



Original article

Frequency of alleles and haplotypes of the human leukocyte antigen system in Bauru, São Paulo, Brazil

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Background: HLA allele identification is used in bone marrow transplant programs as HLA compatibility between the donor and recipient may prevent graft rejection.

Objective: This study aimed to estimate the frequency of alleles and haplotypes of the HLA system in the region of Bauru and compare these with the frequencies found in other regions of the country.

Methods: HLA-A*, HLA-B*, and HLA-DRB1* allele frequencies and haplotypes were analyzed in a sample of 3542 volunteer donors at the National Registry of Voluntary Bone Marrow Donors (REDOME) in Bauru. HLA low resolution typing was performed using reverse line blot with the Dynal Reli™ SSO-HLA Typing Kit and automated Dynal AutoReli™48 device (Invitrogen, USA).

Results: Twenty, 36, and 13 HLA-A*, HLA-B*, and HLA-DRB1* allele groups, respectively, were identified. The most common alleles for each locus were HLA-A*02, HLA-B*35, and HLA-DRB1*07. The most frequent haplotype was A*01-B*08-DRB1*03. Allele and haplotype frequencies were compared to other regions in Brazil and the similarities and differences among populations are shown.

Conclusion: The knowledge of the immunogenic profile of a population contributes to the comprehension of the historical and anthropological aspects of different regions. Moreover, this helps to find suitable donors quickly, thereby shortening waiting lists for transplants and thus increasing survival rates among recipients.

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Introduction

The major histocompatibility complex (MHC) is a system composed by genes that encode the human leukocyte antigen (HLA) molecules.^{1,2} HLA genes are part of a polygenic complex located on the short arm of chromosome 6 in the 6p21.3 region.^{1,3} HLA genes are expressed by all nucleated cells of the human body. The main function of this system is the presentation of antigenic peptides to T lymphocytes in order to trigger the proliferation and differentiation of cells capable of activating a specific immunological response.^{2,3}

The HLA class I molecules (HLA-A, HLA-B, and HLA-C) display endogenous antigens to CD8⁺ cytotoxic T lymphocytes, while the HLA class II molecules (HLA-DR, HLA-DQ and HLA-DP) present exogenous antigens to CD4⁺ T-helper lymphocytes.^{2,3}

HLA molecules differ among individuals, and these differences are associated with solid organ rejection after organ transplantation as well as with graft-versus-host disease (GVHD) after bone marrow transplantation. The identification of HLA alleles is the main tool used in bone marrow transplant programs as HLA compatibility between the donor and recipient is necessary to prevent graft rejection.⁴

In Brazil, the National Voluntary Bone Marrow Donor Registry (REDOME) was founded in 1993 with the aim of gathering data of volunteer donors who want to donate bone marrow to patients requiring a transplant.^{5,6} REDOME, with about 3 million people registered, is currently considered the third largest bone marrow donor program; the United States has the world's largest bank followed by Germany.^{7,8}

Similar to REDOME, the National Registry of Bone Marrow Recipients (REREME) stores information about patients waiting for bone marrow transplants. These patients are added to the registry when a matching donor is unavailable.^{5,6}

REDOME is important because it increases the chance of finding a matching for a particular recipient. In Brazil, the probability of genetic compatibility among unrelated donors is small due to miscegenation.⁸ The probability of finding a recipient compatible with an unrelated donor is about 1:100,000,⁹ and the probability of a sibling being compatible is 25%.¹⁰

The genetic characterization of individuals leads to a rapid selection of unrelated donors for recipients; thus, waiting lists for transplantation and possibly graft rejection decreases, thereby improving patient survival.¹¹

In addition to the role in transplantation and susceptibility or resistance to diseases, HLA is also important for anthropological studies as the frequency of alleles varies between ethnic groups. For example, HLA-B*35 is most often found in Caucasians, whereas HLA-B*15 is common in Africans.

