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Cox-2 inhibition enhances the activity of sunitinib in human renal cell carcinoma xenografts

X Wang1,2, L Zhang2, A O’Neill3, B Bahamon4, D C Alsop1, J W Mier2, S N Goldberg1,5, S Signoretti4, M B Atkins6, C G Wood7 and R S Bhatt*,2

1Department of Radiology, Beth Israel Deaconess Medical Center, Harvard Medical School, 330 Brookline Avenue, Boston, MA 02215, USA; 2Division of Hematology-Oncology and Cancer Biology, Beth Israel Deaconess Medical Center, Harvard Medical School, 330 Brookline Avenue, Boston, MA 02215, USA; 3Department of Biostatistics and Computational Biology, Dana-Farber Cancer Institute, 450 Brookline Avenue, Boston, MA 02215, USA; 4Department of Pathology, Brigham and Women’s Hospital, Harvard Medical School, 75 Francis Street, Boston, MA 02115, USA; 5Department of Radiology, Hadassah Hebrew University Medical Center, PO Box 12000, Jerusalem 91120, Israel; 6Georgetown-Lombardi Comprehensive Cancer Center, 3970 Reservoir Road, NW, Washington, DC 20057, USA and 7Department of Urology, Division of Surgery, The University of Texas MD Anderson Cancer Center, 1515 Holcombe-Unit1373, Houston, TX 77030, USA

Background: Sunitinib (Su), a tyrosine kinase inhibitor of VEGFR, is effective at producing tumour response in clear cell renal cell carcinoma (cRCC), but resistance to therapy is inevitable. As COX-2 is a known mediator of tumour growth, we explored the potential benefit of COX-2 inhibition in combination with VEGFR inhibition in attempts at delaying tumour progression on Su.

Methods: COX-2 expression was compared with areas of hypoxia in tumours that progressed on Su vs untreated tumours. Mice bearing human cRCC xenografts were treated with Su and the COX-2 inhibitor, celecoxib, and the effects on tumour growth were assessed. Sequential vs concurrent regimens were compared.

Results: COX-2 expression was increased in cRCC xenografts in areas of tumour hypoxia. The combination of Su and celecoxib achieved longer times to tumour progression compared to treatment with either agent alone or to untreated control animals in four models. This effect was seen with concurrent but not with sequential therapy.

Conclusion: COX-2 inhibition can extend the effectiveness of VEGFR inhibition. This effect is dependent on the timing of therapy. Clinical trials combining Su and COX-2 inhibitors should be considered as a means delaying time to progression on sunitinib in patients with metastatic cRCC.
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2012). While the beneficial effects of TKIs are significant, strategies to enhance and/or prolong the effect of these agents are of importance as responses are almost exclusively partial and tumours develop tumour progression on therapy at a median of less than 1 year.

The mechanisms of resistance to VEGFR inhibition are not fully understood and have been variably reported to involve restored angiogenesis, inhibition of antiangiogenic pathways, and induction of angiogenesis independent processes such as increased tumour invasiveness and epithelial-to-mesenchymal transition (Casanovas et al, 2005; Rini and Atkins, 2009; Bhatt et al, 2010; Hammers et al, 2010; Huang et al, 2010). Understanding the mechanism of resistance may not only guide subsequent treatment selection, but may also provide insights into optimal upfront combination treatment approaches.

Cyclooxygenase-2 (COX-2) is an enzyme in the pathway leading to production of prostaglandin E2 (PGE2) and arachidonic acid (Ghosh et al, 2010). This pathway is known to play a role in inflammation, tumour growth, invasiveness and metastasis, inhibition of apoptosis and angiogenesis. Inhibition of COX-2 has been shown to be a promising antitumour and antiangiogenic strategy in several tumour types including RCC (Hida et al, 2000; Song et al, 2002; Chen et al, 2004a; Arber et al, 2006). In preclinical models, COX-2 inhibition has been shown to exhibit activity as a single agent as well as in combination with immunotherapy and chemotherapy (Veltman et al, 2010; Legge et al, 2011; Vivaldi et al, 2012). COX-2 inhibitors have also been studied in combination with interferon-α therapy in patients with advanced RCC. Although initial reports suggested improved response rates for patients bearing high COX-2 expressing tumours, a subsequent study of the combination of a COX-2 inhibitor and interferon-α confined to this patient population did not substantiate a significant benefit for the combination relative to interferon alone. The value of COX-2 inhibition in combination with VEGFR TKIs has not been formally studied in patients with RCC.

