The Hypoxic Synovial Environment Regulates Expression of Vascular Endothelial Growth Factor and Osteopontin in Juvenile Idiopathic Arthritis

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ABSTRACT. Objective. Synovial angiogenesis, a critical determinant of juvenile idiopathic arthritis (JIA) pathogenesis, is sustained by various mediators, including vascular endothelial growth factor (VEGF) and osteopontin (OPN). We characterized the contribution of the local hypoxic environment to VEGF and OPN production by monocytic cells recruited to the synovium in JIA.

Methods. Paired synovial fluid (SF) and peripheral blood (PB) samples were collected from 20 patients with JIA. Mononuclear cells (MC) were isolated, and monocytic cells were purified by adherence, maintained in a hypoxic environment, or subjected to reoxygenation. VEGF and OPN protein concentrations were tested in SF, plasma, and culture supernatants by ELISA, and mRNA expression was assessed in freshly purified and cultured cells by reverse transcriptase-polymerase chain reaction. Synovial tissue was obtained at synovectomy from 4 patients with JIA, and analyzed by immunohistochemistry for VEGF, OPN, CD68, and hypoxia-inducible factor-1α (HIF-1α).

Results. VEGF mRNA expression was increased in SFMC and SF monocytic cells compared to matched PBMC and PB monocytic cells or SF lymphocytes, correlating with significantly higher protein levels in SF relative to plasma samples. Accordingly, OPN mRNA expression in SF monocytic cells was associated with significant increase of SF protein. Immunohistochemistry revealed the presence of both factors in synovial tissues at the level of the lining and sublining layers, which colocalized with intense CD68 and HIF-1α staining, suggesting production by hypoxic synovial monocytic cells. VEGF and OPN expression was abrogated upon SF monocytic cell reoxygenation and maintained by exposure to prolonged hypoxia.

Conclusion. Hypoxic synovial monocytic cells are a likely source of VEGF and OPN in JIA. These data point to a role for hypoxia in the perpetuation of synovitis in JIA. (First Release April 15 2009; J Rheumatol 2009;36:1318–29; doi:10.3899/jrheum.080782)

Key Indexing Terms: JUVENILE IDIOPATHIC ARTHRITIS MONOCYTES HYPOXIA ENVIRONMENT SYNOVIAL FLUID VASCULAR ENDOTHELIAL GROWTH FACTOR OSTEOPONTIN

Juvenile idiopathic arthritis (JIA) consists of a clinically heterogeneous group of chronic pediatric arthritides with poorly defined etiology characterized by synovial joint inflammation, which results in synovial tissue hyperplasia, progressive cartilage destruction, and bone erosion. The chronicity of inflammation is crucial to the pathogenesis of JIA, and persists as a consequence of extensive angiogenesis and progressive infiltration and activation of inflammatory cells in synovial tissue and synovial fluid (SF) of affected joints.

Local release of proangiogenic mediators is responsible for the uncontrolled endothelial cell (EC) proliferation that takes place in many angiogenesis-dependent diseases, including rheumatic arthropathies. The EC-specific mitogen, vascular endothelial growth factor (VEGF), plays a central role in driving blood vessel neo-formation in the rheumatoid synovium via stimulation of EC survival, proliferation, and chemotaxis, and in monocytic cell recruitment/activation. Another proangiogenic mediator potentially implicated in the pathogenesis of inflammatory arthritis is the cytokine/extracellular matrix protein, osteopontin (OPN), which is endowed with EC adhesive capacity and represents a chemotactic stimulus for EC, monocytic cells, and T lymphocytes. Abundant VEGF and OPN expression was detected in the inflamed joints of patients with JIA, showing a good correlation with the
degree of vascularization and disease activity measures. However, the contribution of individual cell types and the role of environmental signals to VEGF and OPN production in JIA have not been assessed, hindering understanding of the pathogenetic mechanisms of the disease and the design of targeted therapies.

One of the major promoters of angiogenesis is hypoxia, a local decrease in partial pressure of oxygen (pO₂) that characterizes inflammatory and neoplasic lesions and is mediated by release of proangiogenic factors. The hypoxic microenvironment is a hallmark of the inflamed synovium, and its importance in the pathogenesis of adult arthropathies was documented. We have recently demonstrated the hypoxic nature of the synovium of patients with JIA and the constitutive expression of hypoxia-inducible factor-1α (HIF-1α) and HIF-2α, key regulators of hypoxia transcriptional response, in mononuclear cells recruited to inflamed joints in JIA. Further, we demonstrated hypoxic induction of the chemokine CCL20 in JIA synovial mononuclear cells, suggesting that the hypoxic synovial microenvironment may directly contribute to the persistence of JIA synovitis by increasing inflammatory cell infiltration in the affected joints.

