Identification of an iridium(III) complex with anti-bacterial and anti-cancer activity

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Group 9 transition metal complexes have been widely explored as therapeutic agents due to their unique geometry, their propensity to undergo ligand exchanges with biomolecules and their diverse steric and electronic properties. These metal complexes can offer distinct modes of action in living organisms compared to carbon-based molecules. In this study, we investigated the antimicrobial and anti-proliferative abilities of a series of cyclometallated iridium(III) complexes. The iridium(III) complex 1 inhibited the growth of *S. aureus* with MIC and MBC values of 3.60 and 7.19 μM, respectively, indicating its potent bactericidal activity. Moreover, complex 1 also exhibited cytotoxicity against a number of cancer cell lines, with particular potency against ovarian, cervical and melanoma cells. This cyclometallated iridium(III) complex is the first example of a substitutionally-inert, Group 9 organometallic compound utilized as a direct and selective inhibitor of *S. aureus*.

The design and synthesis of metal complexes for imaging and therapy have attracted considerable attention from inorganic medicinal chemists over the past decade14,15. Increasingly, transition metal complexes have been explored as therapeutic agents for treating human diseases due their prominent advantages. For example, the steric and electronic properties of metal complexes can be fine-tuned by modification of their auxiliary ligands. Moreover, metal complexes possess unique geometries for the attachment of ligands, and they can also undergo ligand exchange reactions with specific biomolecules5. Consequently, metal complexes can offer distinct modes of action compared to carbon-based molecules6–18.

With the global rise of antibiotic resistance and the lack of new antibiotics reaching the market, new antimicrobial drugs with novel mechanisms of action are desperately needed17. Complexes of copper(II)18, silver(I)19–21, ruthenium(II)22, bismuth(III)23 vanadium(IV)24, and iron(II)25, among other complexes26–36, have been reported to disrupt antibiotic-resistant biofilms, exert synergistic bactericidal activity with other biocides, inhibit metabolic pathways in a selective manner or to kill multidrug-resistant bacteria.

A number of mechanisms of metal-induced microbial activity have been extensively studied37. The first is the production of reactive oxygen species (ROS) and antioxidant depletion. Treatment of *E. coli* with exogenous hydrogen peroxide (H2O2) or agents that catalyse the generation of superoxide (O2•−) resulted in DNA damage and the inhibition of enzyme activities that are important for bacterial growth37. Moreover, bacterial and *S. cerevisiae* strains lacking ROS-scavenging enzymes or other cellular antioxidants showed altered sensitivity towards redox-active metals38. Alternatively, certain metals might disrupt cellular donor ligands that coordinate Fe, leading to the release of additional Fenton-active Fe into the bacterial cytoplasm resulting in increased ROS production. Oxidative stress in bacteria can also lead

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to the oxidation of cellular thiols, leading to the formation of protein disulphides and the depletion of
antioxidant reserves such as glutathione in bacterial cells.

Another interesting strategy is the use of “Trojan horse” compounds that trick bacteria into taking up
toxic metals. These ions can be delivered to bacterial cells by conjugation to a siderophore such as
desferrioxamine (DFO), enterochelin or protoporphyrin IX (PPIX). Notably, gallium nitrate (Ga(NO₃)₃), Ga-DFO and Ga-PPIX show broad-spectrum bactericidal activity against Gram-negative and
Gram-positive bacteria in vitro and in vivo of infection.

Only limited examples of kinetically-inert Group 9 metal complexes have been reported as antibacterial and anticancer agents. Recently, three cyclometalated iridium(III) complexes bearing dithiocarbamate derivatives have been reported to interact with calf thymus DNA and possess anti-bacterial activity. The delocalization of the π-electrons over the chelate ring was proposed to increase the lipophilicity of the complexes and facilitate their penetration into bacterial cell membranes, thereby resulting in inhibition of bacterial growth. Highly hydrophobic Ir(III) complexes containing both Cp* and a C,N-chelating ligand have been validated for their significant cytotoxic activity towards A2780 human
ovarian cancer cells. Additionally, an NHC iridium(I) complex with significant antiproliferative properties in vitro has been shown to target cytochrome c with simultaneous oxidation of Ir(I) to Ir(III). In this report, we describe the synthesis and biological evaluation of a substitutionally-inert, organometallic iridium(III) complex that inhibits the growth of Staphylococcus aureus.

