Nasal polyposis is a severe, chronic inflammatory condition of the paranasal sinuses and is frequently associated with asthma and aspirin sensitivity. Mesenchymal stem cells exhibit a potent immunosuppressive effect in several inflammatory conditions, and their role in nasal polyposis remains little explored. Hence, we investigated whether bone marrow-derived mesenchymal stem cells could modulate cell phenotype in the nasal polyp milieu. After coculture with mesenchymal stem cells, the frequency of these inflammatory cells was found to decrease. Furthermore, mesenchymal stem cells promoted strong inhibition of CD4+ and CD8+ T cell proliferation, increased the frequency of CD4+CD25+Foxp3 T cells, and changed the global cytokine profile from an inflammatory to an anti-inflammatory response. We believe that mesenchymal stem cells may be a very useful adjunct for investigation of the inflammatory process in nasal polyposis, contributing to better understanding of the inflammatory course of this condition.

1. Introduction

Nasal polyposis (NP) is a severe, chronic inflammatory condition of the paranasal sinuses, with a prevalence ranging from 1% to 4% in the general population [1]. It is frequently associated with asthma and aspirin sensitivity [1]. NP in combination with aspirin-induced asthma (AIA) represents the most severe form of airway inflammation [2].

NP is characterized by overgrowth of nasal mucosa caused by the influx of a variety of inflammatory cells. The inflammatory process characteristic of NP is defined mainly by T-cell activation and arrest of regulatory T-cell function, with a decrease in Foxp3 expression and concomitant upregulation of T-bet and GATA-3 levels [3]. The predominance of T-effector cells in polyp tissue is closely associated with patient ethnicity. In white European patients a Th2-driven response is predominant, whereas in Chinese patients, a Th1/Th17-driven response has been demonstrated [4]. However, little is known about the inflammatory milieu of nasal polyposis, and understanding of this process can play an important role in defining the course of the disease.

The most important features of NP concern its unique remodeling process, which is characterized by low production of transforming growth factor-β (TGF-β) associated with a lack of collagen as compared with healthy subjects [5, 6]. In NP, matrix metalloprotease-7 (MMP-7) and matrix metalloprotease-9 (MMP-9) levels are increased and tissue inhibitor of metalloproteinases 1 (TIMP-1) levels are, inversely, decreased as compared with normal nasal mucosa [7]. This imbalance can be partially explained by the lack of
TGF-β1 in NP and its inhibitory effect on MMP-9 activity via TIMP-1 release [8]. TGF-β1-activated PAI-1 (plasminogen activator inhibitor-1) is also found decreased in NP, leading to an increase in the levels of plasminogen activator and MMP levels when compared with controls [6, 9].

Due to the similarities between upper and lower airway mucosa, the last one has been used as a model to understand NP [10]. It is demonstrated an epithelium disruption in nasal tissue less extensive than in the lungs [11], so to repair the injured epithelium less TGF-β is produced in nasal mucosa when compared to bronchial mucosa [12]. Supporting these findings it is reported that the basement membrane in nasal mucosa has limited pseudothickening with significant less elastase positive cells comparatively to bronchial mucosa [11].

Furthermore, histological examination of biopsy specimens shows a soft tissue with clear lack of extracellular matrix [13], major edema, albumin-filled pseudocysts, and alpha-2-macroglobulin [14].

Multipotent stromal cells or mesenchymal stem cells (MSC) are adult, adherent, nonhematopoietic stem cells with the ability to differentiate into several mesenchymal cell lines (chondrocytes, adipocytes, and osteocytes) beyond to promote a prominent immunomodulatory effects on inflamed environment. MSCs retain low immunogenicity and exert immunosuppressive effects in allogeneic transplantation [15]. These cells exhibit reduced expression of both major histocompatibility complex (MHCs) and costimulatory molecules (CD80, CD86, and CD40) and have emerged as a very useful tool for therapeutic use, including in regenerative medicine and tissue bioengineering [16, 17].

