PREVALENCE OF *Trypanosoma cruzi* AND *Leishmania chagasi* INFECTION AND RISK FACTORS IN A COLOMBIAN INDIGENOUS POPULATION

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**SUMMARY**

This study was carried out in order to obtain base-line data concerning the epidemiology of American Visceral Leishmaniasis and Chagas’ Disease in an indigenous population with whom the government is starting a dwelling improvement programme. Information was collected from 242 dwellings (1,440 people), by means of house to house interviews about socio-economic and environmental factors associated with *Leishmania chagasi* and *Trypanosoma cruzi* transmission risk. A leishmanin skin test was applied to 385 people and 454 blood samples were collected on filter paper in order to detect *L. chagasi* antibodies by ELISA and IFAT and *T. cruzi* antibodies by ELISA.

*T. cruzi* seroprevalence was 8.7% by ELISA, *L. chagasi* was 4.6% and 5.1% by IFAT and ELISA, respectively. ELISA sensitivity and specificity for *L. chagasi* antibodies were 57% and 97.5% respectively, as compared to the IFAT. Leishmanin skin test positivity was 19%. *L. chagasi* infection prevalence being defined as a positive result in the three-immunodiagnostic tests, was 17.1%. Additionally, 2.7% of the population studied was positive to both *L. chagasi* and *T. cruzi*, showing a possible cross-reaction. *L. chagasi* and *T. cruzi* seropositivity increased with age, while no association with gender was observed. Age (p<0.007), number of inhabitants (p<0.05), floor material (p<0.03) and recognition of vector (p<0.01) were associated with *T. cruzi* infection, whilst age (p<0.007) and dwelling improvement (p<0.02) were associated with *L. chagasi* infection. It is necessary to evaluate the long-term impact of the dwelling improvement programme on these parasitic infections in this community.

**KEYWORDS:** Visceral leishmaniasis; Chagas’ disease; Seroprevalence; Colombia.

**INTRODUCTION**

The Tripanosomatidae which affect man in America are from the *Leishmania* and *Trypanosoma* genera. The first combines different leishmaniasis species, within which *Leishmania chagasi* is found. The American Visceral Leishmaniasis’ (AVL) causative agent; within the second is found *Trypanosoma cruzi*, the causative agent of American trypanosomiasis. These parasites occupy shared ecological niches in many parts of America, from the south of the United States to Argentina, which allow simultaneous evaluation of the two entities.

In Colombia, AVL was described for the first time in 1944 by GAST GALVIS in a patient from the State of Santander. After notification of this AVL case, almost 20 years (1960) passed until two new cases were reported. From 1981 onwards, more than 600 new cases have been recorded, which can be explained by amongst other aspects, the suspension of DDT spraying in *L. chagasi* synanthropic foci and an improvement in the diagnosis and reporting of the disease. The Tolima department is the region of the country where most visceral leishmaniasis cases have been registered since 1981; 37.3% corresponded to the municipality of Coyaima, which makes it one of the country’s most important foci.

On the other hand, the estimated figures for human infection with *T. cruzi* in Colombia oscillate between 1,200,000 and 1,700,000 inhabitants and some 40,000 new cases occur every year, principally in regions situated below 2,000 metres above sea-level and which coincide with the distribution of the principal vector in Colombia, *Rhodnius proliris*, in the departments of Guajira, Cesar, Santander, Boyacá, Arauca, Casanare, Cundinamarca, Tolima, Meta, Huila, Caquetá, Bolívar and Caldas. Some blood-bank seroprevalence studies from endemic zones in the departments of Santander, Norte de Santander, Tolima, Boyaca, Cundinamarca, Meta, Huila, Guajira, Cesar, Arauca, Casanare and Caqueta, show levels oscillating between 3% and 6%.

In the municipality of Coyaima, *R. proliris* was found in habitations and to be infected with *T. cruzi* and *T. rangeli*. Some previous serological surveys revealed active trypanosome transmission. However, until the present, no studies have been initiated to allow the magnitude of American trypanosomiasis in this region to be established.