Activated mononuclear cells are proangiogenic in different pathophysiological conditions, and induction of the mononuclear cell proangiogenic phenotype is associated with expression of HIF-1α and HIF-2α. We investigated the role of mononuclear cells infiltrating the JIA synovium in the production of VEGF and OPN and the contribution of the local hypoxic microenvironment to their expression, using an experimental protocol designed to prevent cell reoxygenation after the harvest from inflamed joints.

MATERIALS AND METHODS

Study population. Twenty patients were enrolled consecutively in this study provided they met the International League of Associations for Rheumatology Durban criteria for classification of JIA and had active knee disease, with clinical effusion and swelling of the joint at the time of therapeutic arthrocentesis. Patients with JIA were classified into 3 subtypes depending on the onset and course of disease: persistent oligoarticular JIA (pOJIA, oligoarticular onset and course); extended oligoarticular JIA (eOJIA, oligoarticular onset and polyarticular course); and polyarticular JIA (pJIA, polyarticular onset and course). Thirteen patients belonging to all disease subtypes had been previously treated with methotrexate (MTX) and/or nonsteroidal antiinflammatory drugs (NSAID), and only one patient with pJIA had been treated with a tumor necrosis factor inhibitor; the remaining patients were naive to any systemic biological agents. Informed consent was obtained prior to sample collection according to the procedure approved by our institution’s ethical committee. Clinical characteristics of patients, various laboratory measures of disease activity, and previous therapeutic regimens are reported in Table 1.

SF samples were collected at the time of therapeutic knee arthrocentesis into sodium-heparin tubes under vacuum, and their pO₂ levels were monitored with a portable trace oxygen analyzer (Oxi 315i/set, WTW, Germany). Paired peripheral blood (PB) samples were obtained on the occasion of routine venepuncture and collected as for SF. PB from 5 age-matched control subjects undergoing routine venepuncture for minor surgical procedures was collected after informed consent of the parents. A history of inflammatory or infection disorders in the 4 weeks before examination, and clinical or laboratory signs of inflammation at the time of blood sampling, was considered as an exclusion criterion. Specimens were centrifuged to prepare cell-free SF and plasma and stored at –80°C. Synovial tissues were obtained from 4 patients with oligoarticular-onset JIA undergoing synovectomy or arthroscopic biopsy after informed consent of the parents, fixed in 10% formalin, and embedded in paraffin.

Monocytic cell purification and culture. SF mononuclear cells (SFMC) and PB mononuclear cells (PBMC) were isolated by density gradient centrifugation over a Ficoll cushion (Amersham, Milan, Italy) and resuspended in RPMI-1640 (Euroclone, Celfio, Milano, Italy) supplemented with 10% heat-inactivated fetal calf serum (HyClone, Celfio). SFMC and PBMC were separated into adherent (monocytic) and nonadherent (lymphocyte) cells by incubation for 1 h at 37°C in tissue culture plates (Costar, Celfio), and their purity was assessed by flow cytometry (see below). For a subset of samples, monocytic cells were purified by negative selection using MACS magnetic beads according to standard methods (Human Monocyte Isolation kit-II, Miltenyi Biotec, Bologna, Italy). To avoid cell reoxygenation, SF-derived samples were handled in a 37°C humidified anaerobic work station incubator (BUG-BOX, ALC-International, Milano) flushed with a gas mixture of 94% N₂, 5% CO₂, and 1% O₂. In a few experiments, SF monocytic cells were reoxygenated for 16 h by incubation at 37°C in a humidified atmosphere containing 75% N₂, 5% CO₂, and 20% O₂ or maintained for 16 h under prolonged hypoxia (1% O₂), pO₂ in culture medium was monitored with the oxygen analyzer, as described. The endotoxin content, determined by the Limulus amoebocyte test (Bio-Whittaker, Walkersville, MD, USA), was <0.125 EU/ml in all reagents.

