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Research article

Suppression of murine collagen-induced arthritis by targeted apoptosis of synovial neovasculature

Danielle M Gerlag*, Eric Borges†§, Paul P Tak*, H Michael Ellerby†¶, Dale E Bredesen†, Renata Pasqualini†***¶, Erkki Ruoslahti† and Gary S Firestein*

*Division of Rheumatology, Allergy and Immunology, University of California, San Diego School of Medicine, La Jolla, CA, USA
†The Burnham Institute, La Jolla, CA, USA
‡Division of Clinical Immunology and Rheumatology, Academic Medical Center/ University of Amsterdam, Amsterdam, The Netherlands
§Present affiliation: MorphoSys AG, Martinsried/München, Germany
¶Present affiliation: The Buck Center for Research in Aging, Novato, CA, USA
**Present affiliation: Department of GU Oncology, The University of Texas MD Anderson Cancer Center, Houston, TX, USA

Correspondence: Gary S Firestein MD, Division of Rheumatology, Allergy and Immunology #0656, UCSD School of Medicine, 9500 Gilman Drive, La Jolla, CA 92093, USA. Tel: +1 858 534 2359; fax: +1 858 534 2606; e-mail: gfirestein@ucsd.edu

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Abstract

Because angiogenesis plays a major role in the perpetuation of inflammatory arthritis, we explored a method for selectively targeting and destroying new synovial blood vessels. Mice with collagen-induced arthritis were injected intravenously with phage expressing an RGD motif. In addition, the RGD peptide (RGD-4C) was covalently linked to a proapoptotic heptapeptide dimer, D(KLAKLAK)2, and was systemically administered to mice with collagen-induced arthritis. A phage displaying an RGD-containing cyclic peptide (RGD-4C) that binds selectively to the αvβ3 and αvβ5 integrins accumulated in inflamed synovium but not in normal synovium. Homing of RGD-4C phage to inflamed synovium was inhibited by co-administration of soluble RGD-4C. Intravenous injections of the RGD-4C–D(KLAKLAK)2 chimeric peptide significantly decreased clinical arthritis and increased apoptosis of synovial blood vessels, whereas treatment with vehicle or uncoupled mixture of the RGD-4C and the untargeted proapoptotic peptide had no effect. Targeted apoptosis of synovial neovasculature can induce apoptosis and suppress clinical arthritis. This form of therapy has potential utility in the treatment of inflammatory arthritis.

Keywords: angiogenesis, apoptosis, collagen-induced arthritis, rheumatoid arthritis

Introduction

In rheumatoid arthritis (RA), the synovium is characterized by hyperplasia of the intimal lining and mononuclear infiltration of the sublining, leading to erosion of cartilage and subchondral bone by invasive pannus [1]. Angiogenesis plays a crucial role in the formation of pannus, and the extensive network of blood vessels facilitates recruitment of mononuclear cells [2]. Proangiogenic mediators found in RA synovial tissue regulate migration and proliferation of endothelial cells. In addition, αvβ3 and αvβ5 integrins are important in angiogenesis [3–5]. αvβ3 is expressed on synovial blood vessels in rheumatoid arthritis [6,7], and αv antagonists injected directly into the joint suppress synovitis in rabbits [7]. Therefore, targeted...
induction of apoptosis in the neovascularure is a potential therapy for RA [8].

We evaluated a novel method of targeted drug delivery to inflamed joints using peptides that selectively bind to neovascularure [9]. Our data show that a constrained cyclic RGD peptide that binds to \(\alpha_v\beta_3\) and \(\alpha_v\beta_5\) integrins [10] homes to blood vessels in inflamed synovium after systemic administration. This peptide was covalently linked to a 14-amino-acid proapoptotic peptide and successfully suppressed arthritis in collagen-induced arthritis (CIA).