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In the municipality of Coyaima, *R. proliris* was found in habitations and to be infected with *T. cruzi* and *T. rangeli*. Some previous serological surveys revealed active trypanosome transmission. However, until the present, no studies have been initiated to allow the magnitude of American trypanosomiasis in this region to be established.
According to the characteristics of the region in question, it is necessary to establish the dynamics of the two entities’ transmission, by means of the evaluation of the different ecological, socio-economic and cultural factors, which influence its transmission chain. This study was intended to learn about *T. cruzi* and *L. chagasi* infection prevalence and their possible relationship with reservoirs, vectors and associated risk factors. Given that these parasitoses are determined, for the greater part, by housing conditions, the study allowed base-line information to be obtained concerning epidemiological aspects in order to evaluate the impact of a housing modification and improvement programme on the prevalence of these diseases. This programme was initiated by the government due to the 1991 epidemic in this municipality and consisted of peridomiciliary changes, latrine, farmyard and kitchen construction and domestic changes such as modifications to walls, roof and/or floor.

**MATERIALS AND METHODS**

The study was carried out in the municipality of Coyaima, located in the Tolima department, covering an area of 667 km². The town centre is located at 03° 48’ 09'' latitude North and 75° 11’ 54” longitude West and 350 m altitude (Fig. 1). The average temperature is 28.2 °C, with an annual rainfall of 1,502 mm. The surrounding land is distributed between thermic, hot and mild levels. Economic activities of greatest importance are agriculture, dairy-farming and mining. Principal crops are coffee, sesame seeds, maize and millo. Precious metal, lime and barita mines are being worked.

In the chosen localities, 242 homes (100% of the total) were surveyed during 1995-1996, concerning risk, socio-economic and environmental factors associated with *L. chagasi* and *T. cruzi* transmission such as: number of years as householders, overcrowding, type of housing, roof, floor and wall materials, public services to the housing, prevention measures taken for vector control such as spraying and use of mosquito-netting, vector presence and identification, reservoir presence, possession of animals and housing modification.

The survey revealed a population of 1,440 people of indigenous ethology, with equal gender proportions (sexes) and showed that of the 242 homes, 124 had been included in the housing modification and improvement programme.

The procedure which was applied will be described next, previous written consent having been obtained from the 538 people who made up the sample. The sample presented the same age and gender distribution as the original population.

The Montenegro test was applied to the whole sample, reading of which was only possible for 385 people, 48 hours after inoculation, individuals with ≥4 mm indurations being considered to be positive. The test was made with *L. chagasi* promastigote soluble extract, containing 25 µg/ml parasite protein, manufactured at the Instituto Nacional de Salud according to previously described methodology.

Blood samples were taken from 454 people, from the sample’s total 538, on Whatman #3 filter paper on which tests to determine *T. cruzi* and *L. chagasi* antibodies were carried out utilizing ELISA and IFAT (Indirect Fluorescence Antibody Test) techniques, according to previously described methodology. IFAT was carried out following CAMARGO; DE SOUZA & CAMARGO and CORREDOR & LÓPEZ’s procedures. The ELISA was made according to BARTLETT et al., using *T. cruzi* and *L. chagasi* soluble antigen in 0.75 µg/ml and 500 µg/ml concentrations respectively and 1:1000 in *T. cruzi* and 1:50 in *L. chagasi*. Optical densities greater than 0.4 and 0.5 were considered to be ELISA positive for *T. cruzi* and *L. chagasi*. Titres equal to or greater than 1:32 for the IFAT test were considered to be positive for *L. chagasi*. All individuals who proved to be positive, in any of the tests, were submitted to a clinical examination looking for possible signs of Chagas’ disease or AVL.

The search for AVL infection in domestic reservoirs was carried out on 22 dogs belonging to houses where seropositive individuals or those reactive to leishmanina for *L. chagasi* were found. These dogs were evaluated by clinical examination and direct smear of the popliteal ganglion haemolymph.

Phlebotomine captures were undertaken for 10 days, inside the housing’s living space as well as in the peridomiciliary area where positive cases were registered. Forest samples were also taken from form-wooded areas close to the housing. These samples were taken during dawn and nightfall hours (approximately 8 hours/day/person) during the month of July, using automatic and manual aspirators on human and animal bait (chicken, pigs, cattle and dogs). Captured Phlebotomines were stored in 70% alcohol and information concerning day, hour and capture site were noted. The insects so collected were identified in the Universidad

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1. Doyare Centro
2. Doyare Esmeralda

**Fig. 1** - Municipality of Coyaima, Tolima located in the centre of the country.

For risk factor analysis, a house was considered positive when one or more individuals living in it were positive by any of the tests performed.