MSCs have been used in experiments involving a very broad range of diseases, including repair of acutely injured tissue, chronic diseases, graft rejection, and autoimmune conditions [18]. Such widespread use of these cells is based on their distinct natural properties, namely, stromal cell differentiation, soluble factor secretion stimulating hematopoiesis, ECM maintenance, and immunoregulatory effects [19]. The immunomodulatory role of MSCs has been demonstrated in many in vitro and in vivo studies and consists essentially of downmodulation of the inflammatory process, inhibiting T-cell, B cell, NK cell, and APC cell proliferation via a paracrine secretory mechanism [20–22].

Most soluble factors produced by MSCs are associated with their immunoregulatory properties, including TGF-β1, prostaglandin-E2 (PGE-2), IDO, IL-10, IL-6, MMP, and TIMP [17], and could interact with severely inflamed tissues (such as the NP environment) to restore a balanced T cell-mediated response. Hence, we hypothesized that MSCs could be able to modulate allergic inflammation in the context of nasal polyps.

Using MSCs as an immunosuppressive tool, we first characterized the infiltrating polyp-derived cells and found that, after coculture with MSCs, the frequency and activation of inflammatory cells had changed. We also observed that MSC-promoted modulation of the cytokine profile, inhibition of T-cell proliferation, and expansion of CD4+CD25+Foxp3 cells in an in vitro assay.

### Table 1

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2. Material and Methods

2.1. Patients and Clinical Diagnosis. Nasal polyp tissue samples of 12 patients with known NP were obtained during functional endoscopic sinus surgery (FESS) performed at the Department of Otorhinolaryngology, University of São Paulo, Brazil. The study was approved by the local Research Ethics Committee and written informed consent was obtained from each patient before sample collection. The diagnosis of NP was based on medical history, clinical examination, nasal endoscopy, and computed tomography (CT) of the paranasal sinuses according to the European Position Paper on the Primary Care Diagnosis and Management of Rhinosinusitis and Nasal Polyps 2012 [23].

All subjects underwent a skin prick test for common inhalant allergens. The diagnosis of asthma was obtained from the Department of Pulmonology at the University of São Paulo. Information on aspirin intolerance was collected from patient histories (Table 1).

2.2. MSC Isolation and Preparation of Nasal Polyp Single-Cell Suspension. Human mesenchymal stem cells (MSCs) were isolated by rinsing of the cells remaining in bone marrow collection tubes, kindly provided by the Children’s Hospital of São Paulo and GRAACC (Support Group for Children and Adolescents with Cancer), with ethical approval and the informed consent of the donors (Protocol no. 45/09). The tubes were washed with PBS, cells were isolated by the Ficoll-Hypaque protocol (Sigma, USA), and culture was performed as previously described by Lennon and Caplan (2006) and Pittenger et al. (1999) [24, 25]. The cells were cultured in basal medium containing DMEM-low glucose medium (Gibco, USA) supplemented with 10% HyClone fetal bovine serum (Thermo, USA), antibiotic (100 U/mL penicillin and 100 μg/mL streptomycin, Gibco), and 100 mM of nonessential amino acids (Gibco). The MSCs were subcultured for 10 passages and then used for coculture experiments with polyp-derived cells. All polyp-derived cells were isolated from polyp tissues by mechanical separation (with surgical scissors and forceps) followed by digestion in collagenase IV (1 mg/mL, Sigma) for 50 minutes at 37°C. The cells were then washed.
in complete medium (10% serum), filtered in a 70μm cell strainer (BD Biosciences, USA), suspended, and seeded onto six well plates containing R10 medium: RPMI-1640 culture medium (Gibco), supplemented with 10% FBS (Gibco) and antibiotic (100 U/mL penicillin and 100 μg/mL streptomycin; Gibco). During coculture assays, both MSCs and polyderived cells were incubated at 37°C in a 5% CO₂ atmosphere for 72 hours.