We have previously developed murine models of acquired tumour resistance to VEGFR TKI therapy using human cRCC xenografts. One of these models utilises tumour tissue directly obtained from a patient at the time of nephrectomy for cRCC. The tumour model MDA-62 is a xenograft model of cRCC developed from a tumour fragment harvested from a patient with locally advanced cRCC. Thus, unlike the cell line models that are of clonal origin, this low passage xenograft may better represent tumour heterogeneity noted in the human condition. In this study we explored the role of COX-2 in these models by assessing tumour expression of COX-2 in tumour that have progressed while on sunitinib and studying the effects of the combination of sunitinib and the selective COX-2 inhibitor, celecoxib.

MATERIALS AND METHODS

Tumour xenograft induction. For subcutaneous xenografts the following models were used: A498, 786-O: human cRCC cell lines (ATCC, Manassas, VA, USA), UMRC-3 (Grossman et al, 1985) and MDA-62 a primary tumour from a patient with locally advanced clear cell RCC that was propagated subcutaneously in mice for three passages prior to cryopreservation (Karam et al, 2011). Female nude beige mice (Charles River Laboratories, Wilmington, MA, USA), 6–8 weeks of age and 20 g average body weight, were used. The mice were housed and maintained in laminar flow cabinets under specific pathogen-free conditions. All experiments were approved by the Institutional Animal Care and Use Committee (IACUC) at Beth Israel Deaconess Medical Center. To produce tumours, renal cancer cells were harvested from subconfluent cultures by a brief exposure to 0.25% trypsin and 0.02% EDTA. Trypsinization was stopped with medium containing 10% FBS, and the cells were washed once in serum-free medium and resuspended in PBS. Only suspensions consisting of single cells with > 90% viability were used for the injections.

Cells were injected subcutaneously (1 × 10⁶ cells) into the flanks of the mice. Tumours developed in > 80% of the mice and were usually visible within a few days of implantation and once they reached a diameter of 3–5 mm, were measured daily to ensure a consistent size at the outset of treatment. Vehicle, sunitinib (additive-free, 53.6 mg kg⁻¹ daily by gavage), celecoxib (50 mg kg⁻¹ daily by gavage) (Farooqui et al, 2007; Park et al, 2008; Li et al, 2011), or the combination was begun when the tumours had grown to a diameter of 12 mm as per our previous reports (Bhatt et al, 2010; Zhang et al, 2011). Tumours were measured daily during therapy to determine growth curves.

In all, 12 mm was used as the pre-specified treatment start size in the 786-O and A498 models in part because this size of tumour may be sufficiently large to be proportionately comparable to a lesion in the human clinical setting, but not too large to prevent a period of several weeks before the mice would need to be sacrificed at the end point of 20 mm tumour size. It is also an optimal size for tumour perfusion imaging, which is difficult to perform on smaller tumours. Growth by 2 mm from this baseline is the minimal reproducible change in tumour size that could be accurately measured by our calipers. Time to increase by 2 mm was defined as time to tumour progression. This approach is roughly equivalent to the increase in tumour size (20% increase in long axis) that would classify a patient as having progressive disease by RECIST criteria (and therefore, treatment resistance) while receiving such therapy, as it represents a 17% increase in diameter and 59% increase in volume. Additionally, in our analysis, the tumour growth curves showed similar results with both tumour volume and long axis measurements (Figures 2A and B). In this model, progression also correlates with the resumption in tumour perfusion that follows an initial reduction in perfusion, seen in our prior studies (Schorbardach et al, 2009; Bhatt et al, 2010).

Treatment was continued until tumours grew to 20 mm in long-axis diameter (i.e., the maximum allowable growth by IACUC) at which point the mice were sacrificed. Tumour tissue from sunitinib, celecoxib, and untreated tumours was obtained at 20 mm for various analyses described below. Sunitinib-treated tumours that grew beyond 14 mm and grew to the end point of 20 mm size despite continued treatment were defined as ‘Resistant to sunitinib’.

In the studies with UMRC-3 and MDA-62, treatment was started when tumours reached a size of 5 mm in maximum diameter. Tumour size was measured daily and mice were sacrificed after 30 days of treatment. The pretreatment tumour size was less in this model because the study design differed from the xenograft model in which imaging was performed at 12 mm for optimal imaging resolution.

