



Uniparental genetic markers in South Amerindians

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Abstract

A comprehensive review of uniparental systems in South Amerindians was undertaken. Variability in the Y-chromosome haplogroups were assessed in 68 populations and 1,814 individuals whereas that of Y-STR markers was assessed in 29 populations and 590 subjects. Variability in the mitochondrial DNA (mtDNA) haplogroup was examined in 108 populations and 6,697 persons, and sequencing studies used either the complete mtDNA genome or the highly variable segments 1 and 2. The diversity of the markers made it difficult to establish a general picture of Y-chromosome variability in the populations studied. However, haplogroup Q1a3a* was almost always the most prevalent whereas Q1a3* occurred equally in all regions, which suggested its prevalence among the early colonizers. The STR allele frequencies were used to derive a possible ancient Native American Q-clade chromosome haplotype and five of six STR loci showed significant geographic variation. Geographic and linguistic factors moderately influenced the mtDNA distributions (6% and 7%, respectively) and mtDNA haplogroups A and D correlated positively and negatively, respectively, with latitude. The data analyzed here provide rich material for understanding the biological history of South Amerindians and can serve as a basis for comparative studies involving other types of data, such as cultural data.

Key words: genetics, language and geography, mitochondrial DNA, Native Americans, South Amerindians, Y-chromosome.

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Introduction

Native Americans have been the subject of a large number of population genetic studies because of particular characteristics: (a) there are groups among them that until recently had a hunter-gatherer way of living with only incipient agriculture, typical of our ancestors, (b) they show considerable interpopulation but low intrapopulation variability, and (c) since until recently they could not write there is no written record of their history, except for those of non-Amerindian colonizers. Biological studies can therefore be used to investigate their past.

The first genetic studies examined the variability in blood groups and proteins and have been summarized in Salzano and Callegari-Jacques (1988) and Crawford (1998). The advent of modern molecular biology, which allows direct, detailed DNA analysis, has opened new possibilities for investigating these populations.

DNA studies can basically be divided into two groups: those involving autosomal markers and those involving uniparental (Y-chromosome, mitochondrial DNA) markers. The latter are important because they can provide

a clear-cut pattern of historical events that is not clouded by recombination factors. For Amerindians, the number of reviews that have dealt with these markers is not large or comprehensive. For the Y-chromosome, Bortolini *et al.* (2003) considered 438 individuals from 23 Southern and one Northern Amerindian populations who were screened for eight single nucleotide polymorphisms (SNPs) and six short tandem repeat/microsatellite (STR) loci, and Zegura *et al.* (2004) studied 63 binary polymorphisms and 10 STR regions in 2,344 persons from 15 Northern and three Southern Amerindian groups. Only a few recent studies have used all known SNPs necessary to identify the major Native American Y-haplogroups and their sublineages in Amerindian populations (Geppert *et al.*, 2011; Jota *et al.*, 2011; Bisso-Machado *et al.*, 2011).

The most recent mtDNA reviews were published four years ago and involved sequence variability in the hypervariable region 1 (Hunley *et al.*, 2007; Lewis Jr *et al.*, 2007). Schurr and Sherry (2004), on the other hand, associated data from Y-chromosome markers with mitochondrial DNA (mtDNA) results, providing a good picture of the information available at the time. No general review considering both data sets has been published since then.

This review provides a detailed, comprehensive survey of Y-chromosome haplogroup frequency variation in 68 populations involving 1,814 individuals. In addition,

specific information on Y-STR markers for 29 populations and 590 subjects is given. The haplogroup mtDNA data included 108 populations involving a total of 6,697 persons. Geographic and linguistic factors that may have influenced this variation were carefully considered, leading to a global, overview of the genetic pattern associated with these markers in South Amerindians. Information on mtDNA sequencing studies is also supplied.

Materials and Methods

The data used in this review were obtained from 17 primary surveys of the Y-chromosome and 66 primary surveys of mtDNA. These studies were retrieved through PubMed and by searching the reference lists of the corresponding papers. Haplogroup frequencies were obtained by direct counting. Intra- and inter-population diversity was calculated with AMOVA (Weir and Cockerham, 1984; Excoffier *et al.*, 1992; Weir, 1996) using Arlequin 3.5.1.2 software (Excoffier and Lischer, 2010). AMOVA was also used to estimate the level of differentiation between and within 17 pre-defined language and 7 geographical categories, respectively.

The distribution patterns of the mtDNA haplogroup frequencies were established by generating isoline maps using IDRISI 16.0 software (IDRISI Taiga) (Eastman, 2006). Spearman's correlation coefficients were calculated with PASW Statistics 18 software. Average heterozygosity (ah) was calculated with Arlequin 3.5.1.2 software.

Results and Discussion

Table 1 gives the distribution of the Q and non-Q-chromosomes (defined by a set of SNPs), as well as linguistic and geographical information for the samples considered. The samples were distributed from latitude 11° North to 45° South and longitude 46° to 76° West, with the individuals involved speaking 23 languages. Sample sizes varied widely from 1 to 151 individuals. Twenty-two of the studies involved less than 10 persons. Unfortunately, there is no standardization on the number of SNPs studied and in most cases only the M242 and M3 markers (which define the Asian/Native American paragroup Q* and its autochthonous Native American sublineage Q1a3a*, respec-

Table 1 - The distribution of Q and non-Q lineages and linguistic and geographical information for the samples considered.

Populations (n) ¹	Haplogroup (%)		Language ²	Geographical coordinates	References
	Q lineages / Amerindian origin	Non-Q lineages			
Wayuu (19)	69	31	Arawakan	11° N; 73° W	Bortolini <i>et al.</i> (2003)
Kogi (17)	100		Chibchan	11° N; 74° W	Rojas <i>et al.</i> (2010)
Barira (12)	100		Chibchan	10° 44' N; 71° 23' W	Bortolini <i>et al.</i> (2003)
Arsario (Wiwa) (6)	100		Chibchan	10° 25' N; 73° 05' W	Rojas <i>et al.</i> (2010)
Arhuaco (Ijka) (19)	100		Chibchan	9° 04' N; 73° 59' W	Rojas <i>et al.</i> (2010)
Warao (12)	100		Warao	9° N; 61° W	Bortolini <i>et al.</i> (2003)
Yukpa (12)	100		Carib	8° 40' N; 72° 41' W	Bortolini <i>et al.</i> (2003)
Zenu (52)	79	21	Spanish ³	8° 30' N; 76° W	Bortolini <i>et al.</i> (2003); Rojas <i>et al.</i> (2010)
Embera (13)	92	8	Choco	7° N; 76° 30' W	Rojas <i>et al.</i> (2010)
Makiritare (25)	68	32	Carib	5° 33' N; 65° 33' W	Lell <i>et al.</i> (2002)
Kali'na (21)	81	19	Carib	5° 31' N; 53° 47' W	Mazières <i>et al.</i> (2008)
Waunana (29)	100		Choco	4° 50' N; 77° W	Rojas <i>et al.</i> (2010)
Palikur (35)	94	6	Arawakan	4° N; 51° 45' W	Mazières <i>et al.</i> (2008)
Macushi (4)	100		Carib	4° N; 60° 50' W	Lell <i>et al.</i> (2002)
Piaroa (6)	100		Salivan	3° 57' N; 66° 22' W	Lell <i>et al.</i> (2002)
Wapishana (2)	50	50	Arawakan	3° 07' N; 60° 03' W	Lell <i>et al.</i> (2002)
Emerillon (9)	100		Tupi	3° N; 53° W	Mazières <i>et al.</i> (2008)
Yanomámi (39)	38	62	Yanomam	2° 50' N; 54° W	Rodriguez-Delfin <i>et al.</i> (1997); Lell <i>et al.</i> (2002)
Tiryiό (4)	100		Carib	2° N; 56° W	Bortolini <i>et al.</i> (2003)
Apalaί (57)	98	2	Carib	1° 20' N; 54° 40' W	Rodriguez-Delfin <i>et al.</i> (1997); Bortolini <i>et al.</i> (2003)
Wayampi (62)	100		Tupi	1° N; 53° W	Rodriguez-Delfin <i>et al.</i> (1997); Bortolini <i>et al.</i> (2003); Mazières <i>et al.</i> (2008)

Table 1 (cont).

Populations (n) ¹	Haplogroup (%)		Language ²	Geographical coordinates	References
	Q lineages / Amerindian origin	Non-Q lineages			
Yagua (7)	100		Peba-Yaguan	0° 51' N; 72° 27' W	Bortolini <i>et al.</i> (2003)
Ingano (108)	80	20	Quechuan	0° 50' N; 77° W	Bortolini <i>et al.</i> (2003); Rojas <i>et al.</i> (2010)
Wai-Wai (9)	100		Carib	0° 40' S; 58° W	Bisso-Machado <i>et al.</i> (2011)
Urubu-Kaapor (16)	100		Tupi	2° -3° S; 46° -47° W	Bortolini <i>et al.</i> (2003)
Huitoto (4)	75	25	Witotoan	2° 14' S; 72° 19' W	Bortolini <i>et al.</i> (2003)
Arara (15)	100		Carib	3° 30' -4° 20' S; 53° 0' -54° 10' W	Rodriguez-Delfin <i>et al.</i> (1997); Bianchi <i>et al.</i> (1998); Bisso-Machado <i>et al.</i> (2011)
Asurini (4)	100		Tupi	3° 35' -4° 12' S; 49° 40' -52° 26' W	Bortolini <i>et al.</i> (2003)
Ticuna (59)	93	7	Ticuna	4° S; 69° 58' W;	Bortolini <i>et al.</i> (2003); Rojas <i>et al.</i> (2010)
Parakanã (20)	100		Tupi	5° 22' S; 51° 17' W	Bortolini <i>et al.</i> (2003)
Xikrin (14)	100		Macro-Ge	5° 55' S; 51° W	Bortolini <i>et al.</i> (2003); Bisso-Machado <i>et al.</i> (2011)
Suruí (24)	96	4	Tupi	5° 58' -10° 50' S; 48° 39' -61° 10' W	Underhill <i>et al.</i> (1996); Bisso-Machado <i>et al.</i> (2011)
Araweté (4)	100		Tupi	5° 9' S; 52° 22' W	Bisso-Machado <i>et al.</i> (2011)
Munduruku (1)	100		Tupi	6° 23' S; 59° 9' W	Bisso-Machado <i>et al.</i> (2011)
Jamamadi (3)	100		Arauan	7° 15' S; 66° 41' W	Bisso-Machado <i>et al.</i> (2011)
Gorotire (19)	100		Macro-Ge	7° 44' S; 51° 10' W	Bortolini <i>et al.</i> (2003); Bisso-Machado <i>et al.</i> (2011)
Krahó (15)	93	7	Macro-Ge	8° S; 47° 15' W	Lell <i>et al.</i> (2002); Bortolini <i>et al.</i> (2003)
Kuben-Kran-Kegn (9)	100		Macro-Ge	8° 10' S; 52° 8' W	Bisso-Machado <i>et al.</i> (2011)
Tenharim (1)	100		Tupi	8° 20' S; 62° W	Bisso-Machado <i>et al.</i> (2011)
Mekranoti (9)	78	22	Macro-Ge	8° 40' S; 54° W	Bortolini <i>et al.</i> (2003)
Kayapó (10)	100		Macro-Ge	9° S; 53° W	Rodriguez-Delfin <i>et al.</i> (1997)
Karitiana (18)	100		Tupi	9° 30' S; 64° 15' W	Underhill <i>et al.</i> (1996); Bisso-Machado <i>et al.</i> (2011)
Cinta-Larga (15)	100		Tupi	9° 50' -12° 30' S; 59° 10' -60° 50' W	Bortolini <i>et al.</i> (2003)
Gavião (7)	100		Tupi	10° 10' S; 61° 8' W	Bisso-Machado <i>et al.</i> (2011)
Karipuna (1)		100	Tupi	10° 14' S; 64° 13' W	Bisso-Machado <i>et al.</i> (2011)
Zoró (6)	100		Tupi	10° 20' S; 60° 20' W	Bisso-Machado <i>et al.</i> (2011)
Matsiguenga (28)	91	9	Arawakan	10° 47' -12° 51' S; 73° 17' -70° 44' W	Mazières <i>et al.</i> (2008)
Pacaás Novos (Wari) (29)	100		Chapacura-Wanham	11° 8' S; 65° W	Bortolini <i>et al.</i> (2003)
Panoa (5)	100		Pano	12° 55' S; 65° 12' W	Lell <i>et al.</i> (2002)
Xavante (15)	100		Macro-Ge	14° S; 52° 30' W	Bisso-Machado <i>et al.</i> (2011)
Quechua (44)	73	27	Quechuan	14° 30' S; 69° W	Gayà-Vidal <i>et al.</i> (2011)
Aymara (59)	97	3	Aymaran	17° 68' S; 69° 16' W	Gayà-Vidal <i>et al.</i> (2011)
Ayoreo (9)	78	22	Zamucoan	19° S; 60° 30' W	Bailliet <i>et al.</i> (2009)
Wichí (Mataco) (151)	48	52	Mataco-Guaicuru	22° 28' S; 62° 70' W	Demarchi and Mitchell (2004); Bailliet <i>et al.</i> (2009)
Lengua (36)	97	3	Mascoian	22° 45' S; 58° 5' W	Bailliet <i>et al.</i> (2009); Bisso-Machado <i>et al.</i> (2011)
Chorote (9)	89	21	Mataco-Guaicuru	22° 90' S; 65° 40' W;	Bailliet <i>et al.</i> (2009)
Aché (54)	98	2	Tupi	23° 30' -24° 10' S; 55° 50' -56° 30' W	Bortolini <i>et al.</i> (2003)

Table 1 (cont).