2.3. Phenotype and Differentiation Capacity of MSCs. To characterize MSC phenotype, the cells were harvested by treatment with trypsin (Gibco), washed, and suspended in phosphate-buffered saline (PBS), and approximately 1 × 10⁵ cells were incubated with conjugated monoclonal antibodies (1:100) against CD73, CD90, CD105, CD45, HLA-DR, PDL-1, PDL-2, and CTLA-4 (conjugated with PE, FITC, or PerCP fluorochrome; BD Biosciences). The FACSCanto II flow cytometer (BD, Biosciences) was used for acquisition, and analysis was performed using the FlowJo software, version 7.2.4 (TreeStar, USA). To assess the ability to differentiate into several mesenchymal cell lines, the cells were subjected to adipogenic, chondrogenic, and osteogenic differentiation and analyzed by the FACSCanto II flow cytometer (BD, Biosciences). The results were presented as cell frequency.

2.4. Cellular Phenotyping of Nasal Polyp-Derived Tissue. To determine the immune cell profile of whole polyp cells, fresh and cultured (for 72 hours) cells were harvested (both adherent and nonadherent) without enzymatic dissociation and labeled with a set of various specific fluorescent antibodies, such as CD4-Pacific Blue, CD8-PECy7, CD11c-PE, CD14-PerCP-Cy5.5, CD19-Violett, CD25-APC, CD40-FITC, CD69-FITC, and NK1.1-PE (all from BD Biosciences). These sets of antibodies were used for both fresh and in vitro coculture analysis. To investigate the CD4+CD25+Foxp3+ T lymphocyte profile, intracellular staining for Foxp3 expression was performed after 72 hours in culture with or without the presence of MSCs. The cells were fixed and permeabilized using the Fix/Perm Buffer Set Kit (BD Biosciences). Staining was performed using Foxp3 (PE), CD4 (APC-Cy7), and CD25 (APC) antibodies at 1:100 dilution (BD Biosciences). Analysis was determined by flow cytometry (FACSCanto II cytometer, BD Biosciences), within FSC and SSC gate using histograms for the T-cell type of interest (CD4+ or CD8+). Phytohemagglutinin A (1μg/mL) (PHA, Invitrogen, USA) was used as a polyclonal-positive stimulus.

2.5. T-Cell Proliferation Assay. To determine the ability of MSCs to modulate the polyp microenvironment, fresh whole polyp cells were labeled with 5μM carboxyfluorescein succinimidyl ester (CFSE, BD Biosciences) and seeded in culture with or without MSCs for 5 days. MSCs from nonrelated donors were cocultured with 1 × 10⁵ whole polyp cells in a 1:3 ratio in R10 medium supplemented with 100 mM of vitamin complex (Thermo), antibiotic (Pen/Strep, 100 U/mL, Gibco), 100 mM L-Glutamine Mixture (Gibco), 100 mM MEM nonessential amino acids (Gibco), and 1 mM HyClone sodium pyruvate (Thermo). After 5 days, all polyp cells were stained with anti-CD4 and anti-CD8 antibodies (1:100 ratio, BD Biosciences), and CFSE dilution was read in FITC channel by flow cytometry analysis (FACSCanto II cytometer, BD Biosciences). This analysis was performed within the FSC and SSC gate using histograms for the T-cell type of interest (CD4+ or CD8+). Phytohemagglutinin A (1μg/mL) (PHA, Invitrogen, USA) was used as a polyclonal-positive stimulus.

2.6. Cytokine Profile of Nasal Polyp-Derived Cells. Isolated polyp culture and coculture (MSC and polyps) supernatants (from 6 out of 12 NP samples) were further harvested after 72 hours for cytokine quantification, using the Cytometric Bead Array (CBA) Th1/Th2/Th17 Cytokine Kit, according to manufacturer recommendations (BD, Becton Dickinson Biosciences). Acquisition was performed in the FACSCanto II cytometer (BD Biosciences) and the samples analyzed using FCAP Array software v3.0 (Soft Flow Inc., HUN).