Immunohistochemistry. For hypoxia analysis, animals were injected intraperitoneally with Hypoxyprobe (Pimonidazole HCl: Hypoxyprobe store, NPI, Cat # HP1-100 kit, Burlington, MA, USA) 100 µl 2 h prior to sacrifice. For COX-2 and Hypoxyprobe analysis, 4 µm thick sections were prepared from formalin-fixed, paraffin-embedded tumour specimens. Sections were deparaffinized, rehydrated and heated with a pressure cooker to 125 °C for 30 s in citrate buffer for antigen retrieval. After cooling to room temperature, sections were incubated in 3% hydrogen peroxide for 5 min to quench endogenous peroxidase, (Dako, Carpinteria, CA, USA). The primary antibodies (anti-COX-2 antibody from CAYman Chemical, Ann Arbor, MI, USA, Cat #160112; anti-Hypoxyprobe antibody from Natural Pharmacia International, Burlington, MA, USA, Cat#FITC-MAb1) were applied at a 1 : 50
dilution, to sections for 1h. Detection was performed by incubation with Dako EnVision + System HRP labelled polymer anti-mouse (Dako, Cat# K4006) for 30 min followed by incubation with DAB chromogen. Slides were slightly counterstained with hematoxylin. Slides were scanned using the Scanscope XT (Aperio Technologies Inc., Visa, CA, USA).

**Tumour perfusion imaging.** Tumour perfusion imaging with arterial spin-labelling MRI (ASL MRI) was performed as previously described (Bhatt *et al*, 2010). Briefly mice were anaesthetized, and placed in the supine position on a 3-cm diameter custom-built surface coil. Images were acquired using a 3.0 T whole-body clinical MRI scanner (3T HD; GE Healthcare Technologies, Waukesha, WI, USA). A single slice ASL image was obtained with a single-shot fast spin echo sequence (SSFSE) using a background-suppressed, flow-sensitive alternating inversion-recovery strategy. Twenty-four label and control pair images were acquired and averaged for the ASL acquisition. A reference proton density image was acquired by turning off all background suppression and labelling pulses in the ASL preparation. T1 measurement was performed by using the same imaging sequence at same slice location but with inversion recovery at different inversion times. The single transverse slice of ASL was carefully positioned at the centre of the tumour, which was marked on the skin with a permanent marker pen for follow-up MRI studies. ASL raw data were reconstructed and converted to quantitative tumour perfusion maps using automated, custom software written within the Interactive Data Language (IDI; ITT visual Information Solutions, Boulder, CO, USA) (Alsop and Detre, 1996). Regional analysis of the perfusion images was performed with Image-J software (Image Processing and Analysis in Java; National Institutes of Health, Bethesda, MD, USA). To determine tumour perfusion, a region of interest was drawn freehand around the peripheral margin of the tumour by using an electronic cursor on the reference image that was then copied to the perfusion image. The mean blood flow for the tumour tissue within the region of interest was derived.

A 16-colour table was applied in 10 ml per 100 g per min increments ranging from 0 to 160 ml per 100 g per min, with flow values represented as varying shades of black, blue, green, yellow, red, and purple, in order of increasing perfusion.

**Statistical analysis.** The primary end point was time to progression (TTP). TTP was defined as time in days to increase by 2 mm in tumour size and compared between groups via the exact Wilcoxon rank sum test. Median TTP in days and ranges are reported throughout.

Figure 1. Sunitinib resistance is accompanied by enhanced COX-2 expression in areas of tumour hypoxia. Representative low (A, C, E, G, I, K, M, O) (scale bar = 300 μm) and high (B, D, F, H, J, L, N, P) (scale bar = 100 μm) magnification microscopic images of cRCC xenografts analysed by immunohistochemistry. 786-O tumours treated with sunitinib (A–D) or vehicle (E–H) and A498 tumours treated with sunitinib (I–L) or vehicle (M–P) were immunostained for Hypoxyprobe (A, B, E, F, I, J, M, N) or COX-2 (C, D, G, H, K, L, O, P).
RESULTS

Sunitinib-treated tumours exhibit enhanced hypoxia which correlates with COX-2 expression. Since sunitinib treatment reduces tumour blood flow (Bhatt et al, 2010; Zhang et al, 2011), we assessed the effect of this agent on tumour hypoxia. We found that sunitinib treatment of either A498 or 786-O tumours leads to tumour hypoxia which is present, as shown in Figure 1, at the time of resistance to sunitinib. Pimonidazole HCl-positive hypoxic tumour cells are consistently observed around areas of tumour necrosis (Figure 1A, B, I, and J) while only scattered hypoxic tumour cells are detected in untreated tumours (Figure 1E, F, M, and N). As COX-2 expression is known to be induced by hypoxia, we also assessed the COX-2 expression in sunitinib resistant tumours. As shown in Figure 1, both 786-O (Figures 1C and D) and A498 (Figure 1K and L) sunitinib resistant tumours exhibit a spatially regulated area of increased COX-2 expression, with maximal expression in the hypoxic tumour cells surrounding areas of necrosis in 786-O (C,D) and A498 (K,L) tumours. In contrast, consistent with the absence of significant tumour hypoxia, COX-2 expression does not occur in such a specific pattern in vehicle-treated tumours of the same size. Untreated tumours did not exhibit hypoxia and showed diffuse COX-2 expression (Figure 1E–H and M–P). Thus, sunitinib resistance is accompanied by an increase in COX-2 expression levels in areas of hypoxic tumour cells.