Results and Discussion

Synthesis. The iridium(III) complexes 1–4 and rhodium(III) complex 5 are kinetically-inert, organo-
metallic complexes with a general structure [M(C^N)₂(N^N)]⁺ (M = Ir or Rh). Complexes 1 and 2
contain a 5-amino-1,10-phenanthroline N^N ligand, while complexes 3–5 contain the bathophenanth-
roline N^N ligand. In regards to the C^N ligand, complexes 3 and 5 possess 2-phenylpyridine ligands,
while complex 1 bears the related 2-(p-tolyl)pyridine ligand. Complexes 2 and 4 carry the bulkier 2-phe
nylquinoline and 1-isouquinoline C^N ligands, respectively.

The syntheses and characterization of the iridium(III) or rhodium(III) complexes 1–5 are described
in the Methods section. The spectroscopic data of the complexes are presented in Table 1. Complexes 2
and 5 are novel whereas complexes 1, 3 and 4 have been previously described. All complexes were
synthesized and tested as racemic mixtures of enantiomers.

Antibacterial activity. The anti-bacterial activity of the complexes 1–5 was investigated using a
disk diffusion assay. In this assay, four different bacterial strains (Staphylococcus aureus, Escherichia coli,
Enterococcus faecalis and Klebsiella pneumoniae) were plated on an agar dish, and disks soaked with
solutions of the metal complexes (11.5μM) or ampicillin were placed onto the dish. The antimicrobial
activity of the metal complexes would be manifested as the absence of bacterial growth around the disk, due to diffusion of the compounds into the surrounding agar. After 24 h, the growth inhibition zones of the test strains were determined. Encouragingly, the results showed that racemic complex 1 (rac-1) exhibited selective anti-bacterial activity towards S. aureus, with an average inhibition zone of 15 mm (Fig. 2). On the other hand, the other metal complexes showed limited antibacterial activity.

**Determination of MIC and MBC values.** We next determined the minimum inhibitory concentration (MIC) value of complex 1 against S. aureus by the broth dilution method. Briefly, S. aureus inoculum in Luria Bertani broth was incubated with serial dilutions of complex 1 at 37°C for 24 h. The MIC was determined as the lowest concentration of complex 1 inhibiting visible bacterial growth. The results showed that the MIC of S. aureus grown with complex 1 was 3.60 μM (Fig. 3).

To determine the MBC (minimum bactericidal concentration), the agar dilution method was used. Briefly, S. aureus inoculum treated with different concentrations of complex 1 were taken aseptically from tubes that had not presented visible turbidity and plated onto agar. The MBC was considered as the lowest concentration of complex 1 that allowed less than 0.1% of the original inoculum treated with complex 1 to grow on the surface of the agar. The results showed that the MBC of S. aureus grown with complex 1 was 7.19 μM. To determine the nature of antibacterial effect of a compound, the MBC/MIC ratio is typically used. When this ratio is lower than 4, the compound is considered as a bactericidal. On the other hand, if this ratio is higher than 4, the compound is considered to be bacteriostatic. As the MBC/MIC ratio of complex 1 is 2, complex 1 is considered to be bactericidal.

From these data, a brief structure-activity relationship analysis can be conducted. Complex 1 contains the 2-(p-tolyl)pyridine ligand C^N, which is relatively smaller compared to the quinoline or isoquinoline-based C^N ligands of complexes 2 and 4, respectively. This suggests that smaller C^N ligands may be beneficial for antimicrobial activity. Additionally, complex 1 also contains an amino substituent on the phenanthroline N^N ligand. The amino group may confer the necessary hydrophilic/hydrophobic balance for the complex to penetrate the thick peptidoglycan cell wall of Gram-positive bacteria such as S. aureus. While complex 2 contains the same 5-amino-1,10-phenanthroline N^N substituent as complex 1, its larger 2-phenylquinoline C^N ligand may have diminished its activity.

