C-type natriuretic peptide in combination with sildenafil attenuates proliferation of rhabdomyosarcoma cells

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Abstract

Rhabdomyosarcoma (RMS) is a malignant mesenchymal tumor and the most common soft tissue sarcoma in children. Because of several complications associated with intensive multimodal therapies, including growth disturbance and secondary cancer, novel therapies with less toxicity are urgently needed. C-type natriuretic peptide (CNP), an endogenous peptide secreted by endothelial cells, exerts antiproliferative effects in multiple types of mesenchymal cells. Therefore, we investigated whether CNP attenuates proliferation of RMS cells. We examined RMS patient samples and RMS cell lines. All RMS clinical samples expressed higher levels of guanylyl cyclase B (GC-B), the specific receptor for CNP, than RMS cell lines. GC-B expression in RMS cells decreased with the number of passages in vitro. Therefore, GC-B stable expression lines were established to mimic clinical samples. CNP increased cyclic guanosine monophosphate (cGMP) levels in RMS cells in a dose-dependent manner, demonstrating the biological activity of CNP. However, because cGMP is quickly degraded by phosphodiesterases (PDEs), the selective PDE5 inhibitor sildenafil was added to inhibit its degradation. In vitro, CNP, and sildenafil synergistically inhibited proliferation of RMS cells stably expressing GC-B and decreased Raf-1, Mitogen-activated protein kinase kinase (MEK), and extracellular signal-regulated kinase (ERK) phosphorylation. These results suggested that CNP in combination with sildenafil exerts antiproliferative effects on RMS cells by inhibiting the Raf/MEK/ERK pathway. This regimen exerted synergistic effects on tumor growth inhibition without severe adverse effects in vivo such as body weight loss. Thus, CNP in combination with sildenafil represents a promising new therapeutic approach against RMS.
Introduction

Rhabdomyosarcoma (RMS), a malignant mesenchymal tumor, is the most common soft tissue sarcoma in children. RMS is classified histopathologically and biologically into two major types: embryonal (ERMS) and alveolar (ARMS) [1]. Although clinical outcomes for RMS patients have improved over the past two decades due to advances in multimodal therapy including surgery, radiotherapy, and chemotherapy, RMS cases with high-risk features such as relapse or systemic metastatic disease have a poor prognosis and limited treatment options [2]. Furthermore, late complications including growth disturbance and secondary cancer remain important problems to be solved. Consequently, novel therapies with reduced toxicity and improved tolerability relative to conventional chemotherapy are urgently required.

C-type natriuretic peptide (CNP) is an endogenous peptide secreted by vascular endothelial cells. The natriuretic peptide family includes atrial and brain natriuretic peptides (ANP and BNP, respectively) [3]. CNP inhibits the proliferation of several types of mesenchymal cells, including vascular smooth muscle cells [4, 5], kidney mesangial cells [6–8], and fibroblasts [9, 10]. However, it remains unknown whether CNP can inhibit proliferation of RMS cells.

CNP binds to and activates the guanylyl cyclase B (GC-B) receptor, resulting in production of cyclic guanosine monophosphate (cGMP) as a second messenger and activation of downstream molecules [11]. Cyclic GMP is quickly degraded to GMP by phosphodiesterases (PDEs). Because PDE5 is a major negative regulator of cGMP, PDE5 inhibitors such as sildenafil, currently used for treatment of erectile dysfunction [12] and pulmonary hypertension [13], have been used in conjunction with natriuretic peptides or other cGMP inducers to prevent degradation of cGMP by PDE5.

In this study, we demonstrated that CNP attenuates proliferation of RMS cells and that sildenafil potentiates the antiproliferative effects of CNP, and examined the underlying mechanisms responsible for this effect.

Materials and Methods

RMS tissue from patients

Tissue from two ERMS patients and three ARMS patients was obtained from the Department of Pediatric Surgery, Osaka University Graduate School of Medicine. Written informed consent to use resected specimens for this study was obtained from all the patients/parents/guardians, and the study was approved by the Institutional Review Board (IRB) of Osaka University Hospital (IRB number: 15022). It conforms to the provisions of the Declaration of Helsinki in 1995. Patient and sample characteristics are provided in the Table 1.

