Short Communication

Local delivery of beta interferon using an adeno-associated virus type 5 effectively inhibits adjuvant arthritis in rats

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Beta interferon (IFN-β) is a cytokine with potent immunomodulatory properties and has been described as a promising therapeutic molecule for the treatment of rheumatoid arthritis (RA). IFN-β was previously overexpressed intra-articularly using an adenoviral vector in rats with adjuvant arthritis (AA) as a model of RA. This effect was powerful, albeit transient due to the vector chosen. Therefore, in the context of pre-clinical development, a delivery vector optimized for intra-articular gene transfer, recombinant adeno-associated virus type 5 (rAAV5), was selected.

To exert an optimal effect, protein production should parallel the course of the disease. For this reason, the gene for IFN-β was placed under the control of an inflammation-responsive [nuclear factor (NF)-κB] promoter. After intra-articular injection of the rAAV5 constructs in rats with AA, local transcription of the transgene and production of the IFN-β protein was found, leading to a pronounced and sustained effect on paw swelling when the expression was under the control of the NF-κB-responsive promoter. Additionally, a significant beneficial effect was observed on proteoglycan depletion and erosions. Thus, intra-articular overexpression of IFN-β using a rAAV5 vector exhibits potential as an innovative therapy for the treatment of RA.

An improved understanding of the pathophysiology of rheumatoid arthritis (RA) has led to the development of several potential new therapeutic approaches based on either cytokines or cytokine inhibitors. Beta interferon (IFN-β) is a potent cytokine that exerts a variety of anti-inflammatory and chondroprotective effects in various animal models of RA (Tak et al., 1999; Triantaphyllopoulos et al., 1999; van Holten et al., 2004). In addition, the absence of IFN-β was recently shown to exacerbate arthritis in IFN-β−/− mice by activation of osteoclasts and stromal cells (Treschow et al., 2005). However, in patients, subcutaneous injections of recombinant IFN-β three times a week did not appear to be clinically effective (van Holten et al., 2004). We speculated that the short half-life of IFN-β protein in vivo and the route of administration may have contributed to insufficiently high levels in the joints for a therapeutic effect.

The feasibility of overexpressing IFN-β by intra-articular delivery of gene therapy vectors was therefore investigated in an animal model of RA (Adriaansen et al., 2006a). In this proof of principle study, an adenoviral vector encoding IFN-β under the control of the cytomegalovirus (CMV) promoter injected into the joints of arthritic rats was observed to induce a robust therapeutic effect, illustrated by a decrease in paw swelling, a significant shift towards an anti-inflammatory cytokine profile in the synovial tissue and pronounced effects on bone degradation. However, this effect was transient, lasting only 8 days. Adenoviral vector delivery can induce a strong host immune response, leading to elimination of transduced cells (Bessis et al., 2004), while the CMV major immediate-early (IE) promoter can also be silenced in vivo, resulting in a rapid decrease of transgene expression (Brooks et al., 2004; van de Loo et al., 2004).

We selected an alternative vector system, derived from adeno-associated virus (AAV) to test in pre-clinical developmental studies. AAV vectors are associated with low immunogenicity, minimal toxicity and are able to transduce both dividing and non-dividing cells (Adriaansen et al., 2006b). Previously, recombinant AAV (rAAV) vectors, packaged in serotype 5 capsids (rAAV5) have been shown to efficiently transduce synovial tissue and they gave rise to stable, long-term intra-articular transgene expression (Adriaansen et al., 2005). In the study presented here, the rat IFN-β gene was inserted between the inverted terminal repeats of AAV2 and the DNA was packaged in AAV5 capsids. Both the CMV major IE and a nuclear factor (NF)-κB-responsive promoter were used. Using the latter
promoter, expression of IFN-β should be disease-inducible, as NF-κB is highly activated in the synovium during inflammation (Tak & Firestein, 2001). We have validated this promoter in vitro for its specific NF-κB inducibility (data not shown). In addition, we attempted to boost IFN-β production by inclusion of an intron in the expression construct.