The population of Brazil is a good example of miscegenation. Colonization and immigration have made the population diverse; the genetic information of native Amerindians has mixed with that of Europeans and Africans brought to Brazil as slaves.^{11,12} Immigrants from other parts of the world, such as Germans, Arabs, Italians, Spanish and Japanese have

since come to Brazil to find better living conditions; these immigrants have helped constitute the new genetic profile of Brazilians.¹³

According to the demographic census carried out by the Brazilian Institute for Geography and Statistics (IBGE), the population of the city of Bauru in 2010 was estimated to be 343,937;¹⁴ the majority of city residents were White (70.66%), followed by Mulatto (22.69%), Black (4.95%), Asian (1.57%), and Amerindian (0.13%).¹⁵ Until 1850, the region was inhabited solely by Amerindians, but pioneers from Minas Gerais and São Paulo explored this land but did not adopt the slavery system which was prevalent in other regions of the state until 1868.¹⁶

The distribution of HLA variants has been analyzed in different Brazilian regions and throughout the world. Monte et al.¹² established the frequencies of HLA alleles in Teresina, Piauí and concluded that multiracial people, the dominant ethnic group in this region, have predominately Caucasian and African genes with a small proportion of Amerindian genes.

Bortolotto et al.¹¹ compared the HLA alleles of ethnic groups from different regions of Brazil and noticed some similarities and differences among them. The HLA-A*02 allele was common in populations from Rio Grande do Sul and Paraná, regions mostly populated by Caucasians. The frequency of this allele was also high in mulattos from Piauí;^{11,12} however, the HLA-B*15 allele, common in Black people, was found in Caucasians, Blacks, and Mulattos.¹¹

Middleton et al.¹⁷ and Williams et al.¹⁸ analyzed the distribution of the HLA-A and HLA-B alleles, respectively, in diverse populations including Brazilians. Both studies showed common alleles among the studied populations, thus indicating the extent of miscegenation possibly due to migration.^{17,18}

The characterization and determination of population allele frequencies is important because this can identify whether these predominant alleles exist in a specific region, if they are shared with other populations and if they are related to disease susceptibility or protection.^{17,19}

Through HLA class I (HLA-A* and -B*) and class II (HLA-DRB1*) allele typing from the registry of volunteers (REDOME) of Bauru, São Paulo, we aimed to estimate the frequencies of HLA alleles and haplotypes prevalent in the region, and to compare the frequencies of these alleles to those of other regions of Brazil.

Methods

Population

Among donors listed in the REDOME from 2008 to 2012, 3542 volunteer bone marrow donors were selected and evaluated at the Immunogenetics Laboratory of the Instituto Lauro de Souza Lima in Bauru. Volunteers were evaluated according to gender, age, marital status, ethnic group and HLA type.

Results from published studies on the populations of Ribeirão Preto (n = 184),²⁰ São Paulo (n = 239),²⁰ Paraná

(n = 2775),²⁰ Rio Grande do Sul (n = 5000),¹¹ Minas Gerais (n = 1000),²⁰ Pernambuco (n = 101),²⁰ and Piauí (n = 97)¹² were used to compare HLA frequencies.

The study was approved by the Ethics Committee of the Instituto Lauro de Souza Lima, Bauru.

Extraction of DNA and HLA typing

DNA was isolated by the salting-out method using venous blood kept in ethylenediaminetetraacetic acid (EDTA) anticoagulant.²¹

The typing of HLA class I alleles (loci A* and B*) and class II alleles (locus DRB1*) was performed by the reverse line blot technique at a low resolution (Dynal Reli™ SSO-HLA Typing Kit and Dynal AutoReli™48, Invitrogen, USA).

The DNA samples were amplified using specific biotinylated primers to each HLA region (0.5 μM) using 20 mM of Tris-HCl solution (with 30% glycerol and 100 mM KCl), DNTPs (400 μM of dATP, dCTP, and dGTP; 800 μM of dUTP), Taq polymerase (100 μ/mL), and sodium azide (0.05%). Samples underwent 35 cycles at 95°C, 60°C, and 72°C denaturation, annealing and extension temperatures, respectively. After amplification by polymerase chain reaction (PCR), the amplicons were chemically denatured and added to a nylon membrane containing sequence-specific oligonucleotide probes. The amplicons marked with biotin hybridized with the corresponding probes with the complementary target sequence and were observed using a colorimetric reaction; conjugated streptavidin, hydrogen peroxide and tetramethylbenzidine (TMB) substrate were added. Perfect matching program (PMP) software was used for interpretation.

Table 1 - Gender, age, marital status and ethnicity of donors from Bauru registered in REDOME (n = 3542).