2.7. Statistical Analysis. Results were assessed for normal distribution by the Kolmogorov-Smirnov test. Categorical variables were expressed as percentages (%), and continuous variables (data) were presented as means ± standard deviations. The Mann-Whitney nonparametric test was used to assess between-group differences. For analysis among three or more groups, the Kruskal-Wallis nonparametric test for one-way analysis of variance was used. For all analyses, a P value ≤ 0.05 was considered statistically significant. All statistical testing and plotting were performed using GraphPad Prism 5 (GraphPad software, Inc., USA).

3. Results

3.1. Characterization of Mesenchymal Stem Cells and Polyp Infiltrating Cells. The bone marrow-derived mesenchymal stem cells used in this study exhibited most important characteristics of MSCs, such as a plastic-adherent growth pattern (Figure 1), fibroblast-like morphology (Figure 1(a)), expression of specific surface antigens (CD105, CD54, CD90, and CD73), and immunoregulatory molecules (CTLA-4, PD-L1, and PD-L2), and no expression of the immunogenic and hematopoietic surface markers HLA-DR and CD45, respectively (Figure 1(k)). Additionally, MSCs demonstrated the ability to differentiate into adipocytes (Figures I(c)-(i)), chondrocytes (Figure 1(f)), and osteocytes (Figure 1(h)) in vitro. The polyp-derived cells spread from the whole polyp in culture and exhibited spheroid-like morphology (Figures 1(i) and 1(j)).
Figure 1: Characterization of bone marrow-derived mesenchymal stem cells and polyp-derived cells. (a) MSCs exhibiting fibroblastic-like morphology in vitro; (b) MSC culture in adipogenic control medium showing absence of lipid vesicles; (c) MSCs containing lipid vesicles (black arrow) in adipogenic differentiation medium without Oil Red stain; (d) MSC culture in adipogenic differentiation medium stained with Oil Red (black arrow); (e) MSC culture in chondrogenic control medium; (f) MSC culture in chondrogenic differentiation medium stained with Safranin O; (g) MSC culture in osteocyte control medium; (h) MSC culture in osteogenic differentiation medium stained with Alizarin Red; ((i)-(j)) representative view of polyp-derived cells spreading and growing in culture (black arrow); and (k) immunophenotypic signature of MSCs in culture, demonstrating absence of hematopoietic (CD45 and CD34) and immunogenic (HLA-DR) markers and expression of CD105, CD90, CD73, and CD54, as well as immunoregulatory receptors such as PDL-1 and -2 and CTLA-4.

The immunophenotype of whole NP-derived cells showed the presence of distinct immune cells after single-cell suspension. NK cells, T cells, dendritic cells, monocytes, and B cells were present (Figures 2(a)–2(h)). The frequency of each subtype of immune cells was as follows: NK cells, 10.60 ± 6.28%; CD4+, 11.26 ± 4.70%; CD4+CD69+, 6.71 ± 4.31%; CD4+CD25+, 6.77 ± 4.35%; CD8+, 11.19 ± 3.49%; CD8+CD69+, 6.73 ± 5.09%; CD11c+, 14.17 ± 4.83%;
We found a significant decrease in the frequency of most inflammatory cells in NP-derived tissue, based on the fact that MSC had promoted a decrease in the frequency of B cells, monocytes, and dendritic cells, as well as cytotoxic T cells (CD8+) showed a significant decrease in frequency when cocultured with MSC; however, no changes in the activated T cell compartment (CD4+CD69+ and CD8+CD69+) were observed (Figures 3(a)–3(d)). The frequency of B cells, monocytes, and dendritic cells also tended to decrease after coculture with MSCs. De novo, the fraction of activated cells included in the monocyte and dendritic cell subpopulations was not altered in comparison with cultures performed in the absence of MSCs (Figures 3(e)–3(g)).