Briefly, cDNA encoding the rat IFN-β gene (kindly provided by Dr. C. Kaynor, Biogen Idec, Cambridge, MA, USA) including its signal peptide was amplified by PCR (Fwd: 5’-CCCGGATCCACATTGCCCAGTGGAAGTTTCTATAAAGTC-3’; Rev: 5’-CCCGGATCCACATTGCCCAGTGGAAGTTTCTATAAAGTC-3’) and cloned into the BamHI site of the pDC315-shuttle plasmid (Microbix Biosystems). This gene was subsequently inserted between Nhel and Xhol sites in the pAAV2-CMV-shuttle plasmid (Applied Viromics). To insert an intron downstream of the rat IFN-β gene, pDC315 was digested with Nhel and SalI and ligated into a pCI (Promega) plasmid containing the intron. The gene cassette, including the intron, was subsequently excised with SnaBI and SalI and inserted in the pAAV2-CMV-shuttle plasmid. To clone the IFN-β gene under the control of the NF-κB-inducible promoter, the CMV promoter was replaced by an NF-κB-responsive promoter. The NF-κB motif was derived from the human immunodeficiency virus long terminal repeat and was cloned into pGmCMV (Promega) using restriction enzymes Nhel and Xhol, forming the construct pGnL2. The vector pGnL6 containing six repeats of the NF-κB motif was constructed by digesting pGnL2 with Nhel and Xhol and further ligation with the phosphorylated NF-κB oligonucleotides. Thereafter, the NF-κB promoter was digested from pGnL6 by Ncol and Xbal to replace the CMV promoter in the previously obtained pAAV2-CMV-IFN-β plasmid. All constructs were confirmed by sequencing.

rAAV5 vectors were subsequently produced at the UNC Vector Core (Chapel Hill, NC, USA), as described previously (Xiao et al., 1998). Briefly, AAV vectors were generated using the triple transfection protocol and purified using a sequential process of nuclei isolation, density-gradient centrifugation and heparin-sulfate affinity column chromatography. Subsequently, vectors were titrated by dot-blot analysis.

All together, three rAAV5 vectors containing the rat IFN-β gene, rAAV5.CMV-IFN-β, rAAV5.NF-κB-IFN-β and rAAV5.CMV-IFN-β-intron were constructed; an empty vector consisting of only the CMV-shuttle vector without the transcription cassette served as a control. Using these vectors, an animal experiment was conducted. Rats (n=7 per group) were immunized at the base of their tail with 1 mg Mycobacterium tuberculosis H37RA (Difco) in 0.1 ml mineral oil on day 0 (Tak et al., 2001). Paw swelling was usually observed by days 10–11 and measured daily by water displacement plethysmometry. The right ankle joints were injected intra-articularly on day 12 after immunization in animals anaesthetized with isoflurane. The skin was disinfected with ethanol and 5 x 10^10 viral molecules (vm) of rAAV5.NF-κB-IFN-β, rAAV5.CMV-IFN-β, rAAV5.CMV-IFN-β-intron or empty vector serving as a control was injected anterolaterally into the right ankle joints in a total volume of 50 μl saline using a 31-gauge needle on a glass syringe (Nguyen et al., 1998). Three weeks later, the animals were sacrificed and hind joints were collected. Hind ankle joints were snap frozen in liquid nitrogen, pulverized using a pestle and mortar, and homogenized in Trizol Reagent (100 mg ml^-1; Invitrogen) using a tissue homogenizer. For both reverse transcriptase and real-time PCR (n=3 per group), total RNA was isolated from the aqueous phase according to the manufacturer’s instructions. RNA was dissolved in DEPC-treated water; the quality was checked by gel electrophoresis and quantified by spectrophotometry. cDNA was synthesized using the iScript cDNA Synthesis kit (Bio-Rad) and 1 μg RNA according to the instruction manual. RT-PCR amplification mixtures (25 μl) contained 25 ng template cDNA, 2 x SYBR Green I Supermix (12.5 μl; Bio-Rad) and 300 nM IFN-β Fwd (5’-CGTTCTGCTGTGGTTCATCC-3’) and Rev (5’-TGTAACCTCTTCACCTGT-TGAC-3’) primers. Glyceraldehyde-3-phosphate dehydrogenase (GAPDH) was used as an internal reference gene (Fwd: 5’-ATGCCATCACTGCCACCCTGCAG-3’; Rev: 5’-GGGTAGGAAACACGGAAGG-3’). Reactions were run on a Mini-Opticon real-time thermal cycler (Bio-Rad). The thermal profile consisted of one cycle at 95 °C for 3 min followed by 40 cycles at 95 °C for 15 s and 59 °C for 45 s. Each assay included (in duplicate): a standard curve of five serial dilutions of IFN-β and GAPDH cDNA, a no-template control and 25 ng sample cDNA. Each run was followed by a melting curve. Single control normalization for internal control gene and correction for primer efficiency were calculated as described earlier (Vandesompele et al., 2002).