Populations (n) ¹	Haplogroup (%)		Language ²	Geographical coordinates	References
	Q lineages / Amerindian origin	Non-Q lineages			
Guarani (78)	77	23	Tupi	23° 6' S; 55° 12' W	Bortolini <i>et al.</i> (2003); Marrero <i>et al.</i> (2007)
Pilagá	47	53	Mataco-Guaicuru	24° S; 59° W	Demarchi and Mitchell (2004)
Colla (63)	35	65	Quechuan ³	24° 10' -24° 43' S; 65° 17' -65° 52' W	Blanco-Verea <i>et al.</i> (2010); Toscanini <i>et al.</i> (2011)
Toba (89)	88	12	Mataco-Guaicuru	26° S; 58° W	Demarchi and Mitchell (2004); Bailliet <i>et al.</i> (2009); Toscanini <i>et al.</i> (2011)
Kaingang (59)	69	31	Macro-Ge	28° S; 51° 20' W	Bortolini <i>et al.</i> (2003); Marrero <i>et al.</i> (2007); Bisso-Machado <i>et al.</i> (2011)
Diaguita (24)	37	63	Quechuan ⁴	28° 20' S; 67° 43' W	Blanco-Verea <i>et al.</i> (2010)
Mocoví (40)	60	40	Mataco-Guaicuru	29° 51' S; 59° 56' W	Bailliet <i>et al.</i> (2009)
Pehuenche (18)	83	17	Araucanian	37° 43' S; 71° 16' W	Bailliet <i>et al.</i> (2009)
Mapuche (105)	36	64	Araucanian	39° 10' -41° 20' S; 68° 37' -70° 22' W	Bailliet <i>et al.</i> (2009); Blanco-Verea <i>et al.</i> (2010)
Huilliche (26)	50	50	Araucanian	41° 16' S; 73° W	Bailliet <i>et al.</i> (2009)
Tehuelche (20)	65	35	Chon	45° S; 71° W	Bailliet <i>et al.</i> (2009)

¹Arranged according to latitude. ²Classification according to Lewis (2009). ³Original language is extinct. ⁴The Diaguita spoke originally Kakán, but this language became extinct and was substituted by Quechua.

tively; Pena *et al.*, 1995; Bortolini *et al.*, 2003; Seielstad *et al.*, 2003) were investigated. This fact precludes a complete, precise view of the distribution of Q1a3a sublineages and other Q clade chromosomes in South America. For this reason, the information in Table 1 was limited to the frequencies of the Q and non-Q-lineages only. Note that non-Q-chromosomes (which, for the reasons given above, could not be identified in sublineages) were identified in ~50% of the tribal groups. For some of these populations admixture with non-Indians is known and could be the source of these non-Q chromosomes (for example, Mapuche and Guarani; Marrero *et al.*, 2007; Bailliet *et al.*, 2009; Blanco-Verea *et al.*, 2010). Overall, the numbers presented in Table 1 indicate a higher presence of non-Q lineages in southern populations than in those of the northern/Amazonian region, probably because of greater admixture with non-Indians in the former than in the latter. However, for some isolated groups such as the Yanomámi, it is unlikely that admixture explains the findings. In these cases other causes are more probable, such as the presence of unknown autochthonous lineages and/or known Q lineages whose defining markers were not tested.

Despite the great variation in the number of Y-SNPs used in these studies, Figure 1 illustrates some of the trends that were observed: The autochthonous Native American Q1a3a* is almost always the most prevalent, whereas its sublineages (Q1a3a1, Q1a3a2, Q1a3a3 and Q1a3a4) seem to have more restricted geographical distributions. The second most prevalent, Q1a3*, appears to occur equally in all regions, suggesting its presence among the first settlers of South America. The other known Q clade chromosomes

(Q1*, Q1a*, Q1a1, Q1a2, Q1a4, Q1a5, Q1a6 and Q1b) have not yet been identified in South America. Only one non-Q-chromosome (C3*) of probable native origin has been described in northwest South Amerindian populations (Figure 1; Geppert *et al.*, 2011).

The nature of some evolutionary and demographic scenarios, mediated by men, in native American populations has also been evaluated by using Y microsatellite markers (Y-STRs), which have a much faster evolutionary rate than SNPs. Y-STRs allow the retrieval of population and chromosome evolutionary histories. For example, STR data have been used to estimate that the mutations that gave rise to the Q1a3a1 and Q1a3a4 sublineages occurred $7,972 \pm 2,916$ and $5,280 \pm 1,330$ years ago, probably in northwest South America and the Andean region, respectively (Bortolini *et al.*, 2003; Jota *et al.*, 2011).

Table 2 shows the STR allele frequencies observed in 29 South Amerindian populations, based only on Q clade chromosomes. In this compilation, we considered only studies containing information on the allele frequencies for each population individually. There was considerable variation in the number of samples tested in each study, the number of tribes, and the number of individuals per tribe. Depending on the locus considered, the number of alleles observed ranged from one to eight, with some of them appearing in only one study while others were present in almost all populations. Based on the most prevalent alleles per locus we reconstructed a probable haplotype of the ancient Native American Q-clade chromosome (ANAQC) as: 13(DYS19)-12(DYS388)-14(DYS389I)-31(DYS389II)-2

Table 2 (a) (cont.)

STR (allele)	Ref. (n)														
	Aché (48) ²	Apalai (9) ¹	Arara (8) ¹	Aymara (57) ⁹	Ayoreo (2) ⁷	Barira (12) ²	Diaguita (9) ⁸	Guarani (47) ^{4,5}	Ingano (8) ²	Kaingang (17) ³	Kayapó (10) ¹	Colla (22) ^{6,8}	Lengua (6) ⁷	Mapuche (24) ^{6,8}	Mekranoti (5) ²
DYS390 (20)															
DYS390 (21)		0.230	0.120	0.070						0.060	0.100	0.040			
DYS390 (22)		0.110						0.130						0.040	
DYS390 (23)	1.000	0.330	0.880	0.560		0.080	0.330	0.080	0.250		0.100	0.640	0.330	0.420	0.400
DYS390 (24)		0.330		0.250	1.000	0.920	0.450	0.680	0.370	0.940	0.700	0.180	0.670	0.330	0.600
DYS390 (25)				0.120			0.220	0.090	0.250		0.100			0.210	
DYS390 (26)								0.020	0.130			0.140			
DYS390 (27)															
DYS390 total															
DYS391 (9)				0.020										0.210	
DYS391 (10)	1.000			0.820	1.000	1.000	0.890	0.210	0.860			0.820	1.000	0.750	1.000
DYS391 (11)				0.140			0.110	0.790	0.140			0.180		0.040	
DYS391 (12)				0.020											
DYS391 total															
DYS392 (11)										0.060					
DYS392 (12)												0.140			
DYS392 (13)		1.000	1.000			0.920		0.070	0.120	0.350	0.700	0.050		0.170	1.000
DYS392 (14)	1.000			0.470	0.500	0.080	0.450	0.720	0.880	0.530	0.300	0.360	0.830	0.710	
DYS392 (15)				0.040	0.500		0.330	0.210		0.060		0.090	0.170	0.120	
DYS392 (16)				0.470			0.220					0.360			
DYS392 (17)				0.020											
DYS392 (18)															
DYS392 total															
DYS393 (11)								0.480							
DYS393 (12)		0.220	0.120			0.080		0.090	0.120	0.060	0.400	0.090			0.400
DYS393 (13)	1.000	0.780	0.880	0.440	1.000		0.670	0.340	0.760	0.590	0.600	0.550	1.000	1.000	0.600
DYS393 (14)				0.560		0.920	0.330	0.090	0.120	0.350		0.360			
DYS393 (15)															
DYS393 (16)															
DYS393 total															
DYS437 (8)															
DYS437 (9)															
DYS437 (11)														0.040	
DYS437 (14)				1.000			1.000	0.930		0.590		1.000		0.920	
DYS437 (15)								0.070		0.410				0.040	
DYS437 total															
DYS438 (9)															
DYS438 (10)				0.050			0.220							0.040	
DYS438 (11)				0.930			0.670	0.860		0.590		1.000		0.920	
DYS438 (12)				0.020			0.110	0.140		0.410					
DYS438 (16)														0.040	
DYS438 total															
DYS439 (9)												0.050			
DYS439 (10)										0.060				0.130	
DYS439 (11)				0.160			0.220	0.070		0.230		0.050		0.220	
DYS439 (12)				0.240			0.450	0.640		0.590		0.360		0.390	
DYS439 (13)				0.370			0.330	0.290		0.060		0.270		0.260	

Table 2 (b) (cont.)

STR (allele)	Ref. (n)														Total (590)
	Mocoví (2) ⁸	Pacaás Novos (15) ²	Parakanã (4) ²	Pilagá (9) ³	Quechua (58) ^{10,11}	Ticuna (36) ²	Toba (70) ^{3,7}	Warao (12) ²	Wayampi (10) ¹	Wayuu (14) ²	Wichi (27) ³	Yanomama (9) ¹	Yukpa (12) ²	Zenu (28) ²	
DYS437 (8)				0.420			0.360				1.000				0.200
DYS437 (9)				0.580			0.030								0.040
DYS437 (11)															0.010
DYS437 (14)					1.000		0.500								0.700
DYS437 (15)							0.110								0.050
DYS437 total															322
DYS438 (9)					0.020										0.010
DYS438 (10)							0.050				0.160				0.050
DYS438 (11)				0.790	0.930		0.850				0.840				0.850
DYS438 (12)				0.210	0.050		0.100								0.080
DYS438 (16)															0.010
DYS438 total															322
DYS439 (9)															0.010
DYS439 (10)															0.010
DYS439 (11)					0.090		0.130				0.230				0.140
DYS439 (12)				0.320	0.240		0.500				0.710				0.400
DYS439 (13)				0.680	0.500		0.290				0.060				0.330
DYS439 (14)					0.170		0.080								0.110
DYS439 total															322
DYS448 (18)															0.010
DYS448 (19)					0.140		0.020								0.090
DYS448 (20)					0.550		0.930								0.740
DYS448 (21)					0.240		0.050								0.140
DYS448 (22)					0.070										0.020
DYS448 total															169
DYS456 (11)															0.010
DYS456 (13)															0.020
DYS456 (14)					0.230		0.020								0.090
DYS456 (15)					0.720		0.730								0.750
DYS456 (16)					0.050		0.250								0.120
DYS456 (17)															0.010
DYS456 total															169
DYS458 (13)															0.020
DYS458 (15)					0.030		0.040								0.040
DYS458 (16)					0.360		0.180								0.380
DYS458 (17)					0.520		0.550								0.410
DYS458 (18)					0.090		0.230								0.140
DYS458 (19)															0.010
DYS458 total															169
DYS635 (22)					0.950		0.950								0.890
DYS635 (23)					0.020		0.050								0.090
DYS635 (24)					0.030										0.010
DYS635 (26)															0.010
DYS635 total															169

¹Rodriguez-Delfin *et al.* (1997); ²Bortolini *et al.* (2003); ³Demarchi and Mitchell (2004); ⁴Altuna *et al.* (2006); ⁵Leite *et al.* (2008); ⁶Toscanini *et al.* (2008); ⁷Bailliet *et al.* (2009); ⁸Blanco-Verea *et al.* (2010); ⁹Gayà-Vidal *et al.* (2011); ¹⁰Jota *et al.* (2011).

