Self-Therapeutic Cobalt Hydroxide Nanosheets ($\text{Co(OH)}_2$ NS) for Ovarian Cancer Therapy

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ABSTRACT: High-grade serous ovarian cancer (HGSOC) is one of the major life-threatening cancers in women, with a survival rate of less than 50%. So far, chemotherapy is the main therapeutic tool to cure this lethal disease; however, in many cases, it fails to cure HGSOC even with severe side effects. Self-therapeutic nanomaterials could be an effective alternative to chemotherapy, facilitated by their diverse physicochemical properties and the ability to generate reactive species for killing cancer cells. Herein, inorganic cobalt hydroxide nanosheets ($\text{Co(OH)}_2$ NS) were synthesized by a simple solution process at room temperature, and morphological, spectroscopic, and crystallographic analyses revealed the formation of $\text{Co(OH)}_2$ NS with good crystallinity and purity. The as-prepared $\text{Co(OH)}_2$ NS showed excellent potency, comparable to the FDA-approved cisplatin drug to kill ovarian cancer cells. Flow cytometric analysis (nexin V) revealed increased cellular apoptosis for $\text{Co(OH)}_2$ NS than cobalt acetate (the precursor). Tracking experiments demonstrated that $\text{Co(OH)}_2$ NS are internalized through the lysosome pathway, although relocalization in the cytoplasm has been observed. Hence, $\text{Co(OH)}_2$ NS could be an effective self-therapeutic drug and open up an area for the optimization of self-therapeutic properties of cobalt nanomaterials for cancer treatment.

INTRODUCTION

High-grade serous ovarian cancer (HGSOC) represents about 70% of all ovarian cancers and is characterized by high lethality.1 The main therapeutic approach of HGSOC includes surgery and chemotherapy.2,3 Chemotherapy based on platinum salts is considered essential for HGSOC treatment4 but has many side effects like other systemic therapies.5–8 and resistance to chemotherapy appears almost in all of the patients with recurrent disease, limiting the effectiveness of treatments.2 Other drugs with clinical benefits approved by FDA are paclitaxel, doxorubicin, capcitabine, topotecan, and PARP inhibitors, and many others are under clinical trials, but still, these treatments are not resolutive.4,9,10

A way to improve efficacy and decrease toxicity is the development of delivery systems using different nanomaterials taking advantage of their different intrinsic properties.10,11 Nanosized materials (e.g., nanoparticles) (NMs) are currently becoming a topic of interest in drug delivery applications for cancer therapy12–14 since they are not expensive, easy to synthesize, and could be functionalized to be selective for and internalized by tumor cells.15,16 Differently from the solubility problems of drugs (many therapeutic drugs fail in clinical trials due to solubility issues),17 nanoparticles could be solubilized in water due to their nanometer size and acquire excellent pharmacokinetic parameters due to their versatility in physicochemical properties.18,19 NMs have a tunable size and could be designed to efficiently take advantage of the enhanced permeability and retention (EPR) effects of some tumors and actively internalize by the membrane of cancer cells or to escape the scavenger macrophages before reaching to the tumor microenvironment.20 Several NMs have been utilized in ovarian cancer therapy like liposomes, micelles, dendrimers, hydrogels, and many other organic and inorganic compounds.21–23 Although a large number of NMs have been developed as delivery systems, so far only up to 1% of the drug reaches the tumoral sites and hence making this approach far from real applications.24,25

