpanzees. *Behavior*, 115, 203–246.
- 7 Riley, A., Samuelson, J., & Huffman, S. (1993). The relationship of age at menarche and fertility  
8 in undernourished adolescents. In: R. Gray, H. Leridon & A. Spira (Eds), *Biomedical and  
9 Demographic Determinants of Reproduction* (pp. 50–64). Oxford: Clarendon.
- 10 Rose, M. (1983). Theories of life-history evolution. *American Zoologist*, 23, 15–23.
- 11 Schaffer, W. (1974). Selection for optimal life histories: The effects of age structure. *Ecology*, 55,  
12 291–303.
- 13 Schwartz, S., Wilson, M., Walker, M., & Collins, D. (1988). Dietary influences on growth  
14 and sexual maturation in premenarcheal rhesus monkeys. *Hormones and Behavior*, 22,  
15 231–251.
- 16 Steckel, R., Rose, J., Larsen, C., & Walker, P. (2002). Skeletal health in the Western Hemisphere from  
17 4000 BC to the present. *Evolutionary Anthropology*, 11, 142–155.
- 18 Sugiyama, L. (1996). *In search of the adapted mind: A study of human cognitive adaptations among  
19 the Shiwiar of Ecuador and the Yora of Peru*. Ph.D. dissertation. Santa Barbara: University of  
20 California.
- 21 Sugiyama, L. (2000). Cooperative foraging among the Shiwiar. Paper presented at the American  
22 Anthropological Association Annual Meeting, San Francisco.
- 23 Sugiyama, L. (2004). Illness, injury, and disability among Shiwiar forager-horticulturalists:  
24 Implications of health-risk buffering for the evolution of human life history. *American Journal  
25 of Physical Anthropology*, 123, 371–389.
- 26 Sugiyama, L., & Chacon, R. (2000). Effects of illness and injury on foraging among the Yora and  
27 Shiwiar: Pathology risk as adaptive problem. In: L. Cronk, N. Chagnon & W. Irons (Eds),  
28 *Human Behavior and Adaptation: An Anthropological Perspective* (pp. 371–395). New York:  
29 Aldine de Gruyter.
- 30 Sugiyama, L., & Chacon, R. (n.d.). Juvenile responses to household ecology among the Yora of Peruvian  
31 Amazonia. In: B. Hewlett & M. Lamb (Eds), *Hunter-Gatherer Childhoods: Evolutionary,  
32 Developmental, and Cultural Perspectives*. New York: Aldine de Gruyter.
- 33 Sutherland, W., Grafen, W., & Harvey, P. (1986). Life history correlations and demography. *Nature*,  
34 320, 88.
- 35 Tooby, J., & DeVore, I. (1987). The reconstruction of hominid behavioral evolution through strategic  
36 modeling. In: W. Kinzey (Ed.), *The Evolution of Human Behavior: Primate Models* (pp.  
37 183–237). Albany: State University of New York Press.
- 38 Tucker, B., & Young, A. (n.d.). Growing up Mikea: Children's time allocation and tuber foraging in  
39 southwestern Madagascar. In: B. Hewlett & M. Lamb (Eds), *Hunter Gatherer Childhoods:  
40 Evolutionary, Developmental and Cultural perspectives*. New York: Aldine de Gruyter.
- Walker, P., Sugiyama, L., & Chacon, R. (1998). Diet, dental health, and cultural change among  
recently contacted South American Indian hunter-horticulturalists. In: J. Lukacs & B.  
Hemphill (Eds), *Human Dental Development, Morphology and Pathology: Essays in  
Honor of Albert Dahlberg* (pp. 355–386). Eugene: University of Oregon Anthropological  
Papers.
- Walker, R., Hill, K., Kaplan, H., & McMillan, G. (2001). Age-dependency in hunting ability among  
the Aché of eastern Paraguay. *Journal of Human Evolution*, 42, 639–657.

Pl. provide year  
for ref.  
"Sugiyama, L., &  
Chacon, R. (n.d.)".

- 1 Watts, D., & Pusey, A. (1993). Behavior of juvenile and adolescent great apes. In: M. Pereira & L.  
2 Fairbanks (Eds), *Juvenile Primates: Life History, Development, and Behavior* (pp. 148–171).  
3 New York: Oxford University Press.
- 4 Williams, G. (1957). Pleiotropy, natural selection, and the evolution of senescence. *Evolution*, *11*,  
5 298–411.
- 6 Williams, G. (1966). *Adaptation and natural selection*. Princeton, NJ: Princeton University Press.
- 7 Winterhalder, B. (1996). Social foraging and the behavioral ecology of intragroup resource transfers.  
8 *Evolutionary Anthropology*, *5*, 46–57.
- 9 Wood, J., Milner, G., Harpending, H., & Weiss, K. (1992). The osteological paradox: Problems of  
10 inferring prehistoric health from skeletal samples. *Current Anthropology*, *33*, 343–371.

### 11 **Uncited reference**

12  
13  
14 References cited in the text must appear in the reference list; conversely, each  
15 entry in the reference list must be cited in the text . . . The author must make  
16 certain that each source referenced appears in both places and that the text citation  
17 and reference list entry are identical in spelling and year.

18 [Tucker and Young \(n.d.\)](#).

19  
20  
21  
22  
23  
24  
25  
26  
27  
28  
29  
30  
31  
32  
33  
34  
35  
36  
37  
38  
39  
40