Combining COX-2 inhibition with sunitinib enhances anti-tumour efficacy and delays disease progression on sunitinib. We hypothesised that the increased COX-2 expression seen in sunitinib resistant tumours might indicate a role for COX-2 inhibition in enhancing efficacy of and/or delaying tumour progression on sunitinib. Mice harbouring A498 and 786-O xenografts were treated with vehicle, sunitinib, celecoxib, or sunitinib + celecoxib. Figure 2A–C show the tumour growth by tumour volume, length, and time to progression in the four treatment arms in the mice bearing A498 tumours. The
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**Concurrent therapy is superior to sequential therapy.** To assess
the temporal relationship between sunitinib and celecoxib admin-
istration, we compared adding celecoxib vs switching to celecoxib
at the time of initial tumour progression on sunitinib in the 786-O
model and determined the time to increase by another 2 mm in
long axis. Figure 4 shows that switching to celecoxib (Sunitinib →
Celecoxib) is significantly worse than continuous sunitinib
(sunitinib continuous) (median (range) 10 (5–10) days, n = 6 vs
12 (10–13) days, n = 7, respectively, P = 0.003) and that delayed
addition of celecoxib (sunitinib → celecoxib + sunitinib) (median
(range) 10 (7–13) days, n = 7) did not improve the efficacy of
continuous sunitinib. Thus, early administration of celecoxib
appears to be necessary for it to enhance the efficacy of sunitinib
therapy in this model.

Celecoxib augments the activity of sunitinib in an angiogenesis-
independent manner. As celecoxib has been shown to exhibit
angiogenic activities in addition to many other antitumour
activities, we first assessed the effects of celecoxib on the
angiogenic activity of sunitinib. Tumour perfusion was
measured in the 786-O tumour model by ASL MRI, a technique
which correlates with microvascular density, at days 0, 2, 4, and 8
(Figure 5). While sunitinib (n = 2) treatment led to a pronounced
drop in tumour perfusion at all time points compared with vehicle
(n = 3), celecoxib (n = 2) did not decrease tumour perfusion as a
single agent or further decrease perfusion when added to sunitinib
(n = 3). Although sample sizes are small, preventing statistical
data accuracy, trends are apparent. Representative perfusion images of a
single tumour slice are shown in Figure 5B.

**DISCUSSION**

In this study we show that resistance to sunitinib is accompanied
by increased COX-2 expression in areas of tumour hypoxia in two
cRCC xenograft models. We also show that the COX-2 inhibitor
celecoxib enhances the effectiveness of sunitinib in four cRCC
xenograft models by delaying time to progression when adminis-
tered early in the course of sunitinib therapy. Celecoxib as a single
agent showed activity only in the 786-O tumour model. However,
in all models the combination of sunitinib and celecoxib showed
greater inhibition of tumour growth than either agent alone. The
MDA-62 model may be more relevant to patients as the xenograft

**Figure 3.** Celecoxib enhances the anti-tumour activity of sunitinib in additional human xenograft clear cell cRCC models. Comparison of volume (mm³) over time (days) following treatment with vehicle, celecoxib, sunitinib, or combination of celecoxib and sunitinib, in two different human xenograft cRCC models: MDA-62 (A) and UMRC-3 (B). N = 3 tumours in each arm. Data are shown as average ± s.e.m.
was obtained directly from a patient undergoing surgery for cRCC, implanted directly into the subcutaneous space of female BALB/c nude mice and serially passaged into new mice without ever being propagated in vitro. Because it is not clonal in origin, but rather developed from a fragment of human tumour, it may be more reflective the heterogeneity in tumour biology associated with the human condition. Although different pretreatment sizes were used in the patient tumour model and xenograft models due to differences in imaging requirements, similar treatment trends were obtained. In general the effects of treatment are more pronounced when tumours are smaller prior to treatment. Further experiments to clarify the differences in effect vs pretreatment sizes should be considered. Experiments with this human tumour model further strengthen the findings noted with the clonally derived cell lines.