4(DYS390)-10(DYS391)-14(DYS392)-13(DYS393)-14(DYS437)-11(DYS438)-12(DYS439)-20(DYS448)-15(DYS456)-16(DYS458)-22(DYS635). Using this information and additional data for these loci (except DYS388) reported in the Y Chromosome Haplotype Reference Database we found no matches in 36,448 haplotypes (245 populations). Although we found no complete identity with our estimated ANAQC, three one-step neighbor haplotypes were encountered, two in individuals with an admixed ancestry living in Latin American countries and one in a Native American individual (Kaqchiquel).

Table 3 shows the results of the molecular analysis of variance for populations structured by language or geography based on the data in Table 2. The estimates were calculated for each STR locus because testing heterogeneity prevented haplotype identification. As expected, most of the diversity was attributable to intrapopulation variation, with one exception (DYS437) that was explained by the fixation of allele 14 in 40% of the populations, whereas only allele 8 was found in the Wichí. In contrast, significant variation among subdivisions was detected for only six loci (DYS398I, DYS391, DYS392, DYS393, DYS437 and DYS456) and in five out of these six it was attributable to geography. There was also considerable inter-population/within subdivision variability (significant in 28 of 30 evaluations), with the average percentage being 16% for geography and 21% for language.

Table 4 summarizes the information on sequencing studies of mitochondrial DNA. The mtDNA genome of representative individuals from 35 populations has been entirely sequenced, as reported in six publications (Ingman *et al.*, 2000; Kivisild *et al.*, 2006; Tamm *et al.*, 2007; Fagundes *et al.*, 2008; Perego *et al.*, 2009, 2010). However, the analyses performed did not consider the within South Amerindian relationships and were mostly concerned with interethnic or interhaplogroup comparisons. Based on 86 complete Amerindian genomes, Fagundes *et al.* (2008) concluded that the prehistoric colonization of the Americas involved a single founding population, with an initial differentiation from Asia occurring in Beringia that ended around 19,000-23,000 years ago, with a moderate bottleneck. Expansion into the New World would have occurred about 18,000 years ago. An extensive 5.76 kb analysis by Dornelles *et al.* (2005) established that haplogroup X is not present in extant South American Indians.

The most extensive set of data involves the highly variable segment I (HVS-I) that has been studied in 92 populations and reported in 30 papers; surveys that have included the HVS-II region are much less common (10 articles) (Table 4). For HVS-I, Merriwether *et al.* (2000) provided an excellent example of how intrapopulation variability in the Yanomámi could be interpreted in a historical and demographical context and relating it to other Amerindian and Asian data. They studied 129 Yanomámi sequences from individuals in eight villages and compared

Table 3 - Analysis of molecular variance of the distinct alleles of the Y-Q STRs in relation to the language and geography of the populations tested.

STR loci (structured by)	Among subdivisions	Among populations within subdivisions	Within populations
DYS19 (Language) ¹	0	0.311*	0.689*
DYS19 (Geography) ²	0	0.332*	0.668*
DYS388 (Language) ¹	0.045	0.317*	0.638*
DYS388 (Geography) ²	0.091	0.240*	0.669*
DYS389I (Language) ¹	0.227*	0.030	0.743*
DYS389I (Geography) ²	0.116	0.148*	0.736*
DYS389II (Language) ¹	0.036	0.077*	0.887*
DYS389II (Geography) ²	0	0.125*	0.875*
DYS390 (Language) ¹	0.008	0.326*	0.666*
DYS390 (Geography) ²	0	0.359*	0.642*
DYS391 (Language) ¹	0	0.385*	0.615*
DYS391 (Geography) ²	0.253*	0.131*	0.616*
DYS392 (Language) ¹	0.005	0.319*	0.676*
DYS392 (Geography) ²	0.007*	0.262*	0.661*
DYS393 (Language) ¹	0	0.331*	0.669*
DYS393 (Geography) ²	0.177*	0.167*	0.656*
DYS437 (Language) ¹	0.289	0.273*	0.438*
DYS437 (Geography) ²	0.390*	0.213*	0.397*
DYS438 (Language) ¹	0.048	0.025	0.927*
DYS438 (Geography) ²	0.023	0.054*	0.923*
DYS439 (Language) ¹	0.019	0.054*	0.929*
DYS439 (Geography) ²	0.055	0.034*	0.911*
DYS448 (Language) ¹	0	0.148*	0.852*
DYS448 (Geography) ²	0	0.073*	0.927*
DYS456 (Language) ¹	0	0.078*	0.922*
DYS456 (Geography) ²	0.017*	0.044*	0.939*
DYS458 (Language) ¹	0	0.144*	0.856*
DYS458 (Geography) ²	0.043	0.069*	0.888*
DYS635 (Language) ¹	0	0.278*	0.722*
DYS635 (Geography) ²	0	0.128*	0.872*

¹Language: Tupi: Aché, Guarani, Parakanã, Wayampi; Carib: Apalaí, Arara, Yukpa; Macro-Ge: Kaingang, Kayapó, Mekranoti; Quechua: Diaguita, Quechua, Ingano; Mataco-Guaicuru: Mocoí, Toba, Wichí, Pilagá; Isolated languages and others with only one population were included as a sixth group.

²Geography: Amazonia/Central Brazilian Plateau: Apalaí, Arara, Kayapó, Mekranoti, Pacaás Novos, Parakanã, Ticuna, Warao, Wayampi, Yanomámi; Southern Brazil: Guarani and Kaingang; Chaco: Ache, Ayoreo, Lengua, Mocoí, Pilagá, Toba, Wichí; Andes: Aymara, Barira, Diaguita, Kolla, Quechua, Wayuu, Yukpa, Zenu.

*Significant values ($p = 0.05$). Negative values were adjusted to zero.

their haplotypes with those of other Asian and New World populations, in a total of 482 unique haplotypes. Interestingly, the pairwise inter-population gene flow estimates were lower between some pairs of Yanomámi villages than between them and four other South Amerindian groups.

With regard to intrapopulation variability, as measured by Θ_k , Fuselli *et al.* (2003) and Corella *et al.* (2007) reported extensive variation for 14 and 27 Central and

Table 4 - Mitochondrial DNA sequencing studies in South Amerindian populations.

Population	HVS-I	HVS-II	Complete mt genome	References
Aché	X	X	X	Schmitt <i>et al.</i> (2004); Dornelles <i>et al.</i> (2005); Fagundes <i>et al.</i> (2008); Yang <i>et al.</i> (2010)
Ancash	X		X	Lewis Jr <i>et al.</i> (2005); Perego <i>et al.</i> (2009)
Andean	X			García-Bour <i>et al.</i> (2004) ¹
Apalaí	X			Lobato-da-Silva <i>et al.</i> (2001); Mazières <i>et al.</i> (2008)
Arara	X		X	Santos <i>et al.</i> (1996); Lobato-da-Silva <i>et al.</i> (2001); Ribeiro-dos-Santos <i>et al.</i> (2001); Silva Jr <i>et al.</i> (2002, 2003) ² ; Fagundes <i>et al.</i> (2008)
Araweté	X			Lobato-da-Silva <i>et al.</i> (2001)
Arequipa	X		X	Fuselli <i>et al.</i> (2003); Perego <i>et al.</i> (2009)
Arhuaco	X	X	X	Melton <i>et al.</i> (2007); Tamm <i>et al.</i> (2007); Yang <i>et al.</i> (2010)
Arsario	X		X	Melton <i>et al.</i> (2007); Tamm <i>et al.</i> (2007)
Asurini	X	X		Lobato-da-Silva <i>et al.</i> (2001); Dornelles <i>et al.</i> (2005)
Auca			X	Kivisild <i>et al.</i> (2006)
Awa-Guajá	X			Santos <i>et al.</i> (1996); Lobato-da-Silva <i>et al.</i> (2001)
Awa-Juriti	X			Lobato-da-Silva <i>et al.</i> (2001)
Aymara	X	X		Corella <i>et al.</i> (2007); Lewis Jr <i>et al.</i> (2007); Yang <i>et al.</i> (2010); Barbieri <i>et al.</i> (2011)
Ayoreo	X	X		Dornelles <i>et al.</i> (2004, 2005)
Catamarca			X	Tamm <i>et al.</i> (2007)
Cayapa	X	X	X	Rickards <i>et al.</i> (1999); Tamm <i>et al.</i> (2007)
Chimane	X			Corella <i>et al.</i> (2007)
Chilean North Coast	X			Moraga <i>et al.</i> (2005) ¹
Cinta Larga	X	X		Lobato-da-Silva <i>et al.</i> (2001); Dornelles <i>et al.</i> (2005)
Coreguaje			X	Tamm <i>et al.</i> (2007)
Coya	X	X		Alvarez-Iglesias <i>et al.</i> (2007)
Cubeo	X	X		Torres <i>et al.</i> (2006)
Curripaco	X	X		Torres <i>et al.</i> (2006)
Desano	X	X		Torres <i>et al.</i> (2006)
Diaguíta			X	Perego <i>et al.</i> (2010)
Embera	X	X	X	Torres <i>et al.</i> (2006); Tamm <i>et al.</i> (2007)
Emerillon	X			Mazières <i>et al.</i> (2008)
Gavião	X		X	Ward <i>et al.</i> (1996); Fagundes <i>et al.</i> (2008)
Gorotire	X	X		Dornelles <i>et al.</i> (2005)
Guahibo	X	X		Vona <i>et al.</i> (2005); Torres <i>et al.</i> (2006)
Guarani	X	X	X	Ingman <i>et al.</i> (2000); Silva Jr <i>et al.</i> (2002, 2003) ² ; Dornelles <i>et al.</i> (2005); Kivisild <i>et al.</i> (2006); Marrero <i>et al.</i> (2007); Fagundes <i>et al.</i> (2008); Sala <i>et al.</i> (2010); Yang <i>et al.</i> (2010)
Huilliche	X	X		Yang <i>et al.</i> (2010)
Huitoto	X	X		Monsalve <i>et al.</i> (1994); Torres <i>et al.</i> (2006)
Içana River Indians	X	X		Dornelles <i>et al.</i> (2005)
Ignaciano	X			Bert <i>et al.</i> (2004)
Inga	X	X		Torres <i>et al.</i> (2006); Yang <i>et al.</i> (2010)
Jaqaru	X			Lewis Jr <i>et al.</i> (2007)
Jamamadi	X			Lobato-da-Silva <i>et al.</i> (2001)
Jebero	X	X		Monsalve <i>et al.</i> (1994); Torres <i>et al.</i> (2006)
Kaingang	X	X		Dornelles <i>et al.</i> (2005); Marrero <i>et al.</i> (2007); Yang <i>et al.</i> (2010)
Kali'na	X			Mazières <i>et al.</i> (2008)
Karitiana	X			Lobato-da-Silva <i>et al.</i> (2001);

Table 4 (cont.)

