Production of monoclonal antibodies against *Streptococcus mutans* antigens

*Produção de anticorpos monoclonais contra antígenos de Streptococcus mutans*

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**ABSTRACT:** Several studies have been conducted in the last decades aiming to obtain an anti-caries vaccine, however some studies have demonstrated cross reactivity between *Streptococcus mutans* surface antigens and the human cardiac tissue. In this work, the reactivity of five anti-*Streptococcus mutans* monoclonal antibodies (MoAb) (24A, 56G, C8, E8 and F6) was tested against oral streptococci, cardiac antigens and skeletal and cardiac myosins, aiming to evaluate the specificity of these MoAb. The hybrid producers of immunoglobulins of the IgG\(_{2b}\) class were cloned by limit dilution and expanded \textit{in vivo}. MoAb were tested by ELISA. The hybrid 24A reacted with *S. mutans* CCT 1910, *S. salivarius* CCT 0365 and *S. pyogenes* T23. No reactivity difference was observed among the tested species. Cross reactivity with heart and cardiac myosin was not confirmed and only reaction with myosin of skeletal muscle was observed (\(p = 0.0381\)). The hybrid 56G reacted with all the tested microorganisms and there was statistically significant difference between *S. mutans* and *S. pyogenes* T23 (\(p < 0.001\)). This hybrid also reacted with myosin of skeletal muscle (\(p = 0.0095\)). C8, E8 and F6 presented low reactivity against oral streptococci strains and no reactivity against cardiac antigens. The data of this study showed that the 24A and 56G anti-*S. mutans* MoAb presented reactivity with *S. pyogenes* and *S. salivarius*, reinforcing the occurrence of common antigens between these species. The tested MoAb presented low cross-reactivity with myosin of skeletal muscle, but anti-heart activity could not be confirmed.

**DESCRIPTORS:** Antibodies, monoclonal; *Streptococcus mutans*.

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INTRODUCTION

*Streptococcus mutans* has been considered the main bacterial species involved in the etiology of dental caries. The main surface antigens are the enzymes glucosyltransferase (GTF) that converts sucrose, present in the human diet, into glucans (extracellular polysaccharides) that are responsible for *S. mutans* adherence to teeth. Another important proteinic antigen was denominated I/II, and it works as an adhesin, being important for the adherence mechanism of *S. mutans* to the receptors of salivary acquired pellicle. Immunizations with antigen I/II and glucosyltransferases protected experimental animals against dental caries. Researches in humans, using passive immunization with MoAb for antigen I/II (185 kDa) showed that this procedure might hinder the early implantation of *S. mutans*, delaying the oral colonization by these microorganisms. Unfortunately, some studies demonstrated some degree of cross reactivity between *S. mutans* surface antigens and human cardiac tissues, making not viable a protocol of active immunization.

The studies on *S. mutans* antigens and immunological responses to these components are important to comprehend the anti-dental caries defenses. Considering this, analysis of the specificity of monoclonal antibodies may help in the selection of interesting antigens for an anti-caries vaccine.

MATERIAL AND METHODS

Production of anti-*S. mutans* monoclonal antibodies

**Cell culture**

Cells of the lineage Sp2-Oag14 (myeloma cells) and the hybrids obtained were cultivated in RPMI-1640 medium (Sigma, Baden-Wuttemberg, Steinheim, Germany) containing 1 mM L-glutamine (Sigma, Baden-Wuttemberg, Steinheim, Germany), 10 mM Heps (Sigma, Baden-Wuttemberg, Steinheim, Germany), 24 mM sodium bicarbonate (Sigma, Baden-Wuttemberg, Steinheim, Germany), 40 mg/ml gentamicin sulphate (Shering, Rio de Janeiro, Rio de Janeiro, Brazil) and supplemented with 10% fetal bovine serum (Sigma, Baden-Wuttemberg, Steinheim, Germany). Both were maintained in humid atmosphere with 5% CO₂ and at 37°C.