The proportions were compared by X² test and Fischer test when necessary. The EPIINFO programme version 6.04 was used to make these tests.

RESULTS

Seroprevalence

The ELISA test was positive in 8.7% for T. cruzi antibodies. For L. chagasi, ELISA and IFAT positivity was 5.1% and 4.6% respectively. Montenegro Skin Test (MST) was 19.0% (Table 1). If all individuals positive for any one of the three immunodiagnostic methods are considered to be L. chagasi positive cases, then positivity was 23.3%.

The positivity of the MST (reactive individuals over population examined) showed a statistically significant progressive increase with age (p< 0.05) (Figure 2).

In the case of T. cruzi infection as evaluated by ELISA, positivity increased with age and then decreased in the 40-49 year old group (Figure 3). L. chagasi antibody detection was carried out by ELISA as well as IFAT, finding a progressive increase with age, however a diminution was observed in the 20-29 and 40-49 year old groups (Figure 3). The ELISA test showed a 57% sensitivity and 97.5% specificity as opposed to IFAT (routine test used for LVA diagnosis by National Institute of Health of Colombia) (Table 2). Serological positivity with both tests (IFAT and ELISA) had an 80.4% agreement with the Montenegro test (Table 3). 2.7% of the people were simultaneously positive for L. chagasi and T. cruzi, indicating a possible crossed reaction or a mixed infection since individuals were encountered who were serologically positive for T. cruzi and negative for L. chagasi as determined by MST.

No differences were observed between positivity with respect to gender for T. cruzi nor for L. chagasi. Serologically reactive individuals did not show important clinical alterations compatible with AVL or Chagas’ disease.

Table 1

<table>
<thead>
<tr>
<th>L. chagasi INFECTION</th>
<th>ELISA</th>
<th>IFAT</th>
<th>Montenegro Skin Test</th>
</tr>
</thead>
<tbody>
<tr>
<td>POSITIVE</td>
<td>23</td>
<td>21</td>
<td>73</td>
</tr>
<tr>
<td>NEGATIVE</td>
<td>431</td>
<td>433</td>
<td>312</td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th>T. cruzi INFECTION</th>
<th>ELISA</th>
</tr>
</thead>
<tbody>
<tr>
<td>POSITIVE</td>
<td>39</td>
</tr>
<tr>
<td>NEGATIVE</td>
<td>409</td>
</tr>
</tbody>
</table>

Total: 454 454 385 448

Table 2
Comparison between serological tests for L. chagasi antibody detection. Coyaima, 1996

<table>
<thead>
<tr>
<th>IFAT</th>
<th>POSITIVE</th>
<th>NEGATIVE</th>
<th>TOTAL</th>
</tr>
</thead>
<tbody>
<tr>
<td>ELISA</td>
<td>12</td>
<td>11</td>
<td>23</td>
</tr>
<tr>
<td>NEGATIVE</td>
<td>9</td>
<td>422</td>
<td>431</td>
</tr>
<tr>
<td>TOTAL</td>
<td>21</td>
<td>433</td>
<td>454</td>
</tr>
</tbody>
</table>

Fig. 2 - Montenegro skin test by age groups. Coyaima. 1996.

Fig. 3 - Seroprevalence by T. cruzi and L. chagasi by age groups.
Housing and risk factors.
1. 100% of the housing studied did not have a drinking-water supply, nor electric light and only housing included in the modification programme had excretion disposal facilities (latrines or fixed sanitary ware).
2. 50.6% of the houses presented earth floors and 90.1% bahareque (wooden-slat) walls.
3. 84.7% and 84.5% of the population did not use repellent nor mosquito netting respectively.
4. 86.3% recognized the vectors of the entities studied and 91% had seen wild reservoirs near their housing.