Interestingly, a slightly increased zone area was observed when the iridium(III) center of complex 1 was replaced with rhodium(III) (as in congener 5), suggesting the importance of the metal center in contributing to antibacterial activity. However, complexes 3–5, containing the relatively large bathophenanthroline N^N ligand, generally showed modest zone areas and were less active compared to complex 1. In particular, complexes 2–5 showed only very limited activity against S. aureus. Additionally, complexes 2–5 at a concentration of 50–60 μM were not able to inhibit bacterial growth in LB broth. These complexes might exhibit antibacterial activity at higher concentrations, but this would not be practical for clinical use. Taken together, these findings suggest that the presence of the relatively bulky N^N ligand detracts from antibacterial activity. We propose that the lipophilicity and/or structure of complex 1 might allow it to effectively pass through the hydrophobic interior of the bacterial cell membrane.

**Partition Coefficients (log P).** The octanol-water partition coefficients (log P) for complex 1 was determined. Complex 1 showed a log P value of 0.629, indicating that it is slightly lipophilic and also satisfies Lipinski's lipophilicity criterion (log P < 5) for drug likeness.

**Evaluation of cell proliferation.** Lipophilicity correlates with cytotoxic potency. Therefore, the cytotoxicity of complex 1 towards different cell lines was evaluated. By using the 3-(4,5-dimethylthiazol-2-yl)-2,5-diphenyltetrazolium bromide (MTT) assay, the in vitro cytotoxicity of complex 1 was evaluated in a panel of human cell lines, including ovarian cancer A2780 and SKOV3 cells, hepatocellular carcinoma HepG2 cells, cervical adenocarcinoma HeLa cells, and malignant melanoma A375 and A2058 cells. Cells were exposed to complex 1 in concentrations ranging from 0.01 to 10 μM and cellular proliferation was assessed after 72 h. The results showed that complex 1 inhibited cellular proliferation of A2780, HepG2, SKOV3, HeLa, A375, and A2058 cells in a dose-dependent manner with IC_{50} values of 7.23 ± 1.30, 17.0 ± 1.39, 1.77 ± 1.70, 1.68 ± 1.29, 1.55 ± 0.15 and 1.24 ± 0.31 μM, respectively (Fig. 4). The results

| Complex | λ_{max}/nm | UV/vis absorption λ_{max}/nm (ε/dm^3mol^-1cm^-1) |
|---------|------------|-----------------------------------------------|
| 1       | 555        | 270 (2.15 × 10^3), 381 (2.47 × 10^2)[36]    |
| 2       | 590        | 274 (7.98 × 10^3), 354 (1.13 × 10^4)        |
| 3       | 587        | 270 (6.1 × 10^4), 337 (1.99 × 10^4)[37]    |
| 4       | 593        | 279 (2.0 × 10^5), 382 (1.56 × 10^4), 439 (6.10^3)[38] |
| 5       | No emission| 284 (1.21 × 10^5), 366 (2.26 × 10^4)        |

Table 1. Photophysical properties of complexes 1–5.
indicate that the cytotoxic effect of complex 1 is cell type-dependent. Complex 1 selectively inhibits growth of ovarian, cervical and melanoma cells, but not hepatocarcinoma cells.

**Stability of complex 1.** The most potent complex 1 was stable in H$_2$O/acetonitrile (400:1) solution at 298 K for at least 24 h as revealed by UV-Vis spectroscopy. No significant change in absorption spectrum of complex 1 was observed over 24 h (Fig. 5). The result indicates that complex 1 is stable at room temperature.

Figure 2. Anti-bacterial activity of complexes 1–5 as determined by the disk diffusion assay. (a) The anti-\textit{S. aureus} activity of complexes 1–5. (b) The anti-\textit{E. coli} activity of complexes 1–5. (c) The anti-\textit{E. faecalis} activity of complexes 1–5. (d) The anti-\textit{K. pneumoniae} activity of complexes 1–5. All complexes used are racemic mixture of enantiomers.
Methods

Materials and cell lines. All chemicals, unless specified, were obtained from Sigma-Aldrich. *Staphylococcus aureus, Enterococcus faecalis, Escherichia coli, Klebsiella pneumonia, HepG2, SKOV3, HeLa, A375* and A2058 were purchased from American Type Culture Collection (ATCC). A2780, HepG2, SKOV3, HeLa, A375 and A2058 cells were routinely grown in suspension in 90% RPMI-1640 containing L-glutamine (2 mM) or DMEM, antibiotics (100 IU penicillin/mL, 100 μg streptomycin/mL).