3.2. Effect of MSCs on Nasal Polyp-Derived Cells. To evaluate the role of MSCs in the modulation of the NP microenvironment, we carried out a coculture assay for 72 hours and assessed the frequency of poly-derived infiltrating cells. We found a significant decrease in the frequency of most inflammatory cells. NK cells (NK1.1+), T helper cells (CD4+), monocytes 20%, and cytotoxic T cells (CD8+) showed a significant decrease in frequency when cocultured with MSC; however, no changes in the activated T cell compartment (CD4+CD69+ and CD8+CD69+) were observed (Figures 3(a)–3(d)). The frequency of B cells, monocytes, and dendritic cells also tended to decrease after coculture with MSCs. De novo, the fraction of activated cells included in the monocyte and dendritic cell subpopulations was not altered in comparison with cultures performed in the absence of MSCs (Figures 3(e)–3(g)).

3.3. Effect of MSCs on the Expansion/Proliferation Capacity of the T-Cell Compartment in Nasal Polyp-Derived Cells. On the basis of the fact that MSC had promoted a decrease in the frequency of most inflammatory cells in NP-derived tissue,
we sought to investigate whether MSCs could promote the functional ability to arrest NP-derived T-cell proliferation in vitro. Surprisingly, we observed significant inhibition of expansion/proliferation of both T helper cells (CD4+) and cytotoxic T cells (CD8+) derived from NP stimulated with PHA in compared with cells stimulated but cultured in the absence of MSCs (Figures 4(a) and 4(b)). The immunosuppressive effect of MSCs on NP-derived T cells was evident, considering that the index of proliferation of CD4+ and CD8+ T cells cocultured with MSCs did not differ from that of control unstimulated T cells.

The immunoregulatory effect of MSCs is almost always associated with an expansion of regulatory T cells (CD4+CD25+Foxp3+). Thus, in the present work, we investigated the presence of CD4+CD25+Foxp3+ T cells in culture with or without the presence of MSCs. A significant increase in

**Figure 3:** Bone marrow-derived mesenchymal stem cells modulating the polyp microenvironment in vitro. (a) NK cell (NK1.1+) frequency alone and in coculture with MSCs; (b) CD4+ and CD8+ frequencies alone and in coculture with MSCs; (c) CD8+ and CD69+ frequencies alone and in coculture with MSCs; (d) CD4+ and CD69+ frequencies alone and in coculture with MSCs; (e) B cell (CD19+) frequency alone and in coculture with MSCs; (f) CD11c+ and CD11c+CD40+ dendritic cell frequencies alone and in coculture with MSCs and CD14+ and CD14+CD40+ monocyte frequencies alone and in coculture with MSCs. The presence of MSCs may immunomodulate the phenotype of polyp-derived cells toward an immunosuppressive profile.
3.4. Effect of MSCs on the Cytokine Profile of Nasal Polyp-Derived Cells. Finally, to elucidate the mechanism underlying the immunosuppressive effect of MSCs in NP, we analyzed six NP samples using the Cytometric Bead Array Technique. Analysis of culture supernatants showed a number of CD4+CD25+Foxp3+ T cells was observed in the presence of MSCs as compared with cultures of T cells alone (Figure 4(c)), suggesting that these cells can play an essential role in the development of the chronic inflammatory process in NP.
prominent shift from an inflammatory to an anti-inflammatory cytokine profile. In coculture, MSCs promoted an increase in anti-inflammatory molecules such as IL-10, with a concomitant decrease in inflammatory cytokines such as IL-2, TNF-α, and IFN-γ (Figure 5).