Table 4 describes the housing’s principal characteristics and different risk factors for risks associated with T. cruzi and L. chagasi positivity. Statistically significant associations were encountered with vector presence and identification, number of inhabitants per house and flooring material in T. cruzi infection, whilst with L. chagasi only association with housing modification was found (Table 4).

Entomology
In the entomological study 42 Lutzomyia samples were found, which were identified as being L. cayennensis and L. longipalpis. L. longipalpis was found with greater frequency, a specie demonstrated to be the principal vector of L. chagasi in America. It is interesting to note that in spite of the low number of samples collected, various of these were found in the housing’s walls, demonstrating intradomiciliary habits. In the majority of the homes farmyards with domestic animals (chicken, pigs) were found near the housing, a situation which increased the lutzomyias population, facilitating transmission. However, it is necessary to carry out vectorial studies in different seasons of the year to determine the months of greatest density.

Regarding the domestic reservoir study, the 22 dogs studied proved to be negative.

**DISCUSSION**

The 19% positivity found in the Montenegro Skin Test, is low when compared to results obtained in other zones of the country. In Córdoba 40%–20 was found and 51.2% in Cundinamarca. However, this positivity is similar to that found in endemic regions of Africa. As this test has high specificity and cutaneous leishmaniasis was not found in the zone studied, the allergic index, as well as the increase in positivity level with age, confirms active L. chagasi infection transmission.

L. chagasi serological prevalence was found to be similar to that observed in other seroepidemiological studies. Serological values diminished over time whilst cellular immunity measured by Montenegro test remained positive. 3.0% of the T. cruzi and L. chagasi serologically positive but MST negative people could be explained by the possible crossed reactions between Leishmania and Trypanosoma as has been previously reported. Additionally, when comparing these serological tests, ELISA and IFAT, with the Montenegro test high concordance is observed, 80.4%, if compared with that published by BADARÓ et al., showed a 47% concordance in Brazil.

L. chagasi infection in the majority of cases is not clinically manifested, which explains the high positivity in immunodiagnostic tests and the low presence of clinical cases of AVL. Thus, in this study, the positive people were not found to be ill and showed that they had not previously presented a picture compatible with AVL. However, it must not be forgotten that infected patients do not present the disease, thanks to the immune system’s efficient action; but, once this system is altered Leishmania behaves as an opportunistic parasite as is shown in AIDS patients, in the world’s different endemic zones.

If the IFAT is considered to be the gold-standard, it is seen that ELISA has low sensitivity and high specificity for detecting L. chagasi antibodies as has been reported by other authors.

When comparing positivity by gender, no difference was shown in any age group, which suggests that the exposure risk is equal amongst women and men.

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Table 3
Comparison of Serology and Montenegro Skin Test in L. chagasi infection detection. Coyaima, 1996

<table>
<thead>
<tr>
<th>SEROLOGY</th>
<th>MONTENEGRO</th>
<th>FREQUENCY</th>
<th>PERCENTAGE (%)</th>
</tr>
</thead>
<tbody>
<tr>
<td>ELISA and/or IFAT</td>
<td>TEST</td>
<td></td>
<td></td>
</tr>
<tr>
<td>+</td>
<td>+</td>
<td>11</td>
<td>3.7</td>
</tr>
<tr>
<td>+</td>
<td>-</td>
<td>10</td>
<td>3.0</td>
</tr>
<tr>
<td>-</td>
<td>+</td>
<td>49</td>
<td>16.5</td>
</tr>
<tr>
<td>-</td>
<td>-</td>
<td>230</td>
<td>76.7</td>
</tr>
<tr>
<td>TOTAL</td>
<td></td>
<td>300</td>
<td>100</td>
</tr>
</tbody>
</table>

Table 4
Relationship between housing characteristics and risk factors associated with T. cruzi and L. chagasi infection. Coyaima, 1996