**Fusion protocol**

The protocol proposed by Goding (1980), with modifications, was followed. Female Balb/c mice were immunized intradermically in the tail basis with 200 µg of *S. mutans* cytoplasmatic antigen (CAg) in Freund’s complete adjuvant, and then restimulated by intraperitoneal via after 21 days with the same antigen in saline solution buffered with phosphate (PBS - Sigma, Missouri, St. Louis, USA). Three days after the booster, the animals were sacrificed and had their spleen removed. This organ was macerated and its cells (10⁶) were fused with the lineages Sp2-Oag14 (5 × 10⁶), using 45% polyethylene glycol (PEG) (Sigma, Baden-Wuttemberg, Steinheim, Germany) as a fusion agent and 10 ml of RPMI. Then, the suspension was cultured for 10 minutes. Polyethylene glycol was eliminated by centrifugation and the cells were resuspended in RPMI medium supplemented with hypoxanthine, aminopterin and 20% thymidine (HAT) (Sigma, Baden-Wuttemberg, Steinheim, Germany). Hybrids were distributed at a 1 × 10⁶ ratio on the feeder layer, prepared the previous day with the spleen of a non-immunized animal (1.6 × 10⁵), feeding them each 5 days. After seven days, the cell culture was examined with the aid of an inverted microscope (Nikon, model 104; Melville, New York, USA). After 10 days, the HAT medium was replaced by 20% HT (hypoxanthine and aminopterin) (Sigma, Baden-Wuttemberg, Steinheim, Germany), and after 12 days the points presenting confluent growth were tested by the ELISA technique for the antigen used in the immunization. The positive points were cloned in the common ratio of 1/2 and the hybrids of interest were expanded *in vitro* and *in vivo* in Balb/c mice.

Antigen obtaining

*S. mutans* CCT 1910 was grown in triptic soy broth (TSB, Difco, Detroit, USA) at 37°C for 24 hours with 5% CO₂, killed by 0.075% formaldehyde and then maintained for 18 hours at 4°C. The suspension was centrifuged and the cells were washed in 125 mM 2,3 dibromopropyl-chlorhydric acid (Tris-HCl, Life Technologies, New York, USA) pH 7.5, 10 mM EDTA, and resuspended in 50 ml of the same buffer. After that, 5 mM phenylmethyl sulphonyl fluoride (PMSF, Sigma, Steinheim, Germany) was added and the suspension was agitated vigorously with glass beads overnight at 4°C. The lysisate was centrifuged and the supernatant was dialysed against distilled water, concentrated and lyophilized, constituting the cytoplasmatic antigen (CAg). The proteinic content was determined by the Bradford (1976) method.
Streptococci whole cells preparation

*S. mutans* (CCT 1910) cells, *S. salivarius* (CCT 0365) and *S. pyogenes* (CCT 1500 and T23) were grown in triptic soy broth (TSB, Difco, Detroit, USA) for 24 hours and killed by 0.075% formaldehyde. After a wash step in PBS and 10,000 g centrifugation at 25°C for 20 min, the suspensions were standardized by spectrophotometry (595 nm and optical density of 0.200).

Self-antigen preparation

Myosin from pig cardiac muscle (Sigma, Baden-Wuttemberg, Steinheim, Germany) and from mice skeletal muscle (6 mg/ml), kindly donated by Dr. Ternynck (Pasteur Institute, Paris), were included in the study.

Heart extract preparation

Twenty Balb/c mice were sacrificed and had their hearts removed. They were washed in PBS, macerated and filtered in sterile gauze. The filtrate was maintained at –20°C. The pellet was transferred to 5 ml of a buffer solution containing 150 mM Tris-HCl, 6 M urea, 20 mM 2-mercaptoethanol and 1% Tween 20 (Merck, Hessen, Darmstadt, Germany). The mixture was boiled for five minutes and then maintained under refrigeration for 24 hours. It was then centrifuged and the supernatant was dialysed against PBS overnight. Both were concentrated by ultra-filtration. The proteinic concentration was estimated in 30 µg/µl by the Bradford (1976) method.