These expectations could be fulfilled through nanomaterials using their self-therapeutic (intrinsic therapeutic properties, without loaded drugs) properties.24 Recently, the self-therapeutic trend of nanomaterials has been increasing continuously because of their versatile intrinsic properties, cost reduction, and ease of use in therapeutic applications.24,26 Inorganic materials have elicited considerable attraction in this regard because of their intrinsic antitumor abilities, and many (like SiO2, Ag, Au, MOFs, composites, etc.) have been used even if they are often not effective in killing aggressive tumor cells.24 In humans, cobalt is a trace element necessary for the biological activity of vitamin B12 and at a daily assumption...
above 1 mg could create adverse reactions including hematological and thyroid problems.27 Starting from the 1950s, cobalt has been available on the market to treat children and adult anemia (now discontinued),28 and many research groups have introduced cobalt-based therapeutic systems for different cancer therapies and in vivo imaging purposes.29 Thamilarasan et al. utilized cobalt(III) complexes as a potential anticancer agent, demonstrating the ability of interacting with DNA.30 Klein et al. prepared magnetite and cobalt ferrite nanoparticles with better biocompatibility and water dispersibility as an enhancer of reactive oxygen species (ROS) formation for radiation therapy.31 Bejarbaneh et al. synthesized cobalt hydroxide nano-flakes functionalized with glutamic acid and conjugated with thiosemicarbazide as an anticancer agent against human breast cancer cells.32 In another report, it was described that cobalt complexes were effective in overcoming multidrug resistance and as an alternative to cisplatin in vivo.33 Hernandez et al. and Munteanu et al. separately reviewed the cobalt-based complexes for therapeutic, imaging, and drug delivery applications.29,34 However, no studies have been reported for the treatment of ovarian cancer with inorganic cobalt or its derivatives.

Here, in this study, we propose for the first time two-dimensional (2D) cobalt hydroxide nanosheets (Co(OH)2 NS) for HGSOC treatment. Very simple and cost-effective solutions have been utilized to synthesize Co(OH)2 NS at room temperature (RT). Co(OH)2 NS were investigated through different characterization techniques and tested as an effective drug for ovarian cancer cells. Different experiments were performed to check the toxic effects, localization, and internalization of Co(OH)2 NS.

## RESULTS AND DISCUSSION

### Synthesis of Co(OH)2 NS

A water–benzene mixed solvent system was employed to prepare Co(OH)2 NS, and the synthesis procedure is schematically shown in Figure 1a. Figure 1b shows the possible reaction pathways for the formation of Co(OH)2 NS. Initially, basic 2-ethylimidazole (EIM) and trimethylamine (TEA) could react with water-forming hydroxyl (OH) ions (eqs 1 and 2).36 Subsequently, the OH in the solution react with Co2+ of Co(CH3CO2)2 and form Co(OH)2 (eq 3). Upon stirring, the water immiscibility and low specifi city of benzene can result in an outward pushing force and facilitate the formation of NS.

#### Morphological Characterizations

Figure 2a,b shows the field emission scanning electron microscopy (FE-SEM) images of the as-synthesized Co(OH)2 NS. The low transparency to the electron beam in the high-resolution transmission electron microscope (HR-TEM) image (Figure 2c) further suggests the formation of interconnected 2D NS, which is consistent with the FE-SEM results. The bright spots with a ringlike pattern having d-spacings of ca. 2.4, 1.45, 1.26, and 0.99 Å with the corresponding hkl reflections of (001), (100), (011), and (102) obtained from the selected area diffraction pattern (SAED) (Figure 2d) suggested the good crystallinity and purity of Co(OH)2.37 All of the structural analyses confirmed the high crystallinity and purity of the as-synthesized Co(OH)2 NS.
Structural Characterizations. Figure 3a shows the XRD powder pattern of the as-prepared Co(OH)$_2$ NS along with the simulated XRD pattern of Co(OH)$_2$. The simulated pattern was perfectly matched with the XRD pattern of Co(OH)$_2$ NS without the presence of impurity peaks. The intense and sharp XRD peaks indicated the high crystallinity of Co(OH)$_2$ NS, consistent with the TEM analyses. The major peaks of Co(OH)$_2$NS at 2$\theta$ angles of ca. 19.30, 32.60, 38.10, 51.35, 57.98, and 61.75° could be ascribed to the $hkl$ reflections of (001), (100), (011), (102), and (110), respectively. A Fourier transform infrared spectroscopy (FTIR) study was performed to investigate the chemical bonding in Co(OH)$_2$ NS (Figure 3b), which showed a strong absorption peak of O–H stretching at ca. 3425 cm$^{-1}$, arising from the Co–OH bonding.
groups and the adsorbed water molecules. The absorption bands that appeared at ca. 1115 and 1575 cm\(^{-1}\) could be ascribed to C=O stretching and C–H bending, respectively. These bands could be originated from the adsorbed or intercalated CH\(_3\)COO\(^{-}\) ions into the Co(OH)\(_2\) NS. The low-intense FTIR absorption band observed at ca. 640 cm\(^{-1}\) could be assigned to both Co–OH and Co–O bending vibrations, as similarly observed for Co(OH)\(_2\), Ni(OH)\(_2\), and Ni(OH)\(_2\)-Co(OH)\(_2\) layered double hydroxide. The stability of Co(OH)\(_2\) NS was examined by creating artificial in vivo body-like conditions in two different pHs (7.4 and 5.5) at 37 °C. The samples were collected at different time intervals after the treatment and analyzed by UV–vis absorption measurements (Figure S1). Results demonstrated the appearance of Co(OH)\(_2\) absorption band at about 230 nm without any noticeable shifting in the absorbance maxima at both pHs for up to 1 week, suggesting the high stability of Co(OH)\(_2\) NS as a self-therapeutic nanomedicine.