For immunohistochemical detection of rat IFN-β protein in synovial tissue (n=4 per group), arthritic paws were fixed in 4 % buffered formalin and decalcified in 15 % EDTA. The paws were then embedded in paraffin and 5 μm sagittal serial sections of the hind ankle joints were cut. Primary IgG antibody (rabbit anti-rat IFN-β; Biomedical Laboratories) was incubated overnight at 4 °C, followed by incubation with horseradish peroxidase (HRP)-conjugated swine anti-rabbit immunoglobulin (Dako) for 30 min. For control sections, the primary antibody was omitted. HRP activity was detected using hydrogen peroxide as the substrate and 3-amino-9-ethylcarbazole (Sigma) as the dye. Sections were briefly counterstained with Mayer’s hemalum solution. The signal was subsequently quantified by digital image analysis. Six randomly selected fields within each section were chosen for digitizing the intensity of the signal. These images were acquired on an Olympus microscope, captured using a Charged Coupled Device video camera (Sony) and digitized with a PV100 multimedia 16-bit colour video digitizer card. In the resultant colour images, the area of positive staining and the mean optical density (MOD) were
measured by a macro program as described earlier (Kraan et al., 2001). The MOD is proportional to the cellular concentration of protein. The integrated optical density (IOD) is equal to the MOD multiplied by the area of positive staining.

Sections were also stained with Safranin O-fast green to determine the loss of proteoglycans. Safranin O staining was scored using a semiquantitative scoring system (0–3), where 0 represents no loss of proteoglycans, while a score of 3 indicates complete loss of staining for proteoglycans (Joosten et al., 1999; van Holten et al., 2004). Differences between groups were determined by the Kruskal–Wallis test, followed by a Mann–Whitney U-rank-sum test. \( P<0.05 \) was considered statistically significant. All analyses were done using SPSS version 11.5 (SPSS).

**Fig. 1.** Evaluation of rAAV5-mediated IFN-\( \beta \) gene delivery in AA in rats. Relative IFN-\( \beta \) mRNA levels in joint tissue. Total RNA was extracted from crushed joints 24 days after intra-articular injection and cDNA was synthesized. RT-PCR (a) and real-time PCR (b) were performed using specific primers for the IFN-\( \beta \) gene. \( C_t \) values were normalized to GAPDH levels and expressed as fold increase compared with matched controls mean ± SEM. All treated animals significantly overexpressed the transgene in their joints, rAAV5.NF-\( \kappa \)B-IFN-\( \beta \) gave rise to the highest IFN-\( \beta \) mRNA levels (*\( P<0.05; n=3 \) per group) and intra-articular IFN-\( \beta \) protein production after local gene therapy. Rat joints were collected 24 days after injecting the virus, decalcified and embedded in paraffin (\( n=4 \) per group). Sections were immunohistochemically stained for rat IFN-\( \beta \) (c) and quantified by digital image analysis (d). Protein production was highest in rAAV5.NF-\( \kappa \)B-IFN-\( \beta \) injected animals, followed by the rAAV5.CMV-IFN-\( \beta \) group. Control and rAAV5.CMV-IFN-\( \beta \)-intron injection did not result in increased IFN-\( \beta \) production.
Increased levels of rat IFN-β mRNA were detected in all animals treated with AAV recombinants. Injection of rAAV5.NF-κB-IFN-β resulted in the highest levels of IFN-β mRNA, whereas in animals treated with the control vector only low amounts of endogenous IFN-β mRNA could be detected by both RT-PCR (Fig. 1a) and real-time PCR (Fig. 1b). IFN-β protein could clearly be detected by immunohistochemical staining (Fig. 1c) and was quantified as described above by computer-assisted image analysis. Injection of rAAV5.NF-κB-IFN-β resulted in the highest synovial production of IFN-β, followed by the rAAV5.CMV-IFN-β injected animals. Control and rAAV5.CMV-IFN-β-intron treated animals showed only endogenous levels of IFN-β (Fig. 1d).

Interestingly, a pronounced reduction in paw swelling was shown after a single intra-articular injection of rAAV5.NF-κB-IFN-β, resulting in sustained amelioration of arthritis until the end of the experiment. In contrast, both rAAV5.CMV-IFN-β and rAAV5.CMV-IFN-β-intron did not lead to significant improvement in paw thickness compared with control animals (Fig. 2a). This is most likely because of the lower intra-articular IFN-β production compared with the rAAV5.NF-κB-IFN-β injected animals, where the promoter is continuously driving expression during inflammation. In addition to the effect on inflammation we investigated the effect on cartilage degradation by Safranin O staining (Fig. 2b), which showed significantly less proteoglycan loss and erosions in treated animals as determined by a semiquantitative scoring system (Joosten et al., 1999; van Holten et al., 2004) (Fig. 2c). Taken together, we demonstrated that the combined use of a low-immunogenic rAAV5 vector and a disease inducible NF-κB-responsive promoter allows prolonged therapeutic effects in an animal model for RA. Thus, application of IFN-β in this context may represent a promising and innovative therapy for the intra-articular treatment of RA.
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