ELISA technique

Polystyrene plates (number 3590, Costar, Cambridge, MA, USA) were coated with whole streptococci cells, with myosin (5 µg/ml) and heart extract (10 µg/ml) in 0.1 M carbonate buffer (pH 9.6) and incubated for 2 hours at 37°C and for 18 hours at 4°C. The wells were blocked with 0.5% gelatin (Merck, Hessen, Darmstadt, Germany) in phosphate buffered saline (PBS-G) for 30 minutes at 37°C. Then the plates were washed with 0.5% Tween 20 PBS (T-PBS). The supernatants of the culture and/or ascites diluted in T-PBS-G were added to the wells, in duplicate, and diluted in the common ratio of 1/2. Plates were incubated for 2 hours at 37°C, and maintained at 4°C overnight. After an additional wash step with T-PBS, 50 µl of anti-mice IgG peroxidase-labelled (Sigma, Baden-Wuttemberg, Steinheim, Germany) were added to the wells (1 µg/ml) and incubated for 1 hour at 37°C. Finally, 100 µl/well of o-phenylenediamine (Merck, Hessen, Darmstadt, Germany) in 0.1 M citrate buffer (pH 5.5) (Merck, Hessen, Darmstadt, Germany) were added at room temperature until a yellow color developed. The reaction was stopped with 2.5 M sulfuric acid (Merck, Hessen, Darmstadt, Germany) and the color was measured at 490 nm with a model 3550 reader (Bio Rad Laboratories, California, Hercules, USA). Three experiments, in duplicate, with their respective controls (Irrelevant MoAb 17C) were performed. The concentration of 5 µg/ml was elected for statistical analysis, as this value was the mean point of the curves.

Statistical analysis

The data obtained were analyzed by Mann-Whitney and Kruskal-Wallis tests, at the level of significance of 5%.

RESULTS

MoAb production by hybrids

All the points with optical densities obtained by ELISA higher than 0.300 were considered positive. This value corresponds to at least three times the mean reactivity of a control serum of mice diluted at 1:100.

First fusion

Twenty-seven hybrids (4.6% of the plated points) presented confluent growth in six plates of 96 wells (576 points) in the first reading, and three points (11.1%) were positive for *S. mutans* in ELISA. Among these hybrids, only two continued to produce IgG antibodies after initial cloning and were recloned by limit dilution. The hybrids that produced antibodies were denominated 24A and 56G.

Second fusion

Sixty-six hybrids (11.4% of the total of plated points) were obtained, seventeen produced antibodies against *S. mutans* and only three (4.5%) maintained the production of IgG (C8, E8 and F6).

Reactivity of MoAb in ELISA

C8, E8 and F6 presented low reactivity against oral streptococci strains (data not shown). Only the antibodies produced by 56G and 24A showed reactivity against heart antigens, skeletal and car-

...cardiac myosins. Graphs 1 and 2 show the reactivity of MoAb 24A and 56G to the different species of streptococci and to the auto-antigens.

Graph 1 shows that the reactivity levels of MoAb 24A to *S. mutans*, *S. salivarius* and *S. pyogenes* T23 were significantly higher when compared to that of the irrelevant MoAb 17C (p = 0.0190, p = 0.0095, p = 0.0095, respectively), but no significant difference was found for *S. pyogenes* CCT 1500 (p = 0.9143). No differences in the reactivity of MoAb 24A to the different species of streptococci were found. Regarding the reactivity of 56G, significant levels were observed against all the tested species (p = 0.0095 in all cases). The values were compared to verify if the antibody might discriminate *S. mutans* among the tested species, and statistically significant difference was observed only between *S. mutans* and *S. pyogenes* T23 (p < 0.001).

Regarding cross reactivity, Graph 2 shows that the level of significance in the case of MoAb 24A was reached only with skeletal muscle myosin (p = 0.0381). Considering the heart antigen and cardiac myosin, the reactivity of MoAb 24A was not statistically significant. The most marked mean reactivity for the heart did not present statistical significance; however, an elevated standard deviation was observed due to the great variability among the results of the analysed strains. It could be observed that MoAb 56G also reacted significantly with skeletal muscle myosin (p = 0.0095), even though not markedly. The values obtained for the reactions with heart and cardiac myosin were not statistically significant.

**DISCUSSION**

The MoAb obtained (56G, 24A, C8, E8 and F6) presented reactivity in ELISA with whole cells of three species of streptococci (*S. mutans*, *S. salivarius* and *S. pyogenes*). These results are in accordance with those of the study of Krisher, Cunningham (1985) that produced a MoAb for *S. pyogenes* with cross reactivity to *S. mutans* and myosin of skeletal muscle. Adsorption of the ascite with M protein of *S. pyogenes* resulted in loss of this cross reactivity. The same occurred when the antibody was adsorbed with *S. mutans*, indicating the existence of homologous epitopes between these species.