Flow cytometry analysis. The purity of the adherent and nonadherent cell fractions isolated from SFMC and PBMC was determined by flow cytometry. Cells were preincubated for 20 min at 4°C with phosphate-buffered saline (PBS) supplemented with 2% human AB serum and 0.05% Na₂CO₃ to block nonspecific labeling. Surface staining was carried out with FITC or phycocerythin (PE)-conjugated anti-CD14 (a pan-monocytic cell marker), PE-conjugated anti-CD3 (a pan-T cell marker), and IgG2a isotype-matched monoclonal antibodies (mAb; BD-Pharmingen, Milano, Italy) for 30 min at 4°C. Intracellular staining with FITC-conjugated anti-CD68 (a pan-monocytic cell marker) or IgG1 isotype-matched mAb (BD-Pharmingen) was performed after cell fixation in 4% paraformaldehyde and permeabilization in PBS containing 1% AB serum and 0.1% saponin (Sigma). Fluorescence was quantified on a FACScan flow cytometer equipped with CellQuest software (BD Biosciences). The percentage of monocytic cells in the adherent cell population ranged between 93% and 95% in the different samples, as determined by positivity for the CD14 and CD68 markers, whereas the nonadherent fractions were negative for CD14 and composed predominantly by CD3+ T cells at a purity of ≥83%. Viability was ≥95%, as determined by propidium iodide staining.

ELISA. Secreted VEGF and OPN were measured in SF, plasma, and cell-free supernatants by ELISA, using specific Quantikine immunoassay kits (R&D Systems, Space, Milano, Italy). Optical density was determined by a Tecan Spectrafluor Plus plate reader (Tecan, Manndorf, Switzerland) at 450 nm. Data were analyzed with the Graph Pad Prism 3 Software (Graph Pad, San Diego, CA, USA).

Reverse transcriptase-polymerase chain reaction (RT-PCR). Total RNA was purified from single-cell suspensions by the Quick RNeasy Mini Kit. Control of RNA integrity was carried out by electrophoresis with an Agilent Bioanalyzer 2100 (Agilent, Waldbronn, Germany). RNA (1 µg) was reverse-transcribed into double-stranded cDNA on a GeneAmp PCR
Table 1. Clinical and laboratory features of patients with juvenile idiopathic arthritis at the time of sampling. Results are expressed as mean (range).

<table>
<thead>
<tr>
<th>Disease Subtype</th>
<th>Age at Sampling, yrs</th>
<th>Disease Duration, yrs</th>
<th>Female/Male</th>
<th>No. Active Joints</th>
<th>ESR, mm/h</th>
<th>CRP, mg/dl</th>
<th>Hb, g/dl</th>
<th>RF+, %</th>
<th>ANA+, %</th>
<th>Treatment</th>
</tr>
</thead>
<tbody>
<tr>
<td>Persistent oligoarticular, n = 9</td>
<td>11.2 (4.9–20.1)</td>
<td>5.2 (0.2–11.1)</td>
<td>8/1</td>
<td>1 (1–2)</td>
<td>25 (5–63)</td>
<td>0.98 (0.46–2.8)</td>
<td>12.8 (12–15)</td>
<td>0</td>
<td>100</td>
<td>MTX (1 patient)</td>
</tr>
<tr>
<td>Extended oligoarticular*, n = 7</td>
<td>11.5 (4.3–18)</td>
<td>6.8 (1.1–13.1)</td>
<td>6/1</td>
<td>3 (1–8)</td>
<td>31 (3–54)</td>
<td>2.1 (0.46–7.5)</td>
<td>12</td>
<td>0</td>
<td>100</td>
<td>MTX (6 patients)</td>
</tr>
<tr>
<td>Polyarticular*, n = 4</td>
<td>12 (5.3–18.1)</td>
<td>7.2 (0.6–12.8)</td>
<td>4/0</td>
<td>6 (1–11)</td>
<td>30 (6–53)</td>
<td>1.6 (0.46–3.2)</td>
<td>11.8</td>
<td>0</td>
<td>100</td>
<td>NSAID alone (1 patient)</td>
</tr>
</tbody>
</table>

† Disease activity was defined by the presence of joint swelling or limitation of movement with either pain on movement or tenderness. * Polyarticular course.

ESR: erythrocyte sedimentation rate; CRP: C-reactive protein; Hb: hemoglobin; RF: rheumatoid factor antibody; ANA: antinuclear antibody; MTX: methotrexate; NSAID: nonsteroidal antiinflammatory drugs.