**Cell Viability**. Cell viability measurements were performed on different cell lines (ovarian and colon cancer cell lines and normal cells). The results are summarized in Table 1. For the IC\(_{50}\) values of Co(OH)\(_2\) NS are in the range of 1.6–11.4 µg/mL, compared to those of the cobalt acetate precursor (5.5–41.6 µg/mL). These results suggest that inside the cells, Co(OH)\(_2\) NS could release Co\(^{2+}\) ions and exert its toxic activity. Compared with cisplatin, in cancer cells, Co(OH)\(_2\) NS is less potent in the range of 2- to 32-fold. Conversely, in human MRC-5 fibroblast cells, the toxicity of Co(OH)\(_2\) NS is 80-fold less. The calculate ratio of potency/toxicity of Co(OH)\(_2\) NS vs cisplatin is 2.5- to 40-fold higher. In mouse NIH-3T3 fibroblast cell line, the Co(OH)\(_2\) NS is still less toxic than cisplatin (5-fold less). The results suggest that Co(OH)\(_2\) NS could be a possible future candidate as an alternative drug effective for late-stage tumors. The versatility of the proposed Co(OH)\(_2\) NS for other cancers was tested on the HCT-116 colon cancer cell line and showed promising potency and comparable results with FDA-approved compounds.

**Evaluation of Cell Death.** Apoptosis (Annexin V) was evaluated using flow cytometric analysis with Co(OH)\(_2\) NS (50 µg/mL), cobalt acetate (50 µg/mL), and cisplatin (6 µg/mL; 20 µM) as the reference drug at different time points. All of the results are summarized in Figure 4. No apoptosis has been seen in the untreated set of experiments, while cisplatin, cobalt acetate, and Co(OH)\(_2\) NS promptly induced apoptosis after 6 h. Both early apoptosis and late apoptosis were observed starting from 6 h and increased with time. The apoptosis is much higher in Co(OH)\(_2\) NS-treated cells compared to cobalt acetate-treated cells at any time point. These results confirmed that Co(OH)\(_2\) NS could be an alternative candidate that could cause earlier apoptosis and cell death in aggressive tumors.