MoAb 56G and 24A exhibited another particularity when compared with C8, E8 and F6. Both were reactive, even with low intensity, with self-components. In previous studies we demonstrated that mice immunized with CAg of *S. mutans* synthesized antibodies that were reactive to myosin of skeletal and cardiac muscles. van de Rijn et al. (1976) also noticed this phenomenon in rabbits immunized with antigens from several *S. mutans* strains that produced polyclonal antibodies that presented cross reactivity.
reactivity with cardiac tissue and skeletal muscle. Also, they observed that the adsorption of rabbit serum with sarcolemma extracts, S. mutans and A group β-haemolyticus streptococci removed completely the reactivity with mammalian tissues.

Since then, several studies demonstrated in a convincing way that antibodies with cross reactivity were present in the sera of immunized animals and in supernatants of unique clones, in the case of hybridomas, always associated to immunization with streptococci and infection by these microorganisms.\textsuperscript{3,7,13,14} In all the studies found in literature, the possibility that these reactions may be induced by rheumatoid factors (anti-immunoglobulin antibodies) was questioned. In fact, in the first attempts of hybridisation, ten antibodies reactive to streptococci were obtained. However, in all these cases no reactivity was detected by Western Blot. At least half of these antibodies presented activity of rheumatoid factor (data not shown). Russel\textsuperscript{13} (1987), after hyperimmunization of rabbits with whole cells of S. mutans, S. sobrinus, S. rattus and S. mitis, observed that the cross reactivity of serum auto-antibodies was not always associated to antigen I/II, the main S. mutans surface antigen. This author concluded, with the aid of a combined ELISA technique and gel diffusion, that the antibodies reactive to cardiac tissue did not bind to antigen I/II. Then, there were clearly two distinct populations of antibodies with different specificities. The author also observed an elevated level of IgM rheumatoid factor that might be responsible for the reactivity to cardiac tissue observed in previous studies.

In our results, anti-heart reactivity of the tested MoAb could not be confirmed by the statistical tests. The reactivity of these antibodies were of low intensity (as in 24A) or highly specific for S. mutans (56G). Also, both of them were IgG\textsubscript{2a} isotypes. This result was expected as Leão\textsuperscript{et al.}\textsuperscript{10} (2000) previously demonstrated an increase of this subclass and also of IgG\textsubscript{2a} after immunization of mice with S. mutans cytoplasmatic antigen.

The data of this study did not exclude totally the reactivity to self-components. As evidenced by ELISA, 24A and 56G presented some degree of reactivity to skeletal muscle myosin. However, it is important to clarify that this reactivity was not intense, and that no experiment of adsorption was performed with the aim of eliminating it. And the hypothesis of reactivity induced by natural auto-antibodies described by other researches for explaining the reaction between polyclonal antibodies and host tissues may not be discharged. These antibodies have been related to the pathogenesis of auto-immune diseases.\textsuperscript{1,6}

Common antigens occur frequently in a bacterial genus, making it difficult to find species-specific monoclonal antibodies. The reactivity between S. mutans and S. pyogenes has been related in the literature and is considered by some authors as an additional factor in the development of rheumatic fever. S. mutans and S. pyogenes present similar antigens in the cell surface, and S. mutans may reach the blood circulation after dental procedures. This relation may constitute a constant antigenic stimulation in the development of rheumatic fever.\textsuperscript{5} This sharing of antigens between both species was reinforced in this study, considering that MoAb against S. mutans recognized superficial epitopes in S. pyogenes.

The monoclonal antibodies 24A and 56G isolated in this study (specially the 56G) will be valuable in future experiments, mainly in the fields of epidemiology and diagnosis of caries risk in children.

**CONCLUSIONS**

The isolated MoAb reacted in ELISA with whole cells of S. mutans. C8, E8 and F6 presented low reactivity against oral streptococci strains and no reactivity with self-components. The MoAb anti-S. mutans 56G and 24A recognized S. pyogenes and S. salivarius. Although anti-heart reactivity of MoAb 56G and 24A was not observed, there was reactivity with self-components (myosin of skeletal muscle).

**REFERENCES**