4. Discussion

Nasal polyposis is characterized by the most severe upper-airway inflammatory process observed in clinical practice. This process is crucial to understand the mechanisms that underlie development of polyposis. It is known that MSCs have immunomodulatory properties, as demonstrated in some organs such as the brain, kidney, heart, bone, and lung [26–30]. MSC-mediated immunomodulation can occur via cell-to-cell contact or by release of soluble factors, which are associated with many regulatory effects of these cells in a tissue inflammation context. The best documented MSC-secreted cytokines are TGF-β1, PGE-2, IDO, IL-10, IL-6, MMP-2,9, TIMP-2,3, nitric oxide (NO), chemokine ligand 2 and 5 (CCL2,5), human leukocyte antigen 5 (HLA-G5), heme oxygenase-1 (HO-1), hepatocyte growth factor (HGF), and leukemia inhibitory factor (LIF) [17]. On the basis of their properties, MSCs have been explored in a wide range of experiments and have been used for therapeutic purposes more extensively than any other subtype of stem cell. These cells also retain further important features, such as low immunogenicity, and promote inhibition of proliferation/activation in allogeneic lymphocytes [18].

Thus, in the present study, we assessed the impact of MSCs on modulation of the inflamed NP microenvironment. Firstly, we demonstrated that several types of inflammatory cells (NK cells, B cells, T cells, monocytes, and dendritic cells) are found in this milieu. After characterization of MSC phenotype and differentiation, these cells were cultured with polyp-derived cells, and a direct immunomodulatory effect on the inflammatory NP cell compartment was observed. There was a significant decrease in the frequency of CD4+, CD8+, CD14+, and NK cells, as well as a significant increase in CD4+CD25+Foxp3+ T cells, when MSCs were cocultured with NP-derived cells. A decrease in regulatory T cells has been described as a feature of the NP disease course. This is an important finding, and the increase in CD4+CD25+Foxp3 cells induced by presence of MSCs in the present study may help in our understanding of the progression of NP. Functionally, we observed that MSCs inhibited both CD4+ and CD8+ polyp-derived T cell proliferation and efficiently changed the local inflammatory pattern, promoting a shift from an inflammatory to an anti-inflammatory cytokine profile.

The stem cells actions are widely dependent on the disease and time of therapy. Contact-dependent immunosuppression is one mechanism of MSC action and can be associated with the expression of immunoregulatory molecules expressed on the surface of MSCs or by delivering microvesicles carrying bioactive molecules [31]. CTLA-4, PDL-1, and PDL-2 were present in the MSCs used in this study, and their role in inhibition of proliferation and activation of T lymphocytes

Figure 5: In vitro cytokine profile of polyp-derived cells cultured in the presence or absence of bone marrow-derived mesenchymal stem cells. A decrease in inflammatory cytokines (IL-2, TNF-α, and IFN-γ) and enhancement of anti-inflammatory cytokine (IL-10 and IL-6) levels was observed when MSCs were cocultured with polyp-derived cells.
has been reported extensively in the stem cell literature [32, 33].

MSC is also capable of exerting influence on inflamed process via paracrine action [34] even far from the injured tissue. The soluble factors released by MSCs are another mechanism that may also have influenced modulation of the polyderyed cell microenvironment. Many studies have demonstrated the effect of TGF-β secreted by MSCs on suppression of peripheral blood mononuclear cell (PBMC) proliferation [35]. TGF-β is also capable of increasing the frequency of regulatory T cells, especially when associated with PGE2 [36–38]. In a model of asthmatic mice, TGF-β secreted by MSCs showed a beneficial effect by decreasing levels of IL-4, IL-5, IL-13, and immunoglobulin in bronchoalveolar lavage fluid [38].

Although the role of TGF-β on remodeling process is still controversial in the literature, there is a strong evidence in NP that the lack of TGF-β is involved in decreasing extracellular matrix formation [5, 6, 39] highlighting the importance of this molecule on the development of inflammation in NP.

In addition, PGE2 is another molecule which is found in low levels in NP tissue [40] and MSCs produce PGE2. PGE2 exerts an immunosuppressive action on T cells and, consequently, promotes a decrease in IFN-γ and TNF-α levels [41, 42]. Our study corroborates these findings, demonstrating decreased IFN-γ and TNF-α expression in NP cells cocultured with MSCs in comparison with cultures of NP-derived cells alone; maybe PGE2 is contributing to decrease these interleukins.