Population	HVS-I	HVS-II	Complete mt genome	References
Katueña	X		X	Santos <i>et al.</i> (1996); Lobato-da-Silva <i>et al.</i> (2001); Silva Jr <i>et al.</i> (2002, 2003) ² ; Fagundes <i>et al.</i> (2008)
Kayapó	X			Lobato-da-Silva <i>et al.</i> (2001); Silva Jr <i>et al.</i> (2002, 2003) ²
Kawésqar	X			García-Bour <i>et al.</i> (2004) ¹
Kogi	X	X	X	Melton <i>et al.</i> (2007); Tamm <i>et al.</i> (2007); Yang <i>et al.</i> (2010)
Kolla			X	Perego <i>et al.</i> (2010)
Krahô	X	X		Dornelles <i>et al.</i> (2005)
Kikretun	X		X	Santos <i>et al.</i> (1996); Fagundes <i>et al.</i> (2008)
Kubenkoqre	X		X	Santos <i>et al.</i> (1996); Fagundes <i>et al.</i> (2008)
Kuben-Kran-Kegn	X	X		Dornelles <i>et al.</i> (2005)
Lengua	X	X		Dornelles <i>et al.</i> (2005)
Mapuche	X	X		Ginther <i>et al.</i> (1993); Moraga <i>et al.</i> (2000); Dornelles <i>et al.</i> (2005)
Mekranoti	X	X		Dornelles <i>et al.</i> (2005)
Mocovi			X	Tamm <i>et al.</i> (2007)
Movima	X			Bert <i>et al.</i> (2004); Melton <i>et al.</i> (2007)
Munduruku	X			Lobato-da-Silva <i>et al.</i> (2001);
Mura	X	X		Dornelles <i>et al.</i> (2005)
Ocaína	X	X		Monsalve <i>et al.</i> (1994); Torres <i>et al.</i> (2006)
Pacaás Novos	X	X		Dornelles <i>et al.</i> (2005)
Paez	X	X		Torres <i>et al.</i> (2006)
Palikur	X			Lobato-da-Silva <i>et al.</i> (2001); Mazières <i>et al.</i> (2008)
Parakanã	X	X		Lobato-da-Silva <i>et al.</i> (2001); Dornelles <i>et al.</i> (2005)
Pehuenche	X	X		Merriwether <i>et al.</i> (1994, 1995); Merriwether and Ferrell (1996); Moraga <i>et al.</i> (1997, 2000); García <i>et al.</i> (2006)
Peruvian Andes	X	X		Shinoda <i>et al.</i> (2006) ¹ ; Fehren-Schmitz <i>et al.</i> (2011) ¹
Peruvian Southern Coast	X			Fehren-Schmitz <i>et al.</i> (2010) ¹
Piapoco	X	X		Torres <i>et al.</i> (2006)
Pilagá	X			Cabana <i>et al.</i> (2006)
Poturuqara	X		X	Santos <i>et al.</i> (1996); Lobato-da-Silva <i>et al.</i> (2001); Silva Jr <i>et al.</i> (2002, 2003) ² ; Fagundes <i>et al.</i> (2008)
Puinave	X	X		Torres <i>et al.</i> (2006)
Puno	X			Lewis Jr <i>et al.</i> (2007)
Quechua	X	X	X	Monsalve <i>et al.</i> (1994); Silva Jr <i>et al.</i> (2002, 2003) ² ; Fuselli <i>et al.</i> (2003); Dornelles <i>et al.</i> (2005); Kivisild <i>et al.</i> (2006); Corella <i>et al.</i> (2007); Lewis Jr <i>et al.</i> (2007); Fagundes <i>et al.</i> (2008); Yang <i>et al.</i> (2010); Barbieri <i>et al.</i> (2011)
Saliva	X	X		Torres <i>et al.</i> (2006)
Salta			X	Tamm <i>et al.</i> (2007)
Sateré Mawé	X	X		Dornelles <i>et al.</i> (2005)
Selknam	X			García-Bour <i>et al.</i> (2004) ¹
Sicán	X			Shimada <i>et al.</i> (2004) ¹
Suruí	X		X	Lobato-da-Silva <i>et al.</i> (2001); Fagundes <i>et al.</i> (2008)
Tayacaja	X			Bert <i>et al.</i> (2004)
Ticuna	X	X		Monsalve <i>et al.</i> (1994); Torres <i>et al.</i> (2006); Rojas <i>et al.</i> (2010); Yang <i>et al.</i> (2010)
Tiriyó	X	X	X	Santos <i>et al.</i> (1996); Lobato-da-Silva <i>et al.</i> (2001); Silva Jr <i>et al.</i> (2002, 2003) ² ; Dornelles <i>et al.</i> (2005); Fagundes <i>et al.</i> (2008)
Toba	X			Cabana <i>et al.</i> (2006)

Table 4 (cont.)

Population	HVS-I	HVS-II	Complete mt genome	References
Trinitario	X			Bert <i>et al.</i> (2004)
Tucuman			X	Tamm <i>et al.</i> (2007)
Tupe	X			Lewis Jr <i>et al.</i> (2007)
Txukahamãe	X	X		Dornelles <i>et al.</i> (2005)
Uro	X			Barbieri <i>et al.</i> (2011)
Urubu Kaapor	X	X		Lobato-da-Silva <i>et al.</i> (2001); Dornelles <i>et al.</i> (2005)
Vaupe			X	Tamm <i>et al.</i> (2007)
Wai-wai			X	Fagundes <i>et al.</i> (2008)
Wayampi	X	X	X	Santos <i>et al.</i> (1996); Lobato-da-Silva <i>et al.</i> (2001); Silva Jr <i>et al.</i> (2002, 2003) ² ; Dornelles <i>et al.</i> (2005); Fagundes <i>et al.</i> (2008); Mazières <i>et al.</i> (2008)
Warao			X	Ingman <i>et al.</i> (2000)
Waunana			X	Tamm <i>et al.</i> (2007)
Wayuu	X	X	X	Torres <i>et al.</i> (2006); Melton <i>et al.</i> (2007); Tamm <i>et al.</i> (2007); Yang <i>et al.</i> (2010)
Wichí	X			Cabana <i>et al.</i> (2006)
Xavante	X	X	X	Ward <i>et al.</i> (1996); Dornelles <i>et al.</i> (2005); Fagundes <i>et al.</i> (2008)
Xikrin	X	X		Lobato-da-Silva <i>et al.</i> (2001); Dornelles <i>et al.</i> (2005)
Yagua	X	X		Monsalve <i>et al.</i> (1994); Torres <i>et al.</i> (2006)
Yámana	X			García-Bour <i>et al.</i> (2004) ¹
Yanomámi	X	X	X	Easton <i>et al.</i> (1996); Santos <i>et al.</i> (1996); Merriwether <i>et al.</i> (2000); Lobato-da-Silva <i>et al.</i> (2001); Silva Jr <i>et al.</i> (2002, 2003) ² ; Williams <i>et al.</i> (2002); Dornelles <i>et al.</i> (2005); Fagundes <i>et al.</i> (2008)
Yungay	X			Lewis Jr <i>et al.</i> (2007)
Yuracare	X			Bert <i>et al.</i> (2004)
Zenu	X	X		Torres <i>et al.</i> (2006)
Zoró	X		X	Ward <i>et al.</i> (1996); Fagundes <i>et al.</i> (2008)

¹Ancient DNA. ²Sequencing included almost half of the genome (sites 7,148-15,976).

Southern Amerindian populations, respectively (*e.g.*, from 0.659 for the Quechua of Peru to 0.011 for the Xavante of the Brazilian Mato Grosso). Intra- and intergroup nucleotide diversity was calculated by Melton *et al.* (2007) for 20 of these Amerindian groups, whereas Barbieri *et al.* (2011) compared the sources of variation among North, Central and South Amerindians in 51 populations; the latter authors observed 3% variation among the three sets, 21% variation among populations within the subcontinent and 76% variation within populations.

To explore the mtDNA data further we compiled the prevalences of haplogroups A-D for 109 populations, in a total of 6,697 individuals distributed between latitude 11° North and 54° South, and longitude 46° to 78° West (Table 5). Sample sizes varied widely, from only one subject tested (Jebero) up to 491 (Yanomámi). The haplogroup frequencies reported in 52 articles also varied widely. The presence of mtDNA genomes of probable non-Amerindian origin was rare in all regions and populations, in contrast to the Y-SNP data (Table 1). Asymmetrical sex-mediated admixture was common during the first centuries of South

American colonization, and involved mostly European men and Amerindian/African women. The main consequences of this historical contact was the formation of mestizos and the present-day national societies; the former are characterized by a composite genome, with the majority of Y-chromosomes being of European origin, while their mtDNA derives from Amerindian or African sources (Bortolini *et al.*, 1999; Alves-Silva *et al.*, 2000; Carvalho-Silva *et al.*, 2001; Salzano and Bortolini, 2002). Asymmetrical mating could also explain the introduction of non-Amerindian Y-chromosomes into the tribes, while the autochthonous mtDNA genomes were preserved. However, the admixture dynamics are probably different from those observed in urban groups since they normally involve Amerindian women who live on reservations and men who live near the border of the reservations. In this situation, the children normally remain with their mothers. This phenomenon has been described for Guarani Indians (Marrero *et al.*, 2007), but the data presented here indicate that it could be much more common than previously thought.

Table 5 - Mitochondrial DNA haplogroup and linguistic and geographical information for the samples considered.

Population (n) ¹	Haplogroups (%)					Language	Geographical coordinates	References
	A	B	C	D	Others ²			
Wayuu (89)	26	28	45	0	1	Arawakan	11° N; 73° W	Mesa <i>et al.</i> (2000); Keyeux <i>et al.</i> (2002); Melton <i>et al.</i> (2007)
Kogi (153)	67	0	33	0	0	Chibchan	11° N; 74° W	Keyeux <i>et al.</i> (2002); Melton <i>et al.</i> (2007); Rojas <i>et al.</i> (2010)
Arsario (Wiwa) (76)	63	0	37	0	0	Chibchan	10° 25' N; 73° 05' W	Keyeux <i>et al.</i> (2002); Melton <i>et al.</i> (2007); Rojas <i>et al.</i> (2010)
Chimila (35)	88	0	3	6	3	Chibchan	10° 16' N; 74° 4' W	Keyeux <i>et al.</i> (2002)
Arhuaco (Ijka) (134)	87	1	12	0	0	Chibchan	9° 04' N; 73° 59' W	Keyeux <i>et al.</i> (2002); Melton <i>et al.</i> (2007); Rojas <i>et al.</i> (2010)
Yukpa (88)	0	100	0	0	0	Carib	8° 40' N; 72° 41' W	Keyeux <i>et al.</i> (2002)
Zenu (107)	19	38	36	5	2	Spanish	8° 30' N; 76° W	Mesa <i>et al.</i> (2000); Keyeux <i>et al.</i> (2002); Torres <i>et al.</i> (2006)
Embera (43)	53	35	2	5	5	Choco	7° N; 76° 30' W	Mesa <i>et al.</i> (2000); Keyeux <i>et al.</i> (2002)
Tule-Cuna (30)	50	27	20	0	3	Chibchan	6° 56' N; 76° 45' W	Keyeux <i>et al.</i> (2002)
Guane-Butaregua (33)	12	64	0	24	0	Chibchan	6° 15' N; 73° 15' W	Keyeux <i>et al.</i> (2002)
Cubeo (22)	27	18	50	5	0	Tucanoan	5° 9' N; 70° 18' W	Torres <i>et al.</i> (2006)
Makiritare (10)	20	0	70	10	0	Carib	5° 33' N; 65° 33' W	Torrioni <i>et al.</i> (1993)
Kali' na (Galibi) (29)	7	41	38	7	7	Carib	5° 31' N; 53° 47' W	Mazières <i>et al.</i> (2008)
Guahibo (99)	52	3	33	0	12	Guahiban	5° N; 69° W	Keyeux <i>et al.</i> (2002); Vona <i>et al.</i> (2005); Torres <i>et al.</i> (2006)
Wanana (161)	21	49	16	14	0	Choco	4° 50' N; 77° W	Keyeux <i>et al.</i> (2002); Tamm <i>et al.</i> (2007); Rojas <i>et al.</i> (2010)
Palikúr (64)	1	47	4	47	1	Arawakan	4° N; 51° 45' W	Lobato-da-Silva <i>et al.</i> (2001); Mazières <i>et al.</i> (2008)
Macushi (10)	10	20	30	40	0	Carib	4° N; 60° 50' W	Torrioni <i>et al.</i> (1993)
Páez (51)	59	12	27	2	0	Páez	3° 9' N; 75° 28' W	Keyeux <i>et al.</i> (2002); Torres <i>et al.</i> (2006)
Ocaína (2)	0	0	100	0	0	Witotoan	3° 58' N; 68° 2' W	Torres <i>et al.</i> (2006)
Jebero (1)	0	0	100	0	0	Cahuapanan	3° 58' N; 68° 2' W	Torres <i>et al.</i> (2006)
Piaroa (28)	36	11	21	32	0	Salivan	3° 57' N; 66° 22' W	Torrioni <i>et al.</i> (1993); Keyeux <i>et al.</i> (2002)
Desano (2)	50	0	0	50	0	Tucanoan	3° 24' N; 69° 40' W	Torres <i>et al.</i> (2006)
Wapishana (12)	0	25	8	67	0	Arawakan	3° 07' N; 60° 03' W	Torrioni <i>et al.</i> (1993)
Emerillon (30)	30	70	0	0	0	Tupi	3° N; 53° W	Mazières <i>et al.</i> (2008)
Guambiano (23)	4	4	79	13	0	Barbacoan	2° 6' N; 76° 23' W	Keyeux <i>et al.</i> (2002)
Yanomámi (491)	2	25	50	19	4	Yanomam	2° 50' N; 54° W	Torrioni <i>et al.</i> (1992, 1993); Easton <i>et al.</i> (1996); Merriwether <i>et al.</i> (2000); Williams <i>et al.</i> (2002); Silva Jr <i>et al.</i> (2003)
Guayabero (30)	50	17	13	0	20	Guahiban	2° 25' N; 71° 4' W	Keyeux <i>et al.</i> (2002)
Curripaco (22)	41	36	23	0	0	Arawakan	2° 10' N; 68° 54' W	Keyeux <i>et al.</i> (2002); Torres <i>et al.</i> (2006)
Tiriyó (32)	9	19	22	47	3	Carib	2° N; 56° W	Lobato-da-Silva <i>et al.</i> (2001); Silva Jr <i>et al.</i> (2003)
Nukak (20)	0	20	80	0	0	Maku	1° 44' N; 70° 44' W	Keyeux <i>et al.</i> (2002)
Apalaí (120)	37	1	30	32	0	Carib	1° 20' N; 54° 40' W	Lobato-da-Silva <i>et al.</i> (2001); Mazières <i>et al.</i> (2008)
Cayapa (120)	29	40	9	22	0	Barbacoan	1° 17' N; 78° 50' W	Rickards <i>et al.</i> (1999)
Wayampi (99)	62	11	8	19	0	Tupi	1° N; 53° W	Santos <i>et al.</i> (1996); Lobato-da-Silva <i>et al.</i> (2001); Silva Jr <i>et al.</i> (2003); Mazières <i>et al.</i> (2008)
Siona (12)	75	17	8	0	0	Tucanoan	0° 6' N; 75° 36' W	Keyeux <i>et al.</i> (2002)

Table 5 (cont.)