System 2700 (Applied Biosystems, Milano, Italy), using the Advantage RT-for-PCR Kit (BD-Pharmingen). For semiquantitative RT-PCR, cDNA was amplified with recombinant TaqDNA polymerase (Invitrogen), as described. Products were electrophoresed on a 1.2% agarose gel and visualized by ethidium bromide staining. Real-time quantitative PCR (qRT-PCR) was performed on a 7500 Real-time PCR System (Applied Biosystems), using SYBR Green PCR Master Mix and 300 nM of the following sense and antisense oligonucleotide primers (TIBMolbiol, Genova, Italy): VEGF, forward, 5'-GCA GCT TGA GTT AAA CGA ACG-3'; reverse, 5'-GCA GCG TGG TTT CTG TAT C-3'; OPN, forward, 5'-TGA CTT TTC GTT GGA CCA GCT TAC-3'; reverse, 5'-ATG GCT TTC GTT GGA CTT AC, as described.

Immunohistochemistry. 5 x 10^4 SF monocytic cells were applied to polylsine glass slides, fixed in 4% PAF, and permeabilized with 0.1% Triton X-100. VEGF protein expression was assessed by immunostaining of cytocentrifuge slides with anti-hVEGF rabbit antibody (Santa Cruz Biotechnology, Santa Cruz, CA, USA), and detection was carried out with anti-rabbit Ig conjugated to HRP-labeled dextran polymer (Dako, Glostrup, Denmark), as described. Isotype-matched mAb was used as control.

Immunohistochemical labeling of tissue specimens was performed by a 3-step immunoperoxidase technique. Paraffin-embedded serial tissue sections were incubated at room temperature for 30 min with anti-Cd68 (Dako), anti-Cd34 (Dako), anti-VEGF (Santa Cruz), and anti-OPN (Abcam, Cambridge, UK) antibody, or at 4°C overnight with anti-HIF-1α mAb (Abcam). Sections were reacted with anti-mouse (for Cd68 and Cd34) and anti-rabbit (for VEGF and OPN) Ig conjugated to peroxidase-labeled dextran polymer (Dako). For HIF-1α detection, the tyramide signal amplification system kit was used (Dako). Isotype-matched antibodies were used as negative controls. Chromogenic diaminobenzidine substrate was then applied. Slides were counterstained with Mayer’s hematoxylin and examined under a phase-contrast microscope (Olympus, Milano, Italy).

Statistical analysis. VEGF and OPN concentrations in SF or plasma were compared among the 3 subgroups of patients and controls using the non-parametric Mann-Whitney U-test (p < 0.05 as statistically significant). Concomitant cytokine determinations in paired SF and plasma specimens were evaluated by the Wilcoxon rank test.

RESULTS

Monocytic cells are an important source of VEGF in JIA SF. Initial experiments were carried out to measure VEGF release into SF from patients with JIA. Elevated amounts of VEGF were detected in the SF from all disease subtypes, although with some degree of patient to patient variability (Figure 1A), with higher concentrations in children with polyarticular onset (pJIA, median 1321 pg/ml) or course of disease (eOJIA, median 1090 pg/ml), compared with patients with an oligoarticular onset and course (pOJIA, median 820 pg/ml). Globally, median protein concentrations were significantly increased (p < 0.0001) in SF relative to paired plasma samples (Figure 1A, inset), ranging from 2.8-, 4.2-, and 5.9-fold in pOJIA, eOJIA, and pJIA subgroups, respectively. When these data were analyzed in relation to the clinical characteristics of patients, we did not find any statistically significant correlation between VEGF levels and disease duration or severity, number of active joints involved, or therapeutic regimen.

The cellular source of VEGF in JIA SF was then investigated. VEGF expression was first assessed by RT-PCR in SFMC from a subset of 8 children with different forms of JIA and compared with that of paired PBMC (Figure 1B). VEGF mRNA was constitutively expressed in all SFMC samples, although it varied among different individuals, with high levels present in 7/8 samples and a lower expression detectable in one sample. The variability in VEGF transcript levels did not correlate with the disease subtype or type of treatment. In contrast, minimal or undetectable mRNA levels were found in matched PBMC samples, suggesting that VEGF expression is upregulated in cells recruited to the inflamed synovium. Equal amounts of RNA purified from the samples shown in Figure 1B were combined according to disease subtype, and RNA pools were subjected to qRT-PCR. VEGF mRNA expression was significantly higher in SFMC than in PBMC from all disease subtypes (Figure 1C), confirming the semiquantitative analysis.