Cell culture and establishment of cells stably expressing GC-B

The human RMS cell lines RD and KYM-1, derived from ERMS, were purchased from Japanese Collection of Research Bioresources Cell Bank (Ibaraki, Japan), RMS-YM cells derived from ERMS were purchased from Riken Cell Bank (Tsukuba, Japan), and RH30 cells derived from ARMS were purchased from American Type Culture Collection (ATCC) (Rockville, MD). All cell lines were cultured in RPMI-1640 medium (Life Technologies, Grand Island, NY) supplemented with 10% fetal calf serum (Invitrogen GIBCO, Tokyo, Japan) in 5% CO₂ in air at 37°C.

The plasmid used for preparing retrovirus vector expressing GC-B-FLAG (pCX4 puro GC-B-FLAG) was constructed by introducing the GC-B-FLAG fragment, amplified by polymerase chain reaction (PCR) from human GC-B obtained from Promega (Madison, WI), into the NotI site of pCX4 puro. The vector was transfected into BOSC cells with pE-eco and pGp (TAKARA BIO, Kyoto, Japan) using FuGENE6 (Promega) to produce ecotropic retrovirus. After transfection, culture supernatants were collected, and filtered. Retroviral infection of RD and RMS-YM cells was performed using the retrovirus and Ecotropic Receptor Booster (TAKARA BIO). After infection, cells were selected with puromycin. These resultant cell lines are referred to as RD-GC-B or RMS-YM-GC-B.

Table 1. Patient and sample characteristics.

| Case | Age | Sex | Type | Primary site | Metastasis | Follow-up | Primary or recurrent |
|------|-----|-----|------|-------------|------------|-----------|---------------------|
| 1    | 2Y11M | M   | Embryonal | Prostae | –          | Alive     | Primary              |
| 2    | 9Y0M   | M   | Alveolar | Abdominal wall | –          | Alive     | Primary              |
| 3    | 13Y3M  | M   | Alveolar | Anterachial region | Axillary LN | Alive     | Primary              |
| 4    | 3Y10M  | F   | Embryonal | Retropertioneum | –          | Expired   | Primary              |
| 5    | 11M    | M   | Alveolar | Gastrocnemius | Retro-peritoneum (recurrence) | Alive | Recurrence         |

LN, lymph node.
Reagents

CNP was purchased from Peptide Institute (Osaka, Japan) and sildenafil was purchased from Sigma-Aldrich (St. Louis, MO). Antibodies were purchased as follows: antiphosphorylated extracellular-signal-regulated kinase (ERK)1/2 (Thr202/Tyr204), anti-total ERK and anti-GAPDH from Santa Cruz Biotechnology (Dallas, TX), antiphosphorylated Raf-1 (Ser338), antiphosphorylated MEK1/2 (Ser217/221), hairy/enhancer of split (HES-1), β-catenin, GLI1, and phosphorylated Akt (Ser473) from Cell Signaling Technology (Beverly, MA), and anti-Ki67 antibody (clone MIB-1), as a marker for cell proliferation, from Dako Japan (Kyoto, Japan). Caspase-3 activity was evaluated using a Nuc-View™ 488 Caspase-3 assay kit (Biotium Inc., Cam-bridge, UK). The cells were incubated at room temperature for 30 min with 1 μmol/L Nuc-View 488 substrate solution.

Gene expression analysis

Total RNA was isolated from cell lines and patient tissue samples homogenates using guanidinium-phenol-chloroform extraction and the RNeasy mini kit (Qiagen, Hilden, Germany). The obtained RNA was reverse-transcribed into complementary DNA using a QuantiTect Reverse Transcription Kit (Qiagen). Quantitative PCR assays were performed using the SYBR Premix Ex Taq kit (TAKARA Bio) on a Light Cycler 480 System II (Roche Applied Science, Indianapolis, IN). The following primers were used: GC-B, sense 5′-TAAGAATGAGCATCCAGATACAG-3′ and antisense 5′-TCATTGTGGGAGCAGACATG-3′ and antisense 5′-GCCGAGCATCCAGATAAGACAAAGG-3′; 36B4, sense 5′-TCATTGTGGGAGCAGACATG-3′ and antisense 5′-AGGTCCTCCTTGGTGTAACACAAAGC-3′. Quantitation of gene expression was normalized to the housekeeping gene 36B4.