<table>
<thead>
<tr>
<th>ASSOCIATED RISK FACTORS</th>
<th>CASE INFECTION T. cruzi</th>
<th>CASE INFECTION L. chagasi</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>X² P</td>
<td>X² P</td>
</tr>
<tr>
<td>Housing Modification</td>
<td>2 0.15</td>
<td>5.32 0.02*</td>
</tr>
<tr>
<td>Wall Material</td>
<td>0 0.99</td>
<td>2.52 0.11</td>
</tr>
<tr>
<td>Flooring Material</td>
<td>4.32 0.03*</td>
<td>1.69 0.19</td>
</tr>
<tr>
<td>Cement</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Earth</td>
<td>0.51 0.47</td>
<td>0.53 0.96</td>
</tr>
<tr>
<td>Housing Fumigation</td>
<td>0.01 0.93</td>
<td>0.56 0.45</td>
</tr>
<tr>
<td>Use of mosquito netting</td>
<td>0.01 0.93</td>
<td>0.48 0.49</td>
</tr>
<tr>
<td>Possession of animals</td>
<td>1.11 0.29</td>
<td>0.03 0.85</td>
</tr>
<tr>
<td>Possession of dogs</td>
<td>5.65 0.01*</td>
<td>3.27 0.07</td>
</tr>
<tr>
<td>Vector Identification</td>
<td></td>
<td></td>
</tr>
<tr>
<td># inhabitants per house</td>
<td>3.86 0.049*</td>
<td>0.43 0.5</td>
</tr>
<tr>
<td>&lt; 2 inhabitants</td>
<td></td>
<td></td>
</tr>
<tr>
<td>&gt; 2 inhabitants</td>
<td></td>
<td></td>
</tr>
</tbody>
</table>
The \( T. cruzi \) positivity found indicates its active transmission in the region. Even though it is true that the crossed reaction with \( L. chagasi \) can explain some seropositives, 6% were only positive for \( T. cruzi \). This figure is low when compared to highly endemic Colombian regions where a 26% positivity has been found\(^1\). However, to evaluate the problem's real magnitude, it is necessary to do further serological studies and to carry out electrocardiographic evaluation on positive individuals.

No correlation between housing modification and \( T. cruzi \) infection was found, whilst in \( L. chagasi \) infection greater infection was observed in the modified houses. This could be attributed to the short time elapsed between the modification of the housing and this survey, which is not useful for measuring the impact of housing modification although it can be employed as baseline data for the long-term assessment of the housing modification program. Additionally a positive relationship between triatomine recognition and \( T. cruzi \) seropositivity was found.

In general, the studied area’s socio-economic conditions are precarious constituting the principal risk factor in the transmission of AVL as well as of Chagas’ disease.

RESUMO

Prevalência da infecção por \textit{Trypanosoma cruzi} e \textit{Leishmania chagasi} e fatores de risco numa população indígena da Colômbia

Este estudo foi realizado para obter a linha de base da epidemiologia da Leishmaniose Visceral Americana e da Doença de Chagas numa comunidade indígena, onde o governo está desenvolvendo um programa de melhoramento da habitação. A coleta de dados referentes aos fatores sócio-econômicos e do meio ambiente associados ao risco de transmissão de \textit{Leishmania chagasi} e \textit{Trypanosoma cruzi} foi feita por meio de respostas a questionário endereçado aos componentes acima mencionados. O inquérito foi realizado em 242 unidades domiciliares (1440 indivíduos). Foi realizada a prova de Montenegro em 385 indivíduos e colhidas 454 amostras de sangue em papel de filtro, para pesquisar o teor de anticorpos contra \( L. chagasi \) por meio das técnicas de ELISA e IFI e o teor de anticorpos contra \( T. cruzi \) por meio de ELISA.

A prevalência sorológica foi de 8,7% para \( T. cruzi \), 4,6% e 5,1% para \( L. chagasi \) por meio de IFI e ELISA, respectivamente. Ao se comparar estas duas provas foi encontrado que por meio de ELISA a sensibilidade e especificidade para detecção de anticorpos contra \( L. chagasi \) foi de 57% e 97% respectivamente. Os resultados da intradermo-reação de Montenegro revelaram uma positividade de 19%. Os resultados dos três testes de imunodiagnóstico mostraram uma prevalência da infecção por \( L. chagasi \) de 17,1%. Além disso, 2,7% da população estudada apresentou reações sorológicas positivas para os dois parasitos, evidenciando uma possível reação cruzada. A soropositividade para \( L. chagasi \) e \( T. cruzi \) aumentou com o tempo, e não houve associação com o gênero. Idade (p<0,007), número de moradores (p<0,05), tipo de piso (p<0,03) e o reconhecimento do vetor (p<0,01) foram associados com a infecção por \( T. cruzi \). Entretanto, na infecção por \( L. chagasi \) foi encontrada associação com a idade (p<0,007) e o melhoramento da habitação (p<0,02). Recomenda-se avaliar o impacto do programa de melhoramento da habitação sobre estas infecções parasitárias nesta comunidade num prazo longo.