In mice, macrophages from septic lungs produced higher levels of IL-10 when treated with MSCs than untreated macrophages. The authors suggested that the EP2 and EP4 receptors (prostaglandin receptors) were responsible for this increase in IL-10 production [43]. In this context, IL-10 is considered to be the main immunosuppressive interleukin, and we found higher levels of IL-10 in NP cells treated with MSCs than in untreated cells.

Consistently with what is found in a variety of other diseases [44, 45], the presence of MSCs in NP cell cultures increased the expression of IL-6 (an interleukin mainly produced by Th1 cells). Paradoxically, although IL-6 is associated with increased production of IL-2 [46], we found decreased levels of this inflammatory interleukin in NP cells treated with MSCs. One plausible explanation is the fact that MSCs can secrete TGF-β and IL-6 and that differentiation of Treg/Th17 depends on the proportion of these two cytokines [47].

In the present study, we did not detect IL-17 production in NP cell cultures, with or without the presence of MSCs. This may suggest that the interaction of distinct soluble factors with different cell types could alter the immune context of NP.

Furthermore, indoleamine 2,3-dioxygenase (IDO), a rate-limiting enzyme that catalyzes the degradation of tryptophan, is found at increased levels during chronic inflammatory diseases induced by inflammatory mediators such as IFNs and IL-6 [48]. Elevated IDO expression has been observed in nasal polyloid tissue as compared with healthy nasal mucosa [49]. IDO is an immunoregulatory enzyme and belongs to the MSC arsenal. The role of MSCs in induction of tolerance in renal allograft recipients was not confirmed in IDO knockout mice, showing the crucial importance of IDO to the immunosuppressive effect of MSCs via regulatory T cells [50].

In addition, we observed an elevated presence of IFN-γ and IL-6 in cultures of NP-derived cells. After coculture with MSCs, IL-6 levels increased whereas IFN-γ declined. It is important to note that a decrease in IFN-γ could indicate a decrease in Th1 cells activation.

Different types of T cells are implicated in the pathogenesis and progression of NP, but, in populations of European descent, Th2-driven disease is still a hallmark of the condition. In the Asian population NP is a Th17/Th1-driven disease with increase of neutrophils. We characterize our patients as descendants of European and none was of Asian origin. In our sample the inflammatory cells profile was similar to the one found in European population (increase of Th2 cells and eosinophils).

Some studies have demonstrated that MSCs are able to promote conversion of the Th1 phenotype into a Th2 response [51]. In this sense, MSCs could contribute to NP, considering the microenvironment already saturated with Th2-dependent interleukins in this setting. In the present study, we did not detect IL-4 which is considered an important interleukin that induces differentiation from Th0 to Th2 immune response in NP-derived cell cultures, regardless of the presence of MSCs. This would suggest that in our coculture experiments with NP-derived cells, MSCs could not intensify a specific Th2 response.

Additionally, in the presence of MSCs, Th1 cytokines profile was altered, Th2 cytokines were not detected, and IL-10 was increased. These results indirectly suggest that the T-cell profile may have been directed to a regulatory pattern, considering the prominent increase of CD4+CD25+Foxp3+ T cell frequency in our MSC cocultures.

The NP treatment is based on two main pillars: oral and topical steroids and surgery, with the recurrence of NP after surgery being usual. The understanding of MSC mechanism in decreasing the inflammatory process in NP could be helpful to reduce the intake of steroids and the surgery indications.

In conclusion, we demonstrated that MSCs can be a useful tool for the investigation of the inflammatory microenvironment of NP. These results were obtained entirely in vitro, and any conclusions about the actual effects of MSCs in NP in vivo remain to be explored. However, our findings clearly demonstrate an immunoregulatory effect of MSCs on immune cells (especially T cells) derived from nasal tissue affected by polyposis. Finally, we hope that further studies will be performed in the search for an understanding of the mechanism of MSC activity in the context of NP inflammation.

**Conflict of Interests**

The authors declare that there is no conflict of interests regarding the publication of this paper.
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