**Intracellular Localization.** Lysosomes have long been known to play a key role in the degradation of extracellular materials, including chemotherapeutics agents. The colocalization experiments were performed to evaluate the amount of Co(OH)\(_2\) NS trafficking to lysosomes. The experiments were performed in the A2780 ovarian cancer cell line (Figure 5). The nucleus was labeled with Hoechst 33342 (blue), lysosome with LysoTracker Green DND-26 (green), and Co(OH)\(_2\) NS was labeled with rhodamine B (red). Co(OH)\(_2\) NS were analyzed with a fluorescence microscope after 1, 6, and 24 h. It was observed that Co(OH)\(_2\) NS were uptaken at each time point. It was demonstrated that Co(OH)\(_2\) NS started internalizing even after 1 h. The colocalization of Co(OH)\(_2\) NS and lysosomes was evaluated with Pearson’s correlation coefficient (R) after 24 h. The “R” value was found to be 0.43 for Co(OH)\(_2\) NS, meaning that Co(OH)\(_2\) NS partially localized to the lysosomes and could relocalize to the cytoplasm avoiding degradation.

**Co(OH)\(_2\) NS Are Effective in Killing Cancer Organoids from High-Grade Ovarian Cancer Patients.** More than 30% of patients at the late stage of ovarian cancer will develop ascites. Ascites are treated indirectly with chemotherapy, and paracentesis is used to alleviate the symptoms. Ascites contain free-floating cells that are responsible for intra-peritoneal metastasis.

To test the effectiveness of Co(OH)\(_2\) NS, cancer organoids were generated. Cancer organoids are the last frontiers for ex vivo testing of drugs that replicate the response of patients in clinic. Two HGSOC-derived organoids from ascites (Pat A, B) and one high-grade endometroid ovarian cancer from a primary tumor (Pat C) were treated with cobalt acetate and Co(OH)\(_2\) NS. Immunohistochemistry (IHC) analysis showed the expression of typical markers of HGSOC, CA125 (cancer antigen 125), WT-1 (Wilms tumor 1), and Pax8 (Paired box 8) (Figure S2). Co(OH)\(_2\) NS are effective in the range of 7–26 µg/mL. The potency of Co(OH)\(_2\) NS is similar for ascites and primary tumors, suggesting a possible application of them in chemotherapy-naïve and heavily treated patients who are resistant to most of the drugs (Table 2).

Compared to cobalt acetate, Co(OH)\(_2\) NS are more effective (p value < 0.05) (Figure S3). The median values for Co(OH)\(_2\) NS and cobalt acetate are 11 and 29, respectively.

**DISCUSSION**

Cobalt is available to humans through vitamin B12 or cyanocobalamin, an essential vitamin for many physiological processes that include DNA and aminoacidic synthesis among others. The active forms of vitamin B12, methylcobalamin and adenosylcobalamin, are the cofactors of essential enzymes methionine synthase (MS) and methylmalonyl-CoA mutase (MCM), respectively. Deficiency of vitamin B12 is responsible for or associated with many diseases, including megaloblastic anemia, methylmalonic aciduria, hyperhomocys-

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**Table 1. IC\(_{50}\) Values of Co(OH)\(_2\) NS, Cobalt Acetate, and Cisplatin**

| Drug          | AV (µg/mL) | SD  (µg/mL) | ratio |
|---------------|------------|-------------|-------|
| A2780 cisplatin | 0.05       | 0.01        | 1     |
| Co(OH)\(_2\) NS | 1.6        | 0.2         | 32    |
| cobalt acetate | 5.5        | 2.6         | 110   |
| OVCAR-3 cisplatin | 0.6       | 0.1         | 1     |
| Co(OH)\(_2\) NS | 11.4       | 2.2         | 19    |
| cobalt acetate | 12.1       | 1.7         | 20    |
| HCT-116 cisplatin | 3.1       | 0.3         | 1     |
| Co(OH)\(_2\) NS | 6.7        | 0.4         | 2     |
| cobalt acetate | 41.6       | 1.6         | 13    |
| NIH-3T3 cisplatin | 13.1       | 5.5         | 1     |
| Co(OH)\(_2\) NS | 69         | 13          | 5     |
| cobalt acetate | 82.4       | 3.8         | 6     |
| MRC-5 cisplatin | 0.4        | 0.1         | 1     |
| Co(OH)\(_2\) NS | 31.9       | 3.6         | 80    |
| cobalt acetate | 29.8       | 11.4        | 75    |