Measurement of intracellular cyclic GMP

Rhabdomyosarcoma cells were seeded onto 24-well plastic plates and incubated for 24 h. After the cells were confluent as a monolayer, they were washed twice with 500 μL of serum-free RPMI-1640 and incubated in serum-free medium for 24 h. CNP was dissolved in 5 mmol/L isobutylmethylxanthine (IBMX) at final concentrations of 1 mmol/L–10 μmol/L. Various concentrations of CNP were added to the medium, and the monolayer was incubated at 37°C for an additional 15 min. The medium was rapidly removed by aspiration, and 400 μL of 70% ethanol containing 100 μmol/L hydrochloric acid (HCl) was added to lyse the cells. The cGMP content of the dried extract was quantitated using a radioimmunoassay (RIA) kit (Yamas Shoyu, Tokyo, Japan).

Cell proliferation assay

A total of 2000 cells per well were seeded into 96-well plates in 100 μL of culture medium. After 24 h, the wells were washed twice with 1% serum-free RPMI-1640 medium and subsequently incubated in this medium. Twenty-four hours after serum deprivation, CNP (1 μmol/L) and/or sildenafil (100 μmol/L) were added, and the cells were incubated for 0–4 days. Cell proliferation was determined in sextuplicate using the Cell Counting Kit-8 (Nacalai Tesque, Kyoto, Japan).

Treatment of cells with CNP and sildenafil, preparation of cell lysate, and western blot analysis

A total of 1 × 10^5 cells per well were plated in 12-well plastic plates in complete media. Cells were serum-starved with RPMI-1640 medium containing 1% serum for 24 h, and then treated with CNP (1 μmol/L) and/or sildenafil (100 μmol/L) for indicated times. Cells were lysed with NP-40 buffer (0.5% NP-40, 10 mmol/L Tris [pH 7.6], 0.15 mol/L NaCl, 1.5 mmol/L MgC92) containing phosphatase inhibitor cocktail (Nacalai Tesque). Samples were electrophoresed on 4–15% gradient gels (Bio-Rad, Hercules, CA, USA) and transferred onto polyvinylidene difluoride transfer membranes (Millipore Bedford, MA). Antibody binding was detected with an enhanced chemiluminescence detection system (Amersham Pharmacia Biotech, Piscataway, NJ).

Xenograft efficacy studies

All animal experiments were performed according to a protocol approved by the Animal Care Ethics Committee of the National Cerebral and Cardiovascular Center Research Institute. Five-week-old male BALB/c nu/nu mice and 5-week-old male nonobese diabetic/severe combined immunodeficiency (NOD)severe combined immunodeficiency (SCID) mice were purchased from Japan SLC (Shizuoka, Japan) and Oriental Yeast (Tokyo, Japan), respectively. RD-GC-B cells (1–3 × 10^7 cells) were subcutaneously implanted into the chest and abdominal area of NOD/SCID mice. When the subcutaneous tumors reached a diameter of 10–12 mm, they were resected, divided into 5-mm pieces, and reimplemented into the chest area of BALB/c nu/nu mice. Ten days after tumor implantation, administration of CNP and/or sildenafil was initiated. CNP was continuously infused using an osmotic mini-pump (Alzet Model 1003D; Duret Corporation, Cupertino, CA), as previously reported [14]. CNP was subcutaneously administered at a rate of 2.5 μg/kg/min, and treatment continued for 4 weeks. Sildenafil was intraperitoneally administered at a dose of 20 mg/kg every other day for...
4 weeks. Tumor volume was calculated by the formula: \( V = 0.5 \times a \times b^2 \), where \( a \) is the longest tumor axis and \( b \) is the shortest tumor axis. Upon termination of the experiment, after blood was withdrawn from the inferior vena cava, tumors were excised and weighed, and tumor samples were processed for immunohistochemical and Western blot analysis.