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“IN VIVO” AND “IN VITRO” DEMONSTRATION OF HEMOGLOBIN C CRYSTALS IN NON-SPLENECTOMIZED PATIENTS

J.T. de ARAÚJO(1), A.C. BATISSOCO(2) & L. BODEMEIER(3)

SUMMARY

We studied 12 Hb C carriers: 4 homozygotic Hb CC and 8 heterozygotic. We observed the presence of free crystals in the peripheral blood of the homozygotes but in none of the heterozygotes. However, after incubation with 3% NaCl we were able to detect crystals in the heterozygotes (Hb AC and Hb SC), and in the homozygotes (Hb CC). In patient 04 (P04) less crystals formation occurred due to inhibition of the process by the presence of elevated levels of Hb F (12.2%). All the homozygotic patients had a splenomegaly of 3 to 6 fingerbreadths. We believe that the spleen wears off with time, thus allowing the passage of crystals to the peripheral blood. This finding might be associated with splenic insufficiency without a reduction of its dimensions. Finally, the finding of crystals in the peripheral blood permitted the diagnosis of Hb C obviating the need for electrophoresis.

KEYWORDS: Hemoglobin C; Crystal; Non-splenectomized

INTRODUCTION

In 1950 ITANO & NEEL.10 described a new hemoglobin and named it hemoglobin C (Hb C). In 1953 SPAET et al.16 described a second case and soon thereafter RANNEY et al.13 report the third one with the curious observation of many target red blood cells in the patient’s peripheral blood smear.

DIGGS et al.5 in 1954 described intra-erythrocytic crystals in the peripheral blood of Hb C carriers, thus emphasizing that target cells were in reality crystals of Hb C and the presence of free rods identified as crystals also caught his attention.

The main defect of Hb C is the substitution of the glutamic acid residue at position 6 of the N-terminal of the beta chain by a lysine, whilst in sickle cell disease the presence of Hb S is due to the substitution of this residue by a valine.

This phenomenon explains the morphological variations of the erythrocytes: forming crystals in Hb C, and sickle cells in Hb S, thus yielding it possible to identify these two different diseases by simply looking at peripheral blood smears.

In our opinion, the red cells described by RANNEY et al.13 contained Hb C crystals within them and were morphologically different from the ones seen in Hb E, thalassemia, and certain liver diseases which are real target cells. Based on our experience it is possible to identify wether an Hb is of the C or E type simply by analyzing peripheral blood smears using a common optical microscope. In Hb E, thalassemia, and certain liver diseases typical target cells can be seen which differ from the ones observed in Hb C were both intra and extra-erythrocytic crystals are found even with the spleen present. FABRY et al.7, mention that only in smears of splenectomized Hb C patients can the crystals be seen, as these crystals would be all removed by the spleen. Therefore, when splenectomy is performed, the resistance of the splenic network certainly disappears and consequently liberates these crystals. It is also hypothesized that as the splenic network wears off with time, splenic function in Hb C patients diminishes which leads to increasing amounts of the crystals in the peripheral blood with age. In sickle cell disease patients we know that fibrosis followed by splenic involution does occur, many times reducing the spleen to a single nodule. However, in patients homozygotic for Hb C, although a reduction in splenic size does not occur, the progressive decrease in function is a fact that explains the finding of crystals in peripheral blood.

ADACHI & ASAKURA, 19799 studied Hb C crystals using a concentrated fosfate buffer and were able to identify numerous intra and extraerythrocytic crystals of this hemoglobin in the blood of Hb C carriers. FABRY et al.,1981 performed the same study using a 3% NaCl solution, and the resulting crystals were named by ROZENBERG14 as sickling for Hb C, in contrast to red cells containing Hb S.