“Values are expressed as µg/mL.”
teinemia, and cobalamin neuropathy. Treatment of anemia with a range of 25–150 mg of CoCl₂/day has been utilized for many years, but adverse effects drive clinicians to alternative therapies. Excess of cobalt in the blood has been linked to hematological (range: 0.2–1 mg Co/kg per day), thyroid (range: 0.5–2.7 mg Co/kg per day), neurological (0.3 mg Co/kg per day), and cardiac (range: 0.04–0.07 mg Co/kg per day) effects. As observed, free cobalt ions are responsible for the manifested toxicity. Cobalt ions could generate ROS (Fenton-like reaction) and interfere with the activity of proteins and ions necessary for cellular and organ physiology, such as calcium, iron, iodine, mitochondrial functions, and erythropoiesis.

Starting from these concepts, cobalt complexes that are sufficiently stable during blood circulation will avoid systemic toxicity, in turn releasing cobalt ions after the accumulation in tumors due to the EPR effects and acidic pH typical of the tumor microenvironment. Inside the tumors, the release of cobalt ions could exert their toxic activity. In this research, we developed highly stable cobalt hydroxide nanosheets by a novel solution synthetic process root with an optimal potency/toxicity ratio. The as-synthesized nanosheets were characterized using different techniques to confirm the morphology and crystal structures and showed a sheetlike structure that was confirmed by SEM and TEM analyses. Furthermore, the purity of the cobalt hydroxide nanosheets was examined for their crystal structure using X-ray and FTIR spectroscopies. The X-ray results having distinct crystal planes of cobalt hydroxide agree with the SAED of TEM results with concentric bright rings. The as-synthesized nanosheets were used on different cancer cell lines. The results showed that the cytotoxicity of the cobalt acetate (the precursor) and Co(OH)₂ NS showed toxic activity in the same range for cancer cells. Compared to cisplatin, Co(OH)₂ NS is less toxic with a favorable potency/toxicity ratio. Flow cytometric analysis (Annexin V) was performed to check cellular apoptosis, and results showed an

Table 1

| Time (hours) | Sample name                  | Late Apoptosis % (Q2) | Early Apoptosis % (Q4) |
|--------------|------------------------------|-----------------------|------------------------|
| 6            | Not treated                  | 0.05 ± 0.02           | 0.08 ± 0.04            |
|              | Cisplatin 20 μΜ              | 0.05 ± 0.01           | 0.10 ± 0.02            |
|              | Cobalt acetate 50 μg/mL      | 0.13 ± 0.02           | 0.18 ± 0.01            |
|              | Co(OH)₂ NP 50 μg/mL          | 0.46 ± 0.06           | 0.23 ± 0.02            |
| 24           | Not treated                  | 0.14 ± 0.04           | 0.14 ± 0.03            |
|              | Cisplatin 20 μΜ              | 8.66 ± 0.05           | 2.40 ± 0.04            |
|              | Cobalt acetate 50 μg/mL      | 1.18 ± 0.04           | 0.58 ± 0.04            |
|              | Co(OH)₂ NP 50 μg/mL          | 8.97 ± 0.33           | 0.79 ± 0.01            |
| 48           | Not treated                  | 0.08 ± 0.02           | 0.17 ± 0.02            |
|              | Cisplatin 20 μΜ              | 18.08 ± 1.24          | 5.29 ± 0.20            |
|              | Cobalt acetate 50 μg/mL      | 3.36 ± 0.04           | 0.89 ± 0.04            |
|              | Co(OH)₂ NP 50 μg/mL          | 9.26 ± 1.23           | 1.13 ± 0.04            |

Figure 4. Top: An example of apoptosis analysis of cobalt acetate, Co(OH)₂ NS, and cisplatin at different time points. Q1, dead cells; Q2, late apoptosis; Q3, healthy cells; Q4, early apoptosis. Bottom: Values (μg/mL) of early apoptosis and late apoptosis from three experiments.
increase in apoptosis that was much more evident for Co(OH)2 NS than cobalt acetate. Colocalization and cellular internalization of Co(OH)2 NS were performed using a LysoTracker marker. The results indicated an active internalization of Co(OH)2 NS that can escape the lysosomal degradation pathway.