COX-2 inhibition has been shown to have antitumour activity in cRCC and is postulated to function via a variety of antitumour and antiangiogenic mechanisms (Chen et al, 2004a and b; Sato et al, 2010). COX-2 expression has also been reported to correlate with cRCC prognosis. Although the precise relationship is not well-understood, most published studies report COX-2 expression to correlate with poor clinicopathologic features and predict a poor prognosis (Miyata et al, 2003; Tuna et al, 2004; Sozen et al, 2008; Li et al, 2009). One report, however, found that COX-2 expression predicted for a longer median overall survival (Kankuri-Tammilehto et al, 2010). While it remains unclear how to reconcile these seemingly disparate findings, it is possible that the discrepancies could relate to the clinical context in which the tissue was obtained. In this regard, our studies showing enhanced

Figure 5. Measurements of perfusion demonstrate that the celecoxib does not affect perfusion despite significant tumour size effect. Bar graphs of average tumour perfusion as measured by ASL MRI with SD are shown at four time points in (A). The treatment arms are as follows: sunitinib (n = 2), celecoxib (n = 2), sunitinib + celecoxib (n = 3), and vehicle (n = 3). Representative images from the four treatment arms are shown in (B). Representative tumour regions are outlined with yellow line in ASL perfusion images (right column of each time point) and the corresponding reference proton density images (left column). The overall tumour size was measured with long and short axes (in mm), one axis is shown in this figure, and the mean blood flow (in ml per 100 g per min) are shown below each image. A colour bar on bottom represents range of perfusion values from 0 to 160 ml per 100 g per min.
COX-2 expression at the time of sunitinib resistance, particularly adjacent to areas of tumour necrosis, suggest that this expression may be both heterogeneous and inducible as a consequence of VEGFR TKI-induced hypoxia within the cRCC xenografts. In this study we cannot rule out the possibility that COX-2 expression appears to be increasing in areas of hypoxia and actually decreasing in non-hypoxic areas. This possibility is less likely since COX-2 is known to be positively regulated by hypoxia.

COX-2 inhibition has also been studied in clinical trials of patients with advanced cRCC. Rini et al (2006) have examined the relationship between COX-2 expression and response to celecoxib and interferon-α in phase II trials (Schwandt et al, 2011). An initial trial in patients with metastatic cRCC introduced the hypothesis that maximal COX-2 immunostaining may directly correlate with response to the combination (Rini et al, 2006). However, a subsequent study of the combination confined to patients with high COX-2 expressing tumours did not substantiate a significant benefit for the combination relative to interferon alone (Schwandt et al, 2011). Another clinical trial combining the COX-2 inhibitor, meloxicam, and IFN-α in 43 patients with metastatic cRCC showed an overall response rate of 37.2% with a median time to progression of 14 months (Shinohara et al, 2009). However, at this point, no clinical studies have examined the role of COX-2 inhibition in combination with a VEGFR TKI.

Evasive resistance to antiangiogenic therapy in cRCC is likely multi-factorial. Several potential mechanisms of resistance have recently been reported (Casanovas et al, 2005; Rini and Atkins, 2009; Bhatt et al, 2010; Hammers et al, 2010; Huang et al, 2010). We have shown that VEGFR blockade results in prompt tumour devascularisation and likely hypoxia-driven tumour necrosis(Schor-Bardach et al, 2009). Thus, upregulation of hypoxia-mediated mechanisms of resistance is likely an early event post-VEGFR TKI therapy. Although our prior studies have shown that resumed angiogenesis and restored perfusion was one mechanism of acquired resistance to sunitinib, we did not find that administration of celecoxib led to decreased perfusion by ASL MRI in our models. Thus, celecoxib is likely acting by a distinct mechanism from sunitinib, which potently reduces tumour perfusion. It is conceivable that as VEGFR2 blockade persists, the mechanisms contributing to resistance become more diverse. Our timing experiments show that the activity of celecoxib depends on dynamic changes induced by sunitinib treatment. Further experiments in tumour models that include tumour cell-endothelial cell interactions are needed and are planned.

Optimally, in the future, application of surrogate biomarkers such as PGE-M, a urinary metabolite of PGE2 (Murphey et al, 2004; Johnson et al, 2006; Reckamp et al, 2011), might enable institution of treatment at the time of COX-2 upregulation, thereby sparing the cost and toxicity associated with upfront combination therapy. If a biomarker for COX-2 activity were available, patients with metastatic RCC being treated with a VEGFR pathway inhibitor could be monitored over the course of therapy and celecoxib added at the time of biomarker elevation and compared with continued VEGFR pathway inhibitor alone. However, in the absence of such biomarkers, our data support the study of the initial combination of celecoxib and sunitinib in patients with advanced cRCC.

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