Figure 3. Inhibition of the growth of *S. aureus* by racemic mixture of complex 1 as determined by the broth dilution method. Results are representative of three independent experiments, and the complete data are shown in Table S2.

Figure 4. Cell viability of racemic mixture of complex 1 against A2780, HepG2, SKOV3, HeLa, A375 and A2058 cells as determined by the MTT assay. Error bars represent the standard deviations of the results from three independent experiments.

Figure 5. UV/Vis absorption of 1 (2.5 μM) in H₂O/acetonitrile (400:1) at t = 0 h and after incubation for 0, 1, 2, 4, 12, and 24 h at 298 K.
and supplemented with 10% (v/v) fetal bovine serum (FBS). The cells were maintained at a cell density of 0.2–1 × 10^5 cells/mL in a 5% CO2 humidified atmosphere at 37°C.

**General experimental.** Mass spectrometry was performed at the Mass Spectroscopy Unit at the Department of Chemistry, Hong Kong Baptist University, Hong Kong (China). Melting points were determined using a Gallenkamp melting apparatus and are uncorrected. Deuterated solvents for NMR purposes were obtained from Armar and used as received.

1H and 13C NMR were recorded on a Bruker Avance 400 spectrometer operating at 400 MHz (1H) and 100 MHz (13C). 1H and 13C chemical shifts were referenced internally to solvent shift (acetone-d_6: 1H δ 2.05, 13C δ 29.8, 206.1). Chemical shifts (δ) are quoted in ppm, the downfield direction being defined as positive. Uncertainties in chemical shifts are typically ±0.01 ppm for 1H and ±0.05 for 13C. Coupling constants are typically ±0.1 Hz for 1H-1H and ±0.5 Hz for 1H-13C couplings. The following abbreviations are used for convenience in reporting the multiplicity of NMR resonances: s, singlet; d, doublet; t, triplet; m, multiplet. All NMR data was acquired and processed using standard Bruker software (Topspin).

**Photophysical measurement.** Emission spectra and lifetime measurements for complex 1 were performed according to the previously described method^51.

**Synthesis.** These complexes were synthesized using a modified literature method^55. MCl_3·3H_2O was heated to 150°C with 2.2 equivalents of C^N ligand in 3:1 methoxymethanol and deionized water under a nitrogen atmosphere for 12 h. The product was filtered off and washed with ether (3 × 50 mL) and then with deionized water (3 × 50 mL). A suspension of [M_2(C^N)Cl_2] (0.2 mmol) and the N^N ligand (0.44 mmol) in a mixture of dichloromethane:methanol (1:1, 20 mL) was refluxed overnight under a nitrogen atmosphere. The resulting solution was then allowed to cool to room temperature, and filtered to remove unreacted cyclometallated dimer. To the filtrate, a solution of ammonium hexafluorophosphate (0.44 mmol) in methanol (5 mL) was added and the filtrate was reduced in volume by rotary evaporation until precipitation of the crude product occurred. The precipitate was then filtered and washed with several portions of water (2 × 50 mL) followed by diethyl ether (2 × 50 mL). The product was recrystallized by acetonitrile:dichethyl ether vapor diffusion to yield the titled compound. All complexes were characterized by 1H-NMR, 13C-NMR, high resolution mass spectrometry (HRMS) and elemental analysis.