Population (n) ¹	Haplogroups (%)					Language	Geographical coordinates	References
	A	B	C	D	Others ²			
Pasto (9)	67	33	0	0	0	Barbacoan	0° 58' N; 77° 44' W	Keyeux <i>et al.</i> (2002)
Yagua (12)	25	0	67	8	0	Peba-Yaguan	0° 51' N; 72° 27' W	Torres <i>et al.</i> (2006)
Ingano (111)	18	38	42	0	2	Quechuan	0° 50' N; 77° W	Mesa <i>et al.</i> (2000); Keyeux <i>et al.</i> (2002); Torres <i>et al.</i> (2006); Rojas <i>et al.</i> (2010)
Tucano (17)	0	18	47	35	0	Tucanoan	0° 42' N; 69° 53' W	Keyeux <i>et al.</i> (2002)
Coreguaje (69)	4	20	66	6	4	Tucanoan	0° 38' N; 76° 8' W	Keyeux <i>et al.</i> (2002); Tamm <i>et al.</i> (2007)
Awa-Juriti (18)	0	72	11	0	17	Tucanoan	0° 16' N; 70° 45' W	Lobato-da-Silva <i>et al.</i> (2001)
Muinane (19)	11	21	37	26	5	Witotoan	0° 11' N; 73° 25' W	Keyeux <i>et al.</i> (2002)
Poturujara (23)	44	0	26	30	0	Tupi	0° 18' S; 55° 18' W	Lobato-da-Silva <i>et al.</i> (2001); Silva Jr <i>et al.</i> (2003)
Katuena (23)	26	9	35	30	0	Carib	0° 40' S; 57° 30' W	Lobato-da-Silva <i>et al.</i> (2001); Silva Jr <i>et al.</i> (2003)
Wai-wai (26)	15	15	43	27	0	Carib	0° 40' S; 58° W	Bonato and Salzano (1997)
Urubu Kaapor (42)	21	31	14	29	5	Tupi	2° -3° S; 46° -47° W	Torrioni <i>et al.</i> (1992, 1993); Lobato-da-Silva <i>et al.</i> (2001); Dornelles <i>et al.</i> (2005)
Huitoto (35)	23	3	25	46	3	Witotoan	2° 14' S; 72° 19' W	Keyeux <i>et al.</i> (2002); Torres <i>et al.</i> (2006)
Arara (70)	54	20	26	0	0	Carib	3° 30' -4° 20' S; 53° 0' -54° 10' W	Lobato-da-Silva <i>et al.</i> (2001); Ribeiro-dos-Santos <i>et al.</i> (2001); Silva Jr <i>et al.</i> (2003); Bisso-Machado (2010, MSc Dissertation, Universidade Federal do Rio Grande do Sul, Porto Alegre, RS, Brazil.)
Awa-Guajá (53)	13	87	0	0	0	Tupi	3° 30' S; 46° 40' W	Lobato-da-Silva <i>et al.</i> (2001)
Asurini (24)	4	54	17	21	4	Tupi	3° 35' -4° 12' S; 49° 40' -52° 26' W	Lobato-da-Silva <i>et al.</i> (2001)
Piapoco (39)	18	3	15	5	59	Arawakan	3° 36' S; 70° 23' W	Torres <i>et al.</i> (2006)
Puinave (19)	5	16	58	16	5	Puinave	3° 36' S; 70° 23' W	Torres <i>et al.</i> (2006)
Sáliba (13)	15	0	55	15	15	Salivan	3° 49' S; 70° 9' W	Torres <i>et al.</i> (2006)
Ticuna (371)	20	11	35	33	1	Ticuna	4° S; 69° 58' W	Schurr <i>et al.</i> (1990); Mesa <i>et al.</i> (2000); Torres <i>et al.</i> (2006); Mendes-Junior and Simões (2009); Rojas <i>et al.</i> (2010)
Parakanã (31)	6	39	32	23	0	Tupi	5° 22' S; 51° 17' W	Lobato-da-Silva <i>et al.</i> (2001); Bisso-Machado (2010, MSc Dissertation)
Xikrin (33)	30	64	3	3	0	Macro-Ge	5° 55' S; 51° W	Lobato-da-Silva <i>et al.</i> (2001); Dornelles <i>et al.</i> (2005); Bisso-Machado (2010, MSc Dissertation)
Suruí (44)	7	4	0	89	0	Tupi	5° 58' -10° 50' S; 48° 39' -61° 10' W	Bonato and Salzano (1997); Lobato-da-Silva <i>et al.</i> (2001)
Araweté (18)	39	0	50	11	0	Tupi	5° 9' S; 52° 22' W	Lobato-da-Silva <i>et al.</i> (2001)
Munduruku (92)	12	17	9	58	4	Tupi	6° 23' S; 59° 9' W	Torrioni <i>et al.</i> (1992, 1993); Lobato-da-Silva <i>et al.</i> (2001); Marrero <i>et al.</i> (2007); Bisso-Machado (2010, MSc Dissertation)
Marubo (10)	10	0	60	30	0	Panoan	6° 47' S; 72° 80' W	Torrioni <i>et al.</i> (1993)
Jamamadi (23)	0	0	96	4	0	Arauan	7° 15' S; 66° 41' W	Lobato-da-Silva <i>et al.</i> (2001); Bisso-Machado (2010, MSc Dissertation)
Yungay (38)	5	45	34	16	0	Quechuan	7° 26' S; 77° 4' W	Lewis Jr <i>et al.</i> (2007)
Ancash (33)	9	52	18	21	0	Quechua	7° 41' S; 77° 6' W	Lewis Jr <i>et al.</i> (2005)
Gorotire (11)	28	18	18	36	0	Macro-Ge	7° 44' S; 51° 10' W	Bisso-Machado (2010, MSc Dissertation)
Krahó (14)	29	57	14	0	0	Macro-Ge	8° S; 47° 15' W	Torrioni <i>et al.</i> (1993)
Kuben-Kran-Kegn (19)	58	26	6	10	0	Macro-Ge	8° 10' S; 52° 8' W	Bisso-Machado (2010, MSc Dissertation)

Table 5 (cont.)

Population (n) ¹	Haplogroups (%)					Language	Geographical coordinates	References
	A	B	C	D	Others ²			
Mekranoti (19)	26	63	11	0	0	Macro-Ge	8° 40' S; 54° W	Dornelles <i>et al.</i> (2005); Bisso-Machado (2010, MSc Dissertation)
Kubenkokre (4)	0	100	0	0	0	Macro-Ge	8° 43' S; 53° 23' W	Marrero <i>et al.</i> (2007)
Kayapó (13)	46	54	0	0	0	Macro-Ge	9° S; 53° W	Lobato-da-Silva <i>et al.</i> (2001)
Karitiana (19)	0	11	0	89	0	Tupi	9° 30' S; 64° 15' W	Lobato-da-Silva <i>et al.</i> (2001)
Cinta-Larga (45)	25	0	20	53	2	Tupi	9° 50' -12° 30' S; 59° 10' -60° 50' W	Lobato-da-Silva <i>et al.</i> (2001); Dornelles <i>et al.</i> (2005); Bisso-Machado (2010, MSc Dissertation)
Gavião (27)	15	15	0	70	0	Tupi	10° 10' S; 61° 8' W	Ward <i>et al.</i> (1996)
Tupe (16)	0	69	31	0	0	Aymaran	10° 16' S; 75° 47' W	Lewis Jr <i>et al.</i> (2007)
Txukahamãe (2)	100	0	0	0	0	Macro-Ge	10° 20' S; 53° 5' W	Dornelles <i>et al.</i> (2005)
Zoró (30)	20	7	13	60	0	Tupi	10° 20' S; 60° 20' W	Ward <i>et al.</i> (1996)
Matsiguenga (38)	5	92	0	3	0	Arawakan	10° 47' -12° 51' S; 73° 17' -70° 44' W	Mazières <i>et al.</i> (2008)
Kokraimoro (2)	50	50	0	0	0	Macro-Ge	10° 49' S; 55° 27' W	Marrero <i>et al.</i> (2007)
Pacaás Novos (Wari) (30)	40	30	27	3	0	Chapacura-Wanh am	11° 8' S; 65° W	Bisso-Machado (2010, MSc Dissertation)
Tayacaja (61)	21	33	13	30	3	Quechuan	12° 24' S; 74° 34' W	Fuselli <i>et al.</i> (2003)
Arequipa (22)	9	68	14	9	0	Quechua	13° 13' S; 72° 11' W	Fuselli <i>et al.</i> (2003)
Trinitario (35)	14	40	37	3	6	Arawakan	14° S; 65° W	Bert <i>et al.</i> (2001)
Xavante (25)	16	84	0	0	0	Macro-Ge	14° S; 52° 30' W	Ward <i>et al.</i> (1996)
Movima (22)	9	9	64	18	0	Movima	14° 26' S; 65° 53' W	Bert <i>et al.</i> (2001)
Quechua (232)	14	62	15	9	0	Quechuan	14° 30' S; 69° W	Merriwether <i>et al.</i> (1995); Bert <i>et al.</i> (2001); Silva Jr <i>et al.</i> (2003); Lewis Jr <i>et al.</i> (2007); Corella <i>et al.</i> (2007); Barbieri <i>et al.</i> (2011); Gayà-Vidal <i>et al.</i> (2011)
Chimane (Moseten) (71)	39	54	3	0	4	Chimane	14° 41' S; 66° 50' W	Bert <i>et al.</i> (2001); Corella <i>et al.</i> (2007);
Ignaciano (22)	18	36	41	0	5	Arawakan	15° 1' S; 66° 4' W	Bert <i>et al.</i> (2001)
Uro (64)	11	69	9	11	0	Uru-Chipaya	15° 45' S; 69° 53' W	Barbieri <i>et al.</i> (2011)
Yuracare (28)	39	32	21	4	4	Yuracare	17° S; 65° W	Bert <i>et al.</i> (2001)
Aymara (411)	4	76	8	11	1	Aymaran	17° 68' S; 69° 16' W	Merriwether <i>et al.</i> (1995); Easton <i>et al.</i> (1996); Bert <i>et al.</i> (2001); Lewis Jr <i>et al.</i> (2007); Corella <i>et al.</i> (2007); Barbieri <i>et al.</i> (2011); Gayà-Vidal <i>et al.</i> (2011)
Ayoreo (91)	0	0	83	17	0	Zamucoan	19° S; 60° 30' W	Dornelles <i>et al.</i> (2004)
Wichí (199)	12	51	7	29	1	Mataco-Guaicuru	22° 28' S; 62° 70' W	Torrioni <i>et al.</i> (1993); Bianchi <i>et al.</i> (1995); Bravi <i>et al.</i> (1995); Demarchi <i>et al.</i> (2001); Cabana <i>et al.</i> (2006)
Chorote (34)	15	44	23	18	0	Mataco-Guaicuru	22° 90' S; 65° 40' W	Bianchi <i>et al.</i> (1995); Bravi <i>et al.</i> (1995)
Humahuaca (46)	11	68	17	4	0	Spanish	23° 11' S; 65° 20' W	Dipierri <i>et al.</i> (1998)
Aché (63)	10	90	0	0	0	Tupi	23° 30' -24° 10' S; 55° 50' -56° 30' W	Schmitt <i>et al.</i> (2004)
Atacameño (79)	13	73	10	4	0	Atacama	23° 50' S; 68° W	Bailliet <i>et al.</i> (1994); Merriwether <i>et al.</i> (1995); Merriwether and Ferrell (1996)
Guarani (249)	77	6	9	6	2	Tupi	23° 6' S; 55° 12' W	Silva Jr <i>et al.</i> (2003); Marrero <i>et al.</i> (2007); García and Demarchi (2009)
Pilagá (41)	5	37	27	29	2	Mataco-Guaicuru	24° S; 59° W	Demarchi <i>et al.</i> (2001); Cabana <i>et al.</i> (2006)
Coya (60)	13	57	23	5	2	Coya	25° 30' S; 67° 28' W	Álvarez-Iglesias <i>et al.</i> (2007)

Table 5 (cont.)