Variable	n	%
<i>Gender</i>		
Female	1941	54.80
Male	1600	45.17
unavailable	1	0.03
Age (years old)	18 to 55	Median 30
<i>Marital status</i>		
Single	1743	49.21
Married	1498	42.29
Divorced	217	6.13
Widower	27	0.76
Stable union	13	0.37
Others	44	1.24
<i>Ethnicity</i>		
White	3056	86.28
Black	165	4.66
Mulatto	158	4.46
Asian	50	1.14
Others	113	3.19

Statistical analysis

Allele frequencies were obtained by direct count and the haplotype construction was performed using a probabilistic computational model. The distribution of gene frequencies in the population was checked using the Hardy-Weinberg equilibrium, and the analysis of haplotypes was performed using the Arlequin software version 3.1.²²

Statistical differences between populations were determined by the chi-square test with the SISA online software.²³ Significance was set at a p-value of ≤ 0.05.

Table 2 - Frequencies of the HLA-A, B and DRB1 alleles in the Bauru population.

HLA-A	AF	HLA-B	AF	HLA-DRB1	AF
A*01	0.088	B*07	0.061	DRB1*01	0.100
A*02	0.263	B*08	0.050	DRB1*03	0.092
A*03	0.094	B*13	0.020	DRB1*04	0.092
A*11	0.055	B*14	0.051	DRB1*07	0.149
A*23	0.045	B*15	0.078	DRB1*08	0.047
A*24	0.098	B*18	0.051	DRB1*09	0.015
A*25	0.012	B*27	0.024	DRB1*10	0.018
A*26	0.034	B*35	0.120	DRB1*11	0.129
A*29	0.042	B*37	0.011	DRB1*12	0.017
A*30	0.058	B*38	0.021	DRB1*13	0.129
A*31	0.042	B*39	0.032	DRB1*14	0.043
A*32	0.034	B*40	0.047	DRB1*15	0.111
A*33	0.029	B*41	0.013	DRB1*16	0.041
A*34	0.006	B*42	0.012		
A*36	0.004	B*44	0.106		
A*66	0.007	B*45	0.014		
A*68	0.055	B*46	0.0004		
A*69	0.001	B*47	0.002		
A*74	0.008	B*48	0.005		
A*80	0.001	B*49	0.028		
		B*50	0.027		
		B*51	0.083		
		B*52	0.019		
		B*53	0.021		
		B*54	0.001		
		B*55	0.009		
		B*56	0.003		
		B*57	0.029		
		B*58	0.024		
		B*59	0.001		
		B*67	0.001		
		B*71	0.001		
		B*73	0.001		
		B*78	0.001		
		B*81	0.002		
		B*82	0.001		

AF: allele frequency.

Table 3 - Statistically significant differences between HLA-A alleles in the Bauru population compared to the populations of Ribeirão Preto, São Paulo, Paraná, Rio Grande do Sul, Minas Gerais and Piauí.

Locus HLA	Bauru (n = 3542) (AF)	Ribeirão Preto (n = 184) (AF)	p-value		São Paulo (n = 239) (AF)	p-value		Paraná (n = 2775) (AF)	p-value		Rio Grande do Sul (n = 5000) (AF)	p-value		Minas Gerais (n = 1000) (AF)	p-value		Pernambuco (n = 101) (AF)	p-value		Piauí (n = 97) (AF)	p-value		
A*02	0.263	0.217			0.272			0.232	0.0001	0.475	0.0010	0.236	0.0139	0.287			0.402						
A*03	0.094	0.074			0.057	0.0074		0.095		0.197		0.098		0.084			0.227						
A*11	0.055	0.091	0.0080		0.073			0.053		0.099		0.055		0.025			0.093						
A*23	0.045	0.041			0.067	0.0364		0.035		0.078		0.059	0.0118	0.129		0.0001	0.113						
A*24	0.098	0.107			0.094			0.106		0.196		0.081	0.0212	0.104			0.186						
A*25	0.012	0.005			0.010			0.017		0.034	0.0281	0.008		0.014			0.021						
A*29	0.042	0.049			0.040			0.040		0.092		0.057	0.0087	0.030			0.072						
A*30	0.058	0.071			0.073			0.032	0.0001	0.070	0.0001	0.073		0.045			0.113						
A*31	0.042	0.044			0.030			0.043		0.100	0.0306	0.032	0.0307	0.049			0.072						
A*33	0.029	0.025			0.030			0.020	0.0008	0.055		0.035		0.035			0.041						
A*34	0.006	0.008			0.020	0.0013		0.003		0.008	0.0299	0.012	0.0237	0.005			0.041						
A*36	0.004	-			0.005			0.002	0.0229	0.007		0.007		0.005			0.031						
A*43	0	-			-			0.001	0.0184	0.001		-		-			-						
A*66	0.007	0.011			0			0.007		0.016		0.014	0.0114	0.030		0.0025	0.041						
A*68	0.055	0.074			0.040			0.052		0.094	0.0100	0.076	0.0010	0.089			0.155						
A*74	0.008	0.016			0.025	0.0007		0.005	0.0286	0.011	0.0248	0.015	0.0133	0.020			0.031						
A*80	0.001	0.003			0			0.002		0.001		0.003		0.010	0.0281		-						