**Complex 1.** Reported^51.

**Complex 2.** Yield: 61%. 1H NMR (400 MHz, Acetone-d_6) δ 9.22 (d, J = 8.4 Hz, 1H), 9.12 (s, 1H), 9.06 (d, J = 8.0 Hz, 1H), 8.98 (d, J = 4.0 Hz, 1H), 8.94 (d, J = 4.0 Hz, 1H), 8.59 (d, J = 8.8 Hz, 2H), 8.52 (d, J = 8.8 Hz, 2H), 8.34 (d, J = 8.0 Hz, 2H), 8.31-8.25 (m, 2H), 7.84 (d, J = 8.0 Hz, 2H), 7.34-7.25 (m, 6H), 6.93 (dd, J = 6.0 Hz, 1.2 Hz, 2H), 6.70 (d, J = 8.0 Hz, 2H); 13C NMR (100 MHz, Acetone-d_6) δ 171.4, 152.6, 152.1, 149.4, 148.6, 147.2, 147.1, 145.3, 144.5, 141.9, 141.1, 141.0, 136.1, 135.6, 135.5, 134.1, 133.8, 131.5, 131.4, 131.3, 130.1, 130.0, 128.6, 128.3, 128.2, 127.6, 127.5, 127.3, 126.4, 125.3, 125.2, 124.7, 123.8, 123.7, 119.0, 118.9, 103.7. HRMS: Calcd. for C_{46}H_{32}N_4RhPF_6·2H_2O: 979.2091. Found: 979.2091. Anal. C, H, N: calcd. 51.64, 3.40, 7.17, found: 51.65, 3.15, 7.41. The elemental analysis result (Figure S1) was shown in the supporting information.

**Complex 3.** Reported^52.

**Complex 4.** Reported^53.

**Complex 5.** Yield: 58%. 1H NMR (400 MHz, Acetone-d_6) δ 8.58 (d, J = 5.2 Hz, 2H), 8.33-8.30 (m, 4H), 8.06-8.02 (m, 6H), 7.80 (d, J = 4.8 Hz, 2H), 7.70-7.66 (m, 10H), 7.19 (t, J = 3.6 Hz, 2H), 7.09 (dd, J = 6.0 Hz, 1.2 Hz, 4H), 6.52 (d, J = 7.6 Hz, 2H); 13C NMR (100 MHz, Acetone-d_6) δ 164.6, 164.3, 153.7, 150.0, 148.9, 145.8, 139.3, 137.4, 135.5, 133.4, 129.8, 129.6, 129.4, 129.0, 128.1, 126.9, 125.7, 124.1, 123.1, 121.4; HRMS: Calcd. for C_{62}H_{53}IrN_4PF_6·3H_2O: 1159.2008. Found: 1159.2009. Anal. C, H, N: calcd. 51.54, 3.71, 6.24, found: 51.51, 3.66, 6.28. The elemental analysis result (Figure S2) was shown in the supporting information.

**Disk diffusion susceptibility test.** The disk diffusion method was used to determine sensitivity of bacteria to complexes 1–5.† S. aureus, E. faecalis, E. coli and K. pneumoniae direct colony suspensions were prepared in Luria Bertain broth for less than 18 h. The turbidity of four bacterial suspensions was standardized to match 1.0 × 10⁸ UFC/mL (or 0.5 McFarland turbidity units). A sterile swab dipped in fresh bacterial suspensions was inoculated to the entire surface of Mueller-Hinton agar and the inoculum was distributed evenly. The plates were incubated with standardizing inoculum suspension for 15 min at room temperature. 10 μL complexes 1–5 (11.5 μM) or ampicillin (28 μM) were impregnated on sterile 6 mm paper disks for 15 min. Subsequently, the paper disks were aseptically placed on and pressed down firmly to the inoculated plates. Plates were incubated for 24 h in a 37°C incubator. After the incubation, the inhibition zones were measured. Ampicillin was used as a positive control for bacterial inhibition.
The presence or absence of growth around the disks was a measure of the ability of complexes 1–5 to inhibit growth of bacterial. All experiments were done in triplicate.

**MIC and MBC testing.** Complex 1 was examined for its antimicrobial activity against *S. aureus*. The MIC was estimated by the broth dilution method in Luria Bertani broth using the standardized method\(^{39–40}\). Briefly, serial dilutions of complex 1 were prepared in distilled water with concentrations ranging from 1.8 to 57.6 μM. 1 mL of an *S. aureus* inoculum (1 × 10^8 UFC/mL) and 0.1 mL of serially diluted complex 1 were added to 2.9 mL of Luria Bertani broth. Controls without complex 1 were also prepared. After 24 h of incubation at 37 °C under agitation in hermetic tubes, the MIC was determined as the lowest concentration of complex 1 that inhibited visible bacterial growth. The experiments were performed in triplicate. The complete data are shown in Table S2 in the Supplementary Information.