Population (n) ¹	Haplogroups (%)					Language	Geographical coordinates	References
	A	B	C	D	Others ²			
Toba (80)	15	43	5	37	0	Mataco-Guaicuru	26° S; 58° W	Bianchi <i>et al.</i> (1995); Demarchi <i>et al.</i> (2001); Goicoechea <i>et al.</i> (2001); Cabana <i>et al.</i> (2006)
Jujuy (19)	16	58	16	10	0	Spanish	27° 27' S; 58° 59' W	Dipierri <i>et al.</i> (1998)
Kaingang (79)	47	4	48	0	1	Macro-Ge	28° S; 51° 20' W	Dornelles <i>et al.</i> (2005); Marrero <i>et al.</i> (2007)
Mocoví (5)	80	0	0	20	0	Mataco-Guaicuru	29° 51' S; 59° 56' W	Tamm <i>et al.</i> (2007)
Pehuenche (205)	2	9	40	49	0	Araucanian	37° 43' S; 71° 16' W	Merriwether <i>et al.</i> (1995); Moraga <i>et al.</i> (1997, 2000)
Mapuche (314)	5	23	32	36	4	Araucanian	39° 10' -41° 20' S; 68° 37' -70° 22' W	Ginther <i>et al.</i> (1993); Horai <i>et al.</i> (1993); Bailliet <i>et al.</i> (1994); Bianchi <i>et al.</i> (1995); Moraga <i>et al.</i> (2000)
Huilliche (207)	4	28	20	48	0	Araucanian	41° 16' S; 73° W	Bailliet <i>et al.</i> (1994); Merriwether <i>et al.</i> (1995); Merriwether and Ferrell (1996)
Aónikenk ^{3,4} (15)	0	0	27	73	0	Chon	45° S; 71° W	Lalueza (1995, PhD thesis, Universitat de Barcelona, Barcelona, Spain)
Tehuelche ⁴ (29)	0	20	24	56	0	Chon	45° S; 71° W	Moraga <i>et al.</i> (2000)
Yámana ³ (Yaghan) (32)	0	0	63	37	0	Yámana	47° S; 74° W	Lalueza (1995, PhD thesis); Moraga <i>et al.</i> (1997, 2000)
Kawéskar ² (Alacaluf) (19)	0	0	16	84	0	Alacalufan	49° S; 74° W	Lalueza (1995, PhD thesis)
Selknam ² (Ona) (16)	0	0	56	38	6	Chon	54° S; 74° W	Lalueza (1995, PhD thesis); García-Bour <i>et al.</i> (2004)

¹Arranged according to latitude. ²Probably of non-Amerindian origin. ³Ancient DNA. ⁴Aónikenk and Tehuelche are the same tribe separated by time. Aónikenk refers to ancient DNA.

Table 6 summarizes the influence of geography. In the seven regions that were defined, 74% of the variation occurred within populations, 6% among geographic divisions and 20% among populations within divisions. To analyze this variability further, the isolated frequencies of haplogroups A to D were plotted as shown in Figure 2. High frequencies of haplogroup C were observed in specific regions along the northwestern portion of the continent, with additional high spots in southern Brazil and

northern Argentina. The prevalences of haplogroups B and D showed a clear east-west separation, while for haplogroup A there were three main high prevalence nuclei in the north, center and south of the continent. Spearman's correlation coefficient between haplogroup frequencies and latitude yielded a positive value (0.27; $p < 0.01$) for haplogroup A, with a corresponding negative one (-0.25; $p < 0.01$) for haplogroup D. The coefficients for haplogroups B and C were not significant.

Table 6 - Mitochondrial DNA haplogroup frequencies by geography¹.

Geographic divisions	No. of populations	No. of individuals	Haplogroups (%)				
			A	B	C	D	Other
Amazonia	55	2410	20	21	31	25	3
Central Plateau	2	39	21	74	5	0	0
Southern Brazil	2	328	70	6	18	4	2
Chaco	6	479	10	43	22	24	1
Southern South America	3	726	4	20	31	43	2
Tierra del Fuego	5	111	0	5	39	55	1
Andes	35	2604	27	45	20	7	1
Total	108	6697					

¹AMOVA results: (a) Among geographic divisions: 6.2%; (b) Among populations within geographic divisions: 19.5%; (c) Within populations: 74.3%. The three values are statistically significant.

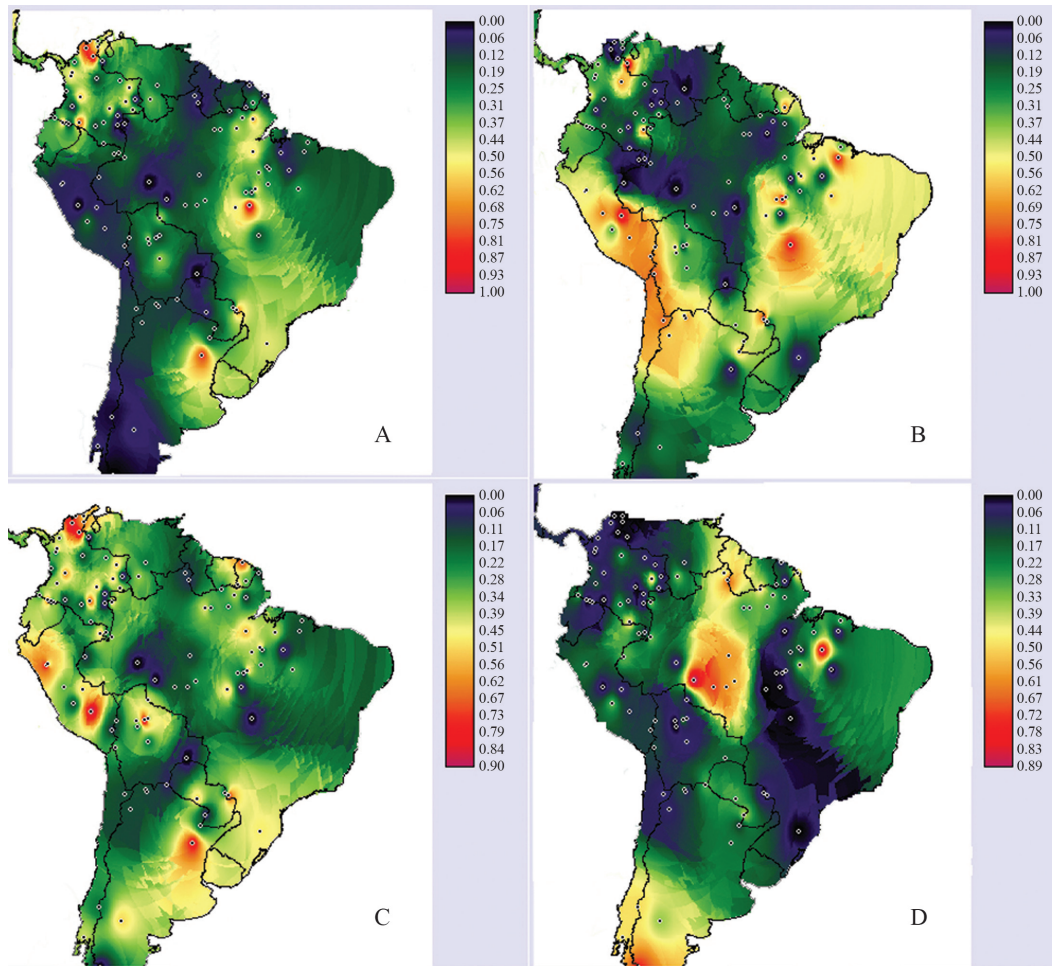


Figure 2 - Isoline map distribution showing the geographic pattern of the four (A-D) mtDNA haplogroups in South Amerindians. The dots indicate the locations of the populations sampled. As indicated in the scales given at right of each map, the colors represent the haplogroup frequencies, from dark blue (0.00) to red (1.00).

Table 7 summarizes the influence of language. Sixteen main language groups were considered, plus a composite set of “others”. The AMOVA results indicated that 73% of the haplogroup prevalence variability occurred

within populations, with 7% of it being attributable to languages. However, there was considerable heterogeneity (20%) within the language categories established. Overall, the variability was similar to that obtained for geography.

Table 7 - Mitochondrial DNA haplogroup frequencies by language¹.

Language	No. of populations	No. of individuals	Haplogroups (%)				
			A	B	C	D	Other
Tupi	16	889	38	24	10	27	1
Macro-Ge	11	221	37	38	21	3	1
Carib	9	408	24	32	25	18	1
Chibchan	6	461	69	6	22	2	1
Mataco-Guaicuru	5	359	13	46	11	29	1
Arawakan	8	321	16	38	24	13	9
Araucanian	3	726	4	20	31	43	2
Choco	2	204	28	46	13	12	1
Chon	3	60	0	10	33	55	2
Tucanoan	6	140	14	26	48	8	4

Table 7 (cont.)

Language	No. of populations	No. of individuals	Haplogroups (%)				
			A	B	C	D	Other
Aymaran	2	427	4	76	9	10	1
Barbacoan	3	152	28	34	19	19	0
Guahiban	2	129	51	6	29	0	14
Witotoan	3	56	18	9	32	37	4
Salivan	2	41	29	7	32	27	5
Quechuan	6	497	14	51	23	11	1
Other	21	1606	13	26	40	19	2
Total	108	6697					

¹AMOVA results: (a) Among language groups: 6.6%; (b) Among populations within language groups: 20.1%; (c) Within populations: 73.3%. The three values are statistically significant.

Conclusion

South Amerindians have been extensively studied with regard to the Y-chromosome, as well as and especially so for mtDNA markers. In agreement with studies from other regions, by far most of the mtDNA variability (73%-74%) is intrapopulation. Geographical and linguistic factors influenced the patterns of mtDNA diversity to a similar extent, while geography was apparently more important than language in explaining the data for the Y chromosome Q clade-STRs. Additional factors that may have influenced these results include distinct male and female migration patterns, as well as cultural and other characteristics. The fact that most studies have generally dealt with small populations, in which genetic drift may be important, could also have influenced the results.

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References

Altuna ME, Modesti NM and Demarchi DA (2006) Y-chromosomal evidence for a founder effect in Mbyá-guaraní Amerindians from northeast Argentina. *Hum Biol* 78:635-639.

Alvarez-Iglesias V, Jaime JC, Carracedo A and Salas A (2007) Coding region mitochondrial DNA SNPs: Targeting East Asian and Native American haplogroups. *Forensic Sci Int Genet* 1:44-55.

Alves-Silva J, da Silva Santos M, Guimarães PE, Ferreira AC, Bandelt HJ, Pena SD and Prado VF (2000) The ancestry of Brazilian mtDNA lineages. *Am J Hum Genet* 67:444-461.

Bailliet G, Rothhammer F, Carnese FR, Bravi CM and Bianchi NO (1994) Founder mitochondrial haplotypes in Amerindian populations. *Am J Hum Genet* 54:27-33.

Bailliet G, Ramallo V, Muzzio M, García A, Santos MR, Alfaro EL, Dipierri JE, Salceda S, Carnese FR, Bravi CM, *et al.* (2009) Brief communication: Restricted geographic distribution for Y-Q* paragroup in South America. *Am J Phys Anthropol* 140:578-582.

Barbieri C, Heggarty P, Castri L, Luiselli D and Pettener D (2011) Mitochondrial DNA variability in the Titicaca basin: Matches and mismatches with linguistics and ethnohistory. *Am J Hum Biol* 23:89-99.

Bert F, Corella A, Gené M, Pérez-Pérez A and Turbón D (2001) Major mitochondrial DNA haplotype heterogeneity in highland and lowland Amerindian populations from Bolivia. *Hum Biol* 73:1-16.

Bert F, Corella A, Gene M, Pérez-Pérez A and Turbón D (2004) Mitochondrial DNA diversity in the Llanos de Moxos: Moxo, Movima and Yuracare Amerindian populations. *Ann Hum Biol* 31:9-28.

Bianchi NO, Bailliet G and Bravi CM (1995) Peopling of the Americas as inferred through the analysis of mitochondrial DNA. *Braz J Genet* 18:161-168.

Bianchi NO, Catanesi CI, Bailliet G, Martínez-Marignac VL, Bravi CM, Vidal-Rioja LB, Herrera RJ and López-Camelo JS (1998) Characterization of ancestral and derived Y-chromosome haplotypes of New World native populations. *Am J Hum Genet* 63:1862-1871.

Bisso-Machado R, Jota MS, Ramallo V, Paixão-Côrtes VR, Lacerda DR, Salzano FM, Bonatto SL, Santos FR and Bortolini MC (2011) Distribution of Y-chromosome Q lineages in Native Americans. *Am J Hum Biol* 23:563-566.