HLA: human leukocyte antigen; AF: allele frequency; -: untyped alleles. Statistical differences set for p -values ≤ 0.05 .

Results

This study included 3542 volunteers registered in REDOME [1941 females (54.80%) and 1600 males (45.17%)]. The age of the donors ranged from 18 to 55 years (median, 30 years) and most of the volunteers were single ($n = 1743 - 49.21\%$). According to the responses of the participants about their ethnic background, 3056 (86.28%) individuals classified themselves as Whites, 165 (4.66%) as Blacks, 79 (2.23%) as Mulattos, 50 (1.14%) as Asians and 113 (3.19%) as others (Table 1). All participants were from cities covered by the Bauru Regional Health Department (DRS-VI).

Twenty, 36, and 13 alleles were identified in the HLA-A, HLA-B, and HLA-DRB1 groups, respectively. The most common characteristics found at each locus are listed in Table 2. In this population, the most frequently identified alleles at locus A, B and DRB1* were A*02 (26.3%), B*35 (12.0%), and DRB1*07 (14.9%) respectively.

Tables 3, 4, and 5 show the differences between the frequencies of HLA alleles found in the population of Bauru compared to those of studies from Ribeirão Preto, São Paulo, Paraná, Rio Grande do Sul, Minas Gerais, Pernambuco, and Piauí.

In descending order, the frequencies of A*01-B*08-DRB1*03 (1.9%), A*29-B*44-DRB1*07 (1.4%), A*02-B*44-DRB1*07 (1.1%), A*02-B*07-DRB1*15 (0.9%), A*02-B*18-DRB1*11(0.8%), and A*11-B*35-DRB1*01 (0.8%) haplotypes were identified in this study (Table 6).

Discussion

The present study evaluated the frequency expression of HLA, class I (HLA-A* and B*) and class II (HLA-DRB1*) alleles and haplotypes of bone marrow donors registered in the REDOME in Bauru.

We found that the donors were predominantly White, female and tended to be young adults. Our results agree with the demographics reported for this city as presented by the IBGE census¹⁵ and by a study of the northern region of Paraná State by Bardi et al.²⁴

The most frequent alleles observed in Bauru were HLA-A*02, HLA-B*35, and HLA-DRB1*07. Similar to this study, the most common allele found in all Brazilian populations (Ribeirão Preto,²⁰ São Paulo,²⁰ Paraná,²⁰ Rio Grande do Sul,¹¹ Minas Gerais,²⁰ Pernambuco²⁰ and Piauí¹²) was HLA-A*02.

The frequency of the HLA-B and HLA-DRB1 alleles differed between certain regions. For HLA-B, the most frequent allele was HLA-B*44 in Ribeirão Preto²⁰ and São Paulo,²⁰ and this was the second most common allele in the current study. In Pernambuco,²⁰ the HLA-B*42 allele was the most common, however this allele was rare in our population. Moreover, the HLA-B*07 allele was the most common in Piauí¹² and the fifth most common in our study.

The frequency of the HLA-DRB1*07 allele was the same for our region as well as for Ribeirão Preto,²⁰ however, the HLA-DRB1*13 allele was the most common in São Paulo,²⁰ Rio

Table 4 - Statically significant differences of the HLA-B alleles in the Bauru population compared with the populations of Ribeirão Preto, São Paulo, Paraná, Rio Grande do Sul, Minas Gerais and Piauí.