The monocytic cell and lymphocyte cell fractions of SFMC (Figure 2A) were then isolated from 3 randomly selected patients, each representative of a specific disease subtype, and tested for VEGF expression by RT-PCR (Figure 2B). Both cell types constitutively expressed VEGF mRNA, but transcript levels were higher in SF monocytic cell
cells (SF-Mn) from all the patients tested, suggesting that VEGF was transcribed predominantly in the monocytic cells than in the lymphocyte cell population present in the JIA SF. These findings were confirmed by qRT-PCR, which demonstrated a 15-fold higher VEGF expression in SF monocytic cell-derived compared to SF lymphocyte (SF-Ly)-derived RNA pools (data not shown). Interestingly, SF monocytic cells expressed increased VEGF mRNA compared to monocytic cells isolated from paired PBMC (PB-Mn; Figure 2C, upper panel). These findings were confirmed by qRT-PCR mRNA quantification, which revealed 3.9-, 4.1-, and 4.4-fold increases in SF monocytic cells relative to matched PB monocytic cell preparations (Figure 2C, lower panel). Comparable VEGF transcript levels were detectable in PB monocytic cells purified from patients with JIA and controls (Figure 2C). A similar pattern of results was observed in monocytic cells purified from the other 5 patients (data not shown). mRNA expression in SF monocytic cells was paralleled by VEGF protein immunostaining (Figure 2D). These data indicate that VEGF was significantly upregulated in monocytic cells recruited to the inflamed joints of patients with JIA, suggesting a likely role for SF monocytic cells as a source for this factor and raising the issue of the stimuli controlling its production in the synovial environment. 

Hypoxia induces VEGF production by SF monocytic cells. We next investigated whether the pathological low O2 levels of the rheumatoid synovial environment \(^{23,25}\) induced VEGF expression in SF monocytic cells from patients with JIA.
Freshly purified SF monocytic cells were exposed for 16 h to ambient pO2 (20% O2, hereafter referred to as reoxygenation) or were maintained under a controlled hypoxic environment (1% O2) for the same length of time, and VEGF transcript levels were then quantified (Figure 3A). Interestingly, reoxygenation caused a pronounced reduction of VEGF mRNA expression relative to fresh cells, and this effect was in part counteracted by culture under hypoxia. Accordingly, VEGF secretion by SF monocytic cells maintained under the hypoxic environment exceeded by 2.3- to 5-fold that of reoxygenated cultures (Figure 3B). A similar pattern of results was observed in SF monocytic cells purified from all the patients listed in Figure 1B (data not shown), suggesting that hypoxia is a stimulus for VEGF production by SF monocytic cells in JIA. The potential role of monocytic cells as a source for VEGF within the hypoxic synovial environment was supported by data showing that VEGF mRNA expression was upregulated by more than 4-fold in PBMC cultured under low O2 concentrations (1%) similar to those present in the JIA SF28, and that this response was completely abrogated upon depletion of monocytic cells (Figure 3C).

OPN production is enhanced by hypoxia in SF monocytic cells from patients with JIA. A crosstalk between VEGF and OPN was previously established in endothelial cells and shown to amplify the angiogenic response9,14. To determine...
whether VEGF overexpression in JIA SF was associated with increased OPN release, OPN protein concentrations were assessed in the same specimens tested for VEGF (Figure 4A). A significant increase in median OPN levels (p < 0.0001) was detected in SF relative to paired plasma samples (Figure 4A, inset), paralleling that of VEGF. Higher OPN concentrations were found in SF from pJIA (median 325.3 ng/ml) and eOJIA (median 319.9 ng/ml) subgroups compared to the pOJIA subtype (median 235.2 ng/ml), with an average increase of 9-, 7.6-, and 6.2-fold with respect to the corresponding plasma specimens (Figure 4A), thus showing a trend for a positive correlation between OPN and VEGF production in JIA SF. As for VEGF, OPN levels were independent of disease duration or activity, number of active joints involved, or type of treatment. Quantification of OPN transcript levels by qRT-PCR in SF monocytic cell and PB monocytic cell-derived RNA pools demonstrated increased OPN expression in SF monocytic cells isolated from patients with JIA compared to paired PB monocytic cells or to PB monocytic cells purified from controls (Figure 4B), reflecting the protein results, and higher OPN transcript levels were quantified in the monocytic cells than in the lymphocyte fractions of SFMC (data not shown), as shown for VEGF.