Although both Hb C and Hb S arise from the African continent, SUTCHARITCHAN et al.13 performing molecular studies, were able to identify the Hb C gene in Thailandes natives, thus demonstrating a non-African origin for Hb C.

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The incidence of Hb C is 17 to 28% in Western Africa, in the vicinities of Northern Ghana. Up to the present date, no reasonable explanation accounts for this high occurrence.

In Brazil, ARAUJO et al., reported an incidence of 0.6 to 1.0% of Hb C by means of crystals demonstration. He also described rare cases of Hb C in Italian and Portuguese descendants.

METHODS AND MATERIALS

During our analysis we identified 4 non-splenectomized carriers of Hb CC, 4 of Hb AC, and also 4 of Hb SC. We studied Hb C crystals in all the carriers, both after incubation with a phosphate buffer and a 3% NaCl solution.

We also analyzed 50 normal blood samples as a negative control for the crystals.

The blood samples were collected with anticoagulant (EDTA), washed 3 times in saline, hemolysed to a 10 g% hemoglobin concentration, and then analyzed by pH 8.6 cellulose acetate electrophoresis, and pH 6.2 agar gel. The fractions were measured after elution, and Hb F was quantitated by the Betke method.

In order to demonstrate the crystals, we used a hypertonic 3% NaCl solution, and also a phosphate buffer with a pH of 7.4 and 1.8M.

The technique consisted of placing one drop of total blood and 2 drops of the 3% NaCl solution in a test tube and then incubating at 37 °C for 4 hours. An identical procedure was performed using the phosphate buffer with a pH of 7.4 and 1.8M.

The blood smears were stained with the Leishman method, and the crystals could be easily identified with the common optical microscope (Figure 1).

RESULTS

In the 50 samples of normal donors (negative controls) no intra or extra-erythrocytic crystals formation was observed.

The highest amount of crystals formation was seen in the homozygotes for Hb C, whilst the lower amount occurred in the heterozygotes for Hb AC. An amount of crystals intermediate between Hb CC and Hb AS was found in the Hb SC cases (Fig. 1 and Fig. 2).

Also, a lower amount of crystals was present in those homozygotes for Hb C that had higher levels of Fetal Hb (Table 1).

Visualization of the crystals was better with the phosphate buffer at pH 7.4 and 1.8M than with the 3% NaCl solution.

The concentration of Hb C measured by electrophoresis was of 30 to 40% for AC heterozygotes and of 40 to 50% for the SC. As for the homozygotes the concentration of Hb C was of 98% with the remaining 2% consisting of Fetal Hb, excepting those cases where there is an increase in Hb F which reaches a concentration of 12.2% (Fig. 3).
The spleen’s clearing function, also known as “culling”, refers to the organ’s capacity of removing aged, dysmorphic, deformed, agglutinated, antibody or complement coated erythrocytes, and erythrocytic corpuscles (or bodies)\(^1\). This is the reason why a normal spleen would remove any free Hb C crystals occasionally present in the circulation. However, even a normal splenic function might permit a few crystals containing erythrocytes to reach the circulation, and this accounts for the finding of these crystals in the peripheral blood of homozygotic Hb C carriers.

When an individual undergoes splenectomy, the peripheral blood may show red cells with crystals similar to sickling forms, free crystals, and also Heinz and Howell-Jolly bodies\(^1\). DIGGS et al.,\(^1\) in 1954, and posteriorly FABRY et al.,\(^1\) 1981\(^1\) demonstrated the presence of Hb C crystals in the peripheral blood of splenectomized and non-splenectomized patients. This is probably due to the fact the splenic network is many times unable to retain all the red cells with crystals, thus permitting their identification in the peripheral blood.

ROZENBERG\(^1\), using photographs, demonstrated the presence of crystals in the peripheral blood of individuals with Hb C, and also confirmed the presence of these in the non-splenectomized carriers of Hb C by means of the 3% NaCl solution technique.

At our laboratory, we routinely examine the peripheral blood smears without prior knowledge of the diagnosis. In all cases, the crystals allowed us to identify the smears as proceeding from Hb C patients, which was further confirmed by electrophoresis. Due to this fact, we concluded that the spleen is unable to retain all the crystals containing red cells or even the free crystals (Figure 2). We believe that over time, as the spleen retains the crystals of Hb C the network becomes insufficient and thus allows the crystals to appear in the peripheral blood.