Materials and methods

Phage and peptides

Insertless fd phage and phage presenting the peptide CDCRGDGCFC (RGD-4C) was prepared as previously described elsewhere [10–12]. The chemical structure of RGD-4C was determined by NMR analysis and is described in detail elsewhere [13]. K91kan bacteria were a gift from G Smith. The peptides RGD-4C, CARAC, \(\text{d}(\text{KLAKLAKKLAKLAK})\) [designated \(\text{d}(\text{KLAKLAK})_2\)], and CDCRGDCFC-GG-\(\text{d}(\text{KLAKLAKKLAKLAK})\) [designated RGD-4C-\(\text{d}(\text{KLAKLAK})_2\)] were synthesized by AnaSpec, Inc (San Jose, CA, USA).

Collagen-induced arthritis in mice

CIA was induced in 6- to 8-week-old male DBA/1J mice (Jackson Laboratory, Bar Harbor, ME, USA), as previously described [14,15].

Phage delivery and detection

To determine homing characteristics of the RGD-4C phage \textit{in vivo}, mice with CIA on day 35 after immunization were anesthetized and injected with 200 \(\mu\)l of medium containing \(10^{10}\) transducing units (TU) of phage into the tail vein. After 5 min, the mice were perfused with 5 ml of medium given through the right atrium. The organs of interest and synovia were pooled, homogenized, incubated with K91kan bacteria for 30 min, and plated on tetracycline plates. Phage enrichment was calculated as the ratio of TU/g of synovial tissue divided by the TU/g of brain tissue. Because animals are perfused and sacrificed within minutes after administration, relative rates of penetration into the target tissue do not influence this assay system (i.e., the target is the intravascular surface of the blood vessels). For the competitive peptide inhibition studies, the phage was injected with either 1 mg of the RGD-4C peptide or with 1 mg of a control peptide, CARAC [16]. Unimmunized mice were used as normal controls.

Treatment protocol for CIA

In the clinical efficacy study, mice with established CIA were anesthetized and injected intravenously on days 35 and 41 with either 500 \(\mu\)g of covalently linked RGD-4C-\(\text{d}(\text{KLAKLAK})_2\) dissolved in 500 \(\mu\)l DMEM \((N=14)\), with a mixture of the equimolar amounts of uncoupled RGD-4C and \(\text{d}(\text{KLAKLAK})_2\) dissolved in 500 \(\mu\)l DMEM \((N=20)\), or with 500 \(\mu\)l DMEM only \((N=20)\). The clinical arthritis scores were assessed daily in a blinded manner using a semiquantitative scoring system from 0 to 4+ for each paw (maximum score) [14,15].

Immunohistochemical staining and antibodies

Mice were sacrificed on day 35 after immunization and their paws and internal organs were fixed in a 10% formalin solution for 24 hours, decalcified, and embedded in paraffin. The primary antibodies (horseradish peroxidase/anti-M13 monoclonal conjugate [Pharmacia Biotech, Piscataway, NJ, USA], rabbit anti-\(\alpha_v\) antibody [17]) and secondary antibody (swine anti-rabbit-AP; Dako, Glostrup, Denmark) were diluted in 2% BSA–PBS. Peroxidase activity was detected using 3,3′-diaminobenzidine, and alkaline phosphatase activity was detected using alkaline phosphatase kit III (Vector Laboratories Inc, Burlingame, CA, USA).

Terminal deoxynucleotidyl transferase-mediated UTP end labeling (TUNEL) assay

The paws of the animals treated with vehicle or 500 \(\mu\)g of RGD-4C coupled to \(\text{d}(\text{KLAKLAK})_2\) were fixed in a 10% formalin solution, decalcified for 14 days in a 15% EDTA–PBS solution at 4°C, and embedded in paraffin. Sections (5 \(\mu\)m) were incubated with proteinase K (20 \(\mu\)g/ml) for 20 minutes. The In situ Death Detection Kit from Boehringer Mannheim GmbH (Mannheim, Germany) was used in accordance with the manufacturer’s instructions.