Locus HLA	Bauru (n = 3542)		Ribeirão Preto (n = 184)		São Paulo (n = 239)		Paraná (n = 2775)		Rio Grande do Sul (n = 5000)		Minas Gerais (n = 1000)		Pernambuco (n = 101)		Piauí (n = 97)	
	(AF)	(AF)	p-value	(AF)	p-value	(AF)	p-value	(AF)	p-value	(AF)	p-value	(AF)	p-value	(AF)	p-value	
B*07	0.061	0.058		0.073		0.069		0.133		0.066		0.099	0.0423	0.206	0.0267	
B*08	0.050	0.049		0.036		0.055		0.118	0.0135	0.047		0.015	0.0339	0.144		
B*13	0.020	0.016		0.010		0.022		0.031	0.0293	0.016		0.020	-	-		
B*14	0.051	0.077	0.0479	0.067		0.036	0.0001	0.103		0.060		0.049		0.052		
B*15	0.078	0.060		0.102		0.080		0.163		0.077		0.124	0.0283	0.186		
B*35	0.120	0.126		0.089	0.0372	0.113		0.236		0.115		0.074		0.176		
B*38	0.021	0.016		0.030		0.028	0.0142	0.048		0.015		0.005		0.052		
B*40	0.047	0.047		0.040		0.048		0.098		0.037	0.0458	0.044		0.134		
B*41	0.013	0.016		0.020		0.008	0.0036	0.021		0.018		0.015		0.001		
B*42	0.012	0.025		0.005		0.004	0.0001	0.011	0.0001	0.025	0.0001	0.198	0.0001	0.052		
B*45	0.014	0.011		0.020		0.010	0.0150	0.023		0.034	0.0001	0.030		0.031		
B*47	0.002	-		0		0.002		0.001	0.0001	0.002		0.015		-		
B*48	0.005	0.005		0.005		0.005		0.013		0.003		0.030	0.0001	-		
B*49	0.028	0.027		0.020		0.021	0.0069	0.054		0.031		0.040		0.052		
B*53	0.021	0.027		0.040	0.0124	0.012	0.0001	0.030	0.0030	0.039	0.0001	0.035		0.083		
B*54	0.001	-		0.005		0	0.0317	-		-		-		-		
B*55	0.009	0.008				0.010		0.021		0.013		0.005		0.031		
B*57	0.029	0.025		0.062	0.0001	0.031		0.058		0.037		0.040		0.001		
B*58	0.024	0.036		0.005	0.0069	0.015	0.0002	0.040		0.030		0.025		0.083		
B*73	0.001	-		0		0		0.001		0.003	0.0463	-		-		
B*81	0.002	0.011	0.0362	0		0.001	0.0192	0.001	0.0001	0.005		-		0.021		
B*82	0.001	-		0		0		0.001		0.001		-		-		

HLA: human leukocyte antigen; AF: allele frequency; -: untyped alleles.
 Statistical differences set for p-values ≤ 0.05.

Table 5 - Statically significant differences of the HLA-DR alleles in the Bauru population compared with the populations of Ribeirão Preto, São Paulo, Paraná, Rio Grande do Sul, Minas Gerais and Piauí.

Locus HLA	Bauru (n = 3542)		Ribeirão Preto (n = 184)		São Paulo (n = 239)		Paraná (n = 2775)		Rio Grande do Sul (n = 5000)		Minas Gerais (n = 1000)		Pernambuco (n = 101)		Piauí (n = 97)	
	(AF)	(AF)	p-value	(AF)	p-value	(AF)	p-value	(AF)	p-value	(AF)	p-value	(AF)	p-value	(AF)	p-value	
DRB1*04	0.092	0.113		0.115		0.122	0.0001	0.233	0.0001	0.115	0.0038	0.268				
DRB1*07	0.149	0.137		0.115		0.199	0.0001	0.246	0.0001	0.142		0.144	0.0040			
DRB1*08	0.047	0.060		0.052		0.048		0.119	0.0004	0.048		0.186	0.0057			
DRB1*09	0.015	0.038	0.0028	0.028		0.011	0.0216	0.031		0.018		0.001				
DRB1*10	0.018	0.025		0.015		0.011	0.0008	0.030		0.023		0.031				
DRB1*11	0.129	0.129		0.125		0.130		0.220	0.0001	0.108	0.0121	0.289				
DRB1*13	0.129	0.137		0.147		0.117	0.0387	0.256		0.157	0.0016	0.258				
DRB1*14	0.043	0.022		0.038		0.038		0.082		0.031	0.0132	0.093				
DRB1*15	0.111	0.110		0.094		0.090	0.0001	0.173	0.0001	0.108		0.165				
DRB1*16	0.041	0.016	0.0220	0.032		0.033	0.0108	0.070	0.0217	0.030	0.0183	0.072				