The spleen undergoes regression in sickle cell disease, in contrast to homozygotic Hb C disease where the spleen remains enlarged, and no regression occurs. This does not mean, however, that damage due to crystals deposition does not occur. In fact, the crystals can be seen in simple X-Rays of the spleen.

Also, lower amounts of crystals are formed when comparing patients with Hb C that present with high levels of Fetal hemoglobin in relation to those with normal levels.

HIRSCH et al.,\(^4\) demonstrated that Hb F inhibits the crystallization of Hb C when compared to Hb A. The same is true for \(\alpha^2\) (\(\alpha^c\delta\)) although to a lesser extent than Hb F (\(\alpha^c\gamma\)). These authors performed a study comparing the inhibitory potentials of the gama (\(\gamma\)) delta (\(\delta\)) chains. These chains both differ from the beta (\(\beta\)) chain at 12 substitution sites, whilst having 4 residues in common located at positions 9, 50, 22, and 87. The two former residues are probably not responsible for the phenomenon, as they involve substitutions exhibiting similar proprieties. Position 22, however, although a common point in both gama (\(\gamma\)) and delta (\(\delta\)) chains, involves different substitutions. The gama (\(\gamma\)) chain possesses an Asp substituting a Glu, whilst at the delta (\(\delta\)) chain the Glu is substituted by a Val. We are left with position 87 were Thr is substituted for Gli in both the gama (\(\gamma\)) and delta (\(\delta\)) chains, and is therefore responsible for the crystallization of Hb C. In addition, this is the same residue that inhibits the polymerization of Hb S by Hb A\(^2\) and Hb F\(^3\). Hb Lepore-Washington originates from a mutation where a fusion between the beta and delta chains occurs, with substitution of only 6 residues. The resulting Hb is also able to inhibit Hb C crystals formation albeit in lesser degrees if compared to Hb A\(^2\) and Hb F. Consequently, this observation suggests that the Gli 87 residue is in fact responsible for the inhibition of Hb C crystals formation. Nevertheless, in agreement with HIRSCH et al.,\(^4\), the varied intensities of inhibition among the different hemoglobins enable us to conclude that there may be two possible explanations: the first is that this variation is probably due to the differences in hemoglobin molecular conformations, and the second is that residue 87 is not the only culprit. In this case, the other residues located between positions 88 and 146 might merit consideration.

Therefore, Fetal Hb with the Gli 87 residue partially inhibits Hb C crystallization, whilst Hb S speeds up the process. This fact definitively demonstrates the existence of intra-erythrocytic crystals in the oxygenated form of the red blood cells of Hb SC individuals. Hb S might speed up crystallization due to the fact the phosphate buffer renders it insoluble\(^6\).

The best visualization of the crystals occurred with the 1.8M 3% NaCl solution technique.
phosphate buffer at a pH of 7.4 using the common optical microscope, due to the fact Hb C is less soluble than Hb A in these conditions.

**RESUMO**

*Cristais de hemoglobina C demonstráveis “in vivo” e “in vitro” em pacientes não esplenectomizados*

Estudamos 12 pacientes portadores de Hb C, sendo 4 homozigotos Hb CC e 8 heterozigotos.

Observamos a presença de cristais livres no sangue periférico dos homozigotos e nenhum nos heterozigotos. Quando incubamos com NaCl 3%, detectamos cristais quer nos heterozigotos, Hb AC e Hb SC, quanto nos homozigotos, Hb CC. No paciente 4 (P04) houve menor formação de cristais devido a inibição pela Hb Fetal que se encontrava aumentada (12,2%).

Todos os pacientes homozigotos mostraram esplenomegalia de 3 a 6 dedos. Emitimos a hipótese de que ocorra no baço um desgaste com o progredir da idade, permitindo a passagem dos cristais para o sangue periférico. Este fato poderia estar associado a uma insuficiência esplênica sem redução do baço.

A presença dos cristais no sangue periférico permitiu o diagnóstico de Hb C mesmo sem a realização da eletroforese de Hb.

**REFERENCES**


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