Blanco-Verea A, Jaime JC, Brión M and Carracedo A (2010) Y-chromosome lineages in native South American populations. *Forensic Sci Int Genet* 4:187-193.

Bonatto SL and Salzano FM (1997) A single and early migration for the peopling of the Americas supported by mitochondrial DNA sequence data. *Proc Natl Acad Sci USA* 94:1866-1871.

Bortolini MC, Da Silva Junior WA, De Guerra DC, Remonato G, Mirandola R, Hutz MH, Weimer TA, Silva MC, Zago MA and Salzano FM (1999) African-derived South American populations: A history of symmetrical and asymmetrical

- matings according to sex revealed by bi- and uni-parental genetic markers. *Am J Hum Biol* 11:551-563.
- Bortolini MC, Salzano FM, Bau CHD, Layrisse Z, Petzl-Erler ML, Tsuneto LT, Hill K, Hurtado AM, Castro-de-Guerra D, Bedoya G, *et al.* (2002) Y-chromosome biallelic polymorphisms and Native American population structure. *Ann Hum Genet* 66:255-259.
- Bortolini MC, Salzano FM, Thomas MG, Stuart S, Nasanen SPK, Bau CHD, Hutz MH, Layrisse Z, Petzl-Erler ML, Tsuneto LT, *et al.* (2003) Y-Chromosome evidence for differing ancient demographic histories in the Americas. *Am J Hum Genet* 73:524-539.
- Bravi CM, Cejas S, Bailliet G, Goicoechea AS, Carnese FR and Bianchi NO (1995) Haplotipos mitocondriales en Amerindios. Abstracts do XXVI Congreso Argentino de Genética, San Carlos de Bariloche, pp 152.
- Cabana GS, Merriwether DA, Hunley K and Demarchi DA (2006) Is the genetic structure of Gran Chaco populations unique? Interregional perspectives on Native South American mitochondrial DNA variation. *Am J Phys Anthropol* 131:108-119.
- Carvalho-Silva DR, Santos FR, Hutz MH, Salzano FM and Pena SDJ (1999) Divergent human Y-chromosome microsatellite evolution rates. *J Mol Evol* 49:204-214.
- Carvalho-Silva DR, Santos FR, Rocha J and Pena SD (2001) The phylogeography of Brazilian Y-chromosome lineages. *Am J Hum Genet* 68:281-286.
- Corella A, Bert F, Pérez-Pérez A, Gené M and Turbón D (2007) Mitochondrial DNA diversity of the Amerindian populations living in the Andean Piedmont of Bolivia: Chimane, Mosen, Aymara and Quechua. *Ann Hum Biol* 34:34-55.
- Crawford MH (1998) *The Origins of Native Americans. Evidence from Anthropological Genetics.* Cambridge University Press, Cambridge, 308 pp.
- Demarchi DA and Mitchell RJ (2004) Genetic structure and gene flow in Gran Chaco populations of Argentina: Evidence from Y-Chromosome markers. *Hum Biol* 76:413-429.
- Demarchi DA, Panzetta-Dutari GM, Colantonio SE and Marcelino AJ (2001) Absence of the 9-bp deletion of mitochondrial DNA in pre-Hispanic inhabitants of Argentina. *Hum Biol* 73:575-582.
- Dipierrri JE, Alfaro E, Martínez-Marignac VL, Bailliet G, Bravi CM, Cejas S and Bianchi NO (1998) Paternal directional mating in two Amerindian subpopulations located at different altitudes in northwestern Argentina. *Hum Biol* 70:1001-1010.
- Dornelles CL, Battilana J, Fagundes NJ, Freitas LB, Bonatto SL and Salzano FM (2004) Mitochondrial DNA and Alu insertions in a genetically peculiar population: The Ayoreo. *Am J Hum Biol* 16:479-488.
- Dornelles CL, Bonatto SL, Freitas LB and Salzano FM (2005) Is haplogroup X present in extant South American Indians? *Am J Phys Anthropol* 127:439-448.
- Eastman JR (2006) *IDRISI 15.0: The Andes edition.* Clark University, Worcester.
- Easton RD, Merriwether DA, Crews DE and Ferrell RE (1996) mtDNA variation in the Yanomami: Evidence for additional New World founding lineages. *Am J Hum Genet* 59:213-225.
- Excoffier L and Lischer HE (2010) Arlequin suite ver 3.5: A new series of programs to perform population genetics analyses under Linux and Windows. *Mol Ecol Resour* 10:564-567.
- Excoffier L, Smouse PE and Quattro JM (1992) Analysis of molecular variance inferred from metric distances among DNA haplotypes: Application to human mitochondrial DNA restriction data. *Genetics* 131:479-491.
- Fagundes NJR, Kanitz R, Eckert R, Valls ACS, Bogo MR, Salzano FM, Smith DG, Silva Jr WA, Zago MA, Ribeiro-dos-Santos AK, *et al.* (2008) Mitochondrial population genomics supports a single pre-Clovis origin with a coastal route for the peopling of the Americas. *Am J Hum Genet* 82:583-592.
- Fehren-Schmitz L, Reindel M, Cagigao ET, Hummel S and Herrmann B (2010) Pre-Columbian population dynamics in coastal southern Peru: A diachronic investigation of mtDNA patterns in the Palpa region by ancient DNA analysis. *Am J Phys Anthropol* 141:208-221.
- Fehren-Schmitz L, Warnberg O, Reindel M, Seidenberg V, Tomasto-Cagigao E, Isla-Cuadrado J, Hummel S and Herrmann B (2011) Diachronic investigations of mitochondrial and Y-chromosomal genetic markers in pre-Columbian Andean highlanders from South Peru. *Ann Hum Genet* 75:266-283.
- Fuselli S, Tarazona-Santos S, Dupanloup I, Soto A, Luiselli D and Pettener D (2003) Mitochondrial DNA diversity in South America and the genetic history of Andean Highlanders. *Mol Biol Evol* 20:1682-1691.
- García A and Demarchi DA (2009) Incidence and distribution of Native American mtDNA haplogroups in central Argentina. *Hum Biol* 81:59-69.
- García F, Moraga M, Vera S, Henríquez H, Llop E, Aspillaga E and Rothhammer F (2006) mtDNA microevolution in Southern Chile's archipelagos. *Am J Phys Anthropol* 129:473-481.
- García-Bour J, Pérez-Pérez A, Álvarez S, Fernández E, López-Parra AM, Arroyo-Pardo E and Turbón D (2004) Early population differentiation in extinct aborigines from Tierra Del Fuego – Patagonia: Ancient mtDNA sequences and Y-chromosome STR characterization. *Am J Phys Anthropol* 123:361-370.
- Gayà-Vidal M, Moral P, Saenz-Ruales N, Gerbault P, Tonasso L, Villena M, Vasquez R, Bravi CM and Dugoujon JM (2011) mtDNA and Y-chromosome diversity in Aymaras and Quechuas from Bolivia: Different stories and special genetic traits of the Andean Altiplano populations. *Am J Phys Anthropol* 145:215-230.
- Geppert M, Baeta M, Núñez C, Martínez-Jarreta B, Zweynert S, Cruz OW, González-Andrade F, González-Solorzano J, Nagy M and Roewer L (2011) Hierarchical Y-SNP assay to study the hidden diversity and phylogenetic relationship of native populations in South America. *Forensic Sci Int Genet* 5:100-104.
- Ginther C, Corach D, Penacino GA, Rey JA, Carnese FR, Hutz MH, Anderson A, Just J, Salzano FM and King M-C (1993) Genetic variation among the Mapuche Indians from the Patagonian region of Argentina: Mitochondrial DNA sequence variation and allele frequencies of several nuclear genes. In: Pena SDJ, Chakraborty R, Epplen JT and Jeffreys AJ (eds) *DNA Fingerprinting: State of the Science.* Birkhäuser Verlag, Berlin, pp 211-219.

- Goicoechea AS, Carnese FR, Dejean C, Avena SA, Weimer TA, Estalote AC, Simões ML, Palatnik M, Salamoni SP, Salzano FM, *et al.* (2001) New genetic data on Amerindians from the Paraguayan Chaco. *Am J Hum Biol* 13:660-667.
- Horai S, Kondo S, Nakagawa-Hattori Y, Hayashi S, Sonoda S and Tajima K (1993) Peopling of the Americas, founded by four major lineages of mitochondrial DNA. *Mol Biol Evol* 10:23-47.
- Hunley KL, Cabana GS, Merriwether DA and Long JC (2007) A formal test of linguistic and genetic coevolution in native Central and South America. *Am J Phys Anthropol* 132:622-631.
- Ingman M, Kaessmann H, Pääbo S and Gyllensten U (2000) Mitochondrial genome variation and the origin of modern humans. *Nature* 408:708-713.
- Jota MS, Lacerda DR, Sandoval JR, Vieira PPR, Santos-Lopes SS, Bisso-Machado R, Paixão-Cortes VR, Revollo S, Pazy-Miño C, Fujita R, *et al.* (2011) A new subhaplogroup of Native American Y chromosomes from the Andes. *Am J Phys Anthropol* 146:553-559.
- Karafet T, Segura SL, Vuturo-Brady J, Posukh O, Osipova L, Wiebe V, Romero F, Long JC, Harihara S, Jin F, *et al.* (1997) Y chromosome markers and trans-Bering Strait dispersals. *Am J Phys Anthropol* 102:301-314.
- Karafet TM, Zegura SL, Posukh O, Osipova L, Bergen A, Long J, Goldman D, Klitz W, Harihara S, de Knijff P, *et al.* (1999) Ancestral Asian source(s) of New World Y-chromosome founder haplotypes. *Am J Hum Genet* 64:817-831.
- Karafet TM, Mendez FL, Meilerman MB, Underhill PA, Zegura SL and Hammer MF (2008) New binary polymorphisms reshape and increase resolution of the human Y chromosomal haplogroup tree. *Genome Res* 18:830-838.
- Keyeux G, Rodas C, Gelvez N and Carter D (2002) Possible migration routes into South America deduced from mitochondrial DNA studies in Colombian Amerindian populations. *Hum Biol* 74:211-233.
- Kivisild T, Shen P, Wall D, Do B, Sung R, Davis K, Passarino G, Underhill PA, Scharfe C, Torroni A, *et al.* (2006) The role of selection in the evolution of human mitochondrial genomes. *Genetics* 172:373-387.
- Leite FP, Callegari-Jacques SM, Carvalho BA, Kommers T, Matte CH, Raimann PE, Schwengber SP, Sortica VA, Tsuneto LT, Petzl-Erler ML, *et al.* (2008) Y-STR analysis in Brazilian and South Amerindian populations. *Am J Hum Biol* 20:359-363.
- Lell JT, Sukernik RI, Starikovskaya YB, Su B, Jin L, Schurr TG, Underhill PA and Wallace DC (2002) The dual origin and Siberian affinities of Native American Y chromosomes. *Am J Hum Genet* 70:192-206.
- Lewis Jr CM, Tito RY, Lizárraga B and Stone AC (2005) Land, language, and loci: mtDNA in Native Americans and the genetic history of Peru. *Am J Phys Anthropol* 127:351-360.
- Lewis Jr CM, Buikstra JE and Stone AC (2007) Ancient DNA and genetic continuity in the South Central Andes. *Lat Am Antiq* 18:1-48.
- Lobato-da-Silva DF, Ribeiro-dos-Santos AKC and Santos SEB (2001) Diversidade genética de populações humanas na Amazônia. In: Guimarães Vieira IC, Cardoso da Silva JM, Oren DC and D'Incao MA (eds) *Diversidade Humana e Cultural na Amazônia*. Museu Paraense Emílio Goeldi, Belém, pp 167-193.
- Marrero AR, Silva-Junior WA, Bravi CM, Hutz MH, Petzl-Erler ML, Ruiz-Linares A, Salzano FM and Bortolini MC (2007) Demographic and evolutionary trajectories of the Guarani and Kaingang natives of Brazil. *Am J Phys Anthropol* 132:301-310.
- Mazières S, Guitard E, Crubézy E, Dugoujon JM, Bortolini MC, Bonatto SL, Hutz MH, Bois E, Tiouka F, Larrouy G, *et al.* (2008) Uniparental (mtDNA, Y-chromosome) polymorphisms in French Guiana and two related populations – Implications for the region's colonization. *Ann Hum Genet* 72:145-156.
- Melton PE, Briceño I, Gómez A, Devor EJ, Bernal JE and Crawford MH (2007) Biological relationship between Central and South American Chibchan speaking populations: Evidence from mtDNA. *Am J Phys Anthropol* 133:753-770.
- Mendes-Junior CT and Simões AL (2009) Mitochondrial DNA variability among eight Tikúna villages: Evidence for an intratribal genetic heterogeneity pattern. *Am J Phys Anthropol* 140:526-531.
- Merriwether DA and Ferrell RE (1996) The four founding lineage hypothesis for the New World: A critical reevaluation. *Mol Phylogenet Evol* 5:241-246.
- Merriwether DA, Rothhammer F and Ferrell RE (1994) Genetic variation in the New World: Ancient teeth, bone, and tissue as sources of DNA. *Experientia* 50:592-601.
- Merriwether DA, Rothhammer F and Ferrell RE (1995) Distribution of the four founding lineage haplotypes in Native Americans suggests a single wave of migration for the New World. *Am J Phys Anthropol* 98:411-430.
- Merriwether DA, Kemp BM, Crews DE and Neel JV (2000) Gene flow and genetic variation in the Yanomama as revealed by mitochondrial DNA. In: Renfrew C (ed) *America Past, America Present: Genes and Languages in the Americas and Beyond*. Oxbow books, Oxford, pp 89-124.
- Mesa NR, Mondragón MC, Soto ID, Parra MV, Duque C, Ortiz-Barrientos D, García LF, Velez ID, Bravo ML, Múnera JG, *et al.* (2000) Autosomal, mtDNA, and Y-chromosome diversity in Amerindians: Pre- and post-Columbian patterns of gene flow in South America. *Am J Hum Genet* 67:1277-1286.
- Monsalve MV, Groot de Restrepo H, Espinel A, Correal G and Devine DV (1994) Evidence of mitochondrial DNA diversity in South American aboriginals. *Ann Hum Genet* 58:265-273.
- Moraga M, Rothhammer F and Carvallo P (1997) Mitochondrial DNA variation in aboriginal populations of southern Chile. In: Barton SA, Rothhammer F and Schull WS (eds) *Patterns of Morbidity in Andean Aboriginal Populations: 8,000 Years of Evolution*. Amphora Editora, Santiago, pp 32-36.
- Moraga ML, Rocco P, Miquel JF, Nervi F, Llop E, Chakraborty R, Rothhammer F and Carvallo P (2000) Mitochondrial DNA polymorphisms in Chilean aboriginal populations: Implications for the peopling of the southern cone of the continent. *Am J Phys Anthropol* 113:19-29.
- Moraga M, Santoro CM, Standen VG, Carvallo P and Rothhammer F (2005) Microevolution in prehistoric Andean populations: Chronologic mtDNA variation in the desert valleys of northern Chile. *Am J Phys Anthropol* 127:170-181.
- Pena S, Santos FR, Bianchi NO, Bravi CM, Carnese RF, Rothhammer F, Gerelsaikhan T, Munkhtuja B and Oyunsuren T