**Immunohistochemistry**

Immunohistochemical staining was performed by the avidin-biotin-peroxidase complex (ABC) method using the Vectastain Elite ABC kit (Vector, Burlingame, CA) as described previously [14]. Because the anti-Ki67 antibody was mouse monoclonal antibody, a mouse on mouse immunodetection kit (Vector) was used. Images of immunohistochemical staining were captured using a FSX100 Bio Imaging Navigator microscope (Olympus, Tokyo, Japan). Ki67-positive and -negative cells were counted using the CellSens Dimension software (Olympus). Ten randomly selected fields per section and 7–8 sections per animal were analyzed at 20 × magnification.

**Plasma CNP measurements**

Osmotic mini-pumps containing CNP were implanted subcutaneously in 6-week-old male C57BL/6 mice purchased from Japan SLC. CNP was administered at a rate of 2.5 μg/kg/min, and blood samples from these treated mice and untreated mice were withdrawn 3 days after the implantation. CNP content was determined using a CNP RIA kit (Phoenix Pharmaceuticals, Belmont, CA).

**Statistical analysis**

Data were analyzed using GraphPad Prism version 6.0 (GraphPad Software, San Diego, CA), and expressed as means ± standard error. Between-group comparisons were performed using the Welch’s \( t \) test or Mann–Whitney \( U \) test. Multiple-group comparisons were performed using one-way or two-way ANOVA, followed by the post hoc Tukey–Kramer (pairwise comparisons) or Dunnett test (comparisons with controls). \( P < 0.05 \) was considered significant.

**Results**

**GC-B expression levels in RMS cell lines are lower than those in RMS tumor samples**

GC-B expressions in tumor samples of RMS patients and RMS cell lines were initially evaluated. All samples from RMS patients and two RMS cell lines, RD and RMS-YM cells, expressed GC-B. In contrast, KYM-1 and RH-30 cells had no detectable GC-B mRNA expression (Fig. 1A). To determine whether the GC-B receptors in RD and RMS-YM cells were functional, we measured cGMP production in response to CNP treatment. CNP increased the intracellular cGMP levels in a dose-dependent manner (Fig. 1B). In these cells, cGMP levels almost reached a plateau at CNP concentration of 1 μmol/L (Fig. 1B). Therefore, in the following in vitro experiments, we used 1 μmol/L CNP.

Next, we compared GC-B expression levels between tumor samples from RMS patients and RMS cell lines. The RMS patient samples had 5–26-fold higher GC-B expression than the RD or RMS-YM cell lines. The recurrent tumor sample (Case 5) had significantly higher GC-B expression than the samples from primary tumors (Fig. 1A). Additionally, we noticed that the amount of GC-B expression in RD cells decreased with the number of passages (Fig. S1), as previously reported for smooth muscle and osteoblast cells [15, 16]. For these reasons, we established RD and RMS-YM cells stably expressing GC-B to mimic the RMS patient samples (Fig. S2). Hereafter, we refer to these cell lines as RD-GC-B and RMS-YM-GC-B, respectively. These cells exhibited robust increases in GC-B expression and CNP-GC-B dependent cGMP production (Fig. 1C).