Next, we investigated the possible relationship between hypoxia and OPN production by SF monocytic cells. As for VEGF, fresh SF monocytic cells constitutively expressed elevated levels of OPN mRNA, which were strongly reduced by reoxygenation, whereas OPN transcript expression was significantly higher in hypoxic relative to reoxygenated cells (Figure 5A). Accordingly, OPN protein secretion was increased in hypoxic relative to reoxygenated SF
monocytic cells, ranging between 2.4- and 4.8-fold in the samples tested (Figure 5B). These data suggest a regulatory role for hypoxia in SF monocytic cell production of OPN in JIA. OPN mRNA expression was upregulated by more than 3-fold in PBMC cultured at 1% O₂, and this response was partially abrogated upon monocytic cell depletion (Figure 5C), confirming the potential role of these cells as OPN producers in response to the low pO₂ present in the JIA synovial environment²³,²⁵,²⁸.

Monocytic cells recruited to the hypoxic JIA synovium express VEGF and OPN proteins. Experiments were carried out to measure VEGF and OPN expression in vivo in monocytic cells infiltrating the hypoxic areas of the JIA synovium. Tissue specimens obtained at synovectomy from a subset of patients with JIA were analyzed by immunohistochemistry. Serial sections were immunostained for the hypoxia marker HIF-1α (Figure 6A), the pan-monocytic cell marker CD68 (Figure 6B), the endothelial cell marker CD34 (Figure 6E), and for VEGF (Figure 6C) and OPN (Figure 6D), and counterstained with hematoxylin-eosin (Figure 6F). Clear positivity for HIF-1α was observed in the lining and, with lower intensity, in the sublining layers of synovial membranes, in areas characterized by intense CD68 staining, suggesting its induction in infiltrating monocytic cells, although a few positive fibroblast-like cells (CD68−) were also detectable. VEGF and OPN immunoreactivity was observed in the same areas of serial tissue sections, confined almost entirely to cells positive for HIF-1α and CD68. No staining was detectable with isotype-matched control antibodies (data not shown). These data suggest that monocytic cells recruited to the JIA synovium respond to the local hypoxic environment with production of VEGF and OPN. Staining with anti-CD34 mAb showed some positivity for VEGF and OPN around areas of vascularization, consistent with the proangiogenic role of both cytokines.

DISCUSSION
Angiogenesis and chronic inflammation are codependent because an inflammatory state can promote new vessel formation, which, in turn, allows the migration and progressive infiltration of other inflammatory cells and perpetuation of the inflammatory state³⁴. Soluble mediators produced by stromal and inflammatory cells drive angiogenesis and are crucial to the perpetuation of synovial inflammation in rheumatoid arthritis²,⁵-⁸,¹²,¹⁵,²¹,³⁵. In our study, we observed abundant release of the proangiogenic and chemotactic mediators VEGF and OPN into the inflamed joints of patients with JIA and identified synovial monocytic cells as a likely cellular source for both factors. Further, we provide the first evidence that the hypoxic synovial environment is a stimulus for their production in patients with JIA.

Elevated VEGF levels accumulated in JIA SF as compared to matched plasma samples, in particular in patients with pJIA and eOJIA relative to the pOJIA subgroup, confirming previous findings¹⁸. A parallel significant increase in the amounts of OPN was measured in SF with respect to matched plasma specimens. These results suggest a trend for a positive correlation between OPN and VEGF production in JIA SF. A possible relationship between the 2 factors is also suggested by the observation of their similar distribution at the level of the lining layer of JIA synovial membranes. These data are in accord with our previous findings...
demonstrating a clear positive correlation between VEGF and OPN synovial expression and degree of angiogenesis, and suggest a functional cooperation between the 2 factors in the control and stimulation of vascularization in the inflamed joint of patients with JIA. A crosstalk between VEGF and OPN was previously established and shown to amplify the angiogenic response. Expression of various angiogenic factors, including VEGF, in fresh JIA membranes was shown to correlate with increased tissue angiogenesis and revascularization of human synovium engrafted into SCID mice. Thus, production of higher amounts of VEGF and OPN in patients with pJIA and eOJIA, relative to the pOJIA subgroup, may be clinically relevant, resulting in a greater extent of joint vascularization and inflammation and in a major severity of the disease.