To determine the MBC, 10 μL of bacterial inoculum treated with different concentrations of complex 1 was taken aseptically from tubes that had not presented visible turbidity and was plated onto Mueller-Hinton agar. The MBC was considered as the lowest concentration of complex 1 that allowed less than 0.1% of the original inoculum treated with complex 1 to grow on the surface of the Mueller-Hinton agar. The experiments were performed in triplicate. The complete data are shown in Table S2 in the Supplementary Information.

To determine the nature of antibacterial effect of complex 1, the MBC/MIC ratio was used; when the ratio was lower than 4, complex 1 was considered as a bactericidal complex 1 and when the ratio was higher than 4, it was considered as a bacteriostatic complex 1.

**Log P Determination.** This experiment was carried out according to a reported method and the partitions were detected by HPLC\(^{61}\). The partition coefficients of the Ir(III) complexes were calculated using the equation: 
$$\log P = \log(\text{[Ir]}_{\text{WSO}}/\text{[Ir]}_{\text{OSW}})$$

where [Ir]\(_{\text{WSO}}\) means the concentration of Ir(III) complex dissolved in octanol-saturated water (WSO), and [Ir]\(_{\text{OSW}}\) means the concentration of Ir(III) complex dissolved in water-saturated octanol (OSW).

**Cell cytotoxicity.** Complex 1 was dissolved in dimethylsulfoxide (DMSO) at a stock concentration of 10 mM and serially diluted with low FBS medium to the working concentrations. The evaluation of cells proliferation was conducted using MTT assay. In brief, the cells were seeded in 96-well plates at a density of 8 × 10^4 cells/mL in complete medium for overnight. Serial dilution of complex 1 in low FBS medium was then added to the wells. After 72 h, 20 μL MTT (5 mg/mL) was added to each well. The plates were then incubated for an additional 4 h at 37 °C while avoiding direct light. Subsequently, the medium in each well was removed and 100 μL DMSO was added to each well. The plates were shaken gently for 10 min at room temperature to completely dissolve the purple formazan in the mitochondria of living cells. The color intensity was measured at 570 nm using a SpectraMax M5 microplate reader (Molecular Devices).

**Conclusion**

In summary, we have identified a metal-based complex that selectively inhibited the growth of *S. aureus*. The iridium(III) or rhodium(III) complexes 1–5 were synthesized by the reaction of MCl\(_3\)·H\(_2\)O with the C\(^{-}\)N ligand to yield the dimeric compound [M\(_2\)(C\(^{-}\)N)\(_2\)]Cl\(_2\), which was treated with the corresponding N\(^{-}\)N ligand to give the targeted complexes. Complexes 2 and 5 are novel and have been characterized for the first time.

Complex rac-1 inhibited the growth of *S. aureus* with MIC and MBC values of 3.60 and 7.19 μM. As the MBC/MIC ratio of complex 1 is approximately 2, complex 1 is considered to be bactericidal. Moreover, complex 1 also exhibited cytotoxicity against a number of cancer cells, with particular potency against ovarian, cervical and melanoma cells but not hepatocarcinoma cells, suggesting that its cytotoxic activity was cell type-dependent. To our knowledge, this cyclometallated iridium(III) complex is the first example of a substitutionally-inert, Group 9 organometallic compound utilized as a direct inhibitor of *S. aureus*.

We anticipate that cyclometallated iridium(III) complex 1 may serve as a useful scaffold for the further development of highly potent inhibitors of *S. aureus* as potential anti-bacterial agents and anti-cancer agents.

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Author Contributions
L.L., L.J.L. and W.C.C. contributed equally to the work. L.L. and M.W. performed the synthesis of metal complexes, photophysical measurement and Log P determination. L.J.L. and H.J.Z. contributed to the cell cytotoxicity experiment. W.C.C. conducted the antibacterial experiments. R.N.L., D.L.M. and C.H.L. designed the experiments and supervised the whole research. All the authors contributed to the results analysis and discussion.

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Supplementary information
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