**CNP/GC-B/cGMP system attenuates RMS cell proliferation, and sildenafil potentiates the antiproliferative effect**

We tested the effects of CNP/GC-B/cGMP signaling system on proliferation of RD-GC-B and RMS-YM-GC-B cells. As shown in Figure 2, cell proliferation was significantly attenuated in both RD-GC-B (Fig. 2A) and RMS-YM-GC-B cells (Fig. 2B) 96 h after CNP treatment. The effects of cGMP can be enhanced and prolonged by suppressing cGMP degradation using sildenafil, a PDE5 inhibitor [25]. Therefore, we examined the effects of sildenafil on cell proliferation of RMS cells by CNP. Combination treatment with sildenafil and CNP significantly decreased the proliferation of RD-GC-B (Fig. 2A) and RMS-YM-GC-B cells (Fig. 2B) relative to treatment with CNP or sildenafil alone. Morphological appearance of cell death was not observed during 96 h in treatment with sildenafil and/or CNP. In addition, caspase-3 activity was not significantly induced in RD-GC-B cells after the treatment with CNP and/or sildenafil, when we evaluated apoptosis under the same conditions of the cell proliferation assay using a Nuc-View™ 488 Caspase-3 substrate (data not shown). Together, these findings demonstrate that the CNP/GC-B/cGMP system attenuates proliferation of GC-B–expressing ERMS cells, and that PDE5 inhibition synergistically potentiates this antiproliferative effect.
Figure 1. GC-B is upregulated in rhabdomyosarcoma (RMS) tumor samples, but is expressed at significantly lower levels in RMS cell lines. (A). GC-B mRNA levels (normalized against 36B4, used as an internal control), as determined as analyzed by quantitative PCR, in five RMS patient samples (white) (case 1 and 4: embryonal type; case 2, 3 and 5: alveolar type, case 1–4: primary tumor; case 5: recurrent tumor) and four RMS cell lines (black). (B). CNP bioactivities in RD and RMS-YM cells, determined by measurement of intracellular cyclic guanosine monophosphate (cGMP) levels. The data represent the average from four wells with the indicated standard error. (C). CNP bioactivities in RD-GC-B and RMS-YM-GC-B cells determined by measurement of intracellular cGMP levels.

CNP inhibits Raf/MEK/ERK signaling, and sildenafil enhances and prolongs the inhibition

Next, we investigated the mechanism underlying inhibition of RMS cell proliferation by sildenafil and CNP. In RMS cells, several signaling pathways including Notch [17, 18], Wnt [19], Hedgehog [20], phosphoinositide 3-kinase (PI3K)/AKT/mTOR [21, 22], and MAPK/ERK [23, 24] are involved in promoting cell proliferation. When we treated RD-GC-B and RMS-YM-GC-B cells with CNP, we observed dephosphorylation of ERK (Figs. 3A and B). Although sildenafil alone did not cause dephosphorylate ERK during 8 h of treatment, it enhanced and prolonged the dephosphorylation of ERK by CNP (Figs. 3A and B). Dephosphorylation of Raf-1 and MEK (the kinase upstream of ERK) by CNP was enhanced by sildenafil (Figs. 3C and D). Expression of HES-1, β-catenin, and GLI1, as well as phosphorylation of Akt, were unchanged by CNP and sildenafil treatment in RD-GC-B cells, suggesting that the Notch, Wnt, Hedgehog, and PI3K/AKT/mTOR signaling pathways were not regulated by CNP in RD-GC-B cells (Fig. S3). Additionally, CNP had no effect on proliferation or reduction in ERK phosphorylation in parental RD and RMS-YM cells (data not shown). Overall, these data suggest that CNP/GC-B/cGMP system attenuates the proliferation of RMS cells by inhibiting the Raf/MEK/ERK signaling pathway.
CNP and sildenafil synergistically inhibit growth of RD-GC-B xenograft tumors

Next, we evaluated the long-term effects of CNP and sildenafil on tumor growth in vivo. As shown in Figure 4A, treatment with CNP or sildenafil alone did not suppress tumor growth significantly; however, the combination treatment significantly suppressed tumor growth and reduced tumor weight relative to treatment with vehicle alone (Fig. 4A). Pharmacokinetic analysis of CNP revealed that the plasma concentration of CNP was significantly increased by continuous subcutaneous infusion of CNP (Fig. 4B). Ki67 labeling index was significantly reduced in the CNP, sildenafil, and sildenafil plus CNP groups.
relative to the control group (Fig. 4C). Phosphorylated ERK levels were equally reduced, to almost 30% of control levels, in both the CNP and sildenafil groups, and were further reduced to almost 10% of control levels in the sildenafil plus CNP group (Fig. 4D). To evaluate tolerability, we assessed the body weights of the mice during treatment (Fig. 4E).
CNP and Sildenafil Attenuate RMS Proliferation