SFMC, and in particular the monocytic cell fraction, were identified as a possible source for both cytokines. Expression of VEGF and OPN was significantly higher in SF-derived than in matched PB-derived cells from all disease subtypes or PB monocyctic cells from control subjects, indicating that recruitment to the local synovial environment is a stimulus for their production. Further, mRNA was expressed at increased levels in SF monocytic cells compared to SF lymphocytes present in the JIA SF, and was paralleled by protein expression, expanding previous evidence on the predominant role of synovial monocytic cells rather than synovial lymphocytes for the production of proinflammatory factors such as CCL2, CCL3, and CXCL10 in JIA. In contrast, findings reported for adult arthritides, such as rheumatoid arthritis (RA) and osteoarthritis (OA), suggest that the distribution of VEGF and OPN in the inflamed synovium was preferentially confined to fibroblasts and T cells.

Hypoxia is a common feature of the rheumatoid synovium and has been implicated in the pathogenesis of adult arthropathies. HIF-1α was found constitutively expressed in the lining and, to a lesser extent, the sublining layers of the JIA synovium, confirming the presence of
reduced pO₂ in the synovia of patients with JIA. Positivity for HIF-1α was mostly colocalized with CD68-specific staining in serial tissue sections, and was associated with intense immunoreactivity for both VEGF and OPN. These findings support an active role of monocytic cells migrating to the JIA synovium as VEGF and OPN producers, and suggest that cytokine overexpression in synovial monocytic cells is probably a result of activation of the HIF-1α pathway by local hypoxia.

The possibility that hypoxia stimulates synovial monocytic cells to produce VEGF and OPN in JIA is supported by the observation that constitutive VEGF and OPN mRNA levels detectable in freshly isolated SF monocytic cells decreased upon normoxic in vitro cell culture, whereas maintenance of the hypoxic environment sustained their expression. Accordingly, cell incubation in prolonged hypoxia resulted in increased production of both cytokines compared to reoxygenated cells. These data extend earlier findings that VEGF can be upregulated by hypoxia in synovial fibroblasts from patients with RA and OA21,24,38, and provide the first evidence that hypoxia can induce OPN expression in synovial cells, linking hypoxia signaling to activation of the angiogenic process in JIA. The observation that VEGF and OPN induction in PBMC cultured under low

Figure 6. Immunohistochemical detection of VEGF and OPN proteins in hypoxic monocytes infiltrating the JIA synovium. Synovial tissue was obtained at synovectomy from a representative patient (Patient 3), and staining was carried out on serial sections with the indicated mAb. Hematoxylin-eosin staining (EE) is shown in panel F as a control. Slides were examined under a phase-contrast microscope (original magnification 10×). Positive cells (stained brown) are detectable at level of lining layer (black arrows) and sublining layer (white arrows). Results are representative of 4 patients.
O₂ concentrations similar to those present in the inflamed JIA synovium²³,²⁵ was almost completely abrogated upon depletion of monocyteic cells is noteworthy, and is consistent with the possible role of hypoxic monocytic cells as a source for both factors. However, the observation of higher levels of VEGF and OPN expression in fresh SF mononuclear cells than in cells maintained in vitro under hypoxic conditions suggests that additional factors, such as proinflammatory cytokines present in the rheumatoid synovial environment¹⁰,¹⁴,²⁶,³⁴,⁴⁰, may contribute to the hypoxia stimulatory effects in vivo. Evidence that hypoxia may interact with immune-derived stimuli in the regulation of gene expression in mononuclear cells and in modulating proinflammatory mediator production in synovial cells was previously reported³⁸,⁴⁰-⁴³. Future studies to examine the effects of hypoxia and proinflammatory cytokines in combination will yield a more complete picture of the functional behavior of mononuclear cells within the pathological intraarticular environment in JIA.