Results

Phage displaying RGD-4C accumulates in inflamed synovium

RGD-4C phage was injected intravenously into DBA/1J mice with active CIA on day 35 \((N=18)\). Control mice with CIA received insertless fd phage \((N=14)\). After circulation of the phage and perfusion with medium, the control organ (the brain) and synovium from arthritic joints were surgically dissected and phage enrichment was calculated as the ratio of TU/g of synovial tissue to TU/g of control tissue. Accumulation of the RGD-4C phage in inflamed synovium was 8.0 ± 1.7 (mean ± standard deviation) fold that in brain \((P<0.01)\). For the control phage, this value was 1.5 ± 0.3 (Fig. 1). Neither the RGD-4C phage nor the control phage accumulated in synovium of normal DBA/1J mice \((N=6\) and 5, respectively; \(P<0.0001\) compared with inflamed synovium). Other organs (lung, liver, spleen, heart, kidney, pancreas, and gut) were also tested for enrichment of the RGD-4C and control phage. There was no selective accumulation of the RGD-4C phage in these organs (data not shown).

Inhibition of RGD-4C phage by a competing peptide

After showing that the RGD-4C phage targets inflamed synovium, we assessed the specificity of homing using the soluble RGD-4C peptide, a competitive inhibitor. The
RGD-4C phage was co-injected with 1 mg of either the RGD-4C peptide (N = 5) or a control peptide (CARAC; N = 5). Figure 2 shows that soluble RGD-4C peptide, but not the control peptide, blocked accumulation of the RGD-4C phage in inflamed synovium.

**Figure 2**

Inhibition of RGD-4C phage homing with the cognate soluble peptide. In competitive peptide inhibition studies in mice with collagen-induced arthritis, RGD-4C phage (‘RGD phage’) was co-injected with 1 mg of either RGD-4C peptide or control peptide (CARAC). Note accumulation of RGD-4C phage in inflamed synovium (P < 0.01).

TUNEL studies were performed on joint specimens collected 72 hours after intravenous injection of the chimeric peptide. Abundant positive cells were detected in small vessels of the inflamed synovium (Fig. 3c) but not in synovium of arthritic mice injected with vehicle (not shown). Positive cells were not found in lung, liver, or spleen of chimera-treated arthritic mice, indicating that apoptosis was selective to inflamed synovium.

**Discussion**

Angiogenesis has been implicated in inflammatory diseases such as RA [2,8]. Synovial tissues from patients with RA express more of the angiogenesis marker, αvβ3 integrin, than control synovium [6,7,18]. An angiostatic compound suppressed arthritis in rat adjuvant arthritis and CIA [19,20], and intra-articular administration of an RGD peptide that binds to αvβ3 decreased synovial inflammation in a rabbit model of arthritis [7].

The homing properties of the cyclic RGD peptide allowed us to deliver a toxic agent – the proapoptotic peptide D(KLAKLAK)2 [16]. The chimeric compound was injected intravenously on days 35 and 41. Control mice with CIA were either injected with the vehicle or were co-injected with similar quantities of the uncoupled RGD-4C and D(KLAKLAK)2 peptides. The mean pretreatment score was 10.3 ± 0.5. Neither the vehicle nor the uncoupled mixture of RGD-4C with D(KLAKLAK)2 had no effect on arthritis, whereas the targeted chimeric compound significantly decreased clinical arthritis on day 44 (P < 0.001; Fig. 4).
The RGD-4C peptide also enhances internalization of RGD-4C–D(KLAKLAK)2 [9]. When covalently linked to the homing peptide RGD-4C, the proapoptotic peptide enters cells that express αvβ3 or αvβ5 [22]. Once in the cytoplasm, it disrupts mitochondrial membranes and initiates apoptosis [15]. Distribution studies in vivo followed by cell fractionation indicate that RGD-4C–D(KLAKLAK)2 localizes to mitochondrial membranes of the target cells but not to endothelial cells in control tissues [15]. Cells that do not express these integrins are spared, because the proapoptotic peptide alone is not internalized. As with our studies, Pasqualini and colleagues showed that αv-directed RGD-

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References