All of these results showed that Co(OH)2 NS are promising to inhibit the growth of aggressive cancer cells and could be a potential candidate to cure many other late-stage tumors. However, further in vivo studies are required to evaluate the pharmacokinetic parameters and the toxic effects on healthy organs.

**CONCLUSIONS**

In summary, Co(OH)2 NS synthesized by a simple solution process are used as a self-therapeutic drug for treating HGSOC. The results revealed that the cytotoxicity of the cobalt acetate (the precursor) and the as-synthesized high-purity and crystalline Co(OH)2 NS showed toxic activity in the same range for cancer cells. Compared to cisplatin, Co(OH)2 NS are less toxic with a favorable potency/toxicity ratio. Flow cytometric analysis (Annexin V) showed an increase in cellular apoptosis that was much more evident for Co(OH)2 NS than cobalt acetate. Furthermore, colocalization and cellular internalization experiments of Co(OH)2 NS using a LysoTracker marker indicated an active internalization of Co(OH)2 NS that can escape the lysosomal degradation pathway.

Although a high concentration of cobalt is toxic for humans, nanoformulations have been demonstrated to be potentially safe at least at the preclinical level. Thus, this study opens up a window to develop cobalt-based nanomaterials as a self-therapeutic nanomedicine for cancer therapy.

**MATERIALS AND METHODS**

**Materials and Reagents.** Cobalt(II) acetate (Co-(CH3CO2)2), (99.99%), trimethylamine (TEA) (99%), 2-ethylimidazole (EIM) (98%), benzene (99.8%), and absolute ethyl alcohol were purchased from Sigma-Aldrich (St. Louis, MO). A Millipore Milli-Q Biocell A10 water purifying system was used to prepare ultrapure water. A2780 (Sigma-Aldrich, St. Louis, MO), OVCAR-3, HCT-116, and MRC-5 (ATCC, Manassas, VA) cell lines were grown accordingly to the manufacturers’ instructions. LysoTracker Green DND-26, Hoechst 33342, and rhodamine B were purchased from Thermo Fisher Scientific (Waltham, Massachusetts).

**Synthesis of Co(OH)2 NS.** A simple solution process was utilized to prepare pure Co(OH)2 NS. In brief, a solution of Co(CH3CO2)2 (0.3 g) was prepared in a beaker using ultrapure water (50 mL). In another beaker, a mixed solution of EIM (0.40 g) and TEA (0.40 g) was prepared using 50 mL of a mixed solvent of water and benzene (90:10, v/v). Then, both solutions were mixed and stirred vigorously at room temperature (RT) for 1 h. After completing the reaction, the precipitate was collected by the centrifuge method at 6000 RPM, washed with water and ethanol, and dried in a vacuum oven at 100 °C for 2 h. The as-synthesized Co(OH)2 NS were stored in an airtight vial for further characterization and application purposes.