The relationship between synovial hypoxia and mononuclear cell expression of VEGF and OPN expands previous observations on the relevance of hypoxia for the activation of mononuclear cell proangiogenic phenotype in other pathological conditions¹⁰,¹³,³⁰-³², and indicates that a dynamic change in production of proangiogenic mediators by synovial mononuclear cells may occur depending on the O₂ availability. These findings also highlight the importance of performing in vitro experiments under conditions that mimic those occurring in vivo within JIA rheumatic joints for understanding the pathogenetic mechanisms of the disease. Because mobilization causes arthritic joints to undergo cycles of hypoxia-reoxygenation²¹,²⁵,⁴⁰,⁴⁴, SF mononuclear cells may be subjected to different degrees of oxygenation within diseased joints, resulting in shifting gradients of secreted VEGF and OPN and consequently in the variable extent of synovial vascularization. Extreme O₂ situations were tested in this study. pO₂ in rheumatic joints ranged between 6 and 50 mm Hg (0.8%–7% O₂), whereas physiological O₂ concentrations in healthy joints range between 50 and 70 mm Hg (7%–10% O₂)²⁳. One percent of O₂, a common experimental condition of hypoxia²¹,²³,三十年, is in the low range of the pathological O₂ levels present in the inflamed synovium²³,²⁸,⁴⁰, whereas 20% O₂, a standard in vitro cell-culture condition³⁵,³⁰, is likely to represent a supraphysiological situation. Future studies will aim at assessing SF mononuclear cell responses at intermediate O₂ concentrations to mimic more closely the in vivo physiopathological O₂ fluctuations in the JIA synovium.

VEGF and OPN promote angiogenesis by directly stimulating EC proliferation and survival, adhesion, and migration³,⁴,⁵. However, their proangiogenic activity is also exerted indirectly through the control of leukocyte trafficking¹⁰,¹²,¹³,¹⁵,⁴⁶,⁴⁷. VEGF is a potent chemoattractant for mononuclear cells¹⁰,⁴⁷, and OPN acts as an adhesive, chemotactic, and activating factor for monocytic cells and T lymphocytes, promoting their recruitment into the inflamed synovium and stimulating proinflammatory cytokine production¹²-¹⁶. The importance of these factors in the pathogenesis of inflammatory synovitis is emphasized by the observations that inhibition of VEGF-R1 signaling⁴⁷,⁴⁸ or OPN production¹¹,⁴⁹ is associated with marked reduction of joint inflammation in murine RA models. Previous studies in mouse models of arthritis have suggested a critical role for synovial monocytic cells in the regulation of arthritic joint inflammation, because their local removal from the knee joints by intraarticular administration of selective depleting agents, such as clodronate-encapsulated liposomes⁵⁰, prior to induction of experimental arthritis results in a substantial decrease of influx of inflammatory cells and arthritis severity⁵¹,⁵². Thus, we can hypothesize that sustained VEGF and OPN induction by intraarticular hypoxia in synovial mononuclear cells may represent an important mechanism of amplification of the inflammatory process through the recruitment of other inflammatory cells, which in turn will enhance the inflammatory response in the arthritic joints, ultimately leading to the chronicity of JIA synovitis.

Angiogenesis is a complex process, involving multiple gene products expressed by different cell types, all contributing to an integrated sequence of events. Elevated levels of other proangiogenic factors in addition to VEGF and OPN, such as interleukin 8 (IL-8), have been detected in synovial fluids from patients with JIA⁴,⁵³ and may contribute to the pathogenesis of the disease. Interestingly, IL-8 is inducible by hypoxia in mononuclear cells⁵⁴, and a positive feedback loop among IL-8, VEGF, and OPN production was previously established. VEGF was reported to stimulate endothelial cells to produce both IL-8⁵⁵ and OPN⁹,¹⁴, and OPN was shown to trigger a proangiogenic phenotype in human monocytes by inducing IL-8 expression both in vitro and in vivo¹²,¹⁴. On the other hand, IL-8 can increase VEGF production⁵⁶,⁵⁷ and enhance VEGF-mediated neovascularization by concomitantly triggering EC proliferation and migration⁵⁸. Hence, it is conceivable that the proangiogenic state of mononuclear cells within the local hypoxic environment of inflamed joints will ultimately be dictated by the crosstalk among these mediators, which synergistically stimulate each other’s production and function as 2-way signaling molecules between mononuclear cells and EC. Further investigations are under way in our laboratory to define the functional cooperation among these factors in JIA.

Our findings point to a role for hypoxia and the HIF-1 pathway in the persistence and amplification of synovial inflammation associated with JIA and as potential pharmacological targets for treatment of the disease.

REFERENCES