- (1995) A major founder Y-chromosome haplotype in Amerindians. *Nat Genet* 11:15-16.
- Perego UA, Achilli A, Angerhofer N, Accetturo M, Pala M, Olivieri A, Kashani BH, Ritchie KH, Scozzari R, Kong Q-P, *et al.* (2009) Distinctive Paleo-Indian migration routes from Beringia marked by two rare mtDNA haplogroups. *Curr Biol* 19:1-8.
- Perego UA, Angerhofer N, Pala M, Olivieri A, Lancioni H, Kashani BH, Carossa V, Ekins JE, Gómez-Carballa A, Huber G, *et al.* (2010) The initial peopling of the Americas: A growing number of founding mitochondrial genomes from Beringia. *Genome Res* 20:1174-1179.
- Ribeiro-dos-Santos AKC, Guerreiro JF, Santos SEB and Zago MA (2001) The split of the Arara population: Comparison of genetic drift and founder effect. *Hum Hered* 51:79-84.
- Rickards O, Martinez-Labarga C, Lum JK, De Stefano GF and Cann RL (1999) mtDNA history of the Cayapa Amerinds of Ecuador: Detection of additional founding lineages for the native American populations. *Am J Hum Genet* 65:519-530.
- Rodriguez-Delfin L, Santos SEB and Zago MA (1997) Diversity of the human Y chromosome of South American Amerindians: A comparison with Blacks, Whites and Japanese from Brazil. *Ann Hum Genet* 61:439-448.
- Rojas W, Parra MV, Campo O, Caro MA, Lopera JG, Arias W, Duque C, Naranjo A, García J, Vergara C, *et al.* (2010) Genetic makeup and structure of Colombian populations by means of uniparental and biparental DNA markers. *Am J Phys Anthropol* 143:13-20.
- Sala A, Argüelles CF, Marino ME, Bobillo C, Fenocchio A and Corach D (2010) Genetic analysis of six communities of Mbyá-Guaraní inhabiting northeastern Argentina by means of nuclear and mitochondrial polymorphic markers. *Hum Biol* 82:433-456.
- Salzano FM and Callegari-Jacques SM (1988) South American Indians: A Case Study in Evolution. Clarendon Press, Oxford, 259 pp.
- Salzano FM and Bortolini MC (2002) Evolution and Genetics of Latin American Populations. Cambridge University Press, Cambridge, 512 pp.
- Santos FR, Hutz MH, Coimbra CEA, Santos RV, Salzano FM and Pena SDJ (1995) Further evidence for the existence of a major founder Y chromosome haplotype in Amerindians. *Braz J Genet* 18:669-672.
- Santos SEB, Ribeiro-dos-Santos AKC, Meyer D and Zago MA (1996) Multiple founder haplotypes of mitochondrial DNA in Amerindians revealed by RFLP and sequencing. *Ann Hum Genet* 60:305-319.
- Schmitt R, Bonatto SL, Freitas LB, Muschner VC, Hill K, Hurtado AM and Salzano FM (2004) Extremely limited mitochondrial DNA variability among the Aché Natives of Paraguay. *Ann Hum Biol* 31:87-94.
- Schurr TG and Sherry ST (2004) Mitochondrial DNA and Y chromosome diversity and the peopling of the Americas: Evolutionary and demographic evidence. *Am J Hum Biol* 16:420-439.
- Schurr TG, Ballinger SW, Gan Y-Y, Hodge JA, Merriwether DA, Lawrence DN, Knowler WC, Weiss KM and Wallace DC (1990) Amerindian mitochondrial DNAs have rare Asian mutations at high frequencies, suggesting they derived from four primary maternal lineages. *Am J Hum Genet* 46:613-623.
- Seielstad M, Yuldasheva N, Singh N, Underhill P, Oefner P, Shen P and Wells RS (2003) A novel Y-chromosome variant puts an upper limit on the timing of first entry into the Americas. *Am J Hum Genet* 73:700-705.
- Shimada I, Shinoda K-I, Farnum J, Corruccini R and Watanabe H (2004) An integrated analysis of pre-Hispanic mortuary practices. *Curr Anthropol* 45:369-402.
- Shinoda K, Adachi N, Guillen S and Shimada I (2006) Mitochondrial DNA analysis of ancient Peruvian highlanders. *Am J Phys Anthropol* 131:98-107.
- Silva Jr WA, Bonatto SL, Holanda AJ, Ribeiro-dos-Santos AK, Paixão BM, Goldman GH, Abe-Sandes K, Rodriguez-Delfin L, Barbosa M, Paçó-Larson ML, *et al.* (2002) Mitochondrial genome diversity of Native Americans supports a single early entry of founder populations into America. *Am J Hum Genet* 71:187-192.
- Silva Jr WA, Bonatto SL, Holanda AJ, Ribeiro-dos-Santos AK, Paixão BM, Goldman GH, Abe-Sandes K, Rodriguez-Delfin L, Barbosa M, Paçó-Larson ML, *et al.* (2003) Correction: Mitochondrial DNA variation in Amerindians. *Am J Hum Genet* 72:1346-1348.
- Tamm E, Kivisild T, Reidla M, Metspalu M, Smith DG, Mulligan CJ, Bravi CM, Rickards O, Martinez-Labarga C, Khusnutdinova EK, *et al.* (2007) Beringian standstill and spread of Native American founders. *PLoS One* 2:e829.
- The Y Chromosome Consortium (2002) A nomenclature system for the tree of human Y-chromosomal binary haplogroups. *Genome Res* 12:339-348.
- Torres MM, Bravi CM, Bortolini MC, Duque C, Callegari-Jacques S, Ortiz D, Bedoya G, Groot de Restrepo H and Ruiz-Linares A (2006) A revertant of the major founder Native American haplogroup C common in populations from northern South America. *Am J Hum Biol* 18:59-65.
- Torrioni A, Schurr TG, Yang C-C, Szathmari EJE, Williams RC, Schanfield MS, Troup GA, Knowler WC, Lawrence DN, Weiss KM, *et al.* (1992) Native American mitochondrial DNA analysis indicates that the Amerind and the Nadene populations were founded by two independent migrations. *Genetics* 130:153-162.
- Torrioni A, Schurr TG, Cabell MF, Brown MD, Neel JV, Larsen M, Smith DG, Vullo CM and Wallace DC (1993) Asian affinities and continental radiation of the four founding Native American mtDNAs. *Am J Hum Genet* 53:563-590.
- Toscanini U, Gusmão L, Berardi G, Amorim A, Carracedo A, Salas A and Raimondi E (2008) Y chromosome microsatellite genetic variation in two Native American populations from Argentina: Population stratification and mutation data. *Forensic Sci Int Genet* 2:274-280.
- Toscanini U, Gusmão L, Berardi G, Gomes V, Amorim A, Salas A and Raimondi E (2011) Male lineages in South American native groups: Evidence of M19 traveling south. *Am J Phys Anthropol* 146:188-196.
- Underhill PA, Jin L, Zeman R, Oefner PJ and Cavalli-Sforza LL (1996) A pre-Columbian Y chromosome-specific transition and its implications for human evolutionary history. *Proc Natl Acad Sci USA* 93:196-200.
- Underhill PA, Jin L, Lin AA, Mehdi SQ, Jenkins T, Vollrath D, Davis RW, Cavalli-Sforza LL and Oefner PJ (1997) Detection of numerous Y chromosome biallelic polymorphisms by denaturing high-performance liquid chromatography. *Genome Res* 7:996-1005.

- Underhill PA, Passarino G, Lin AA, Shen P, Mirazón Lahr M, Foley RA, Oefner PJ and Cavalli-Sforza LL (2001) The phylogeography of Y chromosome binary haplotypes and the origins of modern human populations. *Ann Hum Genet* 65:43-62.
- Vallinoto AC, Cayres-Vallinoto IM, Ribeiro Dos Santos ÂKC, Zago MA, Santos SE and Guerreiro JF (1999) Heterogeneity of Y chromosome markers among Brazilian Amerindians. *Am J Hum Biol* 11:481-487.
- Vona G, Falchi A, Moral P, Calò CM and Varesi L (2005) Mitochondrial sequence variation in the Guahibo Amerindian population from Venezuela. *Am J Phys Anthropol* 127:361-369.
- Ward RH, Salzano FM, Bonatto SL, Hutz MH, Coimbra CEA and Santos RV (1996) Mitochondrial DNA polymorphism in three Brazilian Indian tribes. *Am J Hum Biol* 8:317-323.
- Weir BS (1996) The second National Research Council report on forensic DNA evidence. *Am J Hum Genet* 59:497-500.
- Weir BS and Cockerham CC (1984) Estimating F-statistics for the analysis of population structure. *Evolution* 38:1358-1370.
- Williams SR, Chagnon NA and Spielman RS (2002) Nuclear and mitochondrial genetic variation in the Yanomamo: A test case for ancient DNA studies of prehistoric populations. *Am J Phys Anthropol* 117:246-259.
- Yang NN, Mazières S, Bravi C, Ray N, Wang S, Burley MW, Bedoya G, Rojas W, Parra MV, Molina JA, *et al.* (2010) Contrasting patterns of nuclear and mtDNA diversity in Native American populations. *Ann Hum Genet* 74:525-538.
- Zegura SL, Karafet TM, Zhivotovsky LA and Hammer MF (2004) High-resolution SNPs and microsatellite haplotypes point to a single, recent entry of Native American Y chromosomes into the Americas. *Mol Biol Evol* 21:164-175.

Internet Resources

- PASW Statistics 18, <http://www.spss.com> (June 10, 2011).
- Pubmed, <http://www.ncbi.nlm.nih.gov/pubmed/> (June 10, 2011).
- Y Chromosome Haplotype Reference Database, <http://www.yhrd.org/> (June 10, 2011).
- Lewis MP (2009) *Ethnologue: Languages of the World*. 16th edition. SIL International, Dallas, <http://www.ethnologue.com/>.

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