HLA: human leukocyte antigen; AF: allele frequency.
 Statistical differences set for p-values ≤ 0.05.

Table 6 - Comparison of the most common haplotypes in the population of Bauru with the populations of Paraná and Rio Grande do Sul.

Haplotype	Bauru	Paraná	p-value ^a	Rio Grande do Sul	p-value ^a
A*01 B*08 DRB1*03	0.019			0.028	0.0003
A*03 B*07 DRB1*15	0.006	0.013	0.0001	0.013	0.0001
A*02 B*44 DRB1*01	0.004			0.008	0.0022
A*02 B*15 DRB1*04	0.002			0.007	0.0001
A*02 B*44 DRB1*04	0.004	0.007	0.0248		
A*02 B*44 DRB1*07	0.011	0.007	0.0353		

^a p-value ≤ 0.05.

Grande do Sul¹¹ and Minas Gerais.²⁰ The HLA-DRB1*11 allele was the third most common in Piauí,¹² but the second most common in our population.

The A*01-B*08-DRB1*03 haplotype was most frequently observed in Bauru, Paraná²⁰ and Rio Grande do Sul.¹¹ However, in the study conducted by Bortolotto et al.,¹¹ the frequency of the A*01-B*08-DRB1*03 haplotype was significantly higher in Rio Grande do Sul than in the region of Bauru. The A*03-B*07-DRB1*15, A*02-B*44-DRB1*01, and A*02-B*15-DRB1*04 haplotypes were uncommon in the population from Bauru, which differed significantly from the frequencies present in the population from Rio Grande do Sul. When the frequencies of the haplotypes in Bauru were compared to those of Paraná,²⁰ there were significant differences for the A*03-B*07-DRB1*15 and A*02-B*44-DRB1*04 haplotypes. These haplotypes were uncommon in our region. The A*01-B*44-DRB1*07 haplotype was the third most common in our region and the tenth most common in Paraná.

The Brazilian population is genetically diverse due to miscegenation that occurred during its history.^{8,12} The frequencies of some alleles identified in the population of Bauru were similar to those of populations of other regions of Brazil and some differed. As expected, the population of Bauru resembles the populations of Ribeirão Preto and São Paulo, which are in the same state. Interestingly, the population of Bauru has similarities with the populations of Pernambuco and Piauí in northeast Brazil, distant from our region. Of these two areas, the population of Piauí had the greatest resemblance to that of Bauru. One possible explanation for this could be that both regions had similar colonization with Portuguese and Indians, and little influence of Africans.²⁵ Moreover, some people may have migrated to the northeastern region of São Paulo State looking for better living conditions.^{26,27} Regarding the other populations, Bauru, in terms of alleles and haplotypes, is more different from Paraná, Rio Grande do Sul, and Minas Gerais, even though they are geographically closer. The colonization of the southern region of Brazil was marked by the immigration of Germans,²⁸ Spanish, Italians, and Africans²⁹ and in Minas Gerais, most immigrants were Portuguese, Africans and Italians.³⁰ Differences in the populations that immigrated to the various regions may explain the differences in the HLA system.

Conclusion

The HLA compatibility between donors and recipients is essential for a successful bone marrow transplantation. Therefore, the estimation of the immunogenic profile in our region and other regions in Brazil can assist in targeting, developing, and maintaining the REDOME database. The identification of the frequencies of HLA alleles within a nation is important because it eases the burden of searching compatible donors, decreases the wait for transplants and therefore increases the chances of survival of the recipients. In addition, knowledge of HLA frequencies can improve our understanding of the historical and anthropological composition of populations, that may also allow for areas of interest in each region to be identified.

Conflicts of interest

The authors declare no conflicts of interest.

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