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Discussion

Recent studies showed that several cGMP inducers regulate tumor cell proliferation. Green tea polyphenol (−)-epigallocatechin-3-O-gallate increases cGMP to inhibit tumor growth by activating apoptotic signaling via the Akt/eNOS/NO/cGMP/PKG pathway in multiple myeloma, gastric, pancreatic, and prostate cancers [25]. The nonsteroidal anti-inflammatory drug (NSAID) sulindac sulfide inhibits colon cancer cell growth by suppressing Wnt/β-catenin signaling via the cGMP/protein kinase G (PKG) pathway [26]. In this study, we examined the antiproliferative effects of CNP/GC-B/cGMP system on RMS.

Our data demonstrate that GC-B expression decreases with passage number in RD cells. It is the same in smooth muscle cells and osteoblasts due to oxidative stress resulting in DNA damage accumulation [16]. Meanwhile, in chondrocytes, GC-B expression increases with passage number due to the differentiation into fibroblast-like cells [27]. Therefore, they might depend on the cell types. Additionally, the differentiation may cause the higher levels of GC-B expression in the recurrent tumor sample than in the primary tumor samples.

Phosphorylation of ERK and proliferation of two RMS cell lines were inhibited by CNP. The MAPK/ERK pathway plays a key role in promoting cell proliferation [23, 24]. Activating mutations of N- or K-Ras in RMS clinical samples, and in N-Ras in the RD cell line, lead to uncontrolled cell growth [28–30]. RD cells harbor the N-Ras Q61H mutation, which constitutively activates MEK/ERK. Consistent with this, phosphorylation of ERK was detected in RD cells in the absence of stimulation (Fig. 3A). Similarly, phosphorylation of ERK was observed in unstimulated RMS-YM cells (Fig. 3B). Although N-Ras is active in RD cells, CNP dephosphorylated ERK, suggesting that CNP negatively regulates downstream effectors of N-Ras such as Raf-1 or MEK. Several studies have described the relationship between CNP and the MAPK/ERK pathway. In rat chondrosarcoma chondrocytes, fibroblast growth factor (FGF)-2-mediated activation of MAPK/ERK is inhibited by CNP/GC-B/cGMP signaling at the level of Raf-1 [31]. Moreover, cGMP signaling phosphorylates Raf-1 at Ser43, resulting in the uncoupling of the Ras/Raf-1 interaction and inactivation of the Raf/MEK/ERK pathway [32]. We found that CNP attenuated phosphorylation of Raf-1 in RMS cells (Figs. 3C and D). These findings suggest that CNP/GC-B/cGMP signaling is a potent inhibitor of Raf/MEK/ERK pathway at the level of Raf-1 in RMS cells.

Sildenafil synergistically potentiated the effects of CNP on cell proliferation and ERK phosphorylation in RMS, indicating that cGMP is a negative regulator of cell proliferation and ERK phosphorylation in this cancer. PDE5 is often overexpressed in various types of cancers including gastric, pancreatic, prostate, and breast cancers [25]. Therefore, if both GC-B and PDE5 are overexpressed, the combination of CNP and sildenafil may be more potent against cancer cells than normal tissue.

In tumor xenograft samples, we observed robust dephosphorylation of ERK and inhibition of tumor growth treated with sildenafil plus CNP, consistent with the results of the in vitro study. However, there is some discrepancy between in vitro and in vivo results. Although phosphorylated ERK levels were reduced by treatment of sildenafil alone in vivo, they were not reduced by sildenafil in vitro. In vivo, sildenafil was administered repeatedly; however, in vitro, RMS cells were treated with single-dose administration. This may have caused the differences between ERK phosphorylation by sildenafil in vitro and in vivo. Additionally, there is discrepancy in the in vivo results. Although Ki-67 labeling index and ERK phosphorylation were suppressed by the treatment of either CNP or sildenafil, the tumor growth was not inhibited in vivo. As shown in in vitro results, CNP significantly inhibited ERK phosphorylation in RMS cells; however, antiproliferative effects of CNP alone were relatively weak. This might be one of the causes of the discrepancy. These findings suggest that strong dephosphorylation of ERK induced by treatment with CNP and sildenafil may be required to inhibit the growth of RMS cells in vivo. Therefore, combined treatment with CNP and sildenafil represents a promising approach for therapy against RMS.