**Apparatus and Measurements.** A field emission scanning electron microscope (FE-SEM, Carl Zeiss Sigma VP, Germany) and a high-resolution transmission electron microscope (HR-TEM, JEM-2100 (HRP)) were utilized to analyze the morphology of the synthesized Co(OH)2 NS. HR-TEM was used to characterize the selected area diffraction pattern (SAED) and the lattice fringe of the Co(OH)2 NS. An

Table 2. IC50 Values (μg/mL) of Cobalt Acetate and Co(OH)2 NS in Cancer Organoids Derived from Two HGSOC Ascites Patients (Pat A, B) and One High-Grade Endometroid Tumor Patient (Pat C)*

| patients derived from | cobalt acetate | Co(OH)2 NS | doxorubicin |
|----------------------|----------------|------------|-------------|
| A ascites            | 28 ± 4        | 7 ± 3      | 1.8 ± 0.4   |
| B ascites            | 41 ± 13       | 26 ± 27    | 0.09 ± 0.03 |
| C primary tumor      | 22 ± 12       | 16 ± 5     | 0.9 ± 0.3   |

*Data were obtained from five replicates. The numbers represent mean and standard deviation.
X-ray diffractometer (XRD, Philips) with a Cu Kα radiation (λ = 1.5406 Å) and a Fourier transform infrared (FTIR) spectrophotometer (MIDAC, M4000) were used to characterize the structural properties of Co(OH)2 NS. To calculate IC50 values, luminescence was read with a Tecxan infinite M1000 Pro instrument (Tecan, Männedorf, Switzerland). Annexin V was evaluated with a BD FACS CantoII instrument (BD Biosciences, San Jose, CA). Fluorescence images were capture with a Nikon Ti Eclipse inverted microscope (Nikon, Minato City, Tokyo, Japan).

**Cell Viability.** Co(OH)2 NS were tested on different cell lines that were seeded in 96-multwell plates at a density of 1 × 10^3 (cancer cells) or 8 × 10^3 (MRC-5 cells). Cell viability was measured using the CellTiter-Glo assay system according to the manufacturer’s instructions (Promega, Madison, Wisconsin) after 96 h. Six serial dilutions 1:10 of the compounds were utilized to calculate the IC50 values with GraphPad software utilizing the nonlinear regression method.

Organoids were plated as single cells as possible (around 1000 cells) in five replicates and treated with six serial dilutions 1:10 of the compounds and analyzed after 96 h as cell lines.

**Stability of Co(OH)2 NS.** An artificial in vivo system has been developed by mixing Co(OH)2 NS (3 mg/mL) in physiological solutions having two different pH values (pH 5.5 and pH 7.4). Then, the solutions were placed in an incubator at 37 °C. The samples were collected at different-day intervals (0–7; DO, D1, D2, D3, D5, and D7), centrifuged, and collected. Then, the UV–vis absorption spectra of the samples were measured by dispersing them in DI water.

**Intracellular Localization.** The internalization and localization of Co(OH)2 NS into the cells were evaluated by fluorescence probes. Typically, 150 000 cells were seeded into a transparent microplate containing a glass coverslip. To label Co(OH)2 NS under continuous rotation at room temperature with a Nikon Ti Eclipse inverted microscope (Nikon, Minato City, Tokyo, Japan).

**Histopathology Analysis.** Sections of formalin-fixed, paraffin-embedded ascites and solid tumor organoids were used for histopathological analyses. Organoids were collected, fixed in phosphate-buffered 10% formalin, and embedded in paraffin using Micro NextGen CellBloxing Kit (Cat no: M20; AV BioInnovation) following manufacturer’s instructions. Subsequently, 5 µm sections were stained with hematoxylin and eosin (H&E) and with the Leica ST5020 multitaster and 2 µm sections were cut for IHC analysis. Histopathology staining was performed with UltraVision LP Detection System HRP DAB kit (Thermo Scientific, Waltham). Heat-induced antigen retrieval was performed using 10 mM citrate buffer pH 6.0. The following antibodies were used to characterize patient’s derived organoids and parent tumor: PAX8 (ProteinTech Group, Germany, EU; 10336-1-AP); Ca125 (Santacruz Biotechnology, TX; sc-52095); WT1 (Abcam, U.K.; ab89901). Tissues were analyzed with a light microscope using different magnifications.

**Statistical Analysis.** Statistical analysis was performed using GraphPad Prism software, and a p value <0.05 was considered significant.
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Notes
The authors declare no competing financial interest.

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