In addition to the anticancer effects of CNP and sildenafil, we evaluated the safety of the combined treatment. Side effects of intense chemotherapy, such as growth disturbance and secondary cancer, are major problems for pediatric patients with RMS; therefore, treatment regimens with a less toxic agent are desirable. CNP, an endogenous peptide, is mainly secreted by vascular endothelial cells in a wide variety of tissues, and it exerts a wide range of vasoprotective effects [14, 33–35]. Moreover, we previously reported that CNP protects against cisplatin-induced nephrotoxicity in mice via its anti-inflammatory effects [36]. Therefore, CNP is considered to be a safe alternative to other chemical compounds. Meanwhile, sildenafil is generally used to treat pulmonary arterial hypertension in children, including neonatal patients, in addition to erectile disorders in adults; in both cases, the drug lacks severe side effects [37, 38]. Our results demonstrate that treatment with CNP and/or sildenafil did not induce weight loss (Fig. 4E). Therefore, the treatment with CNP and sildenafil should be well tolerated by pediatric RMS patients, and could improve their quality of life.

treatment. No weight loss was observed in any of the treated groups relative to the control group (Fig. 4E).

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There are some limitations in this study. First, RMS cells stably expressing GC-B expressed higher levels of GC-B than clinical samples. However, given that the recurrent sample expressed a very high level of GC-B, the findings made using RMS cells stably expressing GC-B are still quite valuable. Although further evidence is necessary to confirm this result, it is possible that all recurrent RMS tumor cells express higher levels of GC-B. Because there is neither a standard treatment protocol nor established evidence for therapeutic strategies against recurrent RMS, CNP/GC-B/cGMP system represents a promising target for growth suppression of RMS cells. Second, treatment with CNP and/or sildenafil did not cause severe adverse effects such as body weight loss in vivo; however, further safety evaluation of this treatment is required. Although subcutaneous infusion of CNP at a rate of 2.5 μg/kg/min does not cause hypotension [14], it should be noted that combined treatment with CNP and sildenafil may induce adverse effects, including hypotension.

In addition to the growth-inhibitory effect of CNP, we are currently investigating whether natriuretic peptides have other anticancer effects, for example, antimetastatic effects. We recently found that ANP suppresses E-selectin expression in vascular endothelial cells, resulting in suppression of cancer cell metastasis [39]; consequently, we have initiated a multicenter randomized clinical trial to examine the use of periperaoperative administration of ANP to prevent cancer recurrence after lung cancer surgery. Although ANP has been well characterized as a cardiac hormone and is utilized in clinical settings, CNP has yet to be applied clinically despite abundant evidence of its role as a regulator of cardiac hypertrophy and remodeling (suggesting its use in treatment of cardiovascular disease) [40] and as a stimulator of endochondral bone growth (implying its potential value in patients with skeletal dysplasia) [41]. Our findings suggest that CNP could be used clinically as an anticancer therapy.

In summary, we demonstrated that CNP in combination with sildenafil decreases ERK phosphorylation and suppresses proliferation of RMS cells. Combined treatment with CNP and sildenafil might be advantageous for RMS patients because of this regimen’s lower toxicity and improved tolerability relative to common chemotherapeutic agents.

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Conflict of Interest

The authors have no conflict of interest.

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Supporting Information
Additional supporting information may be found in the online version of this article:

Figure S1. GC-B mRNA expression decreases with the number of passages.

Figure S2. RD and RMS-YM cells were stably expressing GC-B.

Figure S3. Expression of HES-1, β-catenin, GLI1 and phosphorylation of Akt were not changed by CNP and/or sildenafil treatment in RD-GC-B cells.

Figure S4. Phosphorylated ERK levels were reduced in CNP and/or sildenafil groups, and were further reduced in the sildenafil plus CNP group than in the CNP or sildenafil groups in vivo.